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Supplementary appendix 1

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Supplement to: GBD 2017 Risk Factor Collaborators. Global, regional, and national comparative risk assessment of 84 behavioural, environmental and occupational, and metabolic risks or clusters of risks for 195 countries and territories, 1990–2017: a systematic analysis for the Global Burden of Disease Study 2017. *Lancet* 2018; **392:** 1923–45.

Methods appendix to Global, regional, and national comparative risk assessment of 84 behavioural, environmental and occupational, and metabolic risks or clusters of risks, 1990-2017: a systematic analysis for the Global Burden of Disease Study 2017

This appendix provides further methodological detail, supplemental figures, and more detailed results for risk factors. The appendix is organised into broad sections following the structure of the main paper.

Preamble

This appendix provides further methodological detail and more detailed results for "Global, regional, and national comparative risk assessment of 84 behavioural, environmental and occupational, and metabolic risks or clusters of risks, 1990-2017: a systematic analysis for the Global Burden of Disease Study 2017." This study complies with the Guidelines for Accurate and Transparent Health Estimates Reporting (GATHER) recommendations. It includes detailed tables and information on data in an effort to maximise transparency in our estimation processes and provide a comprehensive description of analytical steps. We intend this appendix to be a living document, to be updated with each iteration of the Global Burden of Disease Study.

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Authors' Contributions

Managing the estimation process

Ashkan Afshin, Tahiya Alam, Brigette Blacker, Michael Brauer, Aaron Cohen, Elizabeth Cromwell, Lalit Dandona, Rakhi Dandona, Louisa Degenhardt, Samath Dharmaratne, Charbel El Bcheraoui, Kara Estep, Valery Feigin, Nancy Fullman, Emmanuela Gakidou, Caitlin Hawley, Simon Hay, Spencer L James, Nicholas J Kassebaum, Ibrahim Khalil, Hmwe Kyu, Stephen Lim, Alan Lopez, Rafael Lozano, Ashley Marks, Awoke Misganaw, Ali Mokdad, Christopher Murray, Mohsen Naghavi, Elaine Nsoesie, Helen Olsen, Robert Reiner, Gregory Roth, David Smith, Mari Smith, Jeffrey Stanaway, Stein Emil Vollset, Theo Vos, and Harvey Whiteford

Writing the first draft of the manuscript

Jason Anderson, Kate Causey, William Gardner, Sam Larson, Ashley Marks, Molly Nixon, Jeff Stanaway, Anna Torre.

Providing data or critical feedback on data sources

Aliasghar A Kiadaliri, Kalkidan Hassen Abate, Cristiana Abbafati, Nooshin Abbasi, Ibrahim Abdollahpour, Molla Abebe, Zegeve Abebe, Semaw Ferede Abera, Laith Abu-Raddad, Niveen Abu-Rmeileh, Manfred Accrombessi, Oladimeji Adebayo, Zanfina Ademi, Olatunji Adetokunboh, Mina Adib, José C. Adusar, Mohsen Afarideh, Ashkan Afshin, Sargis Aghayan, Alireza Ahmadi, Mehdi Ahmadi, Muktar Ahmed, Ibtihel Aichour, Mohammad Esmail Akbari, Tomi Akinyemiju, Nadia Akseer, Fares Alahdab, Khurshid Alam, Kefyalew Addis Alene, Syed Aljunid, François Alla, Ubai Alsharif, Dayane Gabriele Alves Silveira, Nelson Alvis-Guzman, Mamusha Aman, Azmeraw T. Amare, Walid Ammar, Catalina Liliana Andrei, Mr. Mustafa Geleto Ansha, Carl Abelardo Antonio, Seth Christopher Y. Appiah, Olatunde Aremu, Johan Ärnlöv, Al Artaman, Krishna Kumar Aryal, Hamid Asayesh, Marcel Ausloos, Ashish Awasthi, Beatriz Paulina Ayala Quintanilla, Rakesh Ayer, Tambe Betrand Ayuk, Peter Azzopardi, Nam Ba Nguyen, Arefeh Babazadeh, Alaa Badawi, Kalpana Balakrishnan, Shoshana Ballew, Maciej Banach, Joseph Banoub, Aleksandra Barac, Till Bärnighausen, Lope Barrero, Sanjay Basu, Bernhard Baune, Neeraj Bedi, Masoud Behzadifar, Meysam Behzadifar, Bayu Begashaw Bekele, Yihalem Abebe Belay, Aminu Bello, Isabela Bensenor, Eduardo Bernabe, Robert Bernstein, Mircea Beuran, Tina Beyranvand, Neeraj Bhala, Suraj Bhattarai, Boris Bikbov, Nigus Bililign, Muhammad Shahdaat Bin Sayeed, Ibrahim Bou-Orm, Rupert Bourne, Michael Brauer, Andrey Briko, Gabrielle Britton, Leah Cahill, Giulia Carreras, Juan J Carrero, Deborah Carvalho Malta, Carlos Castañeda-Orjuela, Jacqueline Castillo Rivas, Franz Castro, Ferrán Catalá-López, Yazan Chaiah, Hsing-Yi Chang, Jung-Chen Chang, Fiona Charlson, Aparajita Chattopadhyay, Miao Li Chee, Ching-Yu Cheng, Peggy Pei-Chia Chiang, Abdulaal Chitheer, Rajiv Chowdhury, Devasahayam Christopher, Sheng-Chia Chung, Massimo Cirillo, Aaron Cohen, Daniel Collado-Mateo, Cyrus Cooper, Josef Coresh, Paolo Angelo Cortesi, Monica Cortinovis, Michael Criqui, Alemneh Kabeta Daba, Albertino Damasceno, Lalit Dandona, Rakhi Dandona, Sarah Darby, Paul Dargan, Ahmad Daryani, José Das Neves, Fernando De La Hoz, Kebede Deribe, Nikolaos Dervenis, Aniruddha Deshpande, Getenet Dessie, Gabrielle Deveber, Samath Dharmaratne, Meghnath Dhimal, Eric L. Ding, Klara Dokova, David Teye Doku, Kerrie Doyle, Manisha Dubey, Eyasu Ejeta Duken, Bruce Duncan, Andre Duraes, Natalie Ebert, Hedyeh Ebrahimi, Soheil Ebrahimpour, Andem Effiong, Anne Elise Eggen, Ziad El-Khatib, Iqbal Elyazar, Benjamin Er, Holly Erskine, Sharareh Eskandarieh, Alireza Esteghamati, Sadaf Esteghamati, Mohammad Fareed, Carla Farinha, Andre Faro, Mohammad Hosen Farzaei, Valery Feigin, Andrea B. Feigl, Seved-Mohammad Fereshtehnejad, Joao Fernandes, Irina Filip, Jonas D. Finger, Nataliya Foigt,

Takeshi Fukumoto, Thomas Fürst, Neal Futran, Seana Gall, Amiran Gamkrelidze, Tigit Gashaw, Abadi Kahsu Gebre, Amanuel Tesfay Gebremedhin, Merhawi Gebremedhin, Teklu Gebremichael, Tilayie Feto Gelano, Johanna Geleijnse, Ayele Geleto, Mamata Ghimire, Simona Giampaoli, Ibrahim Ginawi, Srinivas Goli, Hector Gomez-Dantes, Philimon Gona, Sameer Gopalani, Ayman Grada, Morgan Grams, Giuseppe Grosso, Yuming Guo, Rahul Gupta, Rajat Das Gupta, Rajeev Gupta, Tanush Gupta, Daniela Sarahí Gutiérrez-Torres, Juanita Haagsma, Nima Hafezi-Nejad, Tekleberhan Beyene Hagos, Gessessew Bugssa Hailu, Arya Haj-Mirzaian, Arvin Ha-Mirzaian, Yuantao Hao, Hilda Harb, Sivadasanpillai Harikrishnan, Josep Maria Haro, Hadi Hassankhani, Hamid Y. Hassen, Rasmus Havmoeller, Behnam Heidari, Mohsen Heidari, Delia Hendrie, Andualem Henok, Ileana B. Heredia-Pi, Claudiu Herteliu, Hans Hoek, Howard J. Hoffman, Michael Hole, Praveen Hoogar, H Dean Hosgood, Mehdi Hosseinzadeh, Mihaela Hostiuc, Sorin Hostiuc, Damian Hoy, Mohamed Hsairi, Guoging Hu, Kim Moesgaard Iburg, Nayu Ikeda, Usman Igbal, Farhad Islami, Maria Jackson, Kathryn H. Jacobsen, Mihajlo Jakovljevic, Spencer James, Simerjot Jassal, Achala Jayatilleke, Panniyammakal Jeemon, Vivekanand Jha, Jost B. Jonas, Zahra Jorjoran Shushtari, Jacek Jozwiak, Mikk Jürisson, Zubair Kabir, Amaha Kahsay, Amaha Kahsay, Tanuj Kanchan, André Karch, Corine Karema, Seved M. Karimi, Amir Kasaeian, Getachew Mullu Kassa, Nicholas Kassebaum, Srinivasa Vittal Katikireddi, Anil Kaul, Norito Kawakami, Peter Keiyoro, Andre Keren, Chandrasekharan Nair Kesavachandran, Yousef Khader, Morteza Abdullatif Khafaie, Ibrahim Khalil, Muhammad Shahzeb Khan, Young-Ho Khang, Mona Khater, Habibolah Khazaie, Abdullah T. Khoja, Ardeshir Khosravi, Mohammad Hossein Khosravi, Daniel Kiirithio, Cho-Il Kim, Daniel Kim, Young-Eun Kim, Yun Jin Kim, Adnan Kisa, Luke Knibbs, Ann Kristin Knudsen, Soewarta Kosen, Parvaiz Koul, Ai Koyanagi, Michael Kravchenko, Kewal Krishan, Hans Kromhout, Barthelemy Kuate Defo, Burcu Kucuk Bicer, G Anil Kumar, Manasi Kumar, Ihor Kuzin, Carl Lachat, Deepesh P. Lad, Sheetal Lad, Alessandra Lafranconi, Huong Lan Nguyen, Justin Lang, Van Lansingh, Samantha Larson, Misgan Legesse, Yirga Legesse, James Leigh, Janni Leung, Shanshan Li, Yichong Li, Juan Liang, Xiaofeng Liang, Yu Liao, Lee-Ling Lim, Shiwei Liu, Alan Lopez, Stefan Lorkowski, Paulo Lotufo, Stefan Ma, Erlyn Rachelle Macarayan, Isis Machado, P A Mahesh, Marek Majdan, Reza Majdzadeh, Azeem Majeed, Reza Malekzadeh, Mohammad Ali Mansournia, Lorenzo Mantovani, Joemer Maravilla, Wagner Marcenes, Randall V. Martin, Francisco Rogerlândio Martins-Melo, Winfried März, Melvin Marzan, Benjamin Massenburg, Manu Mathur, Kunihiro Matsushita, Mohsen Mazidi, Colm Mcalinden, John Mcgrath, Abla Mehio Sibai, Varshil Mehta, Toni Meier, Yohannes Adama Melaku, Addisu Melese, Mulugeta Melku, Peter Memiah, Ziad Memish, Walter Mendoza, Gert B.M. Mensink, Atte Meretoja, Tuomo Meretoja, Tomislav Mestrovic, Haftay Berhane Mezgebe, Tomasz Miazgowski, Ted R Miller, Erkin Mirrakhimov, Babak Moazen, Bahram Mohajer, Karzan Mohammad, Moslem Mohammadi, Noushin Mohammadifard, Shafiu Mohammed, Farnam Mohebi, Ali Mokdad, Lorenzo Monasta, Yoshan Moodley, Ghobad Moradi, Maziar Moradi-Lakeh, Mehdi Moradinazar, Lidia Morawska, Joana Morgado-Da-Costa, Marilita Moschos, Seyyed Meysam Mousavi, Dariush Mozaffarian, Achenef Muche, Ulrich Mueller, Kamarul Imran Musa, Ghulam Mustafa, Ashraf Nabhan, Mohsen Naghavi, Seyed Sina Naghibi Irvani, Azin Nahvijou, Bruno Nascimento, Ionut Negoi, Ruxandra Irina Negoi, Charels Newton, Josephine Ngunjiri, Anh Nguyen, Ha Nguyen, Huong Nguyen, Nobuo Nishi, Marzieh Nojomi, Shuhei Nomura, Mehdi Noroozi, Bo Norrving, Jean Jacques Noubiap, Hamid Reza Nouri, Malihe Nourollahpour, Mohammad Reza Nowroozi, Christopher M Odell, Felix Ogbo, In-Hwan Oh, Olanrewaju Oladimeji, Andrew T. Olagunju, Tinuke O. Olagunju, Bolajoko Olusanya, Jacob Olusanya, Kanyin Ong, Sok King Ong, Eyal Oren, Heather Orpana, Alberto Ortiz, Stanislav Otstavnov, Simon Øverland, Mayowa Owolabi, Rosana Pacella, Adrian Pana, Songhomitra Panda-Jonas, Hadi Parsian, Shanti Patel, Sanghamitra Pati, Snehal Patil, Ajay Patle, Deepak Paudel, Wayra Citlali Paz Ballesteros, Neil Pearce,

Alexandre Pereira, David Pereira, Norberto Perico, Max Petzold, Huyen Phuc Do, Meghdad Pirsaheb, Farhad Pishgar, Dietrich Plass, Suzanne Polinder, Kevan Polkinghorne, Hossein Poustchi, Swayam Prakash, Narayan Prasad, Hai Quang Pham, Amir Radfar, Anwar Rafay, Alireza Rafiei, Fakher Rahim, Zohreh Rahimi, Afarin Rahimi-Movaghar, Vafa Rahimi-Movaghar, Mahfuzar Rahman, Mohammad Hifz Ur Rahman, Rajesh Kumar Rai, Usha Ram, Chhabi Lal Ranabhat, Prabhat Ranjan, Goura K Rath, David Laith Rawaf, Salman Rawaf, Kolli Srinath Reddy, Colin D Rehm, Jürgen Rehm, Giuseppe Remuzzi, Andre Renzaho, Luz Myriam Reynales-Shigematsu, Antonio Luiz Ribeiro, Juan A Rivera, Sonia Rodríguez-Ramírez, Leonardo Roever, Luca Ronfani, Gholamreza Roshandel, Ali Rostami, Ambuj Roy, Enrico Rubagotti, Lesley Rushton, Charumathi Sabanayagam, Basema Saddik, Ehsan Sadeghi, Hosein Safari, Yahya Safari, Saeid Safiri, Mohammad Ali Sahraian, Nasir Salam, Payman Salamati, Yahya Salimi, Hamideh Salimzadeh, Devashri Salvi, Abdallah M. Samy, Juan Sanabria, Tania G Sánchez-Pimienta, Yingying Sang, Milena Santric Milicevic, Bruno Sao Jose, Mayank Sardana, Rodrigo Sarmiento-Suárez, Nizal Sarrafzadegan, Shahabeddin Sarvi, Thirunavukkarasu Sathish, Maheswar Satpathy, Monika Sawhney, Mete Saylan, Mehdi Sayyah, Elke Schaeffner, Maria Inês Schmidt, David C Schwebel, Falk Schwendicke, James Scott, Mario Šekerija, Sadaf Sepanlou, Edson Serván-Mori, Sevedmojtaba Seyedmousavi, Gavin Shaddick, Amira Shaheen, Masood Ali Shaikh, Teresa Shamah-Levy, Mehran Shams-Beyranvand, Kiomars Sharafi, Mehdi Sharif, Hamid Sharifi, Sheikh Mohammed Shariful Islam, Jayendra Sharma, Meenakshi Sharma, Aziz Sheikh, Peilin Shi, Min-Jeong Shin, Ivy Shiue, Farhad Shokraneh, Haitham Shoman, Si Si, Soraya Siabani, Tariq Jamal Siddiqi, Diego Augusto Santos Silva, Jasvinder Singh, Virendra Singh, Dhirendra Narain Sinha, Eirini Skiadaresi, Vegard Skirbekk, Soheila Sobhani, Reed Sorensen, Joan B Soriano, Angela Spinelli, Luciano Sposato, Chandrashekhar T Sreeramareddy, Goran Stevanovic, Leo Stockfelt, Mark Stokes, Lela Sturua, Agus Sudaryanto, Muawiyyah Babale Sufiyan, Rizwan Suliankatchi Abdulkader, Patrick Sur, Bryan L. Sykes, Cassandra Szoeke, Rafael Tabarés-Seisdedos, Mesfin Tadese, Eyasu Tamru , Nuno Taveira, Gebre Teklemariam Demoz, Awoke Temesgen, Mohamad-Hani Temsah, Omar Temsah, Belay Tessema, Mebrahtu Teweldemedhin, Hue Thi Mai, Nu Thi Truong, Matthew L Thomas, Nihal Thomas, Giang Thu Vu, Myriam Tobollik, Marcello Tonelli, Miguel Tortajada-Girbés, Mathilde Touvier, Marcos Roberto Tovani-Palone, Bach Tran, Khanh Bao Tran, Thomas Truelsen, Stefanos Tyrovolas, Kingsley Nnanna Ukwaja, Irfan Ullah, Muhammad Sharig Usman, Olalekan A Uthman, Muthiah Vaduganathan, Afsane Vaezi, Pascual Valdez, Aaron Van Donkelaar, Tommi Vasankari, Vidhya Venkateswaran, Narayanaswamy Venketasubramanian, Santos Villafaina, Sergey Vladimirov, Vasiliy Vlassov, Stein Emil Vollset, Fasil Wagnew, Shishay Wahdey, Yasir Waheed, Yanping Wang, Yuan-Pang Wang, Elisabete Weiderpass, Robert Weintraub, Inbal Weiss Salz, Andrea Werdecker, Ronny Westerman, Harvey Whiteford, Justyna Widecka, Tissa Wijeratne, Charles Shey Wiysonge, Charles Wolfe, Tien Yin Wong, Shouling Wu, Denis Xavier, Gelin Xu, Ali Yadollahpour, Hossein Yahyazadeh, Lijing Yan, Mehdi Yaseri, Yasin Jemal Yasin, Alex Yeshaneh, Ebrahim M. Yimer, Naohiro Yonemoto, Seok-Jun Yoon, Marcel Yotebieng, Mustafa Younis, Chuanhua Yu, Zoubida Zaidi, Sojib Bin Zaman, Mohammad Zamani, Luis Zavala-Arciniega, Anthony Lin Zhang, and Sanjay Zodpey.

Developing methods or computational machinery

Cristiana Abbafati, Ibrahim Abdollahpour, Ashkan Afshin, Mohammad Esmail Akbari, Mehran Alijanzadeh, Marcel Ausloos, Doanl Bisanzio, Michael Brauer, Kate Causey, Kelly Cercy, Aaron Cohen, Josef Coresh, Ahmad Daryani, Holly Erskine, Kairsten Fay, Kyle Foreman, Morgan Grams, Hamid Y. Hassen, Claudiu Herteliu, Howard Hu, Chantal Huynh, Spencer James, Nicholas Kassebaum, Young-Eun Kim, Adnan Kisa, Van Lansingh, Misgan Legesse, James Leigh, Tim Lucas, Helena Manguerra, Randall V. Martin, Kunihiro Matsushita, Fantahun Mekonnen, Anoushka Millear, Dariush Mozaffarian, Mohsen Naghavi, Kanyin Ong, Charles Parry, Alireza Rafiei, Chhabi Lal Ranabhat, Jürgen Rehm, Robert Reiner, Marissa Reitsma, Gregory Roth, Lesley Rushton, Abdallah M. Samy, Yingying Sang, Damian Santomauro, Maheswar Satpathy, David C Schwebel, James Scott, Gavin Shaddick, Mehdi Sharif, David Smith, Reed Sorensen, Vinay Srinivasan, Jeff Stanaway, Patrick Sur, Hannah J Thomas, Matthew L Thomas, Hayley Tymeson, Rachel L. Updike, Ronny Westerman, Simon Yadgir, Lijing Yan, and Naohiro Yonemoto.

Applying analytical methods to produce estimates

Ashkan Afshin, Syed Mustafa Ali, Azmeraw T. Amare, Olatunde Aremu, Marcel Ausloos, Bayu Begashaw Bekele, Doanl Bisanzio, Michael Brauer, Kate Causey, Kelly Cercy, Fiona Charlson, Devasahayam Christopher, Aaron Cohen, Ahmad Daryani, Getenet Dessie, Tim Driscoll, Andem Effiong, Aman Endries, Holly Erskine, Kairsten Fay, Alize Ferrari, William Gardner, Segen Gebremeskel, Rakesh Ghosh, Hamid Y. Hassen, Claudiu Herteliu, Caleb Irvine, Manoochehr Karami, Nicholas Kassebaum, Behzad Khafaie , Yun Jin Kim, Adnan Kisa, Samantha Larson, Misgan Legesse, James Leigh, Tim Lucas, Helena Manguerra, Randall V. Martin, Fantahun Mekonnen, Anoushka Millear, Shafiu Mohammed, Ali Mokdad, Mohsen Naghavi, Grant Nguyen, Emma Nichols, Bolajoko Olusanya, Jacob Olusanya, Kanyin Ong, Katherine Paulson , Swayam Prakash, Caroline Purcell, Chhabi Lal Ranabhat, Marissa Reitsma, Gregory Roth, Enrico Rubagotti, Devashri Salvi, Abdallah M. Samy, Yingying Sang, Damian Santomauro, Shahabeddin Sarvi, Maheswar Satpathy, Seyedmojtaba Seyedmousavi, Gavin Shaddick, Mehdi Sharif, Reed Sorensen, Chandrashekhar T Sreeramareddy, Vinay Srinivasan, Jeff Stanaway, Patrick Sur, Bryan L. Sykes, Andrew Theis, Hannah J Thomas, Matthew L Thomas, Rachel L. Updike, Vidhya Venkateswaran, Inbal Weiss Salz, Ronny Westerman, Tissa Wijeratne, and Simon Yadgi.

Providing critical feedback on methods or results

Aliasghar A Kiadaliri, Degu Abate, Kalkidan Hassen Abate, Cristiana Abbafati, Nooshin Abbasi, Hedayat Abbastabar, Jemal Abdela, Ibrahim Abdollahpour, Molla Abebe, Zegeye Abebe, Semaw Ferede Abera, Haftom Abraha, Niveen Abu-Rmeileh, Pawan Acharya, Abdu Adamu, Akilew Awoke Adane, Oladimeji Adebayo, Victor Adekanmbi, Olatunji Adetokunboh, Mina Adib, José C. Adusar, Kossivi Afanvi, Mohsen Afarideh, Ashkan Afshin, Gina Agarwal, Anju Aggarwal, Sargis Aghayan, Anurag Agrawal, Alireza Ahmadi, Mehdi Ahmadi, Hamid Ahmadieh, Muktar Ahmed, Amani Nidhal Aichour, Ibtihel Aichour, Miloud Taki Eddine Aichour, Mohammad Esmail Akbari, Tomi Akinyemiju, Nadia Akseer, Fares Alahdab, Ziyad Al-Aly, Khurshid Alam, Tahiya Alam, Seyed Moayed Alavian, Kefyalew Addis Alene, Ayman Al-Eyadhy, Syed Mustafa Ali, Mehran Alijanzadeh, Reza Alizadeh-Navaei, Syed Aljunid, Ala'a Alkerwi, Hesham Al-Mekhlafi, Ubai Alsharif, Khalid Altirkawi, Dayane Gabriele Alves Silveira, Nelson Alvis-Guzman, Mamusha Aman, Azmeraw T. Amare, Walid Ammar, Mirica Andreea, Catalina Liliana Andrei, Sofia Androudi, Mina Anjomshoa, Mr. Mustafa Geleto Ansha, Josep M Antó, Carl Abelardo Antonio, Palwasha Anwari, Lambert Appiah, Olatunde Aremu, Johan Ärnlöv, Al Artaman, Krishna Kumar Aryal, Hamid Asayesh, Zerihun Ataro, Marcel Ausloos, Euripide Avokpaho, Ashish Awasthi, Beatriz Paulina Ayala Quintanilla, Tambe Betrand Ayuk, Peter Azzopardi, Nam Ba Nguyen, Arefeh Babazadeh, Hamid Badali, Alaa Badawi, Kylie Ball, Shoshana Ballew, Joseph Banoub, Aleksandra Barac, Suzanne Barker-Collo, Till Bärnighausen, Lope Barrero, Sanjay Basu, Bernhard Baune, Shahrzad Bazargan-Hejazi, Neeraj Bedi, Ettore Beghi, Masoud Behzadifar, Meysam Behzadifar, Yannick Béjot, Bayu Begashaw Bekele, Ezrt Belay, Yihalem Abebe Belay, Michelle Bell, Aminu Bello, Derrick Bennett, Isabela Bensenor, Gilles Bergeron, Adugnaw Berhane, Eduardo Bernabe, Robert Bernstein, Mircea Beuran, Tina Beyranvand, Neeraj Bhala, Ashish Bhalla, Suraj Bhattarai, Belete Biadgo, Ali Bijani, Boris Bikbov, Ver Bilano, Nigus Bililign, Muhammad

Shahdaat Bin Sayeed, Doanl Bisanzio, Tuhin Biswas, Tone Bjørge, Archie Bleyer, Rohan Borschmann, Soufiane Boufous, Rupert Bourne, Oliver Brady, Michael Brauer, Alexandra Brazinova, Nicholas Breitborde, Hermann Brenner, Andrey Briko, Gabrielle Britton, Rachelle Buchbinder, Reinhard Busse, Zahid Butt, Leah Cahill, Lucero Cahuana-Hurtado, Ismael Campos-Nonato, Rosario Cárdenas, Juan J Carrero, Felix Carvalho, Deborah Carvalho Malta, Carlos Castañeda-Orjuela, Jacqueline Castillo Rivas, Franz Castro, Ferrán Catalá-López, Ester Cerin, Yazan Chaiah, Jung-Chen Chang, Fiona Charlson, Aparajita Chattopadhyay, Vijay Kumar Chattu, Peggy Pei-Chia Chiang, Odgerel Chimed-Ochir, Ken Chin, Yilma Chisha, Jee-Young Choi, Hanne Christensen, Devasahayam Christopher, Sheng-Chia Chung, Flavia Cicuttini, Massimo Cirillo, Aaron Cohen, Daniel Collado-Mateo, Cyrus Cooper, Josef Coresh, Paolo Angelo Cortesi, Monica Cortinovis, Megan Costa, Ewerton Cousin, Michael Criqui, David Cundiff, Alemneh Kabeta Daba, Berihun Dachew, Abel Dadi, Lalit Dandona, Rakhi Dandona, Sarah Darby, Paul Dargan, Ahmad Daryani, José Das Neves, Tamirat Tesfaye Dasa, Kairat Davletov, Vanessa De La Cruz-Góngora, Fernando De La Hoz, Diego De Leo, Jan-Walter De Neve, Megbaru Debalkie, Robert Dellavalle, Edgar Denova-Gutiérrez, Kebede Deribe, Nikolaos Dervenis, Don Des Jarlais, Aniruddha Deshpande, Getenet Dessie, Subhojit Dey, Samath Dharmaratne, Meghnath Dhimal, Eric L. Ding, Helen Diro, Shirin Dialalinia, Klara Dokova, David Teye Doku, Kerrie Doyle, Tim Driscoll, Manisha Dubey, Eleonora Dubljanin, Eyasu Ejeta Duken, Bruce Duncan, Andre Duraes, Natalie Ebert, Hedyeh Ebrahimi, Soheil Ebrahimpour, David Edvardsson, Andem Effiong, Anne Elise Eggen, Ziad El-Khatib, Igbal Elyazar, Ahmadali Enayati, Aman Endries, Benjamin Er, Holly Erskine, Sharareh Eskandarieh, Alireza Esteghamati, Sadaf Esteghamati, Hamed Fakhim, Mahbobeh Faramarzi, Mohammad Fareed, Talha Farid, Carla Farinha, Andrea Farioli, Andre Faro, Mohammad Hosen Farzaei, Batool Fatima, Valery Feigin, Andrea B. Feigl, Seyed-Mohammad Fereshtehnejad, Eduarda Fernandes, Joao Fernandes, Manuela Ferreira, Irina Filip, Jonas D. Finger, Florian Fischer, Nataliya Foigt, Takeshi Fukumoto, Nancy Fullman, Thomas Fürst, João M. Furtado, Neal Futran, Silvano Gallus, Amiran Gamkrelidze, Morsaleh Ganji, Alberto L. García-Basteiro, Tigit Gashaw, Abadi Kahsu Gebre, Amanuel Tesfay Gebremedhin, Merhawi Gebremedhin, Afewerki Gebremeskel, Teklu Gebremichael, Tilayie Feto Gelano, Johanna Geleijnse, Ayele Geleto, Kebede Embaye Gezae, Reza Ghadimi, Khalil Ghasemi Falavarjani, Maryam Ghasemi-Kasman, Mamata Ghimire, Rakesh Ghosh, Paramjit Gill, Tiffany Gill, Richard Gillum, Giorgia Giussani, Elena Gnedovskaya, Srinivas Goli, Philimon Gona, Aloke Gopal, Sameer Gopalani, Ayman Grada, Morgan Grams, Giuseppe Grosso, Harish Gugnani, Yuming Guo, Rahul Gupta, Rajat Das Gupta, Rajeev Gupta, Tanush Gupta, Reyna Alma Gutiérrez, Juanita Haagsma, Tesfa Dejenie Habtewold, Vladimir Hachinski, Nima Hafezi-Nejad, Tekleberhan Beyene Hagos, Dessalegn Haile, Gessessew Bugssa Hailu, Arya Haj-Mirzaian, Randah Hamadeh, Samer Hamidi, Arvin Ha-Mirzaian, Alexis Jeannine Handal, Graeme Hankey, Yuantao Hao, Hilda Harb, Sivadasanpillai Harikrishnan, Josep Maria Haro, Hadi Hassankhani, Hamid Y. Hassen, Rasmus Havmoeller, Caitlin Hawley, Akbar Hedayatizadeh-Omran, Behzad Heibati, Behnam Heidari, Mohsen Heidari, Delia Hendrie, Ileana B. Heredia-Pi, Claudiu Herteliu, Fatemeh Heydarpour, Sousan Heydarpour, Desalegn Hibstu, Tarig Higazi, Esayas Haregot Hilawe, Michael Hole, Enayatollah Homaie Rad, Praveen Hoogar, H Dean Hosgood, Mostafa Hosseini, Mihaela Hostiuc, Sorin Hostiuc, Damian Hoy, Guoqing Hu, John Huang, Nayu Ikeda, Olayinka Ilesanmi, Usman Igbal, Farhad Islami, Kathryn H. Jacobsen, Nader Jahanmehr, Sudhir Kumar Jain, Mihajlo Jakovljevic, Spencer James, Achala Jayatilleke, Panniyammakal Jeemon, Ravi Prakash Jha, Vivekanand Jha, Jost B. Jonas, Jitendra Jonnagaddala, Zahra Jorjoran Shushtari, Ankur Joshi, Jacek Jozwiak, Mikk Jürisson, Amaha Kahsay, Amaha Kahsay, Rizwan Kalani, Tanuj Kanchan, Chittaranjan Kar, Manoochehr Karami, Behazad Karami Matin, André Karch, Seyed M. Karimi, Amir Kasaeian, Getachew Mullu Kassa, Tesfaye Kassa, Nicholas Kassebaum, Srinivasa Vittal Katikireddi, Zhila Kazemi, Ali Kazemi

Karyani, Adane Kefale, Peter Keiyoro, Andre P Kengne, Chandrasekharan Nair Kesavachandran, Yousef Khader, Morteza Abdullatif Khafaie, Nauman Khalid, Ibrahim Khalil, Gulfaraz Khan, Muhammad Ali Khan, Muhammad Shahzeb Khan, Young-Ho Khang, Mona Khater, Abdullah T. Khoja, Ardeshir Khosravi, Mohammad Hossein Khosravi, Daniel Kiirithio, Daniel Kim, Yun Jin Kim, Ruth Kimokoti, Adnan Kisa, Katarzyna Kissimova-Skarbek, Mika Kivimaki, Luke Knibbs, Ann Kristin Knudsen, Sonali Kochhar, Yoshihiro Kokubo, Tufa Kolola, Jacek Kopec, Parvaiz Koul, Ai Koyanagi, Kewal Krishan, Barthelemy Kuate Defo, G Anil Kumar, Manasi Kumar, Hmwe Kyu, Deepesh P. Lad, Sheetal Lad, Alessandra Lafranconi, Ratilal Lalloo, Tea Lallukka, Faris Lami, Huong Lan Nguyen, Justin Lang, Van Lansingh, Arman Latifi, Jeffrey Lazarus, Paul Lee, Misgan Legesse, Yirga Legesse, James Leigh, Cheru T Leshargie, Janni Leung, Miriam Levi, Sonia Lewycka, Shanshan Li, Yu Liao, Lee-Ling Lim, Shai Linn, Shiwei Liu, Rakesh Lodha, Alan Lopez, Stefan Lorkowski, Paulo Lotufo, Raimundas Lunevicius, Kala M. Mehta, Stefan Ma, Erlyn Rachelle Macarayan, Isis Machado, F Madotto, P A Mahesh, Marek Majdan, Reza Majdzadeh, Azeem Majeed, Reza Malekzadeh, Abdullah Mamun, Mohammad Ali Mansournia, Lorenzo Mantovani, Joemer Maravilla, Wagner Marcenes, Sheila Martins, Francisco Rogerlândio Martins-Melo, Melvin Marzan, Benjamin Massenburg, Manu Mathur, Prashant Mathur, Kunihiro Matsushita, Pallab K Maulik, Mohsen Mazidi, Colm Mcalinden, John Mcgrath, Martin Mckee, Ravi Mehrotra, Varshil Mehta, Toni Meier, Fantahun Mekonnen, Yohannes Adama Melaku, Addisu Melese, Mulugeta Melku, Peter Memiah, Ziad Memish, Walter Mendoza, Desalegn Tadese Mengistu, George Mensah, Gert B.M. Mensink, Seid Tiku Mereta, Atte Meretoja, Tuomo Meretoja, Tomislav Mestrovic, Haftay Berhane Mezgebe, Bartosz Miazgowski, Tomasz Miazgowski, Ted R Miller, Molly Miller-Petrie, Gk Mini, Mojde Mirarefin, Erkin Mirrakhimov, Habtamu Mitiku, Babak Moazen, Moslem Mohammadi, Shafiu Mohammed, Farnam Mohebi, Ali Mokdad, Mariam Molokhia, Fatemeh Momeniha, Lorenzo Monasta, Ghobad Moradi, Maziar Moradi-Lakeh, Mehdi Moradinazar, Paula Moraga, Lidia Morawska, Joana Morgado-Da-Costa, Shane Morrison, Marilita Moschos, Seyyed Meysam Mousavi, Dariush Mozaffarian, Kalayu Brhane Mruts, Achenef Muche, Kindie Fentahun Muchie, Ulrich Mueller, Satinath Mukhopadhyay, Kamarul Imran Musa, Ghulam Mustafa, Ashraf Nabhan, Mohsen Naghavi, Seyed Sina Naghibi Irvani, Aliya Naheed, Azin Nahvijou, Nitish Naik, Farid Najafi, Vinay Nangia, Jobert Richie Nansseu, Bruno Nascimento, Ionut Negoi, Ruxandra Irina Negoi, Subas Neupane, Charels Newton, Josephine Ngunjiri, Emma Nichols, Jing Nie, Nobuo Nishi, Ole F Norheim, Mehdi Noroozi, Bo Norrving, Jean Jacques Noubiap, Hamid Reza Nouri, Malihe Nourollahpour, Mohammad Reza Nowroozi, Dina Nur Anggraini Ningrum, Peter Nyasulu, Richard Ofori-Asenso, Felix Ogbo, In-Hwan Oh, Olanrewaju Oladimeji, Andrew T. Olagunju, Tinuke O. Olagunju, Pedro Olivares, Helen Elizabeth Olsen, Bolajoko Olusanya, Jacob Olusanya, Kanyin Ong, Eyal Oren, Heather Orpana, Alberto Ortiz, Erika Ota, Stanislav Otstavnov, Simon Øverland, Mayowa Owolabi, Rosana Pacella, Abhijit Pakhare, Amir Pakpour, Adrian Pana, Songhomitra Panda-Jonas, Eun-Kee Park, Hadi Parsian, Shanti Patel, Sanghamitra Pati, Snehal Patil, George Patton, Deepak Paudel, Katherine Paulson, Neil Pearce, Alexandre Pereira, David Pereira, Norberto Perico, Huyen Phuc Do, Julian Pillay, Michael Piradov, Meghdad Pirsaheb, Tobias Pischon, Farhad Pishgar, Oleguer Plana-Ripoll, Dietrich Plass, Suzanne Polinder, Kevan Polkinghorne, Maarten Postma, Richie Poulton, Akram Pourshams, Hossein Poustchi, Dorairaj Prabhakaran, Swayam Prakash, Manorama Purwar, Mostafa Qorbani, Hai Quang Pham, Amir Radfar, Anwar Rafay, Alireza Rafiei, Fakher Rahim, Afarin Rahimi-Movaghar, Vafa Rahimi-Movaghar, Mahfuzar Rahman, Muhammad Aziz Rahman, Rajesh Kumar Rai, Fatemeh Rajati, Sasa Rajsic, Usha Ram, Chhabi Lal Ranabhat, Prabhat Ranjan, Goura K Rath, David Laith Rawaf, Salman Rawaf, Kolli Srinath Reddy, Colin D Rehm, Marissa Reitsma, Giuseppe Remuzzi, Andre Renzaho, Serge Resnikoff, Satar Rezaei, Juan A Rivera, Aklilu Abrham Roba, Kedir Teji Roba, Sonia Rodríguez-Ramírez, Leonardo

Roever, Luca Ronfani, Gholamreza Roshandel, Ali Rostami, Gregory Roth, Dietrich Rothenbacher, Ambuj Roy, Enrico Rubagotti, Lesley Rushton, Perminder Sachdev, Basema Saddik, Hosein Safari, Yahya Safari, Roya Safari-Faramani, Mahdi Safdarian, Sare Safi, Saeid Safiri, Rajesh Sagar, Amirhossein Sahebkar, Mohammad Ali Sahraian, Haniye Sadat Sajadi, Nasir Salam, Yahya Salimi, Hamideh Salimzadeh, Joshua A Salomon, Abdallah M. Samy, Juan Sanabria, Maria Dolores Sanchez-Niño, Tania G Sánchez-Pimienta, Taren Sanders, Yingying Sang, Itamar Santos, João Vasco Santos, Milena Santric Milicevic, Bruno Sao Jose, Mayank Sardana, Abdur Razzaque Sarker, Rodrigo Sarmiento-Suárez, Benn Sartorius, Brijesh Sathian, Thirunavukkarasu Sathish, Maheswar Satpathy, Monika Sawhney, Mete Saylan, Mehdi Sayyah, Elke Schaeffner, Maria Inês Schmidt, Ione Schneider, Ben Schöttker, Aletta Schutte, David C Schwebel, Falk Schwendicke, James Scott, Sadaf Sepanlou, Edson Serván-Mori, Seyedmojtaba Seyedmousavi, Hosein Shabaninejad, Azadeh Shafieesabet, Amira Shaheen, Masood Ali Shaikh, Mehran Shams-Beyranvand, Mohammadbagher Shamsi, Heidar Sharafi, Kiomars Sharafi, Mehdi Sharif, Mahdi Sharif-Alhoseini, Hamid Sharifi, Sheikh Mohammed Shariful Islam, Rajesh Sharma, Jun She, Aziz Sheikh, Mika Shigematsu, Min-Jeong Shin, Rahman Shiri, Ivy Shiue, Haitham Shoman, Mark Shrime, Si Si, Soraya Siabani, Tarig Jamal Siddigi, Inga Dora Sigfusdottir, Rannveig Sigurvinsdottir, Diego Augusto Santos Silva, João Pedro Silva, Jasvinder Singh, Virendra Singh, Dhirendra Narain Sinha, Mekonnen Sisay, Eirini Skiadaresi, Badr Sobaih, Soheila Sobhani, Ranjani Somayaji, Moslem Soofi, Reed Sorensen, Joan B Soriano, Ireneous Soyiri, Luciano Sposato, Chandrashekhar T Sreeramareddy, Nadine Steckling-Muschack, Dan J Stein, Murray Stein, Goran Stevanovic, Leo Stockfelt, Mark Stokes, Lela Sturua, Muawiyyah Babale Sufiyan, Rizwan Suliankatchi Abdulkader, Gerhard Sulo, Bruno Sunguya, Patrick Sur, Bryan L. Sykes, Cassandra Szoeke, Rafael Tabarés-Seisdedos, Takahiro Tabuchi, Santosh Tadakamadla, Mesfin Tadese, Ken Takahashi, Eyasu Tamru, Nikhil Tandon, Nuno Taveira, Arash Tehrani-Banihashemi, Gebre Teklemariam Demoz, Awoke Temesgen, Habtamu Temesgen, Mohamad-Hani Temsah, Omar Temsah, Abdullah Terkawi, Tewodros Tesfa, Belay Tessema, Mebrahtu Teweldemedhin, Kavumpurathu Thankappan, Hue Thi Mai, Nu Thi Truong, Nihal Thomas, Giang Thu Vu, George Thurston, Binyam Tilahun, Taavi Tillmann, Quyen G To, Myriam Tobollik, Marcello Tonelli, Roman Topor-Madry, Miguel Tortajada-Girbés, Mathilde Touvier, Marcos Roberto Tovani-Palone, Jeffrey Towbin, Bach Tran, Khanh Bao Tran, Thomas Truelsen, Lorainne Tudor Car, E Murat Tuzcu, Stefanos Tyrovolas, Kingsley Nnanna Ukwaja, Irfan Ullah, Muhammad Sharig Usman, Olalekan A Uthman, Muthiah Vaduganathan, Afsane Vaezi, Pascual Valdez, Santosh Varughese, Tommi Vasankari, Narayanaswamy Venketasubramanian, Santos Villafaina, Francesco S Violante, Sergey Vladimirov, Vasiliy Vlassov, Stein Emil Vollset, Kia Vosoughi, Isidora Vujcic, Fasil Wagnew, Shishay Wahdey, Yasir Waheed, Stephen Waller, Judd Walson, Yafeng Wang, Yuan-Pang Wang, Elisabete Weiderpass, Robert Weintraub, Inbal Weiss Salz, Fitsum Weldegebreal, Andrea Werdecker, Adhena A. Werkneh, Ronny Westerman, Tissa Wijeratne, Andrea Winkler, Alison B. Wiyeh, Charles Shey Wiysonge, Charles Wolfe, Denis Xavier, Gelin Xu, Ali Yadollahpour, Tomohide Yamada, Lijing Yan, Yuichiro Yano, Mehdi Yaseri, Yasin Jemal Yasin, Alex Yeshaneh, Ebrahim M. Yimer, Engida Yisma, Naohiro Yonemoto, Seok-Jun Yoon, Marcel Yotebieng, Mustafa Younis, Mahmoud Yousefifard, Chuanhua Yu, Zoubida Zaidi, Sojib Bin Zaman, Mohammad Zamani, Olifan Zewdie, Anthony Lin Zhang, Hao Zhang, and Kai Zhang.

Drafting the work or revising is critically for important intellectual content

Cristiana Abbafati, Nooshin Abbasi, Olatunji Adetokunboh, Mohsen Afarideh, Sutapa Agrawal, Muktar Ahmed, Mohammad Esmail Akbari, Tomi Akinyemiju, Fares Alahdab, Khurshid Alam, Syed Aljunid, Mamusha Aman, Jason Anderson, Josep M Antó, Marcel Ausloos, Peter Azzopardi, Nam Ba Nguyen, Hamid Badali, Shahrzad Bazargan-Hejazi, Yihalem Abebe Belay, Neeraj Bhala, Oliver Brady, Michael Brauer, Alessandra C Goulart, Franz Castro, Kate Causey, Devasahayam Christopher, Aaron Cohen, Alemneh Kabeta Daba, Ahmad Daryani, Dragos Davitoiu, Jan-Walter De Neve, Selina Deiparine, Getenet Dessie, David Teye Doku, David Edvardsson, Holly Erskine, Sadaf Esteghamati, Andre Faro, Mohammad Hosen Farzaei, Seyed-Mohammad Fereshtehnejad, Joao Fernandes, Nataliya Foigt, Takeshi Fukumoto, Morsaleh Ganji, Segen Gebremeskel, Kebede Embaye Gezae, Maryam Ghasemi-Kasman, Mamata Ghimire, Rakesh Ghosh, Richard Gillum, Rajat Das Gupta, Alexis Jeannine Handal, Hadi Hassankhani, Rasmus Havmoeller, Behzad Heibati, Behnam Heidari, Claudiu Herteliu, Enayatollah Homaie Rad, Farhad Islami, Ravi Prakash Jha, Mikk Jürisson, Manoochehr Karami, Nicholas Kassebaum, Chandrasekharan Nair Kesavachandran, Morteza Abdullatif Khafaie, Behzad Khafaie, Alireza Khaiavi, Gulfaraz Khan, Mona Khater, Yun Jin Kim, Mika Kivimaki, Huong Lan Nguyen, Samantha Larson, James Leigh, Alan Lopez, Azeem Majeed, Mohammad Ali Mansournia, Joemer Maravilla, Ashley Marks, Randall V. Martin, Francisco Rogerlândio Martins-Melo, Melvin Marzan, Benjamin Massenburg, Colm Mcalinden, Fantahun Mekonnen, Tuomo Meretoja, Bahram Mohajer, Karzan Mohammad, Ghobad Moradi, Mehdi Moradinazar, Ghulam Mustafa, Nahid Neamati, Molly Nixon, Olanrewaju Oladimeji, Andrew T. Olagunju, Tinuke O. Olagunju, Bolajoko Olusanya, Jacob Olusanya, Kanyin Ong, Simon Øverland, Konrad Pesudovs, Michael R Phillips, Huyen Phuc Do, Julian Pillay, Richie Poulton, Hai Quang Pham, Alireza Rafiei, Vafa Rahimi-Movaghar, Fatemeh Rajati, Goura K Rath, Marissa Reitsma, Leonardo Roever, Gregory Roth, Saeid Safiri, Juan Sanabria, Damian Santomauro, Milena Santric Milicevic, Bruno Sao Jose, Abdur Razzaque Sarker, Shahabeddin Sarvi, Maheswar Satpathy, James Scott, Seyedmojtaba Seyedmousavi, Mehdi Sharif, Mika Shigematsu, Inga Dora Sigfusdottir, Rannveig Sigurvinsdottir, Diego Augusto Santos Silva, Dhirendra Narain Sinha, Eirini Skiadaresi, Jeff Stanaway, Patrick Sur, Nuno Taveira, Hue Thi Mai, Nu Thi Truong, Hannah J Thomas, Nihal Thomas, Giang Thu Vu, Marcos Roberto Tovani-Palone, Bach Tran, Thomas Truelsen, Olalekan A Uthman, Yuan-Pang Wang, and Robert Weintraub.

Extracting, cleaning, or cataloging data; designing or coding figures and tables

Hedayat Abbastabar, Olatunji Adetokunboh, Muktar Ahmed, Miloud Taki Eddine Aichour, Khurshid Alam, Komal Ali, Jason Anderson, Doanl Bisanzio, Rupert Bourne, Ferrán Catalá-López, Kate Causey, Leslie Cornaby, Ahmad Daryani, Getenet Dessie, Aman Endries, Benjamin Er, Holly Erskine, Andre Faro, Kairsten Fay, Giannina Ferrara, Alize Ferrari, Takeshi Fukumoto, William Gardner, Segen Gebremeskel, Johanna Geleijnse, Ayele Geleto, Rakesh Ghosh, Hadi Hassankhani, Mehdi Hosseinzadeh, Nayu Ikeda, Caleb Irvine, Manoochehr Karami, André Karch, Tesfaye Kassa, Nicholas Kassebaum, Norito Kawakami, Andre Keren, Mohammad Hossein Khosravi, Yun Jin Kim, Barthelemy Kuate Defo, Samantha Larson, Misgan Legesse, Janni Leung, Stefan Ma, Reza Malekzadeh, Helena Manguerra, Mohammad Ali Mansournia, Ashley Marks, Melvin Marzan, Mohsen Mazidi, Anoushka Millear, Lorenzo Monasta, Maziar Moradi-Lakeh, Dariush Mozaffarian, Bruno Nascimento, Minh Nguyen, Emma Nichols, Nobuo Nishi, Kanyin Ong, Katherine Paulson, Hossein Poustchi, Caroline Purcell, Jürgen Rehm, Aklilu Abrham Roba, Yesenia Roman, Luca Ronfani, Enrico Rubagotti, Lesley Rushton, Hosein Safari, Saeid Safiri, Devashri Salvi, Abdallah M. Samy, Damian Santomauro, Maheswar Satpathy, Monika Sawhney, Sadaf Sepanlou, Seyedmojtaba Seyedmousavi, Mehdi Sharif, Farhad Shokraneh, Soraya Siabani, Patrick Sur, Eyasu Tamru, Mohammad Tavakkoli, Awoke Temesgen, Hannah J Thomas, Anna Torre, Irfan Ullah, Rachel L. Updike, Vidhya Venkateswaran, Tissa Wijeratne, Simon Yadgir, Ali Yadollahpour, Hossein Yahyazadeh, Mahmoud Yousefifard, and Stephanie R M Zimsen.

Managing the overall research enterprise

Ashkan Afshin, Tahiya Alam, Brigette F Blacker, Michael Brauer, Deborah Carvalho Malta, Aaron Cohen, Elizabeth Cromwell, Lalit Dandona, Rakhi Dandona, Louisa Degenhardt, Samath Dharmaratne, Charbel El Bcheraoui, Kara Estep, Valery Feigin, Kyle Foreman, Nancy Fullman, Thomas Fürst, Emmanuela Gakidou, Caitlin Hawley, Simon Hay, Spencer James, Nicholas Kassebaum, Ibrahim Khalil, Kristopher Krohn, Hmwe Kyu, Xiaofeng Liang, Stephen Lim, Alan Lopez, Rafael Lozano, Ashley Marks, George Mensah, Molly Miller-Petrie, Awoke Misganaw, Ali Mokdad, Kate Muller, Christopher Murray, Mohsen Naghavi, Molly Nixon, Elaine Nsoesie, Helen Olsen, Robert Reiner, Gregory Roth, Joshua A Salomon, Benn Sartorius, David Smith, Mari Smith, Jeffrey Stanaway, Roman Topor-Madry, Stein Emil Vollset, Theo Vos, Andrea Werdecker, and Harvey Whiteford.

Did not provide contribution information

Dash A P, Dilaram Acharya, Rufus A. Adedoyin, Amha Admasie, Samiah Alam, Nahla Anber, Jalal Arabloo, Traolach Brugha, Maryam Farvid, Ali Abar Fazaeli, John Ji, Narges Karimi, Mohammad Khazaei, Giancarlo Logroscino, Ana Laura Manda, Mousa Mohammadnia-Afrouzi, Simin Mouodi, Sreebhushan Raju, Sahar Saeedi Moghaddam, Zikria Saleem, Soraya Seedat, Mehdi Shahbazi, Reza Shirkoohi, Michelle Subart, and Paul Yip.

Section 1: GBD overview

Section 1.1: Locations of the Analysis

The locations included in Global Burden of Diseases, Injuries, and Risk Factors Study 2017 (GBD 2017) have been arranged into a set of hierarchical categories composed of seven super-regions and a further nested set of 21 regions containing 195 countries and territories. The locations for which GBD estimated global, regional, and national risk exposure, relative risk, theoretical minimum-risk exposure level (TMREL), and population attributable fractions (PAFs), have not expanded following GBD 2015. Subnational estimation in GBD 2017 includes Brazil, China, India, Indonesia, Japan, Kenya, Mexico, South Africa, Sweden, the United Kingdom, and the United States, and new subnational assessments at the administrative one level for Ethiopia, Iran, Norway, and Russia and by Maori ethnicity for New Zealand. For this publication, we present subnational estimates in figures only for all subnational countries with the exception of the new assessments, which will be reported in separate publications. Select subnational estimates are also included in supplementary results appendix. Combined, there are a total of 390 locations at the first subnational unit level. Included in subnational Level 1 locations are countries that have been subdivided into the first subnational level, such as states or provinces, for the GBD analysis; subnational Level 2 only applies to India, England, and Russia. For this paper, we present data at the national and territory level.

Section 1.2: Time Period of the Analysis

A complete set of risk-specific exposure, relative risk, TMREL, and PAFs were computed for the years 1990-2017. All GBD 2017 results and online data visualisations are available at http://vizhub.healthdata.org/gbd-compare with access to results for all GBD metrics.

Section 1.3: Statement of GATHER Compliance

This study complies with the Guidelines for Accurate and Transparent Health Estimates Reporting (GATHER) recommendations. We have documented the steps involved in our analytical procedures and detailed the data sources used. See Appendix Table 1 for the GATHER checklist.

The GATHER recommendations may be found here: <u>http://gather-statement.org/</u>

Section 1.4: GBD risk factor hierarchy

In this analysis, we focus on three groups of risk factors: behavioural, environmental and occupational, and metabolic. The GBD 2017 risk factors hierarchy and Levels are summarised in Appendix Table 2.

The GBD risk list continues to evolve to reflect the policy relevance, public health, and medical care importance of major risk factors. The risk factors list expanded following feedback from GBD 2013 and input from GBD 2015 collaborators. Three risks were added to the list for GBD 2017.

Section 1.5: List of abbreviations

APCSC: Asia-Pacific Cohort Studies Collaboration

BMI: body-mass index BMD: bone mineral density CKD: chronic kidney disease COD: causes of death CODEm: cause of death ensemble modelling COPD: chronic obstructive pulmonary disease CSA: childhood sexual abuse CSMR: cause-specific mortality rate CRA: comparative risk assessment CVD: cardiovascular disease DALY: disability-adjusted life-year DHS: Demographic and Health Survey DRI: data representativeness index EMR: excess mortality rate FAO: Food and Agriculture Organization FPG: fasting plasma glucose GATHER: Guidelines for Accurate and Transparent Health Estimates Reporting GBD: Global Burden of Disease GoF: goodness of fit ID: Iron deficiency IDA: Iron deficiency anaemia IER: integrated exposure response IHD: ischemic heart disease ILO: International Labour Organization IPV: intimate partner violence JMP: Joint Monitoring Project LDI: lag distributed income per capita LMIC: Low and Middle-Income Countries LRI: lower respiratory infection

MCMC: Markov Chain Monte Carlo simulations MDG: Millennium Development Goal MICS: Multiple Indicator Cluster Surveys MoM: method of moments NCD: non-communicable disease OER: observed-to-expected ratio PAF: population attributable fraction PDF: probability density function $PM_{2.5}$: particulate matter <2.5 μ m in diameter **PSC: Prospective Cohort Study** RCT: randomised controlled trial **REDCap:** Research Electronic Data Capture RMSE: root mean square error **RR:** relative risk SBP: systolic blood pressure SD: standard deviation SDG: sustainable development goal SDI: Socio-demographic Index SEER: Surveillance, Epidemiology, and End Results Program SEV: summary exposure value SIR: smoking impact ratio SSB: sugar-sweetened beverages ST-GPR: spatiotemporal Gaussian process regression **TB:** tuberculosis TMREL: theoretical minimum-risk exposure level TSNA: tobacco-specific nitrosamines UI: uncertainty interval WCRF: World Cancer Research Fund WHO: World Health Organization

YLD: years lived with disability

YLL: years of life lost

Section 1.6: GBD results overview

Results from the Global Burden of Disease Study (GBD 2017) are now measured in terabytes. Results are available in an interactive data downloading tool on the Global Health Data exchange (GHDx). Data and underlying code used for this analysis will be made publicly available pending manuscript acceptance.

The current version of the data download tool is available in the GHDx and contains core summary results for the GBD 2017: <u>http://ghdx.healthdata.org/gbd-results-tool</u>. The core summary results include deaths, YLLs, years lived with disability (YLDs), and disability-adjusted life-years (DALYs). The GHDx includes data for causes, risks, cause-risk attribution, aetiologies, and impairments.

In the GBD 2017 version, the GHDx tool also contains measures such as prevalence and incidence as well as rate of change data. Data above a certain size cannot be viewed online but can be downloaded. Depending on the size of the download, users may need to enter an email address; a download location will be sent to them when the files are prepared.

Section 1.7: Data input sources overview

GBD 2017 incorporated a large number and wide variety of input sources to estimate mortality, causes of death and illness, and risk factors for 195 countries and territories from 1990-2017. These input sources are accessible through an interactive citation tool available in the GHDx.

Users can retrieve citations for a specific GBD component, cause or risk, and location by choosing from the available selection boxes. They can then view and access GHDx records for input sources and export a CSV file that includes the GHDx metadata, citations, and information about where the data were used in GBD. Additional metadata for each input source are available through the citation tool, as required by the GATHER statement.

The citation tool is accessible through the GHDx at <u>http://ghdx.healthdata.org/gbd-2016/data-input-</u>sources.

Section 1.8: Funding Sources

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Section 2: Risk factor estimation

Section 2.1: Overview

The comparative risk assessment (CRA) conceptual framework was developed by Murray and Lopez,¹ who established a causal web of hierarchically organised risks or causes that contribute to health outcomes, which allows for quantification of risks or causes at any Level in the framework. In GBD 2017, as in previous iterations of the GBD study, we evaluated a set of behavioural, environmental and occupational, and metabolic risks, where risk-outcome pairs were included based on evidence rules (see Section 2.2.1). These risks were organised in four hierarchical Levels, where Level 1 represents the overarching categories (behavioural, environmental and occupational, and metabolic) nested within Level 1 risks; Level 2 contains both single risks and risk clusters (such as child and maternal malnutrition); Level 3 contains the disaggregated single risks from within Level 2 risk clusters (such as low birthweight and short gestation); and Level 4 details risks with the most granular disaggregation, such as for specific occupational carcinogens, the subcomponents of child growth failure (stunting, wasting, underweight), and suboptimal breastfeeding (discontinued and non-exclusive breastfeeding). At each level of risk, we evaluated whether risk combinations were additive, multiplicative, or shared common pathways for intervention. This approach allows the quantification of the proportion of riskattributable burden shared with another risk or combination of risks and the measurement of potential overlaps between behavioural, environmental and occupational, and metabolic risks. To date in the GBD we have not quantified the contribution of other classes of risk factors illustrated in Appendix Table 3. We do provide some insights into the potential magnitude of distal social, cultural, and economic factors through an analysis of the relationship between risk exposures and development measured using the Socio-demographic Index (SDI) (see Appendix Section 2.9).

Two types of risk assessments are possible within the CRA framework: attributable burden and avoidable burden. Attributable burden is the reduction in current disease burden that would have been possible if past population exposure had shifted to an alternative or counterfactual distribution of risk exposure. Avoidable burden is the potential reduction in future disease burden that could be achieved by changing the current distribution of exposure to a counterfactual distribution of exposure. Murray and Lopez identified four types of counterfactual exposure distributions: (1) theoretical minimum risk; (2) plausible minimum risk; (3) feasible minimum risk; and (4) cost-effective minimum risk.² The theoretical minimum risk level (TMREL) is the level of risk exposure that minimises risk at the population level, or the level of risk that captures the maximum attributable burden. Other possible forms of risk quantification include plausible minimum risk – which reflects the distribution of risk that is conceivably possible and would minimise population-level risk if achieved – while feasible minimum risk describes the lowest risk distribution that has been attained within a population, and the cost-effective minimum risk is the lowest risk distribution for a population that can be attained in a cost-effective manner. Because no robust set of forecasts for all components of GBD is available, in this study we focus on quantifying attributable burden using the theoretical minimum risk counterfactual distribution. Appendix Table 3 shows the eight possible types of risk quantification within the CRA framework, with the hatched box representing the type of CRA currently undertaken by the GBD study. As per the definition of avoidable burden, risk reversibility would be incorporated into this type of assessment, as it would involve reducing risk to the counterfactual for the index year, given a history of past risk exposure. Given the focus in this study on attributable burden, risk reversibility is not a criteria used in estimation here.

In general, this analysis follows the CRA methods used since GBD 2015.³ The methods described here provide a high-level overview of the analytical logic with a focus on areas of notable change from the methods employed in GBD 2015. Here we aim to provide sufficient detail on the methodology and overall structure of the estimation process. This study complies with the GATHER recommendations proposed by the World Health Organization (WHO) and others, which include recommendations on documentation of data sources, estimation methods, and statistical analysis (Appendix Table 1).⁴

Section 2.2: Step 1. Effect size estimation

Section 2.2.1: Collate relative risk data

Criteria for inclusion of risk-outcome pairs

In this study, as in GBD 2016, we have included risk-outcome pairs that we have assessed as meeting the World Cancer Research Fund (WCRF) grades of convincing or probable evidence.⁵ In this framework, convincing evidence consists of biologically plausible associations between exposure and disease established from multiple epidemiological studies in different populations. Evidentiary studies must be substantial, include prospective observational studies, and where relevant, randomised controlled trials (RCTs) of sufficient size, duration, and quality, and showing consistent effects. Probable evidence is similarly based on epidemiological studies with consistent associations between exposure and disease, but for which shortcomings in the evidence exist, such as insufficient trials (or prospective observational studies) available.

The World Cancer Research Fund grading system

Convincing evidence

Evidence based on epidemiological studies showing consistent associations between exposure and disease, with little or no evidence to the contrary. The available evidence is based on a substantial number of studies including prospective observational studies and where relevant, randomised controlled trials of sufficient size, duration, and quality showing consistent effects. The association should be biologically plausible.

Probable evidence

Evidence based on epidemiological studies showing fairly consistent associations between exposure and disease, but for which there are perceived shortcomings in the available evidence or some evidence to the contrary, which precludes a more definite judgment. Shortcomings in the evidence may be any of the following: insufficient duration of trials (or studies); insufficient trials (or studies) available; inadequate sample sizes; or incomplete follow-up. Laboratory evidence is usually supportive. The association should be biologically plausible.

Possible evidence

Evidence based mainly on findings from case-control and cross-sectional studies. Insufficient randomised controlled trials, observational studies, or non-randomised controlled trials are available. Evidence based on non-epidemiological studies, such as clinical and laboratory investigations, is supportive. More trials are needed to support the tentative associations, which should be biologically plausible.

Insufficient evidence

Evidence based on findings of a few studies which are suggestive, but insufficient to establish an association between exposure and disease. Little or no evidence is available from randomised controlled trials. More well designed research is needed to support the tentative association.

Causal criteria

As in GBD 2015 and 2016, to be more objective, consistent, and transparent in our evaluation of the causal relationship, we summarized epidemiologic evidence supporting causality for each risk-outcome pair (Appendix Table 4 and Appendix Table 5). For each pair, we collected data on the following domains:

Domains	Description
RCTs of disease endpoint	Number of independent RCTs evaluating the effect of the risk on the disease endpoint
	Percent of independent RCTs showing significant effect in the opposite direction
	Percent of independent RCTs showing no effect
Prospective observational studies of disease endpoint	Number of independent prospective observational studies evaluating the association of the risk with the disease endpoint
	Percent of independent prospective observational studies with significant association in the opposite direction
Strength	Lower Limit of relative risk (RR) in observational studies> 1.5 (Yes/No)
Dose response	Evidence of the dose-response relationship between the risk and the outcome (Yes/No)
Biologic plausibility	Potential biologic mechanism that could explain the effect of the risk on the disease endpoint (Yes/No)
Analogy	Evidence on the relationship between the risk factor and a disease endpoint from the same category (Yes/No)

For risk-outcome pairs with less than five prospective studies, we summarized evidence from casecontrol studies as well including (a) the number of independent case-control studies evaluating the association of the risk with the disease endpoint and (b) percent of independent case control studies with significant association in the opposite direction.

Section 2.2.2: Determine relative risks

Effect size estimation

The relative risk by level of exposure, or by cause, for mortality or morbidity can be found in published and unpublished primary studies or in secondary studies that summarize relative risks. In Step 1a of the analytical process (Appendix Figure 1), we collated information from randomised controlled trials, cohort, pooled cohort, and case-control studies, and in Step 1b, used these data to determine the relative risk for the risk-outcome pairs included in GBD 2017 (Appendix Table 6). For most risks, data from pooled cohorts, or meta-analyses of cohorts, were used; in the case of the risk of cataracts from household air pollution, cohort data were not available, and instead we used case-control data. We estimated relative risks of mortality and morbidity for 65 risk factors for which we determined attributable burden using relative risk and exposure. We incorporated relative risks from studies that controlled for confounding but not for factors along the causal pathway between exposure and outcome. For risk-outcome pairs with evidence available for only one of mortality or morbidity, we generally assumed that the estimated relative risks applied equally to both. Given evidence of statistically different relative risks for mortality and morbidity, we incorporated different relative risks for each. We did not find that relative risks were consistently higher or lower for mortality compared with morbidity. Details and citation information for the data sources used for relative risks are provided in searchable form through a new web-tool (<u>http://ghdx.healthdata.org/</u>). Available data sources for determining relative risks varied across risks. Details on how relative risks were calculated for each risk can be found in Appendix Section 4: Risk-specific estimates.

For all outcomes related to unsafe sex, the relative risk and exposure framework was not used to estimate attributable burden. For unsafe sex and HIV, we used a direct attribution approach to address the lack of data on unsafe sexual practices in most populations. The proportion of HIV attributable to unsafe sex was modelled directly using DisMod-MR 2.1 from data on the fraction of cases identified as being through sexual transmission, intravenous drug use, or blood transfusion.

For risks estimated from a continuous exposure distribution where the effect size was reported by categories in pooled or meta-analysis studies, we converted those categories to relative risk per unit increase in exposure. This implies a linear increase in the log of the relative risk and exposure; various studies have suggested this is a reasonable approximation of the dose-response curve for many risks. An example of this is high systolic blood pressure, where data from the Prospective Cohort Study (PSC) and the Asia-Pacific Cohort Studies Collaboration (APCSC) were well-described by a linear increase in the logarithm of the relative risk by a 10-unit increase in high systolic blood pressure. This approximately log-linear relationship suggests that the proportional difference in the age-specific risk of stroke death associated with a given absolute difference in exposure is about the same at all levels of risk. Many meta-analyses convert relative risks to per unit increase for convenience, particularly when studies choose different categories that could not otherwise be compared. The log-linear approximation appears plausible⁶ even where there is limited consensus on the appropriate TMREL. Where there were insufficient samples in the primary studies at high levels of exposure to inform the shape of the tail of the distribution, we applied a cap to the maximum relative risk using the midpoint of the last category for which a relative risk was reported.

Section 2.3: Step 2. Exposure estimation

Section 2.3.1: Collate exposure data

Systematic reviews

For GBD 2017, we conducted systematic literature reviews for 23 risks. For other risk factors, only a small fraction of the existing data appears in the published literature and other sources predominate such as survey data and satellite data. Data were systematically screened from household surveys archived in the Global Health Data Exchange (ghdx.healthdata.org), including Demographic and Health Surveys, Multiple Indicator Cluster Surveys, Living Standards Measurement Surveys, and Reproductive Health Surveys. Other national health surveys were identified based on survey series that had yielded usable data for past rounds of GBD, sources suggested to us by in-country collaborators, and surveys identified in major multinational survey data catalogues, such as the International Household Survey Network and the WHO Central Data Catalog, as well as through country Ministry of Health and Central Statistical Office websites. Citations for all data sources used for risk factor estimation in GBD 2017 are

provided in searchable form through a web-tool (http://ghdx.healthdata.org/). A description of the search terms employed for risk-specific systematic reviews are detailed by cause in Appendix Section 4.

Information on systematic reviews were managed using Research Electronic Data Capture (REDCap) electronic data capture tools hosted at the University of Washington.⁷ REDCap is a secure, web-based application designed to support data capture for research studies, providing 1) an intuitive interface for validated data entry; 2) audit trails for tracking data manipulation and export procedures; 3) automated export procedures for seamless data downloads to common statistical packages; and 4) procedures for importing data from external sources

Search terms

Search terms for updates of systematic reviews for GBD 2017 are shown by risk in Appendix Section 4.

Survey data preparation

For GBD 2017, survey data constitutes a substantial part of the underlying data used in the estimation process. During extraction, we concentrate on demographic variables (such as location, gender, age), survey design variables (such as sampling strategy and sampling weights), and the variables used to define the population estimate (such a prevalence or a proportion) and a measure of uncertainty (standard error, confidence interval or sample size and number of cases).

Section 2.3.2: 2b. Adjust exposure data

A number of adjustments were applied to extracted exposure sources in order to make the data more consistent and suitable for modelling. Commonly applied adjustments included age-sex splitting, adding study-level covariates, and bias correction. Age-sex splitting was applied to literature data reported by age or sex but not by age and sex assuring that the total number of cases remained as reported. If a source did not report sample size by age or sex, we applied the age-sex distribution of the population for the same location and year to the reported total sample size. We relied on the metaregression component of DisMod-MR 2.1 for most of the bias correction of data for variations in study attributes such as case definitions and measurement method. DisMod-MR 2.1 calculates a single adjustment that is applied regardless of age, sex, or location. If enough data were available to differentiate these adjustments by age, sex, or location, or if detailed survey data were available to make more precise adjustments between different thresholds on a biochemical measure, we applied bias corrections to the data before entry into DisMod-MR 2.1.

Section 2.3.3: 2c. Estimate exposure

Mean exposure estimation

In Step 2a of the estimation process, we used systematic literature reviews to identify risk factor exposure studies published or identified since GBD 2016 and combined these with existing data from household and health examination surveys, census, morbidity, or satellite imagery and ground sensor data (used for estimation of $PM_{2.5}$ [particulate matter <2.5 µm in diameter]). Certain risks, such as diet and alcohol consumption, also incorporated administrative record systems. Data sources used in estimating risk factor exposure can be accessed through the data source tool at http://ghdx.healthdata.org/.

Once data were collected and compiled, step 2b of the analytical flowchart describes the adjustments applied, where necessary, to correct for bias. Examples of these adjustments include: use of urban studies for lead; crosswalks between different measurements, methods, and definitions, such as for self-

report of obesity and glycated haemoglobin (HbA1C) for diabetes; and age-sex splitting of data, such as for fasting plasma glucose, cholesterol, and systolic blood pressure that may be reported from broad age-groups.

For the GBD, we developed two modelling approaches, a Bayesian meta-regression model (DisMod-MR 2.1) and a spatiotemporal Gaussian process regression model (ST-GPR), to pool data from different sources, control and adjust for bias in data, and incorporate other types of information such as countrylevel covariates. DisMod-MR 2.1 and ST-GPR are mixed effect models that borrow information across age, time, and locations to synthesise multiple data sources into unified estimates of levels and trends. A detailed description of the likelihood used for estimation, and a full description of improvements made for DisMod-MR 2.1, are detailed by Vos and colleagues⁸ with additional detail in the appendix to that paper. The ST-GPR model has three main hyper-parameters that control for smoothing across time, age, and location. Values for these hyper-parameters were selected based on cross-validation. Crossvalidation tests were conducted for different combinations of the hyper-parameters for three types of models: one data-sparse model, one data-moderate model, and one data-dense model. In each test, 20% of the data were held out and the performance of each combination of hyper-parameters evaluated on the held out data. For each hyper-parameter combination, 10 cross-validation tests were conducted. The performance of each model in predicting the withheld 20% of the data was evaluated using a combined measure based on root mean square error (RMSE) and uncertainty interval coverage. A detailed description of the ST-GPR process regression can be found below in Appendix Section 2.3.3.

The main difference between these methods is their power to include unstructured types of data by sex and age group and in their degree of flexibility. Step 2c in Appendix Figure 1 outlines the use of DisMod-MR 2.1 for 10 risk factors where data were available by different age intervals or mixed sex groups; DisMod-MR 2.1 is the preferred tool in these cases because of its ability to integrate over age and adjust for different exposure definitions in the data; however, the use of Bayesian Markov Chain Monte Carlo (MCMC) simulations with large volumes of data renders the analysis computationally intensive and reduces the number of iterations that are possible. If large volumes of standard age-group data are available – as is generally the case for metabolic risks – using ST-GPR becomes the preferred approach.

In some cases, we adapted our methods of modelling exposure to risks where necessary to account for complexities in the risk-outcome relationship or the need for particular handling of data, for example, dietary risks and ambient air pollution (see Appendix Section 4 for more detail). A complete list of risks and the analytical method used is reported in Appendix Table 2. Additional details for adjustments or adaptations to particular risk models are located in Appendix Section 4.

DisMod-MR 2.1 Estimation

DisMod-MR 2.1 description

Until GBD 2010, nonfatal estimates in burden of disease assessments were based on a single data source on prevalence, incidence, remission or a mortality risk selected by the researcher as most relevant to a particular location and time. For GBD 2010, we set a more ambitious goal: to evaluate all available information on a disease that passes a minimum quality standard. That required a different analytical tool that would be able to pool disparate information presented in varying age groupings and from data sources using different methods. The DisMod-MR 1.0 tool used in GBD 2010 evaluated and pooled all available data, adjusted data for systematic bias associated with methods that varied from the reference and produced estimates by world regions with uncertainty intervals using Bayesian statistical methods. For GBD 2013, the improved DisMod-MR 2.0 had increased computational speed allowing computations that were consistent between all disease parameters at the country rather than region Level. The hundred-fold increase in speed of DisMod-MR 2.0 was partly due to a more efficient rewrite of the code in C++ but also by changing to a model specification using log rates rather than a negative binomial model used in DisMod-MR 1.0. In cross-validation tests, the log rates specification worked as well or better than the negative binomial specification.⁹ For GBD 2015, we rewrote the 'wrapper' code that organizes the flow of data and settings at each level of the analytical cascade. The sequence of estimation occurs at five Levels: global, super-region, region, country and, where applicable, subnational location. The super-region priors are generated at the global Level with mixed-effects, nonlinear regression using all available data; the super-region fit, in turn, informs the region fit, and so on down the cascade. The wrapper gives analysts the choice to branch the cascade in terms of time and sex at different levels depending on data density. The default used in most models is to branch by sex after the global fit but to retain all years of data until the lowest Level in the cascade. Appendix Figure 2 summarizes the DisMod-MR process.

In updating the 'wrapper,' we consolidated the code base into a single language, Python, to make the code more transparent and efficient and to better deal with subnational estimation. The computational engine is limited to three levels of random effects; we differentiate estimates at the super-region, region and country Level. In GBD 2013, the subnational units of China, the UK and Mexico were treated as 'countries' such that a random effect was estimated for every location with contributing data. However, the lack of a hierarchy between country and subnational units meant that the fit to country data contributed as much to the estimation of a subnational unit as the fits for all other countries in the region. We found inconsistency between the country fit and the aggregation of subnational level of random effects required a prohibitively comprehensive rewrite of the underlying DisMod-MR engine. Instead, we added a fifth layer to the cascade, with subnational estimation informed by the country fit and country covariates, plus an adjustment based on the average of the residuals between the subnational location's available data and its prior. This mimicked the impact of a random effect on estimates between subnationals.

In GBD 2015, we also improved how country covariates differentiate nonfatal estimates for diseases with sparse data. The coefficients for country covariates are re-estimated at each Level of the cascade. For a given location, country coefficients are calculated using both data and prior information available for that location. In the absence of data, the coefficient of its parent location is used, in order to utilize the predictive power of our covariates in data sparse situations.

For GBD 2016, the computational engine (DisMod-MR 2.1) remained substantively unchanged from GBD 2015. We changed the prediction year set to generate fits for the years 1990, 1995, 2000, 2005, 2010, and 2016. We updated the age prediction sets to include age groups 80-84, 85-89, 90-94, and 95+, to comply with changes across all functional areas of the GBD. We also expanded the set of locations where subnational units are modelled; the set now includes: Brazil, China, England, India, Indonesia, Japan, Kenya, Mexico, Saudi Arabia, South Africa, Sweden, and the United States.

In GBD 2017, we continued to use DisMod-MR 2.1, as there were no substantial changes. Updates to computation include extending the terminal prediction year to 2017 and additional subnational units in

Ethiopia, Iran, New Zealand, Norway, and Russia. Saudi Arabia was also modelled only at the national level in 2017.

The flowchart for the DisMod-MR 2.1 process can be found in Appendix Figure 2.

DisMod-MR 2.1 likelihood estimation

Analysts have the choice of using a Gaussian, log-Gaussian, Laplace or Log-Laplace likelihood function in DisMod-MR 2.1. The default log-Gaussian equation for the data likelihood is:

$$-log[p(y_j|\Phi)] = log(\sqrt{2\pi}) + log(\delta_j + s_j) + \frac{1}{2} \left(\frac{log(a_j + \eta_j) - log(m_j + \eta_j)}{\delta_j + s_j}\right)^2$$

where, y_j is a 'measurement value' (i.e., data point); Φ denotes all model random variables; η_j is the offset value, eta, for a particular 'integrand' (prevalence, incidence, remission, excess mortality rate, with-condition mortality rate, cause-specific mortality rate, relative risk or standardized mortality ratio) and a_j is the adjusted measurement for data point j, defined by:

$$a_j = e^{(-u_j - c_j)} y_j$$

where u_j is the total 'area effect' (i.e., the sum of the random effects at three Levels of the cascade: super-region, region and country) and c_j is the total covariate effect (i.e., the mean combined fixed effects for sex, study level and country level covariates), defined by:

$$c_j = \sum_{k=0}^{K[I(j)]-1} \beta_{I(j),k} \hat{X}_{k,j}$$

with standard deviation

$$s_j = \sum_{l=0}^{L[I(j)]-1} \zeta_{I(j),l} \hat{Z}_{k,j}$$

where k denotes the mean value of each data point in relation to a covariate (also called x-covariate); I(j) denotes a data point for a particular integrand, j; $\beta_{l(j),k}$ is the multiplier of the kth x-covariate for the ith integrand; $\hat{X}_{k,j}$ is the covariate value corresponding to the data point j for covariate k; I denotes the standard deviation of each data point in relation to a covariate (also called z-covariate); $\zeta_{l(j),k}$ is the multiplier of the Ith z-covariate for the ith integrand; and δ_j is the standard deviation for adjusted measurement j, defined by:

$$\delta_j = \log[y_j + e^{(-u_j - c_j)}\eta_j + c_j] - \log[y_j + e^{(-u_j - c_j)}\eta_j]$$

Where m_j denotes the model for the jth measurement, not counting effects or measurement noise and defined by:

$$m_j = rac{1}{B(j) - A(j)} \int_{A(j)}^{B(j)} I_j$$
(a) da

where A(j) is the lower bound of the age range for a data point; B(j) is the upper bound of the age range for a data point; and I_j denotes the function of age corresponding to the integrand for data point j.

Spatiotemporal Gaussian process regression

Spatiotemporal Gaussian process regression (ST-GPR) has been used for risk factors where the data density is sufficient to estimate a very flexible time trend. The flowchart showing the analytic steps can be found in Appendix Figure 3. The approach is a stochastic modelling technique that is designed to detect signals amidst noisy data. It also serves as a powerful tool for interpolating non-linear trends.^{10,11} Unlike classical linear models that assume that the trend underlying data follows a definitive functional form, GPR assumes that the specific trend of interest follows a Gaussian Process, which is defined by a mean function $m(\cdot)$ and a covariance function $Cov(\cdot)$. For example, let $p_{c,a,s,t}$ be the exposure, in normal, log, or logit space, observed in country c, for age group a, and sex s at time t:

$$(p_{c,a,s,t}) = g_{c,a,s}(t) + \epsilon_{c,a,s,t}$$

where

$$\begin{split} & \epsilon_{c,a,s,t} \sim Normal(0,\sigma_p^2), \\ & g_{c,a,s}(t) \sim GP\left(m_{c,a,s}(t), Cov\left(g_{c,a,s}(t)\right)\right) \end{split}$$

The derivation of the mean and covariance functions, $m_{c,a,s}(t)$ and $Cov(g_{c,a,s}(t))$, along with a more detailed description of the error variance (σ_n^2) , is described below.

Estimating mean functions

We estimated mean functions using a two-step approach. To be more specific, $m_{c,a,s}(t)$ can be expressed, depending on the exposure transformation, as:

$$log(p_{c,a,s}(t)) = X_{c,a,s}\beta + h(r_{c,a,s,t})$$
$$logit(p_{c,a,s}(t)) = X_{c,a,s}\beta + h(r_{c,a,s,t})$$
$$p_{c,a,s}(t) = X_{c,a,s}\beta + h(r_{c,a,s,t})$$

where $X\beta$ is the summation of the components of a hierarchical mixed-effects linear regression, including the intercept and the product of covariates with their corresponding fixed effect coefficients. Some models were run as hierarchical mixed-effects linear regressions, with random effects on the levels of the geographic hierarchy. For most mixed-effects models, random effects were only used in the fit, not in the prediction. The second part of the equation, $h(r_{c,a,s,t})$, is a smoothing function for the residuals, $r_{c,a,s,t}$, derived from the linear model.³ Descriptions of exposure transformations and which covariates were used in linear models can be found in Appendix Section 4, which described the riskspecific estimation approaches. Some models used a custom stage-1 estimate – these risks will have detailed information on their mixed-effect estimation process in the risk-specific appendix sections.

While the linear component captures the general trend in exposures over time, much of the data varaibility may still not be adequately accounted for. To address this, we fit a locally weighted polynomial regression (LOESS) function $h(r_{c,a,s,t})$ to systematically estimate this residual variability by

borrowing strength across time, age, and space patterns (the spatiotemporal component of ST-GPR).^{12,13} The time adjustment parameter, defined by λ , aims to borrow strength from neighboring time points (i.e. the exposure in this year is highly correlated with exposure in the previous year but less so further back in time). The age adjustment parameter, defined by ω , borrows strength from data in neighboring age groups. The space adjustment parameter, defined by ξ , aims to borrow strength across the hierarchy of geographical locations. This year, we further combined the spatial and temporal weights into a single space-time weight, to allow the amount of spatial weight given to a particular point $r_{c,a,s,t}$ to fluctuate given the data availability at each time t and location-level I in the location hierarchy.

Let $w_{c,a,s,t}$ be the final weight assigned to observation $r_{c,a,s,t}$ with reference to a focal observation r_{c_0,a_0,s_0,t_0} . We first generated a temporal weight t. $w_{c,a,s,t}$ for smoothing over time, which was based on the scaled distance along the time dimension of the two observations¹⁴:

$$t.w_{c,a,s,t} = \frac{1}{e^{\lambda|t-t_0|}}$$

Next, we generated a spatial weight to smooth over geography. Specifically, we defined a geospatial relationship by categorizing data based on the GBD location hierarchy (Appendix Table 7).

In previous GBD iterations, a vector of spatial weights corresponding to each level of the location hierarchy was derived as $[\xi, \xi * (1 - \xi)^{n_1 - 1}, \dots, \xi * (1 - \xi)^{n_i - 1}, (1 - \xi)^{n_i}]$, where n_i , designated the number of location levels in between the given location and the global level and ζ was typically between .7 - .99. Under the previous spatial weighting system, all country datapoints would receive a weight of ζ , all regional datapoints a weight of $\zeta * (1 - \zeta)$, etc, no matter how much data was available in the country compared to the region. For example, if there was only a single datapoint for a given country and ζ was set to .7, that lone datapoint would receive 70% of the spatial weight.

This year, we reformulated zeta to act as a scalar on a given datapoint given its proximity to the target location:

$$t.w_{c.a.s.t} = \zeta^{|c-c_0|}$$

For example, estimating a country would use the following weighting scheme:

- Country data: $\zeta^0 = 1$
- Regional data not from the country being estimated: ζ^1
- Data from other regions in the same super region: ξ^2
- Global data from other super regions: ζ^3

Under the new spatial weighting specification, typical values of ζ range from [.001, .2], where ζ can be interpreted as the amount to downweight regional datapoints compared to country datapoints for a given estimating country. For example, for a given datapoint $r_{c,a,s,t}$ and $\zeta = .01$, a datapoint not within country *c* but within the same region *r* as $r_{c,a,s,t}$ would be assigned $\frac{1}{100}$ the weight of a datapoint within the country.

The spatial and temporal weights were then multiplied and summed across each level of the location hierarchy, and normalized for each time period t. This allows the space-time weight to implicitly take

into account the amount of data available at the country vs. region vs super-region level and attribute spatial weight accordingly.

Given a normalization constant,

$$K_i = \sum_{c \in C} s. w_{c,t} * t. w_{c.t} + \sum_{c \in R} s. w_{c,t} * t. w_{c.t} + \sum_{c \in SR} s. w_{c,t} * t. w_{c.t}$$

the final space-time weight would then equal

$$w_{c,a,s,t}' = \frac{s.w_{c,t} * t.w_{c,t}}{K_i}$$

Finally, we calculated the weight $w''_{c,a,s,t}$ to smooth over age, which is based on a distance along the age dimension of two observations. For a point between the age a of the observation $r_{c,a,s,t}$ and a focal observation r_{c_0,a_0,s_0,t_0} , the weight is defined as follows:

$$w_{c,a,s,t}^{\prime\prime} = \frac{1}{e^{\omega|a-a_0|}}$$

The final weights would then be computed by simply multiplying the space-time weights and age weights and normalizing so all weights for a given time period *t* sum to 1. A full derivation of weights for each category follow, assuming the location being estimated was a country, follows:

1) If the observation $r_{c,t}$ belongs to the same country c_0 of the focal observation r_{c_0,t_0} :

$$w_{c,a,s,t} = \frac{(w'_{c,a,s,t}w''_{c,a,s,t})}{\sum_{c=c_0} (w'_{c,a,s,t}w''_{c,a,s,t})} \quad \forall c = c_0$$

2) If the observation $r_{c,t}$ belongs to a different country than the focal observation r_{c_0,t_0} , but both belong to the same region R:

$$w_{c,a,s,t} = \frac{(w'_{c,a,s,t}w''_{c,a,s,t})}{\sum_{c \neq c_0} (w'_{c,a,s,t}w''_{c,a,s,t})} \qquad \forall c \neq c_0 \cap R[c] = R[c_0]$$

3) If the observation $r_{c,t}$ belongs to the same super region SR but to a both different country c_0 and region $R[c_0]$ than the focal observation r_{c_0,t_0} :

$$w_{c,a,s,t} = \frac{(w'_{c,a,s,t}w''_{c,a,s,t})}{\sum_{c \neq c_0} (w'_{c,a,s,t}w''_{c,a,s,t})} \qquad \forall c \neq c_0 \cap R[c] \neq R[c_0] \cap SR[c] = SR[c_0]$$

4) If the observation $r_{c,t}$ is from a different super region than the focal observation r_{c_0,t_0} (ie. all other data currently not receiving a weight):

$$w_{c,a,s,t} = \frac{(w'_{c,a,s,t}w''_{c,a,s,t})}{\sum_{c \neq c_0} (w'_{c,a,s,t}w''_{c,a,s,t})} \qquad \forall c \neq c_0 \cap R[c] \neq R[c_0] \cap SR[c] \neq SR[c_0]$$

Observations could be downweighted by a factor of 0.1, usually because they were not geographically representative at the unit of estimation. Details of reasons for downweighting can be found in risk-specific modeling summaries. The final weights were then normalized such that the sum of weights across age, time, and geographic hierarchy for a reference group was 1.

Estimating error variance

 σ_p^2 represents the error variance in normal or transformed space including sampling variance of the estimates and prediction error from any crosswalks performed. First, variance was systematically imputed if the data extraction did not include any measure of uncertainty. When some sample sizes for data were available, missing sample sizes were imputed as the 5th percentile of available sample sizes. Missing variances were then calculated as $\sigma_p^2 = \frac{p*(1-p)}{n}$ for proportions or were predicted from the mean using a regression for continuous values. When sample sizes were entirely missing and could not be imputed, the 95th percentile of available variances at the most granular geographic level (ie, first country, then region, etc.) were used to impute missing variances. For proportions where p*n or (1-p)*n is < 20, variance was replaced using the Wilson Interval Score method.

Next, if the exposure was modelled as a log transformation, the error variance was transformed into logspace using the delta method approximation as follows,

$$\sigma_p^2 \cong \frac{\sigma_{p\prime}^2}{p_{c,a,s,t}^2}$$

where σ_{pr}^2 represents the error variance in normal space. If the exposure was modelled as a logit transformation, the error variance was transformed into logit-space using the delta method approximation as follows,

$$\sigma_p^2 \cong \frac{\sigma_{p'}^2}{(p_{c,a,s,t} * (1 - p_{c,a,s,t}))^2}$$

Finally, prior to GPR, an approximation of non-sampling variance was added to the error variance. Calculations of non-sampling variance were performed on normal-space variances. Non-sampling variance was calculated as the variance of inverse-variance weighted residuals from the space-time estimate at a given location level hierarchy. If there were fewer than 10 data points at a given level of the location hierarchy the non-sampling variance was replaced with that of the next highest geography level with more than 10 data points.

Estimating the covariance function

The final input into GPR is the covariance function, which defines the shape and distribution of the trends. Here, we have chosen the Matern-Euclidian covariance function, which offers the flexibility to model a wide spectrum of trends with varying degrees of smoothness. The function is defined as follows:

$$M(t,t') = \sigma^2 \frac{2^{1-\nu}}{\Gamma(\nu)} \left(\frac{d(t,t')\sqrt{2\nu}}{l}\right)^{\nu} K_{\nu}\left(\frac{d(t,t')\sqrt{2\nu}}{l}\right)$$

where $d(\cdot)$ is a distance function; σ^2 , v, l, and K_v are hyperparameters of the covariance function specifically σ^2 is the marginal variance, v is the smoothness parameter that defines the differentiability of the function, l is the length scale, which roughly defines the distance between which two points become uncorrelated, and K_v is the Bessel function. We approximated σ^2 by taking the normalized median absolute deviation $MADN(r_c')$ of the difference which is the normalized absolute deviation of the difference of the first-stage linear regression estimate from the second-stage spatiotemporal smoothing step for each country. We then took the mean of these country-level MADN estimates for all countries with 10+ country-years of data, to ensure that differences between first- and second-stage estimates had sufficient data to truly convey meaningful information on model uncertainty. We used the parameter specifications v = 2 for all models. The scale parameter l used for each risk is reported in Appendix Section 4.

Prediction using GPR

We integrated over $g_{c,t}(t_*)$ to predict a full time series for country c, age a, sex s, and the prediction time t_* :

$$p_{c,a,s}(t_*) \sim N\left(m_{c,a,s,t}(t_*), \sigma_p^2 I + Cov\left(g_{c,a,s,t}(t_*)\right)\right)$$

Random draws of 1000 samples were obtained from the distributions above for every country for a given indicator. The final estimated mean for each country was the mean of the draws. In addition, 95% uncertainty intervals were calculated by taking the 2.5 and 97.5 percentile of the sample distribution. The linear modelling process was implemented using the Imer4 package in R, and the ST-GPR analysis was implemented through the PyMC2 package in Python.

Subnational Scaling and Aggregation

To ensure internal consistency of the estimates between countries and their respective subnational locations, national estimates were either created by population-weighted aggregation or subnational estimates were adjusted by population-weighted scaling to the national estimates, depending on the data coverage of a given country compared to that of its subnational locations. For example, if there was better data coverage at the national level, relative to its corresponding subnational locations, for a given country and risk across age, sex, and time, estimates were rescaled to be consistent with the national level. Conversely, if there was better data coverage at the oppulation-weighted aggregation of subnational estimates.

This GBD iteration, we incorporated an option to scale estimates within logit space. Scaling in logit space ensures that subnational estimates of proportion models will not exceed one after being rescaled to the national estimate.

Fitting a distribution to exposure data

The most informative data describing the distribution of risk factors within a population come from individual-level data; additional sources of data include reported means and variances. In cases when a risk factor also defines a disease or disease severity cut-off, such as haemoglobin level and mild, moderate, or severe anaemia or diabetes and fasting plasma glucose, the prevalence of disease is also frequently reported. To model the distribution of any particular risk factor, we seek a family of probability density functions (PDFs), a fitting method, and a model selection criterion. To make use of the most commonly available data describing most populations, we used the method of moments

(MoM); the first two empirical moments from a population, the mean and variance, were used to determine the parameters of two-parameter probability distribution families (PDF) describing the distribution of risk within any population. Exceptions to this rule are justified by context. We used the Kolmogorov-Smirnov¹⁵ (KS) test to measure the goodness of fit (GoF), comparing the distance between the empirical and ensemble distributions, but in some cases, the GoF was based on the prediction error for the prevalence of disease.

We used an ensemble technique in which a model selection algorithm is used to choose the best model for each continuous risk factor.¹⁵ We drew the initial set of candidate models from commonly used PDF families, including both right-skewed and left-skewed distributions. These included: beta, exponential, gamma, gumbel, inverse gamma, inverse Weibull, log-logistic, lognormal, mirrored gamma, mirrored gumbel, normal, and Weibull. We fitted each PDF candidate family to each dataset using the MoM, and used the KS test as the measure of GoF. Preliminary analysis showed that the GoF ranking of PDF families varied across datasets for any particular risk factor and that combining the predictions of differently fitted PDF families could dramatically improve the GoF for each dataset. Therefore, we developed a new model for prediction using the ensemble of candidate models, which is a weighted linear combination of all candidate models, $\{f\}$, where a set of weights $\{w\}$ is chosen such that $\sum_i w_i = 1$, and the values of the weights were determined by a second GoF criterion with its own validation process. For each risk, we pooled all available microdata and performed Nelder-Mead numeric optimisation across demographics subsets of data to derive a set of distribution-specific weights such that the average KS statistic across data sets is minimised. The details can be summarised by 1) the summary statistics for each dataset; 2) a table showing the KS statistic for each candidate model; and 3) the weights defining the final ensemble model for each dataset. We then averaged across demographic subsets and data sets to determine the final weights for modelling the distribution of any particular risk factor.

Section 2.4: Step 3. TMREL

In this and all previous GBD studies, the counterfactual level of risk exposure used is the risk exposure that is both theoretically possible and minimizes risk in the exposed population that consequently captures the maximum population attributable burden.² For each risk evaluated in GBD 2017, Step 4 of the analytical flowchart describes the use of the best available epidemiological evidence from published and unpublished relative risks by level of exposure and the lowest observed level of exposure from cohorts, used to select a single level of risk exposure that minimises risk from all causes of DALYs combined to establish the TMREL. In principle, the TMREL for a given risk may vary by age, sex, and location if supported by clear evidence. Based on the available evidence, the TMREL itself can be uncertain, which is reflected in the 95% uncertainty intervals (UIs) in Appendix Table 6. An estimation of uncertainty was derived by resampling from a uniform distribution of TMRELs where evidence supporting the selection of the TMREL was uncertain (for example, elevated systolic blood pressure or cholesterol).

Section 2.5: Step 4. Estimate population attributable fractions

Risks are categorised on the basis of how exposure was measured: dichotomous, polytomous, and continuous. High low-density lipoprotein (LDL) cholesterol is an example of a risk measured on a continuous scale. The population attributable fraction (PAF), which represents the proportion of risk

that would be reduced in a given year if the exposure to a risk factor in the past were reduced to an ideal exposure scenario, is defined for a continuous risk factor as:¹⁶

$$PAF_{joasgt} = \frac{\int_{x=l}^{u} RR_{joasg}(x)P_{jasgt}(x)dx - RR_{joasg}(TMREL_{jas})}{\int_{x=l}^{u} RR_{joasg}(x)P_{jasgt}(x)dx}$$

Where PAF_{joasgt} is the population attributable fraction for cause o due to risk factor j for age group a, sex s, location g, and year t. $RR_{joasg}(x)$ is the relative risk as a function of exposure level x for risk factor j for cause o, age group a, sex s, and location g with the lowest level of observed exposure as l and the highest as u; $P_{jasgt}(x)$ is the distribution of exposure at x for age group a, sex s, location g, and year t; $TMREL_{jas}$ is the TMREL for risk factor j, age group a, and sex s.

The *PAF_{ioasgt}* for dichotomous and polytomous risk factors for every country is defined as:

$$PAF_{joasgt} = \frac{\sum_{x=1}^{u} RR_{joast}(x)P_{jasgt}(x) - RR_{joasg}(TMRE_{jas})}{\sum_{x=1}^{u} RR_{joas}(x)P_{jasgt}(x)}$$

Where PAF_{joasgt} is the population attributable fraction for cause o due to risk factor j for age group a, sex s, location g, and year t. $RR_{joasg}(x)$ is the relative risk as a function of exposure level x for risk factor j for cause o, age group a, sex s, and location g on a plausible range of exposure levels from I to u $P_{jasgt}(x)$ is the proportion of population in risk group (prevalence), for age group a, sex s, location g, and year t; $TMREL_{ias}$ is the TMREL for risk factor j, age group a, and sex s.

Section 2.6: Step 5. Estimate summary exposure values

Summary exposure value (SEV), is the relative risk-weighted prevalence of exposure, a univariate measure of risk-weighted exposure, taking the value zero when no excess risk for a population exists and the value one when the population is at the highest level of risk. We report SEVs on a scale from 0% to 100% where a decline in SEV indicates reduced exposure to a given risk factor and an increase in SEV indicates increased exposure.

We first calculate risk, r, and cause, c, specific SEVs using the following equation,

$$SEV_{rc} = \frac{\frac{PAF_{rc}}{1 - PAF_{rc}}}{RR_{max} - 1}$$

for each most-detailed age, sex, location, year, and outcome. PAF is the YLL (expect for occupational noise, bullying victimization, and occupational ergonomic factors which are YLD only and thus use the YLD) PAF. RR_{max} for categorical risks is the RR at the highest category of exposure. For continuous risks, this is

$$RR_{max} = RR^{\frac{TMREL-1^{st}exposure}{RR_{scalar}}}$$
 if protective, or

$$= RR^{\frac{99^{th}exposure - TMREL}{RR_{scalar}}}$$

otherwise, and for custom modelled risks like ambient particulate matter pollution, household air pollution from solid fuels (HAP), alcohol, smoking, bullying, and activity, the modeller provides draws of RR_{max} . Generally, RRs do not vary across time and space, however, there are exceptions – an example being those risks such as second-hand smoke (SHS) or HAP where the RR is based on the integrated exposure response [IER], curve) and in these cases the RR is averaged across location and year to ensure no time or space variation. If the PAF is negative, signifying a protective effect for that outcome, the PAF is set to 0 and the SEV is then also 0 as the SEV univariate and constrained to be a value between 0 and 1. Once we have a set of risk cause specific SEVs at the most-detailed risk, cause, age, sex, location for all years, we average across causes to produce the final risk specific SEV_r ,

$$SEV_r = \frac{1}{N(c)} \sum_c SEV_{rc}$$

Section 2.7: Step 6. Mediation

Section 2.7.1: Summary

The portion of the burden of disease that is attributable to various combinations of risk factors or to all risk factors combined has been a topic of broad interest.¹⁷ Assumptions about how one risk factor is mediated through other risk factors are needed in order to estimate the joint risk factor burden for combinations of metabolic risks and behavioural or environmental risks. To accomplish this, in Step 6 of the estimation process, for every two risk factors for an outcome, we estimated the fraction of risk that was mediated through the other risk. This resulted in a matrix of parameters containing each possible pairing of risk factors included in the GBD 2017. Using this matrix, we computed the aggregated burden of disease at each level of the GBD 2017 hierarchy and for all risk factors using the following formula:

$$PAF_{Joasgt} = 1 - \prod_{j=1}^{J} \left(1 - PAF_{joasgt} \prod_{i=1}^{J} (1 - MF_{jio}) \right)$$
(5)

where J is a set of risk factors for the aggregation; PAF_{joasgt} is the PAF for risk j for age group a, sex s, location g, and year t; and MF_{jio} is the mediation factor for risk j mediated through i for cause o. Mediation factors can be found in Appendix Table 8.

Section 2.7.2: Additional detail

In GBD 2010, we only aggregated the burden of risk factors for some clusters of risks including access to improved water and sanitation, child and maternal malnutrition, tobacco smoking, alcohol use, dietary risk factors, occupational risk factors, and sexual abuse and violence. We did not aggregate air pollution and metabolic risk factors. For GBD 2013, GBD 2015, GBD 2016, and GBD 2017, we aggregated all risk factors into three large categories: behavioural, environmental and occupational, and metabolic risks -- as well as aggregating all GBD risk factors into a single attributable fraction for each diseases and eventually for all-causes of burden.

Aggregating risk factors at different levels share three essential challenges:
- 1. Risk factor coexistence or aggregation: for example, metabolic risk factors often occur together or high-risk behaviours are related such as drug abuse and unsafe sex.
- Mediation: a risk factor may effect another risk factor that lies in the physiological pathway to a disease outcome. It can be inside a cluster of risk factors such as the effect of obesity through an increase in fasting plasma glucose (FPG) and later cardiovascular disease outcomes, or between clusters of risk factors such as the effect of fibre on cholesterol.
- 3. The formula to calculate the aggregated PAF.

The aggregation method is conceptually applicable to other aggregations such as socioeconomic factors, education, homelessness and refugee status that are being considered for inclusion in future GBD iterations. In the next section, we explain our approach to deal with these challenges.

There are three patterns of associations between risk factors to take into consideration. The first concerns confounding; risk B affects risk A and outcome C (Pattern 1 in *Patterns of associations between risk factors*). In these cases, the relative risk (RR) for A should be adjusted for B, for example, the fruit RR is adjusted for smoking. If part of the effect of A is through B, a mediator, we do not adjust the effect of A for B. For example, we do not adjust the RR of body-mass index (BMI) for cholesterol as cholesterol lies in the biological pathway between BMI and cardiovascular outcomes (Pattern 2 in in *Patterns of associations between risk factors*). The third pattern occurs when risks A and B are proxies of a third variable Z and aggregation aims to estimate the total effect of a latent variable Z, on C. An example is child growth failure, which is measured by stunting, wasting, and underweight as proxies.



Pattern 1

Patterns of associations between risk factors

Section 2.7.3: Calculating burden of multiple risk factors

Validation studies have reported congruency between the true risk associated with multiple risk factors affecting the same outcome and a multiplicative aggregation of the population attributable fractions of the individual risk factors (formula below).¹⁸

$$PAF_{1..i} = 1 - \prod_{i=1}^{n} (1 - PAF_i)$$

Where *PAF* is the population attributable fraction and *i* is each individual risk factor. The same validation studies also found that the overestimation from ignoring the covariance between risk factors is small. This was important to note as there are few data sources from which we can draw information on covariance.

We endeavoured to evaluate RRs that were controlled for confounders. However, as we had to rely on the literature for many RRs we did not always have full control over the choice of confounders controlled for in each study.

Section 2.7.4: Adjusting for mediation

When aggregating the effects of multiple risk factors, we included a mediation factor if a part of the effect of one risk factor was included in the effect estimated for in the mediator. First, we prepared a list of possible mediations especially between behavioural risks and metabolic risk factors with cardiometabolic outcomes. We did not assume any mediation effect between risk factors for cancers.

Danaei and colleagues assumed that part of the effect of BMI on ischemic heart disease (IHD) is through high systolic blood pressure (SBP), cholesterol and FPG.¹⁹ The proportion of the BMI effect that can be explained by other metabolic risk factors is the amount of mediation. The difference between the crude RR of BMI on IHD with the RR adjusted for SBP, FPG, and cholesterol reflects the amount of BMI effect on IHD that is mediated and already included in SBP, FPG, and cholesterol:

$$MF = \frac{RR_{crude} - RR_{adjusted}}{RR_{crude} - 1}$$

We used this approach for estimating mediation factors to adjust PAFs before aggregation.

$$MF = \frac{R_c^+ - R_a^+}{R_c^+ - R_c^-}$$

So:
$$R_a^+ = R_c^+ - MF * (R_c^+ - R_c^-)$$

$$PAF_{c} = \frac{p * (R_{c}^{+} - R_{c}^{-})}{p * R_{c}^{+} + (1 - p) * R_{c}^{-}} = \frac{p * (R_{c}^{+} - R_{c}^{-})}{R_{T}}$$

If R_c^+ : crude risk of outcome in exposed population

 R_c^- : crude risk of outcome in non-exposed population

 R_a^+ : adjusted risk of outcome in exposed population

 R_a^- : adjusted risk of outcome in non-exposed population

 R_T is the overall rate of the outcome in the population. Since we are interested in the part which is from BMI but through cholesterol, the total risk in the population will be the same for the adjusted RR, so the unmediated part of the risk factor would be:

$$PAF_{a} = \frac{p*(R_{a}^{+} - R_{a}^{-})}{R_{T}} = \frac{p*(R_{c}^{+} - MF*(R_{c}^{+} - R_{c}^{-}) - R_{c}^{-})}{R_{T}} = \frac{p*(R_{c}^{+} - R_{c}^{-})*(1 - MF)}{R_{T}} = PAF_{c} * (1 - MF)$$

So for aggregating the PAF of multiple risk factors, we first calculated the part of the effect of every risk factor that is not mediated and then aggregated these assuming they are independent.

Therefore the aggregated PAF would be:

If MF is mediation factor of R2 through R1:

$$PAF_{1,2} = 1 - (1 - PAF_1) * (1 - PAF_2 * (1 - MF_{2/1}))$$

and a generalization for multiple pathways of R1 through other RFs:

$$PAF_{1..i} = 1 - \prod_{i=1}^{n} \left(1 - PAF_i * \left(1 - \prod_{j=1}^{n} (1 - MF_{i/j}) \right) \right)$$

For every risk factor outcome pair, the matrix of possible mediations was calculated and used.

Section 2.7.5: Calculating mediation factor

1 – Comparing crude RR versus mediator-adjusted RR

The best example is the mediation of BMI through SBP, FPG, and cholesterol reported by Danaei et al.¹⁹ In their meta-analysis, they report the adjusted and unadjusted RR of BMI on IHD and stroke based on combined data from individual cohorts. They calculated the mediation factor using the equation below, and we used it directly as mediation factor in risk factor aggregation. Using individual level data from cohort studies, we estimated the mediation factor for other metabolic risk factors and some dietary risks.

$$MF = \frac{RR_{crude} - RR_{adjusted}}{RR_{crude} - 1}$$

2 – Estimating the mediation factor by pathway of the effect

For many other risk factors, there are no data available to use the first method. Instead, we searched studies to estimate the effect of the risk factor on the mediator and finally the expected increase in IHD risk. We pooled available studies to calculate the unit increase in the mediator per unit increase in the risk factor to calculate the size of the IHD RR.



Example of pathway between fruit, high systolic blood pressure, and cardiovascular diseases

We have RRs for the effect of A on C and B on C in GBD from a meta-analysis of studies in the literature. The effect of A on B was estimated by analysis of diet trials.

$$RR_{ABC} = RR_{BC}^{\Delta_{AB}}$$

 RR_{ABC} is expected effect of A through B on C

 RR_{BC} is relative risk of each unit increase in mediator on outcome C

 Δ_{AB} is change in mediator level B per each unit change in A

If RR_{AB} is the overall effect of A on B then:

The mediation factor would be

$$MF = \frac{RR_{ABC} - 1}{RR_{AB} - 1}$$

We kept uncertainty of each parameter by generating and following 1000 draws of the estimates to calculate 1000 draws of the posterior distribution of the mediation factor. We did not include risk-mediator pairs if the mediation factor was not significant at 5% level (more than 50 out of 1000 draws were negative). We truncated the mediation factor distribution at 1 where the whole effect of the risk factor on the outcome would be assumed to be through the mediator pathway.

Some mediation factors equal 1 where the whole effect was calculated through other risk factor, e.g. the effect of salt through SBP, or when we assumed other risk factors are sources of the exposure, for example, fibre is provided by consuming fruit, vegetable, and whole grains and all the beneficial effect of milk on colorectal cancer is mediated through calcium.

Dietary risk factors

For each dietary risk factor, we searched for randomised trials evaluating the effect of the diet component on metabolic risk factors and estimated the change in a given mediator per unit change in the diet component. Only salt outcomes, with the exception of stomach cancer, are mediated through BMI – all other dietary risk factors have their relative risk applied directly.

Physical activity

We found cohort studies on the effect of physical activity on FPG. The data was more on the effect of physical activity on diabetes incidence, so we calculated the shift in FPG using the provided RR value. We used this to calculate the mediated part of effect of physical activity on cardiovascular disease (CVD).^{20–}

Air pollution

We considered mediation for particulate matter pollution, but the evidence was not strong enough to justify inclusion of PM_{2.5} on SBP, FPG, or cholesterol. There are two cohort studies which have published findings of increased risk of hypertension due to long-term exposure to ambient PM_{2.5} and several studies which have found elevated SPB due to household solid fuel use. We found time series studies with different PM_{2.5} lag (by day) that show very short-term and confounded effects. We decided to re-examine these outcomes when more evidence becomes available.

Assumed mediations

For the risk factors with PAFs of 100% such as FPG and diabetes, impaired kidney function and chronic kidney disease, SBP and hypertensive heart disease, alcohol and alcohol use disorders, child underweight and protein-energy malnutrition, and child wasting and protein-energy malnutrition, and drug use and drug use disorders, no mediation is needed.

Section 2.7.6: Piecewise aggregation (Pattern 3)

There are three anthropometric indicators that are highly correlated: child underweight, stunting, and wasting, as demonstrated in *Venn diagram demonstrating the correlation between child underweight, stunting, and wasting*. Available RRs for each indicator are not adjusted for the other two because there is a high correlation between these indicators and also interaction where the majority of the burden occurs. Estimating the total burden due to childe growth failure, a latent variable, is difficult. The three anthropometric indicators are not independent, so the covariance between them should be considered. This was the main reason that GBD 2010 only included child underweight. If covariance between these indicators is significant (as is shown in the Figure below), aggregating these indicators assuming independence would overestimate the total burden significantly.

To use the best available data, we adjusted observed RRs reported by Olofin et al for underweight, stunting and wasting by simulating the joint distribution of the three indicators using the distribution of each indicator and covariance between indicators in the countries included in the meta-analysis (extracted from Demographic and Health Survey (DHS) micro-data).²⁷ Based on the analysis done by McDonald et al, we assumed there is an interaction between the three indicators, and extracted the interaction terms from the corresponding analysis.²⁸ We calculated the adjusted RRs by minimizing the error between observed crude RRs (from meta-analysis) and expected crude RRs derived from adjusted RRs.



Venn diagram demonstrating the correlation between child underweight, stunting, and wasting

After adjusting for the three risk factors, we calculated the PAFs and aggregated underweight, stunting and wasting burden.

Section 2.7.7: Uncertainty of aggregated and mediated PAFs

We generated 1000 draws of posterior distribution of mediation factor calculated by different methods to use beside draws of other inputs to the PAF aggregation.

Section 2.7.8: Important assumptions in aggregating risk factors and including mediation

1 – The mediation factors or PAF adjustments are similar across countries, age, sex, and years. While it is quite likely that the size of mediation is different in different populations, there is little data to inform the covariance between different risk factors or the mediation factor amount by age and countries. For example, in some countries, the size of the mediated BMI-IHD PAF through cholesterol, calculated by the mediation factor, was even bigger than the total burden of cholesterol, indicating that less effect of BMI is mediated through cholesterol and mediation factors are not similar across countries.

2 – For many risk-mediator-outcome pairs, there are no data available, so we assumed the mediation is zero.

3 – Since the covariance between undernutrition indicators is different by location (and across time, results were not reported), and there is an interaction between these indicators, the total burden might be underestimated.

4 – It is assumed that there is no significant covariance between PAFs, which might not be true between some risk factors such as between metabolic risk factors. While this overestimation is controlled by using adjusted RRs, using crude RRs for BMI and other metabolic risk factors may cause significant overestimation of aggregated metabolic risks burden.

Section 2.8: Step 7. Estimate attributable burden

Four key components are included in estimation of the burden attributable to a given risk factor: the metric of burden being assessed (the number of deaths, years of life lost [YLLs], years lived with disability [YLDs], or DALYs [the sum of YLLs and YLDs]); the exposure levels for a risk factor; the relative

risk of a given outcome due to exposure; and the counterfactual level of risk factor exposure. Estimates of attributable burden as DALYs for risk-outcome pairs were generated using the following model:

$$AB_{jasgt} = \sum_{o=1}^{w} DALY_{joasgt} PAF_{joasgt}$$

where AB_{jasgt} is the attributable burden for risk factor j for age group a, sex s, location g, and year t; $DALY_{joasgt}$ is total DALYs for cause o (of w relevant outcomes for risk factor j) for age group a, sex s, location g, and year t; PAF_{joasgt} is the population attributable fraction (PAF) for cause o due to risk factor j for age group a, sex s, location g, and year t. The proportion of deaths, YLLs, or YLDs attributable to a given risk factor or risk factor cluster were analogously computed by sequentially substituting each metric in place of DALYs in the equation above.

Section 2.9: Decomposition analysis of deaths and DALYs

We conducted a decomposition analysis of changes in DALYs from 2007 to 2017, decomposing changes in all-age cause-specific DALYs attributable to all risk factors and individual risk factors due to changes in population growth, population age structure, exposure to the given risk for a disease, and risk-deleted death and DALY rates. In this case, risk-deleted rates are the rates after removing the effect of a risk factor or combination of risk factors; in other words, observed DALY rates multiplied by one minus the PAF for the risk or set of risks. Our decomposition analyses draw from methods developed by Das Gupta²⁹ to provide a computationally tractable solution to isolating drivers of burden changes whereby all combinations of possible pathways are averaged across factors. Attributable burden is determined, following the methods of Das Gupta, as a product of three factors such that:

$$T_{asgt} = (A_{asgt} B_{asgt} C_{asgt})$$

where T_{asgt} represents the attributable burden at year t; A_{sgt} is the age-specific population size for a given age group a, sex s and location g at year t; B_{asgt} is the underlying rate of the outcome unrelated to the risk factor or observed rate, multiplied by 1 - PAF for a given age group a, sex s and location g at year t; and where C_{asgt} is the ratio of attributable burden to the underlying rate, which reflects the risk exposure effect for a given age group a, sex s, and location g at year t defined as PAF/(1 - PAF) in the case of decomposing attributable burden to a risk. Risk exposure effects for individual risk factors are scaled such that they sum to the all risk exposure effect by location, age, sex, and cause accounting for mediation. This allows for aggregation of risks; the exposure for all risks for a disease can be split into exposure to metabolic, behavioural, and environmental risks. The contribution of each factor to total change in attributable burden was determined by changing the level of one factor from time t_0 to t_1 – here 2007 to 2017 – with all other factors held constant. Thus, the effect of any of the three factors, for example A_{asat} on the change of attributable burden between 2007 (A_{07}) and 2017 (A_{17}) is calculated as:

$$E_A = (A_{17} - A_{07}) \left(\frac{B_{07}C_{07} + B_{17}C_{17}}{3} + \frac{B_{07}C_{17} + B_{17}C_{07}}{6} \right)$$

Where E_A is the proportion of change due to factor A, and the subscripts for each factor in the equation denote the year for each estimate. Since the effect depends on the order of entry of the factor, we calculated the average of all combinations of the three factors.²⁹ The proportion of change due to factor A_{sqt} , the age-specific population size for a given age group a, sex s and location g at year t, is

then further split, setting change in population growth equal to the percent change in all-age population from time t_0 to t_1 and change in population age structure to the residual, giving four factors.

This three factor decomposition method does not work for risks where the PAF, by definition, is 100% (such as high fasting plasma glucose and diabetes type 2) or where the PAF is directly estimated (such as for unsafe sex and HIV). In the cases of child underweight and protein-energy malnutrition, child wasting and protein-energy malnutrition, short gestation for birth weight and neonatal preterm birth complications, low birth weight for gestation and neonatal preterm birth complications, iron deficiency and iron-deficiency anaemia, alcohol use and liver cancer due to alcohol use, alcohol use and cirrhosis and other chronic liver diseases due to alcohol use, alcohol use and alcohol use disorders, alcohol use and alcoholic cardiomyopathy, drug use and drug use disorders, occupational particulate matter, gases, and fumes and other pneumoconiosis, occupational particulate matter, gases, and fumes and coal workers pneumoconiosis, occupational exposure to asbestos and asbestosis, and occupational exposure to silica and silicosis, we used a two factor decomposition method, which examines the contribution of population, ageing, and risk exposure. Effectively, we assume trends in these cases are driven by exposure, not change in the risk-deleted rates. Conversely, for unsafe sex and sexually transmitted diseases excluding HIV, we used a two-factor decomposition method, which examines the contribution of population, ageing, and risk-deleted death and DALY rates, assuming trends in these cases are driven by risk-deleted rates, not change in exposure. For high fasting plasma glucose and diabetes mellitus type 1 and 2, high fasting plasma glucose and chronic kidney disease due to diabetes mellitus type 1 and 2, high systolic blood pressure and hypertensive heart disease, high systolic blood pressure and chronic kidney disease due to hypertension, and impaired kidney function and chronic kidney disease, we used GBD estimates of Summary Exposure Values (SEVs) for the given risk and the case-fatality rate decompose trends into the contribution of the three factors. Similarly, for unsafe sex and cervical cancer, we used GBD estimates of the incidence of cervical cancer and the case-fatality rate to decompose trends into the contribution of the three factors. For unsafe sex and HIV, we used spectrum counterfactual and CD4 risk-weighted prevalence.

Section 2.10: SDI Analysis

Section 2.10.1: Development of SDI

The Socio-demographic Index (SDI) is a composite indicator of development status strongly correlated with health outcomes. In short, it is the geometric mean of 0 to 1 indices of total fertility rate under the age of 25 (TFU25), mean education for those aged 15 and older (EDU15+), and lag distributed income (LDI) per capita.

Section 2.10.2: Development of revised SDI indicator

SDI was originally constructed for GBD 2015 using the Human Development Index (HDI) methodology, wherein a 0 to 1 index value was determined for each of the original three covariate inputs (total fertility rate in ages 15 to 49, EDU15+, and LDI per capita) using the observed minima and maxima over the estimation period to set the scales.

In response to feedback from collaborators and the evolution of the GBD, we have refined the indicator with each GBD cycle. For GBD 2017, in conjunction with our expanded estimation of age-specific fertility, we chose to replace the total fertility rate as one of the three component indices with the total fertility rate under 25 (TFU25). The TFU25 provides a better measure of women's status in society, as it focuses on ages where childbearing disrupts the pursuit of education and entrance into the workforce.

During GBD 2016, we moved from using relative index scales to absolute scales to enhance the stability of SDI's interpretation over time, as we noticed that the measure was highly sensitive to the addition of subnational units that tended to stretch the empirical minima and maxima. We selected the minima and maxima of the scales by examining the relationships each of the inputs had with life expectancy at birth and under-5 mortality and identifying points of limiting returns at both high and low values, if they occurred prior to theoretical limits (e.g., a TFU25 of 0).

Thus, an index score of 0 represents the minimum level of each covariate input past which selected health outcomes can get no worse, while an index score of 1 represents the maximum level of each covariate input past which selected health outcomes cease to improve. As a composite, a location with an SDI of 0 would have a theoretical minimum level of development relevant to these health outcomes, while a location with an SDI of 1 would have a theoretical maximum level of development relevant to these health outcomes.

We summarize the final scales for GBD 2017 in the table below.

Input	Lower Bound	Upper Bound
TFU25	0	3
LDI per capita	250 USD (5.5 log USD) ^b	60,000 USD (11.0 log USD)
EDU15+	0 years	17 years

^b The minimum for the LDI scale was originally set at the theoretical limit of 0 USD, as we did not observe an asymptotic relationship between log(LDI) and E₀ or 5q0 at lower values of log(LDI). Empirically, however, we also did not observe an LDI below 350 USD (5.86 log USD) for the estimation period 1970-2016. In log-space, this meant that approximately half of our scale was not being utilized, compressing the observed variation in LDI and diminishing its meaningful contribution to SDI. Accordingly, we set the lower limit on LDI to 250 USD (5.52 log USD) to ensure we were fully utilizing the range of the scale to capture its variation across space and time, as is the case with the other two inputs.

Using scales described above, we computed the index scores underlying SDI as follows:

$$I_{Cly} = \frac{(C_{ly} - C_{low})}{(C_{high} - C_{low})}$$

Where I_{cly} – the index for covariate *C*, location *I*, and year *y* – is equal to the difference between the value of that covariate in that location-year and the lower bound of the covariate divided by the difference between the upper and lower bounds for that covariate. If the values of input covariates fell outside the upper or lower bounds (e.g. LDI per capita greater than 60,000 USD), they were mapped to the respective upper or lower bounds. We also note that the index value for TFU25 was computed as $1 - I_{TFU25ly}$, as lower TFU25s correspond to higher levels of development, and thus higher index scores. For GBD 2017 we expanded the computation of SDI to 890 national and subnational locations spanning the time period 1950-2017.

The composite Socio-Demographic Index is the geometric mean of these three indices for a given locationyear. The cut-off values used to determine quintiles for analysis were then computed using country-level estimates of SDI for the year 2017, excluding countries with populations less than 1 million.

Example calculation

Below we present an example calculation of SDI for "Country X":

$$TFU25 = 1.09$$
; Mean educ yrs $pc = 8.23$; $lnLDI = 9.60$

$$I_{TFU25} = 1 - \frac{1.09 - 0}{3 - 0} = .637$$

$$I_{Educ} = \frac{8.23 - 0}{17 - 0} = .484$$

$$I_{lnLDI} = \frac{9.60 - 5.52}{11.00 - 5.52} = .744$$

$$SDI = \sqrt[3]{I_{TFU25} * I_{Educ} * I_{lnLDI}} = \sqrt[3]{.637 * .484 * .744} = .611$$

$$I_{lnLDI} = \frac{9.58 - 5.52}{11.00 - 5.52} = .741$$

$$SDI = \sqrt[3]{I_{TFR} * I_{Educ} * I_{lnLDI}} = \sqrt[3]{.855 * .543 * .741} = .701$$

SDI grouping by location can be found in Appendix Table 9, and SDI values can be found in Appendix Table 10.

Section 2.11: Epidemiological Transition

Section 2.11.1:Derivation of expected SEVs

In order to evaluate the average relationship between SDI attributable burden, we first quantified that between SDI and population-level exposure. Using the "gam" package in R, we fit a generalized additive model (GAM) with a loess smoother on SDI by age and sex group. For age *a*, sex *s*, and risk *r*.

$$logit(SEV_{asr}) = \beta_0 + s(SDI) + \epsilon$$

Inputs to this model were age-sex specific SEVs for all most-detailed risks in the GBD risk hierarchy corresponding to all national GBD locations and years between 1990 and 2017. The span for the loess term was set at a default of .7, and the model was fit on the middle 95% of the data to mitigate the impact of compositional bias evident in outliers very close to zero or one. Expected age and sex proportions of the population on the basis of SDI produced through an analogous modelling framework were used to generate age and sex aggregates of expected exposure.

Section 2.11.2:Calculating attributable burden

Borrowing from forecasting methods, we generated an estimated expected risk-specific PAF by backtransforming the expected SEV to PAF, rearranging the formula for SEV described in further detail in the appendix to solve to PAF for a given age a, sex s, cause c, risk r, and SDI d,

$$PAF_{ascrd} = 1 - \frac{1}{SEV_{asrd} \times (RR_{ascr}^{max} - 1) + 1}$$

As PAF estimates are derived directly from the SEV which is not cause specific, but averaged across causes, we then calculated a correction factor CF by comparing in logit space the empirical PAF to the SEV derived estimate PAF,

$$CF_{ascr} = logit(PAF_{ascrly}) - logit(PAF_{ascrly})$$

and apply the correction factor to the estimated expected PAF to derived an adjusted expected estimated PAF

$$PAF_{ascrd} = expit(logit(PAF_{ascrd}) + CF_{ascr})$$

for all most-detailed risks in the hierarchy, excluding unsafe sex and occupational injuries, by age, sex, cause, and SDI, adding in risk-outcome

To estimate expected risk-attributable burden, we draw from the CRA methods (see Appendix Section 2.6), first calculating the joint adjusted expected *PAF* for all risks for a cause using mediation factors. We then draw from the methods for observed risk-attributable burden calculation, using expected YLLs, deaths, and YLDs (see Appendix Section 2.7) to generate expected burden for a given SDI, not location year.

$$AB_{rascd} = \sum DALY_{ascd} PAF_{rascd}$$

where AB_{rascd} is the expected attributable burden; $DALY_{ascd}$ is expected total; PAF_{ascrd} is the expected PAF. The proportion of expected deaths, YLLs, or YLDs attributable to a given risk factor or risk factor cluster were analogously computed by sequentially substituting each measure in place of expected DALYs in the equation above. Comparisons of observed to expected attributable burden were made to identify locations exhibiting exceptional deviations relative to what would be expected based on their development status.

Section 2.12: Additional Methods Information

Section 2.12.1: Risk-specific comparisons to other estimates

Low birth weight / Short Gestation:

GBD 2017 estimates of total preterm birth prevalence are generally in line with country-specific reports³⁰ as well as the most recent global analysis completed by Blencowe and colleagues.³¹ GBD 2017 estimates of preterm birth prevalence in 2010 are very similar, 11.3% (95% UI: 11.1% to 11.5%) versus 11.1% of live births, compared to estimates by Blencowe and colleagues. Close agreement is not surprising as most of the same data sources were used as data inputs to our modelling process, although the GBD analysis included almost eight times as many data points. Most reports, like GBD 2017, have assessed temporal trends in preterm birth in many locations to be either static or increasing. Compared to UNICEF estimates of low birth weight,³² GBD 2017 estimates of global birth prevalence of 14.4% (14.0% to 14.8%) are similar but slightly lower than the estimate of 15.5% birth prevalence globally. The geographic variation in low birth weight largely mirrors that of the UNICEF report.

Chewing Tobacco

In GBD 2016, we estimated age-sex specific and aggregate current smokeless tobacco use prevalence for all countries and territories from 1990-2016 using all available data. The estimated prevalence was then

attributed to either all chewing tobacco, or all snus/snuff by country, based on input from smokeless tobacco experts.

For GBD 2017, we have changed the exposure definition from current smokeless tobacco use to current chewing tobacco use, based on the strength of evidence supporting the health effects of chewing tobacco use. By estimating chewing tobacco exposure in all countries and territories, burden is now estimated for locations previously classified as predominantly snus/snuff, but still have non-negligible use of chewing tobacco.

We compared GBD estimates to recent research by Siddiqi et al.³³ on smokeless tobacco prevalence, risk, and burden in 2015. Our methods differed from Siddiqi on some key points:

Prevalence estimation: We estimated both age-sex specific and aggregate current smokeless tobacco use prevalence for all countries and territories included in GBD from 1990-2017 using all available data. Siddiqi et al. used only the single most recent survey available containing data on smokeless tobacco use among adults.

Relative risks and attributable burden: GBD excluded hospital-based case-control studies, while Siddiqi included them. GBD calculated relative risks for chewing tobacco (and by sex for oral cancer only), while Siddiqi calculated separate relative risks by geography, and then pooled these to produce global relative risks. Siddiqi used country- or region-specific relative risks where available, and in the absence of region-specific relative risks assigned global relative risks in countries predominantly using products with moderate to high pH and tobacco-specific nitrosamine (TSNA) levels.

The main differences in attributable burden come from the relative risk exclusion criteria. GBD's exclusion of hospital-based led to very different relative risk outputs: we found significant relative risks for oral cancer and oesophageal cancer. Siddiqi found significant relative risks for oral, pharyngeal, and oesophageal cancers and ischemic heart disease, resulting in higher levels of global burden.

Smoking

We compared GBD estimates to the most recent report on the global tobacco epidemic published by WHO³⁰. Overall, we found marked similarities in estimates. Among the 142 countries and territories included in the WHO report and estimated in GBD, the correlation coefficient for current smoking prevalence estimates among females was 0.91 and among males was 0.85. In cases where estimates diverge, discordance can be attributed to differing modelling methods or data sources. GBD uses ST-GPR to estimate smoking prevalence, whereas WHO uses Bayesian meta-regression (DisMod MR). Additionally, the WHO model was fit on 1,175 country-year data sources, whereas the GBD model was fit on 2,870 country-year data sources. There are no comparable global estimates of the burden of disease attributable to smoking, as GBD 2015 estimates of attributable burden were used in the most recent WHO report.

Ambient air pollution

In the past few years, other researchers have estimated the burden of disease due to air pollution using different data and methods. Other sources of estimates have been compared to those of the GBD. Since their introduction in GBD 2010, satellite-based estimates of PM_{2.5} and the Integrated Exposure Response (IER), have been widely adopted. Recent estimates from WHO³¹ of 3·0 million deaths in 2012 used a

similar exposure estimate model as that presented here, but an earlier (GBD 2013) version of the integrated exposure response (IER) and somewhat different baseline disease burden estimates. Lelieveld and colleagues³² analysed source sector contributions to air pollution and the resulting disease burden in 2010 and estimated the burden in 2050. These estimates used an older (GBD 2010) IER. Furthermore, the coarse spatial resolution (~100 × 100 km) of the exposure estimates introduced errors via spatial misalignment between exposure and population density compared with our estimates. Silva et al. estimated 2.23 million deaths/year due to anthropogenic PM_{2.5} and 493 thousand deaths per year due to ozone,³³ while Butt et al. estimated 12.4% global attributable deaths due to PM_{2.5} in the year 2009,³⁴ both using versions of the IER.

In GBD 2017, we added Type 2 diabetes as an outcome of ambient particulate matter pollution estimating 184 thousand attributable deaths and 10.5 million attributable DALYs globally in the year 2017. An independent group of researchers recently published work estimating the 2016 burden of diabetes attributable to ambient PM_{2.5}.³⁵ Using GBD 2015 PM_{2.5} exposure estimates, GBD 2016 diabetes burden, and generating their own IER the team estimated 206 thousand attributable deaths and 8.23 million attributable DALYs globally for the year 2016.

Occupational

Takala and colleagues³⁶ reported 2.3 million deaths attributable to occupational injury/illness in 2011. In the closest comparison year of 2007, GBD estimated nearly 1.1 million deaths. This discrepancy is largely driven by the cause-outcome pairs that GBD currently has the evidence to include based on the criteria of the CRA framework. For example, 45% of Takala's reported burden is driven by occupational circulatory disease (35%) and occupational communicable disease (10%). Circulatory diseases are linked to occupational risks like shift work and lack of control, but the GBD approach currently has insufficient evidence to estimate the variability in exposure to these factors on a global scale. Additionally, the use of a CRA approach in GBD estimates requires careful consideration of proposed counterfactual in order to derive the TMREL for a given risk. The TMREL for something like occupational lack of control is a challenging concept, and as such, these risks are still being reviewed for possible inclusion in future iterations of the GBD.

Takala also reports higher burden from occupational cancer based on the inclusion of carcinogens that are currently still out of the scope of GBD. For example, the authors use attributable fractions derived from Rushton and colleagues to attribute pairs like breast cancer and shift work or skin cancer and solar radiation. These carcinogens, which form a large part of the cancer burden in Takala/Rushton are currently not included in the GBD based on limited exposure data across the time/space that GBD estimates³⁸.

In terms of fatal occupational injuries, Takala reported 353 000 deaths in 2011. The GBD 2017 estimate for deaths attributable to occupation was approximately 348 000 deaths for 2007. The figures are very similar but the GBD estimates are slightly lower, again due to the selection of risk-cause pairs. The ILO estimation strategy includes some kinds of injuries, such as deaths due to intentional violence that the GBD does not attribute to occupation.

Child growth failure (stunting, wasting, and underweight)

UNICEF et al estimate lower proportion of stunting (height-for-age z-score < -2 standard deviations below the reference median) in children under five in 2017 than GBD 2017.³⁷ The geographic patterns generally agree in identifying sub-Saharan Africa and South Asia as the regions with the largest burden of stunting (prevalence and magnitude, estimated as number of stunted children in UNICEF et al, and as DALYs in GBD 2017), with additional high prevalence in Oceania (excluding Australia and New Zealand) and moderate prevalence in Latin America and the Caribbean. While UNICEF et al estimates highlight minimal or lack of progress in reducing stunting since 2000 in Africa and Oceania, GBD 2017 estimates show moderate decline in sub-Saharan Africa and North Africa and the Middle East, and a small decline in Oceania (compared to UNICEF et al's rise in stunting prevalence in Oceania). GBD 2017 estimates show a downward trend in the prevalence of underweight (weight-for-age z-score < -2 standard deviations below the reference median) among children under 5 in 2016, driven largely by populations in sub-Saharan Africa and South Asia, a trend also reflected in the UNICEF et al estimates.

Impaired Kidney Function

Recently published estimates from a meta-analysis of global data on exposure to impaired kidney function indicate prevalence of chronic kidney disease (CKD) stages 1-5 to be 13.4% (11.7-15.1%). (ref) These estimates are similar to the current GBD 2017 exposure estimates across all four levels of impaired kidney function, which indicate a prevalence for individuals over the age of 25 of 14.0% (13.0-15.1%).

Household Air Pollution

WHO estimated 4.3 million deaths and 146.5 million DALYs attributable to exposure to household air pollution globally in 2012, as compared to GBD 2017 estimates for the year 2017 of 1.6 million deaths and 59 million DALYs. Differences in attributable burden arise between the WHO estimates and GBD 2016 for a number of reasons. First, the IER curve was used for all outcomes (LRI, IHD, cerebrovascular stroke, COPD, Type II Diabetes, and lung cancer) except cataracts in our analysis, while WHO adapted relative risks for COPD based on a meta-analysis of published RR estimates. The excess relative risks for COPD used by WHO are larger than the excess relative risks used in GBD, resulting in a larger PAF. Additionally, this was the first year we included Type II Diabetes as an outcome of particulate matter pollution, including household exposure to solid fuel. WHO has not included this as an outcome in their estimates. Finally, WHO only estimates LRI deaths in children aged 0-4, while we estimate respiratory infections and pneumonia at all ages.

Additionally, by adapting the new proportional PAF approach for particulate matter pollution, beginning in GBD 2017 we are taking into account ambient pollution in the counterfactual of household. This has reduced burden estimates from previously published numbers, which we believe more accurately reflects the exposure-outcome relationship.

Another difference from the WHO estimates is the use of a database which maps solid cooking fuel use to $PM_{2.5}$ exposure, allowing us to model differences in exposure level and relative risk by location, while WHO relies on global relative risks. We also adjust this household $PM_{2.5}$ exposure by ambient air pollution levels since personal $PM_{2.5}$ exposure captures all sources of exposure. WHO makes no

adjustments for ambient air pollution exposure. In addition to the differences in data sources, we estimated the burden of cataract attributable to household air pollution (HAP) only in women while WHO estimated for both sexes.

Below is a comparison between the WHO³⁸ and GBD 2017 of the number of DALYs attributed to HAP contributed by each cause (GBD 2017 on right):

•	48,500,000-Pneumonia/LRI	25,400,000-Lower respiratory infections
•	36,900,000-Stroke	8,010,000-stroke
•	30,500,000-Ischaemic heart disease	9,170,000-IHD
•	22,100,000-COPD	11,700,000-COPD
•	6,720,000-Lung cancer	1,780,000-Lung cancer
•	1,670,000-Cataract	1,230,000-Cataract
•	0-Type 2 diabetes	830,000-Type 2 diabetes

Breastfeeding

Globally, WHO estimates that 40% of infants under six months of age are exclusively breastfed, which is consistent with the GBD 2017 estimate (40.3%). A recent WHO study estimates 73.3% continued breastfeeding at one year of age globally, where the GBD 2017 estimate of continued breastfeeding between 6 and 11 months is 88% (we expect our estimates to be higher, as our estimates measure breastfeeding for any infants age 6-11 months, while the WHO estimates focus on breastfeeding status at the age of 1 year).³⁹ CDC reports 25% of mothers in the United States exclusively breastfeed their child through 6 months of age, while GBD 2017 estimates focusing on all infants 0-5 months are exclusively breastfeed (this difference is again explained by our estimates focusing on all infants 0-5 months, while the CDC reports infants who complete 6 full months of exclusive breastfeeding). Additionally, CDC reports 34% of mothers in the U.S. continue to breastfeed at 12 months of age. GBD estimates 44% of mothers in the U.S. continue to breastfeed from 6 to 11 months of age. Finally, in India, WHO and UNICEF estimate 55% exclusive breastfeeding in the first six months of life, comparable to the GBD 2017 estimate of 55%.

WaSH

The Joint Monitoring Project⁴⁰ (JMP), which is led by WHO and UNICEF, estimates water, sanitation, and handwashing access throughout the world. Globally, JMP estimates that 91% of population had access to an improved water source in 2015, while GBD estimates 88% of the population have access to improved water. Additionally, JMP reported the global prevalence of households with piped water connection to be 57% in 2015, while GBD reports piped prevalence of 53% for that year. JMP reported 68% of population had access to improved sanitation in 2015, whereas GBD estimates improved sanitation prevalence of 74%. The slight discrepancies in these estimates at the global level can be largely attributed to differences in input data. The JMP relies almost exclusively on large-scale household surveys (DHS and Multiple Indicator Cluster Surveys [MICS]), while GBD estimates incorporate exposure data from smaller, yet still nationally representative, survey series such as Reproductive Health Survey and various country specific surveys. Due to the relative dearth of data regarding access to a handwashing facility, the JMP only generates handwashing estimates for a select

number of countries (mostly sub-Saharan Africa) where those data are actually collected. However, we model and predict handwashing facility prevalence for all locations, even in the absence of data, and estimates that 67% of the globe has access to a handwashing facility.

Lead

The most recent external estimates for the burden of lead exposure were conducted by WHO in 2004 and provided disaggregated average exposures for children and adults in different regions of the world.⁴¹ The GBD 2017 exposure estimates for the early 2000s are generally higher than these estimates, as we have observed higher estimates of exposure in GBD 2017 with changes in model covariates and covariate effects. The WHO study calculated a global burden attributable to lead exposure of 13 million DALYs, which includes 229 000 deaths. GBD 2017 estimates global burden attributable to lead exposure to be 24.4 million DALYs, including approximately 1.05 million deaths.

In addition to differences in overall attributable burden, the breakdown of burden from intellectual disability and cardiovascular disease is very different. Both attribute 2% to 3% of global CVD to lead exposure, but our estimates of CVD burden in 2007 are higher than WHO's (such that we estimate 18.1 million DALYs from CVD due to lead in 2007 compared to their estimate of 3.1 million). Additionally, our methodology for intellectual disability differs substantially from theirs. In the WHO study, they used a higher disability weight of 0.361 for intellectual disability, whereas we currently use weights ranging from 0.01 to 0.2 (depending on the severity). However, WHO's estimates of intelligence quotient (IQ) shift from lead exposure are much lower than ours, since recent studies have provided better evidence for notable effects of lead on IQ at low levels of exposure. Still, due to differences in our estimates of the underlying burden of intellectual disability, our estimate of 2.67 million DALYs from intellectual disability attributable to lead exposure in 2007 is much smaller than their estimate of 9.8 million.

Intimate partner violence

WHO reports a global lifetime prevalence of physical and/or sexual intimate partner violence among ever-partnered women of 30.0% (27.8-32.2).(ref) For GBD 2017, the estimated all-age global exposure for intimate partner violence (IPV) in 2017 is 19.7% (16.1-24.3) among all women, which is a smaller estimate than the WHO estimate because WHO estimates are among only ever-partnered women, while the estimates used for GBD risk factor exposure are among all women. After making an adjustment using our model for the proportion of women who have ever been partnered, we estimate global lifetime IPV exposure as 32.5% (25.9-40.9) among ever-partnered women – an estimate that agrees with the WHO report. The regional distribution reported by WHO is in agreement with the distribution by GBD super-region; highest prevalence of IPV in North Africa and Middle East; South Asia; and sub-Saharan Africa and lower prevalence in Southeast Asia, East Asia, and Oceania; Central Europe, Eastern Europe, and Central Asia; high-income; and Latin American and Caribbean.

Iron deficiency

Iron deficiency (ID) was the fourth ranked level 3 risk factor in 1990, decreasing to sixth in 2017 after increasing 15.23% over that time period in terms of attributable YLDs, almost all of which was YLDs due to dietary iron deficiency (IDA). We have not identified any other global, systematic analyses of ID as a risk factor for increased disease burden so we are not able to compare our estimates of ID-attributable health loss. There are a number of other studies that have evaluated the prevalence of ID and IDA, however.⁴² The most comprehensive meta-analysis from Low and Middle-Income Countries (LMIC) estimated a much lower prevalence of ID/IDA than we have for GBD 2017. There are three aspects that

make a direct comparison with GBD 2017 difficult. First, the study by Petry and colleagues likely underestimated ID/IDA somewhat by applying a single cut-off for diagnosing ID of <12 grams per decilitre of plasma ferritin concentration, especially with the acknowledged limitation of not being able to fully account for the effect of inflammation in many of its component studies. Second, Petry and colleagues did not distinguish aetiologies of ID/IDA whereas GBD does distinguish many causes of anaemia (e.g. hookworm, gastritis) that can manifest as ID. Third, whereas Petry and colleagues made direct estimation of ID/IDA from serum measurements, the GBD approach for estimating ID/IDA is indirect and therefore does not have a directly comparable case definition. We began by first estimating overall anaemia then, after reassigning large portions to >25 other underlying causes, used fixed proportion redistribution methods to estimate IDA. The risk exposure for ID was then estimated as a counterfactual haemoglobin concentration in the absence of all the "other" causes rather than an explicit prevalence value. Unless all possible causes of anaemia are included, the GBD approach has potential to overestimate the proportion of anaemia to be redistributed to ID/IDA in places where other causes are important. We have begun work to address this in GBD 2017 by adding Cirrhosis, Crohn's Disease, Ulcerative Colitis, and others as causes of anaemia, but there are still a number of others (e.g. cancers, alpha thalassemia, intestinal infections, and other nutritional deficiencies) that have yet to be included.

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Section 4: Risk-specific estimation

The risk-specific modelling write-ups follow the order of the risk factor hierarchy for GBD 2017. In some cases, multiple risk factors are addressed in a single write-up, for example child underweight, wasting, and stunting are all included in a single detailed write-up.

Unsafe Water Capstone Appendix

Flowchart



Input data & methodological summary

Exposure

Case definition

For GBD 2017, exposure to unsafe water was defined based on reported primary water source used by the household and use of household water treatment (HWT) to improve the quality of drinking water before consumption. Water sources were defined as "improved" based on the JMP designation,¹ which includes piped water as improved water, and households with access to piped water connection to the house, yard, or plot were defined as having access to piped water supply. Solar treatment, chlorine treatment, boiling, or the use of filters were all established as effective point-of-use household water treatments based on effect sizes calculated from network meta-analysis.

Input data

The search for usable household surveys and censuses was conducted using the Global Health Data Exchange (GHDx) database. HWT input data is primarily limited to two large survey series (DHS and MICS) due to data availability. Water source data includes censuses and nationally representative surveys such as DHS, MICS, AIS, and WHS. For each survey, household sample weights were multiplied by the number of household members to produce a weighting scheme that estimates proportion of individuals, not proportion of households, exposed to a given indicator. Surveys and censuses were then tabulated to the two water source and two water treatment categories of interest for each location.

Modelling

Water source data is modelled using an ordinal framework, with two distinct models: prevalence of piped water and proportion of improved water (excluding piped) within the non-piped population. Both models produce results for each unique location, year combination. This ordinal framework allows us to estimate the category with the most data (piped water prevalence) and leverage that estimate to

anchor the estimates for improved and unimproved water categories. The results of the improved proportion model are multiplied by the piped water prevalence to calculate improved water prevalence. The sum of improved and piped water prevalence are subtracted from 1 to yield unimproved water prevalence.

HWT categories are estimated in a similar ordinal framework, by modelling prevalence of individuals using no water treatment methods and proportions of households that boil/filter water within the population of households that engage in treatment methods. The prevalence of individuals that boil/filter drinking water is calculated by multiplying the proportion that boil/filter modelled previously times prevalence of any water treatment (estimated by subtracting prevalence of no treatment from 1). The prevalence of individuals that treat their water using solar/chlorine methods was estimated by subtracting the sum of prevalence of no treatment estimates and prevalence of filter/boil treatment from 1. By year and location, each of the above categories are modelled using a 3-step modelling scheme of mixed effect linear regression followed by spatio-temporal Gaussian process regression (ST-GPR), which produces full time series estimates for each GBD 2017 location. Socio-demographic index (SDI), a composite metric combining education per capita, income per capita, and fertility, was set as a fixed effect in the linear regression since it proved to be a significant predictor. Random effects were set at GBD 2017 region and super-region levels to fit the models but were not used in the predictions.

The process of vetting and validating models was accomplished primarily through an examination of ST-GPR scatter plots by GBD 2017 location from 1990-2017. Any unfitting data points were re-inspected for error at the level of extraction and survey implementation, and subsequently excluded from analysis if deemed appropriate. In addition to SDI, a number of different potential fixed effects were considered, including lag-distributed income and urbanicity, but SDI proved to be the strongest predictor of the unsafe water categories. Uncertainty in the estimates was initially formed based on standard deviation by survey, then propagated through ST-GPR modelling by means of confidence intervals around each data point that reflect the point-estimate specific variance.

Once models are vetted, full time series outputs from ST-GPR modelling are then converted from proportion to prevalence by year and geography and then rescaled to form 9 mutually exclusive categories that sum up to 1. The table below provides the final result of this rescaling.

Category	Definition
Unimproved, no HWT	Proportion of individuals that primarily use unimproved source, and <i>do not</i> use any HWT to purify their drinking water.
Unimproved, chlorine/solar	Proportion of individuals that primarily use unimproved source, and solar or chlorine treatment to purify their drinking water.
Unimproved, boil/filter	Proportion of individuals that primarily use unimproved source, and boil or filter to purify their drinking water.
Improved water except piped, no HWT	Proportion of individuals that primarily use improved sources other than piped water supply, and <i>do not</i> use any HWT to purify their drinking water.

Improved water except piped, chlorine/solar	Proportion of individuals that primarily use improved sources other than piped water supply, and use solar or chlorine treatment to purify their drinking water.
Improved water except piped, boil/filter	Proportion of individuals that primarily use improved sources other than piped water supply, and boil/filter their drinking water.
	Proportion of individuals that primarily use basic piped water supply, and <i>do not</i> use any HWT to purify their drinking water
Basic piped water, no HWT	
Basic piped water, chlorine/solar	Proportion of individuals that primarily use basic piped water supply, and <i>use</i> solar or chlorine water treatment to purify their drinking water.
Basic piped water, boil/filter	Proportion of individuals that primarily use basic piped water supply, and boil or filter to purify their drinking water
High-quality (HQ) piped water, boil/filter	Proportion of individuals that primarily use basic piped water supply, and boil or filter to purify their drinking water

We modelled the microbiological quality of piped water sources primarily using data a review by Bain et al.¹ that measured proportion of piped water sources contaminated with fecal indicators. We use the value generated from this model to split the prevalence of piped water into basic piped water and high quality piped water by location, year, age, and sex.

A substantial limitation in our analysis is the paucity of data on HWT and piped water quality. The inclusion of more location-specific data on water treatment utilisation at the household level can greatly improve our estimates in future iterations.

Theoretical minimum-risk exposure level

The theoretical minimum-risk exposure level for unsafe water is defined as all households have access to high quality piped water that has been boiled or filtered before drinking.

Relative risks

For GBD 2017, unsafe water was paired with one outcome-diarrheal diseases-given evidence provided by relative risk studies. A meta-analysis by Wolf et al.³ provided the bulk of the relative risk evidence for the relationship between unsafe water and diarrheal diseases. This meta-analysis was updated through a literature review that searched for related intervention studies post-2014 conducted in PubMed. Search terms used were identical to those provided by Wolf et al.³ Relative risk values for water-source interventions and point-of-use treatment interventions were calculated using network meta-analysis approach so as to include studies that differ in control groups within the same analysis. This analysis produced distinct relative risks for each water source and water treatment category. The combined effect of a source intervention and point-of-use intervention was assumed to be multiplicative in order to match GBD 2017 exposure definitions. Please refer to appendix tables for more information on relative risk values and citations.

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Unsafe Sanitation Capstone Appendix

Flowchart



Input data & methodological summary

Exposure

Case definition

Exposure to unsafe sanitation is defined based on the primary toilet type used by households. Improved facilities are defined as such based on JMP designation (WHO). Sewer connection toilets included flush toilets or any toilet with connection to the sewer or septic tank.

Input data

The search for usable household surveys and censuses was conducted using the Global Health Data Exchange (GHDx) database. For each survey, household sample weights were multiplied by the number of household members to produce a weighting scheme that estimates proportion of individuals, not proportion of households, exposed to a given indicator. Surveys and censuses were then tabulated to two sanitation categories, sewer connection and improved sanitation, for each location. Data in tabulated form was lower priority to add to models and was only updated when time permitted.

Modeling

A change made for GBD 2017 was to model sanitation categories in an ordinal framework instead of independent models. Two distinct indicators were estimated: the prevalence of individuals using sewer connection or septic tank facilities and the proportion of individuals with improved sanitation within the population not connected to sewer or septic tank. This ordinal framework allows us to estimate the category with the most data (sewer connection/septic tank prevalence) and leverage that estimate to anchor the estimates for improved and unimproved sanitation categories. The results of the improved proportion model are multiplied by the sewer connection/septic tank prevalence to calculate improved sanitation prevalence. The sum of improved and sewer connection/septic tank prevalence are subtracted from 1 to yield unimproved sanitation prevalence.

The two indicators were modeled using a 3-step modeling scheme of mixed effect linear regression followed by spatio-temporal Gaussian process regression (ST-GPR), which produced full time series estimates for each GBD 2017 location. Socio-demographic index (SDI), a composite metric combining education per capita, income per capita, and fertility, was set as a fixed effect in the linear regression since it proved to be a significant predictor. Random effects were set at GBD 2017 region and super-region levels to fit the models but were not used in the predictions.

The process of vetting and validating models was accomplished primarily through an examination of ST-GPR scatter plots by GBD 2017 location from 1990-2017. Any unfitting data points were re-inspected for error at the level of extraction and survey implementation, and subsequently excluded from analysis if deemed appropriate. In addition to SDI, a number of different potential fixed effects were considered, including lag-distributed income and urbanicity, but SDI proved to be the strongest predictor of unsafe sanitation in terms of magnitude of the coefficient. Uncertainty in the estimates was initially constructed based on standard deviation around each survey mean, then propagated through ST-GPR modeling by incorporating the variance of each data point in the Gaussian process regression step. A data point with high variance, for example, would contribute relatively less influence to the model than a data point with lower variance.

Once models are vetted, full time series outputs from ST-GPR modeling are then converted from proportion to prevalence by year and geography and then rescaled to form three mutually exclusive categories that sum up to 1. The table below provides the final result of this rescaling.

Category	Definition
Unimproved sanitation	Proportion of individuals that use unimproved sanitation facilities.
Improved sanitation	Proportion of individuals with access to improved sanitation facilities, excluding sewer connection or septic tank.
Sanitation facilities with sewer connection or septic tank	Proportion of individuals with access to toilet facilities with sewer connection or septic tank.

Theoretical minimum-risk exposure level

The theoretical minimum-risk exposure level for unsafe sanitation was defined as all individuals have access to a sanitation facility with sewer connection.

Relative risks

For GBD 2017, unsafe sanitation was only paired with one outcome, diarrheal diseases. A meta-analysis by Wolf et al. 2014 provides the bulk of the relative risk evidence for the relationship between unsafe sanitation and diarrheal diseases. This meta-analysis was updated through a literature review that searched for related intervention studies post-2014 conducted in PubMed. Search terms used were identical to those provided by Wolf et al. 2014. Please refer to appendix tables for more information on relative risk values and citations.

References

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Unsafe Hygiene Capstone Appendix

Flowchart

Unsafe Handwashing



Input data & methodological summary

Exposure

Case definition

Unsafe hygiene is defined as lack of access to a handwashing station with available soap and water. We estimated the burden of unsafe hygiene in both developed and developing settings.

Input data

Since water and soap availability data are very limited, only country-specific Demographic Health Surveys (DHS) and Malaria Indicator Survey Series (MICS) conducted after 2006 were included as input data.

Modelling strategy

By year and location, proportion of households with handwashing facility is modelled using a 3-step modelling scheme of mixed effect linear regression followed by spatio-temporal Gaussian process regression (ST-GPR), which outputs full time series estimates for each GBD 2017 location. Socio-demographic index (SDI), a composite index that include income per capita, education, and fertility, was set as a fixed effect in the linear regression since it proved to have significant coefficient. Random effects were set at GBD 2017 region and super-region levels to fit the model but were not used in the predictions.

The process of vetting and validating models was accomplished primarily through an examination of ST-GPR scatter plots by GBD 2016 location from 1990-2016. Any data points lacking face validity were reinspected for error at the level of extraction and survey implementation, and subsequently excluded from analysis if deemed appropriate. In addition to SDI, a number of different potential fixed effects were considered, including lag-distributed income and urbanicity. However, SDI proved to be the strongest predictor.

A considerable limitation for when estimating handwashing practices for over 190 independent locations around the world was data sparseness. Even when data were published on handwashing prevalence, the definition was often altered from the GBD 2017 standard definition or it may only have pertained to certain populations (such as hospital patients) and lacked representativeness at the geographic scale we required. The incorporation of questions about soap and water availability in DHS and MICS added much-needed information but there remains a large data gap to be filled if we are to become more certain in handwashing access estimates.

Theoretical minimum-risk exposure level

The theoretical minimum-risk exposure level for unsafe hygiene is defined as all individuals with access to handwashing facility after any contact with excreta, including children's excreta.

Relative risks

A meta-analysis by Cairncross et al.¹ provide relative risk values describing the relationship between lack of facility access and diarrheal diseases. A meta-analysis by Rabie and Curtis² provided relative risk evidence for the relationship between lack of facility access and lower respiratory infection. Please refer to appendix tables for more information on relative risk values and citations.

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Ambient Particulate Matter Pollution Capstone Appendix

Flowchart



Ambient particulate matter pollution

Input data and modeling strategy

Exposure

Definition

Exposure to ambient air pollution is defined as the population-weighted annual average mass concentration of particles with an aerodynamic diameter less than 2.5 micrometers ($PM_{2.5}$) in a cubic meter of air. This measurement is reported in $\mu g/m^3$.

Input Data

The data used to estimate exposure to ambient air pollution is drawn from multiple sources, including satellite observations of aerosols in the atmosphere, ground measurements, chemical transport model simulations, population estimates, and land-use data.

The following details the updates in methodology and input data used in GBD 2017.

PM_{2.5} ground measurement database

Updates of ground measurements used for GBD 2017 include using more recent data than that used previously and the addition of data from new locations. The data from the 2018 update of the WHO Global Ambient Air Quality Database include monitor-specific measurements of concentrations of PM_{10} and $PM_{2.5}$ from 9,960 ground monitors (up from 6,003 in GBD 2016) from 108 countries. The majority of measurements were recorded in 2016 (as there is a lag in reporting measurements, little data from 2017 were available). Annual averages were excluded if they were based on less than 75% coverage within a year. Collection year ranged from 2008 to 2017 in data used. If information on coverage was not available then data were included unless they were already sufficient data within a country (monitor density greater than 0.1).

For locations measuring only PM_{10} , $PM_{2.5}$ measurements were estimated from PM_{10} . This was performed using a hierarchy of conversion factors ($PM_{2.5}/PM_{10}$ ratios): (i) for any location a 'local'

conversation factor was used, constructed as the ratio of the average measurements (of $PM_{2.5}$ and PM_{10}) from within 50km and within the same country, if such were available' (ii) if there was not sufficient local information to construct a conversion factor then a country-wide conversion factor was used; and (iii) if there was no appropriate information within a country then a regional factor was used. In each case, to avoid the possible effects of outliers in the measured data (both $PM_{2.5}$ and PM_{10}), extreme values of the ratios were excluded (defined as being greater/lesser than the 95 and 5% quantiles of the empirical distributions of conversion factors) of the latter two cases for the country measurements were available, for both metrics. As in the GBD 2013 and GBD 2015/GBD 2016 databases, in addition to values of $PM_{2.5}$ and whether they were direct measurement or converted from PM_{10} , the database also included additional information, where available, related to the ground measurements such as monitor geo coordinates and monitor site type.

Satellite-based estimates

The updated satellite-based estimates for years 1998-2016 are described in detail in van Donkelaar et al. 2016.¹ These estimates were available at $0.1^{\circ} \times 0.1^{\circ}$ resolution (~11 x 11 km resolution at the equator) and combine aerosol optical depth retrievals from multiple satellites with the GEOS Chem chemical transport model and land use information.

Population data

A comprehensive set of population data on a high-resolution grid was obtained from the Gridded Population of the World (GPW) database. These estimates are adjusted to match UN2015 Population Prosepectus. These data are provided on a 0.0417°×0.0417° resolution. Aggregation to each 0.1°×0.1° grid cell comprised of summing the central 3 × 3 population cells. As this resulted in a resolution higher than necessary, it was repeated four times, each offset by one cell in a North, South, East and West direction. The average of the resulting five quantities was used as the estimated population for each grid cell. Population estimates for 2000, 2005, 2010, 2015 and 2020 were available from GPW version 4 revision 10. Populations for 2016 and 2017 were obtained by interpolation using natural splines with knots placed at 2000, 2005, 2010, 2015 and 2020. This was performed for each grid cell.

Chemical transport model simulations

Estimates of the sum of particulate sulfate, nitrate, ammonium and organic carbon and the compositional concentrations of mineral dust simulated using the GEOS Chem chemical transport model, and a measure combining elevation and the distance to the nearest urban land surface (as described in van Donkelaar et al. 2016¹) were available for 2000 to 2016 for each 0.1°×0.1° grid cell. These were not included within the GBD 2013 analysis.

Modelling strategy

Significant advances have been made in the methodology used to estimate exposure to ambient particulate matter pollution since GBD 2013. The following is a summary of the modelling approach, known as the Data Integration Model for Air Quality (DIMAQ) used in GBD 2015, 2016, and 2017; further details can be found in Shaddick *et al.* (2017).²

In GBD 2010 and GBD 2013 exposure estimates were obtained using a single global function to calibrate available ground measurements to a 'fused' estimate of $PM_{2.5}$; the mean of satellite-based estimates and those from the TM5 chemical transport model, calculated for each $0.1^{\circ} \times 0.1^{\circ}$ grid cell. This was recognised to represent a trade-off between accuracy and computationally efficiency when utilising all the available data sources. In particular, the GBD 2013 exposure estimates were known

to underestimate ground measurements in specific locations (see discussion in Brauer et al., 2013³). This underestimation was largely due to the use of a single, global, calibration function, whereas in reality the relationship between ground measurements and other variables will vary spatially.

In GBD 2015 and GBD 2016, coefficients in the calibration model were estimated for each country. Where data were insufficient within a country, information can be `borrowed' from a higher aggregation (region) and if enough information is still not available from an even higher level (super-region). Individual country level estimates were therefore based on a combination of information from the country, its region and super-region. This was implemented within a Bayesian Hierarchical modelling (BHM) framework. BHMs provide an extremely useful and flexible framework in which to model complex relationships and dependencies in data. Uncertainty can also be propagated through the model allowing uncertainty arising from different components, both data sources and models, to be incorporated within estimates of uncertainty associated with the final estimates. The results of the modelling comprise a posterior distribution for each grid cell, rather than just a single point estimate, allowing a variety of summaries to be calculated. The primary outputs here are the median and 95% credible intervals for each grid cell. Based on the availability of ground measurement data, modelling and evaluation was focused on the year 2016.

The GBD 2017 model was updated to also include within country calibration variation.⁴ The model used for GBD2017, henceforth referred to as DIMAQ2, provides a number of substantial improvements over the initial formulation of DIMAQ. In DIMAQ, ground measurements from different years were all assumed to have been made in the primary year of interest (i.e. 2014 for GBD2015 before extrapolation) and then regressed against values from other inputs (e.g. satellites etc.) made in that year. In the presence of changes over time therefore, and particularly in areas where no recent measurements were available, there was the possibility of mismatches between the ground measurements and other variables. In DIMAQ2, ground measurements and matched with other inputs (over time) and the possibility of the (global level) coefficients being allowed to vary over time, subject to smoothing that is induced by a second-order random walk process. In addition, the manner in which spatial variation can be incorporated within the model has developed: where there is sufficient data, the calibration equations can now vary (smoothly) both within and between countries, achieved by allowing the coefficients to follow (smooth) Gaussian processes. Where there is insufficient data within a country, to produce accurate equations, as before information is borrowed from lower down the hierarchy and it is supplemented with information from the wider region.

DIMAQ2 is used for all regions except for the North Africa-Middle East and Sub-Saharan superregions and remote islands where there is insufficient data to allow the extra complexities of the new model to be implemented. In the North Africa-Middle East and Sub-Saharan super-regions a simplified version of DIMAQ2 is used in which the temporal component is dropped, and for remote islands the original DIMAQ is used.

Due to both the complexity of the models and the size of the data, notably the number of spatial predictions that are required, recently developed techniques that perform 'approximate' Bayesian inference based on integrated nested Laplace approximations (INLA) were used.⁵ Computation was performed using the R interface to the INLA computational engine (<u>R-INLA</u>). Fitting the models and performing predictions for each of the ca. 1.4 million grid cells required the use of a high performance computing cluster (HPC) making use of high memory nodes.

Model evaluation

Model development and comparison was performed using within- and out-of-sample assessment. In the evaluation, cross validation was performed using 25 combinations of training (80%) and validation (20%) datasets. Validation sets were obtained by taking a stratified random sample, using sampling probabilities based on the cross-tabulation of $PM_{2.5}$ categories (0-24.9, 25-49.9, 50-74.9, 75-99.9, 100+ μ g/m³) and super-regions, resulting in them having the same distribution of $PM_{2.5}$ concentrations and super-regions as the overall set of sites. The following metrics were calculated for each training/evaluation set combination: for model fit - R² and deviance information criteria (DIC, a measure of model fit for Bayesian models); for predictive accuracy - root mean squared error (RMSE) and population weighted root mean squared error (PwRMSE).

All modelling was performed on the log-scale. The choice of which variables were included in the model was made based on their contribution to model fit and predictive ability. The following is a list variables and model structures that were included in DIMAQ.

Continuous explanatory variables:

- \circ (SAT) Estimate of PM_{2.5} (in μ gm⁻³) from satellite remote sensing on the log-scale.
- (POP) Estimate of population for the same year as SAT on the log-scale.
- (SNAOC) Estimate of the sum of sulfate, nitrate, ammonium and organic carbon simulated using the GEOS Chem chemical transport model.
- (DST) Estimate of compositional concentrations of mineral dust simulated using the GEOS Chem chemical transport model.
- (EDxDU) The log of the elevation difference between the elevation at the ground measurement location and the mean elevation within the GEOS Chem simulation grid cell multiplied by the inverse distance to the nearest urban land surface.

Discrete explanatory variables:

- (LOC) Binary variable indicating whether exact location of ground measurement is known.
- o (TYPE) Binary variable indicating whether exact type of ground monitor is known.
- $\circ~$ (CONV) Binary variable indicating whether ground measurement is PM_{2.5} or converted from PM_{10}.

Random Effects:

- Grid cell random effects on the intercept to allow for multiple ground monitors in a grid cell.
- o Country-region-super-region hierarchical random effects for the intercept.
- \circ $\,$ Country-region-super-region hierarchical random effects for the coefficient associated with SAT .
- Country-region-super-region hierarchical random effects for the coefficient associated with the difference between estimates from CTM and SAT.
- Country-region-super-region hierarchical random effects for the coefficient associated with POP.
- Country level random effects for population uses a neighbourhood structure allowing specific borrowing of information from neighbouring countries.
- Within a region, country level effects of SAT and the difference between SAT AND CTM are assumed to be independent and identically distributed.
- Within a super-region, region level random effects are assumed to be independent and identically distributed.

• Super-region random effects are assumed to be independent and identically distributed.

Interactions:

o Interactions between the binary variables and the effects of SAT and CTM.

In addition, DIMAQ2 includes

- o Smoothed, spatially varying, random-effects for the intercept
- Smoothed, spatially varying, random-effects for the coefficient of coefficient associated with SAT
- o Smoothed, temporally varying, random-effect for the intercept

Results

The final model contained the following variables: SAT, POP, SNAOC, DST, EDxDU, LOC, TYPE, and CONV, together with interactions between SAT and each of LOC, TYPE and CONV. The model structure contained grid cell random effects on the intercept to allow for multiple ground monitors in a grid cell, country-region-super-region hierarchical random effects for intercepts and SAT and country level random effects for population using a neighbourhood structure allowing specific borrowing of information from neighbouring countries together with region-super-region hierarchical random effects and GBD 2016, based on the evaluation of candidate models, including estimates from the TM5 chemical transport model (CTM) used in GBD 2013 did not improve the predictive ability of the model and was therefore not included.

Compared to the model used in GBD2013, DIMAQ showed improved predictions of ground measurements in all super regions with improvements in both within-sample fit; with a global population-weighted RMSE of 12.1 μ g/m³ compared to 23.1 μ g/m³ when using the GBD 2013 approach.¹ Using the larger database available for GBD2017, with potentially more variability in measurements, DIMAQ2 shows an additional improvement on DIMAQ: overall population-weighted RMSE reduced from 9.32 to 8.11 (12.12 to 11.17 when using all data, irrespective of within-year coverage). Reductions by super-region can be seen in Figure 1. Reductions can be seen in all super-regions with particular improvement in the Southeast Asia, East Asia and Oceania super-region which is based largely on a substantial increase in accuracy in China, PwRMSE 6 vs 9 μ g/m³


Figure 1: Summary measures of predictive ability, globally and by super-region. Dots denote the median values of population weighted root mean squared error (μ g/m³) from 25 validation sets with vertical lines showing the range of values over those sets.

Estimates for other years

In contrast to the method used previously, where estimates (of PM_{2.5}) were extrapolated to produce estimates for the year of interest (e.g. 2017 where data was available up to and including 2016) due to the extra complexity of the smooth spatial processes in DIMAQ2 this would not be possible in any straightforward manner. With DIMAQ2 it is the input variables that are extrapolated; this allows estimates for 2017 to be produced in the same way as other years and crucially, allows measures of uncertainty to be produced within the BHM framework rather than by using post-hoc approximations.

Satellite estimates and quantities estimated using the GEOS-Chem model were available for 1990, 1995, 2000, 2005, 2010-2016. Estimates of these input variables for 2017 were produced by extrapolating, on a cell-by-cell basis, using natural splines. Population estimates for 2000, 2005, 2010, 2015 and 2020 were available from GPW version 4. For 1990 and 1995 data were extracted from GPW version 3, as in GBD2013.² As with populations for 2015, values for each cell for 2011-2017 were obtained by interpolation using natural splines with knots placed at 2000, 2005, 2010, 2015 and 2020.

These were used as inputs to DIMAQ, enabling estimates of exposures to be obtained for each of these years respectively. For 2017, estimates of exposures were obtained from predictions from locally-varying regression models.⁶ For each cell a model was fit to the values within that cell over time, with a constraint placed on the rate of change between 2016 and 2017 to avoid unrealistic and/or unjustified extrapolation of trends. Measures of uncertainty were obtained by repeating the procedure for the limits of the 95% credible intervals, again on a cell-by-cell basis.

Population-weighted exposure generation

To generate a distribution of the population-weighted ambient particulate matter, we took a weighted sampling strategy, taking samples from all grid cells in a given location. For example, for a country with n grid cells, we randomly sampled 1000 values from the n (grid cells) x 1000 (samples) where the probability of being sampled was proportional to the population of that grid cell.

Theoretical minimum-risk exposure level

The TMREL was assigned a uniform distribution with lower/upper bounds given by the average of the minimum and 5th percentiles of outdoor air pollution cohort studies exposure distributions conducted in North America, with the assumption that current evidence was insufficient to precisely characterise the shape of the concentration-response function below the 5th percentile of the exposure distributions. The TMREL was defined as a uniform distribution rather than a fixed value in order to represent the uncertainty regarding the level at which the scientific evidence was consistent with adverse effects of exposure. The specific outdoor air pollution cohort studies selected for this averaging were based on the criteria that their 5th percentile of 8.2 based on Turner et al. (2016).⁷ This criterion was selected since GBD 2010 used the minimum, 5.8, and 5th percentile solely from the CPS II cohort. The resulting lower/upper bounds of the distribution for GBD 2017 were 2.4 and 5.9. This has not changed since GBD 2015.

Relative risks and population attributable fractions

We estimated the Ambient Air Pollution-attributable burden of disease based on the relation of long-term exposure to PM2.5 with Ischemic Heart Disease, stroke (ischemic and hemorrhagic), COPD, lung cancer and acute lower respiratory infection. These were also the pollutant-outcome pairs used to estimate the Ambient Air Pollution attributable burden since GBD 2010. For GBD 2017 we also added Type II Diabetes as an outcome of ambient air pollution. We used results from all cohort studies published as of July 2018 that reported cause-specific relative risk estimates based on measured or modelled PM2.5 and that adjusted for potential confounding due to other major risk factors such as tobacco smoking using data for each study participant.

Bowe et al. recently published work that assembled the evidence for the relationship between particulate matter and diabetes to generate IER curves and attributable burden estimates based on methodologies similar to those of the GBD.⁸

When generating the IER for Type II Diabetes, we included all eight of the studies summarized by Bowe et al. in addition to six other cohorts. Resulting attributable burden estimates were remarkably similar to GBD 2017 results. All citations for studies used in the fitting of the IER curve can be found using the GBD 17 Data Input Sources Tool.

Integrated exposure response function

The Integrated Exposure Response Function (IER) was created to ascertain the shape of the dose response curve for a variety of health outcomes across a wide range of exposure to PM2.5. The IER model is fit by integrating RR information from studies of outdoor air pollution (OAP), Second hand tobacco smoke (SHS), Household Air Pollution (HAP), and Active Smoking (AS). Because OAP studies are often performed at the lower end of the ambient air pollution range, incorporating other exposures to particulate matter enables RR estimation across the global range of exposure. These methods have been described in detail elsewhere.^{9,10}

Notable changes for GBD 2017 include added studies for OAP, SHS, and HAP, updated literature reviews for AS studies, and more informative priors to stabilize the shape of the IER curves.

- We added all newly published cohorts of long-term exposure to Ambient PM2.5 and incidence or mortality due to IHD, stroke, COPD, lung cancer, and LRI. One notable addition was the China Male Cohort which included mortality due to IHD, Stroke, COPD, Lung Cancer, and Diabetes (unpublished analysis).¹¹ This study represented a higher exposure range than most of our previously incorporated studies with 5th and 95th percentile of 15.5 and 77.1 micrograms/m³. For Type II Diabetes, the new outcome included in GBD 2017, we included all cohorts which measured long-term PM2.5 exposure and incident diabetes or mortality due to diabetes.
- We did not change the SHS input studies with the exception of including all studies from a recent meta-analysis examining the relationship between SHS and Type II Diabetes.¹² We also added seven studies found from a systematic review examining SHS exposure and COPD. We had previously not included SHS in the formation of this curve.
- We added four cohort studies of HAP and any of our measured outcomes. Previously we have only included which measured levels of PM2.5 exposure. To incorporate cohort studies with binary exposure data (presence or absence of solid-fuel use for cooking) we used the PM2.5 mapping function (see Household Air Pollution Appendix for more details) to obtain a PM2.5 level attributed to solid fuel use for cooking for the location-year of the study (Exp_{HAP}). We also used the OAP exposure model to obtain an OAP PM2.5 level for the location-year (Exp_{OAP}). The study RR was used to inform the curve on the range of Exp_{OAP} to (Exp_{OAP} + Exp_{HAP}).
- For all outcomes, we used updated systematic reviews of the literature performed by the GBD smoking team for studies examining cigarettes smoked per day and the six IER outcomes to inform the high exposure range of the curve. The smoking team found that the process of systematic review and inclusion of all acceptable studies led to lower relative risks.
- To help obtain more reasonable curve fits, we added more informative priors to two of three IER function parameters in the MCMC Bayesian fitting process.

Limitations

It is important to recognize the inherent limitations of the IER approach. The use of various sources to construct a risk curve assumes an equitoxicity of particles, consistent with evaluations by US EPA and WHO. However, current evidence suggests there are differences in health impact by source, size, and chemical composition. This is seen when comparing studies of ambient and household particulate matter. As this body of evidence grows, we will continue to re-examine our strategy for the integrated exposure-response curve. For now, the IER is a practical solution to fill gaps in the literature where we do not have sufficient evidence such as household air pollution exposures and ambient in highly polluted areas.

Additionally, currently the exposure concentrations used for both SHS and AS data points when fitting the IER are contrasted with the TMREL and do not take into account ambient particulate matter pollution. In future iterations of fitting the curve, we will test alternate approaches, including a similar approach to HAP, allowing each data point to inform the curve on the range of Exp_{OAP} to $(Exp_{OAP} + Exp_{AS/SHS})$.

Relative risk and proportional PAF approach

For GBD 2017 we developed a new approach to use the IER for obtaining PAFs for both OAP and HAP. Previously, relative risks for both exposures were obtained from the IER as a function of exposure and relative to the same TMREL. In reality, were a country to reduce only one of these risk factors, the other would remain. We failed to consider the joint effects of particulate matter from outdoor exposure and burning solid fuels for cooking.

In GBD 2017, relative risks were still estimated from the output of the IER curve. Everyone is exposed to some level of OAP, but only a proportion of the population in each location-year use solid cooking fuel and are exposed to HAP. For the proportion of the population not exposed to HAP the relative risk was obtained by $RR_{OAP} = IER(z = Exp_{OAP})$ and used to calculate the PAF for each location based on the population-weighted exposure.

For the proportion of the population exposed to both OAP and HAP, we calculated a joint relative risk from the IER by $RR_{OAP+HAP} = IER(z = Exp_{OAP}+Exp_{HAP})$. This joint relative risk is used to calculate a joint PAF for each location. PAF calculation is detailed in the methods appendix. For each location, we proportioned the joint PAF based on the proportion of exposure due to OAP and HAP respectively. See the table below for equations used to calculate proportional PAFs.

PAF Population not exposed to HAP		Population exposed to HAP	
ΟΑΡ	PAF _{OAP}	(Exp _{OAP} /(Exp _{OAP} +Exp _{HAP}))*PAF _{OAP+HAP}	
НАР	0	(Exp _{HAP} /(Exp _{OAP} +Exp _{HAP}))*PAF _{OAP+HAP}	

Generally, as expected, this new strategy led to lower PAFs for both ambient and household particulate matter pollution.

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Household Air Pollution Capstone Appendix

Flowchart



Input Data & Methodological Summary

Exposure

Case definition

Exposure to household air pollution from solid fuels (HAP) is defined as the proportion of households using solid cooking fuels. The definition of solid fuel in our analysis includes coal, wood, charcoal, dung, and agricultural residues.

Input data

Data were extracted from the standard multi-country survey series such as Demographic and Health Surveys (DHS), Living Standards Measurement Surveys (LSMS), Multiple Indicator Cluster Surveys (MICS), and World Health Surveys (WHS), as well as country-specific survey series such as Kenya Welfare Monitoring Survey and South Africa General Household Survey. To fill the gaps of data in surveys and censuses, we also downloaded and updated HAP estimates from WHO Energy Database and extracted from literature through systematic review. Each nationally or sub-nationally representative data point provided an estimate for the percentage of households using solid cooking fuels. Estimates for the usage of solid fuels for non-cooking purpose were excluded, i.e. primary fuels for lighting. The database, with estimates from 1980 to 2017, contained about 680 studies from 150 countries. As updates to systematic reviews are performed on an ongoing schedule across all GBD causes and risk factors, an update for household air pollution will be performed in the next 1-2 iterations.

Modelling strategy

Household air pollution was modelled at household level using a three-step modelling strategy that uses linear regression, spatiotemporal regression and Gaussian Process Regression (GPR). The first step is a mixed-effect linear regression of logit-transformed proportion of households using solid cooking fuels. The linear model contains maternal education, proportion of population living in urban areas, and

lagged-distributed income as covariates and has nested random effect by GBD region, and GBD super region respectively. The full ST-GPR process is specified elsewhere this appendix. No substantial modelling changes were made in this round compared to GBD 2016.

Theoretical minimum-risk exposure level

For cataract, the TMREL is defined as no households using solid cooking fuel. For outcomes that utilise evidence based on the Integrated Exposure Response (IER), the TMREL is defined as uniform distribution between 2.4 and 5.9 ug/m³.

Relative risks

In addition to the previously included outcomes of lower respiratory infections (LRI), stroke, Ischemic Heart Disease (IHD), Chronic Obstructive Pulmonary Disease (COPD), lung cancer, and cataract, in GBD 2017 we added Type II Diabetes as a new outcome of household air pollution. The relative risk for cataracts was extracted from a meta-analysis and is 2.47 with 95% (1.61, 3.73).¹ GBD currently only estimates cataracts as an outcome for females.

In GBD 2017, we adopted a new approach for risk attribution using the Integrated Exposure-Response Function (IER). Updates to the IER and the new joint-estimation PAF approach is described in the Ambient Particulate Matter appendix.

PM_{2.5} mapping value

In order to use the IER curve, we must estimate the exposure to particulate matter with diameter of less than 2.5 micrometers (PM_{2.5}). Since GBD 2015 we have been using a mapping model relying on a database of now almost 90 studies which measures PM_{2.5} exposure in households using solid cooking fuel. Using socio-demographic index and study-level factors as covariates, we predict exposure for all location-years.

In GBD 2017, we updated the model to estimate the individual exposure to $PM_{2.5}$ over and above ambient levels due to the use of solid cooking fuel. We did this by subtracting off the estimated ambient level $PM_{2.5}$ for the location-year of each study in the database before inputting them into the model. By doing this we have independent estimates for $PM_{2.5}$ exposure due to ambient and household solid fuel use.

These exposures are cross-walked to values for men, women, and children by generating the ratio of each group's mean exposure to the overall mean personal exposure. The resulting location, year, sex, and age specific PM_{2.5} exposure values are used as inputs in the IER and attributable burden calculation process.

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Ambient Ozone Pollution Capstone Appendix

Flowchart



Input data and methodological summary

Exposure

Case definition

For GBD 2017, exposure to ozone pollution is defined as the seasonal (6 month period with highest mean) 8 hour daily maximum ozone concentrations, measured in ppb. This was an update from the previous exposure metric in accordance with an update of the American Cancer Society Cancer Prevention Study II (ACS CPS-II).¹

Input data

Previously, exposure estimates were based on a chemical transport model with no measurement database or evaluation. In GBD 2017, exposure estimates incorporated a new comprehensive ozone measurement database (TOAR).² This enabled a continent-specific weighted blend of 6 chemical transport models with grid cell level bias correction. The use of ground measurements also enabled the incorporation of error estimation, where previously we had assumed a +/- 6% error. The output of this model is a global raster of ozone exposure which is a summary for the years 2008-2014.³

Modelling strategy for trends

To estimate ozone concentrations over time, we used the trend from the former GBD model for 1990, 2000, and 2010 and cubic splines for 1995, 2005, and 2011, after applying an adjustment for the difference in trends between the previous (1 hour daily maximum) and current (8 hour daily maximum metrics. Annualised rate of change was used to predict for the years 2012-2017.

Theoretical minimum-risk exposure level

The TMREL of ozone was updated this year based on the exposure distribution from the updated ACS CPS-II study.¹ A uniform distribution was drawn around the minimum and 5th percentile values experienced by the cohort, defined as ~U(29.1, 35.7), in ppb.

Relative risks

Since the inclusion of ozone in GBD 2010 the relative risk of ozone exposure for respiratory COPD mortality has been defined to be 1.029, 95% C.I. (1.01-1.048) per 10 ppb of ozone exposure. Note that this comes from one study that looked at all respiratory mortality.⁴ For GBD 2017, we performed a literature review and included five cohorts from Canada, the UK, and the US which all measured COPD mortality. For cohorts with multiple analyses we chose the most recent analysis. We found a resulting relative risk of 1.06, 95% C.I. (1.02, 1.10).

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Radon Exposure Capstone Appendix

Flowchart



Input data & methodological summary

Exposure

Case definition

Radon is a radioactive gas that is produced as a byproduct of the decay chain of uranium, occurring naturally within the Earth's crust. Some fraction of this natural radon production escapes into the atmosphere, where it forms at low concentration unless build-up is caused by enclosed spaces like homes, mines, or caves. Radon exposure is expressed as average daily exposure to indoor air radon gas levels measured in Becquerels (disintegrations per second) per cubic meter (Bq/m³).

Input data

Exposure to radon is determined using values curated by an expert group. These values are taken from a variety of sources including literature, government agencies, and monitoring stations. Their methodology is then inspected to determine if they are robust enough to be considered as country-level averages. 76 data points were added for GBD 2017 including several from a study which reports on subnational variation in India. Before modelling, a crosswalk is performed from studies measuring geometric to arithmetic mean.

Modelling strategy

Because radon is naturally occurring and is not considered to have much temporal fluctuation,¹ we shifted from a spatio-temporal GPR model to a mixed effects linear model. The model included nested random effects on super-region, region, and location (most detailed) and one fixed effect covariate, long-term mean temperature as a proxy for adequate building ventilation.

We did not have the microdata necessary to use ensemble modelling to inform our radon exposure distribution, so for GBD 2017 we continued to assume a lognormal distribution. Arithmetic mean exposure estimates obtained were used to fit the lognormal distribution before applying relative risks.

Theoretical minimum-risk exposure level

The TMREL was also taken directly from literature values that were not updated for GBD 2017. Given that radon is naturally occurring, zero exposure would be impossible. As such, we continue to use a TMREL of 10 Bq/m³, which is equivalent to the outdoor concentration of radon.²

Relative risks

The relative risk for radon exposure was extracted from literature values – a 2005 meta-analysis of casecontrol studies showing the association of radon with lung cancer.³ This value was used in GBD 2010 and has not been changed since.

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Lead Exposure Capstone Appendix



Flowchart

Definitions

Exposure to lead is defined in two different ways according to the currently known pathways of health loss. Acute lead exposure, relevant to disease burden through IQ loss in children, is measured as the micrograms of lead per deciliter of blood (μ g/dL). Long-term lead exposure, relevant to disease burden in adults given the manifestation of health impact through increased systolic blood pressure and hence a decline of cardiovascular health, is measured as the accumulation of lead in the bone as micrograms of lead per gram of bone (μ g/g).

Input Data

The input data for lead exposure is primarily extracted from literature regarding blood lead, in addition to a few blood lead surveys. Blood lead values are derived from studies that take blood samples and analyze them using various techniques to determine the level of lead present. Our literature review resulted in 3,183 usable data points from 554 different studies, which span the years 1970 to 2017. The database of literature values was modelled for data-sparse countries using spatio-temporal Gaussian process regression (ST-GPR). These values were used as blood lead exposure estimates. The second pathway of burden is related to bone lead exposure, which was estimated by calculating a cumulative

blood lead index for cohorts using estimated blood lead over their lifetime. The cumulative blood lead index is then used to estimate bone lead using a scalar defined in literature.¹

Exposure Modeling

The methodology to estimate lead exposure last underwent significant change in GBD 2013. Global exposure had been previously modelled using age-integrating Bayesian hierarchal modelling (DisMod-MR). The modelling process was updated for GBD 2013 by shifting to spatial-temporal GPR methodology. This allowed for estimates of all country-age-sex-year groups for single years instead of five year periods. This approach improved the granularity of estimates for bone lead, which requires back-estimation of previous blood lead to calculate a cumulative blood lead index.

For GBD 2017, the spatio-temporal Gaussian process regression modelling methodology was updated as detailed in the appendix specific to this analytical technique, which is common to a variety of risk factors. In order to predict blood lead in country-years with insufficient data, covariates that have been produced across time and space relevant to this analysis were used. For blood lead exposure, the covariates determined to have predictive ability were the socio-demographic index (SDI), the proportion of a location's population living in urban settings (logit transformed), the combined number of 2 and 4-wheel vehicles per capita, and a covariate indicating whether leaded gasoline had been phased out in a given country-year (smoothed over the first 5 years of phase-out to reflect its gradual implementation). ST-GPR was used to produce estimates of mean and standard deviation of blood lead for all age groups, for both sexes, and for all GBD locations from 1970 to 2017.

In earlier iterations of GBD, the distribution of lead exposure was assumed to be log-normal. Since GBD 2016, ensemble modelling techniques were used to find an optimal global distribution by fitting a variety of distributions to the available blood lead microdata. This was a common update for all continuous risk factors. The ST-GPR estimates of mean and standard deviation blood lead were used with the global distribution shape to determine distributions for blood lead exposure.

To calculate blood lead over the lifetime of a given cohort, blood lead was assumed to grow linearly from 2.0 ug/dL in 1920 (see TMREL) to the value for that cohort in 1970. Using the exposure distributions of blood lead over time and space, cohorts were constructed such that lifetime blood lead could be expressed as a curve over each year of life. The area under this curve was the cumulative blood lead index, which could be used to estimate bone lead in a given year with the aforementioned scalar.

Estimating Attributable Burden

Assessment of risk-outcome pairs

We included outcomes based on the strength of available evidence supporting a causal relationship. Blood lead level (a measure of acute lead exposure) is paired with idiopathic developmental intellectual disability as modeled through the impact of blood lead levels on IQ in children. Bone lead level (a measure of cumulative lead exposure) is paired with systolic blood pressure as an outcome, and subsequently to all cardiovascular outcomes to which systolic blood pressure is paired, which includes rheumatic heart disease, ischemic heart disease, ischemic stroke, intracerebral hemorrhage, hypertensive heart disease, other cardiomyopathy, atrial fibrillation and flutter, aortic aneurysm, peripheral artery disease, endocarditis, other cardiovascular and circulatory diseases, chronic kidney disease due to hypertension, chronic kidney disease due to glomerulonephritis, and chronic kidney disease due to other and unspecified causes.

Theoretical minimum-risk exposure level

In previous iterations of GBD, the TMREL was taken from literature estimates of pre-industrial blood lead in humans.² That value was estimated at 2.0 ug/dL. The decision was made that the TMREL of blood lead could not be 0 given the ambient sources of lead that would be impossible to eliminate.³

However, average blood lead exposures in a number of countries have fallen below 2.0 ug/dL in the past few years, suggesting that the TMREL ought to be lowered. Unfortunately, we were not able to find literature with statistically significant estimates for relative risk at such low levels of blood lead exposure. As a result, we have continued to use a TMREL of 2.0 ug/dL for GBD 2017.

Relative Risks

Because the relative risk of IQ loss from lead exposure is specific to children, in GBD 2015 no burden of lead via IQ loss was estimated in the population aged 15 and above. To better account for the continued burden of past lead exposure on IQ in older age groups, since GBD 2016 we have constructed cohorts from the entire population. Estimates of a cohort's lead exposure in early childhood (at 24 months of age) were used to determine past IQ loss, and thus calculate burden via the impact on concurrent IQ in the older population.

Blood lead relative risks were previously taken from a 2005 pooled analysis that was first incorporated in GBD 2010.⁴ For GBD 2017, blood lead relative risks have been updated with a 2013 re-analysis of the findings of that 2005 paper, providing slightly adjusted relative risk estimates specific to exposure at 24 months of age.⁵ The bone lead relative risks were taken from a 2008 meta-analysis that was updated for GBD 2010.⁶

Population Attributable Fraction

We used the standard GBD population attributable fraction (PAF) equation to calculate PAFs for bone lead exposure and each of its paired outcomes using exposure estimates and relative risks. We used a similar approach for estimating PAFs for the burden of intellectual disability attributable to blood lead, which uses the estimated distribution of intellectual disability and the modeled shifts in IQ due to blood lead levels to determine the PAF.

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Occupational Risk Factors Capstone Appendix

Exposure definitions

The following definitions were used for occupational risk factor exposures. All exposures were estimated for ages 15 and older.

Occupational Asbestos	Cumulative lifetime exposure to occupational asbestos, using mesothelioma death rate as an analogue
Occupational Asthmagens	Proportion of the working population exposed to asthmagens, based on population distributions across nine occupational categories
Occupational Carcinogens (arsenic, benzene, beryllium, cadmium, chromium, diesel engine exhaust, formaldehyde, nickel, polycyclic aromatic hydrocarbons, silica, sulfuric acid, and trichloroethylene)	Proportion of the population that was ever occupationally exposed to carcinogens at high or low exposure levels, based on population distributions across seventeen economic activities
Occupational Ergonomic Factors	Proportion of the working population exposed to low back pain-inducing work, based on population distributions across nine occupational categories
Occupational Injuries	Proportion of injuries in the working-age population attributable to occupational work, based on fatal injury rates in seventeen economic activities
Occupational Noise	Proportion of the population occupationally exposed to 85+ decibels of noise, based on population distributions across seventeen economic activities
Occupational Particulates	Proportion of the population occupationally exposed to particulates, based on population distributions across seventeen economic activities

Economic activities and occupations were coded according to the following categories:

Economic Activities	Occupations
Agriculture, hunting, forestry	Legislators, senior officials, and managers
Fishing	Professionals
Mining and Quarrying	Technicians and associate professionals
Manufacturing	Clerks
Electricity, gas, and water	Service workers and shop/market sales workers
Construction	Skilled agricultural and fishery workers
Wholesale and retail trade/repair	Plant and machine operators and assemblers
Hospitality	Craft and related workers

Transport, storage, and communication	Elementary occupations
Financial intermediation	
Real estate/renting	
Public administration/defense; compulsory social	
security	
Education	
Health and social work	
Other community/social/personal service	
activities	
Private households	
Extra-territorial organisations/bodies	

Input data

Primary inputs were obtained from the ILO,¹⁻⁴ and included raw data on economic activity proportions, occupation proportions, fatal injury rates, and employment to population ratio estimates. A systematic web review was conducted in order to collect the underlying microdata from the ILO's estimates to aid in re-extraction at greater levels of granularity. Where freely available, survey datasets were downloaded from the survey organisations in question. Other datasets were obtained through submission of requests to agencies and through the GBD collaborator network. Microdata was tabulated in order to create survey-weighted estimates of economic activities and occupations for the GBD geographies and years. Various classification systems were crosswalked to ISIC Rev.3 (for economic activities) and ISCO 1988 (for occupations). Subnational estimates for UK and China were added to the datasets for economic activities and occupations.^{5,6}

For occupational asbestos, primary inputs were obtained through GBD 2017 cause of death estimates and published studies.^{7,13,14}

Uncertainty for inputs where microdata was unavailable was generated by fitting a Loess curve to the data and determining the standard deviation of the data from the fitted curve.

Modelling strategies

A Spatio-temporal Gaussian process regression (ST-GPR) was used to generate estimates for all years and locations for the primary inputs. Study level covariates used in the prior model were education in years per capita, geological covariates (for mining models), the proportion of the population living with access to a coastline (for fishing models), the IHME socio-demographic index (SDI), the mean temperature/latitude (for agriculture models), and the proportion of the population living in urban areas. Space-time parameters were chosen by maximising out-of-sample cross-validation and minimising RMSE. For economic activity and occupation proportions, estimates from ST-GPR were then re-scaled to sum to 1 across categories by dividing each estimate by the sum of all the estimates.

The following sections describe the modelling approaches for each occupational risk's exposure prevalence.

Occupational carcinogens, occupational noise, and occupational particulates

Prevalence of exposure to these risks was determined using the following equation:

 $Prevalence \ of \ Exposure_{c,y,s,a,r,l} = \sum_{EA} Proportion_{EA,c,y} * EAP_{c,y,s,a} * Exposure \ rate_{EA,r,l,d}$

where:

EAP = economically active population	c = country	r = risk
EA = economic activity	d = duration	s = sex
a = age	I = level of exposure	y =year

Exposure rate was provided by expert group recommendations and literature⁸⁻¹¹ (see table 1). The CAREX database was used in order to quantify the association between exposure by industry/carcinogen to SDI across all the countries in the database. This effect was used to predict exposure in countries that were not included in CAREX. Duration was considered for occupational carcinogens through application of occupational turnover factors¹² and for occupational noise and particulates by calculating cumulative exposure as the average exposure over the lifetime (the past 50 years) for each age/sex cohort.

Occupational ergonomic factors and occupational asthmagens

Prevalence of exposure to these risks was determined using the following equation:

$$\begin{array}{l} Prevalence \ of \ Exposure_{c,y,s,a,r} = \sum_{EA} Proportion_{OCC,c,y} * EAP_{c,y,s,a} \\ \\ \text{where:} \\ \text{EAP} = \text{economically active population} \quad c = \text{country} \\ \text{OCC} = \text{occupation} \quad a = \text{age} \quad s = \text{sex} \\ & y = y \text{ear} \end{array}$$

Occupational injuries

Occupational injury counts were estimated using the following equation:

Occupational fatal injuries_{c,y,a,s}

$$= \sum_{EA} Injury \ rate_{EA,c,y,s} * Population_{c,y,a,s} * EAP_{c,y,s,a} * Proportion_{EA,c,y}$$

where:

EAP = economically active population	c = country	y = year
EA = economic activity	a = age	s = sex

Occupational asbestos

Prevalence of exposure to asbestos was estimated using the asbestos impact ratio (AIR), which is equivalent to the excess deaths due to mesothelioma observed in a population divided by excess deaths due to mesothelioma in a population heavily exposed to asbestos. Formally, this is defined using the following equation:

$$AIR = \frac{Mort_{c,y,s} - N_{c,y,s}}{Mort_{c,ys,}^* - N_{c,y,s}}$$

where:

Mort = Mortality rate due to mesotheliomac = countryMort* = Mortality rate due to mesothelioma iny = yearpopulation highly exposed to asbestoss = sexN = Mortality rate due to mesothelioma inpopulation not exposed to asbestos

Mortality rate due to mesothelioma was estimated from GBD 2017 causes of death.⁷ Mortality rate due to mesothelioma in populations not exposed to asbestos was calculated using the model in Lin et al.,¹³ while the mortality rate due to high exposure to asbestos was estimated in Goodman et al.¹⁴ Asbestos exposure prevalence created using the AIR was used to estimate PAFs for all asbestos-associated causes except for mesothelioma. Custom PAFs were calculated for mesothelioma by using the ratio of the excess mortality with respect to an unexposed population (Mort – N) divided by the mortality rate in the population in question (Mort). This calculation assumes that all mesothelioma is a product of occupational asbestos exposure and could potentially over-estimate burden due to occupational asbestos exposure.

Theoretical minimum-risk exposure level

For all occupational risks, with the exception of occupational asbestos, the theoretical minimum-risk exposure level was assumed to be no exposure to that risk.

Relative risk

Relative risks were obtained for all occupational risks by conducting a systematic review of published meta-analysis. The estimates used, as well as the associated studies, are reported by category group in appendix table 5.

PAFs

For all occupational risks, with the exception of injuries (outlined below) and mesothelioma (outlined above), PAFs were calculated using the prevalences estimated above, using the PAF formula in outlined in the GBD 2017 methods appendix.

Occupational injuries PAF

The PAFs for occupational injuries were calculated using the following formula:

$$PAF_{c,y,a,s} = \frac{Occupational\ fatal\ injuries_{c,y,a,s} - TMREL}{Fatal\ injuries_{c,y,a,s}}$$

where:

c = country	a = age
y = year	s = sex

Fatal injury totals were obtained from GBD 2017 causes of death.⁷

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Suboptimal Breastfeeding Capstone Appendix

Flowchart



Definitions

Exposure to suboptimal breastfeeding is composed of 2 distinct categories: non-exclusive breastfeeding and discontinued breastfeeding.

Non-exclusive breastfeeding is defined as the proportion of children under 6 months of age who are not exclusively breastfed. We then parse those not exclusively breastfed into 3 categories – predominant, partial, and no breastfeeding. Exclusive breastfeeding is defined as the proportion of children who receive no other food or drink except breast milk (allowing for ORS, drops, or syrups containing vitamins, minerals, or medicines). Predominant breastfeeding is the proportion of children whose predominant source of nourishment is breastmilk but also receive other liquids. Partial breastfeeding refers to those infants who receive breastmilk as well as food and liquids, including non-human milk and formula. No breastfeeding refers to infants who do not receive breast milk as a source of nourishment.

Discontinued breastfeeding is defined as the proportion of children between 6 to 23 months who receive no breast milk as a source of nourishment.

Input Data

We made substantial exposure data updates for GBD 2017, including extracting identified surveys not included in previous rounds and re-extracting all surveys for new GBD 2017 subnational locations. We searched the Global Health Data Exchange (GHDx) database for sources using the keyword "Breastfeeding." Of 2,026 potential sources identified, we extracted 1,081 unique country-years of data (2,262 unique geography-years, including subnational geographies) that met our inclusion criteria. The

data used in the analysis consists mostly of processed individual-level microdata from surveys; in the cases where microdata was unavailable, we used reported tabulated data from survey reports and scientific literature. Data used to categorize type of non-exclusive breastfeeding (predominant, partial, and none) come from surveys with 24-hour dietary logs based on maternal recall.

Exposure Modelling

Using the processed microdata and tabulated data from reports, we generated a complete time series from 1980 to 2017 for the prevalence of breastfeeding patterns for children 0 to 5 months and 6 to 23 months using a three-step spatio-temporal Gaussian process regression modelling process.

First, we estimated a robust linear regression using each geography's sociodemographic index as a covariate. The following linear model was used for the estimation of breastfeeding indicators:

$$logit(P_{x,c,t}) = \beta_0 + \beta_1 SDI_{c,t} + \alpha_c + \gamma_{R[c]} + \omega_{SR[c]} + \varepsilon_{c,t}$$

where $P_{x,c,t}$ is prevalence for breastfeeding category x in country c and year t; $SDI_{c,t}$ is value of the Sociodemographic Index for country c and year t; α_c , $\gamma_{R[c]}$, and $\omega_{SR[c]}$ are country, region, and superregion random intercepts, respectively.

We then followed this with a spatio-temporal regression that uses the residuals of the predictions from the linear regression to perform a locally-weighted regression that provides a greater weighting factor to those nearer in space and time. The predicted residuals from this step are then added to those created in the linear regression step.

Finally, we run a Gaussian process regression that incorporates the variance of the input data as well as the variance of the model predictions. It uses predictions from the spatio-temporal regression as the mean function and generates draws from a multinomial distribution (based on the data uncertainty in the prior) to generate the final prevalence estimates and their confidence intervals.

We estimated six models to produce each of our categories: the proportion of currently breastfeeding infants 0-5 months of age, the ratio of infants exclusively breastfed to breastfed infants 0-5 months of age, the ratio of infants predominantly breastfed to breastfed infants 0-5 months of age, the ratio of infants partially breastfed to breastfed infants 0-5 months of age, the ratio of currently breastfeed to breastfed infants 0-5 months of age, the ratio of currently breastfeed infants 6-11 months of age, and the proportion of currently breastfeeding infants 12-23 months of age. We convert the ratios of exclusive, predominant, and partial breastfeeding to the total category prevalence proportions by multiplying each ratio by the estimates of any breastfeeding 0-5 months" envelope. We calculate the proportion of infants receiving no breastmilk 0-5 months of age by subtracting the estimates of current breastfeeding from 1. We perform the same operation to estimate discontinued breastfeeding in the 6-11 months and 12-23 months categories.

Estimating Attributable Burden

Assessment of risk-outcome pairs

We included outcomes based on the strength of available evidence supporting a causal relationship. Studies evaluating the causal evidence for our risk-outcome pairs came primarily from articles found in a review published by the World Health Organization.¹ Non-exclusive breastfeeding was paired with diarrhea and lower-respiratory infection as diseases outcomes. Discontinued breastfeeding was paired with diarrhea as an outcome.

Theoretical minimum-risk exposure level

For non-exclusive breastfeeding, those children that received no source of nourishment other than breastmilk ("exclusively breastfed") were considered to be at the lowest risk of any of the disease outcomes. For discontinued breastfeeding, we assumed that children aged 6 to 23 months who received any breastmilk as a source of nourishment to be at the lowest risk of disease outcome.

Relative Risks

We estimate relative risks for both non-exclusive and discontinued breastfeeding in a meta-analysis using relative risks from studies compiled in a published review by the World Health Organization.¹

Population Attributable Fraction

We use the standard GBD population attributable fraction (PAF) equation to calculate PAFs for nonexclusive breastfeeding and discontinued breastfeeding and each of their paired outcomes using exposure estimates and relative risks.

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Child Growth Failure Capstone Appendix

Flowchart



Input data & methodological summary

Exposure

Case definition

Child growth failure is estimated using three indicators, stunting, wasting, and underweight, all of which all of which are based on categorical definitions using the WHO 2006 growth standards for children 0-59 months. Definitions are based on Z scores from the growth standards, which were derived from an international reference population. Mild, moderate, and severe categorical prevalences were estimated for each of the three indicators.

Input data

There are three main inputs for the GBD child growth failure models: microdata from population surveys and tabulated data from reports, published literature, and the WHO Global Database on Child Growth and Malnutrition.¹ The primary data additions in GBD 2017 for child growth failure were from population surveys that include anthropometry. Population surveys include a variety of multi-country and countryspecific survey series such as Multiple Indicator Cluster Surveys (MICS), Demographic and Health Surveys (DHS), Living Standards Measurement Surveys (LSMS), and the China Health and Nutrition Survey (CHNS), as well as other one time country specific surveys such as the Indonesia Family Life Survey and the Brazil National Demographic and Health Survey of Children and Women. These microdata contain information about each individual child's age (from which age in weeks and age in months are calculated), as well as height and/or weight. From that information, a height-for-age z-score (HAZ), weight-for-age z-score (WAZ), and weight-for-height z-score (WHZ) are calculated using the WHO 2006 Child Growth Standards and the LMS method.²

All available data from the WHO Global Database on Child Growth and Malnutrition was extracted for GBD 2016 – much of which is from published studies. Exclusions included examination date prior to 1985, non-population representative studies, and those based on self-report. A systematic literature review was last completed in GBD 2010. We looked for four metrics from all sources with tabulated data: mean Z score, prevalence <-1 Z score (mild), prevalence <-2 Z score (moderate), and prevalence <-3 Z score (severe). All data for each metric was extracted for each of stunting (height-for-age Z score; HAZ), wasting (weight-for-height Z score; WHZ), and underweight (weight-for-age Z score; WAZ).

To maximise internal-consistency and comprehensiveness of the modelling dataset, we performed three data transformations. First, any data that were reported using the National Center for Health Statistics (NCHS) 1978 growth standards were crosswalked to corresponding values on the WHO 2006 Growth Standards curves based on a study that evaluated growth standard concordance.³ Crosswalks from 1978 to 2006 growth standards were performed only on <-2 (i.e. moderate) prevalence data as that is where the concordance was most consistent. Second, for any study that lacked a measure of mean Z score for any of stunting, wasting, or underweight, we predicted a mean value for that study based on an ordinary-least squares regression of mean Z score versus <-2 prevalence for that metric from all sources where both were available. Third, any data that was presented as both sexes combined or for 0-59 months combined, we used the age and sex pattern from all data sources that included that detail to split into corresponding and age- and sex-specific data. All data was uploaded to a database and all inputs are catalogued in the Global Health Data Exchange (http://ghdx.healthdata.org). A representative dataset coverage map for moderate stunting is shown below.



Figure 1: Number of data points in moderate stunting (<-2 HAZ) in males, 1990 to 2017

Modelling strategy

Exposure estimation

The following three-step modelling process was applied to each of stunting, wasting, and underweight.

First, all microdata was fit using an ensemble modelling process, a modelling framework developed for GBD 2016 that is described elsewhere in this appendix. A series of 12 individual distributions (normal, log normal, log logistic, exponential, gamma, mirror gamma, inverse gamma, gumbel, mirror gumbel, Weibull, inverse Weibull, and beta) were fit to the entire set of microdata (approximately 2.5 million individual z-scores) at the individual survey level. A weighting algorithm combined each distribution to find the optimal combination of these distributions for each survey, minimising the absolute prediction error across the entire distribution. Ensemble weights for each survey were then averaged across all surveys to produce a single set of global weights of the ensemble distributions. Weights were different for each sex, but invariant across geography, time, and age group. All component distributions that were used to derive weights were parameterised using "method of moments," meaning that each corresponding probability density function (PDF) could be described as a function of the mean and variance of the quantity of interest.

Second, models were developed for mean Z scores and prevalence of moderate and severe growth failure. Individual level microdata were collapsed to calculate three metrics: mean z-score, moderate prevalence, and severe prevalence. These data were combined with that derived from literature, GHDx review, and the WHO Global Database on Child Growth and Malnutrition. Each of the three metrics was then modelled using spatiotemporal Gaussian process regression (ST-GPR), a common modelling framework used across GBD, generating estimates for each age-group, sex, year, and location. Location-level covariates used in all models included Socio-demographic Index (SDI) and logit-transformed proportion of households with improved sanitation.

Third, we combined estimates of mean, prevalence (moderate and severe) with ensemble weights in an optimisation framework in order to derive the variance that would best correspond to the predicted mean and prevalence. This variance was then paired with the mean and, using the method of moments equation for each of the component distributions of the ensemble, PDF of the distribution of Z-scores were calculated for each location, year, age-group, and sex. PDFs were integrated to determine the prevalence between -1 and -2 Z scores (mild), between -2 and -3 Z scores (moderate), and below -3 Z scores (severe). These were categorical exposures used for subsequent attributable risk analysis.

Ad-hoc data exclusions were limited. In some cases, we identified surveys with evidence of data entry issues (e.g. weights entered in a mixture of pounds and kilograms) that could not be corrected and these data were outliered. We initially ran all models with the complete dataset. Data plausibility inspection began with examination of time trends in stunting. If a given datum was judged to have led to a change in the prevalence of moderate stunting in 1-4 year olds of 50% or greater in 5 years or fewer, and was inconsistent with data prior to and after that year (a change considered implausible), we outliered the offending datum and reran the model. We then further visually-inspected the results of moderate stunting, wasting, and underweight in parallel to look for location-age-sex-years where the results were not internally-consistent (e.g. stunting and wasting decreasing, underweight rapidly increasing). This inspection revealed very few inconsistent data.

Improvements from GBD 2015 to GBD 2016/ 2017

In GBD 2017, the primary changes from GBD 2016 were the 1) addition of a significant volume of new survey data, 2) crosswalking instead of down-weighting data based on NCHS 1978 growth standard, 3) utilisation of updated versions of location-level covariates, and 4) utilisation of an updated version of the ST-GPR modelling framework that empirically derives many of the modelling parameters.

There are several important differences from the GBD 2015 analysis. First, our systematic data searching efforts led to an approximately 30% increase in the number of data sources since GBD 2015, including a significant increase in data sources for Oceania, Latin America, and South Asia. Most notable was the increase in data for India through our collaboration with the India Council for Medical Research (ICMR) and Public Health Foundation of India (PHFI). Second, while GBD 2015 also used ST-GPR to model growth failure, models were completed for a single 0-5 age group, followed by application of a pooled uniform age-sex split which resulted in the implicit assumption that the age pattern of growth failure is invariant over time and geography. GBD 2016 estimates, owing to smaller sample sizes in younger age groups, do have wider uncertainty in those age groups. Third, GBD 2015, like all analyses of growth failure before it, assumption in GBD 2016 as it is not accurate and instead made explicit estimates of growth failure in all locations. Fourth, GBD 2015 did not use an ensemble approach or estimate the entire distribution of Z scores. Fifth, we changed the name of this risk factor category changed from childhood undernutrition that are covered by the three component indicators.

Theoretical minimum-risk exposure level

Theoretical minimum risk exposure level (TMREL) for underweight, stunting, and wasting was assigned to be greater than or equal to -1 SD of the WHO 2006 standard weight-for-age, height-for-age, and weight-for-height curves respectively. This has not changed since GBD 2010.

Relative risks

The final list of outcomes paired with child growth failure risks included lower respiratory infections (LRI), diarrhea, measles, and protein energy malnutrition (PEM) as shown in Table 1. These were derived from a pooled cohort analysis by Olofin and colleagues.⁵

There is a high degree of correlation between stunting, wasting, and underweight. Failing to account for their covariance and assuming independence would overestimate the total burden significantly. This is the main reason that GBD 2010 only included childhood underweight. In GBD 2013, a method was developed to adjust observed RRs of Olofin and colleagues by simulating the joint distribution of the three indicators using the distribution of each indicator and covariance between indicators in the countries included in the meta-analysis (extracted from Demographic and Health Survey (DHS) micro-data).⁴ Based on the analysis done by McDonald and colleagues, we assumed there is an interaction between the three indicators, and extracted the interaction terms from the corresponding analysis. We calculated the adjusted RRs by minimising the error between observed crude RRs (from meta-analysis) and expected crude RRs derived from adjusted RRs.

Of historical note, URI and otitis media were included as outcomes in the GBD 2013 risk analysis, based on the "analogy" causal criterion, assuming there is similar pathway as LRI outcome. However, closer review for GBD 2015 did not find sufficient evidence to support their inclusion and they were excluded, a decision that was carried forward into GBD 2016. We also attributed 100% of PEM to childhood wasting and underweight but not stunting. To build on the existing literature base for GBD on risk-outcome pairs, a literature search was conducted for GBD 2017 searching for case-control studies published after January 1st, 1985; this search did not return any sources that were appropriate for this work.

Outcome	Stunting	Wasting	Underweight
	<-1: 1.111 (1.023-1.273)	<-1: 6.601 (2.158-11.243)	<-1: 1.088 (1.046-1.134)
Diarrhea	<-2: 1.222 (1.067-1.5)	<-2: 23.261 (9.02-35.845)	<-2: 1.23 (1.163-1.314)
	<-3: 1.851 (1.28-2.699)	<-3: 105.759 (42.198-157.813)	<-3: 2.332 (2.076-2.802)
Lower respiratory	<-1: 1.125 (0.998-1.655)	<-1: 5.941 (1.972-11.992)	<-1: 1.145 (1.044-1.364)
infoctions (LPI)	<-2: 1.318 (1.014-2.165)	<-2: 20.455 (70.84-37.929)	<-2: 1.365 (1.215-1.755)
Infections (LKI)	<-3: 2.355 (1.15-5.114)	<-3: 47.67 (15.923-94.874)	<-3: 2.593 (1.908-4.39)
	<-1: 1.103 (0.861-1.719)	<-1: 1.833 (0.569-8.965)	<-1: 0.995 (0.5-1.726)
Measles	<-2: 1.54 (1.029-3.222)	<-2: 8.477 (1.33-42.777)	<-2: 2.458 (1.26-5.118)
	<-3: 2.487 (1.129-6.528)	<-3: 37.936 (5.088-199.126)	<-3: 5.668 (1.767-12.414)
Protein-energy malnutrition	0% PAF	100% PAF	100% PAF

References

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- 5 Olofin I, McDonald CM, Ezzati M, *et al.* Associations of Suboptimal Growth with All-Cause and Cause-Specific Mortality in Children under Five Years: A Pooled Analysis of Ten Prospective Studies. *PLOS ONE* 2013; **8**: e64636.

Low Birth Weight and Short Gestation Capstone Appendix

Flowchart



Low birth weight and Short gestation Risk Factors

Input data and methodological summary

The "Low Birth Weight and Short Gestation" (LBWSG) risk factor and its child risks "Low Birth Weight for Gestation" and "Short Gestation for Birth Weight" first were included as risk factors in GBD 2016.

Although low birth weight for gestation and short gestation for birth weight are separate risk factors, the exposures and relative risks for both are estimated jointly through the low birth weight and short gestation parent risk factor. As of GBD 2017, LBWSG are the only risk factors estimated jointly.

Case definition

The meaning of the "low birth weight" and "short gestation" in GBD have subtle definitional differences compared to other usages of "low birth weight" and "short gestation" in literature. The term "low birth weight" has historically been used to refer to birth weight (BW) less than 2500 grams. However, because the goal of the GBD risk factors analysis is to quantify the entirety of attributable burden due to each risk factor, the GBD definition of "low birth weight" therefore refers to all birth weight below the Theoretical Minimum Risk Exposure Level (TMREL) for birth weight. Likewise, new-borns have been typically been classified into gestational age (GA) categories of "extremely preterm" (32-<37 weeks of gestation), "very preterm" (28-<32 weeks of gestation), and "moderate to late preterm" (32-<37 weeks of gestation). "Short gestation" in GBD refers to all gestational ages below the gestational age TMREL.

Exposures and relative risks for the GBD Low birth weight and short gestation risk factors are divided into joint 500-gram birth weight and 2-week gestational age combinations. The lowest risk overall 500-gram/2-week bin is the overall TMREL. The univariate TMRELs vary with GA and BW. The lowest risk GA varies by BW category and the lowest risk BWs vary with GA category. The latter are used to quantify univariate attributable risk. Under this framework, all attributable burden under the joint TMREL is referred to jointly as burden of LBWSG. All attributable burden to BWs under the TMREL for each GA category are, on aggregate, "low birth weight" and all attributable burden to GAs under the TMREL for each BW category are, on aggregate, "short gestation." Each combination of 500-grams and 2-wks is associated with a relative risk for mortality by neonatal period (early and late neonatal) and by the causes listed in Table 2 and described below, and relative to the joint TMREL.

Exposure

Input data

To model the joint distribution of exposure of low birth weight and short gestation for each location, year, and sex estimated in GBD 2017, three types of information are used:

- Distribution of gestational age for each location, year, and sex
- Distribution of birth weight for each location, year, and sex
- Copula family and parameters, specifying correlation between gestational age and birth weight distributions

Modelling strategy

Distributions of birth weight & gestational age

To model the joint distribution of birth weight and gestational age for every location-sex-year, ensemble model methods standard to GBD risk factors (described elsewhere in the methods appendix), are first used to create separate distributions of birth weight and gestational age for every location-sex-year.

Microdata is the most ideal data source for modelling distributions; however, microdata is not widely available for birth weight and is more scarce for gestational age. Categorical prevalence data is much more readily available, and from a wider range of locations and years, for low birth weight (<2500g), extremely preterm (<28 weeks of gestation), very preterm (28-32 weeks of gestation), moderate to late preterm (32-37 weeks of gestation), and preterm birth (<37 weeks of gestation). From GBD 2010 to GBD 2015, this categorical data has been used model birth prevalence of preterm birth by gestational age (<28 weeks, 28-<32 weeks, and 32-<37 weeks) and low birth weight (<2500g) for every location, sex, and year estimated in GBD. Starting in GBD 2016 with the introduction of the LBWSG risk factors, the full distributions at birth have been modelled for gestational age and birth weight for all GBD locations, estimation years, and both sexes. The gestational age and birth weight distributions are then aggregated into the categorical estimates of <28 weeks, 28-<32 weeks, 32-<37 weeks gestation, and <2500 g birth weight.

Ensemble model methods standard to GBD are used to model the distribution at birth of gestational age and birth weight. Gestational age ensemble distribution models use the prevalence of <37 weeks gestation, the prevalence of <28 weeks gestation, and mean gestational age per each location-year-sex as inputs into the model. Birth weight distribution models use the prevalence of <2500 grams birth weight and mean birth weight per each location-year-sex. Prevalence of <37 weeks gestation and of <2500 grams birth weight was estimated for all location-year-sexes using STGPR modelling processes standard to GBD.

Low birth weight (<2500 grams) data was extracted from literature, vital registration systems, and surveys. DHS survey data were observed to have high missingness; to correct for the missingness, birth weight was imputed using the Amelia package in R. Birth weight was predicted using standard Amelia imputation methods from the following variables also in the DHS surveys: urbanicity, sex, birthweight recorded on card, birth order, maternal education, paternal education, child age, child weight, child height, mother's age at birth, mother's weight, shared toilet facility, and household water treated. Data counts for categorical prevalence models are listed in Table 1.

	<28 weeks	<37 weeks	<2500 grams
Site-years (total)	1872	2420	2980
Number of GBD regions with data (out of 21 regions)	14	21	21
Number of GBD super-regions with data (out of 7 super-regions)	6	7	7

Table 1: Data Counts for Categorical Prevalence Models

Global ensemble weights for gestational age were derived by using a 3 million sample of all available microdata in Table 2 to select the ensemble weights. Of the exponential, gamma, inverse gamma, Weibull, log normal, and normal distributions, the three distribution families that received the highest weights were the Weibull (87%), normal (4%), and inverse gamma (4%) distributions. Global ensemble weights for birth weight were derived using a 3 million sample of all available microdata in Table 2, in addition to birth weight microdata available primarily through the DHS and MICS surveys. Of the exponential, gamma, inverse gamma, Weibull, log normal, and normal distributions, the three

distribution families that received the highest weights were the log normal (38%), normal (32%), and Weibull (20%) distributions.

Ordinary least squares was used to model mean gestational age for all location-year-sexes by regressing mean gestational age on prevalence of <37 weeks gestation per location-year. All available microdata (Table 2) was used to fit the model. OLS was also used to model mean birth weight by regressing prevalence of <2500 g birth weight per location-year. All available joint microdata (Table 2), as well as additional birth weight microdata extracted primarily through DHS and MICS surveys, was used to fit the model. As estimates of prevalence of <37 weeks gestation and prevalence of <2500g birth weight are available for all location-year-sexes through STGPR models, mean gestational age and mean birth weight were predicted for all location-year-sexes.

Copula optimisation

In order to model the joint distribution of gestational age and birth weight from separate distributions, information is needed about the correlation between the two distributions. Distributions of gestational age and birth weight are not independent; the Spearman correlation for each country where joint microdata was available (Table 2), pooling across all years of data available, ranged from 0.25-0.49. The overall Spearman correlation was 0.38, pooling across all countries in the dataset.

Location	Years of data	Total births*	Format of data	Spearman correlation	Used in Ensemble Weight Selection	Used in Copula Parameter Selection	Used in Relative Risk Models
BRA	2016	2,854,380	Microdata	0.37	Yes	Yes	No
ECU	2003-2015	2,473,039	Microdata	0.34	Yes	Yes	No
ESP	1990-2014	8,537,220	Microdata	0.42	Yes	Yes	No
JPN	1995-2015	23,644,506	Tabulations	0.41	No	No	Yes
MEX	2008-2012	10,256,117	Microdata	0.35	Yes	Yes	No
NOR	1990-2014	1,489,210	Microdata	0.44	Yes	Yes	Yes
NZL	1990-2016	1,600,501	Microdata	0.25	Yes	Yes	Yes
SGP	1993-2015	972,775	Tabulations	0.41	No	No	Yes
TWN	1998-2002	1,331,760	Tabulations	0.38	No	No	Yes
URY	1996-2014	698,622	Microdata	0.49	Yes	Yes	No
USA	1990-2014	81,929,879	Microdata	0.38	Yes	Yes	Yes

Table 3: Summary of Data Inputs

* Pooled across all year and sexes, excluding data missing year of birth, gestational age, or birth weight

Copula modelling is used to model joint distributions between the birth weight and gestational age marginal distributions. The Copula and VineCopula packages in R were used to select the optimal copula family and copula parameters to model the joint distribution, using joint microdata from the country-years in Table 2. The copula family selected from the microdata was "Survival BB8", with theta parameter set to 1.75 and delta parameter set to 1.

The joint distribution of birth weight and gestational age per location-year-sex was modelled using the global copula family and parameters selected and the location-year-sex gestational age and birth weight distributions. The joint distribution was simulated 100 times to capture uncertainty. Each simulation consisted of 100,000 simulated joint birth weight and gestational age data points. Each joint distribution

was divided into 500g by 2wk bins to match the categorical bins of the relative risk surface. Birth prevalence was then calculated for each 500g by 2wk bin.

Estimating Early Neonatal Prevalence & Late Neonatal Prevalence from Birth Cohorts

Early neonatal prevalence and late neonatal prevalence was estimated using life table approaches for each 500g & 2wk bin. Using the all-cause early neonatal mortality rate for each location-year-sex, births per location-year-sex-bin, and the relative risks for each location-year-sex-bin in the early neonatal period, the all-cause early neonatal mortality rate was calculated for each location-year-sex-bin. The early neonatal mortality rate per bin was used to calculate the number of survivors at 7 days and prevalence in the early neonatal period. Using the same process, the all-cause late neonatal mortality rate for each location-year-sex was paired with the number of survivors at 7 days and late neonatal relative risks per bin to calculate late neonatal prevalence and survivors at 28 days.

Relative risks & theoretical minimum-risk exposure level

Causes

The available data for deriving relative risk was only for all-cause mortality. The exception was the USA linked infant birth-death cohort data, which contained 3-digit ICD causes of death, but also had nearly 30% of deaths coded to causes that are ill-defined, or intermediate, in the GBD cause classification system. For GBD 2017, like in GBD 2016, we analysed the relative risk of all-cause mortality across all available sources and selected outcomes based on criteria of biologic plausibility. Some causes, most notably congenital birth defects, haemoglobinopathies, malaria, and HIV/AIDS, were excluded based on the criteria that reverse causality could not be excluded. The final list of outcomes included in calculating the attributable burden for LBW/SG are in Table 3.

radic 3. cause list of batcomes for low birth weight and short gestation	Table 3: Cause list o	f outcomes	for low birth	weight and s	hort gestation
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Cause ID	Cause name
302	Diarrheal diseases
322	Lower respiratory infections
328	Upper respiratory infections
329	Otitis media
333	Pneumococcal meningitis
334	H influenzae type B meningitis
335	Meningococcal meningitis
336	Other meningitis
337	Encephalitis
381	Neonatal preterm birth complications
382	Neonatal encephalopathy due to birth asphyxia and trauma
383	Neonatal sepsis and other neonatal infections
384	Hemolytic disease and other neonatal jaundice
385	Other neonatal disorders
686	Sudden infant death syndrome

Input data

In the Norway, New Zealand, and US Linked Birth/Death Cohort microdata datasets, live births are reported with gestational age, birth weight, and an indicator of death at 7 days and 28 days. For this analysis, gestational age was grouped into two-week categories, and birth weight was grouped into 500-gram categories. The Taiwan, Japan, and Singapore datasets were prepared in tabulations of joint 500-gram and two-week categories.

Modelling strategy

For each location, data was pooled across years, and the risk of all-cause mortality at the early neonatal period and late neonatal period at joint birth weight and gestational age combinations was calculated. In all datasets except for the United States, sex-specific data were combined to maximise sample size. The United States analyses were sex-specific. To calculate relative risk at each 500g and 2wk combination, logistic regression was first used to calculate mortality odds for each joint 2-week gestational age and 500-gram birth weight category. Mortality odds were smoothed with Gaussian Process Regression, with the independent distributions of mortality odds by birth weight and mortality odds by gestational age serving as priors in the regression.

A pooled country analysis¹ of mortality risk in the early neonatal period and late neonatal period by SGA category in developing countries in Asia and Sub-Saharan Africa were also converted into 500-gram and 2-week bin mortality odds surfaces. The relative risk surfaces produced from microdata and the Asia and Africa surfaces produced from the pooled country analysis were meta-analyzed, resulting in a meta-analysed mortality odds surface for each location. The meta-analysed mortality odds surface for each location was smoothed using Gaussian Process Regression and then converted into mortality risk. To calculate mortality relative risks, the risk of each joint 2-week gestational age and 500-gram birth weight category were divided by the risk of mortality in the joint gestational age and birth weight category with the lowest mortality risk.

For each of the country-derived relative risk surfaces, the 500 g and 2-week gestational age joint bin with the lowest risk was identified. This bin differed within each country dataset. To identify the universal 500 g and 2-week gestational age category that would serve as the universal TMREL for our analysis, we chose the bins that was identified to be the TMREL in each country dataset to contribute to the universal TMREL. Therefore, the joint categories that served as our universal TMREL for the LBWSG risk factor were "38-40 weeks of gestation and 3500-4000 grams", "38-40 weeks of gestation and 4000-4500 grams". As the joint TMREL, all three categories were assigned to a relative risk equal to 1.

PAF calculations

The total PAF for the low birth weight and short gestation joint risk factor is calculated by summing the PAF calculated from each 500g x 2wk category, with the lowest risk category among all the 500g x 2wk categories serving as the TMREL. The equation for calculating PAF for each 500g x 2wk category is:

$$PAF_{joasgt} = \frac{\sum_{x=1}^{u} RR_{joast}(x)P_{jasgt}(x) - RR_{joasg}(TMRE_{jas})}{\sum_{x=1}^{u} RR_{joas}(x)P_{jasgt}(x)}$$

To calculate the overall PAF for the short gestation for birth weight risk factor, PAF was once again calculated for each joint 500-gram and 2-week category. Unlike the joint PAF calculation, which used only one TMREL for all 500-gram and 2-week categories, the joint 500-gram and 2-week category with the lowest risk for each 500-gram birth weight grouping served as the TMREL for that 500-gram birth weight grouping. For example, the [3000, 3500) grams; [36, 37) weeks and [3000, 3500) grams; [37, 38) weeks and [3000, 3500) grams; [38, 40) weeks and [3000, 3500) grams; and [40, 42) weeks and [3000, 3500) grams joint category has the lowest risk, and so it serves as the TMREL for the [3000, 3500) gram birth weight grouping. In the Relative Risk surface figures, a birth weight grouping is one "column" of the birth weight and gestational age matrix.

The overall PAF for the short gestation for birth weight risk factor was then calculated for all the joint 500-gram and 2-week categories using the formula below:

$$PAF_{1..i} = 1 - \prod_{i=1}^{n} (1 - PAF_i)$$

The same methodology was applied to calculate the total PAF for the low birth weight for gestation risk factor, using two-week gestational age categories (each "row" of the matrix) instead of 500-gram birth weight categories. For example, the [24, 26) weeks gestational age grouping contains three joint categories: [0, 500) grams and [24, 26) weeks; [500, 1000) grams and [24, 26) weeks; and [1000, 1500) grams and [24, 26) weeks. The [1000, 1500) grams and [24, 26) weeks gestational age grouping.

After the short gestation for birth weight PAF and low birth weight for gestational age PAF were calculated, they were then scaled so that the sum of the short gestation for birth weight PAF and low birth weight for gestation PAF equal the low birth weight and short gestation parent PAF calculated for each location/year/sex/age group.

References

1. Katz et al. Mortality risk in preterm and small-for-gestational-age infants in low-income and middleincome countries: a pooled country analysis. 2013. Volume 382, Issue 9890. *The Lancet*.

Iron Deficiency Capstone Appendix



Flowchart

Input data & methodological summary

Exposure

Case definition

For GBD 2017, as with GBD 2016, the anemia model has two main steps: estimation of the anemia envelope and causal attribution. Our analytic strategy began with calculation of an anemia envelope – a determination of mean hemoglobin, as well as a sum total of anemia prevalence, by severity for each country, age group, and both sexes for each year from 1990 through 2017. The envelope approach avoids double-counting while capturing potentially different disease profiles within each population group. We defined a population group as a specific geography, sex, age-group, and year.

Input data

Iron-deficiency anemia (IDA) estimates include acute and chronic hemorrhagic states for which supplementation may be helpful, but poor nutritional intake is not the only underlying problem. A few causes in this category – hookworm, schistosomiasis, upper gastrointestinal bleeding, and gynecologic diseases – were considered separately from IDA in GBD 2016 because there was enough data from GBD prevalence estimation processes to do so. Distribution of anemia burden to IDA only after assignment to "known" causes avoided double counting of these cases. In GBD 2017, we redefined the iron deficiency risk factor to encompass all causes of anemia that would respond to iron supplementation.

For our nonfatal anemia estimates, the envelope approach to the anemia impairment utilises data from a variety of sources. Population-based surveys of hemoglobin concentration were the primary input to our analytic dataset. Examples include the Demographic and Health Survey (DHS) and Multiple Indicator Cluster Survey (MICS) series, along with other national and subnational surveys that completed hemoglobin testing. We supplemented with pertinent sources downloaded from the WHO Vitamin and
Mineral Nutrition Information System (VMNIS) available at

<u>http://www.who.int/vmnis/database/anaemia/countries/en/</u>. A full source list is available elsewhere in this appendix. Most used a HemoCue test, adjusted for altitude, and excluded those with terminal or acute medical conditions. Inclusion, exclusion and diagnostic criteria for other studies were similar and can be found in each study.

Modelling strategy

For GBD 2017, we estimated the mean hemoglobin in g/dL among pregnant women aged 15 to 49 years of age and the implied mean hemoglobin among pregnant women in the absence of iron deficiency anemia, as the risk exposure for maternal iron deficiency anemia.

Theoretical minimum-risk exposure level

The implied mean hemoglobin in the absence of iron deficiency anemia is the theoretical minimum risk exposure level. This was calculated by adding the iron responsive hemoglobin shift back onto the observed hemoglobin concentration for each demographic. For example, if the observed hemoglobin concentration among 30-34 year old pregnant women in Ethiopia was 132.9 g/L, and the shift was 1.6 g/L in that demographic, then the counterfactual was 134.5 g/L. The GBD 2017 anemia modelling strategy provides details on how the iron deficiency shifts were calculated.

Relative risk

We attribute 100% of iron-deficiency anemia to iron deficiency. The other outcomes used in GBD 2016 are maternal hemorrhage, maternal sepsis and other maternal infections. In GBD 2017 we added the following additional maternal outcomes: maternal hypertensive disorders, maternal obstructed labor an uterine rupture, maternal abortion and miscarriage, ectopic pregnancy, indirect maternal deaths, late maternal deaths, maternal deaths aggravated by HIV/AIDS, and other maternal disorders. For these additional maternal outcomes, we assigned the same relative risk as that used for the maternal outcomes from 2016. Sources of evidence for these relative risks are unchanged from GBD 2013.

References

- Centers for Disease Control and Prevention (CDC). Iron deficiency--United States, 1999-2000. MMWR Morb Mortal Wkly Rep 2002; 51: 897–9.
- 2. Looker AC, Dallman PR, Carroll MD, Gunter EW, Johnson CL. Prevalence of iron deficiency in the united states. JAMA 1997; 277: 973–6.

Vitamin A Deficiency Capstone Appendix

Flowchart



Input data & methodological summary

Exposure

Case definition

For GBD 2017, vitamin A deficiency is defined as serum retinol <70 µmol/L. We examined vitamin A deficiency as a risk factor in children aged 6 months to 5 years.

To ensure we were using as much information as possible, and therefore maximize the data basis of our estimates, we modeled Vitamin A deficiency sequentially. The first step was to estimate the coverage of Vitamin A supplementation. Although the typical metric on which supplementation is tracked is 2+ doses of Vitamin A in the previous 12 months for children under 5 years, most existing health surveys do not routinely provide sufficient information to calculate it. Our case definition for the supplementation model was therefore the proportion of children 6-59 months of age who received at least one dose of Vitamin A in the previous 6 months. Supplementation estimates were then used as a location-level covariate to guide exposure models of overall Vitamin A deficiency.

Input data

For GBD 2017, we used data from the WHO Vitamin and Mineral Nutrition Information System, health surveys such as DHS and MICS, and studies identified through literature review. This included updating the dataset to include all ages and all studies available in VMNIS as of April 2018. A separate systematic review was last conducted for GBD 2013. The PubMed search terms were: ((vitamin A deficiency[Title/Abstract] AND prevalence[Title/Abstract]) AND ("2009"[Date – Publication] : "2013"[Date – Publication])). The table below shows the number of data points included in the final datasets. Exclusion criteria were:

- 1. Studies that were not population-based, e.g., hospital or clinic-based studies
- 2. Studies that did not provide primary data on epidemiological parameters, e.g., commentaries
- 3. Review articles
- 4. Case series
- 5. Self-reported cases

Table 1. Geographic representation of datasets used for three stages of Vitamin A deficiency risk factor burden estimation (number of data points per geography)

Geography	Supplementation (proportion)	Deficiency (prevalence)
Global	900	1540
East Asia	12	27
Southeast Asia	102	212
Oceania	24	54
Central Asia	51	66
Central Europe	2	13
Eastern Europe		3
Australasia		1
Western Europe		38
Southern Latin America		16
High-income North America		33
Caribbean	17	34
Andean Latin America	25	70
Central Latin America	33	212
Tropical Latin America	1	52
North Africa and Middle East	49	148
South Asia	61	96
Central Sub-Saharan Africa	60	8
Eastern Sub-Saharan Africa	182	220
Southern Sub-Saharan Africa	49	57
Western Sub-Saharan Africa	232	180

Modeling strategy

All Vitamin A deficiency estimates were made using DisMod-MR 2.1. As described above, we first estimated Vitamin A supplementation coverage. Although all data was from ages 6-59 months, we assumed no difference in age pattern of supplementation coverage and used the natural log of lagdistributed income per capita (LN-LDI) as a location-level covariate to inform estimates where data was absent. DHS and MICS data was cross-walked to the reference data source, which came from UNICEF (<u>http://data.worldbank.org/indicator/SN.ITK.VITA.ZS</u>).

Measure	Covariate	Туре	Value	Exponentiated
Dravalanca	MICS	Study loval	-0.6	0.55
Prevalence	IVIICS	Study-level	(-0.76 — -0.45)	(0.47 — 0.64)
Provalanco	טחט	Study loval	-0.09	0.91
Prevalence	UDS	Study-level	(-0.2 — 0.025)	(0.82 — 1.03)
Prevalence	LDI (I\$ per capita)	Country-level	0.013	1.01
			(0.00033 — 0.042)	(1.00 - 1.04)

Table 2: Covariate effects for Vitamin A supplementation model

Second, we estimated the age- and sex-specific prevalence of Vitamin A deficiency (serum retinol < 0.7 μ mol/L). WHO VMNIS was the primary data source for this model and was supplemented with data from DHS and other health surveys where testing was performed. We assumed the following in our model: no excess mortality, birth prevalence is possible, incidence is decreasing after age 5 and remission is increasing after age 5. Data from subnational locations was crosswalked to the reference data sources of nationally-representative data. Females were found to have 1.09 times higher Vitamin A deficiency, although the uncertainty in that ratio ranged from 0.97 to 1.24. Location-level covariates were used for Vitamin A supplementation coverage from the above model as well as GBD 2016 Socio-demographic Index (SDI) numbers.

Measure	Covariate	Туре	Value	Exponentiated
Provalanca	Sov	Study loval (x.cov)	-0.0091	0.99
Flevalence	Sex	Study-level (X-COV)	(-0.088 — 0.065)	(0.92 — 1.07)
Prevalence	Subnational Study-level (x-co	Study loval (x.cov)	-0.28	0.76
		Study-level (X-COV)	(-0.44 — -0.1)	(0.64 — 0.90)
Prevalence	Vitamin A supplem.	Country-level	-0.028	0.97
	coverage rate		(-0.10.00071)	(0.90 - 1.00)
Prevalence	Socio-demographic	Country-level	-2.98	0.051
	Index		(-3 — -2.92)	(0.050 — 0.054)
Prevalence	vegetables unadjusted(g)	Country-level	-1.36	0.26
			(-1.53 — -1.12)	(0.22 — 0.33)

Table 3: Covariate effects for Vitamin A deficiency model

Theoretical minimum-risk exposure level

The theoretical minimum risk exposure is that the prevalence of vitamin A deficiency is zero.

Relative risks

The relative risks were updated in GBD 2017 to reflect studies included in the most recently published systematic review by Imdad and colleagues.¹ The overall estimation strategy has not changed. For each trial identified by the systematic review, we adjusted the relative risk for the background prevalence of Vitamin A deficiency in 1-4 years from the GBD 2017 model described above. This adjustment assumes the effect of supplementation is observed only in the fraction of the trial population that are Vitamin A deficient. Many studies evaluate either incidence or mortality. A subset of studies evaluated both incidence and cause-specific mortality as outcomes for the same cause. We found no statistical difference between the effect sizes of incidence and mortality in any of these studies so pooled all incidence and mortality observations as independent observations prior to meta-analysis. We then performed a fixed effects meta-analysis of all adjusted RRs to determine final outcomes to be included in GBD risk factor attribution estimates. Forest plots are shown in Figures 1-5; Final RRs are shown in Table 4. Three outcomes – diarrhea, lower respiratory infections (LRI), and measles – were found to be statistically significant after adjustment, pooling, and meta-analysis. Meningitis was non-significant. Malaria was significant, but only a single study was identified that evaluated this outcomes, which does not meet GBD causal criteria.

Cause	GBD 2016 RR	GBD 2017 RR	Include in GBD 2017
Diarrhea	1.6 (1.21 - 2.02)	2.35 (2.17 - 2.54)	Yes
Measles	2.4 (1.61 - 3.48)	2.76 (2.01 - 3.78)	Yes
Lower Respiratory Infections (LRI)		1.23 (1.03 - 1.48)	Yes
Meningitis		3.2 (0.69 - 14.75)	No (not significant)
Malaria		3.65 (2.23 - 5.97)	No (only one study)

Table 4: Pooled relative risks for risk-outcome pairs included in GBD 2017

Figure 1: Forest plot of RR of diarrhea in Vitamin A deficiency

Venkatarao 1996 Diarrhea Mortality ~ 909 Agarwal 1995 Diarrhea Mortality ~ 15247 Chowdhury 2002 Diarrhea Mortality ~ 1520 Rahmathullah 1990 Diarrhea Mortality ~ 15419 Fisker 2014 Diarrhea Mortality ~ 7587 Ross (Survival) 1993 Diarrhea Mortality ~ 21906 Herrera 1992 Diarrhea Mortality ~ 28753 Daulaire 1992 Diarrhea Mortality ~ 7197 Awasthi 2013 Diarrhea Mortality ~ 1000000 Arya 2000 Diarrhea Incidence ~ 256 Fisker 2014 Diarrhea Incidence ~ 256 Fisker 2014 Diarrhea Incidence ~ 387 Florentino 1990 Diarrhea Incidence ~ 387 Florentino 1990 Diarrhea Incidence ~ 387 Cheng 1993 Diarrhea Incidence ~ 198 Herrera 1992 Diarrhea Incidence ~ 28753		$\begin{array}{c} 15.66 \ [0.61, \ 399.35] \\ 1.76 \ [0.08, \ 38.56] \\ 8.46 \ [1.26, \ 56.93] \\ 4.89 \ [1.85, \ 12.95] \\ 1.48 \ [0.54, \ 4.08] \\ 1.98 \ [0.89, \ 4.41] \\ 0.95 \ [0.23, \ 3.98] \\ 2.95 \ [1.50, \ 5.82] \\ 1.19 \ [0.85, \ 1.65] \\ 1.21 \ [0.19, \ 7.72] \\ 1.51 \ [0.79, \ 2.89] \\ 2.75 \ [1.17, \ 6.45] \\ 0.83 \ [0.32, \ 2.11] \\ 1.75 \ [0.63, \ 4.88] \\ 9.14 \ [6.08, \ 13.76] \\ 4.43 \ [2.72, \ 7.21] \end{array}$
Long 2007 Diarrhea Incidence ~ 195		1.45 [0.75, 2.79]
Sempertegui 1999 Diarrhea Incidence ~ 400	F∎4	0.59 [0.19, 1.83]
Dibley 1994 Diarrhea Incidence ~ 1405	⊢ ∎+1	0.84 [0.59, 1.19]
Chowdhury 2002 Diarrhea Incidence ~ 1520		3.33 [2.99, 3.71]
Barreto 1994 Diarrhea Incidence ~ 1240	=	1.23 [1.02, 1.48]
FE Model	•	2.35 [2.17, 2.54]
0.02	0.14 1 7.39 54.6 403.43	
	Group 1 Diarrhea method=FE	

Figure 2: Forest plot of RR of measles in Vitamin A deficiency



Figure 3: Forest plot of RR of LRI in Vitamin A deficiency

Chowdhury 2002 LRTI Mortality ~ 1520	⊢ I	30.88 [1.52, 625.52]
Venkatarao 1996 LRTI Mortality ~ 909	⊢	37.54 [1.98, 712.08]
Rahmathullah 1990 LRTI Mortality ~ 15419	F	2.87 [0.18, 46.91]
Fisker 2014 LRTI Mortality ~ 7587	⊢ I	3.34 [0.61, 18.22]
Daulaire 1992 LRTI Mortality ~ 7197	⊢	1.19 [0.15, 9.66]
Herrera 1992 LRTI Mortality ~ 28753	⊢	7.03 [2.11, 23.38]
Ross (Survival) 1993 LRTI Mortality ~ 21906	⊢	1.00 [0.41, 2.46]
Awasthi 2013 LRTI Mortality ~ 1000000	F ≢ 1	0.94 [0.62, 1.43]
Cheng 1993 LRTI Incidence ~ 198	⊧I	17.36 [1.37, 219.83]
Kartasasmita 1995 LRTI Incidence ~ 269	⊢−−−− 1	0.66 [0.09, 4.59]
Biswas 1994 LRTI Incidence ~ 180	I	2.92 [0.85, 10.06]
Rahmathullah 1990 LRTI Incidence ~ 15419	F	0.96 [0.21, 4.39]
Sempertegui 1999 LRTI Incidence ~ 400	⊢	1.30 [0.34, 4.93]
Chen 2013 b LRTI Incidence ~ 387	⊢ _ ∎	1.35 [0.68, 2.70]
Chen 2013 a LRTI Incidence ~ 387	⊢_	0.89 [0.36, 2.19]
Fisker 2014 LRTI Incidence ~ 7587	H -1	1.15 [0.89, 1.48]
FE Model	•	1.23 [1.03, 1.48]
0.02	0.14 1 7.39 54.6 403.43 2980.96	
	Group 3 LRTI method=FE	

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Figure 4: Forest plot of RR of meningitis in Vitamin A deficiency



Figure 5: Forest plot of RR of malaria in Vitamin A deficiency



References

1. Imdad A, Ahmed Z, Bhutta ZA. Vitamin A supplementation for the prevention of morbidity and mortality in infants one to six months of age. *Cochrane Database of Systematic Reviews* 2016; Sep 28; 9. Art. No: CD007480.

Zinc Deficiency Capstone Appendix

Input data & methodological summary

Exposure

Case definition

Exposure to zinc deficiency is defined as consumption of less than 2.5 milligrams of zinc per day among children between the ages of 1 and 4 years old.

Input data

We used dietary data from nationally and sub-nationally representative nutrition surveys and United Nations FAO Supply and Utilization Accounts to estimate the mean intake of zinc at the population level.

Modelling strategy

For GBD 2016, we first used a spatio-temporal Gaussian process regression (ST-GPR) framework to estimate the mean intake of zinc by age, sex, country, and year. To assist with estimation for locations and years without data, we used the lag-distributed income of that location-year as a covariate. We considered data from 24-hour diet recall as the gold standard, and adjusted data from other sources to the gold standard method. Using the method described in the dietary risks section, we characterised the distribution of zinc intake for children between ages of 1 and 4 years old and estimate the proportion of the children with intake of less than 2.5 milligrams of zinc per day.

Relative risk

Relative risks used for zinc deficiency is based on the results of randomised trials that measured the effect of zinc supplementation.

Theoretical minimum-risk exposure level

The theoretical minimum-risk exposure level for proportion zinc deficient is zero percent deficient.

Smoking Capstone Appendix



We made significant changes to the methods used to estimate smoking attributable burden in GBD 2017. In previous iterations of the GBD, we have used the Peto-Lopez (Smoking Impact Ratio) method to estimate burden attributable to cancers and chronic respiratory diseases. Although this method provides robust estimates of the burden of cancers and chronic respiratory diseases related to tobacco, it is not fully consistent with the GBD approach of estimating exposure independently of the outcomes affected by exposure. For cardiovascular diseases and all other smoking attributable health outcomes, we used five-year lagged daily smoking prevalence as the exposure. With a growing body of evidence on the association between smoking and several types of cancers and with cardiovascular disease, coupled with good estimates of the distribution of cumulative smoking exposure, direct estimation of attributable burden is possible. In GBD 2017, we have transitioned to using continuous measures of exposure that incorporate dose-response effects among daily, occasional, and former smokers for all health outcomes except fractures.

Current and former smoking prevalence

We estimated the prevalence of current smoking and the prevalence of former smoking using data from cross-sectional nationally representative household surveys. We defined current smokers as individuals who currently use any smoked tobacco product on a daily or occasional basis. We defined former

smokers as individuals who quit using all smoked tobacco products for at least 6 months, where possible, or according to the definition used by the survey. Prior to modelling a complete time series for all demographic groups, we made adjustments for alternative case definitions as well as for data reported in non-standard age or sex groups. We modelled current and former prevalence using spatiotemporal Gaussian process regression.

Data extraction

We extracted primary data from individual-level microdata and survey report tabulations. We extracted data on current, former, and/or ever smoked tobacco use reported as any combination of frequency of use (daily, occasional, and unspecified, which includes both daily and occasional smokers) and type of smoked tobacco used (all smoked tobacco, cigarettes, hookah, and other smoked tobacco products such as cigars or pipes), resulting in 36 possible combinations. Other variants of tobacco products, for example hand-rolled cigarettes, were grouped into the four type categories listed above based on product similarities. Only smoked tobacco products are included, smoked drugs are estimated separately as part of the drug use risk factor.

For microdata, we extracted relevant demographic information, including age, sex, location, and year, as well as survey metadata, including survey weights, primary sampling units, and strata. This information allowed us to tabulate individual-level data in the standard GBD five-year age-sex groups and produce accurate estimates of uncertainty. For survey report tabulations, we extracted data at the most granular age-sex group provided.

Crosswalk

Our GBD smoking case definitions were current smoking of any tobacco product and former smoking of any tobacco product. All other data points were adjusted to be consistent with either of these definitions. Some sources contained information on more than one case definition and these sources were used to develop the adjustment coefficient to transform alternative case definitions to the GBD case definition. The adjustment coefficient was the beta value derived from a linear model with one predictor and no intercept.

We generated separate crosswalk coefficients for the 10-14 age group and the 15-19 age group, as we found the relationships between case definitions differed strongly in the younger age groups compared to the 20+ age groups. To account for this, we attempted to generate a global crosswalk coefficient for both the 10-14 and 15-19 age groups, using the same regression as above. Due to data limitations, none of the crosswalk coefficients met the criteria outlined above, so no data covering youths under 20 years old were crosswalked. In other words, all data from these age groups that appear in the model were asked according to our case definition in the survey.

We propagated uncertainty at the survey level from the crosswalk by incorporating both the variance of the errors and the variance of the adjustment coefficients.

For each source that needed adjusting, we assigned space weights based on GBD region and super region to the sources containing more than one case definition. Data from the same region receiving a full weight of 1, and data from the same super-region received a weight of ½. We explored using a time weight, to control for possible changes in the relationship between smokeless tobacco use behaviours over time. We found incorporating temporal information did not significantly change the estimated

coefficients but did undercut sample sizes, and chose to exclude the time weight. Crosswalk coefficients generated from fewer than 20 data sources were dropped

Age and sex splitting

We split data reported in broader age groups than the GBD 5-year age groups or as both sexes combined by adapting the method reported in Ng et al.

(http://jamanetwork.com/journals/jama/fullarticle/1812960) to split using a sex- geography- time specific reference age pattern. We separated the data into two sets: a training dataset, with data already falling into GBD sex-specific 5-year age groups, and a split dataset, which reported data in aggregated age or sex groups. We then used spatiotemporal Gaussian Process Regression (ST-GPR) to estimate sex-geography-time specific age patterns using data in the training dataset. The estimated age patterns were used to split each source in the split dataset.

The ST-GPR model used to estimate the age patterns for age-sex splitting used an age weight parameter value that minimises the effect of any age smoothing. This parameter choice allows the estimated age pattern to be driven by data, rather than being enforced by any smoothing parameters of the model. Because these age-sex split data points will be incorporated in the final ST-GPR exposure model, we do not want to doubly enforce a modelled age pattern for a given sex-location-year on a given aggregate data point.

Smoking prevalence modelling

We used ST-GPR to model current and former smoking prevalence. Full details on the ST-GPR method are reported elsewhere in the Appendix. Briefly, the mean function input to GPR is a complete time series of estimates generated from a mixed effects hierarchical linear model plus weighted residuals smoothed across time, space and age. The linear model formula for current smoking, fit separately by sex using restricted maximum likelihood in R, is:

$$logit(p_{g,a,t}) = \beta_0 + \beta_1 CPC_{g,t} + \sum_{k=2}^{19} \beta_k I_{A[a]} + \alpha_s + \alpha_r + \alpha_g + \epsilon_{g,a,t}$$

Where $CPC_{g,t}$ is the tobacco consumption covariate by geography g and time t, described above, $I_{A[a]}$ is a dummy variable indicating specific age group A that the prevalence point $p_{g,a,t}$ captures, and α_s, α_r , and α_g are super region, region, and geography random intercepts, respectively. Random effects were used in model fitting but not in prediction.

The linear model formula for former smoking is:

$$logit(p_{g,a,t}) = \beta_0 + \beta_1 PctChange_{A[a],g,t} + \beta_3 CSP_{A[a],g,t} + \sum_{k=3}^{20} \beta_k I_{A[a]} + \alpha_s + \alpha_r + \alpha_g + \epsilon_{g,a,t}$$

Where $PctChange_{A[a],g,t}$ is the percent change in current smoking prevalence from the previous year, and $CSP_{A[a],g,t}$ is the current smoking prevalence by specific age group A, geography g, and time t that point $p_{g,a,t}$ captures, both derived from the current smoking ST-GPR model defined above.

Exposure among current and former smokers

We estimated exposure among current smokers for two continuous indicators: cigarettes per smoker per day and pack-years. Pack-years incorporates aspects of both duration and amount. One pack-year represents the equivalent of smoking one pack of cigarettes (assuming a 20 cigarette pack) per day for one year. Since the pack-years indicator collapses duration and intensity into a single dimension, one pack-year of exposure can reflect smoking 40 cigarettes per day for six months or smoking 10 cigarettes per day for two years.

To produce these indicators, we simulated individual smoking histories based on distributions of age of initiation and amount smoked. We informed the simulation with cross-sectional survey data capturing these indicators, modelled at the mean level for all locations, years, ages, and sexes using spatiotemporal Gaussian process regression. We rescaled estimates of cigarettes per smoker per day to an envelope of cigarette consumption based on supply-side data. We estimated pack-years of exposure by summing samples from age- and time-specific distributions of cigarettes per smoker for a birth cohort in order to capture both age trends and time trends and avoid the common assumption that the amount someone currently smokes is the amount they have smoked since they began smoking. All distributions were age-, sex-, and region- specific ensemble distributions, which were found to outperform any single distribution.

We estimated exposure among former smokers using years since cessation. We utilised spatiotemporal Gaussian process regression to model mean age of cessation using cross-sectional survey data capturing age of cessation. Using these estimates, we generated ensemble distributions of years since cessation for every location, year, age group, and sex.

Risk-outcome pairs

We included the following risk-outcome pairs based on evidence supporting a causal relationship: tuberculosis, lower respiratory tract infections, esophageal cancer, stomach cancer, bladder cancer, liver cancer, laryngeal cancer, lung cancer, breast cancer, cervical cancer, colorectal cancer, lip and oral cancer, nasopharyngeal cancer, other pharyngeal cancer, pancreatic cancer, kidney cancer, leukemia, ischemic heart disease, ischemic stroke, hemorrhagic stroke, subarachnoid hemorrhage, atrial fibrillation and flutter, aortic aneurysm, peripheral arterial disease, chronic obstructive pulmonary disease, other chronic respiratory diseases, asthma, peptic ulcer disease, gallbladder and biliary tract diseases, Alzheimer disease and other dementias, Parkinson disease (protective), multiple sclerosis, type-II diabetes, rheumatoid arthritis, low back pain, cataracts, macular degeneration, and fracture.

Dose-response risk curves

We conducted systematic literature reviews for all risk-outcome pairs identified as being caused by smoking. We extracted effect sizes by cigarettes per smoker per day, pack-years, and years since quitting from cohort and case-control studies. We synthesised these data to produce non-linear dose response curves using a Bayesian meta-regression model. For outcomes with significant differences in effect size by sex or age, we produced sex- or age-specific risk curves.

We estimate risk curves of former smokers compared to never smokers taking into account the rate of risk reduction among former smokers seen in the cohort and case-control studies, and the cumulative exposure among former smokers within each age, sex, location and year group.

PAF calculation

We estimated population attributable fractions based on the following equation:

$$PAF = \frac{p(n) + p(f) \int \exp(x) * rr(x) + p(c) \int \exp(y) * rr(y) - 1}{p(n) + p(f) \int \exp(x) * rr(x) + p(c) \int \exp(y) * rr(y)}$$

where p(n) is the prevalence of never smokers, p(f) is the prevalence of former smokers, p(c) is the prevalence of current smokers, exp(x) is a distribution of years since quitting among former smokers, rr(x) is the relative risk for years since quitting, exp(y) is a distribution of cigarettes per smoker per day or pack-years, and rr(y) is the relative risk for cigarettes per smoker per day or pack-years.

We used pack-years as the exposure definition for cancers and chronic respiratory diseases, and cigarettes per smoker per day for cardiovascular diseases and all other health outcomes.

Secondhand Smoke Capstone Appendix

Flowchart



Exposure

Case definition

We define secondhand smoke exposure as current exposure to secondhand tobacco smoke at home, at work, or in other public places. We use household composition as a proxy for non-occupational secondhand smoke exposure and make the assumption that all persons living with a daily smoker are exposed to tobacco smoke. We use surveys to estimate the proportion of individuals exposed to secondhand smoke at work. We only consider non-smokers to be exposed to secondhand smoke. Non-smokers are defined as all persons who are not daily smokers. Ex-smokers and occasional smokers are considered non-smokers in this analysis. Exposure is evaluated for both children and adults.

Input data

To calculate the proportion of non-smokers who live with at least one smoker, we used unit record data on household composition, which included the ages and sexes of all persons living in the same household. Our sources included representative major survey series with a household composition module, including the Demographic Health Surveys (DHS), the Multiple Indicator Cluster Surveys (MICS), and the Living Standards Measurement Surveys (LSMS); and national and subnational censuses, which included those captured in the IPUMS project and identified using the Global Health Data Exchange catalog (GHDx).

To calculate the proportion of individuals exposed to secondhand smoke at work, by age and sex, we used cross-sectional surveys that ask respondents about self-reported occupational secondhand smoke exposure. Sources include the Global Adult Tobacco Surveys, Eurobarometer Surveys, and WHO STEPS Surveys. We identified sources using the GHDx.

Estimates of primary smoking prevalence in each location were also used in our calculations. Further details on the estimation of primary smoking prevalence can be found in the Smoking methods appendix.

Modelling strategy

We estimated the probability that each person is living with a smoker and is also a non-smoker themselves using set theory. First, household composition data were used at the individual level to capture the ages and sexes of each person in the household. Second, we analyzed surveys with both household composition data and tobacco use questions and determined that the distribution of household size, mean age of the household members, and the age distribution were not significantly different between households with and without a self-reported smoker. Since we did not find that household composition varied between smokers and non-smokers, we then used the GBD 2017 primary smoking prevalence model to calculate the probability that each household member is a smoker. Next, we used the probability of the union of sets on each individual household member to calculate the overall probability that at least one of the other household members was a smoker. We incorporated occupational exposure by modelling prevalence of current exposure to secondhand smoke at work, by age, sex, location, and year, using ST-GPR. In order to avoid double counting we calculated the probability that an individual is exposed through either non-occupational exposure or occupational exposure, given their age, sex, and household composition. Finally, we multiplied this probability of exposure by the probability that the individual is not a smoker themselves (i.e. 1 minus primary smoking prevalence for that person's location, year, age, and sex). We then collapse these individual-level probabilities to produce average probabilities of exposure by location, year, age, and sex.

These probabilities were modelled in the GBD ST-GPR framework, which generates exposure estimates from a mixed effects hierarchical linear model plus weighted residuals smoothed across time, space, and age. The linear model formula was fit separately by sex using restricted maximum likelihood in R.

We used the sex-specific overall smoking prevalence for adults (age 15 and older) as a country-level covariate in the model. The overall male adult daily smoking prevalence was used as the covariate for females of all ages and for males under age 15. The overall female adult daily smoking prevalence was used as the covariate for males age 15 and older. This was a modelling change from GBD 2015, in which we used the male age-standardised smoking prevalence for the adult female and children under 15 model, and the female age-standardised smoking prevalence for the adult male model.

All input data points from the probability calculation had a measure of uncertainty (variance and sample size) coming from the uncertainty of the primary smoking prevalence model and the sample size from the unit record data going into the modelling process. Geographic random effects were used in model fitting but were not used in prediction.

Theoretical minimum-risk exposure level

The theoretical minimum-risk exposure level for secondhand smoke is zero exposure among nonsmokers, meaning that non-smokers would not live with any primary smokers.

Relative risks

For children ages 0-14, we estimated the burden of otitis media attributable to secondhand smoke exposure. For all ages we estimated the burden of lower respiratory infections (LRI), and for adults

greater or equal to 25 years of age we estimated the burden of lung cancer, chronic obstructive pulmonary disease (COPD), ischemic heart disease, and cerebrovascular disease attributable to secondhand smoke exposure, breast cancer, and type-II diabetes.

For lung cancer, ischemic heart disease, cerebrovascular disease, and LRI, we used country-specific relative risks created using integrated exposure response curves (IER) for PM2.5 air pollution. The relative risks for otitis media, breast cancer, and diabetes are derived from published meta-analyses.

We used the standard GBD population attributable fraction (PAF) equation to estimate burden based on exposure and relative risks.

Chewing Tobacco Capstone Appendix

Flowchart



Chewing Tobacco: Data and Model Flow Chart

In GBD 2016, we estimated age-sex specific current smokeless tobacco use prevalence for all countries and territories from 1990-2016 using all available data. The estimated prevalence was then attributed to either all chewing tobacco, or all snus/snuff by country, based on input from smokeless tobacco experts.

For GBD 2017, we have changed the exposure definition from current smokeless tobacco use to current chewing tobacco use, based on the strength of evidence supporting the health effects of chewing tobacco use. By estimating chewing tobacco exposure in all countries and territories, burden is now estimated for locations previously classified as predominantly snus/snuff, but still have non-negligible use of chewing tobacco.

Input data

Inclusion criteria

We included sources that reported primary chewing tobacco use among respondents over age 10. To be eligible for inclusion, sources had to be representative for their level of estimation (ie. National sources needed to be nationally representative, subnational sources subnationally representative). We included only self-reported chewing tobacco use data and excluded data from questions asking about others' tobacco use behaviors. We included data collected between 1 January 1980 and 1 January 2018.

Prevalence

We searched the Global Health Data Exchange (GHDx) database for primary data sources with the keyword "Tobacco Use" on January 1, 2018 to ensure all available data sources were captured. Of the 3,318 sources identified in the GHDx, 1,578 country-year sources met inclusion criteria and were included.

In addition to the primary data sources identified through the GHDx, we performed a systematic literature search on PubMed. The search was conducted on January 19, 2017 and returned 5982 hits, of which 267 were eligible for inclusion. Of these 267 sources, 200 had already been identified in the GHDx, so the Pubmed search yielded 67 additional sources overall. The search string is shown below: ("smokeless tobacco"[tiab] OR "Tobacco, Smokeless"[Mesh] OR bajjar[tiab] OR ("betel quid"[tiab] AND tobacco[tiab]) OR "chewing tobacco"[tiab] OR chimó[tiab] OR snuff[tiab] OR snuif[tiab] OR dip[tiab] OR dohra[tiab] OR gudakhu[tiab] OR gul[tiab] OR gutka[tiab] OR gutkha[tiab] OR "hnat hsey"[tiab] OR iq'mik[tiab] OR khaini[tiab] OR kharra[tiab] OR khiwam[tiab] OR khimam[tiab] OR kiwam[tiab] OR kimam[tiab] OR "lal dant manjan"[tiab] OR ("loose leaf"[tiab] AND (chew[tiab] OR tobacco[tiab])) OR mainpuri[tiab] OR maras[tiab] OR mawa[tiab] OR mshri[tiab] OR naffa[tiab] OR nas[Supplementary Concept] OR ((nas[tiab] OR nass[tiab]) AND tobacco[tiab]) OR naswar[tiab] OR nasway[tiab] OR nasvay[tiab] OR neffa[tiab] OR ((pan[tiab] OR paan[tiab]) AND tobacco[tiab]) OR (plug[tiab] AND tobacco[tiab]) OR (rapé[tiab] AND tobacco[tiab]) OR ((red[tiab] OR tobacco[tiab]) AND (toothpowder[tiab] OR toothpaste[tiab])) OR shammah[tiab] OR snus[tiab] OR taaba[tiab] OR tapkeer[tiab] OR tawa[tiab] OR tombol[tiab] OR toombak[tiab] OR tuibur[tiab] OR "tobacco water"[tiab] OR (twist[tiab] AND tobacco[tiab]) OR zarda[tiab]) AND Humans[Mesh] AND English[Language] NOT Case Reports[ptyp]

Prevalence data preprocessing

Data extraction

We extracted primary data from individual-level microdata and survey report tabulations. We extracted data on current, former, and/or ever chewing tobacco use as well as frequency of use (daily, occasional, and unspecified, which includes both daily and occasional smokers). Products that do not include tobacco, such as betel quid without tobacco, were excluded or estimated separately as part of the drug use risk factor, if applicable.

For microdata, we extracted relevant demographic information, including age, sex, location, and year, as well as survey metadata, including survey weights, primary sampling units, and strata. This information allowed us to tabulate individual-level data in the standard GBD five-year age-sex groups and produce accurate estimates of uncertainty. For survey report tabulations, we extracted data at the most granular age-sex group provided.

Crosswalk

Our GBD chewing tobacco case definition is current use of chewing tobacco. All other data points measuring other types of chewing tobacco prevalence were adjusted to be consistent with this definition. Some sources contained information on more than one case definition and these sources were used to develop the adjustment coefficient to transform alternative case definitions to the GBD case definition. The adjustment coefficient was the beta value derived from a linear model with one predictor and no intercept.

We generated separate crosswalk coefficients for the 10-14 age group and the 15-19 age group, as we found the relationships between case definitions differed strongly in the younger age groups compared to the 20+ age groups. To account for this, we attempted to generate a global crosswalk coefficient for both the 10-14 and 15-19 age groups, using the same regression as above. Due to data limitations, none of the crosswalk coefficients met the criteria outlined above, so no data covering youths under 20 years old were crosswalked. In other words, all data from these age groups that appear in the model were asked according to our case definition in the survey.

We propagated uncertainty at the survey level from the crosswalk by incorporating both the variance of the errors and the variance of the adjustment coefficients.

For each source that needed adjusting, we assigned space weights based on GBD region and superregion to the sources containing more than one case definition. Data from the same region receiving a full weight of 1, and data from the same super-region received a weight of ½. We explored using a time weight, to control for possible changes in the relationship between chewing tobacco use behaviors over time. We found incorporating temporal information did not significantly change the estimated coefficients but did undercut sample sizes, and chose to exclude the time weight. Crosswalk coefficients generated from fewer than 20 data sources were dropped.

Age and sex splitting

We split data reported in broader age groups than the GBD 5-year age groups or as both sexes combined by adapting the method reported in Ng et al.

(http://jamanetwork.com/journals/jama/fullarticle/1812960) to split using a sex- geography- time specific reference age pattern. We separated the data into two sets: a training dataset, with data already falling into GBD sex-specific 5-year age groups, and a split dataset, which reported data in aggregated age or sex groups. We then used spatiotemporal Gaussian Process Regression (ST-GPR) to estimate sex-geography-time specific age patterns using data in the training dataset. The estimated age patterns were then used to split each source in the split dataset.

The ST-GPR model used to estimate the age patterns for age-sex splitting used an age weight parameter value that minimizes the effect of any age smoothing. This parameter choice allows the estimated age pattern to be driven by data, rather than being enforced by any smoothing parameters of the model. Because these age-sex split data points will be incorporated in the final ST-GPR exposure model, we do not want to doubly enforce a modelled age pattern for a given sex-location-year on a given aggregate data point.

Chewing tobacco prevalence modelling

We used ST-GPR to model chewing tobacco prevalence. Full details on the ST-GPR method are reported elsewhere in the Appendix. Briefly, the mean function input to GPR is a complete time series of estimates generated from a mixed effects hierarchical linear model plus weighted residuals smoothed across time, space and age. The linear model formula for chewing tobacco, fit separately by sex using restricted maximum likelihood in R, is:

$$logit(p_{g,a,t}) = \beta_0 + \sum_{k=1}^{18} \beta_k I_{A[a]} + \alpha_s + \alpha_r + \alpha_g + \epsilon_{g,a,t}$$

Where $I_{A[a]}$ is a dummy variable indicating specific age group A that the prevalence point $p_{g,a,t}$ captures, and α_s , α_r , and α_g are super region, region, and geography random intercepts, respectively. Random effects were used in model fitting but not in prediction.

The hyperparameters for ST-GPR were as follows: age weight = 0.87 for data density below 10 and 1.48 for data density above 10, time weight of 0.04, space weight of 0.01 for data density below 10 and 0.005 for data density above 10, and scale = 20. Amplitude was calculated at the region level. Hyperparameters were decided based on a random grid search evaluated on out of sample RMSE performed on direct smoking data, but due to computational time was not able to be applied to the chewing tobacco data.

Estimating attributable burden

Assessment of risk-outcome pairs

We included outcomes based on the strength of available evidence supporting a causal relationship. There was sufficient evidence to include oral cancer and oesophageal cancer as health outcomes caused by chewing tobacco use. Appendix Table 4 reports the strength of evidence for included outcomes.

Relative risk

Relative risk estimates were derived from prospective cohort studies and population-based case-control studies. Sources used in relative risk estimation are reported in Appendix Table 4.

Appendix Table 6a reports relative risk estimates and uncertainty for the two outcomes included in the analysis, by sex. We extracted the underlying effect size estimates from prospective cohort studies and population-based case-control studies identified by performing a systematic literature review as well as by reviewing the underlying studies included in published meta-analyses. We did not include hospital-based case control studies due to concerns over representativeness. We only included sources that adequately adjusted for major confounders, especially smoking status. Summary effect size estimates were calculated in R, using the 'metafor' package. We performed a random effects meta-analysis using the DerSimonian and Laird method, which does not assume a true effect size but considers each input study as selected from a random sample of all possible sets of studies, and incorporates this variance into the estimation process. We used an inverse-variance weighting method to determine component study weights. We found significantly different relative risks for oral cancer for males and females, and estimated relative risks separately by sex for oral cancer alone.

Theoretical minimum risk exposure level

The theoretical minimum risk exposure level is that everyone in the population has been a lifelong nonuser of chewing tobacco.

Alcohol Use Capstone Appendix

Flowchart



Input data & methodological summary

Exposure

Case definition

We defined exposure as the grams per day of pure alcohol consumed amongst drinkers. We constructed this exposure using the indicators outlined below:

- 1. Current drinkers, defined as the proportion of individuals who have consumed at least one alcoholic beverage (or some approximation) in a 12-month period.
- 2. Lifetime abstainers, defined as the proportion of individuals who have never consumed an alcoholic beverage.
- 3. Alcohol consumption (in grams per day), defined as grams of alcohol consumed by current drinkers, per day, over a 12-month period.
- 4. Alcohol liters per capita stock, defined in liters per capita of pure alcohol, over a 12-month period.

We also used three additional indicators to adjust alcohol exposure estimates to account for different types of bias:

- 1. Number of tourists within a location, defined as the total amount of visitors to a location within a 12 month period.
- 2. Tourists' duration of stay, defined as the number of days resided in a hosting country.

3. Unrecorded alcohol stock, defined as a percentage of the total alcohol stock produced outside established markets.

Input data

A systematic review of the literature was performed to extract data on our primary indicators. The Global Health Exchange (GHDx), IHME's online database of health-related data, was searched for population survey data containing participant-level information from which we could formulate the required alcohol use indicators on current drinkers, lifetime abstainers, alcohol consumption, and binge drinkers. Data-sources were included if they captured a sample representative of the geographic location under study. We documented relevant survey variables from each data-source in a spreadsheet and extracted using STATA 13.1 and R 3.3 . A total of 2,821 potential data-sources were available in the GHDx across countries with subnational locations, out of which 191 data-sources (corresponding 88,734 tabulated data-points by location/year/sex/age) were included across the four indicators mentioned above.

Within the grams per day, current drinkers, and abstainers model, we had a large amount of data on male drinking but not female drinking. To ensure a balanced dataset between sexes for use within DisMod MR 2.1, we imputed for missing sex observations within locations where data existed on male drinking but not female drinking. We used the following models to do so:

For grams per day:

 $y_i \sim Gamma(\mu, \phi)$ $\mu = log^{-1}(1 + SDI + sex + age + sex * age + (1 + sex|super region) + (1 + sex|region)$

Where y is average amount of grams per day within a demographic, μ a parameter for the mean of the average amounts, and ϕ is a dispersion parameter

For current drinking and abstention:

 $\begin{array}{l} y_i \sim Binomial(\pi_i, n) \\ \pi_i \sim Beta(\mu_i, \theta_i) \\ \mu_i = cloglog^{-1}(1 + SDI + sex + age + sex * age + (1 + sex|super region) + (1 + sex|region) \end{array}$

We then sampled 1000 draws from the above estimator for both sex = male & sex = female, with all other variables fixed by demographic unit. For sampling draws, we assumed the parameters were Gaussian multivariately distributed. For each demographic unit with only male observations, we multiplied male data by the ratio between the draws with sex = male & sex = female to impute for female observations.

To generate estimates of alcohol consumption in liters per capita (LPC), we obtained data from FAOSTAT, and WHO GISAH database.^{1,2} To provide more stable time trends in the model, we transformed FAO sales data (which calculates stock based on primary inputs) to a lagged five-year average. Given WHO uses FAO data in locations where WHO could not find data using their own methods, we removed FAO data in the locations where WHO used FAO data in place of their own. To correct for bias in the underlying data sources, we adjusted the input data (crosswalked), by running a mixed effect model on the log average of the data with dummy variables for the data series, as well as random effects on super region, region, country, and time. We adjusted the data points using the following equation:

Log Average Data = D + (Super Region | D, Region | D, Country | D, Year | D)

Transformed data = data * $e^{\widehat{\beta_1} + \widehat{\beta_3}}$

where:

D is a dummy variable for a data source

None of the data sources on liters per capita provided estimates of uncertainty, which is a component required for our eventual modelling strategy. To generate uncertainty, we ran a Loess model on the adjusted data points and the standard deviation between the difference of the Loess smoothed model and the adjusted data points across a five-year span was used as the standard deviation of the data. (i.e., if the total stock changes more variably in a narrow time frame, we believe the data to be more uncertain).

We obtained data on the number of tourists and their duration of stay from the UNWTO.³ We applied a crosswalk across different tourist categories, similar to the one used for the liters per capita data, to arrive at a consistent definition (i.e. visitors to a country).

We obtained estimates on unrecorded alcohol stock from six published papers,⁴⁻⁹ consisting of 166 locations.

Modelling strategy

While population-based surveys provide accurate estimates of the prevalence of lifetime abstainers and current drinkers, they typically underestimate real alcohol consumption levels.¹⁰⁻¹² As a result, we considered the liter per capita input to be a better estimate of overall volume of consumption. Per capita consumption, however, does not provide age- and sex-specific consumption estimates needed to compute alcohol-attributable burden of disease. Therefore, we use the age-sex pattern of consumption among drinkers modelled from the population survey data and the overall volume of consumption from FAO and GISAH to determine the total amount of alcohol consumed within a location. In the paragraphs we outline how we estimated each primary input in the alcohol exposure model, as well as how we combined these inputs to arrive at our final estimate of grams per day of pure alcohol. We estimated all models below using 1,000 draws.

For data obtained through surveys, we used DisMod-MR 2.1 to construct estimates for each country/year/age/sex. We chose to use DisMod due to its ability to leverage information across the heterogeneous age groups reported in the surveys, through age-integration, as well as the model's ability to leverage information available from data in nearby locations or time-periods.¹³

We modelled the alcohol liters per capita data, as well as the total number of tourists, using a spatiotemporal Gaussian process regression (ST-GPR). We chose parameters, as well as our final model, using out-of-sample 10-fold cross validation.

Given the heterogeneous nature of the estimates on unrecorded consumption, as well as the wide variation across countries and time-periods, we took 1,000 draws from the uniform distribution of the lowest and highest estimates available for a given country. We did this to incorporate the diffuse

uncertainty within the unrecorded estimates reported. We used these 1,000 draws in the above equation. We adjusted LPC only for countries where estimates were available.

We adjusted the alcohol LPC for unrecorded consumption using the following equation:

Alcohol LPC = $\frac{Alcohol LPC}{(1-\% Unrecorded)}$

We then adjusted the estimates for alcohol LPC for tourist consumption by adding in the per capita rate of consumption abroad and subtracting the per capita rate of tourist consumption domestically.

 $\label{eq:alcohol} Alcohol \ LPC_d = Unadjusted \ Alcohol \ LPC_d + Alcohol \ LPC_{Domestic \ consumption \ abroad} \\ - \ Alcohol \ LPC_{Tourist \ consumption \ domestically}$

 $\frac{Alcohol LPC_{i}}{\sum_{l} Tourist Population_{l} * Proportion of tourists_{i,l} * Unadjusted Alcohol LPC_{l} * \frac{Average length of stay_{i,l}}{365} * Population_{d}}$

where:

l is the set of all locations, i is either Domestic consumption abroad or Tourist consumption domestically,

and d is a domestic location

After adjusting alcohol LPC by tourist consumption and unrecorded consumption for all location/years reported, sex-specific and age-specific estimates were generated by incorporating estimates modelled in DisMod for percentage of current drinkers within a location/year/sex/age, as well as consumption trends modelled in the DisMod g/day model. We do this by first making sure the sum of percent current drinkers and percent abstainers sum to one for a given location/year/age/sex. We then calculate the proportion of total consumption for a given location/year by age and sex, using the estimates of alcohol consumed per day, the population size, and the percentage of current drinkers. Lastly, we then multiply this proportion of total stock for a given location/year/sex/age by the total stock for a given location/year/sex/age. We then convert these estimates to be in terms of grams/per day. The following equations describe these calculations:

% Current drinkers $_{l,y,s,a} = \frac{\% Current drinkers _{l,y,s,a}}{\% Current drinkers _{l,y,s,a} + \% Abstainers _{l,y,s,a}}$

Proportion of total consumption $_{l,y,s,a}$

 $= \frac{Alcohol \ g/day \ _{l,y,s,a} * Population \ _{l,y,s,a} * \% Current \ drinkers \ _{l,y,s,a}}{\sum_{s,a} Alcohol \ g/day \ _{l,y,s,a} * Population \ _{l,y,s,a} * \% Current \ drinkers \ _{l,y,s,a}}$

 $Alcohol LPC_{l,y,s,a} = \frac{Alcohol LPC_{l,y} * Population_{l,y} * Proportion of total consumption_{l,y,s,a}}{\% Current drinkers_{l,y,s,a} * Population_{l,y,s,a}}$

 $Alcohol \ g/day \ _{l,y,s,a} = Alcohol \ LPC \ _{l,y,s,a} * \frac{1000}{365}$

where:

l is a location, y is a year, s is a sex, and a is an age group.

We then used the gamma distribution to estimate individual level variation within location, year, sex, age drinking populations, following the recommendations of other published alcohol studies.^{7,8} We chose parameters of the gamma distribution based on the mean and standard deviation of the 1,000 draws of alcohol g/day exposure for a given population.

Theoretical minimum-risk exposure level

We calculated TMREL by first calculating the overall risk attributable to alcohol. We did this by weighting each relative risk curve by the share of overall DALYs for a given cause. We then took the minimum of this overall-risk curve as the TMREL of alcohol-use. More formally,

 $TMREL = argmin average overall risk_{\omega}(g/day)$

Average overall risk_{$$\omega$$}(g/day) = $\sum_{i}^{\omega} RR_{i}(g/day) * \frac{DALY_{i}}{\sum_{i}^{\omega} DALY_{i}}$

Where:

 ω is the set of causes associated with alcohol, i is a given cause from that set, DALY is the global DALY rate in 2010, and RR is the dose response curve for a given cause and exposure level in grams per day.

In other words, we chose TMREL as being the exposure that minimises your risk of suffering burden from any given cause related to alcohol. We weight the risk for a particular cause in our aggregation by the proportion of DALYs due to that cause. (e.g. since more observed people die from IHD, we weight the risk for IHD more in the above calculation of average risk compared to, say, diabetes, even if both have the same relative risk for a given level of consumption)

Relative risks

For GBD 2016, we performed a systematic literature review of all cohort and case-control studies reporting a relative risk, hazard ratio, or odds ratio for any risk-outcome pairs studied in GBD 2016. Studies were included if they reported a categorical or continuous dose for alcohol consumption, as well as uncertainty measures for their outcomes, and the population under study was representative. Relative risk estimates by dose can be found in Appendix Table 6c.

We then used these studies to calculate a dose-response, modelled using DisMod ODE. We chose DisMod ODE rather than a conventional mixed effect meta-regression because of its ability to estimate nonparametric splines over doses (i.e. for most alcohol causes, there is a non-linear relationship with different doses) and incorporate heterogeneous doses through dose-integration (i.e. most studies report doses categorically in wide ranges. DisMod ODE estimates specific doses when categories overlap across studies, through an integration step.) We used the results of the meta-regression to estimate a non-parametric curve for all doses between 0-150 g/day and their corresponding relative risks. For all causes, we assumed the relative risk was the same for all-ages and sexes, with the exception of ischemic heart disease, ischemic stroke, hemorrhagic stroke, and diabetes, which we estimated by sex.

Regarding injuries outcomes, we constructed relative risks based on chronic exposure rather than acute, which has a weaker relationship to the outcome, though still significant.^{15,16,18-21} We decided to use chronic exposure given the lack of available data on acute exposure, as well as, the lack of cohort studies using acute exposure as a metric. Further, using chronic exposure allowed us to construct relative risks curves for unintentional injuries, interpersonal violence, motor vehicle accidents, and self-harm using the same method as reported above.

In the case of motor vehicle accidents, we adjusted the PAF to account for victims of drunk drivers that are involved in accidents. Using data from the Fatality Analysis Reporting System in the US,¹⁷ we calculated the average number of fatalities in a car crash involving alcohol, as well as the percentage of those fatalities distributed by age and sex (figures 1 and 2). We aggregated FARS data across the years 1985-2015, given there was little variation in the data temporally and the number of cases in old age groups had too much variance when constructing estimates by year. To adjust PAFs, we multiplied attributable deaths by the average number of fatalities from FARS and redistributed the PAF amongst each population, based on the probability of being a victim to a certain drunk driver by age and sex, based on the FARS data. The following equation describes this process:

$$Adjusted PAF_{i} = \frac{\sum_{d} PAF_{d} * DALY_{d} * Avg \ Fatalities_{d} * P(i \ is \ a \ victim)_{d}}{DALY_{i}}$$

where:

i is a population by location, year, age, sex and *d* is the set of all age and sex exposed groups within that location and year.





Population attributable fraction

For all causes, we defined PAF as:

$$PAF(x) = \frac{P_A + \int_0^{150} P(x) * RR_C(x) \, dx - 1}{P_A + \int_0^{150} P(x) * RR_C(x) \, dx} \qquad P(x) = P_C * \Gamma(p)$$

where:

 P_c is the prevalence of current drinkers, P_a is the prevalence of abstainers, $RR_c(x)$ is the relative risk function for current drinkers, and p are parameters determined by the mean and sd of exposure

We performed the above equation for 1,000 draws of the exposure and relative risk models. We then used the estimated PAF draws to calculate YLL, YLDs, and DALYs, as per the other risk factors.

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Injecting Drug Use Capstone Appendix



Flowchart

Input data & methodological summary

Exposure

Case definition

Injecting drug users (IDU) are at high risk from blood-borne infections, including human immunodeficiency virus (HIV) and Hepatitis B and C viruses (HBV and HCV, respectively), through the use of shared needles and injection equipment. In GBD 2010, based on the available epidemiological literature and the availability of exposure estimates^{1,2} we measure the burden of disease attributable to HIV, HBV and HCV due to injecting drug use. An injecting drug user was defined as a current or recent user aged 15-64 years old.

Input data

The major burden of mortality from viral hepatitis is due to cirrhosis and liver cancer resulting from chronic hepatitis infection. Cirrhosis mortality was modelled with vital registration data using CODEm. Etiologic proportion models, estimated using DisMod-MR 2.1, were used to split the overarching cirrhosis mortality estimates into cases of cirrhosis attributable to hepatitis B, hepatitis C, alcohol, and other causes.¹⁻⁴

Liver cancer mortality was modelled using cancer registry data. The incidence numbers were transformed into mortality estimates using mortality to incidence ratios. The mortality estimates from cancer registries were then combined with vital registration system data as input data into CODEm, which produced the final mortality estimates for liver cancer. As with cirrhosis mortality, etiologic

proportions for liver cancer due to hepatitis B, and C, alcohol, and other causes were generated using DisMod-MR 2.1.

To estimate the burden of HIV cases attributable to IDU, we extracted data on the proportion of notified HIV cases by transmission route – sexual intercourse, injecting drug use, commercial sex work and other -- from a number of agencies that conduct surveillance of HIV across the globe.⁶⁻¹³

The prevalence of current injecting drug use was estimated using data from a 2008 review conducted by the Reference Group to the UN on HIV and injecting drug use,¹⁵ and a new review currently being conducted by international collaborations and experts. The reviews used a multistage process of systematic review adhering to international guidelines. It involved multiple stages of peer and expert review, with searches of the peer-reviewed literature in addition to an extensive review of online grey literature databases in the drug and alcohol and HIV fields. Additional data on the age and sex distribution of injecting drug use were sourced for this modelling exercise.

In order to generate a pooled incidence rate/absolute relative risk for viral hepatitis among people who inject drugs, we conducted a meta-analysis of longitudinal epidemiological studies that reported a hepatitis B¹⁶⁻²⁰ or hepatitis C¹⁶⁻³¹ incidence rate among PWID. We calculated confidence intervals for the incidence rate (where no CI was reported) from a Poisson distribution around the number of cases.

We excluded studies that focused on non-representative subgroups, such as recent injectors or adolescents or because hepatitis incidence is far higher in those groups than for all people who inject drugs (e.g. Larney et al.³²) We did not vary incidence among active injectors according the availability of blood borne virus prevention strategies (e.g. NSPs, opioid substitution therapy) because too few studies have examined different levels of incidence according to variable coverage, and we were not able to estimate coverage by country over time. In any case, in most countries, coverage of virus prevention strategies remains very low among people who inject drugs,³³ and would have been negligible in most countries until recent years.

Modelling strategy

As part of the GBD 2017 study, we measured the burden of hepatitis B and hepatitis C (including attributable cirrhosis and liver cancer) and HIV at the country, regional, and global level for each age-sex group for the years 1990 to 2017. For HIV, hepatitis B and hepatitis C, disease-specific natural history models were used to estimate deaths and YLDs, because the three-state model in DisMod-MR 2.1(susceptible, cases, dead) did not capture the complexity of the disease processes.

Mortality estimation

Mortality due to overall acute hepatitis was modelled with vital registration data using the Cause of Death Ensemble Modelling tool (CODEm), an analytical tool that tests the predictive power of hundreds of models to estimate trends in causes of death.⁵ Due to poor coverage of cause of death data for each of the acute hepatitis varieties, four natural history models for hepatitis B and C were used to estimate mortality by deriving incidence from measurements of seroprevalence and then multiplying incidence by case fatality to estimate the number of deaths. These four models were then squeezed so as to fit the parent cause of death model.

We estimated HIV mortality using a modified UNAIDS Spectrum model.² This is a compartmental HIV progression model estimates age-specific incidence, prevalence and death rates using methods

described elsewhere.² This modelling approach was adapted according to epidemic type, including concentrated and generalized epidemics. For concentrated epidemics, the Spectrum models were corrected for misclassification of HIV deaths and then calibrated to align with vital registration data. For generalised HIV epidemics, we minimised a loss function to select epidemic curves that were most consistent with the prevalence and all-cause mortality data.²

Estimation of Years Lived with Disability

For non-fatal estimation, we estimated the incidence of hepatitis B and C using seroprevalence data in DisMod-MR 2.1. For both hepatitis B and C, we use data on the seroprevalence of the hepatitis surface antigen (a marker of chronic infection in hepatitis B and a marker of ever-infection in hepatitis C), excess mortality, and remission, to estimate incidence of both hepatitis infections. Incidence of cirrhosis was also estimated in DisMod using cirrhosis hospital data and cause-specific mortality rate (CSMR) data.

Incidence of liver cancer was derived by dividing mortality by the mortality to incidence ratios, which were then used to predict liver cancer survival. Finally, we estimated prevalence as a function of incidence and survival by splitting prevalence into four phases. Each phase had different disability weights, which were used to generate YLDs for that phase.

Finally, incidence of HIV was also estimated using the UNAIDS Spectrum modelling approach described above in the mortality estimation section.

Burden of HIV attributable to injecting drug use

We then estimated the proportion of HIV cases attributable to three transmission categories (sex, IDU and other) for all country-time periods using DisMod-MR 2.1. The only covariate used in the model was one that added variance to the data points derived from data sources that attributed a portion of HIV cases to "unknown" transmission sources. We scaled the proportions from each of the three transmission models (sex, IDU and other) to ensure that they fit the total HIV transmission envelope by country, year, age and sex.

Burden of hepatitis B and hepatitis C attributable to injecting drug use

To estimate the relative contribution of IDU to hepatitis B and C disease burden at the country, regional and global level, we used a cohort method. We re-calibrated individuals according to history of injecting drug use, and their accumulated risk of incident hepatitis B and C due to IDU. We made use of data on prevalence of current injecting drug use, pooled in DisMod-MR 2.1; a meta-analysis of incidence rates of hepatitis B and hepatitis C among people who inject drugs; and estimates of population-level incidence of hepatitis B and C between 1990 and 2017. We used back extrapolations to estimate incidence before 1990. These steps are detailed below.

To estimate the lifetime risk of being infected with hepatitis B or C, we undertook a cohort analysis for each country, year, age, and sex category and estimated the probability of an individual having been infected in each preceding year. One of the main inputs to this cohort method was the probability of having injected drugs in a specific age cohort in a given calendar year. For example, for a cohort of 40-year-olds in 2015, the relevant probability in 2005 is the estimated prevalence of injecting drug use among 30-year-olds.

In addition to a global time series of estimated prevalence of injecting drug use, we also used the incidence of hepatitis B or C and the sero-conversion rate of hepatitis B and hepatitis C among people who inject drugs for each age-sex-country-year from 1960 to 2013 by 5-year age groups.

1. Incidence rate of Hepatitis B and C in the general population

We modelled the annual incidence rate of hepatitis B and hepatitis C using sero-prevalence data in DisMod-MR 2.1. We assumed a low remission (mean 0.015 and standard error 0.0075)¹⁴ in the hepatitis B model to reflect the small proportion of cases who spontaneously clear the infection. We assumed zero remission for hepatitis C.

2. Prevalence of ever-injecting drug use

DisMod-MR 2.1 was used to estimate the prevalence of injecting drug use with year as a covariate to estimate the trends over time. DisMod makes an average estimate of the change in drug use over the time period from 1990-2017 and we took draws from a normal distribution of the coefficient to project IDU prevalence backward in time to 1960 from baseline level in 1990.

3. Pooled seroconversion hazard of hepatitis C and hepatitis B among people who ever injected drugs

This pooled sero-conversion hazard for both hepatitis C and hepatitis B was derived from a metaanalysis of longitudinal epidemiologic studies described above in the input data section.

Theoretical minimum-risk exposure level

The theoretical minimum-risk exposure level is defined as zero exposure to injecting drug use.

Relative risks

For drug use, there were not substantial changes made to the effect sizes from GBD 2015. We used a pooled absolute risk of Hepatitis C and Hepatitis B among those who have ever used injecting drugs.

In addition to assessing IDU as a risk factor for blood-borne infections, the broader category of mental and substance use disorders is assessed as risk factors for suicide. The suicide burden attributable to mental and substance use disorders is estimated by comparing the current health status with a theoretical-minimum-risk exposure defined as the counterfactual status of the absence of mental and substance use disorders (Ferrari, Norman et al 2014).

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Dietary Risks Capstone Appendix



Flowchart

Input data & methodological summary

Exposure

Case definition

For GBD 2017, risk factors associated with diet include: diet low in fruits, vegetables, legumes, whole grains, nuts and seeds, fiber, seafood omega-3 fatty acids, polyunsaturated fatty acids, calcium, milk; and diet high in red meat, processed meat, sugar sweetened beverages, trans fatty acids, and sodium. Exposure to a diet low in fruits is defined as average daily consumption of less than 250 grams per day of fruits (fresh, frozen, cooked, canned, or dried, excluding fruit juices and salted or pickled fruits). Exposure to diet low in vegetables is defined as average daily consumption of less than 360 grams per day of vegetables (fresh, frozen, cooked, canned or dried vegetables excluding legumes and salted or pickled vegetables, juices, nuts and seeds, and starchy vegetables such as potatoes or corn). Exposure to a diet low in legumes is defined as average daily consumption of less than 60 grams per day of legumes. Exposure to diet low in whole grains is defined as average daily consumption of less than 125 grams per day of whole grains (bran, germ, and endosperm in their natural proportion) from breakfast cereals, bread, rice, pasta, biscuits, muffins, tortillas, pancakes and other sources. Exposure to diet low in nuts and seeds is defined as average daily consumption of less than 20.5 grams per day of nuts and seeds. Exposure to diet low in milk is defined as average daily consumption of less than 435 grams per day of milk including non-fat, low-fat, and full-fat milk, excluding soy milk and other plant derivatives. Exposure to diet low in calcium is defined as average daily consumption of less than 1.15 grams per day of calcium

from all sources, including milk, yogurt, and cheese. Exposure to diet low in fiber is defined as average daily consumption of less 23.5 grams per day of than fiber from all sources including fruits, vegetables, grains, legumes and pulses. Exposure to diet low in seafood omega-3 fatty acids is defined as average daily consumption of less than 250 milligrams per day of eicosapentaenoic acid (EPA) and docosahexaenoic acid (DHA). Exposure to diet low in polyunsaturated fatty acids is defined as average daily consumption of less than 11% of total energy intake from polyunsaturated fatty acids. This represents a change from the last round of GBD where the burden associated with a diet low in polyunsaturated fatty acids was estimated as a replacement for consumption of saturated fatty acids. Exposure to diet high in red meat is defined as average daily consumption of greater than 22.5 grams per day of red meat (beef, pork, lamb, and goat but excluding poultry, fish, eggs, and all processed meats). Exposure to diet high in processed meat is defined as average daily consumption of greater than 2 grams of meat preserved by smoking, curing, salting, or addition of chemical preservatives. Exposure to diet high in sugar sweetened beverages is defined as average daily consumption of greater than 2.5 grams per day of beverages with ≥50 kcal per 226.8 gram serving, including carbonated beverages, sodas, energy drinks, fruit drinks, but excluding 100% fruit and vegetable juices. Exposure to diet high in trans-fatty acids is defined as average daily consumption of greater than 0.5% of trans fat from all sources, mainly partially hydrogenated vegetable oils and ruminant products. Exposure to diet high in sodium is defined as average 24 hour urinary sodium greater than 3 grams per day.

Input data

We used dietary data from multiple sources including nationally and sub-nationally representative nutrition surveys, household budget surveys, accounts of national sales, and United Nations FAO Food Balance Sheets and Supply and Utilization Accounts. A specific improvement for this round was a novel approach we developed that allows the incorporation of FAO-based data to assist with our estimation of whole grains consumption. This is a powerful development as it dramatically increases our data coverage across countries and through time. This was achieved through separate estimation of total grain and refined grain availability. With whole grains and refined grains representing the entirety of all grain available, we were then able to calculate the availability of whole grains by difference. Additionally, for sodium and trans-fatty acids, we used data on 24-hour urinary sodium and availability of hydrogenated vegetable oil in packaged foods, respectively. Polyunsaturated and trans-fatty acids were modelled as a percent of total dietary energy. We modelled missing country-year data from FAO using a space-time Gaussian process regression and lag-distributed country income as the covariate. For each dietary factor, we estimated the global age pattern of consumption based on nutrition surveys (i.e., 24-hour diet recall) and applied that age pattern to the FAO data. Substantive changes in input data compared to GBD 2015 are as follows: (a) re-extracting data from all nutrition surveys and standardising the definition of dietary components across sources; (b) incorporating data gathered through a systematic review of literature for each of our dietary risk factors; (c) using sales data for fruit, vegetables, legumes, processed meats, red meats, sugar-sweetened beverages, and milk.

Modelling strategy

We used a spatio-temporal Gaussian process regression (ST-GPR) framework to estimate the intake of each dietary factor by age, sex, country, and year. In GBD 2017, for all dietary factors other than sodium, we considered data from 24-hour diet recall as the gold standard, and cross-walked other methods of assessment to the gold standard method. For sodium, the 24-hour urinary sodium was considered as the

gold standard. To estimate the 24-hour urinary sodium based on dietary sodium, we performed a crosswalk adjustment between these two types of data.

Table 1 summarises the study-level and country-level covariates used in modelling of each dietary factor.

	Data Sources			Country level covariate	
	Sales	FFQ^1	HBS ²	FAO	
Diet low in fruits					Lag-distributed income, total available
					kilocalories per person per day
Diet low in vegetables					Lag-distributed income, total available
			•	•	kilocalories per person per day
Diet low in legumes					Lag-distributed income, total available
		_	•	•	kilocalories per person per day
Diet low in whole grains	-		-		Lag-distributed income
Diet low in nuts and seeds	_	_			Lag-distributed income, total available
			•		kilocalories per person per day
Diet low in milk	•			•	Lag-distributed income, total available
			•		kilocalories per person per day
Diet high in red meat	•			•	Lag-distributed income, total available
		•	•	•	kilocalories per person per day
Diet high in processed meat			●	-	Lag-distributed income
Diet high in sugar-sweetened					National availability of sugar (grams/person/day),
beverages		•		-	Lag-distributed income, total available
					kilocalories per person per day
Diet low in fiber	-	•	_	•	Lag-distributed income, total available
					kilocalories per person per day
Diet suboptimal in calcium	-	•	-	•	Lag-distributed income, total available
					kilocalories per person per day
Diet low in seafood omega-3 fatty	-	_	-	•	Lag-distributed income
acids				-	
Diet low in polyunsaturated fatty acids	-	•	-	•	Lag-distributed income, total available
					kilocalories per person per day
Diet high in trans fatty acids			-	-	-
Diet high in sodium ³	-	-	-	-	-

Table 1. Types of data sources (other than 24-hour dietary recall) and covariates used in modelling of each dietary factor.

¹Food Frequency Questionnaire

² Household Budge Survey

³ For sodium, we used data from the 24-hour urinary sodium and 24-hour dietary recall.

To characterise the distribution of each dietary factor at population level, we use an ensemble approach that separately fit 12 distributions for individual level microdata to specific to each data source's sampled population. The respective goodness of fit of each family was assessed and a weighting scheme was determined to optimise overall fit to the unique distribution of each risk factor. A global mean of the weights for each risk factor's data sources was created. We then determined the standard deviation of each population's consumption through a linear regression that captured the relationship between

the standard deviation and mean of intake in nationally representative nutrition surveys using 24-hour diet recalls:

 $ln (Standard deviation) = \beta_0 + \beta_1 \times ln (Mean_i)$

Then we applied the coefficients of this regression to the outputs of our ST-GPR model to calculate the standard deviation of intake by age, sex, year, and country. We also quantified the within person variation in consumption of each dietary component and adjusted the standard deviations accordingly.

Theoretical minimum-risk exposure level

In GBD 2016, to estimate the TMREL for each dietary factor, we first calculated the level of intake associated with the lowest risk of mortality from each disease endpoint based on the studies included in the meta-analyses of the dietary relative risks. Then, we calculated the TMREL as the weighted average of these numbers using the global number of deaths from each of outcome as the weight (Table 2).

Dietary Factor	GBD 2017
Fruits	200-300 gr/day
Vegetables	290-430 gr/day
Legumes	50-70 gr/day
Whole grains	100-150 gr/day
Nuts	16-25 gr/day
Red meats	18-27 gr/day
Processed meats	0-4 gr/day
Milk	350-520 gr/day
Sugar sweetened beverages	0-5 gr/day
Polyunsaturated fatty acids	9-13% of total daily energy
Seafood omega-3 fatty acids	200-300 mg/day
Trans fatty acids	0-1% of total daily energy
Dietary fiber	19-28 gr/day
Dietary calcium	1.0-1.3 gr/day

Table 2. Theoretical minimum-risk exposure level for dietary factors GBD 2017.

Relative risk

In GBD 2016, we measured the health effects of a diet high in sugar-sweetened beverages (SSBs) through how it changes a population's body-mass index (BMI). All attributed disease burden was then solely through the health outcomes associated with a high BMI. Given the more recent publication of dose-response meta-analyses that quantifies the direct effects of SSB consumption on incidence of and mortality from disease endpoints (i.e., ischemic heart disease, and type-2 diabetes), we have updated

our approach to reflect the best available evidence. In GBD 2017, consistent with other dietary risks, we have estimated the health effects of SSBs on morbidity and mortality from disease endpoints. Considering that the relative risks included in these new meta-analyses were mostly adjusted for BMI, our estimated disease burden reflects the burden of disease attributable to SSB consumption independent of its effect on BMI.

We obtained the relative risk of each disease endpoint per serving of the dietary components from recent dose-response meta-analyses of prospective observational studies, and where available randomised controlled trials. Considering the well-established age trend of the relative risks of metabolic risk factors for cardiovascular disease and diabetes, we conducted a literature review to identify the most important metabolic mediators for each dietary factor and used the age trend of the relative risk for each dietary factor (s) and the disease endpoint to estimate the age-specific relative risk for each dietary factors (Table 3).

	Body Mass Index	Total Serum Cholesterol	Fasting Plasma Glucose	Systolic Blood Pressure
Diet low in fruits	•	•	•	•
Diet low in vegetables	•	•	•	•
Diet low in legumes	•	•	•	•
Diet low in whole grains	•	•	•	-
Diet low in nuts and seeds	•	•	•	•
Diet high in red meats	•	-	•	-
Diet high in processed meats	•	-	•	•
Diet low in fiber	-	•	-	-
Diet low in seafood omega-3 fatty acids	•	-	-	•
Diet low in polyunsaturated fatty acids	-	•	•	-
Diet high in trans fatty acids	•	•	-	-

Table 3. Metabolic mediators used to determine the age trend of the effect of dietary factors on cardiometabolic outcomes.

Intimate Partner Violence Capstone Appendix

Flowchart



Input data & methodological summary

Exposure

Case definition

The case definition for intimate partner violence (IPV) is ever experienced one or more acts of physical and/or sexual violence by a current or former intimate partner since the age of 15 years. Estimated in females only because evidence of risk-outcomes for males does not meet our criteria.

 Physical violence is defined as: 'being slapped or having something thrown at you that could hurt you, being pushed or shoved, being hit with a fist or something else that could hurt, being kicked, dragged, or beaten up, being choked or burnt on purpose, and/or being threatened with or actually having a gun, knife, or other weapon used on you.'

- Sexual violence is defined as: 'being physically forced to have intercourse when you did not want to, having sexual intercourse because you were afraid of what your partner might do, and/or being forced to do something that you found humiliating or degrading' (the definition of humiliating and degrading may vary across studies depending on the regional and cultural setting).
- Intimate partner is defined as: 'a partner to whom you are married or with whom you cohabit.' In countries where people date, dating partners will also be considered (a partner with whom you have an intimate (sexual) relationship with but are not married to or cohabiting).

Input data

No systematic review of the literature was completed for GBD 2017. However, a systematic review of the intimate partner violence prevalence literature was conducted in PubMed for anything published between January 2016 and January 2017 for the GBD 2016 cycle. The following search terms were used to conduct the systematic review:

((("health surveys"[MeSH Terms] AND prevalence[Title/Abstract]) OR ("sentinel surveillance"[MeSH Terms] AND prevalence[Title/Abstract]) OR ("prevalence"[Title/Abstract] AND cross sectional studies[MeSH Terms])) AND (abuse, sexual[MeSH Terms] OR domestic violence[MeSH Terms] OR abuse, partner[MeSH Terms] OR abuse, spousal[MeSH Terms] OR rape[MeSH Terms]) NOT ("comment"[Publication Type] OR "letter"[Publication Type] OR "editorial"[Publication Type]))

We get the proportion of solved homicides that were perpetrated by an intimate partner from crime statistics and police reports.

In GBD 2015, an updated systematic review was done for IPV homicide sources in PubMed through April 2016. The query used for this Pubmed search was:

((IPV[All Fields] OR ("intimate partner violence"[MeSH Terms] OR ("intimate"[All Fields] AND "partner"[All Fields] AND "violence"[All Fields]) OR "intimate partner violence"[All Fields])) AND (("homicide"[MeSH Terms] OR "homicide"[All Fields]) OR femicide[All Fields])) AND ("2013/01/01"[PDAT] : "3000/12/31"[PDAT])

These literature sources were supplemented with sources from the GHDx that were tagged with Intimate partner violence AND Homicide.

Modelling strategy

We use three distinct approaches to estimate burden attributable to IPV, including 1) the traditional exposure and relative risk (RR) to percent attributable fraction (PAF) method for depression and abortion; 2) the direct PAF approach for estimating the proportion of homicides that are perpetrated by an intimate partner; and 3) a cumulative risk approach for estimating the burden of HIV/AIDS attributable to IPV.

Estimating attributable burden to IPV for depression, suicide and abortion

We first adjusted data with variable recall periods (previous 12 months versus lifetime), type of violence (sexual, physical, or both) and severity (severe only versus all levels). To convert data to our reference definition of ever having experienced any physical or sexual IPV, we used data from the WHO Multi-country Study on Women's Health and Domestic Violence against Women to construct crosswalk

regressions. The dependent variable in each of these regressions was ever any IPV (reference), while the key independent variable was one of the 11 alternative metrics of IPV that were represented in our dataset:

- 1. Physical IPV in the past 12 months
- 2. Sexual IPV in the past 12 months
- 3. Severe IPV in the past 12 months
- 4. Severe physical IPV in the past 12 months
- 5. Severe sexual IPV in the past 12 months
- 6. Any IPV (physical and/or sexual) in the past 12 months
- 7. Ever any physical IPV
- 8. Ever any sexual IPV
- 9. Ever any severe IPV
- 10. Ever severe physical IPV
- 11. Ever severe sexual IPV

For alternate metrics 1-6 we included a series of age dummies:

$$logit(REF_{ait}) = \beta_0 + \beta_1 logit(ALT_{ait}) + \beta_2 I_a + \varepsilon$$

For alternate metrics 7-11, we ran the following regression:

$$logit(REF_{it}) = \beta_0 + \beta_1 logit(ALT_{it}) + \varepsilon$$

where REF is the reference metric of IPV prevalence, ALT is the alternate metric of IPV prevalence, *I*_a refers to the complete set of age-group indicators, *a* refers to an age-group, *i* refers to a country, and *t* refers to year. We included age-group indicators in the first six regressions because we expected the prevalence of recent IPV to vary by age. Using the intercepts, coefficients, and variance-covariance matrix from each of these eleven regressions, we were able to convert all of the alternate metrics of IPV prevalence in our dataset to estimates of "ever any IPV." We eliminated observations based on alternate metrics of IPV which came from studies that also provided estimates of IPV based on the reference definition.

After applying crosswalks to the alternate metrics of IPV in the manner described above, we made an additional adjustment to the subset of our data that was based on only ever-partnered, currently partnered women currently married women or ever married women. To adjust these values so that they reflected IPV prevalence in the entire female population, regardless of partnered status, we multiplied estimates from these studies by the age-specific fraction of women who had ever been partnered. An updated time series was generated in GBD 2015 using MICS and DHS data in a single parameter DisMod model to reflect the most recent data on proportion of women that have ever been partnered.

After these pre-DisMod crosswalks and adjustments, a single-parameter prevalence model was run in DisMod-MR 2.1 with age mesh points at 0 14 15 20 30 40 50 60 80 & 100. A study-level fixed effect on integrand variance (z-cov) to indicate whether a study was nationally representative or not was used to account for the heterogeneity introduced by studies that are not generalizable to the entire population. This covariate was first tested as an x-cov and the coefficient indicated no systematic bias.

In addition to the lifetime exposure model as described, a 12-month exposure model was also run in DisMod-MR 2.1, with data collected and processed analogously. This 12-month exposure model was used for the IPV-abortion PAF calculation to match the exposure definition in the risk evidence.

Direct PAF for female homicides

The burden of homicides attributable to intimate partner violence was modelled as a direct PAF.

Input data fed into a single-parameter proportion DisMod-MR 2.1 model, which had age mesh points at 0 10 20 45 & 100. The model had a study-level covariate for sources just including police reported homicides. We also included a study-level fixed effect on integrand variance (z-cov) to indicate whether a study was nationally representative or not. This covariate was first tested as an x-cov and the coefficient indicated no systematic bias.

In GBD 2015, we added prevalence of binge drinking to the model as a country-level covariate, but it was dropped for GBD 2016 because it produced a non-significant coefficient.

Cumulative risk approach for PAF of HIV/AIDS due to IPV

The third and final modelling approach that we used to assess burden attributable to intimate partner violence was a cumulative risk approach to measure the burden of HIV/AIDS attributable to IPV.

The approach itself remained the same in GBD 2017, but included updated intimate partner violence exposure numbers from the DisMod-MR 2.1 model described above, as well as revised HIV incidence numbers.

From two cohort studies (Jewkes et al, Lancet 2010 & Kouyoumdijian, et al AIDS 2013) we pooled incidence rate ratio (IRR) of HIV incidence with a random effects model. As we measure burden based on deaths and prevalence, we needed to quantify attributable fractions for prevalence and death rather than incidence. To get a PAF for prevalence we needed to consider the history of exposure to IPV and the accumulated associated risk of incident HIV due to IPV, relative to the overall risk of HIV at the population level. The ratio of cumulative IPV-attributable HIV incidence to total HIV incidence was an approximation of the relevant PAF for HIV prevalence and we assumed this PAF can also be applied to mortality.

$$\frac{Cumulative \ HIV \ incidence \ due \ to \ IPV}{Cumulative \ HIV \ incidence \ overall} = \frac{1 - \prod_{a=o}^{a=n} (1 - PAF_{ay} * I_{ay})}{1 - \prod_{a=o}^{a=n} (1 - I_{ay})}$$

Where:

I = annual incidence rate of HIV

a = age (15-95)

y = year (1980-2016)

 $PAF_{HIV \ incidence} = \frac{[Prevalence \ of \ IPV]_{ay^*} (IRR-1)}{[Prevalence \ of \ IPV]_{ay^*} (IRR-1)+1}$

Theoretical minimum-risk exposure level

The theoretical minimum-risk exposure level is zero exposure to intimate partner violence, as defined above.

Relative risks

We estimate burden attributable to IPV for abortion, depressive disorders, interpersonal violence (i.e. homicide) and HIV incidence. For GBD 2016, we completed a systematic review of the literature for papers reporting relative risk of IPV and our outcomes. Suicide, which was modelled as an outcome of IPV for GBD 2015, was removed from the analysis based on the availability of relative risk literature using suicide and not attempted suicide as the outcome definition. For GBD 2017, we used the same causal evidence for relative risk as was used for GBD 2016.

For HIV, we use a pooled IRR of 1.59 (95% CI 1.3-1.94) from a random effects inverse variance weighted meta-analysis of the two available prospective studies as of date.

The relative risks for depressive disorders and suicide come came from a systematic review of longitudinal studies assessing intimate partner violence and incident diagnosed major depression. A random effects inverse variance weighted meta-analysis produced a pooled relative risk and 95% confidence interval of 1.44 (1.09, 1.92).

For the relative risk for IPV-abortion, we also performed a random effects, inverse variance metaanalysis, which produced a pooled relative risk and 95% confidence interval of 1.91 (1.15, 3.16). An important methodological note with IPV-abortion is that we must apply the pooled relative risk for abortion to the current prevalence of IPV (in the previous 12 months), rather than lifetime prevalence. This is because the case definition for the relative risk component studies was physical or sexual IPV in the past year.



Meta-analysis for IPV-depressive disorders:

Meta-analysis for IPV-abortion:



Meta-analysis for IPV-HIV:



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Childhood Sexual Abuse Capstone Appendix

Flowchart



Input data & methodological summary

Exposure

Case definition

The case definition for childhood sexual abuse (CSA) is ever having had the experience of intercourse or other contact abuse (i.e. fondling and other sexual touching) when aged 15 years or younger, and the perpetrator or partner was greater than five years older than the victim.

Input data

Currently, we use self-reported survey data to measure CSA prevalence, not data from Child Protection Services (CPS) or other crime data. The reliability and comprehensiveness of CPS and crime statistics varies too much geographically to warrant inclusion.

Although no systematic review of the literature was completed for GBD 2017, an updated systematic review of CSA prevalence literature was conducted for sources published between August 2015 and January 2017 during the GBD 2016 cycle. The following search terms were used:

(((("health surveys"[MeSH Terms] AND prevalence[Title/Abstract]) OR ("sentinel surveillance"[MeSH Terms] AND prevalence[Title/Abstract]) OR ("prevalence"[Title/Abstract] AND cross sectional studies[MeSH Terms])) AND (("child abuse"[MeSH Terms] OR "child abuse, sexual"[MeSH Terms]) OR ("sex offenses"[MeSH Terms] OR "child abuse, sexual"[MeSH Terms]) OR (child*[Title/Abstract] AND sexual[Title/Abstract] AND abuse[Title/Abstract]))) NOT ("comment"[Publication Type] OR "letter"[Publication Type] OR "editorial"[Publication Type]))

We supplemented with data from relevant national health surveys and violence-specific surveys. Several survey series used include the United States Behavioral Risk Factor Surveillance System, the CDC Reproductive Health Surveys, Brazil National Alcohol and Drug Survey, and the Gender, Alcohol, and Culture International Study (GENACIS).

A number of study level covariates were also extracted that were used in the modelling process to adjust for heterogeneous definitions across sources. All crosswalks and adjustments were done in DisMod-MR 2.1.

Modelling strategy

CSA prevalence was modeled as a single parameter prevalence model in DisMod-MR 2.1. CSA exposure is modeled separately for males and females because we observe little correlation between the prevalence of child abuse among females and males, and modeling both sexes together causes unreasonable estimates in countries where we only have data for one sex.

Three study-level covariates were used for alternate definitions of the violence.

- Study asked only about intercourse CSA
- Study asked about contact and non-contact CSA
- Study placed restrictions on the relationship between the perpetrator and the victim (e.g. only asked about CSA committed by a father)

We also included study-level fixed effects for varying age thresholds across studies.

- Study asked about recall for events before ages above 15 years (versus reference age threshold of 15)
- Study asked about recall for events before ages less than 15 years (versus reference age threshold of 15)

Two study-level covariate fixed effects on variance (z-cov) were also included in both the male and female models, including an indicator that the survey was not nationally representative, as well as whether the survey was administered in schools. These study-level covariates were tested as x-covs first, but we did not find coefficients which would indicate systematic bias. We have not included any national-level covariates to date due to lack of knowledge about a covariate (for which we have a time series for all GBD locations) that predicts CSA prevalence.

Theoretical minimum-risk exposure level

The theoretical minimum risk exposure level is zero exposure to contact childhood sexual abuse.

Relative risks

We estimate burden attributable to CSA for the following health outcomes: unipolar depressive disorders (major depressive disorder and dysthymia) and alcohol use disorders.

In GBD 2015, we used one twin study that compared adverse outcome risks in same-sex discordant pairs.¹ This study was deemed reliable given that environmental and contextual factors are inherently controlled for when comparing between twins, avoiding potential confounding. However, to add to the strength of the evidence for GBD 2016, we performed a systematic review and a random effects meta-analysis to produce relative risks for depressive disorders and alcohol use disorders. In a departure from GBD 2015, suicide was not used as an outcome for CSA. This decision was based on the evidence available for the relative risk of suicide given exposure to CSA – not enough studies used suicide as an outcome, but instead used attempted suicide. For GBD 2017, we used the same causal evidence as was used for GBD 2016.

The pooled relative risk figures and 95% confidence intervals were 1.63 (1.41, 1.89) for depressive disorders and 1.54 (1.19, 1.99) for alcohol use disorders. The resulting forest plots are as follows:



CSA and depressive disorders meta-analysis

CSA and alcohol use disorders meta-analysis



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Bullying Victimisation Capstone Appendix



Flowchart

Input data & methodological summary

Exposure

Case definition

Bullying victimisation is commonly conceptualised as the intentional and repeated harm of a less powerful individual by peers.¹ This differentiates bullying victimisation from disagreements, conflicts, or playful teasing. The case definition of bullying victimisation in the GBD context is 'bullying victimisation of children and adolescents attending school by peers.' This definition includes the global concept of bullying victimisation which incorporates combined estimates of subtypes such as physical, verbal, relational, and cyberbullying victimisation. It excludes abuse/harassment by siblings, intimate partners, and adults (e.g. teachers). While bullying can be experienced as either a victim or perpetrator,

perpetration (i.e. those who bully others) is not included in this definition although some victims will also be perpetrators.

Input data

In order for a study to be included, it must report the prevalence of bullying victimisation and 1) have been published since 1980, 2) ask participants about bullying victimisation in the previous year or more recently, 3) use an appropriate frequency threshold to define bullying victimisation (approximating at least once a week or greater than 'occasionally'), 4) be representative of the general population rather than a special population (e.g. ethnic minorities), and 5) report prevalence for bullying victimisation overall rather than a subtype e.g. physical bullying victimisation.

Included studies were sourced from a systematic review of three electronic databases (PubMed, EMBASE, and PsycINFO), covering the period 1980 to 2017. No restriction was set on the language of publication. GhDx was also used to source microdata from survey series meeting the above inclusion criteria. Estimates from the Global School-based Student Health Survey (GSHS), the Health Behavior in School-aged Children (HBSC), and the National Crime Victimization Survey – School Crime Supplement (NCVS-SCS) were extracted and included in the dataset.

Modelling strategy

Bullying victimisation prevalence was modelled as a single parameter prevalence model in DisMod-MR 2.1. We assumed no prevalence prior to 5 years or after 20 years of age. Four study-level covariates were included and are shown in the table below, along with their respective levels. Crosswalks for two of the covariates (low bullying frequency and no bullying definition presented) were calculated using within-study pairs of reference and non-reference estimates (n = 9 pairs and n = 3 pairs, respectively).

Covariate name	Reference	Non-reference	Exponentiated beta
Low bullying	Optimal frequency threshold	Sub-optimal frequency	3.35 (3.35 – 3.35)
frequency	used e.g. 'frequently'	threshold used e.g.	(n = 9 pairs)
		'sometimes + frequently'	
No bullying	Definition of bullying	No definition of bullying	1.12 (1.12 – 1.12)
definition	victimisation presented to	victimisation presented to	(n = 3 pairs)
presented	participants	participants or not specified	
Recall 1 year	Asked about bullying	Asked about bullying	1.47 (1.30 – 1.68)
	victimisation more recently	victimisation in the past year	
	than in the past year		
Single school	Sample was a household	Sample was from a single	1.21 (1.01 – 2.12)
sample	survey or multi-school survey	school	

Adjustment for years of schooling

In order to better represent the prevalence of bullying victimisation, prevalence estimates were adjusted for the proportion of children and adolescents attending school by ages 5-9, 10-14, and 15-19 years by sex, location, and year. Data on the proportion of children and adolescents attending school was sourced from the online database (http://data.uis.unesco.org/) published by the United Nations Educational, Scientific, and Culture Organization (UNESCO). The data covered 18,441 country-years for

age groups 6-11, 12-14, and 15-17 years by sex. This data was modelled in ST-GPR, with average years of education as a country-level covariate, to predict the proportion of children and adolescents attending school by these age groups. This gave estimates of the proportion of children and adolescents attending school by age, sex, year, and location.

Theoretical minimum-risk exposure level

The theoretical minimum risk exposure level was assumed to be zero exposure to bullying victimisation.

Relative risks (RRs)

We estimate burden attributable to bullying victimisation for major depressive disorder and anxiety disorders. Data on the association between bullying victimisation and self-harm was also reviewed but not included due to variation in the definition of 'self-harm' and only one study looking at suicide.

Input data for RRs

Studies reporting the prospective longitudinal association between these outcomes and bullying victimisation were sourced from a systematic review of three electronic databases (PubMed, EMBASE, and PsycINFO), covering the period 1980 to 2017. No restriction was set on the language of publication. Studies had to report RRs, ORs, or sufficient data to calculate RRs (i.e. exposed/non-exposed cases/non-cases).

Meta-analysis

The smaller number of estimates for anxiety disorders (n = 6) led to the decision to combine the major depressive disorder and anxiety disorders RR data into a single dataset. This was considered reasonable as the pooled RRs for major depressive disorder and anxiety disorders were effectively equal following an adjustment for low frequency bullying threshold studies (1.82, 95% CI: 1.62-2.04 vs 1.74, 95% CI: 1.43-2.11, respectively).



Meta-analysis of bullying victimisation and major depressive disorder/anxiety disorders

Meta-regression

A meta-regression was conducted to determine the impact of follow-up time on the relationship between bullying victimisation and major depressive disorder/anxiety disorders.

Prior to this, an initial meta-regression revealed studies with a low bullying frequency threshold reported significantly lower RRs than studies utilising the optimal threshold. Six studies reported RRs for both low and optimal bullying frequency threshold and we determined that the within-study ratios from these studies would better inform an adjustment than between-study comparisons estimated as a covariate. These ratios were pooled and then applied to the low bullying frequency threshold estimates (i.e. suboptimal estimates) prior to analysis. This process could not be done for the other characteristics (i.e. symptoms vs diagnosis or no control vs control for outcome as baseline) as there were no withinstudy comparisons available. In the final meta-regression, these were instead controlled for using covariates.

In the final meta-regression, estimates with a follow-up time of 25 years or more were excluded due to only data from a single cohort being available to inform the effect of follow-up time after this period. In total, 19 studies were included in the analyses.²⁻²⁰ While the meta-regression demonstrated a waning effect of time, this was not statistically significant (p = 0.093). However, the model still predicted plausible estimates of RR over time that matched our strong prior that the effect size would have to diminish over time. Time was also significant in a parsimonious model that did not include the non-significant covariates (p = 0.042).

PAF calculations

For bullying victimisation, the PAF calculations could not be determined by current prevalence and a single value for RR. This is due to the waning effect on outcomes over time (as demonstrated in the meta-regression) and because prevalence estimates were from surveys of young people reporting current bullying victimisation rather than estimates of past exposure at the time the outcomes occur (i.e. retrospective estimates).

A cohort method was subsequently developed to address this issue. The following steps are conducted for each point of estimation (i.e. by age, sex, location, and year), hereafter referred to as a 'cohort':

- 1. Pull current and past bullying victimisation prevalence for the cohort from the DisMod-MR 2.1 exposure model.
- 2. Adjust each bullying victimisation prevalence estimate for the proportion of the cohort attending school in that year.
- Divide the cohort into proportions based on time since first exposed to bullying victimisation. This equates to the incidence of bullying victimisation and is estimated using the following formula:

$$I_k = P_k - \sum_{n=0}^{k-1} (I_n \times r_{k-n})$$

where *I* represents incidence, *P* represents prevalence, *r* represents the estimate of persistence, and *k* represents the time between the incidence estimate and the earliest possible time of exposure in the cohort. I_k requires I_0 through to I_{k-1} to first be calculated and so we complete this process by first estimating I_0 , then I_1 , and so on until we have estimated incidence for the latest possible year of exposure for this cohort. The persistence estimate is based on a separate meta-regression of seven studies.^{11,21-26}

4. We Map RRs to the proportions of the cohort based on the time between the point of estimation and when they were first exposed to bullying victimisation and estimate PAFs via the following formula:

$$PAF = \frac{\sum (p_t \times RR_t) + \sum p_{no \ exposure} - 1}{\sum (p_t \times RR_t) + \sum p_{no \ exposure}}$$

where *t* is the time since first exposed to bullying victimisation, *p* is the proportion of the cohort first exposed to bullying victimisation at time *t* or the proportion not exposed to bullying victimisation, and *RR* is the relative risk for depressive and anxiety disorders given *t*.

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Unsafe Sex Capstone Appendix

Unsafe sex risk factor estimation Final burden estimation UNUDES contry progress reports Under estimation Final burden estimation Final burden estimation Final burden estimation Constitution of HPU Observement epidenoinged surveillance Constitution of HPU Constitution of HPU Observement epidenoinged surveillance Constitution of HPU Constitution of

Flowchart

Input data & methodological summary

Case definition and summary of GBD approach

Unsafe sex is defined as the risk of disease due to sexual transmission. The outcomes associated with unsafe sex that we estimate for GBD include HIV, cervical cancer, and all sexually transmitted diseases (STDs) except for those in neonates from vertical transmission, including HIV, opthalmia neonatorum and neonatal syphilis. We assume 100% of cervical cancer and STDs are attributable to unsafe sex and model the proportion of HIV incidence occurring through sexual transmission to estimate the attributable burden for HIV due to unsafe sex.

Input data

To be used in our models, sources must report HIV cases attributable to various modes of transmission. We screened all UNAIDS country progress reports and searched government epidemiological surveillance records for these data. The primary data sources we used were UNAIDS, the European CDC, and the US CDC.

We excluded all extractions where the "other" category for HIV transmissions accounted for greater than 25 percent of all cases. We believe that such high proportions raise concerns about the quality of reporting.

Modelling strategy

We model the proportion of HIV cases attributable to unsafe sex. To do this we collect and clean data, run three DisMod models (HIV attributable to sex, HIV attributable to injection drug use, HIV

attributable to other routes of transmission), adjust results of the three DisMod models to sum to one, and prepare PAFs.

All of the DisMod models included a study-level covariate fixed effect on integrand variance (z-cov) for sources that include cases of unknown transmission in their "other" category. We assumed that the inclusion of unknown cases in the other category would impact the uncertainty around the point estimates. No country level covariates were included in the models. We tested an injection drug use covariate, an opioid use covariate in the proportion HIV due to drug use model, but found no significant coefficients so excluded them from the final model.

A new approach was introduced for GBD 2016, and used again for GBD 2017, to inform an age-pattern in these HIV transmission models. All-age data points represent the majority of the available data, so we derived an age-pattern for the HIV-IDU transmission model from the age-pattern present in the GBD 2017 population attributable fraction for hepatitis B attributable to intravenous drug use. Assuming the proportion of HIV due to other is constant over time, the age-pattern for the proportion of HIV due to sex was set to be the complement to 1 of the age-pattern for the proportion of HIV due to IDU. The allage data were split according to these age-patterns, and the three HIV transmission DisMod models were run on the age-split data. Additional priors were set to inform an age-pattern: zero proportion HIV transmission due to IDU before age 15, zero proportion HIV transmission due to sex before age 10, and 100% transmission due to other before age 10. The results from these HIV transmission models were adjusted to sum to 100% for a given country-year-age-sex group at each of 1,000 draws.



Squeezed global HIV transmission models by age (females, 2016):

Theoretical minimum-risk exposure level

The theoretical minimum level used for unsafe sex is the absence of disease transmission due to sexual contact.

Population attributable fraction calculations

The outcomes associated with unsafe sex that we report on include HIV, cervical cancer, and all sexually transmitted diseases (STDs) except for those in neonates from vertical transmission, including HIV, opthalmia neonatorum and neonatal syphilis.

Based on evidence in the literature, we attribute 100% of cervical cancer to unsafe sex. These sources state that HPV infection is necessary for cervical cancer to develop and that HPV is only spread through sexual contact. The proportion of STDs attributable to unsafe sex is also 100%.

For HIV, the results from the single parameter proportion DisMod model for HIV transmission due to sex were used directly as the population attributable fraction.

Low Physical Activity Capstone Appendix

Flowchart



Input data and methodological summary

Exposure

Case definition

We measure physical activity performed by adults greater than or equal to 25 years of age, for durations of at least ten minutes at a time, across all domains of life (leisure/recreation, work/household and transport). We use frequency, duration and intensity of activity to calculate total metabolic equivalent-minutes per week. MET (Metabolic Equivalent) is the ratio of the working metabolic rate to the resting metabolic rate. One MET is equivalent to 1 kcal/kg/hour and is equal to the energy cost of sitting quietly. A MET is also defined as the oxygen uptake in ml/kg/min with one MET equal to the oxygen cost of sitting quietly, around 3.5 ml/kl/min.

Input data

We included surveys of the general adult population that captured self-reported physical activity in all domains of life (leisure/recreation, work/household and transport), where random sampling was used.

Data were primarily derived from two standardised questionnaires: The Global Physical Activity Questionnaire (GPAQ) and the International Physical Activity Questionnaire (IPAQ), although we included other survey instruments that asked about intensity, frequency and duration of physical activities performed across all activity domains.

Due to a lack of a consistent relationship on the individual level between activity performed in each domain and total activity, we were not able to use studies that included only recreational/leisure activities.

Physical activity level is categorised by total MET-minutes per week using four categories based on rounded values closest to the quartiles of the global distribution of total MET-minutes/week. The lower limit for the Level 1 category (600 MET-min/week) is the recommended minimum amount of physical activity to get any health benefit. We used four categories with higher thresholds rather than the GPAQ

and IPAQ recommended 3 categories to better capture any additional protective effects from higher activity levels.

- Level 0: < 600 MET-min/week (inactive)
- Level 1: 600-3999 MET-min/week (low-active)
- Level 2: 4000-7,999 MET-min/week (moderately-active)
- Level $3: \geq 8,000$ MET-min/week (highly active)

The GHDx was used to locate all surveys that use the GPAQ or IPAQ questionnaire. Although there were many other surveys that focused specifically on leisure activity, we were unable to use these sources because they did not comprise all three domains (work, transport and leisure). In addition, we excluded any surveys that did not report frequency, duration, and intensity of activity.

Modelling strategy

DisMod modelling

For this round of the GBD, we have chosen to separately model all of the GPAQ data separately from the IPAQ data. We then estimated the proportion of each country/year/age/sex subpopulation in each of the above four activity levels using 12 separate Dismod models (one for each data source). We use six categories of physical activity prevalence rather than four to accommodate the different MET-minute/week cutoffs presented in tabulated data sources where individual unit record data was not available. Since the accepted threshold/definition for inactivity is consistently <600 MET-minutes/week, the vast majority of tabulated data was broken down into proportion inactive (model A) and proportion low, moderate or highly active (model B).

	Label	MET-min/week	Name of sequelae in online visualisation tool
А	inactive	<600	Physical inactivity and low physical activity, inactive
В	low/moderately/highly	≥600	Physical inactivity and low physical activity,
	active		low/moderately/highly active
С	low active	600-3999	Physical inactivity and low physical activity, low active
D	moderately/highly	>4000	Physical inactivity and low physical activity,
	active		moderately/highly active
Е	moderately active	4000-7999	Physical inactivity and low physical activity,
			moderately active
F	highly active	≥8,000	Physical inactivity and low physical activity, highly
			active

These models have mesh points at 0 15 25 35 45 55 65 75 85 100, and a study-level fixed effect on integrand variance (Z-cov) for whether a study was nationally representative or not, to account for the heterogeneity introduced boy studies that are not generalizable to the entire population. They also have national level fixed effects on prevalence of obesity.

After DisMod, we rescale each of the 6 models specific to each data source so that the proportions sum to one. Since we have the most data for models A and B, we rescale the sum of the proportion in each category to be equal to one. Next we rescale the sum of model C and D to be equal to the rescaled value from model B. Then we rescale the sum of models E and F to be equal to the rescaled value from model D. After these three rescales we are left with a proportion for each of the four categories that all sum to

1. Scaled results for each data source are then hybridised to produce only one set of results for the prevalence of the four categories of physical activity.

Similar to the previous round, we have directly estimated total MET-minutes per week globally. Although, this year we made use of two specific machine learning algorithms (Random Forest & XGBoost) that were trained using data that could characterise the relationship between total MET-mins/week and each of the categorical prevalences of physical activity. This resulted in country-year-age-sex specific estimates of total physical activity in the form of MET-minutes per week.

Utilising microdata on total MET-mins per week from individual-level surveys, we characterised the distribution of activity level at the population level. We then used an ensemble approach to distribution fitting, borrowing characteristics from individual distributions to tailor a unique distribution to fit the data using a weighting scheme. We characterised the standard deviation of each population's activity through a linear regression that captured the relationship between standard deviation and mean activity levels in nationally representative IPAQ surveys:

 $ln (Standard deviation) = \beta_0 + \beta_1 \times ln (Mean_i)$

We then applied the coefficients of this regression to the outputs of our estimate of total MET-minutes per week regression outputs to calculate the standard deviation by country, year, age, and sex.

Theoretical minimum-risk exposure level

The theoretical minimum-risk exposure level for physical inactivity is 3000-4500 MET-min per week, which was calculated as the exposure at which minimal deaths across outcomes occurred.³

Relative risks

We used a recently published dose-response meta-analysis of prospective cohort studies to estimate the effect size of the change in physical activity level on breast cancer, colon cancer, diabetes, ischemic heart disease and ischemic stroke.³

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High Fasting Plasma Glucose Capstone Appendix

Flowchart



Case definition

High fasting plasma glucose (FPG) is measured as the mean FPG in a population, where FPG is a continuous exposure in units of mmol/L.

Data seeking

Exposure

We did not conduct a systematic review for FPG in GBD 2017. However, we included data sources found in the systematic review for diabetes. Please see the diabetes mellitus appendix in *Global, regional, and national incidence, prevalence, and years lived with disability for 354 diseases and injuries for 195 countries, 1990–2017: a systematic analysis for the Global Burden of Disease Study 2017* for more information on data sources found through those efforts.

Data inputs

Data inputs come from 3 sources:

- Estimates of mean FPG in a representative population
- Individual-level data of fasting plasma glucose measured from surveys
- Estimates of diabetes prevalence in a representative population

Data sources that did not report mean FPG or prevalence of diabetes are excluded from analysis. When a study reported both mean fasting plasma glucose (FPG) and prevalence of diabetes, we use the mean FPG for exposure estimates. Where possible, individual-level data supersede any data described in a study. Individual-level data are aggregated to produce estimates for each 5-year age group, sex, location, and year of a survey.

Data processing

We perform several processing steps to the data in order to address sampling and measurement inconsistencies that will ensure the data are comparable.

1. <u>Small sample size</u>

Estimates in a sex and age group with a sample size <30 persons is considered a small sample size. In order to avoid small sample size problems that may bias estimates, data are collapsed into the next age group in the same study till the sample size reach at least 30 persons. The intent of collapsing the data is to preserve as much granularity between age groups as possible. If the entire study sample consists of <30 persons and did not include a population-weight, the study is excluded from the modelling process.

2. <u>Time, Age, and Sex Splitting</u>

For more details on how datapoints on mean FPG was processed, please see the diabetes mellitus appendix in *Global, regional, and national incidence, prevalence, and years lived with disability for 354 diseases and injuries for 195 countries, 1990–2017: a systematic analysis for the Global Burden of Disease Study 2017.*

3. <u>Crosswalks</u>

We predicted mean FPG from diabetes prevalence using an ensemble distribution. We characterized the distribution of FPG using individual-level data. Details on the ensemble distribution can be found elsewhere in the Appendix. Before predicting mean FPG from prevalence of diabetes, we ensured that the prevalence of diabetes was based on the reference case definition: fasting plasma glucose (FPG) >126 mg/dL (7 mmol/L) or on treatment. For more details on how the case-definition crosswalk is conducted, please see the diabetes mellitus appendix in *Global, regional, and national incidence, prevalence, and years lived with disability for 354 diseases and injuries for 195 countries, 1990–2017: a systematic analysis for the Global Burden of Disease Study 2017.*

Exposure modelling

Exposure estimates are produced for every year between 1980 to 2017 for each national and subnational location, sex, and for each 5-year age group starting from 25 years. As in GBD 2016, we used a Spatio-Temporal Gaussian Process Regression (ST-GPR) framework to model the mean fasting plasma glucose at the location-, year-, age-, and sex- level. Updates to the ST-GR modelling framework for GBD 2017 are detailed elsewhere in the Appendix.

Fasting plasma glucose is frequently tested or reported in surveys aiming at assessing the prevalence of diabetes mellitus. In these surveys, the case definition of diabetes may include both a glucose test and questions about treatment for diabetes. People with positive history of diabetes treatment may be excluded from the FPG test. Thus, the mean FPG in these surveys would not represent the mean FPG in the entire population. To address this limitation, using the data from the surveys reporting mean FPG in the entire population, we estimated a regression-based correction factor and adjusted the mean FPG to account for diabetics in the population. We also use an ensemble distribution to characterize the distribution of FPG in the population and estimate the standard deviation based on mean FPG and prevalence of diabetics from the non-fatal diabetes mellitus model.

To inform our estimates in data-sparse countries, we systematically tested a range of covariates and selected two covariates based on AIC and adjusted R². These included prevalence of obesity and lag-distributed income per capita (LDI).

Mean FPG iss estimated using a mixed-effects linear regression, run separately by sex:

$$logit(FPG_{c,a,t}) = \beta_0 + \beta_1 log(LDI)_{c,t} + \beta_2 p_{overweight_{c,a,t}} + \sum_{k=2}^{16} \beta_k I_{A[a]} + \alpha_s + \alpha_r + \alpha_c + \epsilon_{c,a,t}$$

where $log(LDI)_{c,t}$ is the log of the lag-distributed income, $p_{overweight_{c,a,t}}$ is the prevalence of overweight, $I_{A[a]}$ is an indicator variable for a fixed effect on a given 5-year age group, and $\alpha_s \alpha_r \alpha_c$ are random effects at the super-region, region, and country level, respectively. The estimates were then propagated through the ST-GPR framework to obtain 1000 draws for each location, year, age, and sex.

Theoretical minimum-risk exposure level

The theoretical minimum-risk exposure level (TMREL) for FPG is 4.5-5.4 mmol/L. This was calculated by taking the person-year weighted average of the levels of FPG that were associated with the lowest risk of mortality in the pooled analyses of prospective cohort studies.¹

Relative risks

Risk	Outcome
Fasting plasma glucose	Ischemic heart disease
Fasting plasma glucose	Ischemic stroke
Fasting plasma glucose	Subarachnoid hemorrhage
Fasting plasma glucose	Intracerebral hemorrhage
Fasting plasma glucose	Peripheral vascular disease
Fasting plasma glucose	Type 1 diabetes
Fasting plasma glucose	Type 2 diabetes
Diabetes mellitus	Drug-resistant tuberculosis
Diabetes mellitus	Drug-susceptible tuberculosis
Diabetes mellitus	Multidrug-resistant tuberculosis
	without extensive drug resistance
Diabetes mellitus	Extensively drug-resistant
	tuberculosis
Diabetes mellitus	Liver cancer due to NASH
Diabetes mellitus	Liver cancer due to other causes
Diabetes mellitus	Pancreatic cancer
Diabetes mellitus	Ovarian cancer
Diabetes mellitus	Colorectal cancer

We estimate 15 outcomes due to high fasting plasma glucose (continuous risk) or diabetes (categorical risk).

Diabetes mellitus	Bladder cancer
Diabetes mellitus	Lung cancer
Diabetes mellitus	Breast cancer
Diabetes mellitus	Glaucoma
Diabetes mellitus	Cataracts
Diabetes mellitus	Dementia

Relative risks for High Fasting Plasma Glucose (continuous risk)

In GBD 2017, diabetes was further split into diabetes type 1 and diabetes type 2, and hemorrhagic stroke was further split into subarachnoid hemorrhage and intracerebral hemorrhage.

Relative risks (RR) were obtained from dose-response meta-analysis of prospective cohort studies. Please see the citation list for a full list of studies that are utilized. For cardiovascular outcomes, we estimated age-specific RRs using DisMod-MR 2.1 with log (RR) as the dependent variable and median age at event as the independent variable with an intercept at age 110. Morbidity and mortality directly caused by diabetes type 1 and diabetes type 2 is considered directly attributable to FPG.

Relative risks for Diabetes mellitus (Categorical risk)

Relative risks were obtained from meta-analysis of cohort studies. Please see the citation list for a full list of studies that are utilized.

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High LDL Cholesterol Capstone Appendix



Flowchart

Input Data & Methodological Summary

Exposure

Case Definition

In the GBD 2016 study, we estimated burden attributable to total cholesterol. For GBD 2017, we modelled blood concentration of low-density lipoprotein (LDL) in units of mmol/L.

Input Data

We utilized data on blood low-density lipoprotein from literature and from household survey microdata and reports. Please see the appendix for a full list of included sources. For the GBD 2016 study, we carried out an updated systematic review for total cholesterol (TC), drawing from the GHDx and Medline via PubMed. For GBD 2017, we reviewed all data sources in our database that reported data on total cholesterol and re-extracted LDL data when it was available. Data on high-density lipoprotein (HDL) and triglycerides (TGL) were also extracted when available.

Inclusion Criteria

Studies were included if they were population-based and measured total low-density lipoprotein using a blood test or calculated using the Friedewald equation. We assumed the data is representative of the location if the geography was not related to the diseases and if it is not an outlier compared to other data in the country or region.

Outliers

Data was utilized in the modeling process unless an assessment of data strongly suggested that the data

was biased. A candidate source was excluded if the quality of study did not warrant a valid estimate because of selection (non-representative populations) or if the study did not provide methodological details for evaluation. In a small number of cases, data point was considered to be an outlier candidate if the level was implausibly low or high based on expert judgement and other country data.

Data Extraction

Where possible, individual level data on LDL estimates were extracted from survey microdata and these were collapsed across demographic groupings to produce mean estimates in the standard GBD 5-year age-sex groups. If microdata were unavailable, information from survey reports or from literature were extracted along with any available measure of uncertainty including standard error, uncertainty intervals, and sample size. Standard deviations were also extracted. Where LDL was reported split out by groups other than age, sex, location, and year (eg, by diabetes status), a weighted mean was calculated.

Lipid crosswalk

Total cholesterol consists of three major components: LDL, HDL, and triglycerides. LDL is often calculated for an individual using the Friedewald equation, shown below:

$$LDL = TC - \left(HDL + \frac{TGL}{2.2}\right)$$

We utilized this relationship at the individual level to impute the mean LDL for a study population when only data on total cholesterol, HDL, and TGL were available. Because studies report different combinations of TC, HDL, and TGL, we constructed a single regression to utilize al available data to evaluate the relationship between each lipid and LDL at the population level. We used the following regression:

$$LDL = ind_{tc}\beta_{1}TC - (ind_{hdl}\beta_{2}HDL + ind_{tgl}\beta_{3}TGL) + \sum \alpha_{l}I_{l}$$

Where ind_{tc} , ind_{hdl} , and ind_{tgl} are indicator variables for whether data is available for a given lipid, I_l is an indicator variable a given set of available lipids l. α_l is a unique intercept for each set of available lipids. For example, for sources that only reported TC and HDL, $\alpha_{l=TC,HDL}$ should account for the missing lipid data, ie, TGL. The form of this regression allows us to estimate the betas for each lipid using all available data. As a sensitivity analysis, we also ran separate regressions for each set of available lipids and found that the single regression method had much lower root-mean squared error. A comparison of the observed vs predicted LDL for each set of available lipids is shown in Figure 1. We found almost no relationship between LDL and HDL or TGL when TC was not available, so only studies that reported TC were crosswalked to LDL.





Incorporating United States prevalence data

Survey reports and literature often report information only about the prevalence, but not the level, of hypercholesterolemia in the population studied. These sources were not used to model LDL, with the exception of data from the Behavioral Risk Factors Surveillance System (BRFSS) because of the availability of a similarly structured exam survey covering the identical population (NHANES). BRFSS is a telephone survey conducted in the United States for all counties. It collects self-reported diagnosis of hypercholesterolemia. These self-reported values of prevalence of raised total cholesterol in each age group, sex, US state, and year were used to predict a mean total cholesterol for the same strata with a regression using data from the National Health and Nutrition Examination Survey, a nationally representative health examination survey of the US adult population. The regression was:

$$TC_{l,a,t,s} = \beta_0 + \beta_1 prev_{l,a,t,s}$$

where $TC_{l,a,t,s}$ is the location, age, time, and sex specific mean total cholesterol and $prev_{l,a,t,s}$ is the location, age, time, and sex specific prevalence of raised total cholesterol. The coefficients for both models are reported in Table 1.
Table 2. Coefficients in the sex-specific US states TC prediction models

Term	Male model	Female model
Intercept	4.23	4.36
Prevalence	6.25	5.22

Out of sample RMSE was used to quantify the predictive validity of the model. The regression was repeated 10 times for each sex, each time randomly holding out 20% of the data. The RMSEs from each holdout analysis were averaged to get the average out of sample RMSE. The results of this holdout analysis are reported in Table 2. Total cholesterol estimates were crosswalked to LDL using the lipid crosswalk reported above.

	Table 3. Out of sa	imple RMSEs of the	sex-specific US states	TC prediction models
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	Male model	Female model
Out of sample RMSE	0.21 mmol/L	0.20 mmol/L

Age and Sex Splitting

Prior to modeling, data provided in age groups wider than the GBD 5-year age groups were processed using the approach outlined in Ng et al.² Briefly, age-sex patterns were identified using person-level microdata (58 sources), and estimate age-sex specific levels of total cholesterol from aggregated results reported in published literature or survey reports. In order to incorporate uncertainty into this process and borrow strength across age groups when constructing the age-sex pattern, we used a model with auto-regression on the change in mean LDL over age groups:

$$u_a = \mu_{a-1} + \omega_a$$
$$\omega_a \sim N(\omega_{a-1}, \tau)$$

Where μ_a is the mean predicted value for age group a, μ_{a-1} is the mean predicted value for the age group previous to age group a, ω_a is the difference in mean between age group a and age group a-1, ω_{a-1} is the difference between age group a-1 and age group a-2, and τ is a user-input prior on how quickly the mean LDL changes for each unit increase in age. We used a τ of 0.09 mmol/L for this model. Draws of the age-sex pattern were combined with draws of the input data needing to be split in order to calculate the new variance of age-sex split data points.

Modeling

Exposure estimates were produced from 1980 to 2017 for each national and subnational location, sex, and for each 5-year age group starting from 25+. As in GBD 2016, we used a Spatio-Temporal Gaussian Process Regression (ST-GPR) framework to model the mean LDL at the location-, year-, age-, sex- level. Updates to the ST-GR modeling framework for GBD 2017 are detailed in the appendix.

Covariate selection

The first step of the ST-GPR framework requires the creation of a linear model for predicting LDL the location-, year-, age-, sex- level. Covariates for this model were selected in two stages. First a list of

variables with an expected causal relationship with LDL was created based on significant association found within high-quality prospective cohort studies reported in the published scientific literature. The second stage in covariate selection was to test the predictive validity of every possible combination of covariates in the linear model, given the covariates selected above. This was done separately for each sex. Predictive validity was measured with out of sample root-mean-squared error.

In GBD 2016, the linear model with the lowest root-mean squared error for each sex was then used in the ST-GPR model. In GBD 2017, we used an ensemble model of the 50 models with the lowest root-mean squared error for each sex. This allows us to utilize covariate information from many plausible linear mixed-effects models. The 50 models were each used to predict the mean LDL for every age, sex, location, and year, and the inverse-RMSE-weighted average of this set of 50 predictions was used as the linear prior. The relative weight in 'draws' contributed by each covariate is plotted by sex in Figure 2.



Figure 2. Results of the ensemble linear model covariate selection

Methodological updates to ST-GPR are reported above. The result of the ST-GPR model are estimates of the mean LDL for each age, sex, location, and year.

Estimate of standard deviation

The standard deviation of LDL within a population was estimated for each national and subnational location, sex, and 5-year age group starting from age 25 using the standard deviation from person-level and some tabulated data sources. Person-level microdata accounted for 3009 of the total 4001 rows of data on standard deviation. The remaining 992 rows came from tabulated data. Tabulated data was only used to model standard deviation if it was sex and 5-year age group specific and reported a population standard deviation LDL. The LDL standard deviation function was estimated using a linear regression:

$$log(SD_{c,a,t,s}) = \beta_0 + \beta_1 log(mean_LDL_{c,a,t,s}) + \beta_4 sex + \sum_{k=2}^{16} \beta_k I_{A[a]}$$

where mean_LDL_{c,a,t,s} is the country, age, time, and sex specific mean LDL estimate from ST-GPR, and $I_{A[a]}$ is a dummy variable for a fixed effect on a given 5-year age group,.

Distribution shape modelling

The shape of the distribution of LDL was estimated using all available person-level microdata sources, which was a subset of the input data into the modelling process. The distribution shape modelling framework for GBD 2017 is detailed in the appendix. Briefly, an ensemble distribution created from a weighted average of distribution families was fit for each individual microdata source, separately by sex. The weights for the distribution families for each individual source were then averaged and weighted to create a global ensemble distribution for each sex.

Theoretical minimum-risk exposure level

A Meta-analysis of randomized trials has shown that outcomes can be improved even at low levels of LDL-cholesterol, below 1.3 mmol/l.³ Recent studies of PCSK-9 inhibitors support these results.⁴ We therefore used a TMREL with a uniform distribution between 0.7-1.3 mmol/l.

Relative risks

After a systematic search, we were unable to find relative risks for LDL that were reported by age and level of LDL. Given this evidence that the relative risks for LDL and TC are very similar⁵ and the strong linear correlation between TC and LDL at the individual level, we used relative risks reported for TC to approximate the relative risks for LDL. We used Dismod-MR 2.1 to pool effect sizes from included studies and generate a dose-response curve for each of the outcomes associated with LDL. The tool enabled us to incorporate random effects across studies and include data with different age ranges. RRs were used universally for all countries and produce RRs with uncertainty and covariance across ages, taking into account the uncertainty of the data points. As in GBD 2016, RRs for IHD and ischemic stroke are obtained from meta-regressions of pooled epidemiological studies: the Asia Pacific Cohort Studies Collaboration (APCSC) and the Prospective Studies Collaboration (PSC).⁶ RRs for IHD were modeled with log (RR) as the dependent variable and median age at event as the independent variable with an age intercept (RR equals 1) at age 110. For LDL and ischemic stroke, a similar approach was used, except that there was no age intercept at age 110, due to the fact that there was no statistically significant

relationship between LDL and stroke after age 70 with a mean RR less than one. We assumed that there is not a protective effect of LDL and therefore did not include an RR for ages 80+.

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High Systolic Blood Pressure Capstone Appendix

Flowchart



Input Data & Methodological Summary

Exposure

Case Definition

Brachial systolic blood pressure in mmHg.

Input Data

We utilised data on mean systolic blood pressure from literature and from household survey microdata and reports (e.g. STEPS, NHANES). Please see the appendix for a full list of included sources. For GBD 2017, we did not carry out a systematic review of the literature for new data. In total, we have utilised 934 sources corresponding to 49,690 unique data points.

Inclusion Criteria

Studies were included if they were population-based and measured systolic blood pressure using a blood test. We assumed the data is representative of the location if the geography was not selected because it was related to the diseases.

Outliers

Data was utilised in the modelling process unless an assessment of data strongly suggested that the data was biased. A candidate source was excluded if the quality of study did not warrant a valid estimate because of selection (non-representative populations) or if the study did not provide methodological details for evaluation. In a small number of cases, a data point was considered to be an outlier candidate if the level was implausibly low or high based on expert judgement and data from other country data.

Data Extraction

Where possible, individual level data on blood pressure estimates were extracted from survey microdata and these were collapsed across individuals and collapsed across demographic groupings to produce mean estimates in the standard GBD 5-year age-sex groups. If microdata were unavailable, information from survey reports or from literature were extracted along with any available measure of uncertainty including standard error, uncertainty intervals, and sample size. Standard deviations were also extracted. Where mean systolic blood pressure was reported split out by groups other than age, sex, location, and year (eg, by hypertensive status), a weighted mean was calculated.

Incorporating United States prevalence data

Survey reports and literature often report information only about the prevalence, but not the level, of hypertension in the population studied. These sources were not used to model systolic blood pressure, with the exception of data from the Behavioral Risk Factors Surveillance System (BRFSS) because of the availability of a similarly structured exam survey that is representative of the same population (NHANES). BRFSS is a telephone survey conducted in the United States for all US counties. It collects self-reported diagnosis of hypertension. These self-reported values of prevalence of raised blood pressure were adjusted for self-report bias and tabulated by age group, sex, US state, and year. These prevalences were used to predict a mean systolic blood pressure for the same strata with a regression using data from the National Health and Nutrition Examination Survey, a nationally representative health examination survey of the US adult population. The regression was run separately by sex, and was specified as:

$$SBP_{l,a,t,s} = \beta_0 + \beta_1 prev_{l,a,t,s}$$

where $SBP_{l,a,t,s}$ is the location, age, time, and sex specific mean systolic blood pressure and $prev_{l,a,t,s}$ is the location, age, time, and sex specific prevalence of raised blood pressure. The coefficients for both models are reported in Table 1.

Term	Male model	Female model
Intercept (β_0)	114.65	108.28
Prevalence (β_1)	51.86	68.87

rabic 1. Coefficients in the sex specific of states blood pressure prediction model	Table 1.	Coefficients	in the sex-	specific US	S states blood	pressure	prediction	models
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Out of sample RMSE was used to quantify the predictive validity of the model. The regression was repeated 10 times for each sex, each time randomly holding out 20% of the data. The RMSEs from each holdout analysis were averaged to get the average out of sample RMSE. The results of this holdout analysis are reported in Table 2.

Table 2. Out of sample RMSEs of the sex-specific US states blood pressure prediction models

	Male model	Female model
Out of sample RMSE	2.37 mmHg	3.27 mmHg

Age and Sex Splitting

Prior to modelling, data provided in age groups wider than the GBD 5-year age groups were processed using the approach outlined in Ng et al.² Briefly, an age-sex pattern were identified using 115 sources of microdata with multiple age-sex groups, and these patterns were applied to estimate age-sex specific levels of mean systolic blood pressure from aggregated results reported in published literature or survey reports. In order to incorporate uncertainty into this process and borrow strength across age groups when constructing the age-sex pattern, we used a model with auto-regression on the change in mean SBP over age groups:

$$\mu_a = \mu_{a-1} + \omega_a$$
$$\omega_a \sim N(\omega_{a-1}, \tau)$$

Where μ_a is the mean predicted value for age group a, μ_{a-1} is the mean predicted value for the age group previous to age group a, ω_a is the difference in mean between age group a and age group a-1, ω_{a-1} is the difference between age group a-1 and age group a-2, and τ is a user-input prior on how quickly the mean SBP changes for each unit increase in age. We used a τ of 1.5 mmHg for this model. Draws of the age-sex pattern were combined with draws of the input data needing to be split in order to calculate the new variance of age-sex split data points.

Modelling

Exposure estimates were produced from 1980 to 2016 for each national and subnational location, sex, and for each 5-year age group starting from 25+. As in GBD 2016, we used a Spatio-Temporal Gaussian Process Regression (ST-GPR) framework to model the mean systolic blood pressure at the location-, year-, age-, sex- level.

Covariate selection

The first step of the ST-GPR framework requires the creation of a linear model for predicting SBP at the location-, year-, age-, sex- level. Covariates for this model were selected in two stages. First a list of variables with an expected causal relationship with SBP was created based on significant association found within high-quality prospective cohort studies reported in the published scientific literature. The second stage in covariate selection was to test the predictive validity of every possible combination of covariates in the linear model, given the covariates selected above. This was done separately for each sex. Predictive validity was measured with out of sample root-mean-squared error. In GBD 2016, the linear model with the lowest root-mean squared error for each sex was then used in

the ST-GPR model. In GBD 2017, we used an ensemble model of the 50 models with the lowest rootmean squared error for each sex. This allows us to utilise covariate information from many plausible linear mixed-effects models. The 50 models were each used to predict the mean SBP for every age, sex, location, and year, and the inverse-RMSE-weighted average of this set of 50 predictions was used as the linear prior. The relative weight in 'draws' contributed by each covariate is plotted by sex in Figure 2.



Figure 2. Results of the ensemble linear model covariate selection

The results of the ensemble linear model were used for the first stage in an ST-GPR model. Methodological updates to ST-GPR are reported above. The result of the ST-GPR model are estimates of the mean SBP for each age, sex, location, and year.

Estimate of Standard Deviation

Currently, the ST-GPR model only produces an estimate of mean exposure level without standard deviation. Therefore, the standard deviation of systolic blood pressure within a population was estimated for each national and subnational location, sex, and 5-year age group starting from age 25 using the standard deviation from person-level and some tabulated data sources. Person-level microdata accounted for 10375 of the total 12570 rows of data on standard deviation. The remaining 2195 rows came from tabulated data. Tabulated data was only used to model standard deviation if it was sex and 5-year age group specific and reported a population standard deviation of systolic blood pressure. The systolic blood pressure standard deviation function was estimated using a linear regression:

$$\log(SD_{l,a,t,s}) = \beta_0 + \beta_1 \log(\text{mean}_SBP_{l,a,t,s}) + \beta_4 \text{sex} + \sum_{k=2}^{16} \beta_k I_A$$

where mean_SBP_{1,a,t,s} is the location, age, time, and sex specific mean SBP estimate from ST-GPR, and I_A is a dummy variable for a fixed effect on a given 5-year age group.

Adjustment for Usual Levels of Blood Pressure

To account for in-person variation in systolic blood pressure, a 'usual blood pressure' adjustment was done. The need for this adjustment has been described elsewhere.5 Briefly, measurements of a risk factor taken at a single time point may not accurately capture an individual's true long-term exposure to that risk. Blood pressure readings are highly variable over time due to measurement error as well as diurnal, seasonal, or biological variation. These sources of variation result in an over-estimation of the variation in cross-sectional studies of the distribution of SBP.

To adjust for this overestimation, we applied a correction factor to each location-, age-, time-, and sexspecific standard deviation. These correction factors were age-specific, and represented the proportion of the variation in blood pressure within a population that would be observed if there were no withinperson variation across time. Four longitudinal surveys were used to estimate these factors: the China Health and Retirement Longitudinal Survey (CHRLS), the Indonesia Family Life Survey (IFLS), the National Health and Nutrition Examination Survey I Epidemiological Follow-up Study (NHANES I/EFS), and the South Africa National Income Dynamics Survey (NIDS). The sample size and number of blood pressure measurements at each measurement period for each survey is reported in Table 5.

Source	Measurement periods	Number of measurements	Sample size
CHRLS	2008	3	1967
	2012	3	1419
IFLS	1997	1	19418
	2000	1	16626
	2007	3	14136
NIDS	1997	2	14084
	2000	2	9612
	2007	2	9098
NHANES I/EFS	1971-1976	2	20716
	1982-1984	3	9932

Table 3. Characteristics of longitudinal surveys used for the usual blood pressure adjustment

For each survey, the following regression was created for each age group:

$$SBP_{i,a} = \beta_0 + \beta_1 sex + \beta_3 age + + \upsilon_i$$

where SBP_{i,a} is the systolic blood pressure of an individual i at age a, sex is a dummy variable for the sex of an individual, age is a continuous variable for the age of an individual, and v_i is a random intercept for each individual. Then, a blood pressure value $\widehat{SBP}_{i,b}$ was predicted for each individual i for his/her age at baseline b. The correction factor cf for each age group within each survey was calculated as variation in these predicted blood pressures was divided by the variation in the observed blood pressures at baseline, SBP_{i,b}:

$$cf = \sqrt{\frac{var(S\widehat{BP}_{b})}{var(SBP_{b})}}$$

The average of the correction factors was taken over the three surveys to get one set of age-specific correction factors, which were then multiplied by the square of the modelled standard deviations to estimate standard deviation of the 'usual blood pressure' of each age, sex, location, and year. Because of low sample sizes, the correction factors for the 75-79 age group was used for all terminal age groups. The final correction factors for each age group are reported in Table 6. Figure 1 shows the correction factors by survey and age group ID.

Age group	Correction factor
25-29	.665
30-34	.713
35-39	.737
40-44	.733
45-49	.798
50-54	.771
55-59	.764
60-64	.753
65-69	.719
70-74	.689
75+	.678

Table 4. Age-specific usual blood pressure correction factors



Figure 1: Correction factor by survey and age group id. The correction factor is equal to the variance of the predictions divided by the variance of the raw dataset. In pink is the average correction factor for each age group, summarised in Table 6.

A visualisation of how the uncorrected blood pressure measurements overestimate the 'usual' blood pressure variation is shown in Figure 1. This image shows the density of the distribution of the observed blood pressure values $SBP_{i,b}$ in participants in the Indonesian Family Life Study survey in red, and the density of the predicted blood pressure values $SBP_{i,b}$ in blue. The ratio of the variance of the blue distribution to the variance of the red distribution is an example of the scalar adjustment factor being applied to the modelled standard deviations.



Figure 2: Raw and predicted distributions of blood pressure in the Indonesia Family Life Survey

Estimating the exposure distribution shape

The shape of the distribution of systolic blood pressure was estimated using all available person-level microdata sources, which was a subset of the input data into the modelling process. The distribution shape modelling framework for GBD 2017 is detailed in the appendix. Briefly, an ensemble distribution created from a weighted average of distribution families was fit for each individual microdata source, separately by sex. The weights for the distribution families for each individual source were then averaged and weighted to create a global ensemble distribution for each sex.

Theoretical minimum-risk exposure level

No changes were made to TMREL used in the GBD2015 study. We estimated that the TMREL of SBP ranges from 110 to 115 mm Hg based on pooled prospective cohort studies that show risk of mortality increases for SBP above that level.^{3,4} Our selection of a TMREL of 110-115 mmHg is consistent with the GBD study approach of estimating all attributable health loss that could be prevented even if current interventions do not exist that can achieve such a change in exposure level, for example a tobacco smoking prevalence of zero percent. To include the uncertainty in the TMREL, we took a random draw from the uniform distribution of the interval between 110 mm and 115 mm Hg each time the population attributable burden was calculated.

Relative risks

No change was made to RR for blood pressure outcomes used in the GBD2016 study. RRs for chronic kidney disease are from the Renal Risk Collaboration meta-analysis of 2.7 million individuals in 106 cohorts. For other outcomes, we used data from two pooled epidemiological studies: the Asia Pacific Cohort Studies Collaboration (APCSC) and the Prospective Studies Collaboration (PSC)^{-4,5} Additional estimates of RR for cardiovascular outcomes were used from the CALIBER study, a health-record linkage cohort study from the UK.⁶

For cardiovascular disease, epidemiological studies have shown that the RR associated with SBP declines with age, with the log (RR) having an approximately linear relationship with age and reaching a value of 1 between the ages of 100 and 120. RRs were reported per 10 mm Hg increase in SBP above the TMREL value (115 mm Hg), calculated as in the equation below:

 $RR(x) = RR_0^{\frac{(x - TMREL)}{10 \ mmHg}}$

Where RR(x) is the RR at exposure level x and RR_0 is the increase in RR for each 10 mmHg above the TMREL. We used Dismod-MR 2.1 to pool effect sizes from included studies and generate a dose-response curve for each of the outcomes associated with high SBP. The tool enabled us to incorporate random effects across studies and include data with different age ranges. RRs were used universally for all countries and the meta-regression only helped to pool the three major sources and produce RRs with uncertainty and covariance across ages taking into account the uncertainty of the data points.

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High Body-Mass Index Capstone Appendix

Flowchart



Adult (Ages 20+) High Body-Mass Index: Data and Model Flow Chart

Childhood (Ages 2-19) High Body-Mass Index: Data and Model Flow Chart ng the distribution of BMI by age, sex, cos try, and tim weight Cross-validation and hyperparameter selection ST-GPR - saului Survey microdata Self-report Age and sex Survey reports ages 15-19 Cross-validation and hyperparameter selection far ng those weight): ST-GPR Systematic literature review Linear Model antifying the effect size of BMI on disease endpoints Published meta-analyses Assessment of strength of evidence supporting causality Published system Relative risks Meta-analysis PAF calculatio DALYnator Published pooled analyses etermining the optimal level of BMI Legend Published meta-TMREL: BMI < 25 kg/m² Inipia Process analyses

Case definitions

High body-mass index (BMI) for adults (ages 20+) is defined as BMI greater than 20 to 25 kg/m². High BMI for children (ages 1-19) is defined as being overweight or obese based on International Obesity Task Force standards.

Input data and methodological summary

Data sources

We systematically searched Medline to identify studies providing nationally or subnationally representative estimates of overweight prevalence, obesity prevalence or mean body-mass index (BMI). We limited the search to literature published between January 1, 2016 and December 31, 2016 to update the systematic literature search previously performed as part of GBD 2015.

The search for adults was conducted on 4 January 2017 using the following terms:

((("Body Mass Index"[Mesh] OR "Overweight"[Mesh] OR "Obesity"[Mesh]) AND ("Geographic Locations"[Mesh] NOT "United States"[Mesh]) AND ("humans"[Mesh] AND "adult"[MeSH]) AND ("Data Collection"[Mesh] OR "Health Services Research"[Mesh] OR "Population Surveillance"[Mesh] OR "Vital statistics"[Mesh] OR "Population"[Mesh] OR "Epidemiology"[Mesh] OR "surve*"[TiAb]) NOT (Comment[ptyp] OR Case Reports[ptyp] OR "hospital"[TiAb])) AND ("2016/01/01"[Date - Publication] : "2016/12/31"[Date - Publication]))

The search for children was conducted on 4 August 2016 using the following terms: ((("Body Mass Index"[Mesh] OR "Overweight"[Mesh] OR "Obesity"[Mesh]) AND ("Geographic Locations"[Mesh] NOT "United States"[Mesh]) AND ("humans"[Mesh] AND "child"[MeSH]) AND ("Data Collection"[Mesh] OR "Health Services Research"[Mesh] OR "Population Surveillance"[Mesh] OR "Vital statistics"[Mesh] OR "Population"[Mesh] OR "Epidemiology"[Mesh] OR "surve*"[TiAb]) NOT (Comment[ptyp] OR Case Reports[ptyp] OR "hospital"[TiAb])) AND ("2016/01/01"[Date - Publication] : "2016/12/31"[Date - Publication]))

Our search for adult estimates identified 456 abstracts, of which 25 met inclusion criteria and were extracted. The search for childhood estimates identified 137 articles, of which 4 were extracted. Including sources from the previous GBD systematic literature searches, a total of 11,220 articles were identified, of which 845 were included. Additionally, we searched the Global Health Data Exchange (GHDx) database for individual-level data from major multinational survey series or country-specific surveys and identified 5,385 location-year sources meeting the inclusion criteria.

Eligibility criteria

We included representative studies providing data on mean BMI or prevalence of overweight or obesity among adults or children. For adults, studies were included if they defined overweight as BMI≥25 kg/m² and obesity as BMI≥30 kg/m², or if estimates using those cutoffs could be back-calculated from reported categories. For children (children ages 2-18), studies were included if they used International Obesity Task Force (IOTF) standards to define overweight and obesity thresholds. We only included studies reporting data collected between January 1, 1980 and December 31, 2016. Studies were excluded if they used nonrandom samples (e.g. case-control studies or convenience samples), conducted among specific subpopulations (e.g. pregnant women, racial or ethnic minorities, immigrants, or individuals with specific diseases), used alternative methods to assess adiposity (e.g. waist-circumference, skin-fold thickness, or hydrodensitometry), had sample sizes of less than 20 per age-sex group, or provided inadequate information on any of the inclusion criteria. We also excluded review articles and non-English language articles.

Data collection process

Where individual-level survey data were available, we computed mean BMI using weight and height. We then used BMI to determine the prevalence of overweight and obesity. For individuals aged over 18 years, we considered them to be overweight if their BMI was greater than or equal to 25 kg/m², and obese if their BMI was greater than or equal to 30 kg/m². For individuals aged 2 to 18 years, we used monthly IOTF cutoffs² to determine overweight and obese status when age in months was available. When only age in years was available, we used the cutoff for the midpoint of that year. Obese individuals were also considered to be overweight. We excluded studies using the World Health Organization (WHO) standards or country-specific cutoffs to define childhood overweight and obesity. At the individual-level, we considered BMI<10 kg/m² and BMI>70 kg/m² to be biologically implausible and excluded those observations.

The rationale for choosing to use the IOTF cutoffs over the WHO standards has been described elsewhere.¹ Briefly, the IOTF cutoffs provide consistent child-specific standards for ages 2-18 derived surveys covering multiple countries. On the other hand, the WHO growth standards apply to children under age 5 and the WHO growth reference applies to children ages 5 to 19. The WHO growth reference for children ages 5 to 19 was derived from United States data which is less representative than the multinational data used by IOTF. Additionally, the switch between references at age 5 can produce artificial discontinuities. Given that we estimate global childhood overweight and obesity for ages 2-19 (with ages 19 using standard adult cutoffs), the IOTF cutoffs were preferable. Additionally, we found that IOTF cutoffs were more commonly used in scientific literature covering childhood obesity.

From report and literature data, we extracted data on mean BMI, prevalence of overweight, and prevalence of obesity, measures of uncertainty for each, and sample size, by the most granular age and sex groups available. Additionally, we extracted the same study-level covariates as were extracted from microdata (measurement, urbanicity, and representativeness), as well as location and year.

In addition to the primary indicators described above, we extracted relevant survey-design variables, including primary sampling unit, strata, and survey weights, which were used to tabulate individual-level microdata and produce accurate measures of uncertainty. We extracted three study-level covariates: 1) whether height and weight data were measured or self-reported; 2) whether the study was predominantly conducted in an urban area, rural area or both; and 3) the level of representativeness of the study (national or subnational).

Finally, we extracted relevant demographic indicators, including location, year, age and sex. We estimated the standard error of the mean from individual-level data, where available, and used the reported standard error of the mean for published data. When multiple data sources were available for the same country, we included all of them in our analysis. If data from the same data source were available in multiple formats such individual-level data and tabulated data, we used individual-level data.

Self-report bias adjustment

We included both measured and self-reported data. We tested for bias in self-report data compared to measured data, which is considered to be the gold-standard. There was no clear direction of bias for children ages 2 to 14, so for these age groups we only included measured data. For individuals ages 15

and above, we adjusted self-reported data for overweight prevalence, obesity prevalence and mean BMI using the following nested hierarchical mixed-effects regression models, fit using restricted maximum likelihood separately by sex:

$$\begin{split} \text{logit}(\text{overweight})_{c,a,t} &= \beta_0 + \beta_1 m + \sum_{k=2}^{19} \beta_k I_{A[a]} + \sum_{j=20}^{55} \beta_j I_{A[a]} I_{M[m]} + \alpha_s + \alpha_s m + \alpha_r + \alpha_r m + \alpha_c + \alpha_c m + \alpha_t + \alpha_t m + \varepsilon_{c,a,t} \\ \text{logit}(\text{obesity})_{c,a,t} &= \beta_0 + \beta_1 m + \sum_{k=2}^{19} \beta_k I_{A[a]} + \sum_{l=20}^{55} \beta_l I_{A[a]} I_{M[m]} + \alpha_s + \alpha_s m + \alpha_r + \alpha_r m + \alpha_c + \alpha_c m + \alpha_t + \alpha_t m + \varepsilon_{c,a,t} \\ \text{log}(\text{BMI})_{c,a,t} &= \beta_0 + \beta_1 m + \sum_{k=2}^{19} \beta_k I_{A[a]} + \sum_{l=20}^{55} \beta_l I_{A[a]} I_{M[m]} + \alpha_s + \alpha_s m + \alpha_r + \alpha_r m + \alpha_c + \alpha_c m + \alpha_t + \alpha_t m + \varepsilon_{c,a,t} \end{split}$$

Where m is a fixed effect on measurement (binary, either measured (1) or self-report (0)), $I_{A[a]}$ is an indicator variable for specific age group A, $I_{A[a]}I_{M[m]}$ is an interaction term between age and measurement, α_s , α_r , and α_c are random effects at the super region, region, and country, respectively, and α_t is a random effect by time-period (1980-1989, 1990-1999, 2000-2009, 2010-2017). Random effects at the country level and time-period level were used to fit the models, but were taken as noise and were not used in adjustment of self-reported data. We propagated the uncertainty in the self-report adjustment model by adding the variance of each of the regression coefficients used in adjustment to the data variance in delta-transformed space. After adjustment, regressions confirmed that self-reported data was no longer significantly different from measured data.

Age and sex splitting

Any report or literature data provided in age groups wider than the standard 5-year age groups or as both sexes combined were split using the approach used by Ng et al.² Briefly, age-sex patterns were identified using sources with data on multiple age-sex groups and these patterns were applied to split aggregated report and literature data. Uncertainty in the age-sex split was propagated by multiplying the standard error of the data by the square root of the number of splits performed. We did not propagate the uncertainty in the age pattern used to split the data as they seemed to have small effect.

Prevalence estimation for overweight and obesity

After adjusting for self-report bias and splitting aggregated data into 5-year age-sex groups, we used spatiotemporal Gaussian process regression (ST-GPR) to estimate the prevalence of overweight and obesity. This modelling approach has been described in detail elsewhere.

The linear model, which when added to the smoothed residuals forms the mean prior for GPR is as follows:

$$logit(overweight)_{c,a,t} = \beta_0 + \beta_1 energy_{c,t} + \beta_2 SDI_{c,t} + \beta_3 vehicles_{c,t} + \beta_4 agriculture_{c,t} + \sum_{k=5}^{21} \beta_k I_{A[a]} + \alpha_s + \alpha_r + \alpha_c \\ logit(obesity/overweight)_{c,a,t} = \beta_0 + \beta_1 energy_{c,t} + \beta_2 SDI_{c,t} + \beta_3 vehicles_{c,t} + \sum_{k=4}^{21} \beta_k I_{A[a]} + \alpha_s + \alpha_r + \alpha_c \\ logit(obesity/overweight)_{c,a,t} = \beta_0 + \beta_1 energy_{c,t} + \beta_2 SDI_{c,t} + \beta_3 vehicles_{c,t} + \sum_{k=4}^{21} \beta_k I_{A[a]} + \alpha_s + \alpha_r + \alpha_c \\ logit(obesity/overweight)_{c,a,t} = \beta_0 + \beta_1 energy_{c,t} + \beta_2 SDI_{c,t} + \beta_3 vehicles_{c,t} + \beta_4 agriculture_{c,t} + \sum_{k=4}^{21} \beta_k I_{A[a]} + \alpha_s + \alpha_r + \alpha_c \\ logit(obesity/overweight)_{c,a,t} = \beta_0 + \beta_1 energy_{c,t} + \beta_2 SDI_{c,t} + \beta_3 vehicles_{c,t} + \beta_4 agriculture_{c,t} + \beta_4 agricu$$

where *energy* is ten-year lag-distributed energy consumption per capita, *SDI* is a composite index of development including lag-distributed income per capita, education, and fertility, *vehicles* is is the number of two or four-wheel vehicles per capita, and *agriculture* is the proportion of the population working in agriculture. $I_{A[a]}$ is a dummy variable indicating specific age group A that the prevalence point

captures, and α_s , α_r , and α_c are super region, region, and country random intercepts, respectively. Random effects were used in model fitting but were not used in prediction.

We tested all combinations of the following covariates to see which performed best in terms of in-sample AIC for the overweight linear model and the obesity as a proportion of overweight linear model: ten-year lag distributed energy per capita, proportion of the population living in urban areas, SDI, lag-distributed income per capita, educational attainment (years) per capita, proportion of the population working in agriculture, grams of sugar adjusted for energy per capita, grams of sugar not adjusted for energy per capita, and the number of two or four-wheeled vehicles per capita. We selected these candidate covariates based on theory as well as reviewing covariates used in other publications. The final linear model was selected based on: 1) if the direction of covariates matched what is expected from theory, 2) all the included covariates were significant, and 3) minimising in-sample AIC. The covariate selection process was performed using the dredge package in R.

The new version of ST-GPR for GBD 2017 incorporates information about data density into the process for smoothing over space and time. Estimates in areas/years with few observations have more weight on regional observations. To specify the distribution of time weights and space weights, we used values of lambda=0.2 and zeta=0.05, respectively. We used a value of omega=1.0 for the distribution of age weights. We set the GPR scale parameter to 20, and used the default global cutoff setting for amplitude.

Estimating mean BMI

To estimate the mean BMI for adults in each country, age, sex, and time period 1980-2017, we first used the following nested hierarchical mixed-effects model, fit using restricted maximum likelihood on data from sources containing estimates of all three indicators (prevalence of overweight, prevalence of obesity, and mean BMI), in order to characterise the relationship between overweight, obesity, and mean BMI:

$$\begin{split} log(BMI_{c,a,s,t}) &= \beta_0 + \beta_1 ow_{c,a,s,t} + \beta_2 ob_{c,a,s,t} + \beta_3 sex + \sum_{k=4}^{20} \beta_k I_{A[a]} + \alpha_s (1 + ow_{c,a,s,t} + ob_{c,a,s,t}) + \alpha_r (1 + ow_{c,a,s,t} + ob_{c,a,s,t}) + \alpha_c (1 + ow_{c,a,s,t} + ob_{c,a,s,t}) + \varepsilon_{c,a,s,t} \end{split}$$

where $ow_{c,a,s,t}$ is the prevalence of overweight in country c, age a, sex s, and year t, $ob_{c,a,s,t}$ is the prevalence of obesity in country c, age a, sex s, and year t, sex is a fixed effect on sex, $I_{A[a]}$ is an indicator variable for age, and α_s , α_r , and α_c are random effects at the super region, region, and country, respectively. The model was run in Stata 13.

We applied 1,000 draws of the regression coefficients to the 1,000 draws of overweight prevalence and obesity prevalence produced through ST-GPR to estimate 1,000 draws of mean BMI for each country, year, age, and sex. This approach ensured that overweight prevalence, obesity prevalence, and mean BMI were correlated at the draw level and uncertainty was propagated.

Estimating BMI distribution

We used the ensemble distribution approach described in the manuscript. We fit ensemble weights by source and sex, with source- and sex-specific weights averaged across all sources included to produce the final global weights. The ensemble weights were fit on measured microdata. The final ensemble weights were: exponential = 0.002, gamma = 0.028, inverse gamma = 0.085, log-logistic = 0.187, Gumbel = 0.220,

Weibull = 0.011, log-normal = 0.058, normal = 0.012, beta = 0.136, mirror gamma = 0.008, and mirror Gumbel = 0.113.

One thousand draws of BMI distributions for each location, year, age group, and sex estimated were produced by fitting an ensemble distribution using 1,000 draws of estimated mean BMI, 1,000 draws of estimated standard deviation, and the ensemble weights. Estimated standard deviation was produced by optimising a standard deviation to fit estimated overweight prevalence draws and estimated obesity prevalence draws.

Assessment of risk-outcome pairs

Risk-outcome pairs were defined based on strength of available evidence supporting a causal effect. We performed a systematic review of published meta-analyses, pooled analyses, and systematic reviews available through PubMed using the following search string: ("Body Mass Index"[Mesh] OR "Overweight"[Mesh] OR "Obesity"[Mesh]) AND (Meta-Analysis[ptyp] OR "systematic review"[tiab] OR "pooled analysis"[tiab]). Inclusion criteria are 1) the health outcome is included in GBD, 2) at least one prospective cohort is included, and 3) that the summary effect size is statistically significant. For outcomes meeting inclusion criteria we completed causal criteria tables to evaluate the strength of evidence supporting a causal relationship (see Appendix Table 4). Gallbladder disease, cataract, multiple myeloma, gout, non-Hodgkin lymphoma, asthma, Alzheimer disease, and atrial fibrillation were added as new outcomes in GBD 2016, resulting in a total of 38 outcomes.

Theoretical minimum risk exposure level

For adults (ages 20+), the theoretical minimum risk exposure level (TMREL) of BMI (20-25 kg/m²) was determined based on the BMI level that was associated with the lowest risk of all-cause mortality in prospective cohort studies.³

For children (ages 2-19), the TMREL is "normal weight," that is, not overweight or obese, based on IOTF cutoffs.

Relative risk

The relative risk per 5-unit change in BMI for each disease endpoint was obtained from meta-analyses, and where available, pooled analyses of prospective observational studies. In cases where a relative risk per 5-unit change in BMI was not available we computed our own dose-response meta-analysis using two-step generalised least squares for time trends estimation methods.

For childhood outcomes (ages 2-19), we computed categorical relative risks for overweight and obesity using a random effects meta-analysis.

Relative risks for all 38 outcomes, by age and sex, are reported in Table 6a.

References

- 1.) Cole, TJ, and T Lobstein. "Extended International (IOTF) Body Mass Index Cut-Offs for Thinness, Overweight and Obesity." Pediatric Obesity 2012; 7(4): 284–94.
- 2.) Ng M, Fleming T, Robinson M, et al. Global, regional, and national prevalence of overweight and obesity in children and adults during 1980–2013: a systematic analysis for the Global Burden of Disease Study 2013. The Lancet 2014; 384: 766–81.
- 3.) Angelantonio ED, Bhupathiraju SN, Wormser D, et al. Body-mass index and all-cause mortality: individual-participant-data meta-analysis of 239 prospective studies in four continents. The Lancet 2016; 388: 776–86.

Bone Mineral Density Capstone Appendix



Flowchart

Input data & methodological summary

Exposure

Case definition

Bone mineral density (BMD) is a continuous variable measured by dual-x-ray-absorptiometry (DXA) at the femoral neck (FN) and is presented in g/cm² after standardizing for the brand of densitometer (sBMD). Low BMD is measured in terms of the difference between BMD of a population and the 99th percentile of a reference population at the same age and sex (theoretical minimum of risk exposure level, TMREL). The burden attributed to low bone mineral density is estimated for adults 20 years and older.

Input data

A systematic review (search string at the end of document) was conducted in GBD 2015 but it was not scheduled for systematic review in GBD 2016 or 2017. Inclusion criteria that informed the search are:

- o Representative, population-based surveys
- o Reporting of quantitative BMD
 - measured by DXA
 - performed at the FN region
 - measured in g/cm²

Mean BMD was occasionally reported in stratified groups, e.g. by fracture status but not for total sample. In these cases, the stratified means were aggregated to obtain a total mean BMD at population level for an age or sex category.

The data availability by GBD super-region is shown in table below.

Super region	The number of data points
Southeast Asia, East Asia, and Oceania	314
Central Europe, Eastern Europe, and Central Asia	36
High-income	682
Latin America and Caribbean	97
North Africa and Middle East	110
South Asia	39
Sub-Saharan Africa	3

Modelling strategy

We modelled mean BMD in DisMod-MR 2.1 as a single 'continuous' parameter model by age and sex, and all GBD locations for years 1990 to 2017. The model had age mesh points at 0 10 20 25 30 40 50 60 70 80 90 & 100, a time window of 10 years for fitting data, and a minimum coefficient of variation of 0.1 for global, 0.06 super region and 0.08 for the region level.

The country covariates of alcohol consumption (litres per capita), tobacco consumption (cigarettes per capita), mean BMI, and adjusted calcium intake (g) were included in modelling. The country covariates total physical activity and milk consumption did not have a significant effect on BMD so we excluded them from our final model.

The uncertainty of BMD was modelled using various distributions. We tested goodness-of-fit in NHANES III data, the only survey for which we had unit record data available. We applied a weighting ensemble on those distributions. The weights were calculated in an optimisation model with an objective function that minimised Kolmogorov-Smirnov statistics. The weights of the distributions in the ensemble were calculated separately for males and females. Distribution weights are shown below.

Figure 1: Distribution of weights used for Females in GBD 2017



Figure 2: Distribution of weights used for Males in GBD 2017



We consider the risk of fatal and non-fatal outcomes for hip non-hip fractures, separately, as relative risk data provide different estimates. Thus, there were various steps after DisMod-MR 2.1 exposure modelling to arrive at attributable fractions that can be applied to fatal and non-fatal fracture outcomes. These osteoporotic non-hip fractures include fractures of vertebrae, clavicle, scapula, humerus, skull, sternum, rib, face bone, radius or ulna, femur, patella, tibia, fibula, ankle, pelvis and vertebrae.

First, we calculated the proportion of injury deaths that are due to fractures. This proportion of death caused by fracture is the envelope that we use to attribute death to BMD. In order to do this, we assumed that hip fracture and some non-hip fractures (any fractures apart from fingers and toes) are potentially fatal fractures. As cause of death data from vital registration and verbal autopsy attributes injury deaths to causes of death (e.g. fall or road injury) and not nature of injury (such as fractures), we used available hospital data to estimate the proportion of injury deaths during admission that could be ascribed to fractures. We restricted our analysis to cases that were dual-coded with both the cause of injury ("E-code") and nature of injury ("N-code"). As injury cases may have multiple forms of trauma, we applied a severity hierarchy to the fatal hospital data to determine the proportion of the deaths that could be attributed to the chosen fracture types but were not accompanied by more severe fatal trauma such as head trauma, spinal cord lesion, and intra-abdominal or thoracic organ damage. We collapsed all deaths over E-code to determine the ratio of deaths attributable to fracture versus non-fracture injuries. We applied this ratio to the YLL.

We restricted non-fatal estimates of low BMD to a list of causes that were deemed to cause osteoporotic fractures. Below is the list of injuries for which a PAF was calculated:

- Transport injuries
- Road injuries
- Pedestrian road injuries
- Cyclist road injuries
- Motorcyclist road injuries
- Motor vehicle road injuries
- Other road injuries
- Other transport injuries
- Unintentional injuries
- Falls
- Exposure to mechanical forces
- Other exposure to mechanical forces
- Non-venomous animal contact
- Interpersonal violence
- Assault by other means

We made use of the E to N-code matrix generated from dual-coded (E-code/N-code) patient level data in our injury analyses to determine the proportion of each E-code that results in a certain N-code. The hip and non-hip fracture population attributable fractions were applied to the appropriate combinations of external cause and fracture estimates of YLD and then summed together to produce a single estimate.

Figure 3: Plot of hospital fractions used by hip and non-hip



Theoretical minimum-risk exposure level

The theoretical minimum of risk exposure level or TMREL was chosen as the age-sex specific 99th percentile of BMD from 5 cycles of NHANES study as the reference population. Below is a descriptive table of the 5 NHANES cycles used.

NHANES cycle	Age range (years)	Number of people tested	BMD range (g/cm2)
1988	20 – 90	14,646	0.23 – 1.84
2005	20 – 85	3,494	0.40 – 1.50
2007	20 – 80	4,726	0.34 – 1.46
2009	20 – 80	5,052	0.33 – 1.63
2013	40 - 80	3,127	0.39 – 1.36



Figure 4: Plot of 99th percentile of BMD at femoral neck in each cycle of NHANES

Relative risks

Relative risks must be reported per standard deviation or per unit bone mass density in order for us to use the data. Many studies report relative risk based on a z-score or the relative risks in the osteoporotic group versus the non-osteoporotic group; neither of these relative risks are usable.

For GBD 2017, we did not update the systematic review for the RR of BMD that was done in GBD 2013. In the GBD 2013 review, twelve prospective observational studies were found, but one meta-analysis of 12 studies¹ reported the dose-response relationship between low BMD and high relative risk of hip and other fractures that are prone to osteoporosis, as shown in the below table.

BMD	Any	v fracture	Oste fi	eoporotic racture	Hip	fracture
z score	RR	95% CI	RR	95% CI	RR	95% CI
-4	1.79	1.44-2.23	2.10	1.63-2.71	2.14	1.40-3.26
-3	1.71	1.44-2.02	1.96	1.61-2.39	2.12	1.54-2.92
-2	1.63	1.45-1.84	1.84	1.60-2.12	2.11	1.70-2.62
-1	1.56	1.45-1.69	1.73	1.59-1.89	2.11	1.86-2.39
0	1.50	1.44-1.56	1.62	1.54-1.71	2.08	1.91-2.26
1	1.39	1.32-1.46	1.42	1.34-1.51	2.04	1.78-2.34
2	1.32	1.21-1.45	1.33	1.19-1.48	2.03	1.60-2.56
3	1.26	1.10 - 1.45	1.25	1.06 - 1.47	2.01	1.44-2.81
4	1.21	1.00 - 1.45	1.17	0.93-1.46	1.99	1.28-3.10

The z score ranged from -5.1 to +5.8.

Search string from GBD 2015 systematic review:

Search	Query	ltems found	Time
#11	Search (#8 AND #10) Filters: Humans	326	12:37:09
#10	Search ("Cross-Sectional Studies"[Mesh] OR "cross-sectional"[title/abstract] OR "Health Surveys"[Mesh] OR Survey[title/abstract] OR cohort[title/abstract] OR "Diet Surveys"[Mesh] OR "Longitudinal Studies"[Mesh] OR "Nutrition Surveys"[Mesh] OR "Surveys and Questionnaires"[Mesh]) Filters: Humans	1324376	12:36:29
#8	Search (#7 AND #6) Filters: Humans	622	12:33:16
#7	Search ("Absorptiometry, Photon"[Mesh] OR "dual-energy x-ray absorptiometry" OR "dual energy x-ray absorptiometry") Filters: Humans	21368	12:32:34
#6	Search (#5 AND ("2010"[Date - Publication] : "3000"[Date - Publication])) Filters: Humans	1387	12:30:26
#5	Search ((#1 OR #2) AND #3) Filters: Humans	3702	12:29:47
#4	Search ((#1 OR #2) AND #3)	4015	12:29:33
#3	Search (((("bone mineral density"[title/abstract] OR "bone mineral densities"[title/abstract]) OR "Bone Density"[Mesh]) AND (mean[title/abstract] OR average[title/abstract])))	12892	12:29:00
#2	Search ((((Initiational TAB) OK International TAB) OK Inational TAB) OK Ination (TAB) OK equitorial (TIAB) OR equitorial (TIAB) OR global [TIAB) OR global [TIAB] OR global [TIAB] OR continental [TIAB] OR continents [TIAB] OR global burden [TIAB] OR burden of disease [TIAB] OR topics (TIAB) OR tropics (TIAB) OR tropical (TIAB) OR Ccennia (TIAB) OR South America [TIAB) OR continents [TIAB] OR pan-america (TIAB) OR Ccentral America [TIAB] OR Mesoamerica [TIAB] OR Americas [TIAB] OR Latin America [TIAB] OR pan-american (TIAB) OR Australasian (TIAB) OR developing countries [TIAB] OR developing nations [TIAB] OR developed countries [TIAB] OR developed nations [TIAB] OR commonwealth [TIAB] OR industrialized [TIAB] OR non-industrialized [TIAB] OR underdeveloped nations [TIAB] OR under-developed country [TIAB] OR under-developed countries [TIAB] OR under-developed country [TIAB] OR under-developed country [TIAB] OR under-developed country [TIAB] OR under-developed nations [TIAB] OR non-developed nations [TIAB] OR non-developed country [TIAB] OR non-developed country [TIAB] OR low-income country [TIAB] OR non-developed nations [TIAB] OR non-developed nations [TIAB] OR non-developed country [TIAB] OR non-developed nations [TIAB] OR non-developed country [TIAB] OR Northern Hemisphere [TIAB	3585538	12:28:22

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Impaired Kidney Function Capstone Appendix

Flowchart



Impaired Kidney Function

1. The Chronic Kidney Disease Prognosis Consortium collects population-level cohort data on CKD from around the world for the purpose of collective meta-analyses

Input Data & Methodological Summary

Exposure

Case Definition

The impaired kidney function risk factor exposure is divided into four categories of renal function defined by urinary albumin to creatinine ratio (ACR) and estimated glomerular filtration rate (GFR): albuminuria with preserved GFR (ACR >30 mg/g & GFR >=60 ml/min/1.73m²), chronic kidney disease (CKD) stage 3 (GFR of 30-59 ml/min/1.73m²), CKD stage 4 (GFR of 15-29 ml/min/1.73m²), and CKD stage 5 (GFR <15ml/min/1.73m², not yet on renal replacement therapy). The modelling of renal function prevalence estimates is described in detail in the appendix to the GBD 2017 non-fatal capstone paper as these are also disease sequelae.

Input data

For GBD 2010, a systematic review of the prevalence of low glomerular filtration rate throughout the world was conducted. This search was updated for GBD 2013, GBD 2015, and 2016. Exclusion criteria included surveys that were not population-representative, studies not reporting on CKD by stage, and studies not reporting on albuminuria with preserved GFR (GFR >=60 ml/min/1.73m²).

Disease	Number of sources	Number of countries	Number of new sources
			for GBD 2016
Albuminuria	72	31	72
CKD Stage III	112	47	49
CKD Stage IV	94	40	45
CKD Stage V	92	38	49

Modeling strategy

Estimates of exposure to albuminuria and CKD were obtained from the GBD 2017 non-fatal burden of disease analysis, which includes stage-specific prevalence estimates at the country level across twenty-three age-groups for both genders. The modeling strategy for these estimates is detailed in the appendix to the GBD 2017 non-fatal capstone paper.

Relative risks were calculated by the Chronic Kidney Disease Prognosis Consortium, a consortium composed of population-level cohorts with prospective data collection from several countries (details below). YLDs and YLLs for cardiovascular diseases and gout were obtained from the GBD 2017 Study for the same geographic, time-period, and age-groups as detailed above.

Theoretical minimum-risk exposure level

The theoretical minimum risk is a diagnosis of albuminuria or CKD stages 3, 4, or 5. An ACR above 30 mg/g and eGFR below 60ml/min/1.73m² have been demonstrated in the literature to be the thresholds at which increased cardiovascular and gout events occur secondary to impaired kidney function. (1-10)

Relative risk

A two-stage pooled meta-analysis was used to calculate relative risks for ischemic heart disease, stroke, and peripheral vascular disease. The relative risk of these conditions was first determined within each cohort, and then a pooled analysis of cohort-level relative risks was performed using a random effects meta-analysis approach. Uncertainty intervals largely overlapped for the relative risks of fatal and nonfatal cardiovascular events from impaired kidney function exposure. Thus, we decided to use the relative risks from the combined analysis for fatal and nonfatal cardiovascular outcomes. Gout relative risk was determined by meta-analysis of a literature review performed for GBD 2013. Search terms included "gout" and "chronic kidney disease". Exclusion criteria for search results included special populations, reversal of exposure and outcome categories, or unclear exposure category definition. This search resulted in four eligible studies; no new studies indicated an increased risk of gout with albuminuria.

Population Attributable Fraction

We calculated the cardiovascular and gout fatal and nonfatal burden attributable to the categorical exposure to impaired kidney function using the following equation:

$$PAF = \frac{\sum_{i=1}^{n} P_i (RR_i - 1)}{\sum_{i=1}^{n} P_i (RR_i - 1) + 1}$$

Equation 1. PAF based on categorical exposure

where RR_i is the relative risk for exposure level *i*, P_i is the proportion of the population in that exposure category, and *n* is the number of exposure categories.(11)

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Section 5: Tables and Figures

Appendix Figure 1. Analytical flowchart of the comparative risk assessment for the estimation of population attributable fractions by geography, age, sex, and year for GBD 2017. Ovals represent data inputs, rectangular boxes represent analytical steps, cylinders represent databases, and parallelograms represent intermediate and final results. GBD=Global Burden of Disease. SEVs=Summary exposure values. TMREL=Theoretical minimum-risk exposure level. PAFs=Population attributable fractions. YLLs=years of life lost. YLDs=years lived with disability. DALYs=disability-adjusted life-years.







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Appendix Figure 3: Spatiotemporal Gaussian Process Regression


Appendix Table 1. GATHER checklist of information that should be included in reports of global health estimates, with description of compliance and location of information for GBD 2017.

#	GATHER checklist item	Description of compliance	Reference
Obj	ectives and funding	compliance	
1	Define the indicators, populations, and time periods for which estimates were made.	Narrative provided in paper and methods appendix describing indicators, definitions, and populations	Main text (Methods— Overview, Geographic units and time periods) and methods appendix
2	List the funding sources for the work.	Funding sources listed in paper	Summary (Funding)
Data	a Inputs		
For	all data inputs from multiple sources that are synthesized as part	t of the study:	1
3	Describe how the data were identified and how the data were accessed.	Narrative description of data seeking methods provided	Main text (Methods) and methods appendix
4	Specify the inclusion and exclusion criteria. Identify all ad-hoc exclusions.	Narrative about inclusion and exclusion criteria by data type provided	Main text (Methods) and methods appendix
5	Provide information on all included data sources and their main characteristics. For each data source used, report reference information or contact name/institution, population represented, data collection method, year(s) of data collection, sex and age range, diagnostic criteria or measurement method, and sample size, as relevant.	An interactive, online data source tool that provides metadata for data sources by component, geography, cause, risk, or impairment has been developed	Online data citation tools
6	Identify and describe any categories of input data that have potentially important biases (e.g., based on characteristics listed in item 5).	Summary of known biases by cause included in methods appendix	Methods appendix
For	data inputs that contribute to the analysis but were not synthesiz	zed as part of the study:	
7	Describe and give sources for any other data inputs.	Included in online data source tool, <u>http://ghdx.healthdata.</u> <u>org/gbd-2017</u>	Online data citation tools
For	all data inputs:	Γ	I
8	Provide all data inputs in a file format from which data can be efficiently extracted (e.g., a spreadsheet as opposed to a PDF), including all relevant meta-data listed in item 5. For any data inputs that cannot be shared due to ethical or legal reasons, such as third-party ownership, provide a contact name or the name of the institution that retains the right to the data.	Downloads of input data available through online tools, including data visualization tools and data query tools, <u>http://ghdx.healthdata.</u>	Online data visualization tools, data query tools, and the Global Health Data Exchange, <u>http://ghdx.healthdata.</u> org

		org/gbd-2017; input	
		data not	
		available in tools will	
		be made available	
		upon request	
Data	a analysis		
9	Provide a conceptual overview of the data analysis method. A	Flow diagrams of the	Main text (Methods)
	diagram may be helpful.	overall methodological	and methods appendix
		processes, as well as	
		cause-specific	
		modelling processes,	
		have been provided	
10	Provide a detailed description of all steps of the analysis,	Flow diagrams and	Main text (Methods)
	including mathematical formulae. This description should	corresponding	and methods appendix
	cover, as relevant, data cleaning, data pre-processing, data	methodological write-	
	aujustiments and weighting of data sources, and	as well as the	
	mathematical of statistical model(s).	demographics and	
		causes of death	
		databases and	
		modelling processes	
		have been provided	
11	Describe how candidate models were evaluated and how the	Provided in the	Methods appendix
	final model(s) were selected.	methodological write-	
		ups	
12	Provide the results of an evaluation of model performance, if	Provided in the	Methods appendix
	done, as well as the results of any relevant sensitivity	methodological write-	
	analysis.	ups	
13	Describe methods for calculating uncertainty of the	Provided in the	Methods appendix
	estimates. State which sources of uncertainty were, and were	methodological write-	
	not, accounted for in the uncertainty analysis.	ups	
14	State how analytic or statistical source code used to generate	Access statement	Code is provided in an
	estimates can be accessed.	provided	online repository
Res	ults and Discussion	Deculto and available	Main tout matter de
12	Provide published estimates in a file format from which data	through	iviain text, methods
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		(http://ghdx.healthdata	http://ghdx.healthdata.
		.org/gbd-2017)	org/gbd-2017)
16	Report a quantitative measure of the uncertainty of the	Uncertainty intervals	Main text, methods
	estimates (e.g. uncertainty intervals).	are provided with all	appendix, and online
		results	data tools (data
			visualization tools, data
			query tools, and the

			Global Health Data
			Exchange,
			http://ghdx.healthdata.
			org/gbd-2017)
17	Interpret results in light of existing evidence. If updating a	Discussion of	Main text (Methods
	previous set of estimates, describe the reasons for changes in	methodological	and Discussion) and
	estimates.	changes between GBD	methods appendix
		rounds provided in the	
		narrative of the Article	
		and methods appendix	
18	Discuss limitations of the estimates. Include a discussion of	Discussion of	Main text (Limitations)
	any modelling assumptions or data limitations that affect	limitations provided in	and methods appendix
	interpretation of the estimates.	the narrative of the	
		main paper, as well as	
		in the methodological	
		write-ups in the	
		methods appendix	

Appendix Table 2. GBD 2017 risk factor hierarchy with levels GBD=Global Burden of Disease.	s, modeli	ing strategies, and the main type of data sources used to e	estimate exposure levels				
Risk factor	Leve] Model type	Main data source for exposure				
All risk factors	0		•				
Environmental/occupational risks	1						
Unsafe water, sanitation, and handwashing	2						
Unsafe water source	3	Spatiotemporal Gauissian process regression (ST-GPR)	Population surveys and censuses				
Unsafe sanitation	3	ST-GPR	Population surveys and censuses				
No handwashing with soap	3	ST-GPR	Population surveys, censuses, and epidemiological studies				
Air pollution	2						
Particulate matter pollution	3						
Ambient particulate matter pollution	4	Regression crosswalk between grid-level fusion of satellite/chemical transport models and ground level monitoring data	Atmospheric chemical transport models, satellite measurements of aerosols in the atmosphere, data from ground-level monitoring sites				
Household air pollution from solid fuels	4	ST-GPR	Population surveys and censuses				
Ambient ozone pollution	3	Chemical transport model	Atmospheric chemical transport models				
Other environmental risks	2						
Residential radon	3	ST-GPR	Literature review				
Lead exposure	3	ST-GPR	Literature review				
Occupational risks	2						
Occupational carcinogens	3	Ashering Turner Deriver and the	CDD				
Occupational exposure to asbestos	4	Asbestos impact Ratio approach	GBD cause-specific mortainty data for mesothelioma, epidemiological studies				
Occupational exposure to arsenic	4	ST-GPR	Labor force surveys, censuses, and international information system on occupational exposure to carcinogens				
Occupational exposure to benzene	4	ST-GPR	Labor force surveys, censuses, and international information system on occupational exposure to carcinogens				
Occupational exposure to beryllium	4	ST-GPR	Labor force surveys, censuses, and international information system on occupational exposure to carcinogens				
Occupational exposure to cadmium	4	ST-GPR	Labor force surveys, censuses, and international information system on occupational exposure to carcinogens				
Occupational exposure to chromium	4	ST-GPR	Labor force surveys, censuses, and international information system on occupational exposure to carcinogens				
Occupational exposure to diesel engine exhaust	4	ST-GPR	Labor force surveys, censuses, and international information system on occupational exposure to carcinogens				
Occupational exposure to formaldehyde	4	ST-GPR	Labor force surveys, censuses, and international information system on occupational exposure to carcinogens				
Occupational exposure to nickel	4	ST-GPR	Labor force surveys, censuses, and international information system on occupational exposure to carcinogenes				
Occupational exposure to polycyclic aromatic	4	ST-GPR	Labor force surveys, censuses, and international information system on occupational exposure to carcinogene				
Occupational exposure to silica	4	ST-GPR	Labor force surveys, censues, and international information system on				
Occupational exposure to sulfphuric acid	4	ST-GPR	Labor force surveys, censuses, and international information system on				
Occupational exposure to trichloroethylene	4	ST-GPR	Labor force surveys, censuses, and international information system on				
Occupational acthmagans	2	ST CDP	occupational exposure to carcinogens				
Occupational particulate matter, gases, and fumes	3	ST-GPR	Labor force surveys and censuses				
Occupational noise	3	ST-GPR	Labor force surveys and censuses, industry-based surveys of noise				
Occupational injuries	2	ST_GPB	International Labor Organization injury database				
Occupational ergonomic factors	3	ST-GPR	Labor force surveys and censuses				
Behavioural risks	1	STOR	Eabor force surveys and consuses				
Child and maternal malnutrition	2						
Suboptimal breastfeeding	3						
Non-exclusive breastfeeding	4	ST-GPR	Population surveys				
Discontinued breastfeeding	4	ST-GPR	Population surveys				
Childhood undernutrition	3						
Childhood underweight	4	ST-GPR	Examination surveys and epidemiological studies				
Childhood wasting	4	ST-GPR	Examination surveys and epidemiological studies				
Childhood stunting	4	ST-GPR	Examination surveys and epidemiological studies				
Iron deficiency	3	Mixed effect regression	Examination surveys and epidemiological studies				
Vitamin A deticiency Zinc deficiency	3	DisMod-MR 2.1 Mixed effect regression based on stunting prevalence and	Examination surveys and epidemiological studies FAO food balance sheets				
Telesco melo	•	dietary composition					
10Dacco smoke	2	• Smaking Impact Patio (SIP) salaylated from h	SID input data: mortality and agues of death data including vit-1				
JHOKIIY	3	• Smoking prevalence estimated using ST-GPR	registration and verbal autopsy Smoking prevalence input data: nationally representative survey and report data				
Second-hand smoke	3	DisMod-MR 2.1	Household surveys and national health surveys				

Risk factor	Leve	1 Model type	Main data source for exposure
Alcohol use	2	Alcohol consumption per capita obtained from the FAO and the WHO Global Information System on Alcohol and Health (GISAH) ST-GPR used to integrate the data and to derive coherent time series for each country Prevalence of current alcohol drinkers, lifetime abstainers, former drinkers, and binge drinkers estimated using DisMod-MR 2.1 DisMod-MR 2.1 used to estimate the relative sex- and age-specific pattern of alcohol consumption in current drinkers	Population surveys, alcohol sales, production, and other economic statistics
Drug Use	2	DisMod-MR 2.1	Systematic review of published literature, reports from governments and international organizations, which include data from: school surveys, population surveys, registration data, and indirect estimates of prevalence
Dietary risks	2		
Diet low in fruits	3	DisMod-MR 2.1	Nutrition and health surveys, FAO food balance sheets
Diet low in vegetables	3	DisMod-MR 2.1	Nutrition and health surveys, FAO food balance sheets
Diet low in whole grains	3	DisMod-MR 2.1	Nutrition and health surveys
Diet low in nuts and seeds	3	DisMod-MR 2.1	Nutrition and health surveys, FAO food balance sheets
Diet low in milk	3	DisMod-MR 2.1	Nutrition and health surveys, FAO food balance sheets
Diet high in red meat	3	DisMod-MR 2.1	Nutrition and health surveys, FAO food balance sheets
Diet high in processed meat	3	DisMod-MR 2.1	Nutrition and health surveys
Diet high in sugar-sweetened beverages	3	DisMod-MR 2.1	Nutrition and health surveys
Diet low in fibre	3	DisMod-MR 2.1	Nutrition and health surveys, FAO SUA/USDA
Diet suboptimal in calcium	3	DisMod-MR 2.1	Nutrition and health surveys, FAO SUA/USDA
Diet low in seafood omega-3 fatty acids	3	DisMod-MR 2.1	Nutrition and health surveys, FAO SUA/USDA
Diet low in polyunsaturated fatty acids	3	DisMod-MR 2.1	Nutrition and health surveys, FAO SUA/USDA
Diet high in trans fatty acids	3	DisMod-MR 2.1	Nutrition and health surveys
Diet high in sodium	3	DisMod-MR 2.1	Nutrition and health surveys
Intimate partner violence	2	DisMod-MR 2.1	Systematic review of published literature, national health surveys, violence-specific surveys
Childhood maltreatment	2		
Childhood sexual abuse	3	DisMod-MR 2.1	Systematic review of published literature, national health surveys, violence-specific surveys
Bullying victimization	3	DisMod-MR 2.1	Systematic review of published literature and surveys reporting bullying victimization, including Global School-based Student Health Survey (GSHS), the Health Behavior in School-aged Children (HBSC), and the National Crime Victimization Survey – School Crime Supplement (NCVS-SCS).
Unsafe sex	2	DisMod-MR 2.1	UNAIDS country progress reports, disease surveillance reports
Low physical activity	2	DisMod-MR 2.1	Surveys of the adult population that capture reported frequency, duration and intensity of physical activity undertaken in the past seven days across all domains of life (work, transport, recreation or house/yard work)
Metabolic risks	1		
High fasting plasma glucose	2	ST-GPR	Examination surveys and epidemiological studies
High LDL cholesterol	2	ST-GPR	Examination surveys and epidemiological studies
High systolic blood pressure	2	ST-GPR	Examination surveys and epidemiological studies
High body-mass index	2	ST-GPR	Examination surveys and epidemiological studies
Low bone mineral density	2	DisMod-MR 2.1	Examination surveys and epidemiological studies
Low glomerular filtration rate	2	DisMod-MR 2.1	Examination surveys and epidemiological studies

Appendix Table 3. Types of Comparative Risk Assessments (CRA) based on the time perspective and the nature of the counterfactual level or distribution of exposure. The shaded box represents the type of CRA currently undertaken in GBD 2017. GBD=Global Burden of Disease.

		Counterfactual distr	ibutions of exposure	
	Theoretical minimum	Plausible minimum	Feasible minimum	Cost-effective
	risk: level of risk with	risk: level of risk with	risk: level of risk with	minimum risk: lowest
	the lowest level of	the lowest level of	the lowest level of	level of risk that can
	burden	burden that could be	burden that has been	be achieved cost-
		imagined with current	achieved in any	effectively in a given
		technology and	population	population
Construct		knowledge		
Attributable burden: burden of				
disease today that would be avoided				
if each individual in the past had	Currently in GBD			
been exposed to the counterfactual				
level of exposure				
Avoidable burden: burden of disease				
in the future that would be avoided				
if each individual today was shifted				
to the counterfactual level of				
exposure				

ix rable 4. Descriptive cataloguing of the	ne epidemiological evidence used to as	sess wheth	ter each ris	sk-outcol	ne paper	meets the	causal c	riteria for i	nclusion	In the Gi	obal Burd	en of D
			he		s	se	n)**					
			E.	•	ndie	in th	ip (8 a				
			fect	%) (%	l stı	l stu on i	ssin nsh	the		.e		
			it ef	sgui	ona	ona ciati	asse atio	n in	1.5	nsh	×+	
			can (%	ibu	/ati	/atio	rels s	(% tion	^	ation	jit	
			tion fi	ll fi	serv	serv it a: tion	air	ocis	RR	rels	usit	
			sig	nu	e ob	e ob Ican	ol si ne p	ol si ass irect	it of	nse	pla	
		Ē	vith te di	vith	Stive	stive gnif fe di	teor	ant ant fe di	in i	spo	cal	y #
		Ts (Ts y	Ls v	spe	spe i sij	-ou	e-co nific	ver	e-re	logi	alog
Risk	Outcome	RC	epp CC	RC	(II)*	Pro: opp	Cas risk	Cas sign opp	Lov	Dos	Bio	Ans
Unsafe water, sanitation, and												
handwashing												
Unsafe water source - chlorination or solar	Diarrhoeal diseases	25	0	44	6	0	-	-	Yes	-	Yes	No
(point of use treatment) Unsafe water source - piped	Diarrhoeal diseases	1	0	0	9	11	-	_	Ves	_	Ves	No
Unsafe water source - filter	Diarrhoeal diseases	11	0	45	2	0	-	-	Yes	_	Yes	No
Unsafe water source - improved water	Diarrhoeal diseases	0	-	-	5	Ő	-	-	Yes	-	Yes	No
Unsafe sanitation - piped	Diarrhoeal diseases	0	-	-	7	0	-	-	Yes	-	Yes	No
Unsafe sanitation - improved sanitation	Diarrhoeal diseases	1	-	100	11	0	-	-	Yes	-	Yes	No
No access to handwashing facility	Diarrhoeal diseases	19	0	42	0	-	-	-	No	-	Yes	No
No access to handwashing facility	Lower respiratory infections	8	0	50	11	0	-	-	No	-	Yes	No
Air pollution												
Particulate matter pollution		0				0				••		
Ambient particulate matter pollution	Lower respiratory infections	0	-	-	17	0	-	-	No	Yes	Yes	No
Ambient particulate matter pollution	I racheal, bronchus, and lung cancer	0	-	-	30	0	-	-	No	Yes	Yes	Yes
Ambient particulate matter pollution	Ischaemic neart disease	0	-	-	10	0	-	-	No	Yes	Yes	Yes
Ambient particulate matter pollution	Intracerebral hemorrhage	0	-	-	30	0	-		No	Ves	Ves	Ves
Ambient particulate matter pollution	Subarachnoid hemorrhage	0	_	-	30	0	-	-	No	Yes	Yes	Yes
Ambient particulate matter pollution	Chronic obstructive pulmonary disease	0	-	-	12	Ő	-	-	No	Yes	Yes	Yes
Ambient particulate matter pollution	Diabetes mellitus type 2	0	-	-	8	0	-	-			Yes	Yes
Household air pollution from solid fuels	Lower respiratory infections	2	0	1	8	0	17	0	No	Yes	Yes	No
Household air pollution from solid fuels	Tracheal, bronchus, and lung cancer	0	-	-	0	-	28	0	No	Yes	Yes	Yes
Household air pollution from solid fuels	Ischaemic heart disease	0	-	-	2	-	2	-	No	Yes	Yes	Yes
Household air pollution from solid fuels	Ischaemic stroke	0	-	-	1	-	-	-	No	Yes	Yes	Yes
Household air pollution from solid fuels	Intracerebral hemorrhage	0	-	-	1	-	-	-	No	Yes	Yes	Yes
Household air pollution from solid fuels	Subarachnoid hemorrhage	0	-	-	1	-	-	-	No	Yes	Yes	Yes
Household air pollution from solid fuels	Chronic obstructive pulmonary disease	0	-	-	0	-	11	0	No	Yes	Yes	Yes
Household air pollution from solid fuels	Diabetes mellitus type 2	0	-	-	1	-	-	-			Yes	Yes
Household air pollution from solid fuels	Cataract	0	-	-	0	-	8	0	No	Yes	Yes	No
Ambient ozone pollution	Chronic obstructive pulmonary disease	0	-	-	3	0	-	-	No	Yes	Yes	No
Other environmental risks												
Residential radon	Tracheal, bronchus, and lung cancer	0	-	-	1	0	29	0	No	Yes	Yes	No
Lead exposure	Idiopathic developmental intellectual	0	-	-	8	0	-	-	No	Yes	Yes	No
Land avposure	disability Systelia blood pressure	0			2	0	1	0	No	Vac	Vac	No
Occupational risks	Systeme blood pressure	0	-	-	5	0	1	0	INU	105	105	INU
Occupational exposure to ashestos	Larvnx cancer	0	-	-	27	0	-		No	-	Yes	Yes
Occupational exposure to asbestos	Tracheal, bronchus, and lung cancer	0	-	-	18	0	-	-	Yes	-	Yes	Yes
Occupational exposure to asbestos	Ovarian cancer	0	-	-	15	0	-	-	No	-	Yes	Yes
Occupational exposure to asbestos	Mesothelioma	0	-	-	5	0	-	-	Yes	-	Yes	Yes
Occupational exposure to arsenic	Tracheal, bronchus, and lung cancer	0	-	-	9	0	-	-	No	-	Yes	No
Occupational exposure to benzene	Leukaemia	0	-	-	12	0	-	-	Yes	-	Yes	No
Occupational exposure to beryllium	Tracheal, bronchus, and lung cancer	0	-	-	3	0	2	0	No	-	Yes	No
Occupational exposure to cadmium	Tracheal, bronchus, and lung cancer	0	-	-	7	0	-	-	No	-	Yes	No
Occupational exposure to chromium	Tracheal, bronchus, and lung cancer	0	-	-	26	0	-	-	No	-	Yes	No
Occupational exposure to diesel engine	Tracheal, bronchus, and lung cancer	0	-	-	17	0	-	-	No	-	Yes	No
exhaust Occupational exposure to formaldehyde	Nasopharynx cancer	0	_	-	2	0	6	0	No	_	Yes	Yes
Occupational exposure to formaldehyde	Leukaemia	0	-	-	13	0	-	-	No	-	Yes	Yes
Occupational exposure to nickel	Tracheal, bronchus, and lung cancer	0	-	-	6	0	-	-	No	-	Yes	No
Occupational exposure to polycyclic aromatic	Tracheal, bronchus, and lung cancer	0	-	-	39	0	-	-	No	-	Yes	No
Occupational exposure to silica	Tracheal, bronchus, and lung cancer	0	-	-	17	0	-	-	No	-	Yes	No
Occupational exposure to sulfphuric acid	Larynx cancer	0	-	-	14	0	-	-	Yes	-	Yes	No
Occupational exposure to trichloroethylene	Kidney cancer	0	-	-	20	0	-	-	No	-	Yes	No
Occupational asthmagens	Asthma	0	-	-	16	0	-	-	No	-	Yes	No
Occupational particulate matter, gases, and	Chronic obstructive pulmonary disease	0	-	-	9	0	-	-	No	-	Yes	No
Occupational noise	Age-related and other hearing loss	0	-	-	5	0	-	-	Yes	-	Yes	No
Child and maternal malnutritian	Low back pain	0	-	-	10	0	-	-	NO	-	Y es	NO
Non-evolutive broastfooding	Diarrhoeal diseases	0			Ę	0	7	0	Vac		Vac	No
Non-exclusive breastfeeding	Lower respiratory infections	0	-	-	5 6	0	/ R	0	I US Vac	-	I CS Vac	INO No
Discontinued breastfeeding	Diarrhoeal diseases	0	-	-	2	0	-	-	No	-	Yes	No
Child underweight	Diarrhoeal diseases	0	-	-	7	0	-		Yes	-	Yes	No
Child underweight	Lower respiratory infections	0	-	-	7	0	-	-	Yes	-	Yes	No
Child underweight	Measles	0	-	-	7	0	-	-	Yes	-	Yes	No
Child wasting	Diarrhoeal diseases	0	-	-	7	0	-	-	Yes	-	Yes	No
Child wasting	Lower respiratory infections	0	-	-	7	0	-	-	Yes	-	Yes	No
Child wasting	Measles	0	-	-	7	0	-	-	Yes	-	Yes	No
Child stunting	Diarrhoeal diseases	0	-	-	7	0	-	-	No	-	Yes	No
Child stunting	Lower respiratory infections	0	-	-	7	0	-	-	No	-	Yes	No
Child stunting	Measles	0	-	-	7	0	-	-	No	-	Yes	No

wel			(1	ith significant effect in the e direction (%)	ith null findings (%)	tive observational studies	tive observational studies nificant association in the e direction (%)	ntrol studies assessing the come pair relationship (n)**	ntrol studies that show int association in the e direction (%)	imit of RR > 1.5	ponse relationship	al plausibility +	*	
isk Le	D. 1		CTs (I	CTs w pposit	CTs w	rospec	rospec ith sig pposit	ase-col sk-out	ase-col gnifica ppositi	ower l	0se-re	iologic	nalogy	
2	Risk Short gestation for birth weight	Diarrhoeal diseases	2 0	- R	ž	20	0 b	Ö.≊	0 10	- Yes	Yes	Yes	₹ Yes	
4	Short gestation for birth weight	Lower respiratory infections	0	-	-	20	0	-	-	Yes	Yes	Yes	Yes	
4	Short gestation for birth weight	Upper respiratory infections	0	-	-	20	0		-	Yes	Yes	Yes	Yes	
4	Short gestation for birth weight	Otitis media Preumococcel meningitis	0	-	-	20	0	-	-	Yes	Y es Ves	Yes	Yes	
4	Short gestation for birth weight	H influenzae type B meningitis	0		-	20	0	-		Yes	Yes	Yes	Yes	
4	Short gestation for birth weight	Meningococcal infection	0	-	-	20	0	-	-	Yes	Yes	Yes	Yes	
4	Short gestation for birth weight	Other meningitis	0	-	-	20	0	-	-	Yes	Yes	Yes	Yes	
4	Short gestation for birth weight	Encephalitis	0	-	-	20	0	-	-	Yes	Yes	Yes	Yes	
4	Short gestation for birth weight	Neonatal preterm birth complications	0	-	-	20	0	-	-	Yes	Yes	Yes	Yes	
4	Short gestation for birth weight	Neonatal sensis and other neonatal	0	-	-	20	0	-	-	Yes	Yes	Yes	Yes	
4	Short gestation for birth weight	Hemolytic disease and other neonatal	0	-	-	20	0	-	-	Yes	Yes	Yes	Yes	
4	Short gestation for birth weight	Other neonatal disorders	0	-	-	20	0	-	-	Yes	Yes	Yes	Yes	
4	Short gestation for birth weight	Sudden infant death syndrome	0	-	-	20	0	-	-	Yes	Yes	Yes	Yes	
4	Low birth weight for gestation	Diarrhoeal diseases	0	-	-	20	0	-	-	Yes	Yes	Yes	Yes	
4	Low birth weight for gestation	Upper respiratory infections	0		-	20	0	-	-	Yes	Yes	Yes	Yes	
4	Low birth weight for gestation	Otitis media	0	-	-	20	0	-	-	Yes	Yes	Yes	Yes	
4	Low birth weight for gestation	Pneumococcal meningitis	0	-	-	20	0	-	-	Yes	Yes	Yes	Yes	
4	Low birth weight for gestation	H influenzae type B meningitis	0	-	-	20	0	-	-	Yes	Yes	Yes	Yes	
4	Low birth weight for gestation	Meningococcal infection	0	-	-	20	0	-	-	Yes	Yes	Yes	Yes	
4	Low birth weight for gestation	Encephalitis	0	-	-	20 20	0		-	Yes	Yes	Yes	Yes	
4	Low birth weight for gestation	Neonatal preterm birth complications	0	-	-	20	0		-	Yes	Yes	Yes	Yes	
4	Low birth weight for gestation	Neonatal encephalopathy due to birth	0	-	-	20	0	-	-	Yes	Yes	Yes	Yes	
4	Low birth weight for gestation	Neonatal sepsis and other neonatal	0	-	-	20	0	-	-	Yes	Yes	Yes	Yes	
4	Low birth weight for gestation	Hemolytic disease and other neonatal	0	-	-	20	0	-	-	Yes	Yes	Yes	Yes	
4	Low birth weight for gestation	Sudden infant death syndrome	0	-	-	20	0		-	Yes	Yes	Yes	Yes	
3	Vitamin A deficiency	Diarrhoeal diseases	19	0	63	0	-	-	-	No	-	Yes	No	
3	Vitamin A deficiency	Measles	12	0	83	0	-	-	-	Yes	-	Yes	No	
3	Zinc deficiency	Diarrhoeal diseases	14	0	29	0	-	-	-	No	-	Yes	No	
3	Zinc deficiency	Lower respiratory infections	6	0	17	0	-	-	-	No	-	Yes	No	
2	Smoking	Tuberculosis	0	-		4	0	10	0	No		Yes	Yes	
3	Smoking	Lip and oral cavity cancer	0	-	-	5	0	-	-	Yes	-	Yes	Yes	
3	Smoking	Nasopharynx cancer	0	-	-	4	0	28	0	Yes	-	Yes	Yes	
3	Smoking	Oesophageal cancer	0	-	-	5	0	-	-	Yes	-	Yes	Yes	
3	Smoking	Colon and rectum cancer	0	-	-	19	0	-	-	No	-	Yes	Yes	
3	Smoking	Gastric cancer	0		-	34 19	0	-	-	No	-	Yes	Yes	
3	Smoking	Pancreatic cancer	0	-	-	19	0	-	-	Yes	-	Yes	Yes	
3	Smoking	Larynx cancer	0	-	-	5	0	-	-	Yes	-	Yes	Yes	
3	Smoking	Tracheal, bronchus, and lung cancer	0	-	-	38	0	-	-	Yes	-	Yes	Yes	
3	Smoking	Breast cancer	0	-	-	19	0	-	-	No	-	Yes	Yes	
3	Smoking	Prostate cancer	0	-	-	15	0		-	No	-	Yes	Yes	
3	Smoking	Kidney cancer	0	-	-	8	0	-	-	Yes	-	Yes	Yes	
3	Smoking	Bladder cancer	0	-	-	37	0	-	-	Yes	-	Yes	Yes	
3	Smoking	Leukaemia	0	-	-	22	0	-	-	No	-	Yes	Yes	
3	Smoking	Ischaemic heart disease	0	-	-	86	-	-	-	No	-	Yes	Yes	
3	Smoking	Stroke Atrial fibrillation and flutter	0	-	-	60 16	-		-	N0 No	-	Y es Y es	Y es Y es	
3	Smoking	Peripheral vascular disease	0	-	-	10	0	-	-	No	-	Yes	Yes	
3	Smoking	Chronic obstructive pulmonary disease	0	-	-	42	0	-	-	Yes	-	Yes	Yes	
3	Smoking	Asthma	0	-	-	8	12	-	-	No	-	Yes	Yes	
3	Smoking	Peptic ulcer disease	0	-	-	7	0	-	-	No	-	Yes	No	
3	Smoking	Alzheimer's disease and other dementias	0		-	10	8	-		No	-	Yes	Yes	
3	Smoking	Parkinson's disease	0	-	-	8	0	-	-	Yes	-	Yes	Yes	
3	Smoking	Multiple sclerosis	0	-	-	6	0	-	-	No	-	Yes	Yes	
3	Smoking	Diabetes mellitus type 2	0	-	-	88	0	-	-	No	-	Yes	No	
3	Smoking	Rheumatoid arthritis	0	-	-	5	0	-	-	No	-	Yes	No	
3	Smoking	Low back pain	0	-	-	13	0	-	-	N0 No	-	Y es Vec	Y es	
3	Smoking	Macular degeneration	0	-	-	5	0	-	-	No	-	Yes	No	
3	Smoking	Non-Hip Fracture	0	-	-	14	14	-	-	No	-	Yes	Yes	
3	Smoking	Abdominal aortic aneurysm	0	-	-	10	0	-	-	No	-	Yes	Yes	
3	Smoking	Hip fracture	0	-	-	15	20	-	-	No	-	Yes	Yes	
3	Chewing tobacco	Lip and oral cavity cancer	0	-	-	4	0	21	5	Yes	-	Yes	Yes	
5		pringen enner	0			-	0	10	0	. 00		1.03	105	

vel			•	ith significant effect in the direction (%)	ith null findings (%)	ive observational studies	ive observational studies aificant association in the direction (%)	ttrol studies assessing the come pair relationship (n)**	trol studies that show nt association in the direction (%)	mit of RR > 1.5	ponse relationship	al plausibility +	++	
sk Le			CTs (n	Ts w posite	Ts w	ospect *	ospect th sig posite	se-con k-outo	se-con nifica posite	wer li	se-res	ologic	alogy	
Ris	Risk	Outcome	RC	RC op]	RC	L I	Prd wit	ris]	Ca sig op]	F	Ď	Bic	UN I	
3	Second-hand smoke	Stroke Lower respiratory infections	0	-		4	0	-	-	No No	Yes	Y es Y es	Yes	
3	Second-hand smoke	Otitis media	0	-	-	1	0	4	0	No	-	Yes	Yes	
3	Second-hand smoke	Tracheal, bronchus, and lung cancer	0	-	-	13	0	-	-	No	Yes	Yes	Yes	
3	Second-hand smoke	Breast cancer	0	-	-	21	0	-	-	No	-	Yes	Yes	
3	Second-hand smoke	Ischaemic heart disease	0	-	-	5	0	-	-	No	Yes	Yes	Yes	
3	Second-hand smoke	Diabetes mellitus type 2	0	-	-	5	0	-	-	No	-	Yes	Yes	
2	Alcohol use	Tuberculosis	0	-	-	9	0	18		Yes	Yes	Yes	Yes	
2	Alcohol use	Lower respiratory infections	0	-	-	3	0	2		Yes	Yes	Yes	Yes	
2	Alcohol use	Lip and oral cavity cancer	0	-	-	26	0	-	-	Yes	Yes	Yes	Yes	
2	Alcohol use	Nasopharynx cancer	0	-	-	6	0	-	-	Yes	Yes	Yes	Yes	
2	Alcohol use	Other pharynx cancer Oesonhageal cancer	0	-	-	19 57	0	-	-	Yes Yes	Yes	Yes Yes	Yes	
2	Alcohol use	Colon and rectum cancer	0	-	-	56	0	-	-	Yes	Yes	Yes	Yes	
2	Alcohol use	Liver cancer	0	-	-	32	0	-	-	Yes	Yes	Yes	Yes	
2	Alcohol use	Larynx cancer Breast cancer	0	-	-	35 123	0	-	-	Yes Yes	Yes	Yes Yes	Yes	
2	Alcohol use	Ischaemic heart disease	Ő	-	-	93	0	-	-	Yes	Yes	Yes	Yes	
2	Alcohol use	Ischaemic stroke	0	-	-	24	0	-	-	Yes	Yes	Yes	Yes	
2	Alcohol use	Haemorrhagic stroke Hypertensive heart disease	0	-	-	13	0	-	-	Yes Yes	Yes Yes	Y es Y es	Yes	
2	Alcohol use	Atrial fibrillation and flutter	0	-	-	9	0	-	-	Yes	Yes	Yes	Yes	
2	Alcohol use	Cirrhosis	0	-	-	14	0	-	-	Yes	Yes	Yes	Yes	
2	Alcohol use	Epilepsy	0	-	-	4	0	2	0	No	Yes	Yes	No	
2	Alcohol use	Diabetes mellitus type 2	0	-	-	43	0	-	-	Yes	Yes	Yes	No	
2	Alcohol use	Motor vehicle road injuries	0	-	-	3	0	-	-	Yes	Yes	Yes	Yes	
2	Alcohol use	Self-harm	0	-	-	4	0	4	0	Yes	Yes	Yes	Yes	
2	Alcohol use	Interpersonal violence	0	-	-	3	0	1	0	Yes	Yes	Yes	Yes	
2	Drug use	Hepatitis B	1	-	-	5	0	-	-	No	-	Yes	Yes	
2	Drug use	Self-harm	0	-	-	10	0	0	0	No	-	Yes	No	
2	Dietary risks													
3	Diet low in fruits	Lip and oral cavity cancer	0	-	-	2	0	15	0	No	Yes	Yes	Yes	
3	Diet low in fruits	Nasopharynx cancer	0	-	-	2	0	15	0	No	Yes	Yes	Yes	
3	Diet low in fruits	Other pharynx cancer	0	-	-	2	0	15	0	No No	Yes	Yes	Yes	
3	Diet low in fruits	Larvnx cancer	0	-	-	2	0	15	0	No	Yes	Yes	Yes	
3	Diet low in fruits	Tracheal, bronchus, and lung cancer	0	-	-	22	0	-	-	No	Yes	Yes	Yes	
3	Diet low in fruits	Ischaemic heart disease	0	-	-	9	0	-	-	No	Yes	Yes	Yes	
3	Diet low in fruits	Ischaemic stroke	0	-	-	9	0	-	-	No	Yes	Yes	Yes	
3	Diet low in fruits	Diabetes mellitus	0	-	-	9	0	-	-	No	Yes	Yes	No	
3	Diet low in vegetables	Oesophageal cancer	0	-	-	5	0	-	-	No	Yes	Yes	No	
3	Diet low in vegetables	Ischaemic stroke	0	-	-	9	0	-	-	No	Y es	Y es	Ves	
3	Diet low in vegetables	Haemorrhagic stroke	0		-	5	0	-	_	No	Yes	Yes	Yes	
3	Diet low in legumes	Ischaemic heart disease	0	-	-	5	0	-	-	No	Yes	Yes	No	
3	Diet low in whole grains	Ischaemic heart disease	0	-	-	7	0	-	-	No	Yes	Yes	Yes	
3	Diet low in whole grains	Ischaemic stroke	0	-	-	6	0	-	-	No	Yes	Yes	Yes	
3	Diet low in whole grains	Haemorrhagic stroke	0	-	-	6	0	-	-	No	Yes	Yes	Yes	
3	Diet low in whole grains	Diabetes mellitus	0	-	-	10	0	-	-	No	Yes	Yes	No	
3	Diet low in nuts and seeds	Diabetes mellitus	1	0	100	5	0	-	-	No	Ves	Ves	No	
3	Diet low in milk	Colon and rectum cancer	0	-	-	7	0	-	_	No	Yes	Yes	No	
3	Diet high in red meat	Colon and rectum cancer	0	-	-	8	0	-	-	No	Yes	Yes	No	
3	Diet high in red meat	Diabetes mellitus	0	-	-	9	11	-	-	No	Yes	Yes	No	
3	Diet high in processed meat	Colon and rectum cancer	0	-	-	9	11	-	-	No	Yes	Yes	No	
3	Diet high in processed meat	Ischaemic heart disease	0	-	-	5	0	-	-	No	Yes	Yes	No	
3	Diet high in sugar-sweetened heverages	Diabetes mellitus	0	-	-	8	0	-	-	N0 No	Y es Ves	Y es Ves	No	
3	Diet high in sugar-sweetened beverages	Ischemic heart disease	0	-	-	4	0	-	-	No	Yes	Yes	No	
3	Diet low in fibre	Colon and rectum cancer	0	-	-	15	0	-	-	No	Yes	Yes	No	
3	Diet low in fibre	Ischaemic heart disease	0	-	-	12	0	-	-	No	Yes	Yes	No	
3	Diet low in calcium	Colon and rectum cancer	0	-	-	13	0	-	-	No	Yes	Yes	No	
3	Diet low in seafood omega-3 fatty acids	Ischaemic heart disease	17	0	94	16	0	-	-	No	Yes	Yes	No	
3	Diet low in polyunsaturated fatty acids	Ischaemic heart disease	8	0	75	11	0	-	-	No N-	Yes	Yes	No	
3	Diet high in sodium	Ischaemic neart disease Stomach cancer	0	-	-	13	0	-	-	NO No	Y es Vec	Y es Vec	INO No	
3	Diet high in sodium	Systolic blood pressure	45	-	- 73	0	-	-	-	No	Yes	Yes	No	
2	Child maltreatment	,		-		-								
3	Childhood sexual abuse	Alcohol use disorders	0	-	-	2	0	3	0	No	-	Yes	Yes	
3	Childhood sexual abuse	Depressive disorders	0	-	-	7	0	-	-	No	-	Yes	Yes	
3	Bullying victimization	Anxiety disorders	0	-	-	19	0	0	0	No	Yes	Yes	No	

				nt effect in the 6)	lings (%)	ional studies	ional studies ciation in the 6)	assessing the lationship (n)**	that show in in the 6)	1.5	nship	ty +		
(Level			ſs (n)	Fs with significal osite direction (%	fs with null find	spective observat	spective observat significant asso osite direction (%	e-control studies -outcome pair re	e-control studies ificant associatio osite direction (%	/er limit of RR >	e-response relatio	logical plausibili	llogy ‡	
Risl	Risk	Outcome	RC	RC' opp	RC	Pro: (n)*	Pro: with op p	Cas risk	Cas sign opp	Lov	Dos	Bio	Ana	
3	Bullying victimization	Depressive disorders	0	-	-	19	0	0	0	No	Yes	Yes	No	
2	Intimate partner violence	HIV/AIDS Maternal abortion and miscarriage	0	-	-	2	0	0	0	N0 Ves	-	Y es Ves	No No	
2	Intimate partner violence	Depressive disorders	0	-	-	4	0	0	0	No	-	Yes	Yes	
2	Low physical activity	Colon and rectum cancer	0	-	-	20	15	-	-	No	Yes	Yes	Yes	
2	Low physical activity	Breast cancer	0	-	-	35	0	-	-	No	Yes	Yes	Yes	
2	Low physical activity	Ischaemic heart disease	0	-	-	45 27	9	-	-	No	Yes	Yes	Yes	
2	Low physical activity	Diabetes mellitus	0	-	-	57	7	-	-	No	Yes	Yes	No	
2	High fasting plasma glucose	Tuberculosis	0	-	-	18	0	-	-	Yes	Yes	Yes	No	
2	High fasting plasma glucose	Colon and rectum cancer	0	-	-	21	0	-	-	No	-	-	Yes	
2	High fasting plasma glucose	Liver cancer	0	-	-	28	0	-	-	Yes	-	-	No	
2	High fasting plasma glucose High fasting plasma glucose	Pancreatic cancer	0	-	-	35 16	0	-	-	Y es	-	-	Y es Y es	
2	High fasting plasma glucose	Breast cancer	0	-	-	39	0	-	-	No	-	-	Yes	
2	High fasting plasma glucose	Ovarian cancer	0	-	-	11	0	-	-	No	-	-	Yes	
2	High fasting plasma glucose	Bladder cancer	0	-	-	14	0	-	-	No	-	-	Yes	
2	High fasting plasma glucose	Ischaemic heart disease	8	0	100	150	-	-	-	Yes	Yes	Yes	Yes	
2	High fasting plasma glucose	Ischaemic stroke	9	0	100	150	-	-	-	Y es Vec	Y es Ves	Y es Ves	Y es Ves	
2	High fasting plasma glucose	Intracerebral hemorrhage	9	0	100	150	-	-	-	Yes	Yes	Yes	Yes	
2	High fasting plasma glucose	Alzheimer's disease and other dementias	0	-	-	17	0	-	-	No	-	-	No	
2	High fasting plasma glucose	Peripheral vascular disease	14	-	-	4	0	-	-	Yes	Yes	Yes	Yes	
2	High fasting plasma glucose	Chronic kidney disease	5	-	-	32	-	-	-	Yes	Yes	Yes	No	
2	High fasting plasma glucose	Glaucoma	0	-	-	5	0	-	-	No	-	-	Yes	
2	High Iasting plasma glucose High I DL cholesterol	Cataract Ischaemic heart disease	21	-	- 57	1	0	1	0	N0 Ves	- Ves	- Ves	Y es Ves	
2	High LDL cholesterol	Ischaemic stroke	21	0	57	88	-	-	-	Yes	Yes	Yes	Yes	
2	High systolic blood pressure	Rheumatic heart disease	0	-	-	62	-	-	-	Yes	Yes	Yes	Yes	
2	High systolic blood pressure	Ischaemic heart disease	56	0	-	88	-	-	-	Yes	Yes	Yes	Yes	
2	High systolic blood pressure	Ischaemic stroke	54	0	-	150	-	-	-	Yes	Yes	Yes	Yes	
2	High systolic blood pressure	Intracerebral hemorrhage	54	0	-	150	-	-	-	Yes	Yes	Yes	Yes	
2	High systolic blood pressure	Cardiomyonathy and myocarditis	54 0	0	-	62	-	-		Yes	Yes	r es Ves	Yes	
2	High systolic blood pressure	Other cardiomyopathy	0	-	-	62	-	-	-	Yes	Yes	Yes	Yes	
2	High systolic blood pressure	Atrial fibrillation and flutter	20	5	60	88	-	-	-	Yes	Yes	Yes	Yes	
2	High systolic blood pressure	Aortic aneurysm	0	-	-	62	-	-	-	Yes	Yes	Yes	Yes	
2	High systolic blood pressure	Peripheral vascular disease	0	-	-	88	-	-	-	Yes	Yes	Yes	Yes	
2	High systolic blood pressure	Endocarditis	0	-	-	62	-	-	-	Yes	Yes	Yes	Yes	
2	High systolic blood pressure	Non-rheumatic calcific aortic valve disease	0	-	-	2	-	-	-	No	Yes	Yes	Yes	
2	High systolic blood pressure	Chronic kidney disease	8	-	-	88	-	-	-	Yes	Yes	Yes	No	
2	High body-mass index (adult)	Oesophageal cancer	0	-	-	6	0	-	-	No	Yes	Yes	Yes	
2	High body-mass index (adult)	Non-hodgkin lymphoma	0	-	-	18	0	-	-	No	Yes	Yes	Yes	
2	High body-mass index (adult)	Colon and rectum cancer	0	-	-	38	0	-	-	No	Yes	Yes	Yes	
2	High body-mass index (adult)	Gallbladder and biliary tract cancer	0		-	54 10	0	-		No	Yes	r es Ves	Yes	
2	High body-mass index (adult)	Pancreatic cancer	0	-	-	20	0	-	-	No	Yes	Yes	Yes	
2	High body-mass index (adult)	Breast cancer (pre-menopausal)	0	-	-	44	2	-	-	No	Yes	Yes	Yes	
2	High body-mass index (adult)	Breast cancer (post-menopausal)	0	-	-	25	8	-	-	No	Yes	Yes	No	
2	High body-mass index (adult)	Uterine cancer	0	-	-	37	0	-	-	No	Yes	Yes	Yes	
2	High body-mass index (adult)	Ovarian cancer	0	-	-	31	3	-	-	N0 No	Yes	Yes	Yes	
2	High body-mass index (adult)	Thyroid cancer	0	-	-	28 16	0	-	-	No	Yes	Yes	Yes	
2	High body-mass index (adult)	Multiple myeloma	0	-	-	20	-	-	-	-	Yes	Yes	Yes	
2	High body-mass index (adult)	Leukaemia	0	-	-	17	0	-	-	No	Yes	Yes	Yes	
2	High body-mass index (adult)	Ischaemic heart disease	0	-	-	129	-	-	-	No	Yes	Yes	Yes	
2	High body-mass index (adult)	Ischaemic stroke	0	-	-	102	-	-	-	No	Yes	Yes	Yes	
2	High body-mass index (adult)	Haemonnagic stroke	0		-	85	-	-		No	Yes	r es Ves	Yes	
2	High body-mass index (adult)	Atrial fibrillation and flutter	0	-	-	5	0	-	-	-	No	Yes	Yes	
2	High body-mass index (adult)	Asthma	0	-	-	7	0	-	-	-	Yes	Yes	No	
2	High body-mass index (adult)	Alzheimer's disease and other dementias	0	-	-	6	0	-	-	-	No	Yes	No	
2	High body-mass index (adult)	Gallbladder disease	0	-	-	16	0	-	-	-	Yes	Yes	Yes	
2	High body-mass index (adult)	Diabetes mellitus	0	-	-	85 57	-	-	-	Yes	Yes	Yes	No	
2	High body-mass index (adult)	Osteoarthritis	0	-	-	32	- 0	-	-	No	Yes	Yes	Yes	
2	High body-mass index (adult)	Low back pain	0	-	-	5	õ	-	-	No	Yes	Yes	Yes	
2	High body-mass index (adult)	Gout	0	-	-	10	0	-	-	-	Yes	Yes	No	
2	High body-mass index (adult)	Cataract	0	-	-	17	0	-	-	-	Yes	Yes	No	
2	High body-mass index (child)	Asthma Injurios	0	-	-	5	0	-	-	No	Yes	Yes	No	
2	Low bone initicial density	injunto	U	-	-	12	-	-	-	1NO	1 05	1 es	1 es	

Risk Level	Risk	Outcome	RCTs (n)	RCTs with significant effect in the opposite direction (%)	RCTs with null findings (%)	Prospective observational studies (n)*	Prospective observational studies with significant association in the opposite direction (%)	Case-control studies assessing the risk-outcome pair relationship (n)**	Case-control studies that show significant association in the opposite direction (%)	Lower limit of RR > 1.5	Dose-response relationship	Biological plausibility +	Analogy ‡	
2	Impaired kidney function	Ischaemic heart disease	0	-	-	6	0	-	-	Yes	-	Yes	Yes	
2	Impaired kidney function	Ischaemic stroke	0	-	-	6	0	-	-	Yes	-	Yes	Yes	
	T 1111 C C	In the second well to see a sufficient of	0		-	8	0	-	-	Yes	-	Yes	Ves	
2	Impaired kidney function	intracerebrai nemorrnage	0	-	-	0	0			100		1 65	105	
2 2	Impaired kidney function	Peripheral vascular disease	0	-	-	5	0	-	-	Yes	-	Yes	Yes	

A. Citations		
Risk	Outcome	Citation/Note
Unsafe water	Diarrhoeal diseases	Cairncross S, Valdmanis V. Water Supply, Sanitation, and Hygiene Promotion. In: Jamison DT, Breman JG, Measham AR, et al., eds. Disease Control Priorities in Developing Countries, 2nd edn. Washington (DC): World Bank, 2006.
		Wolf J, Prüss-Ustün A, Cumming O, et al. Assessing the impact of drinking water and sanitation on diarrhoeal disease in low- and middle-income settings: systematic review and meta-regression. Trop Med Int
Unsafe water	Diarrhoeal diseases	Health 2014; 19: 928–42. Fewtrall L. Kaufmann P. B. Kay, D. Enanoria, W. Haller, L. & Colford, I. M. (2005). Water sanitation
Unsafe water	Diarrhoeal diseases	and hygiene interventions to reduce diarrhoea in less developed countries: a systematic review and meta- analysis. The Lancet Infectious Diseases, 5(1), 42-52. doi:10.1016/s1473-3099(04)01253-8
Unsafe sanitation - improved sanitation	Diarrhoeal diseases	Wolf J, Prüss-Ustün A, Cumming O, et al. Assessing the impact of drinking water and sanitation on diarrhoeal disease in low- and middle-income settings: systematic review and meta-regression. Trop Med Int Health 2014; 19: 928–42.
Unsafe sanitation - improved sanitation	Diarrhoeal diseases	Fewtrell, L., Kaufmann, R. B., Kay, D., Enanoria, W., Haller, L., & Colford, J. M. (2005). Water, sanitation, and hygiene interventions to reduce diarrhoea in less developed countries: a systematic review and meta- analysis. The Lancet Infectious Diseases, 5(1), 42-52. doi:10.1016/s1473-3099(04)01253-8
Unsafe sanitation - improved sanitation	Diarrhoeal diseases	Baker, K. K., O'Reilly, C. E., Levine, M. M., Kotloff, K. L., Nataro, J. P., Ayers, T. L., Mintz, E. D. (2016). Sanitation and Hygiene-Specific Risk Factors for Moderate-to-Severe Diarrhea in Young Children in the Global Enteric Multicenter Study, 2007–2011: Case-Control Study. PLOS Medicine,13(5). doi:10.1371/journal.pmed.1002010
		Genser, B., Strina, A., Santos, L. A., Teles, C. A., Prado, M. S., Cairncross, S., & Barreto, M. L. (2008). Impact of a city-wide sanitation intervention in a large urban centre on social, environmental and behavioural determinants of childhood diarrhoea: analysis of two cohort studies. International Journal of Epidemiology,
Unsafe sanitation - piped	Diarrhoeal diseases	37(4), 831-840. doi:10.1093/ije/dyn101
Unsafe sanitation - nined	Diarthoeal diseases	Norman, G., Pedley, S., & Takkouche, B. (2010). Effects of sewerage on diarrhoea and enteric infections: a systematic review and meta-analysis. The Lancet Infectious Diseases, 10(8), 536-544. doi:10.1016/s1473-3099(10)70123-7
ensure sumation piped	Diamocal diseases	Wolf J, Prüss-Ustün A, Cumming O, et al. Assessing the impact of drinking water and sanitation on diarrhoeal disease in low- and middle-income settings: systematic review and meta-regression. Trop Med Int
Unsafe sanitation - piped	Diarrhoeal diseases	Health 2014; 19: 928–42.
facility	Diarrhoeal diseases	Ejemot-Nwadiaro RI, Eniri JE, Arikpo D, Meremikwu MM, Critchiey JA. Hand wasning promotion for preventing diarrhoea. Cochrane Database Syst Rev 2015; : CD004265.
No access to handwashing facility	Lower respiratory infections	Rabie T, Curtis V. Handwashing and risk of respiratory infections: a quantitative systematic review. Trop Med Int Health Tropical Medicine and International Health. 2006;11(3):258–67.
No access to handwashing facility	Lower respiratory infections	Aiello, A. E., Coulborn, R. M., Perez, V., & Larson, E. L. (2008). Effect of Hand Hygiene on Infectious Disease Risk in the Community Setting: A Meta-Analysis. American Journal of Public Health, 98(8), 1372- 1381. doi:10.2105/ajph.2007.124610
Ambient particulate matter pollution	Lower respiratory infections	Hoek G, Krishnan RM, Beelen R, et al. Long-term air pollution exposure and cardio- respiratory mortality: a review. Environ Health 2013; 12: 43.
Ambient particulate matter pollution	Lower respiratory infections	Mehta S, Shin H, Burnett R, North T, Cohen AJ. Ambient particulate air pollution and acute lower respiratory infections: a systematic review and implications for estimating the global burden of disease. Air Qual Atmos Health 2013; 6: 69–83.
Ambient particulate matter	Lower respiratory infections	MacIntyre EA, Gehring U, Mölter A, Fuertes E, Klümper C, Krämer U, et al. Air Pollution and Respiratory Infections during Early Childhood: An Analysis of 10 European Birth Cohorts within the ESCAPE Project. Environ Health Perspect 2014 Jan 1:122(1):107–13
Ambient particulate matter	Tracheal, bronchus and lung	Raaschou-Nielsen O, Beelen R, Wang M, Hoek G, Andersen ZJ, Hoffmann B, et al. Particulate matter air
pollution	cancer	pollution components and risk for lung cancer. Environ Int. 2016 Feb;87:66–73.
pollution	cancer	and mortality: A meta-analysis. Oncotarget. 2017 Jun 27;8(26):43322–31.
Ambient particulate matter pollution	Ischaemic heart disease	Hoek G, Krishnan RM, Beelen R, et al. Long-term air pollution exposure and cardio- respiratory mortality: a review. Environ Health 2013; 12: 43.
Ambient particulate matter pollution	Ischaemic heart disease	Beelen R, Stafoggia M, Raaschou-Nielsen O, Andersen ZJ, Xun WW, Katsouyanni K, et al. Long-term exposure to air pollution and cardiovascular mortality: an analysis of 22 European cohorts. Epidemiology. 2014 May;25(3):368–78.
Ambient particulate matter pollution	Stroke	Beelen R, Stafoggia M, Raaschou-Nielsen O, Andersen ZJ, Xun WW, Katsouyanni K, et al. Long-term exposure to air pollution and cardiovascular mortality: an analysis of 22 European cohorts. Epidemiology. 2014 May:25(3):368–78.
Ambient particulate matter	Stroko	Scheers H, Jacobs L, Casas L, Nemery B, Nawrot TS. Long-Term Exposure to Particulate Matter Air Pollution Is a Pick Footor for Strake: Mate Analytical Evidence. Strake, 2015 New 44(11):2058, 66
ponution	Suoke	Cohen AJ, Brauer M, Burnett R, Anderson HR, Frostad J, Estep K, et al. Estimates and 25-year trends of the
Ambient particulate matter pollution	Chronic obstructive pulmonary disease	global burden of disease attributable to ambient air pollution: an analysis of data from the Global Burden of Diseases Study 2015. The Lancet. 2017 May 13;389(10082):1907–18.
Ambient particulate matter pollution	Diabetes mellitus type 2	He D, Wu S, Zhao H, Qiu H, Fu Y, Li X, et al. Association between particulate matter 2.5 and diabetes mellitus: A meta-analysis of cohort studies. J Diabetes Investig. 2017 Sep;8(5):687–96.
Household air pollution from solid fuels	Lower respiratory infections	Dherani M, Pope D, Mascarenhas M, Smith KR, Weber M, Bruce N. Indoor air pollution from unprocessed solid fuel use and pneumonia risk in children aged under five years: a systematic review and meta-analysis. Bull World Health Organ 2008; 86: 390–398C.
Household air pollution from solid fuels	Lower respiratory infections	Jary, H., Simpson, H., Havens, D., Manda, G., Pope, D., Bruce, N., & Mortimer, K. (2016). Household Air Pollution and Acute Lower Respiratory Infections in Adults: A Systematic Review. Plos One, 11(12). doi:10.1371/journal.pone.0167656

Appendix Table 5. Epidemiol Citations and B. Additional in A. Citations	ogical evidence supporting causal nformation	ity between risk-outcome pairs included in the Global Burden of Disease 2017 study including A.
Risk	Outcome	Citation/Note
Household air pollution from	Tracheal, bronchus and lung	Kurmi OP, Arya PH, Lam K-BH, Sorahan T, Ayres JG. Lung cancer risk and solid fuel smoke exposure: a
solid fuels	cancer	systematic review and meta-analysis. Eur Respir J. 2012 Nov;40(5):1228-37.
Household air pollution from		Fatmi Z, Coggon D. Coronary heart disease and household air pollution from use of solid fuel: a systematic
solid fuels	Ischaemic heart disease	review. Br Med Bull. 2016 Jun;118(1):91–109.
solid fuels	Stroke	review Br Med Bull 2016 Jun 118(1):91–109
Household air pollution from	Chronic obstructive pulmonary	Kurmi OP, Arya PH, Lam K-BH, Sorahan T, Ayres JG. Lung cancer risk and solid fuel smoke exposure: a
solid fuels	disease	systematic review and meta-analysis. Eur Respir J. 2012 Nov;40(5):1228-37.
Household air pollution from	Dishetes mellitus tune 2	Kim C, Seow WJ, Shu X-O, Bassig BA, Rothman N, Chen BE, et al. Cooking Coal Use and All-Cause and Cause-Specific Mortality in a Prospective Cohort Study of Women in Shanghai, China. Environ Health
solid fuels	Diabetes mentus type 2	West, S., Bates, M., Lee, J., Schaumberg, D., Lee, D., Adair-Rohani, H., Arai, H. (2013). Is Household
Household air pollution from solid fuels	Cataract	Air Pollution a Risk Factor for Eye Disease? International Journal of Environmental Research and Public Health, 10(11), 5378-5398. doi:10.3390/ijerph10115378
		Weichenthal S, Pinault LL, Burnett RT. Impact of Oxidant Gases on the Relationship between Outdoor Fine
Ambient ozone pollution	disease	Particulate Air Pollution and Nonaccidental, Cardiovascular, and Respiratory Mortality. Scientific Reports
Ambient ozone ponution	Chronic obstructive pulmonary	Turner MC. Jerrett M. Pope CA. et al. Long-Term Ozone Exposure and Mortality in a Large Prospective
Ambient ozone pollution	disease	Study. Am J Respir Crit Care Med 2016; 193: 1134–42.
Ambient ozone pollution	Chronic obstructive pulmonary disease	Carey IM, Atkinson RW, Kent AJ, van Staa T, Cook DG, Anderson HR. Mortality Associations with Long- Term Exposure to Outdoor Air Pollution in a National English Cohort. Am J Respir Crit Care Med. 2013 Apr 3;187(11):1226–33.
Å	Tracheal, bronchus and lung	Torres-Durán MCAD, Barros-Dios JM, Fernández-Villar A, Ruano-Ravina A. Residential radon and lung
Residential radon	cancer	cancer in never smokers. A systematic review. Cancer Letters. 2014;345(1):21-6.
	Trachael knowskus and hung	Turner MC Kenneli D. Cher V. Deze CA. Constan S. Thur MI. Deden and Lune Constant the American
Residential radon	cancer	Cancer Society Cohort Cancer Epidemiology Biomarkers & Prevention 20111un 20(3):438–48
	Tracheal, bronchus and lung	Krewski D, Lubin JH, Zielinski JM, Alavanja M, Catalan VS, Field RW, et al. Residential Radon and Risk of
Residential radon	cancer	Lung Cancer. Epidemiology. 2005;16(2):137-45.
	Tracheal, bronchus and lung	Darby S. Radon in homes and risk of lung cancer: collaborative analysis of individual data from 13 European
Residential radon	cancer	case-control studies. Bmj. 2005;330(7485):223.
Residential radon	cancer	2003Jan:104(4):315–9.
	Tracheal, bronchus and lung	Kreuzer M, Gerken M, Kreienbrock L, Wellmann J, Wichmann HE. Lung cancer in lifetime nonsmoking men
Residential radon	cancer	- results of a case-control study in Germany. British Journal of Cancer. 2001May;84(1):134-40.
Desidential and an	Tracheal, bronchus and lung	Kreuzer M, Heinrich J, Kreienbrock L, Rosario AS, Gerken M, Wichmann HE. Risk factors for lung cancer
Residential radon	cancer	Field R Steck DI Smith BI Brus CP Fisher FL Neuberger IS et al. The Jowa radon lung cancer study —
Residential radon	Tracheal, bronchus and lung cancer	phase I: residential radon gas exposure and lung cancer. Science of The Total Environment. 2001;272(1-3):67–72.
Desidential and an	Tracheal, bronchus and lung	Wang Z. Residential Radon and Lung Cancer Risk in a High-exposure Area of Gansu Province, China.
Residential fadoli	Tracheal bronchus and lung	Lagarde F Axelsson G Damber I. Mellander H Nyberg F Pershagen G Residential Radon and Lung
Residential radon	cancer	Cancer among Never-Smokers in Sweden. Epidemiology. 2001;12(4):396–404.
	Tracheal, bronchus and lung	Pershagen G, Akerblom G, Axelson O, Clavensjo B, Damber L, Desai G, et al. Residential Radon Exposure
Residential radon	cancer	and Lung Cancer in Sweden. New England Journal of Medicine. 1994;330(3):159–64.
Residential radon	Tracheal, bronchus and lung cancer	Alavanja MCR, Brownson RC, Lubin JH, Berger E, Chang J, Bolce JD. Residential Radon Exposure and Lung Cancer Among Nonsmoking Women. JNCI Journal of the National Cancer Institute. 1994;86(24):1829–37.
Residential radon	Tracheal, bronchus and lung cancer	Schoenberg JB, Klotz JB, Wilcox HB, Nicholls GP, Gil-del-Real MT, Stemhagen A. Case-control study of residential radon and lung cancer among New Jersey women. Cancer Research. 1990Oct15;50(20):6520–4.
Lead exposure	Systolic blood pressure	Navas-Acien A, Schwartz BS, Kothenberg SJ, Hu H, Silbergeld EK, Guallar E. Bone lead levels and blood
Lead exposure	Systeme blood pressure	Glenn BS, Stewart WF, Links JM, Todd AC, Schwartz BS. The Longitudinal Association of Lead with Blood
Lead exposure	Systolic blood pressure	Pressure. Epidemiology. 2003;14(1):30-6.
		Glenn BS, Bandeen-Roche K, Lee B-K, Weaver VM, Todd AC, Schwartz BS. Changes in Systolic Blood
Lead exposure	Systolic blood pressure	Pressure Associated With Lead in Blood and Bone. Epidemiology. 2006;17(5):538–44.
Lead exposure	Systolic blood pressure	women. American Journal of Public Health. 1999:89(3):330–5.
Loud onposulo	Systeme erood pressure	Cheng Y, Schwartz J, Sparrow D, Aro A, Weiss ST, Hu H. Bone Lead and Blood Lead Levels in Relation to
Lead exposure	Systolic blood pressure	Baseline Blood Pressure and the Prospective Development of Hypertension The Normative Aging Study. American Journal of Epidemiology. 2001;153(2):164–71.
Lead exposure	Idiopathic intellectual disability	Lanphear BP, Hornung R, Khoury J, et al. Low-level environmental lead exposure and children's intellectual function; an international pooled analysis. Environ Health Perspect 2005: 113: 894–9.
	1	Liu J, Li L, Wang Y, Yan C, Liu X. Impact of low blood lead concentrations on IQ and school performance
Lead exposure	Idiopathic intellectual disability	in Chinese children. PLoS ONE 2013; 8: e65230.
Lead exposure	Idionathic intellectual disability	Lanphear BP, Hornung R, Khoury J, et al. Low-level environmental lead exposure and children's intellectual function: an international pooled analysis. Environ Health Perspect 2005; 113: 894–9

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Risk	Outcome	Citation/Note
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Lead exposure	Idiopathic intellectual disability	Doses of Lead in Childhood. New England Journal of Medicine. 1990Nov;322(2):83-8.
Occupational exposure to		Goodman M, Morgan RW, Ray R, Malloy CD, Zhao K. Cancer in asbestos-exposed occupational cohorts: a
asbestos	Larynx cancer	meta-analysis. Cancer Causes Control 1999; 10: 453-65.
		Lenters V, Vermeulen R, Dogger S, et al. A meta-analysis of asbestos and lung cancer: is better quality
Occupational exposure to	Tracheal, bronchus and lung	exposure assessment associated with steeper slopes of the exposure-response relationships? Environ Health
asbestos	cancer	Perspect 2011; 119: 1547–55.
Occupational exposure to		Camargo MC, Stayner LT, Straif K, et al. Occupational exposure to asbestos and ovarian cancer: a meta-
asbestos	Ovarian cancer	analysis. Environ Health Perspect 2011; 119: 1211-7.
Occupational exposure to		Bourdès V, Boffetta P, Pisani P. Environmental exposure to asbestos and risk of pleural mesothelioma:
asbestos	Mesothelioma	review and meta-analysis. Eur J Epidemiol 2000; 16: 411-7.
		Lenters V, Vermeulen R, Dogger S, et al. A meta-analysis of asbestos and lung cancer: is better quality
Occupational exposure to	Tracheal, bronchus and lung	exposure assessment associated with steeper slopes of the exposure-response relationships? Environ Health
arsenic	cancer	Perspect 2011; 119: 1547–55.
Occupational exposure to		Khalade A, Jaakkola MS, Pukkala E, Jaakkola JJK. Exposure to benzene at work and the risk of leukemia: a
benzene	Leukaemia	systematic review and meta-analysis. Environ Health 2010; 9: 31.
Occupational exposure to	Tracheal, bronchus and lung	Boffetta P, Fryzek JP, Mandel JS. Occupational exposure to beryllium and cancer risk: a review of the
beryllium	cancer	epidemiologic evidence. Crit Rev Toxicol 2012; 42: 107-18.
Occupational exposure to	Tracheal, bronchus and lung	Verougstraete V, Lison D, Hotz P. Cadmium, lung and prostate cancer: a systematic review of recent
cadmium	cancer	epidemiological data. J Toxicol Environ Health B Crit Rev 2003; 6: 227-55.
Occupational exposure to	Tracheal, bronchus and lung	Denis Ambroise, Pascal Wild and Jean-Jacques Moulin, Scandinavian Journal of Work, Environment &
chromium	cancer	Health, Vol. 32, No. 1 (February 2006), pp. 22-31
Occupational exposure to diesel	Tracheal, bronchus and lung	Lipsett M, Campleman S. Occupational exposure to diesel exhaust and lung cancer: a meta-analysis. Am J
engine exhaust	cancer	Public Health 1999; 89: 1009–17.
Occupational exposure to		Hauptmann M, Lubin JH, Stewart PA, Hayes RB, Blair A. Mortality from solid cancers among workers in
formaldehyde	Nasopharynx cancer	formaldehyde industries. Am J Epidemiol 2004; 159: 1117-30.
Occupational exposure to	^ · ·	Collins JJ, Lineker GA. A review and meta-analysis of formaldehyde exposure and leukemia. Regul Toxicol
formaldehyde	Leukaemia	Pharmacol 2004; 40: 81–91.
	Tracheal, bronchus and lung	Grimsrud TK, Berge SR, Haldorsen T, Andersen A. Can lung cancer risk among nickel refinery workers be
Occupational exposure to nickel	cancer	explained by occupational exposures other than nickel? Epidemiology 2005; 16: 146-54.
Occupational exposure to		
polycyclic aromatic	Tracheal, bronchus and lung	Armstrong B, Hutchinson E, Unwin J, Fletcher T. Lung cancer risk after exposure to polycyclic aromatic
hydrocarbons	cancer	hydrocarbons: a review and meta-analysis. Environ Health Perspect 2004; 112: 970-8.
		Liu Y, Steenland K, Rong Y, Hnizdo E, Huang X, Zhang H, et al. Exposure-Response Analysis and Risk
	Tracheal, bronchus and lung	Assessment for Lung Cancer in Relationship to Silica Exposure: A 44-Year Cohort Study of 34,018 Workers.
Occupational exposure to silica	cancer	Am J Epidemiol. 2013 Nov 1;178(9):1424–33.
Occupational exposure to		Soskolne CL, Jhangri GS, Siemiatycki J, et al. Occupational exposure to sulfuric acid in southern Ontario,
sulfuric acid	Larynx cancer	Canada, in association with laryngeal cancer. Scand J Work Environ Health 1992; 18: 225-32.
Occupational exposure to		Kelsh MA, Alexander DD, Mink PJ, Mandel JH. Occupational trichloroethylene exposure and kidney cancer:
trichloroethylene	Kidney cancer	a meta-analysis. Epidemiology 2010; 21: 95-102.
		Karialainan A. Kurnna K. Martikainan P. Klaukka T. Karialainan I. Work is related to a substantial portion
		of adult onset asthma incidence in the Finnish population. Am I Bespir Crit Care Med 2001: 164: 565.8
Occupational asthmagens	Asthma	of adult-onset asumna meldence in the r ministi population. And 5 Kespir erit eare Med 2001, 104. 505–6.
Occupational particulate matter,	Chronic obstructive pulmonary	Blanc PD, Iribarren C, Trupin L, et al. Occupational exposures and the risk of COPD: dusty trades revisited.
gases, and fumes	disease	Thorax 2009; 64: 6–12.
		Agrawal Y, Platz EA, Niparko JK. Prevalence of hearing loss and differences by demographic characteristics
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Occupational noise	Age-related and other hearing loss	Med 2008; 168: 1522–30.
		Davis A. The prevalence of hearing impairment and reported hearing disability among adults in Great Britain.
Occupational noise	Age-related and other hearing loss	International Journal of Epidemiology 1989,18: 911-917.
		Wilson D, Walsh P, Sanchez L, Davis A, Taylor A, Tucker G, Meagher I. The epidemiology of hearing
Occupational noise	Age-related and other hearing loss	impairment in an Australian adult population. International Journal of Epidemiology 1999;28:247-252.
		International Labour Organization. Resolution concerning statistics of occupational injuries (resulting from
		occupational accidents). 1998; published online Oct. http://www.ilo.org/global/statistics-and-
		databases/standards-and-guidelines/resolutions-adopted-by-international-conferences-of-labour-
Occupational injuries	Injuries	statisticians/WCMS_087528/langen/index.htm.
		Eurostat. Accidents at work statistics. http://ec.europa.eu/eurostat/statistics-
Occupational injuries	Injuries	explained/index.php/Accidents_at_work_statistics.
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Occupational ergonomic factors	Low back pain	from the Global Burden of Disease 2010 study. Ann Rheum Dis 2014; 73: 975–81.
		Horta BL, Victora CG. Short-term effects of breastfeeding: a systematic review on the benefits of
		breastfeeding on diarrhoea and pneumonia mortality. World Health Organization, 2013
Non-exclusive breastfeeding	Lower respiratory infections	http://allattamento.sip.it/wp-content/uploads/2014/03/WHO_breve-termine.pdf.
		Horta BL, Victora CG. Short-term effects of breastfeeding: a systematic review on the benefits of
		breastfeeding on diarrhoea and pneumonia mortality. World Health Organization, 2013
Non-exclusive breastfeeding	Diarrhoeal diseases	nttp://allattamento.sip.it/wp-content/uploads/2014/03/WHO_breve-termine.pdf.
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D'accella a 11 - 22 - 11		mortainty in children under five years: a pooled analysis of ten prospective studies. PLoS ONE 2013; 8:
Discontinued breastfeeding	Diarrnoeal diseases	e04030.

Risk	Outcome	Citation/Note
Discontinued breastfeeding	Diarrhoeal diseases	Genser, B., Strina, A., Santos, L. A., Teles, C. A., Prado, M. S., Cairncross, S., & Barreto, M. L. (2008). Impact of a city-wide sanitation intervention in a large urban centre on social, environmental and behavioural determinants of childhood diarrhoea: analysis of two cohort studies. International Journal of Epidemiology, 37(4) 831-840. doi:10.1093/iie/dw101
Discontinued oredstreeding		Olofin I, McDonald CM, Ezzati M, et al. Associations of suboptimal growth with all-cause and cause-specific mortality in children under five years: a pooled analysis of ten prospective studies. PLoS ONE 2013; 8:
Childhood underweight	Diarrhoeal diseases	e64636. Olofin I, McDonald CM, Ezzati M, et al. Associations of suboptimal growth with all-cause and cause-specific
Childhood underweight	Lower respiratory infections	mortality in children under five years: a pooled analysis of ten prospective studies. PLoS ONE 2013; 8: e64636.
Childhood underweight	Measles	Olofin I, McDonald CM, Ezzati M, et al. Associations of suboptimal growth with all-cause and cause-specific mortality in children under five years: a pooled analysis of ten prospective studies. PLoS ONE 2013; 8: e64636.
Childhood wasting	Diarrhoeal diseases	Olofin I, McDonald CM, Ezzati M, et al. Associations of suboptimal growth with all-cause and cause-specific mortality in children under five years: a pooled analysis of ten prospective studies. PLoS ONE 2013; 8: e64636.
Childhood wasting	Lower respiratory infections	Olofin I, McDonald CM, Ezzati M, et al. Associations of suboptimal growth with all-cause and cause-specific mortality in children under five years: a pooled analysis of ten prospective studies. PLoS ONE 2013; 8: e64636.
Childhood wasting	Measles	Olofin I, McDonald CM, Ezzati M, et al. Associations of suboptimal growth with all-cause and cause-specific mortality in children under five years: a pooled analysis of ten prospective studies. PLoS ONE 2013; 8: e64636.
Childhood stunting	Diarrhoeal diseases	Olofin I, McDonald CM, Ezzati M, et al. Associations of suboptimal growth with all-cause and cause-specific mortality in children under five years: a pooled analysis of ten prospective studies. PLoS ONE 2013; 8: e64636.
Childhood stunting	Lower respiratory infections	Olofin I, McDonald CM, Ezzati M, et al. Associations of suboptimal growth with all-cause and cause-specific mortality in children under five years: a pooled analysis of ten prospective studies. PLoS ONE 2013; 8: e64636.
Childhood stunting	Measles	Olofin I, McDonald CM, Ezzati M, et al. Associations of suboptimal growth with all-cause and cause-specific mortality in children under five years: a pooled analysis of ten prospective studies. PLoS ONE 2013; 8: e64636.
Iron deficiency	Maternal hemorrhage	Murray-Kolb LE, Chen L, Chen P, Shapiro M, Caulfield L. CHERG Iron Report: Maternal Mortality, Child Mortality, Perinatal Mortality, Child Cognition, and Estimates of Prevalence of Anemia due to Iron Deficiency. Baltimore, USA: CHERG, 2012.
Vitamin A deficiency	Diarrhoeal diseases	Awasthi S, Peto R, Read S, et al. Vitamin A supplementation every 6 months with retinol in 1 million pre- school children in north India: DEVTA, a cluster-randomised trial. Lancet 2013; 381: 1469–77.
Vitamin A deficiency	Diarrhoeal diseases	Diness BR, Christoffersen D, Pedersen UB, Rodrigues A, Fischer TK, Andersen A, Whittle H, Yazdanbakhsh M, Aaby P, Benn CS. The effect of high-dose vitamin A supplementation given with bacille Calmette-Guérin vaccine at birth on infant rotavirus infection and diarrhea: a randomized prospective study from Guinea-Bissau. J Infect Dis. 2010; S243-251.
Vitamin A deficiency	Diarrhoeal diseases	Imdad A, Herzer K, Mayo-Wilson E, Yakoob MY, Bhutta ZA. Vitamin A supplementation for preventing morbidity and mortality in children from 6 months to 5 years of age. Cochrane Database Syst Rev. 2010; CD008524.
Vitamin A deficiency	Diarrhoeal diseases	Imdad A, Yakoob MY, Sudfeld C, Haider BA, Black RE, Bhutta ZA. Impact of vitamin A supplementation on infant and childhood mortality. BMC Public Health. 2011; S20.
Vitamin A deficiency	Measles	Awasthi S, Peto R, Read S, et al. Vitamin A supplementation every 6 months with retinol in 1 million pre- school children in north India: DEVTA, a cluster-randomised trial. Lancet 2013; 381: 1469–77.
Vitamin A deficiency	Measles	Imdad A, Herzer K, Mayo-Wilson E, Yakoob MY, Bhutta ZA. Vitamin A supplementation for preventing morbidity and mortality in children from 6 months to 5 years of age. Cochrane Database Syst Rev. 2010; CD008524.
Vitamin A deficiency	Measles	Imdad A, Yakoob MY, Sudfeld C, Haider BA, Black RE, Bhutta ZA. Impact of vitamin A supplementation on infant and childhood mortality. BMC Public Health. 2011; S20.
Zinc deficiency	Diarrhoeal diseases	Shankar AH, Prasad AS. Zinc and immune function: the biological basis of altered resistance to infection. Am J Clin Nutr 1998; 68: 447S–463S.
		Yakoob MY, Theodoratou E, Jabeen A, et al. Preventive zinc supplementation in developing countries: impact on mortality and morbidity due to diarrhea, pneumonia and malaria. BMC Public Health 2011; 11
Zinc deficiency	Diarrhoeal diseases	Suppl 3: S23. Shankar AH, Prasad AS. Zinc and immune function: the biological basis of altered resistance to infection.
Zinc deficiency	Lower respiratory infections	Am J Clin Nutr 1998; 68: 447S–463S. Yakoob MY, Theodoratou E, Jabeen A, et al. Preventive zinc supplementation in developing countries:
Zinc deficiency	Lower respiratory infections	impact on mortality and morbidity due to diarrhea, pneumonia and malaria. BMC Public Health 2011; 11 Suppl 3: S23.
Smoking	Lower respiratory infections	Surgeon General's Report - The Health Consequences of Smoking. U.S. Department of Health & Human Services, 2004 http://www.cdc.gov/tobacco/data_statistics/sgr/2004/.
Smoking	Larynx cancer	Carter BD, Abnet CC, Feskanich D, et al. Smoking and mortalitybeyond established causes. N Engl J Med 2015; 372: 631–40.
Smoking	Lip and oral cavity cancer	Carter BD, Abnet CC, Feskanich D, et al. Smoking and mortalitybeyond established causes. N Engl J Med 2015; 372: 631–40.

Appendix Table 5. Ep Citations and B. Addi A. Citations	idemiological evidence supporting causa itional information	lity between risk-outcome pairs included in the Global Burden of Disease 2017 study including A.
Risk	Outcome	Citation/Note
Smoking	Tracheal, bronchus and lung	International Agency for Research on Cancer Working Group on the Evaluation of Carcinogenic Risks to Humans. IARC monographs on the evaluation of carcinogenic risks to humans: Tobacco Smoke and Involuntary Smoking, Lyon: IARC, 2004
Shloking	cancer	Ordóñez-Mena JM, Schöttker B, Mons U, et al. Quantification of the smoking-associated cancer risk with rate advancement periods: meta-analysis of individual participant data from cohorts of the CHANCES
Smoking	Breast cancer	consortium. BMC Medicine 2016; 14: 62.
Smoking	Cervical cancer	International Agency for Research on Cancer Working Group on the Evaluation of Carcinogenic Risks to Humans. IARC monographs on the evaluation of carcinogenic risks to humans: Tobacco Smoke and Involuntary Smoking. Lyon: IARC, 2004.
C C		Islami F, Moreira DM, Boffetta P, Freedland SJ. A Systematic Review and Meta-analysis of Tobacco Use and
Smoking	Prostate cancer	Prostate Cancer Mortality and Incidence in Prospective Cohort Studies. European Urology 2014; 66: 1054–64.
Smoking	Kidney cancer	Humans. IARC monographs on the evaluation of carcinogenic risks to humans: Tobacco Smoke and Involuntary Smoking. Lyon: IARC, 2004.
Smoking	Nasopharynx cancer	Xue W-Q, Qin H-D, Ruan H-L, Shugart YY, Jia W-H. Quantitative Association of Tobacco Smoking With the Risk of Nasopharyngeal Carcinoma: A Comprehensive Meta-Analysis of Studies Conducted Between 1979 and 2011. Am J Epidemiol 2013; 178: 325–38.
Smoking	Bladder cancer	Cumberbatch MG, Rota M, Catto JWF, La Vecchia C. The Role of Tobacco Smoke in Bladder and Kidney Carcinogenesis: A Comparison of Exposures and Meta-analysis of Incidence and Mortality Risks. Eur Urol 2016; 70: 458–66.
Smoking	Laukaamia	Colamesta V, D'Aguanno S, Breccia M, Bruffa S, Cartoni C, Torre GL. Do the smoking intensity and duration, the years since quitting, the methodological quality and the year of publication of the studies affect the results of the meta-analysis on cigarette smoking and Acute Myeloid Leukemia (AML) in adults? Critical Paviane in Oneology (Hemetaleon 2016: 00: 376–88
Shloking	Leukaeinna	Surgeon General's Report - The Health Consequences of Smoking. U.S. Department of Health & Human
Smoking	Oesophageal cancer	Services, 2004 http://www.cdc.gov/tobacco/data_statistics/sgr/2004/.
Smoking	Stomach cancer	rate advancement periods: meta-analysis of individual participant data from cohorts of the CHANCES consortium. BMC Medicine 2016; 14: 62.
Smoking	Colon and rectum cancer	Surgeon General's Report - The Health Consequences of Smoking. U.S. Department of Health & Human Services 2004 http://www.cdc.gov/tobacco/data_statistics/sgr/2004/
Smoning		Surgeon General's Report - The Health Consequences of Smoking. U.S. Department of Health & Human
Smoking	Liver cancer	Services, 2004 http://www.cdc.gov/tobacco/data_statistics/sgr/2004/. Ordóñez-Mena JM, Schöttker B, Mons U, et al. Quantification of the smoking-associated cancer risk with
Smoking	Pancreatic cancer	consortium. BMC Medicine 2016; 14: 62.
Smoking	Ischaemic heart disease	Huxley RR, Woodward M. Cigarette smoking as a risk factor for coronary heart disease in women compared with men: a systematic review and meta-analysis of prospective cohort studies. Lancet 2011; 378: 1297–305.
		Peters SAE, Huxley RR, Woodward M. Smoking as a risk factor for stroke in women compared with men: a systematic review and meta-analysis of 81 cohorts, including 3,980,359 individuals and 42,401 strokes.
Smoking	Cerebrovascular disease	Stroke 2013; 44: 2821–8.
Smoking	Atrial fibrillation and flutter	A meta-analysis of prospective studies. Int J Cardiol 2016; 218: 259–66.
Smoking	Abdominal aortic aneurysm	Lederle FA, Nelson DB, Joseph AM. Smokers' relative risk for aortic aneurysm compared with other smoking-related diseases: a systematic review. J Vasc Surg 2003; 38: 329–34.
Smoking	Perinheral vascular disease	Lu L, Mackay DF, Pell JP. Meta-analysis of the association between cigarette smoking and peripheral arterial disease. Heart 2014: 100: 414–23
Shioking	Chronic obstructive pulmonary	Forey BA, Thornton AJ, Lee PN. Systematic review with meta-analysis of the epidemiological evidence
Smoking	disease	relating smoking to COPD, chronic bronchitis and emphysema. BMC Pulm Med 2011; 11: 36. Jayes L, Haslam PL, Gratziou CG, et al. SmokeHaz: Systematic Reviews and Meta-analyses of the Effects of
Smoking	Asthma	Smoking on Respiratory Health. Chest 2016; 150: 164–79. Kurata IH, Nogawa AN, Meta-analysis of risk factors for peptic ulcer. Nonsteroidal antiinflammatory drugs
Smoking	Peptic ulcer disease	Helicobacter pylori, and smoking. J Clin Gastroenterol 1997; 24: 2–17.
Smoking	Gallbladder and biliary tract disease	Aune D, Vatten LJ, Boffetta P. Tobacco smoking and the risk of gallbladder disease. Eur J Epidemiol 2016; 31: 643–53.
Smoking	Alzheimer disease and other dimentias	Zhong G, Wang Y, Zhang Y, Guo JJ, Zhao Y. Smoking Is Associated with an Increased Risk of Dementia: A Meta-Analysis of Prospective Cohort Studies with Investigation of Potential Effect Modifiers. PLoS One 2015; 10. DOI:10.1371/journal.pone.0118333.
Smoking	Parkinson disease	Li X, Li W, Liu G, Shen X, Tang Y. Association between cigarette smoking and Parkinson's disease: A meta- analysis. Arch Gerontol Geriatr 2015; 61: 510–6.
Smoking	Multiple sclerosis	O'Gorman C, Broadley SA. Smoking and multiple sclerosis: evidence for latitudinal and temporal variation. J Neurol 2014: 261: 1677-83
Shloking	wumple scierosis	Nonioi 2014, 201. 10/7-03.
Smoking	Diabetes mellitus type 2	Pan A, Wang Y, Talaei M, Hu FB, Wu T. Relation of active, passive, and quitting smoking with incident type 2 diabetes: a systematic review and meta-analysis. The Lancet Diabetes & Endocrinology 2015; 3: 958–67.
Smoking	Rheumatoid arthritis	Sugiyama D, Nishimura K, Tamaki K, et al. Impact of smoking as a risk factor for developing rheumatoid arthritis: a meta-analysis of observational studies. Ann Rheum Dis 2010; 69: 70–81.

Appendix Table 5. Epider Citations and B. Addition A. Citations	niological evidence supporting causa al information	lity between risk-outcome pairs included in the Global Burden of Disease 2017 study including A.
Risk	Outcome	Citation/Note Shiri R, Karppinen J, Leino-Arjas P, Solovieva S, Viikari-Juntura E. The Association between Smoking and
Smoking	Low back pain	Low Back Pain: A Meta-analysis. The American Journal of Medicine 2010; 123: 87.e7-87.e35.
Smoking	Cataract	Sci 2012; 53: 3885–95.
Smoking	Macular degeneration	Chakravarthy U, Wong TY, Fletcher A, et al. Chinical risk factors for age-related macular degeneration: a systematic review and meta-analysis. BMC Ophthalmol 2010; 10: 31.
Smoking	Injuries	Vestergaard P, Mosekhoe L. Fracture fisk associated with smoking: a meta-analysis. J intern Med 2005; 254: 572–83.
Chewing Tobacco	Lip and oral cavity cancer	Siddiqi K, Shah S, Abbas SM, et al. Global burden of disease due to smokeless tobacco consumption in adults: analysis of data from 113 countries. BMC Med 2015; 13: 194.
Chewing Tobacco	Oesophageal cancer	adults: analysis of data from 113 countries. BMC Med 2015; 13: 194.
Second-hand smoke	Breast cancer	Macacu A, Autier P, Boniol M, Boyle P. Active and passive smoking and risk of breast cancer: a meta- analysis. Breast Cancer Res Treat 2015; 154: 213–24.
Second-hand smoke	Chronic obstructive pulmonary disease	Fischer F, Kraemer A. Meta-analysis of the association between second-hand smoke exposure and ischaemic heart diseases, COPD and stroke. BMC Public Health 2015; 15: 1202.
Second-hand smoke	Diabetes mellitus type 2	Zhu B, Wu X, Wang X, Zheng Q, Sun G. The association between passive smoking and type 2 diabetes: a meta-analysis. Asia Pac J Public Health 2014; 26: 226–37.
Second-hand smoke	Tuberculosis	Dogar OF, Pillai N, Safdar N, Shah SK, Zahid R, Siddiqi K. Second-hand smoke and the risk of tuberculosis: a systematic review and a meta-analysis. Epidemiol Infect 2015; 143: 3158–72.
		Baker RJ, Hertz-Picciotto I, Dostal M, Keller JA, Nozicka J, Kotesovec F, Dejmek J, Loomis D, Sram RJ. Coal home heating and environmental tobacco smoke in relation to lower respiratory illness in Czech
Second-hand smoke	Lower respiratory infections	children, from birth to 3 years of age. Environ Health Perspect. 2006; 1126-32.
Second-hand smoke	Lower respiratory infections	Blizzard L, Ponsonby A-L, Dwyer T, Venn A, Cochrane JA. Parental smoking and infant respiratory infection: how important is not smoking in the same room with the baby?. Am J Public Health. 2003; 482-8.
Second-hand smoke	Lower respiratory infections	exploratory study from India. Health Policy. 2004; 67-83.
Second-hand smoke	Lower respiratory infections	Broor S, Pandey RM, Ghosh M, Maitreyi RS, Lodha R, Singhal T, Kabra SK. Risk factors for severe acute lower respiratory tract infection in under-five children. Indian Pediatr. 2001; 1361-9.
Second-hand smoke	Lower respiratory infections	chen Y, Li WX, Yu SZ, Qian WH. Chang-Ning epidemiological study of children's health: I: Passive smoking and children's respiratory diseases. Int J Epidemiol. 1988; 348-55.
		Duijts L, Jaddoe VWV, Hofman A, Steegers EAP, Mackenbach JP, de Jongste JC, Moll HA. Maternal smoking in pre-natal and early post-natal life and the risk of respiratory tract infections in infancy. The
Second-hand smoke	Lower respiratory infections	Ekwo EE, Weinberger MM, Lachenbruch PA, Huntley WH. Relationship of parental smoking and gas
Second-hand smoke	Lower respiratory infections	cooking to respiratory disease in children. Chest. 1983; 662-8. Etiler N. Velipasaoglu S. Aktekin M. Incidence of acute respiratory infections and the relationship with some
Second-hand smoke	Lower respiratory infections	factors in infancy in Antalya, Turkey. Pediatr Int 2002; 44: 64–9.
Second-hand smoke	Lower respiratory infections	rems BG, ware JH, Berkey CS, Dockery DW, Spiro A, Speizer FE. Effects of passive smoking on nearth of children. Environ Health Perspect. 1985; 289-95. Forestiere F. Corbo GM, Michelozzi P. Pistelli R. Agabiti N. Brancato G. Ciappi G. Perucci CA. Effects of
Second-hand smoke	Lower respiratory infections	environment and passive smoking on the respiratory health of children. Int J Epidemiol. 1992; 66-73.
Second-hand smoke	Lower respiratory infections	Gardner G, Frank AL, Taber LH. Effects of social and family factors on viral respiratory infection and illness in the first year of life. J Epidemiol Community Health. 1984; 42-8.
Second-hand smoke	Lower respiratory infections	Hassan MK, Al-Sadoon I. Risk factors for severe pneumonia in children in Basrah. Trop Doct. 2001; 139-41. Koch A. Molhak K. Homos P. Soransan P. Hiuler T. Olesan ME. Pail I. Pedersan FK. Olean OP. Melhya M.
Second-hand smoke	Lower respiratory infections	Risk factors for acute respiratory tract infections in young Greenlandic children. Am J Epidemiol. 2003; 374- 84.
Second-hand smoke	Lower respiratory infections	Kristensen IA, Olsen J. Determinants of acute respiratory infections in Sowetoa population-based birth cohort. S Afr Med J. 2006: 633-40.
		Margolis PA, Keyes LL, Greenberg RA, Bauman KE, LaVange LM. Urinary cotinine and parent history (questionnaire) as indicators of passive smoking and predictors of lower respiratory illness in infants. Pediatr
Second-hand smoke	Lower respiratory infections	Pulmonol. 1997; 417-23. Nuesslein TG, Beckers D, Rieger CH. Cotinine in meconium indicates risk for early respiratory tract
Second-hand smoke	Lower respiratory infections	infections. Hum Exp Toxicol. 1999; 283-90. Ogston SA, Florey CD, Walker CH. The Tayside infant morbidity and mortality study: effect on health of
Second-hand smoke	Lower respiratory infections	using gas for cooking. BMJ. 1985; 957-60. Ooston SA, Florey CD, Walker CH, Association of infant alimentary and respiratory illness with parental
Second-hand smoke	Lower respiratory infections	smoking and other environmental factors. J Epidemiol Community Health. 1987; 21-5.
Second-hand smoke	Lower respiratory infections	redretra FA, Guandolo VL, Feroli EJ, Mella GW, Weiss IP. Involuntary smoking and incidence of respiratory illness during the first year of life. Pediatrics. 1985; 594-7.
Second-hand smoke	Lower respiratory infections	Rylander E, Pershagen G, Eriksson M, Bermann G. Parental smoking, urinary cotinine, and wheezing bronchitis in children. Epidemiology. 1995; 289-93.
Second-hand smoke	Lower respiratory infections	Suzuki M, Thiem VD, Yanai H, Matsubayashi T, Yoshida LM, Tho LH, Minh TT, Anh DD, Kilgore PE, Ariyoshi K. Association of environmental tobacco smoking exposure with an increased risk of hospital admissions for pneumonia in children under 5 years of age in Vietnam. Thorax, 2009 484-9

Appendix Table 5. Epidemiological evidence supporting causality between risk-outcome pairs included in the Global Burden of Disease 2017 study including A. **Citations and B. Additional information** A. Citations Risk Outcome Citation/Note Taylor B, Wadsworth J. Maternal smoking during pregnancy and lower respiratory tract illness in early life. Second-hand smoke Lower respiratory infections Arch Dis Child. 1987; 786-91. Victora CG, Fuchs SC, Flores JA, Fonseca W, Kirkwood B. Risk factors for pneumonia among children in a Brazilian metropolitan area. Pediatrics. 1994; 977-85. Second-hand smoke Lower respiratory infections Jones LL, Hassanien A, Cook DG, Britton J, Leonardi-Bee J. Parental smoking and the risk of middle ear Second-hand smoke Otitis media disease in children: a systematic review and meta-analysis. Arch Pediatr Adolesc Med 2012; 166: 18-27. Tracheal, bronchus, and lung Jayes L, Haslam PL, Gratziou CG, et al. 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Vitamin C and other compounds in vitamin C rich Alcohol use Tuberculosis food in relation to risk of tuberculosis in male smokers. Am J Epidemiol 1999; 150: 632-41. Brown KE, Campbell AH. Tobacco, alcohol and tuberculosis. British Journal of Diseases of the Chest 1961; Tuberculosis Alcohol use 55: 150-8. Buskin SE, Gale JL, Weiss NS, Nolan CM. Tuberculosis risk factors in adults in King County, Washington, Alcohol use Tuberculosis 1988 through 1990. Am J Public Health 1994; 84: 1750-6. Crampin AC, Glynn JR, Floyd S, et al. Tuberculosis and gender: exploring the patterns in a case control study Alcohol use Tuberculosis in Malawi. Int J Tuberc Lung Dis 2004; 8: 194-203. Lewis JG, Chamberlain DA. Alcohol consumption and smoking habits in male patients with pulmonary Tuberculosis tuberculosis Br J Prev Soc Med 1963: 17: 149-52 Alcohol use Rosenman KD, Hall N. Occupational risk factors for developing tuberculosis. Am J Ind Med 1996; 30: Alcohol use Tuberculosis 148-54. Tekkel M, Rahu M, Loit HM, Baburin A. Risk factors for pulmonary tuberculosis in Estonia. Int J Tuberc Alcohol use Tuberculosis Lung Dis 2002; 6: 887-94. Tocque K, Bellis MA, Beeching NJ, Syed Q, Remmington T, Davies PD. A case-control study of lifestyle risk Alcohol use Tuberculosis factors associated with tuberculosis in Liverpool, North-West England. Eur Respir J 2001; 18: 959-64. Zaridze D, Brennan P, Boreham J, et al. Alcohol and cause-specific mortality in Russia: a retrospective case-Alcohol use Tuberculosis control study of 48,557 adult deaths. Lancet 2009; 373: 2201-14. Samokhvalov AV, Irving HM, Rehm J. Alcohol consumption as a risk factor for pneumonia: a systematic Alcohol use Lower respiratory infections review and meta-analysis. Epidemiol Infect 2010; 138: 1789-95. Baik I, Curhan GC, Rimm EB, Bendich A, Willett WC, Fawzi WW. A prospective study of age and lifestyle factors in relation to community-acquired pneumonia in US men and women. Arch Intern Med 2000; 160: Alcohol use 3082-8. Lower respiratory infections Kornum JB, Due KM, Nørgaard M, et al. Alcohol drinking and risk of subsequent hospitalisation with Alcohol use Lower respiratory infections pneumonia. Eur Respir J 2012; 39: 149-55. Shen C, Ni MY, Schooling CM, Chan WM, Lee SY, Lam TH. Alcohol use and death from respiratory disease Alcohol use Lower respiratory infections in a prospective Chinese elderly cohort study in Hong Kong. Prev Med 2013; 57: 819-23. Almirall J, Bolíbar I, Serra-Prat M, et al. New evidence of risk factors for community-acquired pneumonia: a Alcohol use Lower respiratory infections population-based study. Eur Respir J 2008; 31: 1274-84. Almirall J, Bolíbar I, Balanzó X, González CA. Risk factors for community-acquired pneumonia in adults: a Alcohol use Lower respiratory infections population-based case-control study. Eur Respir J 1999; 13: 349-55. Bagnardi V, Blangiardo M, La Vecchia C, Corrao G. 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Lifetime and baseline alcohol intake and risk of cancer of the upper aero-digestive tract in the European Prospective Investigation into Cancer and Nutrition (EPIC) study. Int J Alcohol use Lip and oral cavity cancer Cancer 2009; 125: 406-12. Martinez I. Factors associated with cancer of the esophagus, mouth, and pharynx in Puerto Rico. J Natl Alcohol use Lip and oral cavity cancer Cancer Inst 1969; 42: 1069-94. Bagnardi V, Blangiardo M, La Vecchia C, Corrao G. A meta-analysis of alcohol drinking and cancer risk. Br Alcohol use Nasopharynx cancer J Cancer 2001; 85: 1700-5. Bagnardi V, Blangiardo M, La Vecchia C, Corrao G. A meta-analysis of alcohol drinking and cancer risk. Br Alcohol use Other pharynx cancer J Cancer 2001; 85: 1700-5. Bagnardi V, Blangiardo M, La Vecchia C, Corrao G. A meta-analysis of alcohol drinking and cancer risk. Br Alcohol use Oesophageal cancer J Cancer 2001; 85: 1700-5. Allen NE, Beral V, Casabonne D, et al. Moderate alcohol intake and cancer incidence in women. J Natl

Cancer Inst 2009; 101: 296-305.

Alcohol use

Oesophageal cancer

A. Citations		
Risk	Outcome	Citation/Note
Alcohol use	Oesophageal cancer	Boffetta P, Garfinkel L. Alcohol drinking and mortality among men enrolled in an American Cancer Society prospective study. Epidemiology 1990; 1: 342–8.
Alcohol use	Oesophageal cancer	Fan Y, Yuan J-M, Wang R, Gao Y-T, Yu MC. Alcohol, tobacco, and diet in relation to esophageal cancer: the Shanghai Cohort Study. Nutr Cancer 2008; 60: 354–63.
Alcohol use	Oesophageal cancer	Freedman ND, Abnet CC, Leitzmann MF, et al. A prospective study of tobacco, alcohol, and the risk of esophageal and gastric cancer subtypes. Am J Epidemiol 2007; 165: 1424–33.
		Ishiguro S, Sasazuki S, Inoue M, Kurahashi N, Iwasaki M, Tsugane S. Effect of alcohol consumption, cigarette smoking and flushing response on esophageal cancer risk: a population-based cohort study (JPHC
Alcohol use	Oesophageal cancer	study). Cancer Lett 2009; 275: 240–6. Kim MK, Ko MJ, Han JT. Alcohol consumption and mortality from all-cause and cancers among 1.34
Alcohol use	Oesophageal cancer	million Koreans: the results from the Korea national health insurance corporation's health examinee cohort in 2000. Cancer Causes Control 2010; 21: 2295–302.
Alcohol use	Oesophageal cancer	Kimm H, Kim S, Jee SH. The independent effects of cigarette smoking, alcohol consumption, and serum aspartate aminotransferase on the alanine aminotransferase ratio in korean men for the risk for esophageal cancer. Yonsei Med J 2010; 51: 310–7.
Alcohol use	Oesophageal cancer	Kono S, Ikeda M, Tokudome S, Nishizumi M, Kuratsune M. Cigarette smoking, alcohol and cancer mortality: a cohort study of male Japanese physicians. Jpn J Cancer Res 1987; 78: 1323–8.
Alcohol use	Oesophageal cancer	Nakaya N, Tsubono Y, Kuriyama S, et al. Alcohol consumption and the risk of cancer in Japanese men: the Miyagi cohort study. Eur J Cancer Prev 2005; 14: 169–74.
Alcohol use	Oesophageal cancer	Ozasa K. Alcohol use and mortality in the Japan Collaborative Cohort Study for Evaluation of Cancer (JACC). Asian Pac J Cancer Prev 2007: 8 Suppl: 81–8.
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Alcohol use	Oesophageal cancer	Smith M, Zhou M, Whitlock G, et al. Esophageal cancer and body mass index: results from a prospective study of 220,000 men in China and a meta-analysis of published studies. Int J Cancer 2008; 122: 1604–10.
Alcohol use	Oesophageal cancer	Wu M, Zhang Z-F, Kampman E, et al. Does family history of cancer modify the effects of lifestyle risk factors on esophageal cancer? A population-based case-control study in China. Int J Cancer 2011; 128: 2147–57.
Alcohol use	Colon and rectum cancer	Bagnardi V, Blangiardo M, La Vecchia C, Corrao G. A meta-analysis of alcohol drinking and cancer risk. Br J Cancer 2001; 85: 1700–5.
		Akhter M. Kuriyama S. Nakaya N. et al. Alcohol consumption is associated with an increased risk of distal
Alcohol use	Colon and rectum cancer	colon and rectal cancer in Japanese men: the Miyagi Cohort Study. Eur J Cancer 2007; 43: 383–90. Allen NE, Beral V, Casabonne D, et al. Moderate alcohol intake and cancer incidence in women. J Natl
Alcohol use	Colon and rectum cancer	Cancer Inst 2009; 101: 296–305.
Alcohol use	Colon and rectum cancer	frequency and cancer-specific mortality in the US population. Am J Epidemiol 2011; 174: 1044–53.
Alcohol use	Colon and rectum cancer	Study in China. Eur J Epidemiol 2005; 20: 149–54.
Alcohol use	Colon and rectum cancer	Chyou PH, Nomura AM, Stemmermann GN. A prospective study of colon and rectal cancer among Hawaii Japanese men. Ann Epidemiol 1996; 6: 276–82. Crockett SD, Long MD, Dellon ES, Martin CF, Galanko JA, Sandler RS. Inverse relationship between
Alcohol use	Colon and rectum cancer	moderate alcohol intake and rectal cancer: analysis of the North Carolina Colon Cancer Study. Dis Colon Rectum 2011; 54: 887–94.
Alcohol use	Colon and rectum cancer	Ferrari P, Jenab M, Norat T, et al. Lifetime and baseline alcohol intake and risk of colon and rectal cancers in the European prospective investigation into cancer and nutrition (EPIC). Int J Cancer 2007; 121: 2065–72. Flood A, Caprario L, Chaterjee N, Lacey JVJ, Schairer C, Schatzkin A. Folate, methionine, alcohol, and
Alcohol use	Colon and rectum cancer	colorectal cancer in a prospective study of women in the United States. Cancer Causes Control 2002; 13: 551-61.
Alcohol use	Colon and rectum cancer	Gaziano JM, Gaziano TA, Glynn RJ, et al. Light-to-moderate alcohol consumption and mortality in the Physicians' Health Study enrollment cohort. J Am Coll Cardiol 2000; 35: 96–105.
Alcohol use	Colon and rectum cancer	Kabat GC, Miller AB, Jain M, Rohan TE. Dietary intake of selected B vitamins in relation to risk of major cancers in women. Br J Cancer 2008: 99: 816–21.
		Kim MK, Ko MJ, Han JT. Alcohol consumption and mortality from all-cause and cancers among 1.34 million Koreans: the results from the Korea national health insurance corporation's health examinee cohort in
Alcohol use	Colon and rectum cancer	2000. Cancer Causes Control 2010; 21: 2295–302. Klatsky AL Armstrong MA Friedman GD, Hiatt RA. The relations of alcoholic beverage use to colon and
Alcohol use	Colon and rectum cancer	rectal cancer. Am J Epidemiol 1988; 128: 1007–15. Kono S. Ikoda M. Takudoma S. Nichianni M. Kuratauna M. Cigaratta smaking, alaohal and cancer
Alcohol use	Colon and rectum cancer	mortality: a cohort study of male Japanese physicians. Jpn J Cancer Res 1987; 78: 1323–8.
Alcohol use	Colon and rectum cancer	cancer in the Korean elderly]. J Prev Med Public Health 2008; 41: 23–9.
Alcohol use	Colon and rectum cancer	Miyagi cohort study. Eur J Cancer Prev 2005; 14: 169–74.
Alaphal was	Colon and mature errors	Otani 1, Iwasaki M, Yamamoto S, et al. Alcohol consumption, smoking, and subsequent risk of colorectal cancer in middle-aged and elderly Japanese men and women: Japan Public Health Center-based prospective study. Cancer Endomial Riemarkows Peru 2002; 12: 1402–500.
Alcohol use	Colon and rectum cancer	Ozasa K. Alcohol use and mortality in the Japan Collaborative Cohort Study for Evaluation of Cancer (JACC) Asian Pac J Cancer Prev 2007: 8 Suppl: 81–8

Appendix Table 5. Epidemiological evidence supporting causality between risk-outcome pairs included in the Global Burden of Disease 2017 study including A. **Citations and B. Additional information** A. Citations Risk Outcome Citation/Note Pedersen A, Johansen C, Gronbaek M. Relations between amount and type of alcohol and colon and rectal Alcohol use Colon and rectum cancer cancer in a Danish population based cohort study. Gut 2003; 52: 861-7. Sanjoaquin MA, Appleby PN, Thorogood M, Mann JI, Key TJ. Nutrition, lifestyle and colorectal cancer incidence: a prospective investigation of 10998 vegetarians and non-vegetarians in the United Kingdom. Br J Alcohol use Colon and rectum cancer Cancer 2004; 90: 118-21. Thun MJ, Peto R, Lopez AD, et al. Alcohol consumption and mortality among middle-aged and elderly U.S. adults. N Engl J Med 1997; 337: 1705-14. Alcohol use Colon and rectum cancer Thygesen LC, Wu K, Gronbaek M, Fuchs CS, Willett WC, Giovannucci E. Alcohol intake and colorectal cancer: a comparison of approaches for including repeated measures of alcohol consumption. Epidemiology Alcohol use Colon and rectum cancer 2008: 19: 258-64 Toriola AT, Kurl S, Laukanen JA, Mazengo C, Kauhanen J. Alcohol consumption and risk of colorectal Alcohol use Colon and rectum cancer cancer: the Findrink study. Eur J Epidemiol 2008; 23: 395-401. Wakai K, Kojima M, Tamakoshi K, et al. Alcohol consumption and colorectal cancer risk: findings from the Alcohol use Colon and rectum cancer JACC Study. J Epidemiol 2005; 15 Suppl 2: S173-179. Wu AH, Paganini-Hill A, Ross RK, Henderson BE. Alcohol, physical activity and other risk factors for colorectal cancer: a prospective study. Br J Cancer 1987; 55: 687-94. Alcohol use Colon and rectum cancer Yamamoto S, Nakagawa T, Matsushita Y, et al. Visceral fat area and markers of insulin resistance in relation Colon and rectum cancer Alcohol use to colorectal neoplasia. Diabetes Care 2010; 33: 184-9. Yi S-W, Sull JW, Linton JA, Nam CM, Ohrr H. Alcohol consumption and digestive cancer mortality in Alcohol use Colon and rectum cancer Koreans: the Kangwha Cohort Study. J Epidemiol 2010; 20: 204-11. Yuan JM, Ross RK, Gao YT, Henderson BE, Yu MC. Follow up study of moderate alcohol intake and Alcohol use Colon and rectum cancer mortality among middle aged men in Shanghai, China. BMJ 1997; 314: 18-23. Bagnardi V, Blangiardo M, La Vecchia C, Corrao G. A meta-analysis of alcohol drinking and cancer risk. Br

Alcohol use	Larynx cancer	J Cancer 2001; 85: 1700–5.
		Allen NE, Beral V, Casabonne D, et al. Moderate alcohol intake and cancer incidence in women. J Natl
Alcohol use	Larynx cancer	Cancer Inst 2009; 101: 296-305.
		Freedman ND, Schatzkin A, Leitzmann MF, Hollenbeck AR, Abnet CC. Alcohol and head and neck cancer
Alcohol use	Larynx cancer	risk in a prospective study. Br J Cancer 2007; 96: 1469-74.
		Garavello W, Bosetti C, Gallus S, et al. Type of alcoholic beverage and the risk of laryngeal cancer. Eur J
Alcohol use	Larynx cancer	Cancer Prev 2006; 15: 69–73.
		Kim MK, Ko MJ, Han JT. Alcohol consumption and mortality from all-cause and cancers among 1.34
		million Koreans: the results from the Korea national health insurance corporation's health examinee cohort in
Alcohol use	Larynx cancer	2000. Cancer Causes Control 2010; 21: 2295–302.
		Bagnardi V, Blangiardo M, La Vecchia C, Corrao G. A meta-analysis of alcohol drinking and cancer risk. Br
Alcohol use	Breast cancer	J Cancer 2001; 85: 1700–5.
		Allen NE, Beral V, Casabonne D, et al. Moderate alcohol intake and cancer incidence in women. J Natl
Alcohol use	Breast cancer	Cancer Inst 2009; 101: 296–305.
	_	Baglietto L, English DR, Gertig DM, Hopper JL, Giles GG. Does dietary folate intake modify effect of
Alcohol use	Breast cancer	alcohol consumption on breast cancer risk? Prospective cohort study. BMJ 2005; 331: 807.
1.1.1.1	Devert	Breslow RA, Chen CM, Graubard BI, Mukamal KJ. Prospective study of alcohol consumption quantity and
Alconol use	Breast cancer	requency and cancer-specific mortality in the US population. Am J Epidemiol 2011; 1/4: 1044–55.
lachal usa	Presst senser	chen WY, Colditz GA, Kosner B, et al. Use of postmenopausal normones, alconol, and risk for invasive
Alconol use	Breast cancer	breast cancer. Ann Intern Med 2002; 157: 798–804.
Alcohol use	Ischaemic heart disease	disease: a systematic review and meta-analysis. Addiction 2012: 107: 1246-60
Alcohol use	Isenaenne neart uisease	Ponkslav PE Brian SE Turner PL Mukamal KL Chali WA Association of electral consumption with
loobol use	Ischaemic heart disease	selected cardiovascular disease outcomes: a systematic review and meta-analysis BMI 2011: 342: d671
ileonor use	isenaemie neure disease	Corrao G Rubbiati L Bagnardi V Zambon A Poikolainen K Alcohol and coronary heart disease: a meta-
Alcohol use	Ischaemic heart disease	analysis. Addiction 2000: 95: 1505–23.
		Zhao J, Stockwell T, Roemer A, Naimi T, Chikritzhs T. Alcohol Consumption and Mortality From Coronary
Alcohol use	Ischaemic heart disease	Heart Disease: An Updated Meta-Analysis of Cohort Studies. J Stud Alcohol Drugs 2017; 78: 375-86.
		Zhang X-Y, Shu L, Si C-J, et al. Dietary Patterns, Alcohol Consumption and Risk of Coronary Heart Disease
Alcohol use	Ischaemic heart disease	in Adults: A Meta-Analysis. Nutrients 2015; 7: 6582-605.
		Zheng Y-L, Lian F, Shi Q, et al. Alcohol intake and associated risk of major cardiovascular outcomes in
		women compared with men: a systematic review and meta-analysis of prospective observational studies.
Alcohol use	Ischaemic heart disease	BMC Public Health 2015; 15: 773.
		Patra J, Taylor B, Irving H, et al. Alcohol consumption and the risk of morbidity and mortality for different
Alcohol use	Stroke	stroke typesa systematic review and meta-analysis. BMC Public Health 2010; 10: 258.
		Briasoulis A, Agarwal V, Messerli FH. Alcohol consumption and the risk of hypertension in men and
Alcohol use	Hypertensive heart disease	women: a systematic review and meta-analysis. J Clin Hypertens (Greenwich) 2012; 14: 792-8.
		Taylor B, Irving HM, Baliunas D, et al. Alcohol and hypertension: gender differences in dose-response
Alcohol use	Hypertensive heart disease	relationships determined through systematic review and meta-analysis. Addiction 2009; 104: 1981–90.
		Kodama S, Saito K, Tanaka S, et al. Alcohol consumption and risk of atrial fibrillation: a meta-analysis. J
Alcohol use	Atrial fibrillation and flutter	Am Coll Cardiol 2011; $5/:427-36$.

Samokhvalov AV, Irving HM, Rehm J. Alcohol consumption as a risk factor for atrial fibrillation: a Atrial fibrillation and flutter systematic review and meta-analysis. Eur J Cardiovasc Prev Rehabil 2010; 17: 706-12. Rehm J, Taylor B, Mohapatra S, et al. Alcohol as a risk factor for liver cirrhosis: a systematic review and meta-analysis. Drug Alcohol Rev 2010; 29: 437-45.

Alcohol use

Alcohol use

Cirrhosis

Appendix Table 5. Epi Citations and B. Addit A. Citations	idemiological evidence supporting ca tional information	usality between risk-outcome pairs included in the Global Burden of Disease 2017 study including A.
Risk	Outcome	Citation/Note
Alcohol use	Cirrhosis	Becker U, Grønbaek M, Johansen D, Sørensen TIA. Lower risk for alcohol-induced cirrhosis in wine drinkers. Hepatology 2002; 35: 868–75.
Alcohol use	Cirrhosis	Becker U, Deis A, Sørensen TI, et al. Prediction of risk of liver disease by alcohol intake, sex, and age: a prospective population study. Hepatology 1996; 23: 1025–9.
Alcohol use	Cirrhosis	Bellentani S, Saccoccio G, Costa G, et al. Drinking habits as cofactors of risk for alcohol induced liver damage. The Dionysos Study Group. Gut 1997; 41: 845–50.
Alcohol use	Cirrhosis	Blackwelder WC, Yano K, Rhoads GG, Kagan A, Gordon T, Palesch Y. Alcohol and mortality: the Honolulu Heart Study. Am J Med 1980: 68: 164–9.
Alcohol use	Cirrhosis	Boffetta P, Garfinkel L. Alcohol drinking and mortality among men enrolled in an American Cancer Society prospective study. Enidemiology 1990: 1: 342–8
	Ci l i	Fuchs CS, Stampfer MJ, Colditz GA, et al. Alcohol consumption and mortality among women. N Engl J Med
Alcohol use	Cirrhosis	1995; 332: 1245–50.
Alcohol use Alcohol use	Cirrhosis Cirrhosis	Garfinkel L, Boffetta P, Stellman SD. Alcohol and breast cancer: a cohort study. Prev Med 1988; 17: 686–93.Gordon T, Doyle JT. Drinking and mortality. The Albany Study. Am J Epidemiol 1987; 125: 263–70.
Alcohol use	Cirrhosis	Gordon T, Kannel WB. Drinking and mortality. The Framingham Study. Am J Epidemiol 1984; 120: 97–107.
Alcohol use	Cirrhosis	Kono S, Ikeda M, Tokudome S, Nishizumi M, Kuratsune M. Alcohol and mortality: a cohort study of male
Alashal	Cinhosis	Thun MJ, Peto R, Lopez AD, et al. Alcohol consumption and mortality among middle-aged and elderly U.S.
Alcohol use	Cirrhosis	adults. N Engl J Med 1997; 337: 1705–14. Yuan JM, Ross RK, Gao YT, Henderson BE, Yu MC. Follow up study of moderate alcohol intake and
Alcohol use	Cirrhosis	mortality among middle aged men in Shanghai, China. BMJ 1997; 314: 18–23. Bagnardi V, Rota M, Botteri E, et al. Alcohol consumption and site-specific cancer risk: a comprehensive
Alcohol use	Liver cancer	dose-response meta-analysis. Br J Cancer 2015; 112: 580–93. Allen NE, Beral V, Casabonne D, et al. Moderate alcohol intake and cancer incidence in women. J Natl
Alcohol use	Liver cancer	Cancer Inst 2009; 101: 296–305.
Alcohol use	Liver cancer	hepatocellular carcinoma in Korea. J Natl Cancer Inst 2004; 96: 1851–6.
Alcohol use	Liver cancer	Joshi S, Song Y-M, Kim T-H, Cho S-I. Socio-economic status and the risk of liver cancer mortality: a prospective study in Korean men. Public Health 2008; 122: 1144–51.
		Kim MK, Ko MJ, Han JT. Alcohol consumption and mortality from all-cause and cancers among 1.34 million Koreans: the results from the Korea national health insurance corporation's health examinee cohort in
Alcohol use	Liver cancer	2000. Cancer Causes Control 2010; 21: 2295–302.
Alcohol use	Liver cancer	Koh W-P, Robien K, Wang R, Govindarajan S, Yuan J-M, Yu MC. Smoking as an independent risk factor for hepatocellular carcinoma: the Singapore Chinese Health Study. Br J Cancer 2011: 105: 1430–5
Alcohol use	Liver concer	Kono S, Ikeda M, Tokudome S, Nishizumi M, Kuratsune M. Cigarette smoking, alcohol and cancer
Alcohol use		Nakaya N, Tsubono Y, Kuriyama S, et al. Alcohol consumption and the risk of cancer in Japanese men: the
Alcohol use	Liver cancer	Miyagi cohort study. Eur J Cancer Prev 2005; 14: 169–74. Ozasa K. Alcohol use and mortality in the Japan Collaborative Cohort Study for Evaluation of Cancer
Alcohol use	Liver cancer	(JACC). Asian Pac J Cancer Prev 2007; 8 Suppl: 81–8. Yi S-W, Sull JW, Linton JA, Nam CM, Ohrr H. Alcohol consumption and digestive cancer mortality in
Alcohol use	Liver cancer	Koreans: the Kangwha Cohort Study. J Epidemiol 2010; 20: 204–11. Alsamarrai A. Das SI M. Windsor IA. Petrov MS. Factors that affect risk for nancreatic disease in the general
Alcohol use	Pancreatitis	population: a systematic review and meta-analysis of prospective cohort studies. Clin Gastroenterol Hepatol 2014: 12: 1635–1644 e5: mir e103
Alashalwas	Danamatikia	Irving HM, Samokhvalov AV, Rehm J. Alcohol as a risk factor for pancreatitis. A systematic review and
Alconol use	Pancreattus	Samokhvalov AV, Rehm J, Roerecke M. Alcohol Consumption as a Risk Factor for Acute and Chronic
Alcohol use	Pancreatitis	 Pancreatitis: A Systematic Review and a Series of Meta-analyses. EBioMedicine 2015; 2: 1996–2002. Samokhvalov AV, Irving H, Mohapatra S, Rehm J. Alcohol consumption, unprovoked seizures, and epilepsy:
Alcohol use	Epilepsy	a systematic review and meta-analysis. Epilepsia 2010; 51: 1177–84. Dworetzky BA, Bromfield EB, Townsend MK, Kang JH. A prospective study of smoking, caffeine, and
Alcohol use	Epilepsy	alcohol as risk factors for seizures or epilepsy in young adult women: data from the Nurses' Health Study II. Epilepsia 2010; 51: 198–205.
Alcohol use	Diabetes mellitus type 2	Carlsson S, Hammar N, Grill V. Alcohol consumption and type 2 diabetes Meta-analysis of epidemiological studies indicates a Lisbaned relationship. Diabetologia 2005: 48: 1051–4
Alashalwas	Diabetes mellitus type 2	Li X-H, Yu F-F, Zhou Y-H, He J. Association between alcohol consumption and the risk of incident type 2 distance and the risk of incident type 2
Aconor use	Diabetes menitus type 2	Cherpitel CJ, Ye Y, Bond J, et al. Alcohol Attributable Fraction for Injury Morbidity from the Dose-Response
Alcohol use	Unintentional Injuries	Relationship of Acute Alcohol Consumption: Emergency Department Data from 18 Countries. Addiction 2015; 110: 1724–32.
Alcohol use	Unintentional Injuries	Corrao G, Bagnardi V, Zambon A, La Vecchia C. A meta-analysis of alcohol consumption and the risk of 15 diseases. Prev Med 2004; 38: 613–9.
		Taylor B, Irving HM, Kanteres F, et al. The more you drink, the harder you fall: a systematic review and meta- analysis of how acute alcohol consumption and injury or collision risk increase together. Drug Alcohol
Alcohol use	Unintentional Injuries	Depend 2010; 110: 108–16.

Appendix Table 5. Epidemic Citations and B. Additional A. Citations	ological evidence supporting causa information	lity between risk-outcome pairs included in the Global Burden of Disease 2017 study including A.
Risk	Outcome	Citation/Note Beghi M, Rosenbaum JF, Cerri C, Cornaggia CM. Risk factors for fatal and nonfatal repetition of suicide
Alcohol use	Self-harm	attempts: a literature review. Neuropsychiatr Dis Treat2013;9: 1725–36. Borges G, Bagge CL, Cherpitel CJ, Conner KR, Orozco R, Rossow I, A meta-analysis of acute use of alcohol
Alcohol use	Self-harm	and the risk of suicide attempt. Psychol Med 2017; 47: 949–57.
Alcohol use	Self-harm	Alcohol Clin Exp Res 2004; 28: 18S–28S.
Alcohol use	Self-harm	Cherpitel CJ, Ye Y, Bond J, et al. Alcohol Attributable Fraction for Injury Morbidity from the Dose-Response Relationship of Acute Alcohol Consumption: Emergency Department Data from 18 Countries. Addiction 2015; 110: 1724–32.
Alcohol use	Self-harm	Corrao G, Bagnardi V, Zambon A, La Vecchia C. A meta-analysis of alcohol consumption and the risk of 15 diseases. Prev Med 2004; 38: 613–9.
Alcoholuse	Self harm	Devries KM, Mak JY, Bacchus LJ, et al. Intimate partner violence and incident depressive symptoms and suicide attempts: a systematic review of longitudinal studies. PLoS Med 2013; 10: e1001/39
neonor use	Sen-harm	
Alcohol use	Self-harm	Haw C, Hawton K, Casey D, Bale E, Shepherd A. Alcohol dependence, excessive drinking and deliberate self- harm: trends and patterns in Oxford, 1989-2002. Soc Psychiatry Psychiatr Epidemiol 2005; 40: 964–71.
		Taylor B, Irving HM, Kanteres F, et al. The more you drink, the harder you fall: a systematic review and meta- analysis of how acute alcohol consumption and injury or collision risk increase together. Drug Alcohol
Alcohol use	Self-harm	Depend 2010; 110: 108–16. Cherpitel CJ, Ye Y, Bond J, et al. Alcohol Attributable Fraction for Injury Morbidity from the Dose-Response
Alcohol use	Interpersonal violence	Relationship of Acute Alcohol Consumption: Emergency Department Data from 18 Countries. Addiction 2015: 110: 1724–32.
		Taylor B, Irving HM, Kanteres F, et al. The more you drink, the harder you fall: a systematic review and meta-
Alcohol use	Interpersonal violence	analysis of how acute alcohol consumption and injury or collision risk increase together. Drug Alcohol Depend 2010; 110: 108–16.
Alcohol use	Ischaemic and haemorrhagic stroke	Patra J, Taylor B, Irving H, et al. Alcohol consumption and the risk of morbidity and mortality for different stroke typesa systematic review and meta-analysis. BMC Public Health 2010; 10: 258.
Alcohol use	Ischaemic and haemorrhagic stroke	Ronksley PE, Brien SE, Turner BJ, Mukamal KJ, Ghali WA. Association of alcohol consumption with selected cardiovascular disease outcomes: a systematic review and meta-analysis. BMJ 2011; 342: d671.
Alcohol use	Ischaemic and haemorrhagic	Bazzano LA, Gu D, Reynolds K, et al. Alcohol consumption and risk for stroke among Chinese men. Ann Neurol 2007; 62: 569–78
	Ischaemic and haemorrhagic	Berger K, Ajani UA, Kase CS, et al. Light-to-moderate alcohol consumption and the risk of stroke among
Alcohol use	stroke Ischaemic and haemorrhagic	U.S. male physicians. N Engl J Med 1999; 341: 1557–64. Chiuve SE, Rexrode KM, Spiegelman D, Logroscino G, Manson JE, Rimm EB. Primary prevention of stroke
Alcohol use	stroke Ischaemic and haemorrhagic	by healthy lifestyle. Circulation 2008; 118: 947–54. Djoussé L, Ellison RC, Beiser A, Scaramucci A, D'Agostino RB, Wolf PA. Alcohol consumption and risk of
Alcohol use	stroke	ischemic stroke: The Framingham Study. Stroke 2002; 33: 907–12.
Alcohol use	stroke	Program. JAMA 1986; 255: 2311–4.
Alcohol use	Ischaemic and haemorrhagic stroke	Elkind MSV, Sciacca R, Boden-Albala B, Rundek T, Paik MC, Sacco RL. Moderate alcohol consumption reduces risk of ischemic stroke: the Northern Manhattan Study. Stroke 2006; 37: 13–9.
Alcohol use	Ischaemic and haemorrhagic stroke	Hansagi H, Romelsjö A, Gerhardsson de Verdier M, Andréasson S, Leifman A. Alcohol consumption and stroke mortality. 20-year follow-up of 15,077 men and women. Stroke 1995; 26: 1768–73.
Alcohol use	Ischaemic and haemorrhagic stroke	Higashiyama A, Wakabayashi I, Ono Y, et al. Association with serum gamma-glutamyltransferase levels and alcohol consumption on stroke and coronary artery disease: the Suita study. Stroke 2011; 42: 1764–7.
Alcohol use	Ischaemic and haemorrhagic	Ikehara S, Iso H, Toyoshima H, et al. Alcohol consumption and mortality from stroke and coronary heart disease among Japanese men and women: the Japan collaborative cohort study. Stroke 2008; 39: 2936-42
	Ischaemic and haemorrhagic	Ikehara S, Iso H, Yamagishi K, et al. Alcohol consumption and risk of stroke and coronary heart disease among Japanese women: the Japan Public Health Center-based prospective study. Prev Med 2013; 57:
Alcohol use	stroke Ischaemic and haemorrhagic	505–10. Ikehara S, Iso H, Yamagishi K, et al. Alcohol consumption, social support, and risk of stroke and coronary
Alcohol use	stroke Ischaemic and haemorrhagic	heart disease among Japanese men: the JPHC Study. Alcohol Clin Exp Res 2009; 33: 1025–32. Iso H. Kitamura A. Shimamoto T. et al. Alcohol intake and the risk of cardiovascular disease in middle-aged
Alcohol use	stroke	Japanese men. Stroke 1995; 26: 767–73.
Alcohol use	stroke	Atherosclerosis Risk in Communities Study. Stroke 2015; 46: 3124–30.
Alcohol use	Ischaemic and haemorrhagic stroke	Klatsky AL, Armstrong MA, Friedman GD, Sidney S. Alcohol drinking and risk of hemorrhagic stroke. Neuroepidemiology 2002; 21: 115–22.
Alcohol use	Ischaemic and haemorrhagic stroke	Klatsky AL, Armstrong MA, Friedman GD, Sidney S. Alcohol drinking and risk of hospitalization for ischemic stroke. Am J Cardiol 2001; 88: 703–6.
Alcohol use	Ischaemic and haemorrhagic stroke	Klatsky AL, Armstrong MA, Friedman GD. Alcohol use and subsequent cerebrovascular disease hospitalizations. Stroke 1989; 20: 741–6.
Alcoholuse	Ischaemic and haemorrhagic	Kono S, Ikeda M, Tokudome S, Nishizumi M, Kuratsune M. Alcohol and mortality: a cohort study of male Japanese physicians. Int J Enidemiol 1986: 15: 527–32
Alashalas	Ischaemic and haemorrhagic	Leppäla JM, Paunio M, Virtamo J, et al. Alcohol consumption and stroke incidence in male smokers.
Alcohol use	Ischaemic and haemorrhagic stroke	Mukamal KJ, Ascherio A, Mittleman MA, et al. Alcohol and risk for ischemic stroke in men: the role of drinking patterns and usual beverage. Ann Intern Med 2005; 142: 11–9.

Appendix Table 5. Epide Citations and B. Additio	emiological evidence supporting caus onal information	ality between risk-outcome pairs included in the Global Burden of Disease 2017 study including A.
Risk	Outcome	Citation/Note
Alcohol use	Ischaemic and haemorrhagic stroke	Mukamal KJ, Chung H, Jenny NS, et al. Alcohol use and risk of ischemic stroke among older adults: the cardiovascular health study. Stroke 2005; 36: 1830–4.
Alcohol use	Ischaemic and haemorrhagic stroke	Nielsen NR, Truelsen T, Barefoot JC, et al. Is the effect of alcohol on risk of stroke confined to highly stressed persons? Neuroepidemiology 2005; 25: 105–13.
Alcohol use	Ischaemic and haemorrhagic stroke	Sankai T, Iso H, Shimamoto T, et al. Prospective study on alcohol intake and risk of subarachnoid hemorrhage among Japanese men and women. Alcohol Clin Exp Res 2000; 24: 386–9.
Alcohol use	Ischaemic and haemorrhagic stroke	Stampfer MJ, Colditz GA, Willett WC, Speizer FE, Hennekens CH. A prospective study of moderate alcohol consumption and the risk of coronary disease and stroke in women. N Engl J Med 1988; 319: 267–73.
Alcohol use	stroke Ischaemic and haemorrhagic	men: Korea Medical Insurance Corporation Study. Lancet 2001; 357: 922–5. Yamada S. Kojzumi A. Joo H. et al. Risk factors for fatal subarachooid hemorrhage: the Japan Collaborative
Alcohol use	stroke	Cohort Study. Stroke 2003; 34: 2781–7. Blomé MA Biörkman P. Flambolc L. Jacobsson H. Molnegren V. Widell A. Minimal transmission of HIV.
Drug use	Hepatitis B	despite persistently high transmission of hepatitis C virus in a Swedish needle exchange program. J Viral Hepat 2011; 18: 831–9.
Drug use	Hepatitis B	Crofts N, Aitken CK. Incidence of bloodborne virus infection and risk behaviours in a cohort of injecting drug users in Victoria, 1990-1995. Med J Aust 1997; 167: 17–20.
Drug use	Hepatitis B	Hagan H, McGough JP, Thiede H, Weiss NS, Hopkins S, Alexander ER. Syringe Exchange and Risk of Infection with Hepatitis B and C Viruses, Am. J. Epi. 1999; 149(3): 203–213.
Drug use	Hepatitis B	Jackson JB, Wei L, Liping F, et al. Prevalence and seroincidence of hepatitis B and hepatitis C infection in high risk people who inject drugs in china and Thailand. Hepat Res Treat 2014; 2014: 296958.
Drug use	Hepatitis B	Månsson AS, Moestrup T, Nordenfelt E, Widell A. Continued transmission of hepatitis B and C viruses, but no transmission of human immunodeficiency virus among intravenous drug users participating in a syringe/needle exchange program. Scand J Infect Dis 2000; 32: 253–8.
Drug use	Hepatitis C	Abou-Saleh M, Davis P, Rice P, et al. The effectiveness of behavioural interventions in the primary prevention of hepatitis C amongst injecting drug users: a randomised controlled trial and lessons learned. Harm Reduct J 2008; 5: 25.
Drug use	Hepatitis C	Blomé MA, Björkman P, Flamholc L, Jacobsson H, Molnegren V, Widell A. Minimal transmission of HIV despite persistently high transmission of hepatitis C virus in a Swedish needle exchange program. J Viral Hepat 2011; 18: 831–9.
Drug use	Hepatitis C	Craine N, Hickman M, Parry JV, et al. Incidence of hepatitis C in drug injectors: the role of homelessness, opiate substitution treatment, equipment sharing, and community size. Epidemiol Infect 2009; 137: 1255–65.
Drug use	Hepatitis C	Crofts N, Aitken CK. Incidence of bloodborne virus infection and risk behaviours in a cohort of injecting drug users in Victoria, 1990-1995. Med J Aust 1997; 167: 17–20.
Drug use	Hepatitis C	Foley SB, Abou-Saleh MT. Risk Behaviors and Transmission of Hepatitis C in Injecting Drug Users. Addictive Disorders & Their Treatment 2009; 8: 13–21.
Drug use	Hepatitis C	Grebely J, Lima VD, Marshall BDL, et al. Declining incidence of hepatitis C virus infection among people who inject drugs in a Canadian setting, 1996-2012. PLoS ONE 2014; 9: e97726. Hagan H, McGough JP, Thiede H, Weiss NS, Honkins S, Alexander ER, Svringe exchange and risk of
Drug use	Hepatitis C	infection with hepatitis B and C viruses. Am J Epidemiol 1999; 149: 203–13. Lackson IB Wai L Linning E et al. Prevalence and serving incidence of hepatitis B and hepatitis C infection in
Drug use	Hepatitis C	high risk people who inject drugs in china and Thailand. Hepat Res Treat 2014; 2014: 296958.
Drug use	Hepatitis C	intravenous drug users in the North and East of France. Epidemiol Infect 2004; 132: 699–708.
Drug use	Hepatitis C	drug users in Australia. Addiction 2006; 101: 1499–508.
Drug use	Hepatitis C	Mansson AS, Moestrup I, Nordenfelt E, Wideli A. Continued transmission of nepatitis B and C viruses, but no transmission of human immunodeficiency virus among intravenous drug users participating in a syringe/needle exchange program. Scand J Infect Dis 2000; 32: 253–8.
Drug use	Hepatitis C	Partanen A, Malin K, Perälä R, Harju O, Holopainen A, Holmström P, et al. Riski-tutkimus 2000-2003. Pistämällä huumeita käyttävien seurantatutkimus. A-Klinikkasäätiön Raporttisarja nro 52. Helsinki: A- Klinikkasäätiön, 2006.
Drug use	Hepatitis C	Roy KM, Goldberg D, Taylor A, et al. A method to detect the incidence of hepatitis C infection among injecting drug users in Glasgow 1993-98. J Infect 2001; 43: 200–5.
Drug use	Hepatitis C	Turner KME, Hutchinson S, Vickerman P, et al. The impact of needle and syringe provision and opiate substitution therapy on the incidence of hepatitis C virus in injecting drug users: pooling of UK evidence. Addiction 2011; 106: 1978–88.
Drug use	Henatitis C	Van Den Berg C, Smit C, Van Brussel G, Coutinho R, Prins M, Amsterdam Cohort. Full participation in harm reduction programmes is associated with decreased risk for human immunodeficiency virus and hepatitis C virus: evidence from the Amsterdam Cohort Studies among drug users. Addiction 2007; 102: 1454–62
Drug uso	Hopatitis C	Villano SA, Vlahov D, Nelson KE, Lyles CM, Cohn S, Thomas DL. Incidence and risk factors for hepatitis C among injection days users in Paltimore Mandand J Clin Minschiel 1007, 25, 2274.7
Diet low in fruits	Lip and oral cavity cancer	Key TI Fruit and vegetables and cancer rick British Journal of Cancer 2011, 104, 6-11
Diet low in fruits	Lip and oral cavity cancer	Jin, Jian, Zhiguo Ouyang, and Zhaoyan Wang. 2014. "Association of Fruit and Vegetables with the Risk of Nasopharyngeal Cancer: Evidence from a Meta-Analysis." Scientific Reports4 (July): srep05229. doi:10.1038/srep05229

Risk	Outcome	Citation/Note
		Pavia, Maria, Claudia Pileggi, Carmelo GA Nobile, and Italo F. Angelillo. 2006. "Association between Fruit
		and Vegetable Consumption and Oral Cancer: A Meta-Analysis of Observational Studies." The American
Diet low in fruits	Lip and oral cavity cancer	Journal of Clinical Nutrition83 (5): 1126–34.
Diet low in fruits	Nasopharynx cancer	American Institute for Cancer Research and World Cancer Research Fund eds 2007 Food Nutrition
		Physical Activity and the Prevention of Cancer: A Global Perspective: A Project of World Cancer Research
Diet low in fruits	Nasopharynx cancer	Fund International. Washington, D.C: American Institute for Cancer Research.
		Pavia, Maria, Claudia Pileggi, Carmelo GA Nobile, and Italo F. Angelillo. 2006. "Association between Fruit
D' 1 ' C '		and Vegetable Consumption and Oral Cancer: A Meta-Analysis of Observational Studies." The American
Diet low in fruits	Nasopharynx cancer	Journal of Clinical Nutrition83 (5): 1126–34.
Diet low in fruits	Other pharyinx cancer	American Institute for Cancer Research and World Cancer Research Fund eds 2007 Food Nutrition
		Physical Activity and the Prevention of Cancer: A Global Perspective: A Project of World Cancer Research
Diet low in fruits	Other pharynx cancer	Fund International. Washington, D.C: American Institute for Cancer Research.
		Pavia, Maria, Claudia Pileggi, Carmelo GA Nobile, and Italo F. Angelillo. 2006. "Association between Fruit
D' (1) ()		and Vegetable Consumption and Oral Cancer: A Meta-Analysis of Observational Studies." The American
Diet low in fruits	Uther pharynx cancer	Journal of Clinical Nutrition83 (5): 1126–34.
Dict low in funs	Larynx cancer	American Institute for Cancer Research and World Cancer Research Fund eds 2007 Food Nutrition
		Physical Activity and the Prevention of Cancer: A Global Perspective: A Project of World Cancer Research
Diet low in fruits	Larynx cancer	Fund International. Washington, D.C: American Institute for Cancer Research.
		Pavia, Maria, Claudia Pileggi, Carmelo GA Nobile, and Italo F. Angelillo. 2006. "Association between Fruit
Dist laws in fastic	T	and Vegetable Consumption and Oral Cancer: A Meta-Analysis of Observational Studies." The American
Diet low in truits	Larynx cancer	Journal of Clinical Nutrition83 (5): 1120–54. Liu I. Wang I. Leng V. Ly C. Inteke of fruit and vagatables and risk of econhageal squamous cell carcinoma
Diet low in fruits	Oesophageal cancer	a meta-analysis of observational studies. Int J Cancer 2013; 133: 473–85.
	Tracheal, bronchus and lung	Vieira AR, Abar L, Vingeliene S, et al. Fruits, vegetables and lung cancer risk: a systematic review and meta
Diet low in fruits	cancer	analysis. Ann Oncol 2016; 27: 81–96.
		Wang X, Ouyang Y, Liu J, et al. Fruit and vegetable consumption and mortality from all causes,
Dist low in fruits	Ischaomic hoart disease	cardiovascular disease, and cancer: systematic review and dose-response meta-analysis of prospective cohort studies. PMI 2014: 340: g4400
Diet low in fruits	ischaenne neart disease	Hu D. Huang I. Wang Y. Zhang D. Ou Y. Fruits and vegetables consumption and risk of stroke: a meta-
Diet low in fruits	Ischaemic stroke	analysis of prospective cohort studies. Stroke 2014; 45: 1613–9.
		Hu D, Huang J, Wang Y, Zhang D, Qu Y. Fruits and vegetables consumption and risk of stroke: a meta-
Diet low in fruits	Hemorrhagic stroke	analysis of prospective cohort studies. Stroke 2014; 45: 1613–9.
Dist low in fruits	Disbatas mallitus	Li M, Fan Y, Zhang X, Hou W, Tang Z. Fruit and vegetable intake and risk of type 2 diabetes mellitus: meta
Diet low in fruits	Diabetes menitus	Liu J. Wang J. Leng Y. Ly C. Intake of fruit and vegetables and risk of esophageal squamous cell carcinoma
Diet low in vegetables	Oesophageal cancer	a meta-analysis of observational studies. Int J Cancer 2013; 133: 473–85.
		Wang X, Ouyang Y, Liu J, et al. Fruit and vegetable consumption and mortality from all causes,
		cardiovascular disease, and cancer: systematic review and dose-response meta-analysis of prospective cohort
Diet low in vegetables	Ischaemic heart disease	studies. BMJ 2014; 349: g4490.
Diet low in vegetables	Ischaemic stroke	nu D, Huang J, wang T, Zhang D, Qu T. Fruits and vegetables consumption and fisk of stroke: a meta- analysis of prospective cohort studies. Stroke 2014: 45: 1613–9
in the generics		Hu D, Huang J, Wang Y, Zhang D, Qu Y. Fruits and vegetables consumption and risk of stroke: a meta-
Diet low in vegetables	Hemorrhagic stroke	analysis of prospective cohort studies. Stroke 2014; 45: 1613-9.
		Afshin A, Micha R, Khatibzadeh S, Mozaffarian D. Consumption of nuts and legumes and risk of incident
Diat low in lagrees	Isahaamia haart disaasa	ischemic heart disease, stroke, and diabetes: a systematic review and meta-analysis. Am J Clin Nutr 2014;
Dict low in leguines	ischaemic neart uisease	Aune D. Norat T. Romundstad P. Vatten I.J. Whole grain and refined grain consumption and the risk of type
		2 diabetes: a systematic review and dose-response meta-analysis of cohort studies. Eur J Epidemiol 2013; 28
Diet low in whole grains	Diabetes mellitus	845–58.
		Aune D, Keum N, Giovannucci E, et al. Whole grain consumption and risk of cardiovascular disease, cancer
Dist law in 1, 1, 1	Tanka and the state	and all cause and cause specific mortality: systematic review and dose-response meta-analysis of prospective
Diet low in whole grains	Ischaemic heart disease	studies. BMJ 2010; 353: 12/10.
		ischemic heart disease stroke and diabetes: a systematic review and meta-analysis Am I Clin Nutr 2014:
Diet low in nuts and seeds	Ischaemic heart disease	100: 278–88.
		Afshin A, Micha R, Khatibzadeh S, Mozaffarian D. Consumption of nuts and legumes and risk of incident
	D	ischemic heart disease, stroke, and diabetes: a systematic review and meta-analysis. Am J Clin Nutr 2014;
Diet low in nuts and seeds	Diabetes mellitus	100: 278–88.
		world Cancer Research Fund, American Institute for Cancer Research, Imperial College London. WCRE/AICR Systematic Literature Review Continuous Undate Project Peport: The Associations between
Diet low in milk	Colon and rectum cancer	Food, Nutrition and Physical Activity and the Risk of Colorectal Cancer. Oct 2010.
		World Cancer Research Fund, American Institute for Cancer Research, Imperial College London.
		WCRF/AICR Systematic Literature Review Continuous Update Project Report: The Associations between
Diet high in red meat	Colon and rectum cancer	Food, Nutrition and Physical Activity and the Risk of Colorectal Cancer. Oct 2010.
Diet high in red meat	Diabetes mellitus	ran A, Sun Q, Bernstein AM, et al. Ked meat consumption and risk of type 2 diabetes: 3 cohorts of US adult and an undated meta-analysis. Am I Clin Nutr 2011: 94: 1088–96
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Appendix Table 5. Epidemiolo Citations and B. Additional in A. Citations	gical evidence supporting cause formation	ality between risk-outcome pairs included in the Global Burden of Disease 2017 study including A.
Risk	Outcome	Citation/Note World Cancer Research Fund, American Institute for Cancer Research, Imperial College London. WCRF/AICR Systematic Literature Review Continuous Update Project Report: The Associations between
Diet high in processed meat	Colon and rectum cancer	Food, Nutrition and Physical Activity and the Risk of Colorectal Cancer. Oct 2010. Micha R, Wallace SK, Mozaffarian D, Red and processed meat consumption and risk of incident coronary
Diet high in processed meat	Ischaemic heart disease	heart disease, stroke, and diabetes mellitus: a systematic review and meta-analysis. Circulation 2010; 121: 2271–83.
Diet high in processed meat	Diabetes mellitus	Pan A, Sun Q, Bernstein AM, et al. Red meat consumption and risk of type 2 diabetes: 3 cohorts of US adults and an updated meta-analysis. Am J Clin Nutr 2011: 94: 1088–96.
Diet high in sugar-sweetened beverages and high body-mass		Malik VS. Pan A. Willett WC. Hu FB. Sugar-sweetened beverages and weight gain in children and adults: a
index	n/a	systematic review and meta-analysis. Am J Clin Nutr 2013; 98: 1084–102. World Cancer Research Fund, American Institute for Cancer Research, Imperial College London
Diet low fibre	Colon and rectum cancer	WCRF/AICR Systematic Literature Review Continuous Update Project Report: The Associations between Food. Nutrition and Physical Activity and the Risk of Colorectal Cancer. Oct 2010.
Diet low fibre	Ischaemic heart disease	Threapleton DE, Greenwood DC, Evans CE, et al. Dietary fibre intake and risk of cardiovascular disease: systematic review and meta-analysis. BMJ (Clinical research ed) 2013: 347: f6879.
		World Cancer Research Fund, American Institute for Cancer Research, Imperial College London. WCRF/AICR Systematic Literature Review Continuous Undate Project Report: The Associations between
Diet low in calcium	Colon and rectum cancer	Food, Nutrition and Physical Activity and the Risk of Colorectal Cancer. Oct 2010. Chowdbury R. Stevens S. Gorman D. et al. Association between fish consumption long chain omega 3 fatty
Diet low in seafood omega-3 fats	Ischaemic heart disease	acids, and risk of cerebrovascular disease: systematic review and meta-analysis. BMJ (Clinical research ed) 2012; 345: e6698.
Diet low in polyunsaturated fats	Ischaemic heart disease	Farvid MS, Ding M, Pan A, et al. Dietary linoleic acid and risk of coronary heart disease: a systematic review and meta-analysis of prospective cohort studies. Circulation 2014; 130: 1568–78.
Diet low in polyunsaturated fats	Ischaemic heart disease	Mozaffarian D, Micha R, Wallace S. Effects on coronary heart disease of increasing polyunsaturated fat in place of saturated fat: a systematic review and meta-analysis of randomized controlled trials. PLoS Med 2010; 7: e1000252.
Diet high in trans fats	Ischaemic heart disease	Mozaffarian D, Clarke R. Quantitative effects on cardiovascular risk factors and coronary heart disease risk of replacing partially hydrogenated vegetable oils with other fats and oils. Eur J Clin Nutr. 2009; 63(Suppl 2): S22-33.
Diet high in trans fats	Ischaemic heart disease	http://www.bmj.com/content/bmj/suppl/2015/08/11/bmj.h3978.DC1/sour025275.ww2_default.pdf; pg. 44
Diet high in sodium and high systolic blood pressure	n/a	Aburto NJ, Ziolkovska A, Hooper L, Elliott P, Cappuccio FP, Meerpohl JJ. Effect of lower sodium intake on health: systematic review and meta-analyses. BMJ 2013; 346: f1326.
Diet high in sodium	Stomach cancer	World Cancer Research Fund, American Institute for Cancer Research. Food, Nutrition, Physical Activity, and the Prevention of Cancer: a Global Perspective. Washington DC: AICR, 2007.
Diet high in sodium	Stomach cancer	D'Elia, Lanfranco, Giovanni Rossi, Renato Ippolito, Francesco P. Cappuccio, and Pasquale Strazzullo. 2012. "Habitual Salt Intake and Risk of Gastric Cancer: A Meta-Analysis of Prospective Studies." Clinical Nutrition31 (4): 489–98. doi:10.1016/j.clnu.2012.01.003.
Childhood sexual abuse	Depressive disorders	Brown J, Cohen P, Johnson JG, Smailes EM. Childhood abuse and neglect: specificity of effects on adolescent and young adult depression and suicidality. J Am Acad Child Adolesc Psychiatry 1999; 38: 1490–6.
Childhood sexual abuse	Depressive disorders	Chapman, D.P., Whitfield, C.L., Felitti, V.J., Dube, S.R., Edwards, V.J. and Anda, R.F., 2004. Adverse childhood experiences and the risk of depressive disorders in adulthood. Journal of affective disorders, 82(2), pp.217-225.
Childhood sexual abuse	Depressive disorders	Cheasty, M., Clare, A.W. and Collins, C., 1998. Relation between sexual abuse in childhood and adult depression: case-control study. Bmj, 316(7126), pp.198-201.
Childhood sexual abuse	Depressive disorders	Dinwiddie S, Heath AC, Dunne MP, Bucholz KK, Madden PA, Slutske WS, Bierut LJ, Statham DB, Martin NG. Early sexual abuse and lifetime psychopathology: a co-twin-control study. Psychol Med. 2000; 30(1): 41–52.
Childhood sexual abuse	Depressive disorders	Dube, S.R., Anda, R.F., Whitfield, C.L., Brown, D.W., Felitti, V.J., Dong, M. and Giles, W.H., 2005. Long-term consequences of childhood sexual abuse by gender of victim. American journal of preventive medicine, 28(5), pp.430-438.
Childhood sexual abuse	Depressive disorders	Ernst C, Angst J, Földényi M. The Zurich Study. XVII. Sexual abuse in childhood. Frequency and relevance for adult morbidity data of a longitudinal epidemiological study. Eur Arch Psychiatry Clin Neurosci. 1993; 242(5): 293–300.
Childhood sexual abuse	Depressive disorders	Jaffee SR, Moffitt TE, Caspi A, Fombonne E, Poulton R, Martin J. Differences in early childhood risk factors for juvenile-onset and adult-onset depression. Arch Gen Psychiatry. 2002; 59(3): 215-22.
Childhead on the	Deserving l'auto	Kendler KS, Bulik CM, Silberg J, Hettema JM, Myers J, Prescott CA. Childhood sexual abuse and adult psychiatric and substance use disorders in women: an epidemiological and cotwin control analysis. Arch Gen
	Depressive disorders	Molnar, B.E., Buka, S.L. and Kessler, R.C., 2001. Child sexual abuse and subsequent psychopathology:
Childhood sexual abuse	Depressive disorders	results from the National Comorbidity Survey. American journal of public health, 91(5), p.753. Nelson EC, Heath AC, Madden PA, Cooper ML, Dinwiddie SH, Bucholz KK, Glowinski A, McLaughlin T, Dunne MP, Statham DJ, Martin NG, Association between self-reported sexual abuse and adverse
Childhood sexual abuse	Depressive disorders	psychosocial outcomes: results from a twin study. Arch Gen Psychiatry. 2002; 59(2): 139-45. Peleikis D.F. Mykletun A and Dahl A Δ 2004 The relative influence of childhood eavual abure and
Childhood sexual abuse	Depressive disorders	other family background risk factors on adult adversities in female outpatients treated for anxiety disorders and depression Child Abuse & Neelect 28(1) nn 61-76
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Appendix Table 5. Epidem Citations and B. Additiona	iiological evidence supporting c al information	ausality between risk-outcome pairs included in the Global Burden of Disease 2017 study including A.
Risk	Outcome	Citation/Note Silverman, A.B., Reinherz, H.Z. and Giaconia, R.M., 1996, The long-term sequelae of child and adolescent
Childhood sexual abuse	Depressive disorders	abuse: A longitudinal community study. Child abuse & neglect, 20(8), pp.709-723.
		Widom, C.S., DuMont, K. and Czaja, S.J., 2007. A prospective investigation of major depressive disorder
Childhood sexual abuse	Depressive disorders	and comorbidity in abused and neglected children grown up. Archives of general psychiatry, 64(1), pp.49-56.
		NG. Early sexual abuse and lifetime psychopathology: a co-twin-control study. Psychol Med. 2000; 30(1):
Childhood sexual abuse	Alcohol use disorders	41-52. Dube S.D. Ande D.E. Whitfield C.L. Brown D.W. Felitti V.L. Dong M. and Ciles W.H. 2005 Long
		term consequences of childhood sexual abuse by gender of victim. American journal of preventive medicine,
Childhood sexual abuse	Alcohol use disorders	28(5), pp.430-438.
		Fleming, J., Mullen, P.E., Sibthorpe, B., Attewell, R. and Bammer, G., 1998. The relationship between
Childhood sexual abuse	Alcohol use disorders	childhood sexual abuse and alcohol abuse in women-a case-control study. Addiction, 93(12), pp.1787-1798. Kendler KS, Bulik CM, Silberg I, Hettema IM, Myers I, Prescott CA, Childhood sexual abuse and adult
		psychiatric and substance use disorders in women: an epidemiological and cotwin control analysis. Arch Gen
Childhood sexual abuse	Alcohol use disorders	Psychiatry. 2000; 57(10): 953–9. Molnar, B.E., Buka, S.L. and Kessler, R.C., 2001. Child sexual abuse and subsequent psychopathology:
Childhood sexual abuse	Alcohol use disorders	results from the National Comorbidity Survey. American journal of public health, 91(5), p.753.
		Nelson EC, Heath AC, Madden PA, Cooper ML, Dinwiddle SH, Bucholz KK, Glowinski A, McLaughlin T, Dunne MP, Statham DJ, Martin NG. Association between self-reported sexual abuse and adverse
Childhood sexual abuse	Alcohol use disorders	psychosocial outcomes: results from a twin study. Arch Gen Psychiatry. 2002; 59(2): 139-45.
		Sartor CE, Lynskey M1, Bucholz KK, McCutcheon VV, Nelson EC, Waldron M, Heath AC. Childhood sexual abuse and the course of alcohol dependence development: findings from a female twin sample. Drug
Childhood sexual abuse	Alcohol use disorders	Alcohol Depend. 2007; 89(2-3): 139–44. Silvarman A. P. Painherz, H.Z. and Giaconia, P.M. 1006. The long term sequeles of child and adolescent
Childhood sexual abuse	Alcohol use disorders	abuse: A longitudinal community study. Child abuse & neglect, 20(8), pp.709-723.
		Bowes L. Joinson C. Wolke D. Lewis G. Peer victimisation during adolescence and its impact on depression
Bullying victimization	Depressive disorders	in early adulthood: prospective cohort study in the United Kingdom. <i>BMJ</i> . 2015; 350: h2469.
		Fahy AE, Stansfeld SA, Smuk M, Smith NR, Cummins S, Clark C. Longitudinal Associations Between
Bullying victimization	Depressive disorders	Cyberbullying Involvement and Adolescent Mental Health. <i>J Adolesc Health</i> . 2016; 59(5): 502–9.
		Farrington DP, Loeber R, Stallings R, Ttoh MM. Bullying perpetration and victimization as predictors of delinquency and depression in the Pittsburgh Youth Study. <i>Journal of Aggression Conflict and Peace</i>
Bullying victimization	Depressive disorders	Research. 2011; 3(2): 74–81.
		children get bullied? A prospective cohort study on the relationship between bullying and health-related
Bullying victimization	Depressive disorders	symptoms. <i>Pediatrics</i> . 2006; 117(5): 1568–74.
		Gibb SJ, Horwood LJ, Fergusson DM. Bullying victimization/perpetration in childhood and later adjustment:
Bullying victimization	Depressive disorders	findings from a 30 year longitudinal study. <i>Jnl Aggress Conflict Peace Res</i> . 2011; 3(2): 82–8. Hemphill SA, Kotevski A, Heerde JA, Longitudinal associations between cyber-bullying perpetration and
		victimization and problem behavior and mental health problems in young Australians. <i>Int J Public</i>
Bullying victimization	Depressive disorders	Health. 2015; 60(2): 227–37. Hemphill SA, Kotevski A, Herrenkohl TI, Bond L, Kim MJ, Toumbourou JW, Catalano RF. Longitudinal
		consequences of adolescent bullying perpetration and victimisation: A study of students in Victoria, Australia.
Bullying vicumization	Depressive disorders	Kaltiala-Heino R, Fröjd S, Marttunen M. Involvement in bullying and depression in a 2-year follow-up in
Bullying victimization	Depressive disorders	middle adolescence. <i>Eur Child Adolesc Psychiatry</i> . 2010; 19(1): 45–55.
		symptoms and deviance in adolescence: An epidemiological sample. <i>Child Abuse Negl</i> . 2000; 24(12):
Bullying victimization	Depressive disorders	1567–77. Lereva ST Copeland WE Zammit S Wolke D Bully/victims: a longitudinal_population-based cohort study
Bullying victimization	Depressive disorders	of their mental health. <i>Eur Child Adolesc Psychiatry</i> . 2015; 24(12): 1461–71.
		Moore SE, Norman RE, Sly PD, Whitehouse AJO, Zubrick SR, Scott J. Adolescent peer aggression and its association with mental health and substance use in an Australian cohort. <i>J Adolesc</i> , 2014: 37(1):
Bullying victimization	Depressive disorders	11–21.
		Patton GC, Olsson C, Bond L, Toumbourou JW, Carlin JB, Hemphill SA, Catalano RF. Predicting female depression across puberty: a two-nation longitudinal study. <i>J Am Acad Child Adolesc Psychiatry</i> .
Bullying victimization	Depressive disorders	2008; 47(12): 1424–32.
		Kotnon C, Head J, Klineberg E, Stansteld S. Can social support protect bullied adolescents from adverse outcomes? A prospective study on the effects of bullying on the educational achievement and mental health of
Bullying victimization	Depressive disorders	adolescents at secondary schools in East London. <i>J Adolesc</i> . 2011; 34(3): 579–88.
		a bully in adolescence on externalizing and internalizing mental health problems in adulthood. <i>Child</i>
Bullying victimization	Depressive disorders	Adolesc Psychiatry Ment Health
Dullying visting in the	Dopposition diagonal	Silberg JL, Copeland W, Linker J, Moore AA, Roberson-Nay R, York TP. Psychiatric outcomes of bullying

Risk	Outcome	Citation/Note
Bullying victimization	Depressive disorders	Sourander A, Jensen P, Rönning JA, Niemelä S, Helenius H, Sillanmäki L, Kumpulainen K, Piha J, Tamminen T, Moilanen I, Almqvist F. What is the early adulthood outcome of boys who bully or are bullied in childhood? The Finnish "From a Boy to a Man" study. <i>Pediatrics</i> . 2007; 120(2): 397–404.
Bullying victimization	Depressive disorders	Vassallo S, Edwards B, Renda J, Olsson CA. Bullying in Early Adolescence and Antisocial Behavior and Depression Six Years Later: What Are the Protective Factors?. <i>J Sch Violence</i> . 2014; 13(1): 100–24.
Bullying victimization	Depressive disorders	Wichstrøm L, Belsky J, Berg-Nielsen TS. Preschool predictors of childhood anxiety disorders: a prospective community study. <i>Journal of Child Psychology and Psychiatry</i> . 2013; 54(12): 1327–36.
Bullying victimization	Depressive disorders	Zwierzynska K, Wolke D, Lereya TS. Peer victimization in childhood and internalizing problems in adolescence: a prospective longitudinal study. <i>J Abnorm Child Psychol</i> . 2013; 41(2): 309–23.
Bullying victimization	Anxiety disorders	Bowes L, Joinson C, Wolke D, Lewis G. Peer victimisation during adolescence and its impact on depression in early adulthood: prospective cohort study in the United Kingdom. <i>BMJ</i> . 2015; 350: h2469.
Bullying victimization	Anxiety disorders	Fahy AE, Stansfeld SA, Smuk M, Smith NR, Cummins S, Clark C. Longitudinal Associations Between Cyberbullying Involvement and Adolescent Mental Health. <i>J Adolesc Health</i> . 2016; 59(5): 502–9.
Bullying victimization	Anxiety disorders	Farrington DP, Loeber R, Stallings R, Ttofi MM. Bullying perpetration and victimization as predictors of delinquency and depression in the Pittsburgh Youth Study. <i>Journal of Aggression Conflict and Peace Research</i> . 2011; 3(2): 74–81.
Bullying victimization	Anxiety disorders	Fekkes M, Pijpers FIM, Fredriks AM, Vogels T, Verloove-Vanhorick SP. Do bullied children get ill, or do il children get bullied? A prospective cohort study on the relationship between bullying and health-related symptoms. <i>Pediatrics</i> . 2006; 117(5): 1568–74.
Bullying victimization	Anxiety disorders	Gibb SJ, Horwood LJ, Fergusson DM. Bullying victimization/perpetration in childhood and later adjustment findings from a 30 year longitudinal study. <i>Jnl Aggress Conflict Peace Res</i> . 2011; 3(2): 82–8.
Bullying victimization	Anxiety disorders	Hemphill SA, Kotevski A, Heerde JA. Longitudinal associations between cyber-bullying perpetration and victimization and problem behavior and mental health problems in young Australians. <i>Int J Public Health</i> . 2015; 60(2): 227–37.
Bullying victimization	Anxiety disorders	Hemphill SA, Kotevski A, Herrenkohl TI, Bond L, Kim MJ, Toumbourou JW, Catalano RF. Longitudinal consequences of adolescent bullying perpetration and victimisation: A study of students in Victoria, Australia <i>Crim Behav Ment Health</i> . 2011; 21(2): 107–16.
Bullying victimization	Anxiety disorders	Kaltiala-Heino R, Fröjd S, Marttunen M. Involvement in bullying and depression in a 2-year follow-up in middle adolescence. <i>Eur Child Adolesc Psychiatry</i> . 2010; 19(1): 45–55.
Bullying victimization	Anxiety disorders	Kumpulainen K, Räsänen E. Children involved in bullying at elementary school age: their psychiatric symptoms and deviance in adolescence: An epidemiological sample. <i>Child Abuse Negl</i> . 2000; 24(12) 1567–77.
Bullying victimization	Anxiety disorders	Lereya ST, Copeland WE, Zammit S, Wolke D. Bully/victims: a longitudinal, population-based cohort study of their mental health. <i>Eur Child Adolesc Psychiatry</i> . 2015; 24(12): 1461–71.
		Moore SE, Norman RE, Sly PD, Whitehouse AJO, Zubrick SR, Scott J. Adolescent peer aggression and its association with mental health and substance use in an Australian cohort. <i>J Adolesc</i> . 2014; 37(1):
Bunying vicunitzation	Anxiety disorders	Patton GC, Olsson C, Bond L, Toumbourou JW, Carlin JB, Hemphill SA, Catalano RF. Predicting female depression across puberty: a two-nation longitudinal study. <i>J Am Acad Child Adolesc Psychiatry</i> .
Bullying victimization	Anxiety disorders	2008; 47(12): 1424–32. Rothon C, Head J, Klineberg E, Stansfeld S. Can social support protect bullied adolescents from adverse outcomes? A prospective study on the effects of bullying on the educational achievement and mental health of
Bullying victimization	Anxiety disorders	adolescents at secondary schools in East London. <i>J Adolesc</i> . 2011; 34(3): 579–88. Sigurdson JF, Undheim AM, Wallander JL, Lydersen S, Sund AM. The long-term effects of being bullied or
Bullying victimization	Anxiety disorders	a bully in adolescence on externalizing and internalizing mental health problems in adulthood. <i>Child Adolesc Psychiatry Ment Health</i> . 2015; 9(1): 1-13.
Bullying victimization	Anxiety disorders	Silberg JL, Copeland W, Linker J, Moore AA, Roberson-Nay R, York TP. Psychiatric outcomes of bullying victimization: a study of discordant monozygotic twins. <i>Psychol Med</i> . 2016; 46(9): 1875–83.
Bullying victimization	Anxiety disorders	Sourander A, Jensen P, Rönning JA, Niemelä S, Helenius H, Sillanmäki L, Kumpulainen K, Piha J, Tamminen T, Moilanen I, Almqvist F. What is the early adulthood outcome of boys who bully or are bullied in childhood? The Finnish "From a Boy to a Man" study. <i>Pediatrics</i> . 2007; 120(2): 397–404.
Bullying victimization	Anxiety disorders	Vassallo S, Edwards B, Renda J, Olsson CA. Bullying in Early Adolescence and Antisocial Behavior and Depression Six Years Later: What Are the Protective Factors?. <i>J Sch Violence</i> . 2014; 13(1): 100–24.
Bullying victimization	Anxiety disorders	Wichstrøm L, Belsky J, Berg-Nielsen TS. Preschool predictors of childhood anxiety disorders: a prospective community study. <i>Journal of Child Psychology and Psychiatry</i> . 2013; 54(12): 1327–36.
Bullying victimization	Anxiety disorders	Zwierzynska K, Wolke D, Lereya TS. Peer victimization in childhood and internalizing problems in adolescence: a prospective longitudinal study. <i>J Abnorm Child Psychol</i> . 2013; 41(2): 309–23.
Intimate partner violence	HIV/AIDS	Jewkes RK, Dunkle K, Nduna M, Shai N. Intimate partner violence, relationship power inequity, and incidence of HIV infection in young women in South Africa: a cohort study. Lancet 2010; 376: 41–8.

Appendix Table 5. Epidemiological evidence supporting causality between risk-outcome pairs included in the Global Burden of Disease 2017 study including A. **Citations and B. Additional information** A. Citations Outcome Citation/Note Risk Kouyoumdjian FG, Calzavara LM, Bondy SJ, et al. Intimate partner violence is associated with incident HIV HIV/AIDS Intimate partner violence infection in women in Uganda. AIDS 2013; 27: 1331-8. Chowdhary N, Patel V. The effect of spousal violence on women's health: findings from the Stree Arogya Depressive disorders Shodh in Goa, India. J Postgrad Med. 2008; 54(4): 306-12. Intimate partner violence Lipsky S, Caetano R, Roy-Byrne P. Racial and ethnic disparities in police-reported intimate partner violence Intimate partner violence Depressive disorders and risk of hospitalization among women. Womens Health Issues. 2009; 19(2):109-118. Ouellet-Morin I, Fisher HL, York-Smith M, Fincham-Campbell S, Moffitt TE, Arseneault L. Intimate partner violence and new-onset depression: a longitudinal study of women's childhood and adult histories of abuse. Intimate partner violence Depressive disorders Depression and anxiety. 2015;32(5):316-324. Suglia SF, Duarte CS, Sandel MT. Housing quality, housing instability, and maternal mental health. J Urban Health. 2011; 88(6): 1105-16. Intimate partner violence Depressive disorders Bourassa D, Bérubé J. The prevalence of intimate partner violence among women and teenagers seeking Intimate partner violence Maternal abortion and miscarriage abortion compared with those continuing pregnancy. J Obstet Gynaecol Can 2007; 29: 415-23. Leung TW, Leung WC, Chan PL, Ho PC. A comparison of the prevalence of domestic violence between patients seeking termination of pregnancy and other general gynecology patients. Int J Gynaecol Obstet 2002; Intimate partner violence Maternal abortion and miscarriage 77: 47-54. Romito P, Escribà-Agüir V, Pomicino L, Lucchetta C, Scrimin F, Molzan Turan J. Violence in the lives of Maternal abortion and miscarriage women in Italy who have an elective abortion. Womens Health Issues 2009; 19: 335-43. Intimate partner violence Taft AJ, Watson LF. Termination of pregnancy: associations with partner violence and other factors in a Intimate partner violence Maternal abortion and miscarriage national cohort of young Australian women. Aust N Z J Public Health 2007; 31: 135-42. Bostick RM, Potter JD, Kushi LH, et al. Sugar, meat, and fat intake, and non-dietary risk factors for colon Low physical activity Colon and rectum cancer cancer incidence in Iowa women (United States). Cancer Causes Control 1994; 5: 38-52. Calton BA, Lacey JV, Schatzkin A, et al. Physical activity and the risk of colon cancer among women: a prospective cohort study (United States). Int J Cancer 2006; 119: 385-91. Low physical activity Colon and rectum cancer Chao A, Connell CJ, Jacobs EJ, et al. Amount, type, and timing of recreational physical activity in relation to colon and rectal cancer in older adults: the Cancer Prevention Study II Nutrition Cohort. Cancer Epidemiol Low physical activity Colon and rectum cancer Biomarkers Prev 2004; 13: 2187-95. Colbert LH, Hartman TJ, Malila N, et al. Physical activity in relation to cancer of the colon and rectum in a Low physical activity Colon and rectum cancer cohort of male smokers. Cancer Epidemiol Biomarkers Prev 2001; 10: 265-8. Fraser G, Pearce N. Occupational physical activity and risk of cancer of the colon and rectum in New Zealand Low physical activity Colon and rectum cancer males. Cancer Causes Control 1993; 4: 45-50. Friedenreich C, Norat T, Steindorf K, et al. Physical activity and risk of colon and rectal cancers: the European prospective investigation into cancer and nutrition. Cancer Epidemiol Biomarkers Prev 2006; 15: Low physical activity Colon and rectum cancer 2398-407. Garabrant DH, Peters JM, Mack TM, Bernstein L. Job activity and colon cancer risk. Am J Epidemiol 1984; Low physical activity Colon and rectum cancer 119: 1005-14. Gerhardsson M, Norell SE, Kiviranta H, Pedersen NL, Ahlbom A. Sedentary jobs and colon cancer. Am J Low physical activity Colon and rectum cancer Epidemiol 1986; 123: 775-80. Giovannucci E, Ascherio A, Rimm EB, Colditz GA, Stampfer MJ, Willett WC. Physical activity, obesity, and Low physical activity Colon and rectum cancer risk for colon cancer and adenoma in men. Ann Intern Med 1995; 122: 327-34. Howard RA, Freedman DM, Park Y, Hollenbeck A, Schatzkin A, Leitzmann MF. Physical activity, sedentary behavior, and the risk of colon and rectal cancer in the NIH-AARP Diet and Health Study. Cancer Causes Low physical activity Colon and rectum cancer Control 2008; 19: 939-53. Larsson SC, Rutegård J, Bergkvist L, Wolk A. Physical activity, obesity, and risk of colon and rectal cancer in Low physical activity Colon and rectum cancer a cohort of Swedish men. Eur J Cancer 2006; 42: 2590-7. Lee IM, Manson JE, Ajani U, Paffenbarger RS, Hennekens CH, Buring JE. Physical activity and risk of colon Colon and rectum cancer Low physical activity cancer: the Physicians' Health Study (United States). Cancer Causes Control 1997; 8: 568-74. Lee IM, Paffenbarger RS. Physical activity and its relation to cancer risk: a prospective study of college Low physical activity Colon and rectum cancer alumni. Med Sci Sports Exerc 1994; 26: 831-7. Lee K-J, Inoue M, Otani T, et al. Physical activity and risk of colorectal cancer in Japanese men and women: Low physical activity Colon and rectum cancer the Japan Public Health Center-based prospective study. Cancer Causes Control 2007; 18: 199-209. Mai PL, Sullivan-Halley J, Ursin G, et al. Physical activity and colon cancer risk among women in the California Teachers Study. Cancer Epidemiol Biomarkers Prev 2007; 16: 517-25. Low physical activity Colon and rectum cancer Moradi T, Gridley G, Björk J, et al. Occupational physical activity and risk for cancer of the colon and rectum Low physical activity Colon and rectum cancer in Sweden among men and women by anatomic subsite. Eur J Cancer Prev 2008; 17: 201-8. Nilsen TIL, Romundstad PR, Petersen H, Gunnell D, Vatten LJ. Recreational physical activity and cancer risk in subsites of the colon (the Nord-Trøndelag Health Study). Cancer Epidemiol Biomarkers Prev 2008; 17: Low physical activity Colon and rectum cancer 183-8. Severson RK, Nomura AM, Grove JS, Stemmermann GN. A prospective analysis of physical activity and Low physical activity Colon and rectum cancer cancer. Am J Epidemiol 1989; 130: 522-9. Thune I, Lund E. Physical activity and risk of colorectal cancer in men and women. Br J Cancer 1996; 73: Colon and rectum cancer Low physical activity 1134-40. Wolin KY, Lee I-M, Colditz GA, Glynn RJ, Fuchs C, Giovannucci E. Leisure-time physical activity patterns Low physical activity Colon and rectum cancer and risk of colon cancer in women. Int J Cancer 2007; 121: 2776-81. Bardia A, Hartmann LC, Vachon CM, et al. Recreational physical activity and risk of postmenopausal breast Low physical activity Breast cancer cancer based on hormone receptor status. Arch Intern Med 2006; 166: 2478-83. Borch KB, Lund E, Braaten T, Weiderpass E. Physical activity and the risk of postmenopausal breast cancer -Low physical activity Breast cancer the Norwegian Women and Cancer Study. J Negat Results Biomed 2014; 13: 3.

Citations and B. Addition	nal information	
Risk	Outcome	Citation/Note
		Breslow RA, Ballard-Barbash R, Munoz K, Graubard BI. Long-term recreational physical activity and breast cancer in the National Health and Nutrition Examination Survey I epidemiologic follow-up study. Cancer
Low physical activity	Breast cancer	Epidemiol Biomarkers Prev 2001; 10: 805–8.
Low physical activity	Breast cancer	prospective study among elderly women. J Gerontol A Biol Sci Med Sci 1998; 53: M251-256.
		Chang S-C, Ziegler RG, Dunn B, et al. Association of energy intake and energy balance with postmenopausal breast cancer in the prostate, lung, colorectal, and ovarian cancer screening trial. Cancer Epidemiol
Low physical activity	Breast cancer	Biomarkers Prev 2006; 15: 334–41.
Low physical activity	Prosst concor	Colditz GA, Feskanich D, Chen WY, Hunter DJ, Willett WC. Physical activity and risk of breast cancer in
Low physical activity	Breast cancer	Dallal CM, Sullivan-Halley J, Ross RK, et al. Long-term recreational physical activity and risk of invasive
Low physical activity	Breast cancer	and in situ breast cancer: the California teachers study. Arch Intern Med 2007; 167: 408–15.
Low physical activity	Breast cancer	Study. Am J Epidemiol 1994; 139: 662–9.
Low physical activity	Breast cancer	Eliassen AH, Hankinson SE, Rosner B, Holmes MD, Willett WC. Physical activity and risk of breast cancer among postmenopausal women. Arch Intern Med 2010; 170; 1758. 64
Low physical activity	bleast calleel	among posumenopausar women. Aren men med 2010, 170. 1758–04.
	_	Frisch RE, Wyshak G, Witschi J, Albright NL, Albright TE, Schiff I. Lower lifetime occurrence of breast
Low physical activity	Breast cancer	cancer and cancers of the reproductive system among former college athletes. Int J Fertil 1987; 32: 217–25.
		prevention recommendations and risk of postmenopausal breast cancer. Cancer Epidemiol Biomarkers Prev
Low physical activity	Breast cancer	2013; 22: 1498–508.
		Hildebrand JS, Gapstur SM, Campbell PT, Gaudet MM, Patel AV. Recreational physical activity and leisure-
Low physical activity	Breast cancer	1906–12.
		Howard RA, Leitzmann MF, Linet MS, Freedman DM. Physical activity and breast cancer risk among pre-
Low physical activity	Prost concor	and postmenopausal women in the U.S. Radiologic Technologists cohort. Cancer Causes Control 2009; 20:
Low physical activity	breast cancer	Leitzmann MF, Moore SC, Peters TM, et al. Prospective study of physical activity and risk of postmenopausal
Low physical activity	Breast cancer	breast cancer. Breast Cancer Res 2008; 10: R92.
Low physical activity	Breast cancer	Luoto R, Latikka P, Pukkala E, Hakulinen T, Vihko V. The effect of physical activity on breast cancer risk: a cohort study of 30.548 women. Eur J Epidemiol 2000: 16: 973–80.
		Margolis KL, Mucci L, Braaten T, et al. Physical activity in different periods of life and the risk of breast
Low physical activity	Prost concor	cancer: the Norwegian-Swedish Women's Lifestyle and Health cohort study. Cancer Epidemiol Biomarkers
Low physical activity	Dreast cancer	110/ 2003, 14. 27-32.
Low physical activity	Breast cancer	Mertens AJ, Sweeney C, Shahar E, Rosamond WD, Folsom AR. Physical activity and breast cancer incidence in middle-aged women: a prospective cohort study. Breast Cancer Res Treat 2006; 97: 209–14.
Low physical activity	Breast cancer	Ministry of Health (Benin), National Institute of Statistics and Economic Analysis (INSAE) (Benin). Benin Health Statistical Vearbook 2005, Porto-Novo, Benin: Ministry of Health (Benin), 2006
Low physical activity	Breast culleer	Ministry of Health (Burkina Faso). Burkina Faso Health Statistical Yearbook 2007. Ouagadougou, Burkina
Low physical activity	Breast cancer	Faso: Ministry of Health (Burkina Faso), 2008.
Low physical activity	Breast cancer	Ministry of Health (Burkina Faso). Burkina Faso Health Statistical Yearbook 2008. Ouagadougou, Burkina Faso: Ministry of Health (Burkina Faso), 2009.
	_	Moradi T, Adami HO, Bergström R, et al. Occupational physical activity and risk for breast cancer in a
Low physical activity	Breast cancer	nationwide cohort study in Sweden. Cancer Causes Control 1999; 10: 423–30. Moradi T. Adami H-O. Ekbom A. et al. Physical activity and risk for breast cancer a prospective cohort study.
Low physical activity	Breast cancer	among Swedish twins. Int J Cancer 2002; 100: 76–81.
Low physical activity	Breast cancer	Peters TM, Schatzkin A, Gierach GL, et al. Physical activity and postmenopausal breast cancer risk in the NIH_AAPP diet and health study. Cancer Epidemiol Biomarkers Prev 2009: 18: 289–96
Low physical activity	breast current	Pronk A, Ji B-T, Shu X-O, et al. Physical activity and breast cancer risk in Chinese women. Br J Cancer
Low physical activity	Breast cancer	2011; 105: 1443–50.
Low physical activity	Breast cancer	Rintala by PE, Pukkala E, Paakkulainen HT, Vihko VJ. Self-experienced physical workload and risk of breast cancer. Scand J Work Environ Health 2002; 28: 158–62.
, j		Rockhill B, Willett WC, Hunter DJ, Manson JE, Hankinson SE, Colditz GA. A prospective study of
Low physical activity	Breast cancer	recreational physical activity and breast cancer risk. Arch Intern Med 1999; 159: 2290–6.
		physical activity and breast cancer incidence in African-American women. Cancer Epidemiol Biomarkers
Low physical activity	Breast cancer	Prev 2014; 23: 2522–31.
Low physical activity	Breast cancer	Sesso HD, Paffenbarger RS, Lee IM. Physical activity and breast cancer risk in the College Alumni Health Study (United States). Cancer Causes Control 1998; 9: 433–9.
		Silvera SAN, Jain M, Howe GR, Miller AB, Rohan TE. Energy balance and breast cancer risk: a prospective
Low physical activity	Breast cancer	cohort study. Breast Cancer Res Treat 2006; 97: 97–106. Suzuki R. Juagaki M. Vamamoto S. et al. Laigure time physical activity and breast cancer rick defined by
		estrogen and progesterone receptor statusthe Japan Public Health Center-based Prospective Study. Prev Med
Low physical activity	Breast cancer	2011; 52: 227–33.
Low physical activity	Breast cancer	Suzuki S, Kojima M, Tokudome S, et al. Effect of physical activity on breast cancer risk: findings of the Japan collaborative cohort study. Cancer Epidemiol Biomarkers Prev 2008: 17: 3396–401
···· F, stear activity		Thune I, Brenn T, Lund E, Gaard M. Physical activity and the risk of breast cancer. N Engl J Med 1997; 336:
Low physical activity	Breast cancer	1269–75.

Appendix Table 5. Epide Citations and B. Addition A. Citations	miological evidence supporting nal information	causality between risk-outcome pairs included in the Global Burden of Disease 2017 study including A.
Risk	Outcome	Citation/Note Wyrwich KW, Wolinsky FD. Physical activity, disability, and the risk of hospitalization for breast cancer
Low physical activity	Breast cancer	among older women. J Gerontol A Biol Sci Med Sci 2000; 55: M418-421.
Low physical activity	Breast cancer	follow-up. Br J Cancer 2000; 82: 726–30.
Low physical activity	Ischaemic stroke	Abbott RD, Rodriguez BL, Burchfiel CM, Curb JD. Physical activity in older middle-aged men and reduced risk of stroke: the Honolulu Heart Program. Am J Epidemiol 1994; 139: 881–93.
Low physical activity	Ischaemic stroke	Agnarsson U, Thorgeirsson G, Sigvaldason H, Sigfusson N. Effects of leisure-time physical activity and ventilatory function on risk for stroke in men: the Revkiavík Study. Ann Intern Med 1999: 130: 987–90
Low physical activity	iselikeline stoke	Autenrieth CS, Evenson KR, Yatsuya H, Shahar E, Baggett C, Rosamond WD. Association between physical activity and risk of stroke subtypes: the atherosclerosis risk in communities study. Neuroepidemiology 2013;
Low physical activity	Ischaemic stroke	40: 109–16.
Low physical activity	Ischaemic stroke	mortality from cardiovascular diseases and all causes: The Zutphen Elderly Study. Arch Intern Med 1998; 158: 1499–505.
		Calling S, Hedblad B, Engström G, Berglund G, Janzon L. Effects of body fatness and physical activity on cardiovascular risk: risk prediction using the bioelectrical impedance method. Scand J Public Health 2006;
Low physical activity	Ischaemic stroke	34: 568–75.
Low physical activity	Ischaemic stroke	Chiuve SE, Rexrode KM, Spiegelman D, Logroscino G, Manson JE, Rimm EB. Primary prevention of stroke by healthy lifestyle. Circulation 2008; 118: 947–54.
Low physical activity	Ischaemic stroke	Ellekjaer H, Holmen J, Ellekjaer E, Vatten L. Physical activity and stroke mortality in women. Ten-year follow-up of the Nord-Trondelag health survey, 1984-1986. Stroke 2000; 31: 14–8.
Low physical activity	Ischaemic stroke	Gulsvik AK, Thelle DS, Samuelsen SO, Myrstad M, Mowé M, Wyller TB. Ageing, physical activity and mortalitya 42-year follow-up study. Int J Epidemiol 2012; 41: 521–30.
Low physical activity	Ischaemic stroke	Häheim LL, Holme I, Hjermann I, Leren P. Risk factors of stroke incidence and mortality. A 12-year follow- up of the Oslo Study. Stroke 1993; 24: 1484–9.
Low physical activity	Ischaemic stroke	Hu FB, Stampfer MJ, Colditz GA, et al. Physical activity and risk of stroke in women. JAMA 2000; 283: 2961–7.
Low physical activity	Ischaemic stroke	Hu G, Sarti C, Jousilahti P, Silventoinen K, Barengo NC, Tuomilehto J. Leisure time, occupational, and commuting physical activity and the risk of stroke. Stroke 2005; 36: 1994–9.
		Lapidus L, Bengtsson C. Socioeconomic factors and physical activity in relation to cardiovascular disease and death. A 12 year follow up of participants in a population study of women in Gothenburg, Sweden. Br Heart J
Low physical activity	Ischaemic stroke	1986; 55: 295–301. Lee IM, Hennekens CH, Berger K, Buring JE, Manson JE. Exercise and risk of stroke in male physicians.
Low physical activity	Ischaemic stroke	Stroke 1999; 30: 1–6.
Low physical activity	Ischaemic stroke	Lee IM, Pattenbarger RS. Physical activity and stroke incidence: the Harvard Alumni Health Study. Stroke 1998; 29: 2049–54.
Low physical activity	Ischaemic stroke	Lindenstrøm E, Boysen G, Nyboe J. Lifestyle factors and risk of cerebrovascular disease in women. The Copenhagen City Heart Study. Stroke 1993; 24: 1468–72.
Low physical activity	Ischaemic stroke	Myint PK, Luben RN, Wareham NJ, et al. Combined work and leisure physical activity and risk of stroke in men and women in the European prospective investigation into Cancer-Norfolk Prospective Population Study, Neuroepidemiology 2006; 27: 122–9
Low physical activity	isolatellite subject	Okada H, Horibe H, Yoshiyuki O, Hayakawa N, Aoki N. A prospective study of cerebrovascular disease in
Low physical activity	Ischaemic stroke	Japanese rural communities, Akabane and Asahi. Part 1: evaluation of risk factors in the occurrence of cerebral hemorrhage and thrombosis. Stroke 1976; 7: 599–607.
Low physical activity	Ischaemic stroke	Paffenbarger RS, Brand RJ, Sholtz RI, Jung DL. Energy expenditure, cigarette smoking, and blood pressure level as related to death from specific diseases. Am J Epidemiol 1978: 108: 12–8
		Paganini-Hill A, Perez Barreto M. Stroke risk in older men and women: aspirin, estrogen, exercise, vitamins,
Low physical activity	Ischaemic stroke	and other factors. J Gend Specif Med 2001; 4: 18–28. Salonen JT, Puska P, Tuomilehto J, Physical activity and risk of myocardial infarction, cerebral stroke and
Low physical activity	Ischaemic stroke	death: a longitudinal study in Eastern Finland. Am J Epidemiol 1982; 115: 526–37.
Low physical activity	Ischaemic stroke	Saterman JK, Kuru I, Burng JE, Lee I-M. Physical activity and risk of stroke in women. Stroke 2010; 41: 1243–50.
Low physical activity	Ischaemic stroke	Simonsick EM, Lafferty ME, Phillips CL, et al. Kisk due to inactivity in physically capable older adults. Am J Public Health 1993; 83: 1443–50.
Low physical activity	Ischaemic stroke	Wannamethee G, Shaper AG. Physical activity and stroke in British middle aged men. BMJ 1992; 304: 597–601.
Low physical activity	Ischaemic stroke	Willey JZ, Moon YP, Paik MC, Boden-Albala B, Sacco RL, Elkind MSV. Physical activity and risk of ischemic stroke in the Northern Manhattan Study. Neurology 2009; 73: 1774–9.
Low physical activity	Ischaemic stroke	Zhang Q, Zhou Y, Gao X, et al. Ideal cardiovascular health metrics and the risks of ischemic and intracerebral hemorrhagic stroke. Stroke 2013; 44: 2451–6.
Low physical activity	Diabetes mellitus	Baan CA, Stolk RP, Grobbee DE, Witteman JC, Feskens EJ. Physical activity in elderly subjects with impaired glucose tolerance and newly diagnosed diabates molliture. Am J Emidemiol 1000; 140: 210: 27
Low physical activity	Diabetes mentus	Bonora E, Kiech S, Willeit J, et al. Population-based incidence rates and risk factors for type 2 diabetes in white in livid whether Population-based incidence rates and risk factors for type 2 diabetes in
Low physical activity	Diabetes mellitus	white individuals: the Bruneck study. Diabetes 2004; 53: 1/82–9. Burchfiel CM, Sharp DS, Curb JD, et al. Physical activity and incidence of diabetes: the Honolulu Heart
Low physical activity	Diabetes mellitus	Program. Am J Epidemiol 1995; 141: 360–8. Carlsson S. Ahlbom A. Lichtenstein P. Andersson T. Shared genetic influence of RML physical activity and
Low physical activity	Diabetes mellitus	type 2 diabetes: a twin study. Diabetologia 2013; 56: 1031–5.

Appendix Table 5. Epidemiological evidence supporting causality between risk-outcome pairs included in the Global Burden of Disease 2017 study including A. **Citations and B. Additional information** A. Citations Risk Outcome Citation/Note Carlsson S, Midthjell K, Tesfamarian MY, Grill V. Age, overweight and physical inactivity increase the risk of latent autoimmune diabetes in adults: results from the Nord-Trøndelag health study. Diabetologia 2007; Low physical activity Diabetes mellitus 50: 55 - 8Chien K-L, Chen M-F, Hsu H-C, Su T-C, Lee Y-T. Sports activity and risk of type 2 diabetes in Chinese. Low physical activity Diabetes mellitus Diabetes Res Clin Pract 2009; 84: 311-8. Demakakos P, Hamer M, Stamatakis E, Steptoe A. Low-intensity physical activity is associated with reduced risk of incident type 2 diabetes in older adults: evidence from the English Longitudinal Study of Ageing. Low physical activity Diabetes mellitus Diabetologia 2010; 53: 1877-85. Doi Y, Ninomiya T, Hata J, et al. Two risk score models for predicting incident Type 2 diabetes in Japan. Diabetes mellitus Diabet Med 2012; 29: 107-14. Low physical activity Dotevall A, Johansson S, Wilhelmsen L, Rosengren A. Increased levels of triglycerides, BMI and blood pressure and low physical activity increase the risk of diabetes in Swedish women. A prospective 18-year Diabetes mellitus Low physical activity follow-up of the BEDA study. Diabet Med 2004; 21: 615-22. Elwood P, Galante J, Pickering J, et al. Healthy lifestyles reduce the incidence of chronic diseases and Low physical activity Diabetes mellitus dementia: evidence from the Caerphilly cohort study. PLoS ONE 2013; 8: e81877. Fan S, Chen J, Huang J, et al. Physical activity level and incident type 2 diabetes among Chinese adults. Med Low physical activity Diabetes mellitus Sci Sports Exerc 2015; 47: 751-6. Folsom AR, Kushi LH, Hong CP. Physical activity and incident diabetes mellitus in postmenopausal women. Low physical activity Diabetes mellitus Am J Public Health 2000; 90: 134-8. Fretts AM, Howard BV, Kriska AM, et al. Physical activity and incident diabetes in American Indians: the Diabetes mellitus Strong Heart Study. Am J Epidemiol 2009; 170: 632-9. Low physical activity Grøntved A, Pan A, Mekary RA, et al. Muscle-strengthening and conditioning activities and risk of type 2

Low physical activity

Diabetes mellitus

Diabetes mellitus

Diabetes mellitus

Diabetes mellitus

Diabetes mellitus

Diabetes mellitus

Low physical activity	Diabetes mellitus	Hsia J, Wu L, Allen C, et al. Physical activity and diabetes risk in postmenopausal women. Am J Prev Med 2005; 28: 19–25.
Low physical activity	Diabetes mellitus	Hu FB, Leitzmann MF, Stampfer MJ, Colditz GA, Willett WC, Rimm EB. Physical activity and television watching in relation to risk for type 2 diabetes mellitus in men. Arch Intern Med 2001; 161: 1542–8.
Low physical activity	Diabetes mellitus	Hu FB, Sigal RJ, Rich-Edwards JW, et al. Walking compared with vigorous physical activity and risk of type 2 diabetes in women: a prospective study. JAMA 1999; 282: 1433–9.
Low physical activity	Diabetes mellitus	Hu G, Qiao Q, Silventoinen K, et al. Occupational, commuting, and leisure-time physical activity in relation to risk for Type 2 diabetes in middle-aged Finnish men and women. Diabetologia 2003; 46: 322–9.
Low physical activity	Diabetes mellitus	James SA, Jamjoum L, Raghunathan TE, Strogatz DS, Furth ED, Khazanie PG. Physical activity and NIDDM in African-Americans. The Pitt County Study. Diabetes Care 1998; 21: 555–62.
Low physical activity	Diabetes mellitus	Jefferis BJ, Whincup PH, Lennon L, Wannamethee SG. Longitudinal associations between changes in physical activity and onset of type 2 diabetes in older British men: the influence of adiposity. Diabetes Care 2012; 35: 1876–83.
Low physical activity	Diabetes mellitus	Joseph J, Svartberg J, Njølstad I, Schirmer H. Incidence of and risk factors for type-2 diabetes in a general population: the Tromsø Study. Scand J Public Health 2010; 38: 768–75.

1997; 26: 739-47.

Public Health 2007; 7: 154.

 biabetes mellitus
 Koloverou E, Panagiotakos DB, Pitsavos C, et al. 10-year incidence of diabetes and associated risk factors in

 Diabetes mellitus
 Greece: the ATTICA study (2002-2012). Rev Diabet Stud 2014; 11: 181–9.

 Krishnan S, Rosenberg L, Palmer JR. Physical activity and television watching in relation to risk of type 2

 Diabetes mellitus
 diabetes: the Black Women's Health Study. Am J Epidemiol 2009; 169: 428–34.

Laaksonen MA, Knekt P, Rissanen H, et al. The relative importance of modifiable potential risk factors of type 2 diabetes: a meta-analysis of two cohorts. Eur J Epidemiol 2010; 25: 115–24.

diabetes: a prospective study in two cohorts of US women. PLoS Med 2014; 11: e1001587.

treatment in the elderly. J Am Geriatr Soc 1994; 42: 1235-40.

physical activity. Med Sci Sports Exerc 1994; 26: 824-30.

Gurwitz JH, Field TS, Glynn RJ, et al. Risk factors for non-insulin-dependent diabetes mellitus requiring

Helmrich SP, Ragland DR, Paffenbarger RS. Prevention of non-insulin-dependent diabetes mellitus with

Holme I, Tonstad S, Sogaard AJ, Larsen PGL, Haheim LL. Leisure time physical activity in middle age predicts the metabolic syndrome in old age: results of a 28-year follow-up of men in the Oslo study. BMC

Haapanen N, Miilunpalo S, Vuori I, Oja P, Pasanen M. Association of leisure time physical activity with the risk of coronary heart disease, hypertension and diabetes in middle-aged men and women. Int J Epidemiol

Lee D, Park I, Jun T-W, et al. Physical activity and body mass index and their associations with the development of type 2 diabetes in korean men. Am J Epidemiol 2012; 176: 43–51.

Low physical activity	Diabetes mellitus	Longo-Mbenza B, On'kin JBKL, Okwe AN, Kabangu NK, Fuele SM. Metabolic syndrome, aging, physical inactivity, and incidence of type 2 diabetes in general African population. Diab Vasc Dis Res 2010; 7: 28–39.
Low physical activity	Diabetes mellitus	Lucke J, Waters B, Hockey R, et al. Trends in women's risk factors and chronic conditions: findings from the Australian Longitudinal Study on Women's Health. Womens Health (Lond) 2007; 3: 423–32.
Low physical activity	Diabetes mellitus	Magliano DJ, Barr ELM, Zimmet PZ, et al. Glucose indices, health behaviors, and incidence of diabetes in Australia: the Australian Diabetes, Obesity and Lifestyle Study. Diabetes Care 2008; 31: 267–72.
Low physical activity	Diabetes mellitus	Manson JE, Nathan DM, Krolewski AS, Stampfer MJ, Willett WC, Hennekens CH. A prospective study of exercise and incidence of diabetes among US male physicians. JAMA 1992; 268: 63–7.
Low physical activity	Diabetes mellitus	Manson JE, Rimm EB, Stampfer MJ, et al. Physical activity and incidence of non-insulin-dependent diabetes mellitus in women. Lancet 1991; 338: 774–8.

Appendix Table 5. Epidemio Citations and B. Additional i A. Citations	logical evidence supporting causa nformation	lity between risk-outcome pairs included in the Global Burden of Disease 2017 study including A.
Risk	Outcome	Citation/Note
		Meisinger C, Löwel H, Thorand B, Döring A. Leisure time physical activity and the risk of type 2 diabetes in men and women from the general population. The MONICA/KORA Augsburg Cohort Study. Diabetologia
Low physical activity	Diabetes mellitus	2005; 48: 27–34. Mozaffarian D. Kaminani A. Carnethon M. Dioussá I. Mukamal KI. Siscovick D. Lifestyle rick factors and
Low physical activity	Disbatas mellitus	new-onset diabetes mellitus in older adults: the cardiovascular health study. Arch Intern Med 2009; 169:
Low physical activity	Diabetes mennus	Okada K, Hayashi T, Tsumura K, Suematsu C, Endo G, Fujii S. Leisure-time physical activity at weekends
Low physical activity	Diabatas mallitus	and the risk of Type 2 diabetes mellitus in Japanese men: the Osaka Health Survey. Diabet Med 2000; 17:
Low physical activity	Diabetes menitus	Panagiotakos DB, Pitsavos C, Skoumas Y, Lentzas Y, Stefanadis C. Five-year incidence of type 2 diabetes mellitus among cardiovascular disease-free Greek adults: findings from the ATTICA study. Vasc Health Rick
Low physical activity	Diabetes mellitus	Manag 2008; 4: 691–8.
		Detheren W. Strashurzz V. Heim M. et al. Insidence of True 2 diskets in the aldede Common production
Low physical activity	Diabetes mellitus	Rathmann W, Strassburger K, Heier M, et al. Incidence of Type 2 diabetes in the elderly German population and the effect of clinical and lifestyle risk factors: KORA S4/F4 cohort study. Diabet Med 2009: 26: 1212–9.
		Reis JP, Loria CM, Sorlie PD, Park Y, Hollenbeck A, Schatzkin A. Lifestyle factors and risk for new-onset
Low physical activity	Diabetes mellitus	diabetes: a population-based cohort study. Ann Intern Med 2011; 155: 292–9.
Low physical activity	Diabetes mellitus	Shi L, Shu X-O, Li H, et al. Physical activity, smoking, and alconic consumption in association with incidence of type 2 diabetes among middle-aged and elderly Chinese men. PLoS ONE 2013; 8: e77919.
Low physical activity	Diabetes mellitus	Siegel LC, Sesso HD, Bowman TS, Lee I-M, Manson JE, Gaziano JM. Physical activity, body mass index, and diabetes risk in men: a prospective study. Am J Med 2009: 122: 1115–21.
		Simonsick EM, Lafferty ME, Phillips CL, et al. Risk due to inactivity in physically capable older adults. Am J
Low physical activity	Diabetes mellitus	Public Health 1993; 83: 1443–50. Steinbrecher A. Erber F. Grandinetti A. Nigg C. Kolonel I.N. Maskarinec G. Physical activity and risk of type
Low physical activity	Diabetes mellitus	2 diabetes among Native Hawaiians, Japanese Americans, and Caucasians: the Multiethnic Cohort. J Phys Act Health 2012; 9: 634–41.
Low physical activity	Diabetes mellitus	Stringhini S, Tabak AG, Akbaraly TN, et al. Contribution of modifiable risk factors to social inequalities in type 2 diabetes: prospective Whitehall II cohort study. BMJ 2012; 345: e5452.
T	Dishetes mellitus	Sun F, Tao Q, Zhan S. An accurate risk score for estimation 5-year risk of type 2 diabetes based on a health
Low physical activity	Diabetes mennus	Tsai AC, Lee S-H. Determinants of new-onset diabetes in older adults—Results of a national cohort study.
Low physical activity	Diabetes mellitus	Clin Nutr 2015; 34: 937–42.
Low physical activity	Diabetes mellitus	Villegas R, Shu X-O, Li H, et al. Physical activity and the incidence of type 2 diabetes in the Shanghai women's health study. Int J Epidemiol 2006; 35: 1553–62.
Low physical activity	Diabetes mellitus	among middle-aged Japanese: a population-based prospective study in the JPHC study cohort I. Diabet Med 2005; 22: 323–31.
		Waller K, Kaprio J, Lehtovirta M, Silventoinen K, Koskenvuo M, Kujala UM. Leisure-time physical activity
Low physical activity	Diabetes mellitus	and type 2 diabetes during a 28 year follow-up in twins. Diabetologia 2010; 53: 2531–7.
Low physical activity	Diabetes mellitus	heart disease and type 2 diabetes. Arch Intern Med 2000; 160: 2108–16.
Low physical activity	Diabetes mellitus	Weinstein AR, Sesso HD, Lee IM, et al. Relationship of physical activity vs body mass index with type 2 diabetes in women. JAMA 2004; 292: 1188–94.
Low physical activity	Diabetes mellitus	Williams PT, Thompson PD. Walking versus running for hypertension, cholesterol, and diabetes mellitus risk reduction. Arterioscler Thromb Vasc Biol 2013; 33: 1085–91
Low physical activity	Diabetes menitus	
Low physical activity	Diabetes mellitus	Xu F, Ware RS, Tse LA, et al. Joint associations of physical activity and hypertension with the development of type 2 diabetes among urban men and women in Mainland China. PLoS ONE 2014; 9: e88719.
		Boussageon R, Bejan-Angoulvant T, Saadatian-Elahi M, et al. Effect of intensive glucose lowering treatment on all cause mortality, cardiovascular death, and microvascular events in type 2 diabetes: meta-analysis of
High fasting plasma glucose	Ischaemic heart disease	randomised controlled trials. BMJ 2011; 343: d4169.
High fasting plasma glucose	Ischaemic heart disease	cardiovascular diseases and diabetes: a pooled analysis. PLoS ONE 2013; 8: e65174.
High fasting plasma glucose	Ischaemic stroke	Singh GM, Danaei G, Farzadfar F, et al. The age-specific quantitative effects of metabolic risk factors on cardiovascular diseases and diabetes: a pooled analysis. PLoS ONE 2013; 8: e65174.
High fasting plasma glucose	Ischaemic stroke	Zhang C, Zhou Y-H, Xu C-L, Chi F-L, Ju H-N. Efficacy of intensive control of glucose in stroke prevention: a meta-analysis of data from 59,197 participants in 9 randomized controlled trials. PloS One 2013: 8: e54465.
0 01 0		Singh GM, Danaei G, Farzadfar F, et al. The age-specific quantitative effects of metabolic risk factors on
High fasting plasma glucose	Hemorrhagic stroke	cardiovascular diseases and diabetes: a pooled analysis. PLoS ONE 2013; 8: e65174.
		Zhang C, Zhou Y-H, Xu C-L, Chi F-L, Ju H-N. Efficacy of intensive control of glucose in stroke prevention:
High fasting plasma glucose	Hemorrhagic stroke	a meta-analysis of data from 59,197 participants in 9 randomized controlled trials. PloS One 2013; 8: e54465.
	Chronic kidney disease due to	development of renal end points in type 2 diabetes mellitus: systematic review and meta-analysis intensive
High fasting plasma glucose	diabetes mellitus	glucose control in type 2 diabetes. Arch Intern Med 2012; 172: 761–9.
High fasting plasma glucose	diabetes mellitus	O Seagnona Civi, reiković v, Lani i n, et al. Blood rressure is a Major Kisk Factor for Kenal Death An Analysis of 560 352 Particinants From the Asia-Pacific Region Hypertension 2000: 54: 509–15

A. Citations		
Risk	Outcome	Citation/Note
	Chronic kidney disease due to	development of renal end points in type 2 diabetes mellitus: systematic review and meta-analysis intensive
High fasting plasma glucose	hypertension	glucose control in type 2 diabetes. Arch Intern Med 2012: 172: 761–9.
88 F 88	Chronic kidney disease due to	O'Seaghdha CM, Perkovic V, Lam TH, et al. Blood Pressure Is a Major Risk Factor for Renal Death An
High fasting plasma glucose	hypertension	Analysis of 560 352 Participants From the Asia-Pacific Region. Hypertension 2009; 54: 509-15.
		Coca SG, Ismail-Beigi F, Haq N, Krumholz HM, Parikh CR. Role of intensive glucose control in
	Chronic kidney disease due to	development of renal end points in type 2 diabetes mellitus: systematic review and meta-analysis intensive
High fasting plasma glucose	glomerulonephritis	glucose control in type 2 diabetes. Arch Intern Med 2012; 172: 761–9.
High fasting plasma glucosa	chronic kidney disease due to	O'Seaghdha CM, Perkovic V, Lam TH, et al. Blood Pressure is a Major Risk Factor for Renal Death An Analysis of 560 352 Participants From the Asia Pacific Pagion, Hypertansion 2000; 54: 500, 15
Tingii fasting plasma glucose	giomeratorepintus	Coca SG Ismail-Beigi F Hao N Krumbolz HM Parikh CR Role of intensive glucose control in
	Chronic kidney disease due to	development of renal end points in type 2 diabetes mellitus: systematic review and meta-analysis intensive
High fasting plasma glucose	other causes	glucose control in type 2 diabetes. Arch Intern Med 2012; 172: 761–9.
	Chronic kidney disease due to	O'Seaghdha CM, Perkovic V, Lam TH, et al. Blood Pressure Is a Major Risk Factor for Renal Death An
High fasting plasma glucose	other causes	Analysis of 560 352 Participants From the Asia-Pacific Region. Hypertension 2009; 54: 509–15.
		Young F, Wotton CJ, Critchley JA, Unwin NC, Goldacre MJ. Increased risk of tuberculosis disease in people
High fasting plasma glucose	Tuberculosis	with diabetes mellitus: record-linkage study in a UK population. J Epidemiol Community Health. 2012;
Then fasting plasma glucose	Tuberculosis	Kuo MC, Lin SH, Lin CH, Mao IC, Chang SI, Hsieh MC, Type 2 diabetes: an independent risk factor for
High fasting plasma glucose	Tuberculosis	tuberculosis: a nationwide population-based study. PLoS One. 2013; 8((11):e78924).
		Dobler CC, Flack JR, Marks GB. Risk of tuberculosis among people with diabetes mellitus: an Australian
High fasting plasma glucose	Tuberculosis	nationwide cohort study. BMJ Open. 2((1):e000666).
		Baker MA, Lin HH, Chang HY, Murray MB. The risk of tuberculosis disease among persons with diabetes
High fasting plasma glucose	Tuberculosis	mellitus: a prospective cohort study. Clin Infect Dis. 2012; 54(6): 818-25.
High fasting plasma glucose	Tuberculosis	Diabetic control and risk of tuberculosis: a cohort study. Am J Epidemiol. 2008: 167(12): 1486-94
88 F 88		Kim SJ, Hong YP, Lew WJ, Yang SC, Lee EG. Incidence of pulmonary tuberculosis among diabetics. Tuber
High fasting plasma glucose	Tuberculosis	Lung Dis. 1995; 76(6): 529-33.
		Pealing L, Wing K, Mathur R, Prieto-Merino D, Smeeth L, Moore DA. Risk of tuberculosis in patients with
		diabetes: population based cohort study using the UK Clinical Practice Research Datalink. BMC Med. 2015;
High fasting plasma glucose	Tuberculosis	13(135). There I, Chen C, Hue S, Lies H, Weng M, Vieng V, Cee F. An undeted mote analysis of exhert studies.
High fasting plasma glucose	Dementia/Alzheimer's	diabetes and risk of Alzheimer's disease Diabetes Res Clin Pract 2017: Feb (124): 41-47
nigh hasting phasma graeose		Li L, Wan XH, Zhao GH. Meta-analysis of the risk of cataract in type 2 diabetes. BMC Ophthalmol. 2014;
High fasting plasma glucose	Cataracts	14(94).
		Zhao D, Cho J, Kim MH, Friedman DS, Guallar E. Diabetes, fasting glucose, and the risk of glaucoma: a
High fasting plasma glucose	Glaucoma	meta-analysis. Ophthalmology. 2015; 122(1): 72-8.
		znu L, Cao H, Znang T, Snen H, Dong W, Wang L, Du J. The effect of diabetes mentus on lung cancer prognosis: a PRISMA-compliant meta-analysis of cohort studies. Medicine (Baltimore), 2016; 95(17):
High fasting plasma glucose	Lung cancer	e3528.
0 01 0		Boyle P, Boniol M, Koechlin A, et al. Diabetes and breast cancer risk: a meta-analysis. Br J Cancer 2012;
High fasting plasma glucose	Breast cancer	107: 1608–17.
		Fang H, Yao B, Yan Y, Xu H, Liu Y, Tang H, Zhou J, Cao L, Wang W, Zhang J, Zhao L, Chen X, Zhang F,
Uigh fasting plasma glusses	Pladdar annaar	Zhao Y. Diabetes mellitus increases the risk of bladder cancer: an updated meta-analysis. Diabetes Technol There 2012; 15(11): 014 22
Tingii fasting plasma glucose	Bladder cancer	Shi I Xiong I Li I Cao H Jiang W Liu B Chen X Liu C Liu K Wang G Cai K A linear dose-response
		relationship between fasting plasma glucose and colorectal cancer risk: systematic review and meta-analysis.
High fasting plasma glucose	Colon cancer	Sci Rep. 2015; 5(17591).
High fasting plasma glucosa	Overien concor	Lee JY I, Jeon I, Kim JW, Song YS, Yoon JM, Park SM. Diabetes mellitus and ovarian cancer risk: a systematic raview and meta analysis of observational studies. Int L Compared Obstat. 2013; 23(2): 402-12
Then fasting plasma glucose	ovarian cancer	Ben O. Xu M. Ning X. Liu J. Hong S. Huang W. Zhang H. Li Z. Diabetes mellitus and risk of pancreatic
High fasting plasma glucose	Pancreatic cancer	cancer: A meta-analysis of cohort studies. Eur J Cancer. 2011; 47(13): 1928-1937.
		Yang WS, Va P, Bray F, Gao S, Gao J, Li HL, Xiang YB. The role of pre-existing diabetes mellitus on
		hepatocellular carcinoma occurrence and prognosis: a meta-analysis of prospective cohort studies. PLoS One.
High fasting plasma glucose	Liver cancer	2011; 6(12): :e27326.
		Rapsomaniki E, Timmis A, George J, Pujades-Rodriguez M, Snan AD, Denaxas S, White IK, Caulifield MJ, Deapfield IF. Smeeth I. Williams B. Hinggrani A. Hemingway H. Blood pressure and incidence of twelve
		cardiovascular diseases: lifetime risks, healthy life-years lost, and age-specific associations in 1.25 million
High fasting plasma glucose	Peripheral vascular disease	people. Lancet. 2014; 31(383): 1899-911.
		Rapsomaniki E, Timmis A, George J, Pujades-Rodriguez M, Shah AD, Denaxas S, White IR, Caulfield MJ,
		Deanfield JE, Smeeth L, Williams B, Hingorani A, Hemingway H. Blood pressure and incidence of twelve
High fasting plasma glucoso	Ischaemic stroke	cardiovascular diseases: lifetime risks, healthy life-years lost, and age-specific associations in 1.25 million people. Lancet 2014; 31(383): 1899-011
ringii rasung piasina giucose	Isenaenne suoke	Ransomaniki E. Timmis A. George I. Puiades-Rodriguez M. Shah AD. Denaxas S. White IR. Caulfield MI
		Deanfield JE, Smeeth L, Williams B, Hingorani A, Hemingway H. Blood pressure and incidence of twelve
		cardiovascular diseases: lifetime risks, healthy life-years lost, and age-specific associations in 1.25 million
High fasting plasma glucose	Hemorrhagic stroke	people. Lancet. 2014; 31(383): 1899-911.

A. Citations		
Risk	Outcome	Citation/Note
High LDL cholesterol	Ischaemic heart disease	cholesterol Treatment Trialists' (CTT) Collaboration, Baigent C, Blackwell L, et al. Efficacy and safety of more intensive lowering of LDL cholesterol: a meta-analysis of data from 170,000 participants in 26 randomised trials. Lancet Lond Engl 2010: 376: 1670–81.
8		Singh GM, Danaei G, Farzadfar F, et al. The age-specific quantitative effects of metabolic risk factors on
High LDL cholesterol	Ischaemic heart disease	cardiovascular diseases and diabetes: a pooled analysis. PLoS ONE 2013; 8: e65174.
		Cholesterol Treatment Trialists' (CTT) Collaboration, Baigent C, Blackwell L, et al. Efficacy and safety of more intensive lowering of LDL cholesterol: a meta-analysis of data from 170,000 participants in 26
High LDL cholesterol	Ischaemic stroke	randomised trials. Lancet Lond Engl 2010; 3/6: 16/0–81.
High LDL cholesterol	Ischaemic stroke	cardiovascular diseases and diabetes: a pooled analysis. PLoS ONE 2013; 8: e65174.
High systolic blood pressure	Rheumatic heart disease	Singh GM, Danael G, Farzadiar F, et al. The age-specific quantitative effects of metabolic risk factors on cardiovascular diseases and diabetes: a pooled analysis. PLoS ONE 2013; 8: e65174.
High systolic blood pressure	Ischaemic heart disease	cardiovascular diseases and diabetes; a pooled analysis, PLoS ONE 2013; 8: e65174.
		Thomopoulos C, Parati G, Zanchetti A. Effects of blood pressure lowering on outcome incidence in hypertension 1. Overview meta-analyses and meta-regression analyses of randomized trials. I Hypertens
High systolic blood pressure	Ischaemic heart disease	2014; 32: 2285–95.
High systolic blood pressure	Ischaemic stroke	Singh GM, Danaei G, Farzadfar F, et al. The age-specific quantitative effects of metabolic risk factors on cardiovascular diseases and diabetes: a pooled analysis. PLoS ONE 2013; 8: e65174.
High systolic blood pressure	Ischaemic stroke	Thomopoulos C, Parati G, Zanchetti A. Effects of blood pressure lowering on outcome incidence in hypertension. 1. Overview, meta-analyses, and meta-regression analyses of randomized trials. J Hypertens 2014: 32: 2285–95.
		Singh GM, Danaei G, Farzadfar F, et al. The age-specific quantitative effects of metabolic risk factors on
High systolic blood pressure	Hemorrhagic stroke	cardiovascular diseases and diabetes: a pooled analysis. PLoS ONE 2013; 8: e65174.
High systolic blood pressure	Hemorrhagic stroke	hypertension. 1. Overview, meta-analyses, and meta-regression analyses of randomized trials. J Hypertens 2014; 32: 2285–95.
		Singh GM, Danaei G, Farzadfar F, et al. The age-specific quantitative effects of metabolic risk factors on
High systolic blood pressure	Hypertensive heart disease	cardiovascular diseases and diabetes: a pooled analysis. PLoS ONE 2013; 8: e65174. Thomopoulos C, Parati G, Zanchetti A. Effects of blood pressure lowering on outcome incidence in
High systolic blood pressure	Hypertensive heart disease	hypertension. 1. Overview, meta-analyses, and meta-regression analyses of randomized trials. J Hypertens 2014; 32: 2285–95.
High systolic blood pressure	Cardiomyopathy and myocarditis	Singh GM, Danaei G, Farzadfar F, et al. The age-specific quantitative effects of metabolic risk factors on cardiovascular diseases and diabetes: a pooled analysis. PLoS ONE 2013; 8: e65174.
		Emdin CA, Callender T, Cao J, Rahimi K. Effect of antihypertensive agents on risk of atrial fibrillation: a
High systolic blood pressure	Atrial fibrillation and flutter	meta-analysis of large-scale randomized trials. Eur Eur Pacing Arrhythm Card Electrophysiol J Work Groups Card Pacing Arrhythm Card Cell Electrophysiol Eur Soc Cardiol 2015; 17: 701–10.
High systolic blood pressure	Atrial fibrillation and flutter	Singh GM, Danaei G, Farzadfar F, et al. The age-specific quantitative effects of metabolic risk factors on cardiovascular diseases and diabetes: a pooled analysis. PLoS ONE 2013; 8: e65174.
		Singh GM, Danaei G, Farzadfar F, et al. The age-specific quantitative effects of metabolic risk factors on
High systolic blood pressure	Aortic aneurysm	cardiovascular diseases and diabetes: a pooled analysis. PLoS ONE 2013; 8: e65174. Singh GM, Danaei G, Farzadfar F, et al. The age-specific quantitative effects of metabolic risk factors on
High systolic blood pressure	Peripheral vascular disease	cardiovascular diseases and diabetes: a pooled analysis. PLoS ONE 2013; 8: e65174.
High systolic blood pressure	Endocarditis	Singh GM, Danaei G, Farzadiar F, et al. The age-specific quantitative effects of metabolic risk factors on cardiovascular diseases and diabetes: a pooled analysis. PLoS ONE 2013; 8: e65174.
High systolic blood pressure	Other cardiovascular and circulatory diseases	Singh GM, Danaei G, Farzadfar F, et al. The age-specific quantitative effects of metabolic risk factors on cardiovascular diseases and diabetes: a pooled analysis. PLoS ONE 2013: 8: e65174.
High systolic blood pressure	Chronic kidney disease due to diabetes mellitus	The Renal Risk Collaboration, Foote C, Lin J, et al. The effect of Blood Pressure on Kidney Failure: a systematic review and meta-analysis in 2.7 million participants (unpublished).
	Chronic kidney disease due to	Xie X, Atkins E, Lv J, et al. Effects of intensive blood pressure lowering on cardiovascular and renal
High systolic blood pressure	diabetes mellitus Chronic kidney disease due to	outcomes: updated systematic review and meta-analysis. Lancet Lond Engl 2016; 387: 435–43. The Renal Risk Collaboration. Foote C. Lin J. et al. The effect of Blood Pressure on Kidney Failure: a
High systolic blood pressure	hypertension	systematic review and meta-analysis in 2.7 million participants (unpublished).
High systolic blood pressure	Chronic kidney disease due to hypertension	Xie X, Atkins E, Lv J, et al. Effects of intensive blood pressure lowering on cardiovascular and renal outcomes: updated systematic review and meta-analysis. Lancet Lond Engl 2016; 387: 435–43.
High systolic blood pressure	Chronic kidney disease due to glomerulonephritis	The Renal Risk Collaboration, Foote C, Lin J, et al. The effect of Blood Pressure on Kidney Failure: a systematic review and meta-analysis in 2.7 million participants (unpublished).
	Chronic kidney disease due to	Xie X, Atkins E, Lv J, et al. Effects of intensive blood pressure lowering on cardiovascular and renal
High systolic blood pressure	Chronic kidney disease due to	The Renal Risk Collaboration, Foote C, Lin J, et al. The effect of Blood Pressure on Kidney Failure: a
High systolic blood pressure	other causes	systematic review and meta-analysis in 2.7 million participants (unpublished).
High systolic blood pressure	Chronic kidney disease due to	Xie X, Atkins E, Lv J, et al. Effects of intensive blood pressure lowering on cardiovascular and renal outcomes: undated systematic review and meta-analysis. Lancet Lond Engl 2016; 387: 435–43
	Non-rheumatic calcific aortic	Bahler RC, Desser DR, Finkelhor RS, Brener SJ, Youssefi M. Factors leading to progression of valvular
High systolic blood pressure	valve disease	aortic stenosis. Am J Cardiol. 1999;84(9):1044-8. Hoadand PM, Cook FE, Flatlay M, Walker C, Coldman L, Cosa control analysis of rick feature for any second
High systolic blood pressure	valve disease	of aortic stenosis in adults (age 50 years or older). Am J Cardiol. 1985;55(6):744-7.
High systolic blood pressure	Non-rheumatic calcific aortic valve disease	Stewart BF, Siscovick D, Lind BK, et al. Clinical factors associated with calcific aortic valve disease. Cardiovascular Health Study, J Am Coll Cardiol, 1997;29(3):630-4
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Appendix Table 5. Epidemiological evidence supporting causality between risk-outcome pairs included in the Global Burden of Disease 2017 study including A. **Citations and B. Additional information** A. Citations Risk Outcome Citation/Note Schlesinger S, Lieb W, Koch M, et al. Body weight gain and risk of colorectal cancer: a systematic review High body-mass index Colon and rectum cancer and meta-analysis of observational studies. Obes Rev 2015; 16: 607-19. Chen Y, Wang X, Wang J, Yan Z, Luo J. Excess body weight and the risk of primary liver cancer: an updated High body-mass index Liver cancer meta-analysis of prospective studies. Eur J Cancer 2012; 48: 2137-45. Renehan AG, Tyson M, Egger M, Heller RF, Zwahlen M. Body-mass index and incidence of cancer: a High body-mass index Liver cancer systematic review and meta-analysis of prospective observational studies. Lancet 2008; 371: 569-78. Rui R, Lou J, Zou L, et al. Excess body mass index and risk of liver cancer: a nonlinear dose-response meta-High body-mass index Liver cancer analysis of prospective studies. PLoS ONE 2012; 7: e44522. Tanaka K, Tsuji I, Tamakoshi A, et al. Obesity and liver cancer risk: an evaluation based on a systematic High body-mass index Liver cancer review of epidemiologic evidence among the Japanese population. Jpn J Clin Oncol 2012; 42: 212-21 Wang Y, Wang B, Shen F, Fan J, Cao H. Body mass index and risk of primary liver cancer: a meta-analysis High body-mass index Liver cancer of prospective studies. Oncologist 2012; 17: 1461-8. Gallbladder and biliary tract Park M, Song DY, Je Y, Lee JE. Body mass index and biliary tract disease: a systematic review and meta-High body-mass index analysis of prospective studies. Prev Med 2014; 65: 13-22 cancer Renehan AG, Tyson M, Egger M, Heller RF, Zwahlen M. Body-mass index and incidence of cancer: a Gallbladder and biliary tract High body-mass index cancer systematic review and meta-analysis of prospective observational studies. Lancet 2008; 371: 569-78. Alsamarrai A, Das SLM, Windsor JA, Petrov MS. Factors that affect risk for pancreatic disease in the general population: a systematic review and meta-analysis of prospective cohort studies. Clin Gastroenterol Hepatol High body-mass index Pancreatic cancer 2014; 12: 1635-1644.e5; quiz e103. Renehan AG, Tyson M, Egger M, Heller RF, Zwahlen M. Body-mass index and incidence of cancer: a High body-mass index Pancreatic cancer systematic review and meta-analysis of prospective observational studies. Lancet 2008; 371: 569-78. Renehan AG, Tyson M, Egger M, Heller RF, Zwahlen M. Body-mass index and incidence of cancer: a High body-mass index Breast cancer (Pre-menopause) systematic review and meta-analysis of prospective observational studies. Lancet 2008; 371: 569-78. Xia X, Chen W, Li J, et al. Body mass index and risk of breast cancer: a nonlinear dose-response meta-High body-mass index analysis of prospective studies. Sci Rep 2014; 4: 7480. Breast cancer (Pre-menopause) Renehan AG, Tyson M, Egger M, Heller RF, Zwahlen M. Body-mass index and incidence of cancer: a High body-mass index Breast cancer (Post-menopause) systematic review and meta-analysis of prospective observational studies. Lancet 2008; 371: 569-78. Xia X, Chen W, Li J, et al. Body mass index and risk of breast cancer: a nonlinear dose-response meta-High body-mass index Breast cancer (Post-menopause) analysis of prospective studies. Sci Rep 2014; 4: 7480. Aune D, Greenwood DC, Chan DSM, et al. Body mass index, abdominal fatness and pancreatic cancer risk: a systematic review and non-linear dose-response meta-analysis of prospective studies. Ann Oncol 2012; 23: High body-mass index Uterine cancer 843-52 Jenabi E, Poorolajal J. The effect of body mass index on endometrial cancer: a meta-analysis. Public Health High body-mass index Uterine cancer

 Uterine cancer
 2015; 129: 872–80.

 Renehan AG, Tyson M, Egger M, Heller RF, Zwahlen M. Body-mass index and incidence of cancer: a

 Uterine cancer
 systematic review and meta-analysis of prospective observational studies. Lancet 2008; 371: 569–78.

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 systematic review and nonlinear dose-response meta-analysis of prospective studies. Int J Cancer 2015; 136:

 Ovarian cancer
 1888–98.

 Collaborative Group on Epidemiological Studies of Ovarian Cancer. Ovarian cancer and body size: individual

		participant meta-analysis including 25,157 women with ovarian cancer from 47 epidemiological studies.
High body-mass index	Ovarian cancer	PLoS Med 2012; 9: e1001200.
		Liu Z, Zhang T-T, Zhao J-J, et al. The association between overweight, obesity and ovarian cancer: a meta-
High body-mass index	Ovarian cancer	analysis. Jpn J Clin Oncol 2015; 45: 1107–15.
		Renehan AG, Tyson M, Egger M, Heller RF, Zwahlen M. Body-mass index and incidence of cancer: a
High body-mass index	Ovarian cancer	systematic review and meta-analysis of prospective observational studies. Lancet 2008; 371: 569–78.
		Renehan AG, Tyson M, Egger M, Heller RF, Zwahlen M. Body-mass index and incidence of cancer: a
High body-mass index	Kidney cancer	systematic review and meta-analysis of prospective observational studies. Lancet 2008; 371: 569–78.
		Wang F, Xu Y. Body mass index and risk of renal cell cancer: a dose-response meta-analysis of published
High body-mass index	Kidney cancer	cohort studies. Int J Cancer 2014; 135: 1673–86.
		Ma J, Huang M, Wang L, Ye W, Tong Y, Wang H. Obesity and risk of thyroid cancer: evidence from a meta-
High body-mass index	Thyroid cancer	analysis of 21 observational studies. Med Sci Monit 2015; 21: 283–91.
		Renehan AG, Tyson M, Egger M, Heller RF, Zwahlen M. Body-mass index and incidence of cancer: a
High body-mass index	Thyroid cancer	systematic review and meta-analysis of prospective observational studies. Lancet. 2008; 371(9612): 569-78.
		Larsson SC, Wolk A. Body mass index and risk of non-Hodgkin's and Hodgkin's lymphoma: A meta-
High body-mass index	Non-hodgkin lymphoma	analysis of prospective studies. European Journal of Cancer 2011; 47: 2422–30.
		Castillo JJ, Reagan JL, Ingham RR, et al. Obesity but not overweight increases the incidence and mortality of
High body-mass index	Leukaemia	leukemia in adults: a meta-analysis of prospective cohort studies. Leuk Res 2012; 36: 868-75.
		Renehan AG, Tyson M, Egger M, Heller RF, Zwahlen M. Body-mass index and incidence of cancer: a
High body-mass index	Leukaemia	systematic review and meta-analysis of prospective observational studies. Lancet. 2008; 371(9612): 569-78.
		Teras LR, Kitahara CM, Birmann BM, et al. Body Size and Multiple Myeloma Mortality: a pooled analysis of
High body-mass index	Multiple myeloma	20 prospective studies. Br J Haematol 2014; 166: 667-76.
		Singh GM, Danaei G, Farzadfar F, et al. The age-specific quantitative effects of metabolic risk factors on
High body-mass index	Ischaemic heart disease	cardiovascular diseases and diabetes: a pooled analysis, PLoS ONE 2013; 8: e65174

High body-mass index

High body-mass index

High body-mass index

Cerebrovascular disease

Singh GM, Danaei G, Farzadfar F, et al. The age-specific quantitative effects of metabolic risk factors on cardiovascular diseases and diabetes: a pooled analysis. PLoS ONE 2013; 8: e65174.

A. Citations		
Risk	Outcome	Citation/Note Sinch CM, Dancei C, Ferradfer F, et al. The age appeific quantitative offects of matchelie rick fectors on
High body-mass index	Hypertensive heart disease	cardiovascular diseases and diabetes: a pooled analysis. PLoS ONE 2013; 8: e65174.
High body-mass index	Atrial fibrillation and flutter	Wanahita N, Messerli FH, Bangalore S, Gami AS, Somers VK, Steinberg JS. Atrial fibrillation and obesity—results of a meta-analysis. American Heart Journal 2008; 155: 310–5.
High body-mass index	Asthma	Beuther DA, Sutherland ER. Overweight, Obesity, and Incident Asthma. Am J Respir Crit Care Med 2007; 175: 661–6.
High body-mass index	Gallbladder diseases	Aune D, Norat T, Vatten LJ. Body mass index, abdominal fatness and the risk of gallbladder disease. Eur J Epidemiol 2015; 30: 1009–19.
High body-mass index	Alzheimer disease and other dementias	Profenno LA, Porsteinsson AP, Faraone SV. Meta-Analysis of Alzheimer's Disease Risk with Obesity, Diabetes, and Related Disorders. Biological Psychiatry 2010; 67: 505–12.
High body-mass index	Diabetes mellitus	Singh GM, Danaei G, Farzadfar F, et al. The age-specific quantitative effects of metabolic risk factors on cardiovascular diseases and diabetes: a pooled analysis. PLoS ONE 2013; 8: e65174.
High body-mass index	Chronic kidney disease	Emerging Risk Factors Collaboration, Wormser D, Kaptoge S, et al. Separate and combined associations of body-mass index and abdominal adiposity with cardiovascular disease: collaborative analysis of 58 prospective studies. Lancet 2011; 377: 1085–95.
High body-mass index	Chronic kidney disease	Ni Mhurchu C, Rodgers A, Pan WH, Gu DF, Woodward M, Asia Pacific Cohort Studies Collaboration. Body mass index and cardiovascular disease in the Asia-Pacific Region: an overview of 33 cohorts involving 310 000 participants. Int J Epidemiol 2004; 33: 751–8.
High body-mass index	Chronic kidney disease	Prospective Studies Collaboration, Whitlock G, Lewington S, et al. Body-mass index and cause-specific mortality in 900 000 adults: collaborative analyses of 57 prospective studies. Lancet 2009; 373: 1083–96. Jiang L, Rong J, Wang Y, et al. The relationship between body mass index and hip osteoarthritis: a systematic
High body-mass index	Osteoarthritis	review and meta-analysis. Joint Bone Spine 2011; 78: 150–5.
High body-mass index	Osteoarthritis	review and meta-analysis. Joint Bone Spine 2012; 79: 291–7.
High body-mass index	Osteoarthritis	Silverwood V, Blagojevic-Buckhall M, Jinks C, Jordan JL, Protneroe J, Jordan KP. Current evidence on risk factors for knee osteoarthritis in older adults: a systematic review and meta-analysis. Osteoarthr Cartil 2015; 23: 507–15.
High body-mass index	Low back pain	back pain: a meta-analysis. Am J Epidemiol 2010; 171: 135–54.
High body-mass index	Gout	Aune D, Norat T, Vatten LJ. Body mass index and the risk of gout: a systematic review and dose-response meta-analysis of prospective studies. Eur J Nutr 2014; 53: 1591–601.
High body-mass index	Cataract	Ye J, Lou L-X, He J-J, Xu Y-F. Body Mass Index and Risk of Age-Related Cataract: A Meta-Analysis of Prospective Cohort Studies. PLOS ONE 2014; 9: e89923.
High body-mass index (child)	Asthma	Mebrahtu TF, Feltbower RG, Greenwood DC, Parslow RC. Childhood body mass index and wheezing disorders: a systematic review and meta-analysis. Pediatr Allergy Immunol 2015; 26: 62–72.
Low bone mineral density	Injuries	Johnell O, Kanis JA, Oden A, et al. Predictive value of BMD for hip and other fractures. J Bone Miner Res 2005; 20: 1185–94.
Impaired kidney function	Ischaemic heart disease	 Chronic Kidney Disease Prognosis Consortium (CKD-PC). Chronic Kidney Disease Prognosis Consortium GBD 2016 Impaired Kidney Function Relative Risk Meta-Analysis. National Heart, Lung, and Blood Institute, National Institutes of Health (NIH). United States Atherosclerosis
Impaired kidney function	Ischaemic heart disease	Risk in Communities Study. Bethesda, United States: National Heart, Lung, and Blood Institute, National Institutes of Health (NIH). International Diabetes Institute (IDD, Australia Diabetes, Obesity and Lifestyle Study 1999-2000, Melbourne,
Impaired kidney function	Ischaemic heart disease	Australia: International Diabetes Institute (IDI).
Impaired kidney function	Ischaemic heart disease	National Institutes of Health (NIH). United States Framingham Heart Study . Association for Cardiac Research, Rome (Italy). The Cubbic Population Study .
Impaired kidney function	Ischaemic heart disease	National Heart, Lung, and Blood Institute, National Institutes of Health, University of California, Los Angeles (UCLA), University of Minnesota. United States Multi-Ethnic Study of Atherosclerosis First Examination 2000-2002. Bethesda, United States: National Heart, Lung, and Blood Institute, National Institutes of Health.
Impaired kidney function	Ischaemic heart disease	Uppsala University. Sweden Uppsala Longitudinal Study of Adult Men. Chronic Kidney Disease Prognosis Consortium (CKD-PC). Chronic Kidney Disease Prognosis Consortium
Impaired kidney function	Cerebrovascular disease	GBD 2016 Impaired Kidney Function Relative Risk Meta-Analysis. National Heart, Lung, and Blood Institute, National Institutes of Health (NIH). United States Atherosclerosis
Impaired kidney function	Cerebrovascular disease	Risk in Communities Study. Bethesda, United States: National Heart, Lung, and Blood Institute, National Institutes of Health (NIH).
Impaired kidney function	Cerebrovascular disease	International Diabetes Institute (IDI). Australia Diabetes, Obesity and Lifestyle Study 1999-2000. Melbourne, Australia: International Diabetes Institute (IDI).
		National Heart, Lung, and Blood Institute, National Institutes of Health, University of California, Los Angeles (UCLA), University of Minnesota. United States Multi-Ethnic Study of Atherosclerosis First Examination
Impaired kidney function	Cerebrovascular disease	2000-2002. Bethesda, United States: National Heart, Lung, and Blood Institute, National Institutes of Health.
Impaired kidney function	Cerebrovascular disease	Uppsala University. Sweden Uppsala Longitudinal Study of Adult Men.
Impaired kidney function	Peripheral vascular disease	GBD 2016 Impaired Kidney Function Relative Risk Meta-Analysis.
Impaired kidney function	Peripheral vascular disease	National Heart, Lung, and Blood Institute, National Institutes of Health (NIH). United States Atherosclerosis Risk in Communities Study. Bethesda, United States: National Heart, Lung, and Blood Institute, National Institutes of Health (NIH).
Appendix Table 5. Epidemiolo Citations and B. Additional in	gical evidence supporting causali formation	ty between risk-outcome pairs included in the Global Burden of Disease 2017 study including A.
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Risk Impaired kidney function	Outcome Perinheral vascular disease	Citation/Note Association for Cardiac Research Rome (Italy) The Gubbio Population Study
Impaired kidney function	Perinheral vascular disease	National Heart, Lung, and Blood Institute, National Institutes of Health, University of California, Los Angeles (UCLA), University of Minnesota. United States Multi-Ethnic Study of Atherosclerosis First Examination 2000-2002 Bethesda United States: National Heart, Lung, and Blood Institute. National Institutes of Health
Imparted kidney function		Cea Soriano L, Rothenbacher D, Choi HK, García Rodríguez LA. Contemporary epidemiology of gout in the UK screar a subject of the state
	Gout	Krishnan E. Chronic kidney disease and the risk of incident gout among middle-aged men: a seven-year
Impaired kidney function	Gout	prospective observational study. Arthritis Rheum 2013; 65: 3271–8. McAdams-DeMarco MA, Maynard JW, Baer AN, Coresh J. Hypertension and the risk of incident gout in a population-based study: the atherosclerosis risk in communities cohort. J Clin Hypertens (Greenwich) 2012;
Impaired kidney function	Gout	14: 675–9. Trifirò G, Morabito P, Cavagna L, et al. Epidemiology of gout and hyperuricaemia in Italy during the years
Impaired kidney function	Gout	2005-2009: a nationwide population-based study. Ann Rheum Dis 2013; 72: 694–700. Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for- Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-
Short gestation for birth weight	Diarrheal diseases	25. Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for- Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-
Short gestation for birth weight	Lower respiratory infections	25. Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-
Short gestation for birth weight	Upper respiratory infections	25. Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for- Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-
Short gestation for birth weight	Otitis media	25. Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-26.
Short gestation for birth weight	Pneumococcal meningitis	25. Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-
Short gestation for birth weight	H influenzae type B meningitis	25. Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-25.
Short gestation for hirth weight	Other meningitis	Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-25

Appendix Table 5. Epidemiolo Citations and B. Additional in A. Citations	gical evidence supporting causali formation	ty between risk-outcome pairs included in the Global Burden of Disease 2017 study including A.
Risk	Outcome	Citation/Note
		Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low income and middle income countries: a pooled country analysis. Lancet. 2013: 382(9800): 417
Short gestation for birth weight	Encephalitis	25.
Short gestation for birth weight	Neonatal preterm birth complications	Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-25.
Short gestation for birth weight	Neonatal encephalopathy due to birth asphyxia and trauma	Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for- Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417- 25.
Short gestation for hirth weight	Neonatal sepsis and other neonatal	Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-25
Short gestation for birth weight	Hemolytic disease and other	Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for- Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013: 382(9890): 417-
Short gestation for birth weight	neonatal jaundice	25.
	Other second directory	Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-
		Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for- Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-
Short gestation for birth weight	Sudden infant death syndrome	25. Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for- Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-
Low birth weight for gestation	Diarrheal diseases	25. Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for- Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417- 25.
Low birth weight for sestation	Upper respiratory infections	Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for- Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417- 25.

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Risk	Outcome	Citation/Note Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for- Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013: 382(9890): 417-
Low birth weight for gestation	Otitis media	25.
Low birth weight for gestation	Pneumococcal meningitis	Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for- Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417- 25.
Low birth weight for gestation	H influenzae type B meningitis	Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-25.
I ow birth weight for gestation	Meningococcal meningitis	Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-25
		Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-
Low birth weight for gestation	Other meningitis	25. Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for- Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-
Low birth weight for gestation	Neonatal preterm birth	25. Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for- Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-
Low birth weight for gestation	complications Neonatal encephalopathy due to birth asphyxia and trauma	25. Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for- Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417- 25.
	Neonatal sepsis and other neonatal	Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for- Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-
Low birth weight for gestation	Infections Hemolytic disease and other neonatal jaundice	25. Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for- Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417- 25.

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Risk	Outcome	Citation/Note Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for- Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-
Low birth weight for gestation	Other neonatal disorders	25. Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-27.
Low birth weight for gestation	Sudden infant death syndrome	25. Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-25
Low birth weight and short gestation	Lower respiratory infections	Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-25.
Low birth weight and short		Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-25
Low birth weight and short	Otitis media	Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-25
Low birth weight and short gestation	Pneumococcal meningitis	Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-25.
Low birth weight and short gestation	H influenzae type B meningitis	Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-25.
Low birth weight and short gestation	Meningococcal meningitis	Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for- Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417- 25.
Low birth weight and short gestation	Other meningitis	Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-25.

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Risk	Outcome	Citation/Note
Low birth weight and short gestation	Encephalitis	Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-25.
		Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA,
Low birth weight and short gestation	Neonatal preterm birth complications	Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for- Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417- 25.
Low birth weight and short gestation	Neonatal encephalopathy due to birth asphyxia and trauma	Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-25.
Low birth weight and short gestation	Neonatal sepsis and other neonatal infections	Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-25.
Low birth weight and short gestation	Hemolytic disease and other neonatal jaundice	Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for- Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417- 25.
Low birth weight and short gestation	Other neonatal disorders	Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-25.
Low birth weight and short gestation	Sudden infant death syndrome	Katz J, Lee AC, Kozuki N, Lawn JE, Cousens S, Blencowe H, Ezzati M, Bhutta ZA, Marchant T, Willey BA, Adair L, Barros F, Baqui AH, Christian P, Fawzi W, Gonzalez R, Humphrey J, Huybregts L, Kolsteren P, Mongkolchati A, Mullany LC, Ndyomugyenyi R, Nien JK, Osrin D, Roberfroid D, Sania A, Schmiegelow C, Silveira MF, Tielsch J, Vaidya A, Velaphi SC, Victora CG, Watson-Jones D, Black RE; CHERG Small-for-Gestational-Age-Preterm Birth Working Group. Mortality risk in preterm and small-for-gestational-age infants in low-income and middle-income countries: a pooled country analysis. Lancet. 2013; 382(9890): 417-25.

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A. Citations		
Risk	Outcome	Citation/Note
6B. Supplemental information	1	

"RCTs (Number)" represents the total number of independent randomized controlled trials evaluating the relationship of each risk-outcome pair. "RCTs with significant effect in the opposite direction (%)" represents the percentage of randomized controlled trials showing a significant effect in the opposite direction. "Prospective observational studies (Number)" shows the total number of independent prospective cohort studies or non-randomized interventions evaluating the relationship of the risk-outcome pair. "Prospective observational studies with significant association in the opposite direction (%)" represents the percentage of prospective cohort studies or non-randomized interventions evaluating the relationship of the risk-outcome pair. "Prospective observational studies with significant association in the opposite direction (%)" represents the percentage of prospective cohort studies or non-randomized interventions reporting a significant association in the opposite direction. "Lower limit of RR > 1.5" shows whether the lower limit of the 95% confidence interval for the relative risk of the risk-outcome pair is greater than 1.5. "Dose-response relationship" shows whether there is any evidence of linear or non-linear dose-response relationship between the risk and the outcome. "Biologic plausibility" shows whether there is any biologic or mechanistic pathway that could potentially explain the relationship of the risk-outcome pair. "Analogy" shows whether the risk is associated with another outcome from the same category and there is evidence that it can cause the current outcome through the same pathway. The numbers in the table represent the independent RCTs and prospective observational studies evaluated the relationship between each risk-outcome pairs. If there were multiple reports from one study, they were counted as one study. Dose-response relationship was only assessed for continuous risks. To evaluate the magnitude of the effect size for continuous risks, we evaluated the RR comparing the 75th per

Unsafe water, sanitation, and	I yphoid and paratyphoid fever	i ypnoid and paratypnoid were included as outcome for unsafe water and sanitation by analogy to diarrhoeal diseases
Household air pollution from solid	Cataract	Evidence on the relationship between household air pollution and cataract was from 6 case-control and 1 cross-sectional
fuels		studies
Air pollution		The relationships of cerebrovascular disease, chronic obstructive pulmonary disease, ischaemic heart disease, and lung cancer with ambient air pollution, second-hand smoke, and active smoking were used to interpolate their relationship with household air pollution. We considered the biological pathway for health impact of all four sources to be PM2.5 exposure, with the effect size being a function of the level of PM2.5. As such, we presented data from cohorts reporting on ambient PM2.5 and the outcome was used to inform the strength of evidence for household air pollution.
Other environmental risks and	Cardiovascular diseases and chronic	The health effects of lead and sodium on cardiousscular outcomes and chronic kidney disease were assessed through
dietary risks	kidney disease	systolic blood pressure and the health effects of sugar sweetened beverages were assessed through body mass index.
Residential Radon	Tracheal, bronchus, and lung cancer	In evaluation of evidence on the relationship of residential radon and lung cancer, we excluded evidence from cohorts of miners as they were not from a representative population. Evidence on this risk-outcome pair mostly comes from case-control studies
Occupational injuries	Injuries	Evidence from International Labour Organization Safety and Health and Eurostat Safety and Health was used to establish causality between occupational injuries and injuries
Child and maternal malnutrition		Evidence on the causal relationship of childhood stunting, underweight, and wasting was from a pooled analysis of 7 prospective cohorts
Child and maternal malnutrition		For the following risk-outcome pairs, the risk factor was considered as the necessary cause: childhood underweight and protein-energy malnutrition; childhood wasting and protein-energy malnutrition; vitamin A deficiency and vitamin A deficiency; alcohol use and cirrhosis due to alcohol use; alcohol use and alcohol use disorders; drug use and cannabis use disorders; drug use and cocaine use disorders; drug use and opioid use disorders; drug use and other drug use disorders; iron deficiency and iron-deficiency anemia; unsafe sex and cervical cancer; unsafe sex and sphilis; unsafe sex and chlamydial infection; unsafe sex and trichomoniasis; unsafe sex and genital herpes; unsafe sex and other sexually transmitted diseases; high systolic blood pressure and hypertensive heart disease; high systolic blood pressure and chronic kidney disease due to hypertension; high fasting plasma glucose and chronic kidney disease mellitus; high fasting plasma glucose and diabetes mellitus; high fasting plasma glucose and diabetes mellitus; high fasting plasma glucose and specification; and specification; and specification; and specification; and glucose and diabetes mellitus; high fasting plasma glucose and chronic kidney disease
Iron deficiency	Maternal haemorrhage	Evidence on the relationship of iron deficiency with maternal haemorrhage and maternal sepsis mainly came 10 observational studies evaluating the association between low hemoglobin and maternal mortality using hospital records
Smoking, alcohol use, and high body mass index		For smoking, alcohol use, and high body mass index evidence from risk reduction trials has not been included
Smoking, alcohol use, and high body mass index	Liver cancer	Liver cancer included liver cancer due to alcohol use, hepatitis B, hepatitis C, and other causes
Smoking	Lower respiratory infections	Evidence on the relationship between smoking and lower respiratory infections comes 10 case-control or cross-sectional studies
Smoking, alcohol use	Nasopharynx cancer	The evidence on causal relationship of alcohol and smoking with nasopharynx cancer was from the studies evaluating oral cavity and pharyngeal cancers as outcome
Smoking	Bladder cancer	The evidence on causal relationship of smoking and bladder cancer was based on the studies evaluating the lower urinary tract as outcome
Smoking	Asbestosis	Asbestosis, coal workers pneumoconiosis, other pneumoconiosis, silicosis were included as outcomes for smoking as they were included in the other chronic respiratory diseases category
Alcohol use	Ischaemic heart disease, cerebrovascular disease, hypertensive heart disease, and diabetes mellitus	Alcohol was included as both a protective and harmful risk factor for ischaemic heart disease, cerebrovascular disease, hypertensive heart disease, and diabetes mellitus
Alcohol use	Cirrhosis	Cirrhosis included cirrhosis due to alcohol use, hepatitis B, hepatitis C, and other causes
Alcohol use	Self-harm	Self-harm was included as an outcome for alcohol use by analogy to injury
Alcohol use	Injuries	Injuries included pedestrian road injuries, cyclist road injuries, motorcyclist road injuries, motor vehicle road injuries, drowning, falls, fire, heat, hot substances, poisonings, unintentional firearm injuries, unintentional suffocation, other exposure to mechanical forces
Alcohol use	Interpersonal violence	Interpersonal violence included assault by firearm, sharp object, other means
Diet low in nuts and seeds	Ischaemic heart disease and diabetes mellitus	Experimental evidence on the relationship of nuts with ischaemic heart disease and diabetes mellitus come from the PREDIMED trial; a randomized trial consisting of three arms: a Mediterranean diet with extra-virgin olive oil, a Mediterranean diet with nuts, and a control diet. Given that the intake of dietary factors other than nuts changed in the intervention arms of this trial, the observed effect might be fully attributable to nuts.

Appendix Table 5. Epidemiol	ogical evidence supporting causali	ty between risk-outcome pairs included in the Global Burden of Disease 2017 study including A.
A. Citations		
Risk	Outcome	Citation/Note
Diet high in sugar sweetened		Evidence on the relationship between sugar-sweetened beverages and body mass index comes from the interventional
beverages and body mass index		and prospective observational studies evaluating the relationship of sugar-sweetened beverages with weight change among children and adults.
Diet high in sodium	Cardiovascular diseases	Evidence on the direct effect of sodium on cardiovascular disease mainly comes from prospective cohort studies. Considering that, in GBD, we have only evaluated the effect of sodium mediated through systolic blood pressure, we did not present epidemiologic evidence on the direct effect of sodium on cardiovascular disease in this table. Evidence on the effect of sodium on systolic blood pressure mostly comes from randomized controlled trials. While some cohort studies evaluated the relationship between sodium and systolic blood pressure, we did not identify a systematic evaluation of these studies.
Drug use	Hepatitis B and C	We included liver cancer due to Hepatitis B and Hepatitis C and cirrhosis due to Hepatitis B and Hepatitis C as outcomes for drug use because these were considered secondary outcomes of Hepatitis B and Hepatitis C.
Drug use, unsafe sex	HIV/AIDS	For the following risk-outcome pairs, the risk factor was considered as the sufficient cause: drug use and HIV/AIDS and unsafe sex and HIV/AIDS
Metabolic risks	Chronic kidney disease	Chronic kidney disease included chronic kidney disease due to diabetes mellitus, hypertension, glomerulonephritis, or other causes
High fasting plasma glucose	Cerebrovascular disease, chronic kidney disease, ischaemic heart disease	Evidence on the relationship of high fasting plasma glucose with stroke (DECODE, APCSC, ERFC); chronic kidney disease (APCSC), and ischaemic heart disease (DECODE, APCSC, ERFC) was from pooled analysis of cohorts
High systolic blood pressure	Atrial fibrillation and flutter, peripheral vascular disease	Evidence on the relationship of high systolic blood pressure with atrial fibrillation and peripheral vascular disease was from two pooled cohort analysis (APCSC and PSC)
High systolic blood pressure	Rheumatic heart disease, cardiomyopathy and myocarditis, aortic aneurysm, endocarditis, and other cardiovascular diseases	Evidence on the relationship of high systolic blood pressure with rheumatic heart disease, cardiomyopathy and myocarditis, aortic aneurysm, endocarditis, and other cardiovascular diseases came from a pooled cohort analysis (PSC)
High body-mass index	Ischaemic heart disease	Evidence on the relationship of high body-mass index with ischaemic heart disease (APCSC, ERFC, PSC) and stroke (ischaemic: APCSC, ERFC, PSC; hemorrhagic: PSC and ERFC) came from three pooled cohort analysis
High body-mass index	Diabetes mellitus, hypertensive heart disease	Evidence on the relationship of high body-mass index with diabetes mellitus and hypertensive heart disease came from two pooled cohort analysis (APCSC and PSC)
High body-mass index	Chronic kidney disease	Evidence on the relationship of high body-mass index with chronic kidney disease was from a pooled cohort analysis (PSC)
High LDL cholesterol	Ischaemic heart disease, ischaemic stroke	Evidence on the relationship of high LDL cholesterol with ischaemic heart disease and ischaemic stroke came from two pooled cohort analysis (APCSC and PSC)
Impaired kidney function	Tuberculosis	Glycemic Control and the Risk of Tuberculosis: A Cohort Study.

Appendix Table 6a. Relative ris	ks used by age and sex for each or	tcome for all risk factors excep	ot for ambient air	pollution alcohol,	and smoking.									,	Ages											
			All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Risk - Outcome Unsafe water source	Category / Units	Morbidity / Mortality Sex		anj -				,																		
Displayed Surger	Driver and estimated	Push Push		11.501	11.501	11.501	11.501	11.501	11.501	11.501	11.501	11.501	11.501	11.501	11.501	11.501	11.501	11.501	11.501	11.501	11.501	11.501	11.501	11.501	11.501	11.501
	Chiliporte, unicated			(2.761 to 31.282) 7.914																						
Diamisea diseises	Unimproved, chiorinated	Boll Boll		(1.971 to 21.188) 4.789																						
Diarrhoral diseases	Unimproved, filter	Both Both	-	(1.204 to 12.752)																						
Diarrhoeal diseases	Improved, untreated	Both Both	h	9.428 (2.782 to 23.028)	(2.782 to 23.028)	9.428 (2.782 to 23.028)	9.428 (2.782 to 23.028)	9.428 (2.782 to 23.028)	9.428 (2.782 to 23.028)	9.428 (2.782 to 23.028)	9.428 (2.782 to 23.028)	9.428 (2.782 to 23.028)	9.428 (2.782 to 23.028)	9.428 (2.782 to 23.028)	(2.782 to 23.028)	9.428 (2.782 to 23.028)	9.428 (2.782 to 23.028)	9.428 (2.782 to 23.028)	9.428 (2.782 to 23.028)	9.428 (2.782 to 23.028)	9.428 (2.782 to 23.028)	(2.782 to 23.028)	(2.782 to 23.028)	9.428 (2.782 to 23.028)	9.428 (2.782 to 23.028)	(2.782 to 23.028)
Diarrhoeal diseases	Improved, chlorinated	Both Both	-	6.488 (1.997 to 15.677)	6.438 (1.997 to 15.677)	6.488 (1.997 to 15.677)	6.488 (1.997 to 15.677)	6.488 (1.997 to 15.677)	6.438 (1.997 to 15.677)	6.488 (1.997 to 15.677)	6.488 (1.997 to 15.677)	6.488 (1.997 to 15.677)	6.488 (1.997 to 15.677)													
Diarrhoeal diseases	Improved, filtered	Both Both	h	3.926 (1.22 to 9.401)	3.926 (1.22 to 9.401)	3.926 (1.22 to 9.401)	(1.22 to 9.401)	3.926 (1.22 to 9.401)	3.926 (1.22 to 9.401)	3.926 (1.22 to 9.401)	3.926 (1.22 to 9.401)	3.926 (1.22 to 9.401)	3.926 (1.22 to 9.401)	3.926 (1.22 to 9.401)	3.926 (1.22 to 9.401)	3.926 (1.22 to 9.401)	3.926 (1.22 to 9.401)	3.926 (1.22 to 9.401)	3.926 (1.22 to 9.401)	3.926 (1.22 to 9.401)	3.926 (1.22 to 9.401)	3.926 (1.22 to 9.401)	3.926 (1.22 to 9.401)	3.926 (1.22 to 9.401)	3.926 (1.22 to 9.401)	3.926 (1.22 to 9.401)
Diarrhoeal diseases	Piped, untreated	Both Both	h	8.431 (2.533 to 20.446)																						
Diamboral diseases	Piped, chlorinated	Both Both	-	5.802 (1.807 to 13.843)																						
Diarrhoeal diseases	Piped, filtered	Both Both		3.511 (1.107 to 8.331)																						
Diarrhoeal diseases	High quality (HQ) piped, untreated	Both Both	h	2.401 (2.037 to 2.818)																						
Diarrhocal diseases	HQ piped, chlorinated	Both Both		1.653 (1.56 to 1.748)																						
Diarrhoeal diseases	HQ piped, filtered	Both Both		1.0	1.0	1.0	1.0 (1.0 to 1.0)	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
Unsafe sanitation																										
Diarrhoeal diseases	Unimproved & untreated	Both Both		3.242	3.242	3.242	3.242	3.242	3.242	3.242	3.242	3.242	3.242	3.242	3.242	3.242	3.242	3.242	3.242	3.242	3.242	3.242	3.242	3.242	3.242	3.242
Diarrhogal diseases	Improved	Both Both		2.595	2.595	2.595	(2.528 to 4.067) 2.595	2.595	2.595	2.595	2.595	2.595	2.595	(2.52810.4.067) 2.595	2.595	(2.528 10 4.067) 2.595	2.595	2.595	2.595	(2.528 10 4.067) 2.595	2.595	2.595	2.595	2.595	2.595	2.595
Distributed distances	Sensore	Both Both		(2.044 to 3.285) 1.0																						
No accurr to handunching fa	scility		-	(1.0 to 1.0)																						
to acces to handwashing ha				1.191	1.191	1.191	1.191	1.191	1.191	1.191	1.191	1.191	1.191	1.191	1.191	1.191	1.191	1.191	1.191	1.191	1.191	1.191	1.191	1.191	1.191	1.191
Lower respiratory infections	No handwashing w/soap & water	Both Both	•	(1.119 to 1.266)																						
Lower respiratory infections	Handwashing wiscop & water	Both Both	h	(1.0 to 1.0)																						
Diarrhoeal diseases	No handwashing w/soap & water	Both Both	-	(1.32 to 2.668)	1.508 (1.32 to 2.668)	1.908 (1.32 to 2.668)	1.908 (1.32 to 2.668)	1.908 (1.32 to 2.668)	1.308 (1.32 to 2.668)	(1.32 to 2.668)	(1.32 to 2.668)	1.908 (1.32 to 2.668)	1.908 (1.32 to 2.668)	1.908 (1.32 to 2.668)	1.908 (1.32 to 2.668)	1.308 (1.32 to 2.668)	(1.32 to 2.668)	(1.32 to 2.668)	1.908 (1.32 to 2.668)	1.908 (1.32 to 2.668)	(1.32 to 2.668)	1.908 (1.32 to 2.668)				
Diarrhoeal diseases	Handwashing w/soap & water	Both Both	h	1.0 (1.0 to 1.0)																						
Household air pollution from	m solid fuels																									
Cataract	Exposed	morbidity Both	h 2.47 (1.61 to 3.73)																							
Ambient ozone pollution																										
Chronic obstructive pulmonary disease	y 10 ppb	Mortality Both	-									1.06 (1.017 to 1.101)														
Residential radon																										
Tracheal, bronchus, and lung ca	ancer Bq/m3	Both Both	h.	1.002 (1.0 to 1.003)																						
Lead exposure in blood																										
IQ shift	10 µg/dL	Morbidity Both	4.688 (1.719 to 7.656)																							
IQ shift	12 µg/dL	Morbidity Both	5.014																							
IQ shift	15 µg/dL	Morbidity Both	5.42																							
IO shift	2 unidi.	Morbidity Both	0.0																							
10 shift	20 unidL	Morbidity Botl	5.952																							
10.114	25	Madridian Real	(2.183 to 9.721) 6.37																							
10.1/2	2.5 pg at.	Matrice Date	(2.336 to 10.403) 6.713																							
aj sme	so py at.	Morekary Bon	(2.462 to 10.965) 7.006																							
1Q shift	35 µg/dL	Morbidity Boll	h (2.56) to 11.442)																							
IQ shift	4 µg/dl.	Morbidity Both	(1.154 to 5.139)																							
IQ shift	40 µg/dL	Morbidity Both	(2.662 to 11.857)																							
IQ shift	6 µg/dL	Morbidity Both	h (1.395 to 6.213)																							
IQ shift	8 µg/dl.	Morbidity Both	4.296 (1.575 to 7.016)																							
Lead exposure in bone																										<u> </u>
Rheamatic heart disease	10 µg/g	Both Both	•									1.03 (1.01 to 1.052)	1.024 (1.01 to 1.04)	1.017 (1.008 to 1.028)	1.013 (1.005 to 1.022)	1.012 (1.006 to 1.019)	1.011 (1.006 to 1.017)	1.01 (1.006 to 1.016)	1.009 (1.005 to 1.014)	1.008 (1.003 to 1.014)	1.007 (1.003 to 1.013)	1.007 (1.004 to 1.013)	1.006 (1.002 to 1.015)	1.006 (1.002 to 1.015)	1.006 (1.002 to 1.015)	1.006 (1.002 to 1.015)
Ischaemic heart disease	10 µg/g	Both Both	h									1.042 (1.022 to 1.06)	1.037 (1.023 to 1.049)	1.032 (1.023 to 1.04)	1.028 (1.021 to 1.036)	1.026 (1.02 to 1.033)	1.024 (1.02 to 1.03)	1.023 (1.019 to 1.026)	1.021 (1.018 to 1.025)	1.019 (1.014 to 1.023)	1.018 (1.012 to 1.022)	1.016 (1.012 to 1.021)	1.014 (1.008 to 1.022)	1.014 (1.008 to 1.022)	1.014 (1.008 to 1.022)	1.014 (1.008 to 1.022)
Ischaemic stroke	$10 \mu g/g$	Both Both										1.038 (1.021 to 1.06)	1.036 (1.022 to 1.051)	1.033 (1.021 to 1.044)	1.03 (1.019 to 1.042)	1.028 (1.019 to 1.037)	1.026 (1.019 to 1.033)	1.024 (1.018 to 1.029)	1.021 (1.016 to 1.026)	1.019 (1.012 to 1.025)	1.017 (1.01 to 1.023)	1.015 (1.01 to 1.02)	1.011 (1.006 to 1.019)	1.011 (1.006 to 1.019)	1.011 (1.006 to 1.019)	1.011 (1.006 to 1.019)
Intracerebral hemorrhage	10 µg/g	Both Both	h									1.047 (1.027 to 1.068)	1.045 (1.029 to 1.061)	1.042 (1.029 to 1.057)	1.039 (1.025 to 1.052)	1.036 (1.024 to 1.047)	1.032 (1.023 to 1.041)	1.028 (1.021 to 1.035)	1.024 (1.018 to 1.03)	1.02 (1.012 to 1.027)	1.017 (1.009 to 1.025)	1.017 (1.011 to 1.023)	1.015 (1.007 to 1.026)	1.015 (1.007 to 1.026)	1.015 (1.007 to 1.026)	1.015 (1.007 to 1.026)
Subarachnoid hemorrhage	10 µg/g	Both Both										1.047 (1.027 to 1.068)	1.045 (1.029 to 1.061)	1.042 (1.029 to 1.057)	1.039 (1.025 to 1.052)	1.036 (1.024 to 1.047)	1.032 (1.023 to 1.041)	1.028 (1.021 to 1.035)	1.024 (1.018 to 1.03)	1.02 (1.012 to 1.027)	1.017 (1.009 to 1.025)	1.017 (1.011 to 1.023)	1.015 (1.007 to 1.026)	1.015 (1.007 to 1.026)	1.015 (1.007 to 1.026)	1.015 (1.007 to 1.026)
Hypertensive heart disease	10 µg/g	Both Both										1.066 (1.038 to 1.09)	1.066 (1.038 to 1.091)	1.065 (1.037 in 1.094)	1.063 (1.035 to 1.091)	1.058	1.052 (1.036 to 1.079)	1.046	1.04 (1.023 to 1.073)	1.033	1.03 (1.003 to 1.072)	1.031 (1.005 to 1.071)	1.033	1.033 (1.006 to 1.075)	1.033 (1.006 to 1.075)	1.033
Non-rheumatic calcific acetic v	valve 10 µg/g	Both Both										1.035	1.029	1.023	1.019	1.018	1.016	1.015	1.014	1.012	1.011	1.01	1.007	1.007	1.007	1.007
other cardiomyopathy	10 µg/g	Both Both										1.035	1.029	1.023	1.019	(101210-1023)	1.016	1.012101.018)	1.014	1.012	1.011	1.01	1.007	1.007	1.007	1.007
Atrial fibrillation and floater	10 µg/g	Both Roal										(1.015 to 1.055)	(1.016 to 1.044) 1.03	(1.015 to 1.051) 1.025	(1.013 to 1.025) 1.022	(1.012 to 1.023)	(1.012 to 1.02)	(1.012 to 1.018)	(1.01 to 1.016)	(1.008 to 1.015) 1.015	(1.007 to 1.014) 1.013	(1.007 to 1.013) 1.012	(1.004 to 1.013) 1.008	(1.004 to 1.013) 1.008	(1.00+101.013)	(1.00+10.1.013)
Antic		Book D.										(1.018 to 1.056) 1.027	(1.02 to 1.044) 1.024	(1.021 to 1.031) 1.02	(1.018 to 1.025) 1.018	(1.017 to 1.023) 1.017	(1.017 to 1.021) 1.016	(1.016 to 1.019) 1.015	(1.014 to 1.018) 1.014	(1.013 to 1.017) 1.012	(1.011 to 1.015) 1.011	(1.01 to 1.013) 1.01	(1.005 to 1.01) 1.007	(1.005 to 1.01) 1.007	(1.005 to 1.01) 1.007	(1.005 to 1.01) 1.007
PROFESSION AND AND AND AND AND AND AND AND AND AN	10 (Boll Boll										(1.014 to 1.048) 1.034	(1.016 to 1.037) 1.025	(1.016 to 1.026) 1.014	(1.013 to 1.023) 1.008	(1.013 to 1.021) 1.008	(1.013 to 1.019) 1.008	(1.012 to 1.017) 1.009	(1.011 to 1.016) 1.009	(1.009 to 1.015) 1.009	(1.008 to 1.014) 1.009	(1.007 to 1.013) 1.008	(1.004 to 1.01) 1.006	(1.004 to 1.01) 1.006	(1.004 to 1.01) 1.006	(1.004 to 1.01) 1.006
Perspheral vascular disease	10 µĝ/g	Both Both										(1.011 to 1.056)	(1.012 to 1.039)	(1.01 to 1.018) 1.023	(1.001 to 1.014)	(1.003 to 1.013)	(1.004 to 1.012)	(1.005 to 1.012) 1.015	(1.006 to 1.011) 1.014	(1.007 to 1.012)	(1.006 to 1.011)	(1.006 to 1.01) 1.01	(1.003 to 1.009) 1.007	(1.003 to 1.009) 1.007	(1.003 to 1.009) 1,007	(1.003 to 1.009) 1.007
Endocarditis	10 µg/g	Both Both	`									(1.015 to 1.055)	(1.016 to 1.044)	(1.015 to 1.031)	(1.013 to 1.025)	(1.012 to 1.023)	(1.012 to 1.02)	(1.012 to 1.018)	(1.01 to 1.016)	(1.008 to 1.015)	(1.007 to 1.014)	(1.007 to 1.013)	(1.004 to 1.013)	(1.004 to 1.013)	(1.004 to 1.013)	(1.004 to 1.013)
Other cardiovascular and circul diseases	10 µg/g	Both Both	*									(1.034 (1.018 to 1.055)	1.03 (1.02 to 1.043)	(1.025 (1.021 to 1.03)	(1.022 (1.019 to 1.025)	(1.018 to 1.023)	(1.019 (1.017 to 1.021)	(1.018 (1.016 to 1.019)	1.016 (1.015 to 1.018)	(1.014 (1.013 to 1.016)	1.013 (1.011 to 1.015)	1.012 (1.01 to 1.013)	1.008 (1.006 to 1.011)	1.008 (1.006 to 1.011)	(1.006 to 1.011)	1.008 (1.006 to 1.011)
Chronic kidney disease due to diabetes mellitus type 1	$10 \mu g/g$	Both Both	*									1.015 (1.01 to 1.021)														
Chronic kidney disease due to diabetes mellitus type 2	$10 \mu g/g$	Both Both	- I									1.015 (1.01 to 1.021)														

Appendix Table 6a. Relative risks used by age and sex for each outcome for all	risk factors excep	t for ambient air pollution a	cohol, and smokir	ag.								A	ges											
Pick - Outcome Category / Unite Markidi	v/Mortality Say	All-age 0-6 d	ays 7-27 d	ays 28-	-364 days 1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Chronic Lidney discuse due to brostrension 10 µg/g	Both Both		I		1					1.015 (1.01 to 1.02)														
Chronic kidney diwasse due to ntomerulomerubritis 10 µg/g	Both Both									1.015 (1.01 to 1.02)	1.015	1.015 (1.01 to 1.02)												
Chronic bidney disease due to other and unspecified causes	Both Both									1.015 (1.01 to 1.021)														
Occupational exposure to asbestos																								
Larynx cancer High exposure	Both Males		I		1			1.38 (1.188 to 1.612)																
Larynx cancer High exposure	Both Female	58						1.385 (1.187 to 1.598)																
Larynx cancer Low exposure	Both Males							1.0 (1.0 to 1.0)																
Larynx cancer Low exposure	Both Female	58						1.0 (1.0 to 1.0)																
Tracheal, bronchus, and hung cancer High exposure	Both Males							2.279 (1.741 to 2.936)																
Tracheal, bronchus, and hung cancer High exposure	Both Female	58						1.875 (1.589 to 2.208)																
Tracheal, bronchus, and lung cancer Low exposure	Both Males							1.0 (1.0 to 1.0)																
Tracheal, bronchus, and lung cancer Low exposure	Both Female							1.0 (1.0 to 1.0)																
Ovarian cancer High exposure	Both Both							1.811 (1.385 to 2.306)																
Ovarian cancer Low exposure	Both Both							1.0 (1.0 to 1.0)																
Occupational exposure to arsenic																								
Tracheal, bronchus, and lung cancer High exposure	Both Both							2.061 (1.521 to 2.553)																
Tracheal, bronchus, and lung cancer Low exposure	Both Both							1.749 (0.698 to 2.775)																
Tracheal, bronchus, and lung cancer No exposure	Both Both							1.0 (1.0 to 1.0)																
Occupational exposure to benzene																								
Acute lymphoid leukaemia High exposure	Both Both							2.623 (1.222 to 3.975)																
Acute lymphoid leukaemia Low exposure	Both Both							1.626 (0.998 to 2.256)																
Acute lymphoid leukaemia No exposure	Both Both							1.0 (1.0 to 1.0)																
Chronic lymphoid kuukaemia High exposure	Both Both							2.623 (1.222 to 3.975)																
Chronic lymphoid leukaemia Low exposure	Both Both							1.626 (0.998 to 2.256)																
Chronic lymphoid leukaemia No exposure	Both Both							1.0 (1.0 to 1.0)																
Acute myeloid leukaemia High exposure	Both Both							2.623 (1.222 to 3.975)																
Acute myeloid leukaemia Low exposure	Both Both							1.626 (0.998 to 2.256)																
Acute myeloid leukaemia No exposure	Both Both							1.0 (1.0 to 1.0)																
Chronic myeloid leukaemia High exposure	Both Both							2.623 (1.222 to 3.975)																
Chronic myeloid leukaemia Low exposure	Both Both							1.626 (0.998 to 2.256)																
Chronic myeloid leukaemia No exposure	Both Both							1.0 (1.0 to 1.0)																
Other leukaemia High exposure	Both Both							2.623 (1.222 to 3.975)																
Other leukaemia Low exposure	Both Both							1.626 (0.998 to 2.256)																
Other leukaemia No exposure	Both Both							1.0 (1.0 to 1.0)																
Occupational exposure to beryllium																								
Tracheal, bronchus, and lung cancer High exposure	Both Males							1.174 (1.086 to 1.269)	1.169 (1.065 to 1.276)	1.17 (1.073 to 1.274)	1.169 (1.073 to 1.269)	1.172 (1.081 to 1.271)	1.168 (1.07 to 1.276)	1.171 (1.071 to 1.274)	1.171 (1.075 to 1.273)	1.17 (1.073 to 1.271)	1.171 (1.072 to 1.271)	1.172 (1.081 to 1.276)	1.171 (1.076 to 1.27)	1.174 (1.079 to 1.276)	1.171 (1.072 to 1.273)	1.171 (1.072 to 1.273)	1.171 (1.072 to 1.273)	1.171 (1.072 to 1.273)
Tracheal, bronchus, and hung cancer High exposure	Both Female							1.17 (1.082 to 1.262)	1.17 (1.076 to 1.277)	1.169 (1.072 to 1.277)	1.17 (1.077 to 1.275)	1.172 (1.075 to 1.274)	1.171 (1.078 to 1.276)	1.173 (1.074 to 1.276)	1.172 (1.074 to 1.273)	1.173 (1.081 to 1.277)	1.169 (1.074 to 1.269)	1.173 (1.077 to 1.282)	1.167 (1.069 to 1.269)	1.169 (1.073 to 1.274)	1.171 (1.068 to 1.279)	1.171 (1.068 to 1.279)	1.171 (1.068 to 1.279)	1.171 (1.068 to 1.279)
Tracheal, bronchus, and hung cancer Low exposure	Both Males							1.0 (1.0 to 1.0)																
Tracheal, bronchus, and lung cancer Low exposure	Both Female							1.0 (1.0 to 1.0)																
Tracheal, bronchus, and lung cancer No exposure	Both Males							1.0 (1.0 to 1.0)																
Tracheal, bronchus, and hung cancer No exposure	Both Female							1.0 (1.0 to 1.0)																
Occupational exposure to cadmium																								
Tracheal, bronchus, and lung cancer High exposure	Both Males							1.192 (1.097 to 1.292)	1.188 (1.083 to 1.287)	1.19 (1.102 to 1.295)	1.191 (1.092 to 1.304)	1.19 (1.091 to 1.298)	1.19 (1.092 to 1.297)	1.192 (1.096 to 1.298)	1.193 (1.096 to 1.296)	1.19 (1.089 to 1.293)	1.192 (1.089 to 1.297)	1.193 (1.088 to 1.3)	1.187 (1.095 to 1.29)	1.191 (1.095 to 1.289)	1.19 (1.088 to 1.296)	1.19 (1.088 to 1.296)	1.19 (1.088 to 1.296)	1.19 (1.088 to 1.296)
Tracheal, bronchus, and hung cancer High exposure	Both Female	58						1.191 (1.087 to 1.295)	1.191 (1.099 to 1.29)	1.188 (1.093 to 1.293)	1.19 (1.092 to 1.302)	1.191 (1.089 to 1.292)	1.194 (1.098 to 1.297)	1.188 (1.091 to 1.3)	1.19 (1.095 to 1.291)	1.19 (1.096 to 1.294)	1.189 (1.095 to 1.29)	1.192 (1.095 to 1.296)	1.192 (1.09 to 1.298)	1.193 (1.097 to 1.297)	1.193 (1.1 to 1.301)	1.193 (1.1 to 1.301)	1.193 (1.1 to 1.301)	1.193 (1.1 to 1.301)
Tracheal, bronchus, and lung cancer Low exposure	Both Males							1.0 (1.0 to 1.0)																
Tracheal, bronchus, and hung cancer Low exposure	Both Female							1.0 (1.0 to 1.0)																
Tracheal, bronchus, and hung cancer No exposure	Both Males							1.0 (1.0 to 1.0)																
Tracheal, bronchus, and lung cancer No exposure	Both Female							1.0 (1.0 to 1.0)																
Occupational exposure to chromium																								
Tracheal, bronchus, and hung cancer High exposure	Both Males							1.179 (1.114 to 1.245)	1.181 (1.117 to 1.25)	1.181 (1.118 to 1.249)	1.18 (1.117 to 1.244)	1.183 (1.117 to 1.249)	1.18 (1.117 to 1.239)	1.18 (1.117 to 1.249)	1.182 (1.12 to 1.246)	1.181 (1.119 to 1.246)	1.179 (1.115 to 1.242)	1.182 (1.121 to 1.247)	1.18 (1.118 to 1.248)	1.18 (1.113 to 1.245)	1.181 (1.117 to 1.244)	1.181 (1.117 to 1.244)	1.181 (1.117 to 1.244)	1.181 (1.117 to 1.244)
Tracheal, bronchus, and lung cancer High exposure	Both Female	x.						1.179 (1.116 to 1.247)	1.18 (1.115 to 1.248)	1.18 (1.115 to 1.244)	1.179 (1.118 to 1.243)	1.18 (1.116 to 1.248)	1.181 (1.115 to 1.245)	1.181 (1.119 to 1.25)	1.181 (1.118 to 1.245)	1.181 (1.117 to 1.248)	1.181 (1.117 to 1.244)	1.179 (1.121 to 1.245)	1.181 (1.116 to 1.254)	1.181 (1.116 to 1.245)	1.179 (1.113 to 1.247)	1.179 (1.113 to 1.247)	1.179 (1.113 to 1.247)	1.179 (1.113 to 1.247)
Tracheal, bronchus, and lung cancer Low exposure	Both Males							1.0 (1.0 to 1.0)																
Tracheal, bronchus, and lung cancer Low exposure	Both Female							1.0 (1.0 to 1.0)																
Tracheal, bronchus, and lung cancer No exposure	Both Male							1.0 (1.0 to 1.0)																
Tracheal, bronchas, and lung cancer No exposure	Both Female	5						1.0 (1.0 to 1.0)																
Occupational exposure to diesel engine exhaust																								
Tracheal, bronchus, and hung cancer High exposure	Both Males							1.469	1.477	1.473	1.474 (1.293 to 1.675)	1.472	1.47	1.475 (1.3 to 1.667)	1.469	1.474	1.477 (1.202 = 1.471)	1.477	1.476	1.473	1.477	1.477	1.477	1.477
- I.		1						(1.4.79 10 1.056)	(1.591-0 1.003)	(1.27.00 1.008)	(1.477-00 1.073)	(1	(1.379 10 1.002)	(1.001.007)	(1.701 10 1.044)	(*	(1.474.001.071)	(1	(1.70= 00 1.009)	(1.009)	(1.00a s0 1.000)	(1.704 40 1.000)	(1.004 00 1.000)	(1.000)

Appendix Table 6a. Relative risks used by age and sex for each of	utcome for all risk factors ex	cept for ambient air	pollution alcohol,	and smoking.									А	ges											
Risk - Outcome Category / Units	Morbidity / Mortality	Sex All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Tracheal, bronchus, and lung cancer High exposure	Both Fe	emales							1.473 (1.287 to 1.682)	1.476 (1.303 to 1.681)	1.469 (1.288 to 1.67)	1.467 (1.282 to 1.67)	1.473 (1.288 to 1.66)	1.475 (1.294 to 1.68)	1.477 (1.3 to 1.679)	1.475 (1.302 to 1.674)	1.476 (1.292 to 1.68)	1.481 (1.309 to 1.681)	1.478 (1.295 to 1.683)	1.473 (1.291 to 1.661)	1.473 (1.29 to 1.676)	1.476 (1.289 to 1.662)	1.476 (1.289 to 1.662)	1.476 (1.289 to 1.662)	1.476 (1.289 to 1.662)
Tracheal, bronchus, and lung cancer Low exposure	Both	Males							1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Tracheal, bronchus, and lung cancer Low exposure	Both Fe	emales							1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Tracheal, bronchus, and lung cancer No exposure	Both 3	Malex							(1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	1.0 (1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	1.0 (1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	(1.0 to 1.0)
Tracheal, bronchus, and lung cancer No exposure	Both Fe	emales							1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Occupational exposure to formaldehyde									2 222	2 294	2 204	2.211	223	2.241	2.265	2.198	2 232	2.215	2.251	2.233	2.234	2 204	2 204	2 204	2 204
Nasopharynx cancer High exposure	Both	Malex							(1.026 to 4.233) 2.202	(1.056 to 4.409) 2.246	(1.023 to 4.036) 2.227	(0.994 to 4.221) 2.269	(1.042 to 4.13) 2.264	(1.077 to 4.068) 2.217	(1.056 to 4.33) 2.201	(1.041 to 4.081) 2.221	(1.104 to 4.091) 2.225	(1.022 to 4.191) 2.276	(1.056 to 4.236) 2.261	(1.022 to 4.329) 2.258	(1.02 to 4.325) 2.241	(1.064 to 4.14) 2.229	(1.054 to 4.14) 2.229	(1.064 to 4.14) 2.229	(1.064 to 4.14) 2.229
Neuropearynx cancer rigin exposure	Bollin Pi	emaaes Malar							(1.04 to 4.059) 1.0	(1.039 to 4.184) 1.0	(1.05 to 4.207) 1.0	(1.084 to 4.182) 1.0	(1.069 to 4.385) 1.0	(1.041 to 4.21) 1.0	(1.035 to 4.162) 1.0	(1.035 to 4.234) 1.0	(1.025 to 4.212) 1.0	(1.072 to 4.204) 1.0	(1.04 to 4.208) 1.0	(1.042 to 4.371) 1.0	(1.031 to 4.156) 1.0	(1.025 to 4.237) 1.0	(1.025 to 4.237) 1.0	(1.025 to 4.237) 1.0	(1.025 to 4.237) 1.0
Namehouse anter Low exposure	Bolh P								(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0
Nuorhorezzarezz No exposure	Both	Malex							(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0
Nasonharyux cancer No exposure	Both Fu	emales							(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0
Acute hyperboid leakacraia High exposure	Both	Males							(1.0 to 1.0) 1.483	(1.0 to 1.0) 1.479	(1.0 to 1.0) 1.474	(1.0 to 1.0) 1.48	(1.0 to 1.0) 1.479	(1.0 to 1.0) 1.467	(1.0 to 1.0) 1.481	(1.0 to 1.0) 1.49	(1.0 to 1.0) 1.47	(1.0 to 1.0) 1.487	(1.0 to 1.0) 1.485	(1.0 to 1.0) 1.48	(1.0 to 1.0) 1.482	(1.0 to 1.0) 1.488	(1.0 to 1.0) 1.488	(1.0 to 1.0) 1.488	(1.0 to 1.0) 1.488
Acute lymphoid leukaemia High exposure	Both Fe	emales							1.485	1.485	1.479	1.471	1.469	1.486	1.485	1.481	1.48	(1.2 to 1.846)	1.473	1.471	(1.202101.839)	(1.198101.827)	(1.198 15 1.827)	1.464	1.464
Acute lymphoid leukaemia Low exposure	Both	Males							(1.199 to 1.843) 1.0 (1.0 to 1.0)	1.0	1.0	1.0	1.0	1.0	1.0	1.0	(1.199101.814) 1.0 (1.0 m 1.0)	(1.184 05 1.812) 1.0 (1.0 m 1.0)	1.0	1.0	(1.209101.859) 1.0 (1.0 m 1.0)	1.0	1.0	(1.18 15 1.788) 1.0 (1.0 to 1.0)	1.0
Acute lymphoid leukaemia Low exposure	Both Fe	emales							1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
Acute lymphoid leukaemia No exposure	Both	Malex							1.0 (1.0 to 1.0)	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
Acute lymphoid leukaemia No exposure	Both Fe	emales							1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0
Chronic lymphoid leukaemia High exposure	Both	Malex							1.483 (1.191 to 1.818)	1.479 (1.183 to 1.83)	1.474	1.48 (1.193 to 1.827)	1.479 (1.197 to 1.804)	1.467 (1.174 to 1.843)	1.481 (1.178 to 1.839)	1.49 (1.199 to 1.831)	1.47 (1.197 to 1.842)	1.487 (1.2 to 1.846)	1.485 (1.188 to 1.803)	1.48 (1.18 to 1.836)	1.482 (1.202 to 1.839)	1.488 (1.198 to 1.827)	1.488 (1.198 m 1.827)	1.488 (1.198 to 1.827)	1.488 (1.198 to 1.827)
Chronic lymphoid leukaemia High exposure	Both Fe	emales							1.485 (1.199 to 1.845)	1.485 (1.183 to 1.855)	1.479 (1.184 to 1.844)	1.471 (1.181 to 1.774)	1.469 (1.183 to 1.819)	1.486 (1.196 to 1.823)	1.485 (1.19 to 1.848)	1.481 (1.191 to 1.856)	1.48 (1.199 to 1.814)	1.47 (1.184 to 1.812)	1.473 (1.19 to 1.814)	1.471 (1.194 to 1.789)	1.49 (1.209 to 1.859)	1.464 (1.18 to 1.788)	1.464 (1.18 to 1.788)	1.464 (1.18 to 1.788)	1.464 (1.18 to 1.788)
Chronic lymphoid leukaemia Low exposure	Both	Males							1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Chronic lymphoid leukaemia Low exposure	Both Fe	emales							1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Chronic lymphoid leukaemia No exposure	Both	Malex							1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Chronic lymphoid leukaemia No exposure	Both Fe	emales							1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Acute myeloid leukaemia High exposure	Both	Males							1.483 (1.191 to 1.818)	1.479 (1.183 to 1.83)	1.474 (1.182 to 1.815)	1.48 (1.193 to 1.827)	1.479 (1.197 to 1.804)	1.467 (1.174 to 1.843)	1.481 (1.178 to 1.839)	1.49 (1.199 to 1.831)	1.47 (1.197 to 1.842)	1.487 (1.2 to 1.846)	1.485 (1.188 to 1.803)	1.48 (1.18 to 1.836)	1.482 (1.202 to 1.839)	1.488 (1.198 to 1.827)	1.488 (1.198 to 1.827)	1.488 (1.198 to 1.827)	1.488 (1.198 to 1.827)
Acute myeloid leukaemia High exposure	Both Fe	emales							1.485 (1.199 to 1.845)	1.485 (1.183 to 1.855)	1.479 (1.184 to 1.844)	1.471 (1.181 to 1.774)	1.469 (1.183 to 1.819)	1.486 (1.196 to 1.823)	1.485 (1.19 to 1.848)	1.481 (1.191 to 1.856)	1.48 (1.199 to 1.814)	1.47 (1.184 to 1.812)	1.473 (1.19 to 1.814)	1.471 (1.194 to 1.789)	1.49 (1.209 to 1.859)	1.464 (1.18 to 1.788)	1.464 (1.18 to 1.788)	1.464 (1.18 to 1.788)	1.464 (1.18 to 1.788)
Acute myeloid leukaemia Lew exposure	Both	Males							1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Acute myeloid leukaemia Low exposure	Both Fe	emales							1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Acute myeloid leukaemia No exposure	Both	Males							1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Acute myeloid leukaemia No exposure	Both Fe	emales							1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Chronic myeloid leukaemia High exposure	Both	Males							1.483 (1.191 to 1.818)	1.479 (1.183 to 1.83)	1.474 (1.182 to 1.815)	1.48 (1.193 to 1.827)	1.479 (1.197 to 1.804)	1.467 (1.174 to 1.843)	1.481 (1.178 to 1.839)	1.49 (1.199 to 1.831)	1.47 (1.197 to 1.842)	1.487 (1.2 to 1.846)	1.485 (1.188 to 1.803)	1.48 (1.18 to 1.836)	1.482 (1.202 to 1.839)	1.488 (1.198 to 1.827)	1.488 (1.198 to 1.827)	1.488 (1.198 to 1.827)	1.488 (1.198 to 1.827)
Chronic myeloid leukaemia High exposure	Both Fe	emales							(1.199 to 1.845)	(1.183 to 1.855)	(1.184 to 1.844)	(1.181 to 1.774)	(1.183 to 1.819)	1.486 (1.196 to 1.823)	1.485 (1.19 to 1.848)	(1.191 to 1.856)	1.48 (1.199 to 1.814)	1.47 (1.184 to 1.812)	(1.19 to 1.814)	1.471 (1.194 to 1.789)	(1.209 to 1.859)	(1.18 to 1.788)	1.464 (1.18 to 1.788)	1.464 (1.18 to 1.788)	1.464 (1.18 to 1.788)
Chronic myeloid leukaemia Low exposure	Both	Males							(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)
Chronic myeloid leukaemia Low exposure	Both Fe	emales							(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)
Chronic myeloid leukaemia No exposure	Both	Malex							(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0)
Chronic myeloud leukaemua No exposure	Both Fo	emakes							(1.0 to 1.0) 1.483	(1.0 to 1.0) 1.479	(1.0 to 1.0) 1.474	(1.0 to 1.0) 1.48	(1.0 to 1.0) 1.479	(1.0 to 1.0) 1.467	(1.0 to 1.0) 1.481	(1.0 to 1.0) 1.49	(1.0 to 1.0) 1.47	(1.0 to 1.0) 1.487	(1.0 to 1.0) 1.485	(1.0 to 1.0) 1.48	(1.0 to 1.0) 1.482	(1.0 to 1.0) 1.488	(1.0 to 1.0) 1.488	(1.0 to 1.0) 1.488	(1.0 to 1.0) 1.488
Other statema right exposure	Boll 7	vines -							(1.191 to 1.818) 1.485	(1.183 to 1.83) 1.485	(1.182 to 1.815) 1.479	(1.193 to 1.827) 1.471	(1.197 to 1.804) 1.469	(1.174 to 1.843) 1.486	(1.178 to 1.839) 1.485	(1.199 to 1.831) 1.481	(1.197 to 1.842) 1.48	(1.2 to 1.846) 1.47	(1.188 to 1.803) 1.473	(1.18 to 1.836) 1.471	(1.202 to 1.839) 1.49	(1.198 to 1.827) 1.464	(1.198 to 1.827) 1.464	(1.198 to 1.827) 1.464	(1.198 to 1.827) 1.464
Oter ladornia I recomme	Both	Malex							(1.199 to 1.845) 1.0	(1.183 to 1.855) 1.0	(1.184 to 1.844) 1.0	(1.181 to 1.774) 1.0	(1.183 to 1.819) 1.0	(1.196 to 1.823) 1.0	(1.19 to 1.848) 1.0	(1.191 to 1.856) 1.0	(1.199 to 1.814) 1.0	(1.184 to 1.812) 1.0	(1.19 to 1.814) 1.0	(1.194 to 1.789) 1.0	(1.209 to 1.859) 1.0	(1.18 to 1.788) 1.0	(1.18 to 1.788) 1.0	(1.18 to 1.788) 1.0	(1.18 to 1.788) 1.0
Other leukaernia Lew exposure	Both Fr	emales							(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0
Other leukaernia No exposure	Both	Malex							(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0)
Other leukaerria No exposure	Both Fe	emales							(1.0 to 1.0) 1.0 (1.0 to 1.0)	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	(1.0 is 1.0)	1.0
Occupational exposure to nickel									(10 0 10)	(10.00.10)	(11111)	(10 10 10)	(101010)	(1111)	(100010)	(10 10 10)	(10.0.10)	((10 10 10)	(in the fully	(101010)	(10 11 14)	(10.0.10)		(14.11.14)
Tracheal, bronchus, and lung cancer High exposure	Both	Both							2.148 (1.296 to 3.297)	2.148 (1.296 to 3.297)	2.148 (1.296 to 3.297)	2.148 (1.296 to 3.297)	2.148 (1.296 to 3.297)	2.148 (1.296 to 3.297)	2.148 (1.296 to 3.297)	2.148 (1.296 to 3.297)	2.148 (1.296 to 3.297)	2.148 (1.296 to 3.297)	2.148 (1.296 to 3.297)	2.148 (1.296 to 3.297)	2.148 (1.296 to 3.297)	2.148 (1.296 to 3.297)	2.148 (1.296 to 3.297)	2.148 (1.296 to 3.297)	2.148 (1.296 to 3.297)
Tracheal, bronchus, and lung cancer Low exposure	Both	Both							1.538 (0.606 to 3.355)	1.538 (0.606 to 3.355)	1.538 (0.606 to 3.355)	1.538 (0.606 to 3.355)	1.538 (0.606 to 3.355)	1.538 (0.606 to 3.355)	1.538 (0.606 to 3.355)	1.538 (0.606 to 3.355)	1.538 (0.606 to 3.355)	1.538 (0.606 to 3.355)	1.538 (0.606 to 3.355)	1.538 (0.606 to 3.355)	1.538 (0.606 to 3.355)	1.538 (0.606 to 3.355)	1.538 (0.606 to 3.355)	1.538 (0.606 to 3.355)	1.538 (0.606 to 3.355)
Tracheal, bronchus, and lang cancer No exposure	Both	Both							1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Occupational exposure to polycyclic aromatic hydrocarbons																									
Tracheal, bronchus, and lung cancer High exposure	Both	Males							1.31 (1.165 to 1.468)	1.304 (1.146 to 1.466)	1.313 (1.164 to 1.485)	1.314 (1.167 to 1.477)	1.315 (1.17 to 1.474)	1.314 (1.16 to 1.475)	1.315 (1.154 to 1.477)	1.312 (1.16 to 1.477)	1.318 (1.166 to 1.483)	1.311 (1.158 to 1.479)	1.313 (1.166 to 1.485)	1.314 (1.154 to 1.483)	1.313 (1.161 to 1.478)	1.309 (1.176 to 1.477)	1.309 (1.176 to 1.477)	1.309 (1.176 to 1.477)	1.309 (1.176 to 1.477)
Tracheal, bronchus, and lung cancer High exposure	Both Fe	emales							1.31 (1.154 to 1.486)	1.311 (1.157 to 1.469)	1.313 (1.154 to 1.472)	1.313 (1.155 to 1.469)	1.316 (1.162 to 1.483)	1.313 (1.156 to 1.476)	1.314 (1.155 to 1.483)	1.314 (1.171 to 1.483)	1.311 (1.146 to 1.477)	1.312 (1.163 to 1.481)	1.309 (1.147 to 1.489)	1.316 (1.168 to 1.475)	1.315 (1.15 to 1.479)	1.315 (1.166 to 1.481)	1.315 (1.166 to 1.481)	1.315 (1.166 to 1.481)	1.315 (1.166 to 1.481)
Tracheal, bronchus, and lung cancer Low exposure	Both	Males							1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Tracheal, bronchus, and lung cancer Low exposure	Both Fe	emakes							1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Tracheal, bronchus, and lung cancer No exposure	Both 3	Males							1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Tracheal, bronchus, and lung cancer No exposure	Both Fi	emakes							1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	(1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	(1.0 to 1.0)
Occupational exposure to silica									1.698	1.698	1.698	1.628	1.698	1.698	1.698	1.698	1.678	1.698	1.698	1.698	1.678	1.698	1.698	1.698	1.698
Tracheal, bronchus, and lung cancer High exposure	Both	Both							(1.164 to 2.259) 1.537	(1.164 to 2.259) 1.537	(1.164 to 2.259) 1.537	(1.164 to 2.259) 1.537	(1.164 to 2.259) 1.537	(1.164 to 2.259) 1.537	(1.164 to 2.259) 1.537	(1.164 to 2.259) 1.537	(1.164 to 2.259) 1.537	(1.164 to 2.259) 1.537	(1.164 to 2.259) 1.537	(1.164 to 2.259) 1.537	(1.164 to 2.259) 1.537	(1.164 to 2.259) 1.537	(1.164 to 2.259) 1.537	(1.164 to 2.259) 1.537	(1.164 to 2.259) 1.537
Trached brotelins and lang cancer. Low exposure	Both	Beth							(1.063 to 1.986) 1.0	(1.063 to 1.986) 1.0	(1.063 to 1.986) 1.0	(1.063 to 1.986) 1.0	(1.063 to 1.986) 1.0	(1.063 to 1.986) 1.0	(1.063 to 1.986) 1.0	(1.063 to 1.986) 1.0	(1.063 to 1.986) 1.0	(1.063 to 1.986) 1.0	(1.063 to 1.986) 1.0	(1.063 to 1.986) 1.0	(1.063 to 1.986) 1.0	(1.063 to 1.986) 1.0	(1.063 to 1.986) 1.0	(1.063 to 1.986) 1.0	(1.063 to 1.986) 1.0
	-Area -	- 1 I							(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)

| Appendix Table 6a. Relat | tive risks used by age and sex for each

 | outcome for all risk factors exce | pt for ambient air | pollution alcohol, | and anosang | | | | | |
 | | | Δ | 145
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---|---|--|--------------------|-------------|-------------|-----------|-----------|-------------|---|---
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NI G.		

 | N 117-111-17-0 | All-age | 0-6 days | 7-27 days | 28-364 days | 1-4 years | 5-9 years | 10-14 years | 15-19 years | 20-24 years
 | 25-29 years | 30-34 years | 35-39 years | 40-44 years
 | 45-49 years | 50-54 years | 55-59 years
 | 60-64 years | 65-69 years | 70-74 years | 75-79 years
 | 80-84 years | 85-89 years | 90-94 years | 95+ years
 |
| Occupational exposur | e Category / Units

 | Morbidity / Mortanty Se | x | | | | | | | |
 | | | |
 | | |
 | | | |
 | | | |
 |
| Larynx cancer | High exposure

 | Both Bo | њ. | | 1 | | | | | 4.566
(2.122 m 8.328) | 4.566
(2.122 to 8.328)
 | 4.566
(2.122 to 8.328) | 4.566
(2.122 to 8.328) | 4.566
(2.122 to 8.328) | 4.566
(2.122 to 8.328)
 | 4.566
(2.122 to 8.328) | 4.566
(2.122 to 8.328) | 4.566
(2.122 to 8.328)
 | 4.566
(2.122 to 8.328) | 4.566
(2.122 to 8.328) | 4.566
(2.122 to 8.328) | 4.566
(2.122 to 8.328)
 | 4.566
(2.122 to 8.328) | 4.566
(2.122 to 8.328) | 4.566
(2.122 to 8.328) | 4.566
(2.122 to 8.328)
 |
| Larynx cancer | Low exposure

 | Both Bo | њ. | | | | | | | 2.024
(0.944 m 3.782) | 2.024
(0.944 to 3.787)
 | 2.024
(0.944 to 3.787) | 2.024
(0.944 to 3.782) | 2.024
(0.944 to 3.782) | 2.024
(0.944 to 3.787)
 | 2.024
(0.944 to 3.782) | 2.024
(0.944 to 3.782) | 2.024
(0.944 to 3.787)
 | 2.024
(0.944 to 3.787) | 2.024
(0.944 to 3.782) | 2.024
(0.944 to 3.787) | 2.024
(0.944 to 3.782)
 | 2.024
(0.944 to 3.787) | 2.024
(0.944 to 3.782) | 2.024
(0.944 to 3.787) | 2.024
(0.944 to 3.787)
 |
| Larynx cancer | No exposure

 | Both Bo | њ. | | | | | | | 1.0 | 1.0
 | 1.0 | 1.0 | 1.0 | 1.0
 | 1.0 | 1.0 | 1.0
 | 1.0 | 1.0 | 1.0 | 1.0
 | 1.0 | 1.0 | 1.0 | 1.0
 |
| Occupational exposu | are to trichloroethylene

 | | | | | | | | | (1.0 10 1.0) | (1.0 10 1.0)
 | (12012) | (10 10 1.0) | (1.010 1.0) | (1.0 10 1.0)
 | (1.0 10 1.0) | (10 0 10) | (101010)
 | (1.5 10 1.55) | (1/10/1/0) | (10010) | (1.0101.0)
 | (1.5 10 1.5) | (13/12/13) | (1.0 00 1.0) | (1.0101.0)
 |
| Kidney cancer | High exposure

 | Both Bo | њ. | | 1 | | | | | 1.245 | 1.245
 | 1.245 | 1.245 | 1245 | 1.245
 | 1.245 | 1.245 | 1.245
 | 1.245 | 1.245 | 1.245 | 1.245
 | 1.245 | 1.245 | 1.245 | 1.245
 |
| Kidney cancer | Low esposare

 | Both Bo | њ. | | | | | | | 1.0 | 1.0
 | 1.0 | 1.0 | 1.0 | 1.0
 | 1.0 | 1.0 | 1.0
 | 1.0 | 1.0 | 1.0 | 1.0
 | 1.0 | 1.0 | 1.0 | 1.0
 |
| Kidney cancer | No exposure

 | Both Bo | њ. | | | | | | | 1.0 | 1.0
 | 1.0 | 1.0 | 1.0 | 1.0
 | 1.0 | 1.0 | 1.0
 | 1.0 | 1.0 | 1.0 | 1.0
 | 1.0 | 1.0 | 1.0 | 1.0
 |
| Occupational asthma | agens

 | | | | | | | | | (1.0 10 1.0) | (1.0 10 1.0)
 | (120012) | (10 10 1.0) | (1.010 1.0) | (15/01/3)
 | (13/01/0) | (13 6 13) | (101010)
 | (1.5 10 1.55) | (13/01/0) | (100010) | (1.0101.0)
 | (1.5 10 1.5) | (13/12/13) | (130213) | (1.0101.0)
 |
| Asthma | Admin

 | Both Mal | a | | 1 | | | | | 1.0
(1.0 m 1.0) | 1.0
 | 1.0 | 1.0 | 1.0 | 1.0
(1.0 m 1.0)
 | 1.0 | 1.0 | 1.0
(1.0 to 1.0)
 | 1.0
(1.0 to 1.0) | 1.0 | 1.0 | 1.0
 | 1.0
(1.0 m 1.0) | | |
 |
| Asthena | Admin

 | Both Fem | iles | | | | | | | 1.0 | 1.0
 | 1.0 | 1.0 | 1.0 | 1.0
 | 1.0 | 1.0 | 1.0
 | 1.0 | 1.0 | 1.0 | 1.0
 | 1.0 | | |
 |
| Asthena | Technical

 | Both Mal | a | | | | | | | 1.05 | 1.051
 | 1.051 | 1.051 | 1.05 | 1.05
 | 1.051 | 1.051 | 1.05
 | 1.05 | 1.051 | 1.051 | 1.051
 | 1.051 | | |
 |
| Asthma | Technical

 | Both Fem | dex | | | | | | | 1.06 | 1.06
 | 1.059 | 1.061 | 1.06 | 1.06
 | 1.059 | 1.06 | 1.06
 | 1.061 | 1.06 | 1.061 | 1.06
 | 1.06 | | |
 |
| Asthma | Sales

 | Both Mal | a | | | | | | | 1.14 | (1.024 to 1.099)
1.14
 | 1.144 | 1.142 | 1.14 | (1.026161.094)
 | 1.143 | (1.028 16 1.054) | (1.0271811.092)
 | 1.138 | (1.027 65 1.096)
1.139 | (1.023 to 1.096) | (1.027181.096)
 | (102781096) | | |
 |
| Aethroa | Sales

 | Both Fem | des | | | | | | | (1.047 to 1.237)
1.131 | (1.049 to 1.254)
1.13
 | (1.055 to 1.233)
1.129 | (1.048 to 1.254)
1.13 | (1.046 to 1.234)
1.131 | (1.053 to 1.235)
1.13
 | (1.052 to 1.238) | (1.047 to 1.256)
1.13 | (1.056 to 1.235)
1.131
 | (1.046 to 1.256)
1.13 | (1.057 to 1.226)
1.129 | (1.052 to 1.243)
1.131 | (1.058 to 1.239)
 | (1.058 to 1.2.59)
1.131 | | |
 |
| Asthrea | Arriculture

 | Both Mal | ~ | | | | | | | (1.083 to 1.182)
1.519 | (1.083 to 1.178)
1.527
 | (1.083 to 1.178)
1.524 | (1.081 to 1.182)
1.513 | (1.084 to 1.18)
1.523 | (1.08 to 1.181)
1.516
 | (1.083 to 1.181)
1.522 | (1.079 to 1.178)
1.52 | (1.077 to 1.184)
1.519
 | (1.081 to 1.182)
1.531 | (1.082 to 1.181)
1.52 | (1.083 to 1.184)
1.508 | (1.083 to 1.183)
1.498
 | (1.083 to 1.183)
1.498 | | |
 |
| Authors | Amindum

 | Rude Emm | | | | | | | | (1.1 to 2.029)
1.52 | (1.119 to 2.063)
1.506
 | (1.103 to 2.019)
1.52 | (1.127 to 1.999)
1.514 | (1.108 to 2.023)
1.509 | (1.083 to 2.043)
1.513
 | (1.106 to 2.038)
1.508 | (1.122 to 2.023)
1.526 | (1.103 to 2.011)
1.519
 | (1.122 to 2.047)
1.53 | (1.106 to 2.041)
1.519 | (1.083 to 2.019)
1.518 | (1.081 to 1.965)
1.502
 | (1.081 to 1.965)
1.502 | | |
 |
| |

 | | | | | | | | | (1.125 to 2.022)
1.959 | (1.099 to 1.997)
1.959
 | (1.115 to 2.05)
1.971 | (1.109 to 2.048)
1.966 | (1.101 to 1.969)
1.963 | (1.128 to 2.026)
1.959
 | (1.108 to 2.0)
1.956 | (1.108 to 2.07)
1.964 | (1.117 to 1.981)
1.969
 | (1.105 to 2.017)
1.954 | (1.107 to 2.024)
1.965 | (1.115 to 2.018)
1.955 | (1.094 to 2.03)
1.953
 | (1.094 to 2.03)
1.953 | | |
 |
| Asima | Milling

 | Boll Mill | | | | | | | | (1.576 to 2.413)
1.956 | (1.568 to 2.381)
1.961
 | (1.602 to 2.414)
1.967 | (1.571 to 2.396)
1.952 | (1.601 to 2.395)
1.959 | (1.588 to 2.385)
1.959
 | (1.595 to 2.406)
1.962 | (1.57 to 2.397)
1.955 | (1.616 to 2.417)
1.948
 | (1.574 to 2.406)
1.964 | (1.558 to 2.393)
1.965 | (1.577 to 2.39)
1.959 | (1.551 to 2.418)
1.973
 | (1.551 to 2.418)
1.973 | | |
 |
| Asima | Milling

 | BOIN PAR | lacs | | | | | | | (1.58 to 2.408) | (1.574 to 2.395)
 | (1.586 to 2.417) | (1.567 to 2.371) | (1.575 to 2.416) | (1.567 to 2.417)
 | (1.585 to 2.386) | (1.591 to 2.393) | (1.57 to 2.381)
 | (1.597 to 2.409) | (1.59 to 2.414) | (1.588 to 2.39) | (1.586 to 2.392)
 | (1.586 to 2.392) | | |
 |
| Asthra | Transport

 | Both Mal | a | | | | | | | (1.22 to 1.402) | (1.222 to 1.406)
 | (1.225 to 1.399) | (1.218 to 1.398) | (1.225 to 1.404) | (1.225 to 1.401)
 | (1.226 to 1.402) | (1.223 to 1.398) | (1.221 to 1.396)
 | (1.224 to 1.408) | (1.228 to 1.397) | (1.225 to 1.409) | (1.226 to 1.407)
 | (1.226 to 1.407) | | |
 |
| Asthra | Transport

 | Both Femi | des | | | | | | | (1.132 to 1.312)
1.559 | (1.138 to 1.31)
 | (1.136 to 1.303) | (1.138 to 1.313) | (1.137 to 1.317) | (1.132 to 1.314)
 | (1.132 to 1.312) | (1.134 to 1.313) | (1.14 to 1.315)
 | (1.139 to 1.309)
1.562 | (1.133 to 1.312) | (1.135 to 1.316) | (1.133 to 1.313)
1.561
 | (1.133 to 1.313) | | |
 |
| Asthra | Manufact

 | Both Mal | 0 | | | | | | | (1.477 to 1.647) | (1.474 to 1.652)
 | (1.474 to 1.657) | (1.472 to 1.658) | (1.473 to 1.65) | (1.471 to 1.655)
 | (1.471 to 1.649) | (1.473 to 1.654) | (1.461 to 1.655)
 | (1.479 to 1.655) | (1.475 to 1.652) | (1.476 to 1.657) | (1.474 to 1.654)
 | (1.474 to 1.654) | | |
 |
| Asthma | Manufact

 | Both Fema | des | | | | | | | (1.272 to 1.392) | (1.27 to 1.39)
 | (1.271 to 1.39) | (1.271 to 1.388) | (1.268 to 1.392) | (1.269 to 1.39)
 | (1.269 to 1.391) | (1.272 to 1.394) | (1.273 to 1.389)
 | (1.272 to 1.391) | (1.268 to 1.392) | (1.268 to 1.394) | (1.272 to 1.39)
 | (1.272 to 1.39) | | |
 |
| Asthma | Services

 | Both Mal | a | | | | | | | (1.415 to 1.646) | (1.416 to 1.652)
 | (1.409 to 1.649) | 1.532
(1.413 to 1.653) | (1.424 to 1.648) | (1.416 to 1.645)
 | 1.533
(1.418 to 1.646) | 1.535
(1.427 to 1.649) | (1.416 to 1.655)
 | 1.528
(1.411 to 1.651) | 1.53
(1.416 to 1.652) | 1.53
(1.411 to 1.658) | 1.531
(1.415 to 1.655)
 | (1.415 to 1.655) | | |
 |
| Asthma | Services

 | Both Femi | ules | | | | | | | 1.41
(1.352 to 1.467) | 1.41
(1.357 to 1.469)
 | 1.412
(1.357 to 1.467) | 1.41
(1.356 to 1.464) | 1.409
(1.353 to 1.463) | 1.409
(1.354 to 1.467)
 | 1.41
(1.354 to 1.467) | 1.411
(1.357 to 1.464) | 1.409
(1.357 to 1.46)
 | 1.411
(1.355 to 1.466) | 1.41
(1.354 to 1.465) | 1.41
(1.357 to 1.467) | 1.411
(1.357 to 1.464)
 | 1.411
(1.357 to 1.464) | | |
 |
| Asthma | Other

 | Both Mal | α | | | | | | | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0)
 | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0)
 | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0)
 | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0)
 | 1.0
(1.0 to 1.0) | | |
 |
| Asthma | Other

 | Both Fem | iles | | 1 | | | | | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0)
 | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0)
 | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0)
 | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0)
 | 1.0
(1.0 to 1.0) | Ļ | |
 |
| Occupational particu | ulate matter, gases, and fumes

 | | | | | | | | | |
 | | | |
 | | |
 | | | |
 | | | |
 |
| Chronic obstructive p
disease | palmonary High

 | Both Mal | a | | | | | | | 2.364
(1.463 to 3.64) | 2.341
(1.426 to 3.555)
 | 2.37
(1.41 to 3.596) | 2.391
(1.455 to 3.756) | 2.387
(1.409 to 3.791) | (1.4 to 3.672)
 | (1.439 to 3.833) | 2.361
(1.447 to 3.684) | 2.367
(1.454 to 3.654)
 | 2.373
(1.375 to 3.64) | 2.387
(1.438 to 3.685) | 2.402
(1.434 to 3.763) | 2.398
(1.441 to 3.813)
 | 2.359
(1.429 to 3.695) | 2.359
(1.429 to 3.695) | 2.359
(1.429 to 3.695) | 2.359
(1.429 to 3.695)
 |
| Chronic obstructive p
disease | palmonary High

 | Both Femi | des | | | | | | | 2.371
(1.459 to 3.719) | 2.395
(1.483 to 3.669)
 | 2.364
(1.473 to 3.694) | (1.431 to 3.763) | 2.375
(1.446 to 3.698) | 2.326
(1.438 to 3.576)
 | 2.38
(1.432 to 3.702) | 2.35
(1.425 to 3.609) | 2.364
(1.461 to 3.677)
 | 2.364
(1.475 to 3.64) | 2.395
(1.476 to 3.715) | 2.336
(1.434 to 3.673) | 2.363
(1.431 to 3.741)
 | 2.404
(1.462 to 3.619) | 2.404
(1.462 to 3.619) | 2.404
(1.462 to 3.619) | 2.404
(1.462 to 3.619)
 |
| Chronic obstructive p
disease | palmonary Low

 | Both Mal | a | | | | | | | 1.462
(1.057 to 1.915) | 1.457
(1.052 to 1.935)
 | 1.464
(1.085 to 1.91) | 1.45
(1.067 to 1.947) | (1.06 to 1.963) | 1.454
(1.075 to 1.96)
 | 1.446
(1.055 to 1.929) | 1.457
(1.076 to 1.933) | 1.453
(1.052 to 1.903)
 | 1.46
(1.067 to 1.954) | 1.443
(1.057 to 1.916) | 1.462
(1.097 to 1.954) | (1.079 to 1.952)
 | 1.462
(1.096 to 1.965) | 1.462
(1.096 to 1.965) | 1.462
(1.096 to 1.965) | 1.462
(1.096 to 1.965)
 |
| chronic obstructive p
disease | palmonary Low

 | Both Femi | des | | | | | | | 1.446
(1.063 to 1.904) | (1.09 to 1.954)
 | 1.453
(1.055 to 2.003) | (1.072 to 1.931) | 1.456
(1.056 to 1.968) | 1.47
(1.089 to 1.96)
 | 1.44
(1.046 to 1.932) | (1.097 to 1.925) | (1.061 to 1.921)
 | 1.451
(1.082 to 1.912) | (1.102 to 1.941) | 1.448
(1.077 to 1.929) | 1.467
(1.072 to 1.971)
 | 1.456
(1.056 to 1.927) | (1.056 to 1.927) | (1.056 to 1.927) | 1.456
(1.056 to 1.927)
 |
| Chronic obstructive p
disease | palmonary None

 | Both Mal | a | | | | | | | (1.0 to 1.0) | (1.0 to 1.0)
 | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0) | (1.0 to 1.0)
 | (1.0 to 1.0) | 1.0
(1.0 to 1.0) | (1.0 to 1.0)
 | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0)
 | (1.0 to 1.0) | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0) | (1.0 to 1.0)
 |
| disease | palmonary None

 | Both Fem | ıles | 1 | 1 | | | | | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0)
 | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0)
 | (1.0 to 1.0) | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0)
 | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0)
 | (1.0 to 1.0) | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0) | 1.0
(1.0 to 1.0)
 |
| Occupational noise * |

 | | | | | | | | | |
 | | | |
 | | |
 | | | |
 | | | |
 |
| Mild hearing loss
and other hearing loss | er to age-related High exposure, >90dB
as

 | Morbidity Bor | th. | | | | | | | 7.477
(5.049 to 11.132) | 7.423
 | | | |
 | | |
 | | | |
 | | | | (1.048 to 1.211)
 |
| Muld hearing loss with
to age-related and oth | th ringing due High exposure, >90dB her hearing loss

 | Morbidity Boy | a. | | | | | | | | (4.787 to 10.956)
 | (4.293 to 6.843) | 5.456
(4.339 to 6.828) | 3.07
(2.692 to 3.51) | 3.081
(2.715 to 3.498)
 | 2.546
(2.332 to 2.774) | 2.554
(2.367 to 2.752) | 1.849
(1.712 to 1.995)
 | 1.85
(1.707 to 1.995) | 1.45
(1.372 to 1.526) | 1.454
(1.378 to 1.529) | 1.131
(1.048 to 1.211)
 | 1.131
(1.048 to 1.211) | 1.131
(1.048 to 1.211) | (1.048 to 1.211) |
 |
| Moderate hearing loss
related and other hear
Moderate hearing loss | ss date to age-
tring loss High exposure, >90dB

 | | | | | | | | | 7.477
(5.049 to 11.132) | (4.787 to 10.956)
7.423
(4.787 to 10.956)
 | 3.504
(4.293 to 6.843)
5.504
(4.293 to 6.843) | 5.456
(4.339 to 6.828)
5.456
(4.339 to 6.828) | 3.07
(2.692 to 3.51)
3.07
(2.692 to 3.51) | 3.081
(2.715 to 3.498)
3.081
(2.715 to 3.498)
 | 2.546
(2.332 to 2.774)
2.546
(2.332 to 2.774) | 2.554
(2.367 to 2.752)
2.554
(2.367 to 2.752) | 1.849
(1.712 to 1.995)
1.849
(1.712 to 1.995)
 | 1.85
(1.707 to 1.995)
1.85
(1.707 to 1.995) | 1.45
(1.372 to 1.526)
1.45
(1.372 to 1.526) | 1.454
(1.378 to 1.529)
1.454
(1.378 to 1.529) | 1.131
(1.048 to 1.211)
1.131
(1.048 to 1.211)
 | 1.131
(1.048 to 1.211)
1.131
(1.048 to 1.211) | 1.131
(1.048 to 1.211)
1.131
(1.048 to 1.211) | (1.048 to 1.211)
1.131
(1.048 to 1.211) | 1.131
(1.048 to 1.211)
 |
| chae to age-related and
loss | as with ringing

 | Morbidity Bo | da . | | | | | | | 7.477
(5.049 to 11.132)
8.175
(4.62 to 13.221) | (4.787 to 10.956)
7.423
(4.787 to 10.956)
8.31
(4.828 to 13.462)
 | 5.504
(4.293 to 6.843)
5.504
(4.293 to 6.843)
6.71
(4.785 to 9.239) | 3.456
(4.339 to 6.828)
5.456
(4.339 to 6.828)
6.756
(4.705 to 9.168) | 3.07
(2.692 to 3.51)
3.07
(2.692 to 3.51)
5.992
(4.34 to 8.307) | 3.081
(2.715 to 3.498)
3.081
(2.715 to 3.498)
5.987
(4.304 to 8.107)
 | 2.546
(2.332 to 2.774)
2.546
(2.332 to 2.774)
5.624
(3.954 to 7.821) | 2.554
(2.367 to 2.752)
2.554
(2.367 to 2.752)
5.61
(3.96 to 7.69) | 1.849
(1.712 to 1.995)
1.849
(1.712 to 1.995)
3.643
(2.576 to 5.019)
 | 1.85
(1.707 to 1.995)
1.85
(1.707 to 1.995)
3.601
(2.494 to 4.936) | 1.45
(1.372 to 1.526)
1.45
(1.372 to 1.526)
2.17
(1.597 to 2.84) | 1.454
(1.378 to 1.529)
1.454
(1.378 to 1.529)
2.146
(1.656 to 2.862) | 1.131
(1.048 to 1.211)
1.131
(1.048 to 1.211)
1.291
(1.06 to 1.539)
 | 1.131
(1.048 to 1.211)
1.131
(1.048 to 1.211)
1.291
(1.06 to 1.539) | 1.131
(1.048 to 1.211)
1.131
(1.048 to 1.211)
1.291
(1.06 to 1.539) | (1.048 to 1.211)
1.131
(1.048 to 1.211)
1.291
(1.06 to 1.539) | 1.131
(1.048 to 1.211)
1.291
(1.06 to 1.539)
 |
| 1 | ss with ringing
al other hearing High exposure, >90dB

 | Morbidity Bo
Morbidity Bo | a
a | | | | | | | 7,477
(5.049 to 11.132)
8.175
(4.62 to 13.221)
8.175
(4.62 to 13.221) | (4.787 to 10.956)
7.423
(4.787 to 10.956)
8.31
(4.828 to 13.462)
8.31
(4.828 to 13.462)
 | 5.504
(4.293 to 6.843)
5.504
(4.293 to 6.843)
6.71
(4.785 to 9.239)
6.71
(4.785 to 9.239) | 5.456
(4.339 to 6.828)
5.456
(4.339 to 6.828)
6.756
(4.705 to 9.168)
6.756
(4.705 to 9.168) | 3.07
(2.692 to 3.51)
3.07
(2.692 to 3.51)
5.992
(4.34 to 8.307)
5.992
(4.34 to 8.307) | 3.081
(2.715 to 3.498)
3.081
(2.715 to 3.498)
5.987
(4.304 to 8.107)
5.987
(4.304 to 8.107)
 | 2.546
(2.332 to 2.774)
2.546
(2.332 to 2.774)
5.624
(3.954 to 7.821)
5.624
(3.954 to 7.821) | 2.554
(2.367 to 2.752)
2.554
(2.367 to 2.752)
5.61
(3.96 to 7.69)
5.61
(3.96 to 7.69) | 1.849
(1.712 to 1.995)
1.849
(1.712 to 1.995)
3.643
(2.576 to 5.019)
3.643
(2.576 to 5.019)
 | 1.85
(1.707 to 1.995)
1.85
(1.707 to 1.995)
3.601
(2.494 to 4.936)
3.601
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(4.007 to 8.022)
5.704
(4.007 to 8.022) | $\begin{array}{c} 1.849\\ (1.712\ to\ 1.995)\\ 1.849\\ (1.712\ to\ 1.995)\\ 3.643\\ (2.576\ to\ 5.019)\\ 3.648\\ (2.576\ to\ 5.019)\\ 3.638\\ (2.576\ to\ 4.959)\\ 3.638\ to\ 4.959\\ (2.576\ to\ 4.959)\\ 3.638\ to\ 4.959\\ (2.576\ to\ $ | $\begin{array}{c} 1.85 \\ (1.307 \ \mathrm{bt} \ 1.995) \\ 1.85 \\ (1.307 \ \mathrm{bt} \ 1.995) \\ 3.601 \\ (2.494 \ \mathrm{bt} \ 4.936) \\ 3.591 \\ (2.583 \ \mathrm{bt} \ 4.981) \\ 3.591 \\ (2.583 \ \mathrm{bt} \ 4.981) \\ \end{array}$
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(4.73 to 9.34) | $\begin{array}{c} 3.846\\ (4.339 to 6.828)\\ 5.456\\ (4.339 to 6.828)\\ 6.756\\ (4.705 to 9.168)\\ 6.756\\ (4.705 to 9.168)\\ 6.726\\ (4.762 to 9.201)\\ 6.726\\ (4.762 to 9.201)\\ 6.637\\ (4.739 to 9.181)\end{array}$ | 3.07
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(2.692 to 3.51)
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5.992
(4.34 to 8.307)
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(4.114 to 8.374)
6.007
(4.114 to 8.374)
6.075
(4.315 to 8.372) | $\begin{array}{c} 3.081\\ (2.715 \pm 3.408)\\ 3.081\\ (2.715 \pm 3.408)\\ (2.715 \pm 3.408)\\ (4.304 \pm 8.8107)\\ 5.987\\ (4.304 \pm 8.8107)\\ 6.987\\ (4.34 \pm 8.8175)\\ 6.072\\ (4.34 \pm 8.475)\\ 6.072\\ (4.34 \pm 8.475)\\ 5.983\\ (4.247 \pm 8.221)\end{array}$
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(3.954 to 7.821)
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(2.513 to 5.117) | $\begin{array}{c} 1.45\\ (1.372\ to 1.526)\\ 1.45\\ (1.372\ to 1.526)\\ 2.17\\ (1.597\ to 2.34)\\ 2.17\\ (1.597\ to 2.34)\\ 2.162\\ (1.64\ to 2.316)\\ 2.162\\ (1.64\ to 2.316)\\ 2.163\\ (1.64\ to 2.316)\\ 2.138\\ (1.606\ to 2.302)\end{array}$ | $\begin{array}{c} 1.454\\ (1.378\mathrm{is}\ 1.529)\\ 1.454\\ (1.378\mathrm{is}\ 1.529)\\ 2.146\\ (1.656\mathrm{is}\ 2.862)\\ 2.146\\ (1.656\mathrm{is}\ 2.862)\\ 2.168\\ (1.631\mathrm{is}\ 2.827)\\ 2.168\\ (1.631\mathrm{is}\ 2.827)\\ 2.173\\ (1.645\mathrm{is}\ 2.807)\end{array}$ | $\begin{array}{c} 1.131\\ (1.048\ \mathrm{in}\ 1.211)\\ 1.131\\ (1.048\ \mathrm{in}\ 1.211)\\ 1.291\\ (1.06\ \mathrm{in}\ 1.539)\\ 1.291\\ (1.06\ \mathrm{in}\ 1.539)\\ 1.297\\ (1.06\ \mathrm{in}\ 1.561)\\ 1.297\\ (1.06\ \mathrm{in}\ 1.561)\\ 1.294\\ (1.05\ \mathrm{in}\ 1.561)\\ 1.294\end{array}$
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(4.73 to 9.94) | $\begin{array}{c} 3, 366\\ (4.396\ m 6.828)\\ 5, 466\\ (4.396\ m 6.828)\\ 6, 756\\ (4.705\ m 9.168)\\ 6, 756\\ (4.705\ m 9.168)\\ 6, 726\\ (4.705\ m 9.201)\\ 6, 687\\ (4.736\ m 9.201)\\ 6, 687\\ (4.739\ m 9.181)\\ 6, 687\\ (4.739\ m 9.181)\\ \end{array}$ | 3.07
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(3.946 to 7.534) | $\begin{array}{c} 1549\\ (1.712u1.995)\\ 1.849\\ (1.712u1.995)\\ 3.643\\ (2.576u5.019)\\ 3.643\\ (2.576u5.019)\\ 3.638\\ (2.576u5.019)\\ 3.638\\ (2.576u4.959)\\ 3.62\\ (2.541u5.01)\\ 3.62\\ (2.541u5.01)\\ 3.62\end{array}$ | $\begin{array}{c} 1.85 \\ (1.707\mathrm{m}\;1.995) \\
1.85 \\ (1.207\mathrm{m}\;1.995) \\ 3.601 \\ (2.494\mathrm{m}\;4.936) \\ 3.601 \\ (2.494\mathrm{m}\;4.936) \\ 1.591 \\ (2.583\mathrm{m}\;4.981) \\ 3.591 \\ (2.583\mathrm{m}\;4.981) \\ 3.583 \\ (2.513\mathrm{m}\;5.117) \\ 3.583 \\ (2.513\mathrm{m}\;5.117) \end{array}$ | $\begin{array}{c} 1.45\\ (1.372\ {\rm m}\ 1.526)\\ 1.45\\ (1.372\ {\rm m}\ 1.526)\\ 2.17\\ (1.597\ {\rm m}\ 2.34)\\ 2.16\\ (1.64\ {\rm m}\ 2.316)\\ 2.162\\ (1.64\ {\rm m}\ 2.316)\\ 2.138\\ (1.666\ {\rm m}\ 2.802)\\ \end{array}$ | $\begin{array}{c} 1.454\\ (1.378 \approx 1.529)\\ 1.454\\ (1.378 \approx 1.529)\\ 2.146\\ (1.556 \approx 2.862)\\ 2.146\\ (1.656 \approx 2.862)\\ 2.146\\ (1.656 \approx 2.862)\\ 2.168\\ (1.631 \approx 2.827)\\ 2.168\\ (1.631 \approx 2.827)\\ 2.173\\ (1.645 \approx 2.807)\\ 2.173\end{array}$ | $\begin{array}{c} 1.131\\ (1.048\ {\rm in}\ 1.211)\\ 1.131\\ (1.048\ {\rm in}\ 1.211)\\ 1.291\\ (1.06\ {\rm in}\ 1.539)\\ 1.297\\ (1.06\ {\rm in}\ 1.539)\\ 1.297\\ (1.06\ {\rm in}\ 1.561)\\ 1.297\\ (1.06\ {\rm in}\ 1.561)\\ 1.294\\ (1.059\ {\rm in}\ 1.569)\\ 1.294\end{array}$
 | $\begin{array}{c} 1.131\\ (1.048 \text{ to } 1.211)\\ 1.531\\ (1.048 \text{ to } 1.211)\\ 1.391\\ (1.06 \text{ to } 1.539)\\ 1.291\\ (1.06 \text{ to } 1.539)\\ 1.297\\ (1.06 \text{ to } 1.561)\\ 1.297\\ (1.06 \text{ to } 1.561)\\ 1.294\\ (1.059 \text{ to } 1.569)\\ 1.294\\ \end{array}$ | $\begin{array}{c} 1.31\\ (1.08 \pm 0.211)\\ 1.33\\ (1.08 \pm 0.211)\\ 1.291\\ (1.08 \pm 0.539)\\ 1.291\\ (1.08 \pm 0.539)\\ 1.297\\ (1.08 \pm 0.561)\\ 1.297\\ (1.08 \pm 0.561)\\ 1.294\\ (1.099 \pm 1.599)\\ 1.294\end{array}$ | $\begin{array}{c} (1.048 \pm 1.211) \\ 1.131 \\ (1.048 \pm 1.211) \\ 1.291 \\ (1.06 \pm 1.239) \\ 1.291 \\ (1.06 \pm 1.539) \\ 1.291 \\ (1.06 \pm 1.561) \\ 1.297 \\ (1.06 \pm 1.561) \\ 1.297 \\ (1.06 \pm 1.561) \\ 1.297 \\ (1.06 \pm 1.561) \\ 1.294 \\ (1.059 \pm 1.569) \end{array}$ | 1.131
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Machiday Bo | 42
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44 | | | | | | | $\begin{array}{c} 7.477\\ (5.649 to 11.132)\\ 8.175\\ (4.62 to 13.221)\\ 8.77\\ (4.62 to 13.221)\\ 8.27\\ (4.807 to 13.224)\\ 8.249\\ (4.2801 to 13.224)\\ 8.249\\ (4.72 to 13.18)\\ 8.239\\ (4.72 to 13.18)\\ 8.233\\ (4.77 to 13.62)\end{array}$ | $\begin{array}{c} (4.787 \ m \ 10.056)\\ 7.423\\ (4.787 \ m \ 10.056)\\ 8.31\\ (4.528 \ m \ 13.462)\\ 8.213\\ (4.709 \ m \ 13.126)\\ 8.223\\ (4.709 \ m \ 13.126)\\ 8.292\\ (4.754 \ m \ 13.144)\\ 8.247\\ (4.945 \ m \ 13.59)\\ \end{array}$ | 3.504 42.93 to 6.843) 5.504 (4.293 to 6.843) (4.293 to 6.843) (4.293 to 6.843) 6.71 (4.785 to 9.239) 6.71 (4.785 to 9.239) 6.764 (4.814 to 9.271) 6.764 (4.814 to 9.271) 6.707 (4.73 to 9.34) 6.707 (4.739 to 9.24) 6.701 (4.739 to 9.26)
 | $\begin{array}{c} 3, 366\\ (4.39) = 6.828)\\ 5, 466\\ (4.39) = 6.828)\\ 6.756\\ (4.708 \pm 9.168)\\ 6.756\\ (4.708 \pm 9.168)\\ 6.726\\ (4.708 \pm 9.201)\\ 6.726\\ (4.762 \pm 9.201)\\ 6.687\\ (4.778 \pm 9.201)\\ 6.687\\ (4.739 \pm 9.181)\\ 6.687\\ (4.739 \pm 9.181)\\ 6.637\\ (4.739 \pm 9.181)\\ 6.734\\ (4.811 \pm 9.238)\\ \end{array}$ | 3.07
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(4.34 to 8.475)
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(4.34 to 8.4 | $\begin{array}{c} 2.546\\ (2321 w 2.774)\\ 3.546\\ (2321 w 2.774)\\ 5.624\\ (3954 w 7.821)\\ 5.624\\ (3954 w 7.821)\\ 5.638\\ (3954 w 7.821)\\ 5.638\\ (3943 w 7.822)\\ 5.638\\ (3943 w 7.822)\\ 5.638\\ (4059 w 7.875)\\ 5.66\\ (4059 w 7.875)\\ 5.66\\ (4059 w 7.875)\\ 5.66\\ (3965 w 7.879)\end{array}$ | 2.554
(2.367 le 2.752)
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(3.96 le 7.69)
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(4.007 le 8.022)
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(3.94 le 7.534)
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(3.94 le 7.534)
5.838 le 7.693)
 | $\begin{array}{c} 1.849\\ (1.712 {\rm m} 1.995)\\ 1.849\\ (1.712 {\rm m} 1.995)\\ 3.5648\\ (2.576 {\rm m} 5.019)\\ 3.648\\ (2.576 {\rm m} 5.019)\\ 3.638\\ (2.576 {\rm m} 4.959)\\ 3.638\\ (2.576 {\rm m} 4.959)\\ 3.638\\ (2.576 {\rm m} 4.959)\\ 3.62\\ (2.441 {\rm m} 5.01)\\ 3.62\\ (2.441 {\rm m} 5.01)\\ 3.62\\ (2.441 {\rm m} 5.01)\\ 3.59\\ (2.489 {\rm m} 5.087)\\ \end{array}$ | $\begin{array}{c} 1.85\\ (1.701\ {\rm m}\ 1.995)\\ 1.23\\ (1.701\ {\rm m}\ 1.995)\\ 3.601\\ (2.494\ {\rm m}\ 4.936)\\ 3.601\\ (2.494\ {\rm m}\ 4.936)\\ 3.591\\ (2.531\ {\rm m}\ 4.936)\\ 3.581\\ (2.531\ {\rm m}\ 5.117)\\ 3.583\\ (2.511\ {\rm m}\ 5.117)\\ 3.583\\ (2.511\ {\rm m}\ 5.117)\\ 3.607\\ (2.48\ {\rm m}\ 5.948)\\ \end{array}$ | $\begin{array}{c} 1.45\\ (1.372 \approx 1.526)\\ 1.45\\ (1.372 \approx 1.536)\\ 2.17\\ (1.597 \approx 2.34)\\ 2.162\\ (1.64 \approx 2.316)\\ 2.162\\ (1.64 \approx 2.316)\\ 2.163\\ (1.66 \approx 2.302)\\ 2.138\\ (1.66 \approx 2.302)\\ 2.138\\ (1.66 \approx 2.302)\\ 2.138\\ (1.66 \approx 2.302)\\ 2.144\\ (1.619 \approx 2.83)\\ \end{array}$
 | $\begin{array}{c} 1.454\\ (1.378\pm 0.1529)\\ 1.454\\ (1.378\pm 0.1529)\\ 2.146\\ (1.558\pm 0.2862)\\ 2.166\\ (1.658\pm 0.2862)\\ 2.168\\ (1.651\pm 0.2877)\\ 2.178\\ (1.651\pm 0.2877)\\ 2.173\\ (1.648\pm 0.2807)\\ 2.173\\ (1.658\pm 0.2807)\\ 2.173\\ (1.638\pm 0.2827)\\ 2.173\\ (1.638\pm 0.2822)\\ \end{array}$ | $\begin{array}{c} 1.131\\ (1.0481e121)\\ 1.131\\ (1.0481e121)\\ 1.291\\ (1.061e1539)\\ 1.291\\ (1.061e1539)\\ 1.297\\ (1.061e1551)\\ 1.297\\ (1.061e1551)\\ 1.294\\ (1.051e1551)\\ 1.294\\ (1.051e1559)\\ 1.294\\ (1.051e1552)\\ 1.294\\ (1.051e1552)\\ 1.294\\ \end{array}$ | $\begin{array}{c} 1.131\\ (1.948 \text{ to } 1.211)\\ 1.131\\ (1.948 \text{ to } 1.211)\\ 1.291\\ (1.06 \text{ to } 1.539)\\ 1.291\\ (1.06 \text{ to } 1.539)\\ 1.297\\ (1.06 \text{ to } 1.539)\\ 1.297\\ (1.06 \text{ to } 1.561)\\ 1.297\\ (1.96 \text{ to } 1.561)\\ 1.294\\ (1.951 \text{ to } 1.562)\\ 1.294\end{array}$ | $\begin{array}{c} 1.31\\ (1.08 \pm 0.211)\\ 1.31\\ (1.08 \pm 0.211)\\ 1.291\\ (1.06 \pm 0.539)\\ 1.291\\ (1.06 \pm 0.539)\\ 1.297\\ (1.06 \pm 0.561)\\ 1.297\\ (1.06 \pm 0.561)\\ 1.294\\ (1.089 \pm 0.569)\\ 1.294\\ (1.081 \pm 0.562)\end{array}$
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(5.6.49 to 11.322)
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8,23 | $\begin{array}{c} (4.38) \approx 10.056)\\ 7.423\\ (4.38) \approx 10.056)\\ 8.11\\ (4.324 \approx 13.462)\\ 8.23\\ (4.324 \approx 13.462)\\ 8.233\\ (4.709 \approx 13.126)\\ 8.223\\ (4.709 \approx 13.126)\\ 8.223\\ (4.704 \approx 13.144)\\ 8.292\\ (4.754 \approx 13.154)\\ 8.292\\ (4.754 \approx 13.519)\\ 8.437\\ (4.945 \approx 13.539)\\ \end{array}$ | 3.3.04
(4.203 to 6.843)
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(4.203 to 6.843)
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(4.285 to 9.239)
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(4.814 to 9.271)
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(4.739 to 9.26)
 | $\begin{array}{c} 3.546\\ (4.139 \mbox{ to 6233})\\ 5.646\\ (4.139 \mbox{ to 6233})\\ 6.756\\ (4.705 \mbox{ to $9,168$})\\ 6.756\\ (4.705 \mbox{ to $9,168$})\\ 6.726\\ (4.762 \mbox{ to $9,201$})\\ 6.687\\ (4.739 \mbox{ to $9,181$})\\ 6.687\\ (4.739 \mbox{ to $9,181$})\\ 6.734\\ (4.811 \mbox{ to $9,238$})\\ 6.734\\ (4.811 \mbox{ to $9,238$})\\ 6.734\\ (4.811 \mbox{ to $9,238$})\\ \end{array}$ | 3.07
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(4.273 to 8.413) | $\begin{array}{c} 2.446\\ (2322 w 2.774)\\ 2.546\\ (2322 w 2.774)\\ 5.624\\ (3.954 w 7.821)\\ 5.624\\ (3.954 w 7.821)\\ 5.638\\ (3.954 w 7.821)\\ 5.638\\ (3.943 w 7.822)\\ 5.638\\ (3.943 w 7.822)\\ 5.66\\ (4.039 w 7.875)\\ 5.66\\ (3.965 w 7.879)\\ 5.629\\ (3.965 w 7.879)\\ 5.629\\ (3.965 w 7.879)\\ \end{array}$
 | $\begin{array}{c} 2.554\\ (2.167\ w > 2.752)\\ 2.564\\ (2.167\ w > 2.752)\\ 5.61\\ (3.96\ w > 7.69)\\ 5.64\\ (3.96\ w > 7.69)\\ 5.764\\ (4.007\ w > 8.022)\\ 5.64\\ (4.007\ w > 8.022)\\ 5.64\\ (3.96\ w > 7.534)\\ 5.54\\ (3.96\ w > 7.534)\\ 5.589\\ (3.338\ w > 7.693)\\ 5.589\\ (3.338\ w > 7.693)\\ 5.589\\ (3.338\ w > 7.693)\end{array}$ | $\begin{array}{c} 1.849\\ (1.712\ {\rm m}\ 1.995)\\ 1.849\\ (1.712\ {\rm m}\ 1.995)\\ 3.643\\ (2.576\ {\rm m}\ 5.019)\\ 3.638\\ (2.576\ {\rm m}\ 5.019)\\ 3.638\\ (2.576\ {\rm m}\ 4.959)\\ 3.638\\ (2.576\ {\rm m}\ 4.959)\\ 3.62\\ (2.541\ {\rm m}\ 5.01)\\ 3.62\\ (2.484\ {\rm m}\ 5.01)\\ 3.591\\ (2.489\ {\rm m}\ 5.087)\\ 3.594\\ (2.489\ {\rm m}\ 5.087)\\ \end{array}$ | $\begin{array}{c} 1.85\\ (1.701\ m\ 1.995)\\ 1.23\\ (1.701\ m\ 1.995)\\ 3.601\\ (2.494\ m\ 4.936)\\ 3.691\\ (2.494\ m\ 4.936)\\ 3.591\\ (2.581\ m\ 4.981)\\ 3.591\\ (2.581\ m\ 4.981)\\ 3.583\\ (2.511\ m\ 5.117)\\ 3.583\\ (2.511\ m\ 5.117)\\ 3.583\\ (2.511\ m\ 5.117)\\ 3.607\\ (2.48\ m\ 5.948)\\ 3.607\\ (2.48\ m\ 5.948)\\ \end{array}$ | $\begin{array}{c} 1.45\\ (1.372 \approx 1.526)\\ 1.45\\ (1.372 \approx 1.526)\\ 2.17\\ (1.597 \approx 2.34)\\ 2.162\\ (1.64 \approx 2.316)\\ 2.162\\ (1.64 \approx 2.316)\\ 2.163\\ (1.66 \approx 2.302)\\ 2.153\\ (1.66 \approx 2.302)\\ 2.144\\ (1.619 \approx 2.33)\\ 2.144\\ (1.619 \approx 2.33)\\ \end{array}$
 | $\begin{array}{c} 1.454\\ (1.378 = 1.529)\\ 1.454\\ (1.378 = 1.529)\\ 2.146\\ (1.568 = 2.862)\\ 2.146\\ (1.568 = 2.862)\\ 2.168\\ (1.511 = 2.827)\\ 2.168\\ (1.511 = 2.827)\\ 2.173\\ (1.645 = 2.807)\\ 2.173\\ (1.638 = 2.822)\\ 2.173\\ (1.638 = 2.822)\\ 2.173\\ (1.638 = 2.822)\\ 2.173\\ 1.638 = 2.8222)\end{array}$ | $\begin{array}{c} 1.131\\ (1.488\ is\ 1.211)\\ 1.131\\ (1.048\ is\ 1.211)\\ 1.291\\ (1.66\ is\ 1.59)\\ 1.291\\ (1.66\ is\ 1.59)\\ 1.297\\ (1.66\ is\ 1.561)\\ 1.297\\ (1.66\ is\ 1.561)\\ 1.294\\ (1.69\ is\ 1.562)\\ 1.294\\ (1.681\ is\ 1.562)\\ 1.294\\ (1.681\ is\ 1.562)\end{array}$ | $\begin{array}{c} 1.131\\ (1.948 \ {\rm m}\ 1.211)\\ 1.131\\ (1.948 \ {\rm m}\ 1.211)\\ 1.291\\ (1.948 \ {\rm m}\ 1.211)\\ 1.291\\ (1.94 \ {\rm m}\ 1.539)\\ 1.291\\ (1.94 \ {\rm m}\ 1.539)\\ 1.291\\ (1.94 \ {\rm m}\ 1.561)\\ 1.294\\ (1.95 \ {\rm m}\ 1.562)\\ 1.294\\ (1.95 \ {\rm m}\ 1.56$ | $\begin{array}{c} 1.31\\ (1.08 \pm 0.2(1)\\ 1.31\\ (1.08 \pm 0.2(1)\\ 1.59)\\ (1.08 \pm 0.2(1)\\ 1.59)\\ (1.08 \pm 0.159)\\ 1.29\\ (1.08 \pm 0.59)\\ 1.291\\ (1.08 \pm 0.56)\\ 1.294\\ (1.099 \pm 1.569)\\ 1.294\\ (1.081 \pm 0.562)\\ 1.294\\ $
 | (1.948 s 211)
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(1.061 s. 1562) | $\begin{array}{c} 1.131\\ (1.048\ {\rm m}\ 1.211)\\ 1.291\\ (1.06\ {\rm m}\ 1.539)\\ 1.291\\ (1.06\ {\rm m}\ 1.539)\\ 1.297\\ (1.06\ {\rm m}\ 1.561)\\ 1.297\\ (1.06\ {\rm m}\ 1.561)\\ 1.294\\ (1.05\ {\rm m}\ 1.569)\\ 1.294\\ (1.081\ {\rm m}\ 1.562)\\ 1.294\\ (1.081\ {\rm m}\ 1.562)\\ 1.294\\ (1.081\ {\rm m}\ 1.562)\\ 1.294\\ \end{array}$ |
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(4.556 wa 13.285)</td> <td>$\begin{array}{c} (4.38) \approx 10.056)\\ 7.423\\ (4.38) \approx 10.056)\\ 8.11\\ (4.328 \approx 13.462)\\ 8.23\\ (4.328 \approx 13.462)\\ 8.233\\ (4.709 \approx 13.126)\\ 8.233\\ (4.709 \approx 13.126)\\ 8.232\\ (4.758 \approx 13.126)\\ 8.292\\ (4.754 \approx 13.144)\\ 8.497\\ (4.945 \approx 13.59)\\ 8.437\\ (4.945 \approx 13.519)\\ 8.437\\ (4.712 \approx 13.3137)\\ \end{array}$</td> <td>3.3.94
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4.2.93 to 6.3.43)
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(4.735 to 9.3.4)
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(4.730 to 9.3.4)
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(4.739 to 9.2.6)
6.771
(4.736 to 9.4)</td> <td>$\begin{array}{c} 3.566\\ (4.39) = 6.6233)\\ 5.456\\ (4.39) = 6.6233)\\ 6.756\\ (4.30) = 0.6233)\\ 6.756\\ (4.705 = 9.168)\\ 6.756\\ (4.705 = 9.201)\\ 6.726\\ (4.705 = 9.201)\\ 6.726\\ (4.705 = 9.201)\\ 6.637\\ (4.739 = 9.181)\\ 6.637\\ (4.739 = 9.181)\\ 6.637\\ (4.739 = 9.181)\\ 6.74\\ (4.811 = 9.228)\\ 6.73\\ (4.731 = 9.238)\\ 6.73\\ (4.731 = 9.419)\end{array}$</td> <td>3.07
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(4.20 tso 8.003)</td> <td>$\begin{array}{c} 3.081\\ (2.715 + 3.498)\\ 1.081\\ (2.715 + 3.498)\\ 5.987\\ (4.304 + 8.407)\\ (4.304 + 8.407)\\ (4.304 + 8.407)\\ (4.304 + 8.8107)\\ (4.344 + 8.847)\\ (4.344 + 8.847)\\ (4.347 + 8.821)\\ 5.981\\ (4.273 + 8.8413)\\ 5.941\\ (4.273 + 8.8413)\\ 5.941\\ (4.176 + 8.222)\end{array}$</td> <td>$\begin{array}{c} 2.446\\ (2.312 \pm 2.774)\\ 3.246\\ (2.352 \pm 2.774)\\ 5.624\\ (3.954 \pm 7.821)\\ 5.624\\ (3.954 \pm 7.821)\\ 5.638\\ (3.943 \pm 7.822)\\ 5.638\\ (3.943 \pm 7.822)\\ 5.638\\ (3.943 \pm 7.822)\\ 5.66\\ (4.079 \pm 7.875)\\ 5.629\\ (3.965 \pm 7.879)\\ 5.62\\ ($</td> <td>$\begin{array}{c} 2.554\\ (2.167 \approx 2.752)\\ 2.554\\ (2.367 \approx 2.752)\\ 5.541\\ (3.96 \approx 7.69)\\ 5.541\\ (3.96 \approx 7.69)\\ 5.504\\ (4.007 \approx 8.022)\\ 5.504\\ (4.007 \approx 8.022)\\ 5.541\\ (3.46 \approx 7.534)\\ 5.599\\ (3.338 \approx 7.693)\\ 5.549\\ (3.338 \approx 7.693)\\ 5.528\\ (4.002 \approx 7.742)\end{array}$</td> <td>$\begin{array}{c} 1.849\\ (1,712 \mbox{in} 1.998)\\ 1.849\\ (1,712 \mbox{in} 1.998)\\ 3.643\\ (2,576 \mbox{in} 5.019)\\ 3.643\\ (2,576 \mbox{in} 5.019)\\ 3.638\\ (2,576 \mbox{in} 4.959)\\ 3.638\\ (2,576 \mbox{in} 4.959)\\ 3.62\\ (2,484 \mbox{in} 5.01)\\ 3.62\\ (2,489 \mbox{in} 5.087)\\ 3.591\\ (2,489 \mbox{in} 5.087)\\ 3.625\\ (2,98 \mbox{in} 5.132)\\ \end{array}$</td> <td>$\begin{array}{c} 1.85\\ (1.207 = 1.995)\\ 1.25\\ (1.270 = 1.995)\\ 3.60\\ (2.494 = 4.956)\\ 3.60\\ (2.494 = 4.956)\\ 1.594\\ (2.585 = 4.984)\\ 3.586\\ (2.513 = 5.117)\\ 3.586\\ (2.513 = 5.117)\\ 3.607\\ (2.48 = 5.948)\\ 3.625\\ (2.558 = 5.046)\\ 3.625\\ (2.558 = 5.056)\\ \end{array}$</td> <td>$\begin{array}{c} 1.45\\ (1.372\pm1528)\\ 1.46\\ (1.372\pm1528)\\ 2.17\\ (1.579\pm2.84)\\ 2.17\\ (1.579\pm2.84)\\ 2.162\\ (1.64\pm2.816)\\ 2.162\\ (1.64\pm2.816)\\ 2.162\\ (1.64\pm2.816)\\ 2.138\\ (1.669\pm2.802)\\ 2.138\\ (1.669\pm2.802)\\ 2.144\\ (1.619\pm2.83)\\ 2.17\\ (1.614\pm2.802)\\ \end{array}$</td> <td>$\begin{array}{c} 1.454\\ (1.378 = 1.529)\\ 1.454\\ (1.378 = 1.529)\\ 2.146\\ (1.568 = 2.562)\\ 2.168\\ (1.518 = 2.827)\\ 2.168\\ (1.518 = 2.827)\\ 2.168\\ (1.518 = 2.827)\\ 2.173\\ (1.638 = 2.822)\\ 2.173\\ (1.638 = 2.822)\\ 2.17\\ (1.638 = 2.822)\\ 2.17\\ (1.640 = 2.881)\\ \end{array}$</td> <td>$\begin{array}{c} 1.131\\ (1.408\ m-1.211)\\ 1.131\\ (1.408\ m-1.21)\\ 1.291\\ (1.60\ m-1.59)\\ 1.291\\ (1.60\ m-1.591)\\ 1.297\\ (1.60\ m-1.561)\\ 1.294\\ (1.659\ m-1.569)\\ 1.294\\ (1.659\ m-1.562)\\ 1.294\\ (1.681\ m-1.$</td> <td>$\begin{array}{c} 1.11\\ (1.086 n - 1.211)\\ (1.086 n - 1.211)\\ 1.231\\ (1.046 n - 1.211)\\ 1.231\\ (1.046 n - 1.539)\\ 1.237\\ (1.066 n - 1.561)\\ 1.237\\ (1.066 n - 1.561)\\ 1.234\\ (1.051 n - 1.562)\\ 1.234\\ (1.061 n - 1.548)\\ 1.541\\ 1$</td> <td>$\begin{array}{c} 1.31\\ (1.088 = 1.211)\\ 1.31\\ (1.088 = 1.211)\\ (1.081 = 1.231)\\ (1.081 = 1.339)\\ 1.291\\ (1.081 = 1.339)\\ 1.291\\ (1.081 = 1.591)\\ 1.291\\ (1.081 = 1.562)\\ 1.294\\ (1.081 =
1.562)\\ 1.294\\ (1.081 = 1.562)\\ 1.294\\ (1.081 =$</td> <td>(1.045 s- 1.21)
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 | $\begin{array}{c} (4.38) \approx 10.056)\\ 7.423\\ (4.38) \approx 10.056)\\ 8.11\\ (4.328 \approx 13.462)\\ 8.23\\ (4.328 \approx 13.462)\\ 8.233\\ (4.709 \approx 13.126)\\ 8.233\\ (4.709 \approx 13.126)\\ 8.232\\ (4.758 \approx 13.126)\\ 8.292\\ (4.754 \approx 13.144)\\ 8.497\\ (4.945 \approx 13.59)\\ 8.437\\ (4.945 \approx 13.519)\\ 8.437\\ (4.712 \approx 13.3137)\\ \end{array}$ | 3.3.94
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 | $\begin{array}{c} 3.081\\ (2.715 + 3.498)\\ 1.081\\ (2.715 + 3.498)\\ 5.987\\ (4.304 + 8.407)\\ (4.304 + 8.407)\\ (4.304 + 8.407)\\ (4.304 + 8.8107)\\ (4.344 + 8.847)\\ (4.344 + 8.847)\\ (4.347 + 8.821)\\ 5.981\\ (4.273 + 8.8413)\\ 5.941\\ (4.273 + 8.8413)\\ 5.941\\ (4.176 + 8.222)\end{array}$ | $\begin{array}{c} 2.446\\ (2.312 \pm 2.774)\\ 3.246\\ (2.352 \pm 2.774)\\ 5.624\\ (3.954 \pm 7.821)\\ 5.624\\ (3.954 \pm 7.821)\\ 5.638\\ (3.943 \pm 7.822)\\ 5.638\\ (3.943 \pm 7.822)\\ 5.638\\ (3.943 \pm 7.822)\\ 5.66\\ (4.079 \pm 7.875)\\ 5.629\\ (3.965 \pm 7.879)\\ 5.62\\ ($ | $\begin{array}{c} 2.554\\ (2.167 \approx 2.752)\\ 2.554\\ (2.367 \approx 2.752)\\ 5.541\\ (3.96 \approx 7.69)\\ 5.541\\ (3.96 \approx 7.69)\\ 5.504\\ (4.007 \approx 8.022)\\ 5.504\\ (4.007 \approx 8.022)\\ 5.541\\ (3.46 \approx 7.534)\\ 5.599\\ (3.338 \approx 7.693)\\ 5.549\\ (3.338 \approx 7.693)\\ 5.528\\ (4.002 \approx 7.742)\end{array}$ | $\begin{array}{c} 1.849\\ (1,712 \mbox{in} 1.998)\\ 1.849\\ (1,712 \mbox{in} 1.998)\\ 3.643\\ (2,576 \mbox{in} 5.019)\\ 3.643\\ (2,576 \mbox{in} 5.019)\\ 3.638\\ (2,576 \mbox{in} 4.959)\\ 3.638\\ (2,576 \mbox{in} 4.959)\\ 3.62\\ (2,484 \mbox{in} 5.01)\\ 3.62\\ (2,489 \mbox{in} 5.087)\\ 3.591\\ (2,489 \mbox{in} 5.087)\\ 3.625\\ (2,98 \mbox{in} 5.132)\\ \end{array}$
 | $\begin{array}{c} 1.85\\ (1.207 = 1.995)\\ 1.25\\ (1.270 = 1.995)\\ 3.60\\ (2.494 = 4.956)\\ 3.60\\ (2.494 = 4.956)\\ 1.594\\ (2.585 = 4.984)\\ 3.586\\ (2.513 = 5.117)\\ 3.586\\ (2.513 = 5.117)\\ 3.607\\ (2.48 = 5.948)\\ 3.625\\ (2.558 = 5.046)\\ 3.625\\ (2.558 = 5.056)\\ \end{array}$ | $\begin{array}{c} 1.45\\ (1.372\pm1528)\\ 1.46\\ (1.372\pm1528)\\ 2.17\\ (1.579\pm2.84)\\ 2.17\\ (1.579\pm2.84)\\ 2.162\\ (1.64\pm2.816)\\ 2.162\\ (1.64\pm2.816)\\ 2.162\\ (1.64\pm2.816)\\ 2.138\\ (1.669\pm2.802)\\ 2.138\\ (1.669\pm2.802)\\ 2.144\\ (1.619\pm2.83)\\ 2.17\\ (1.614\pm2.802)\\ \end{array}$ | $\begin{array}{c} 1.454\\ (1.378 = 1.529)\\ 1.454\\ (1.378 = 1.529)\\ 2.146\\ (1.568 = 2.562)\\ 2.168\\ (1.518 = 2.827)\\ 2.168\\ (1.518 = 2.827)\\ 2.168\\ (1.518 = 2.827)\\ 2.173\\ (1.638 = 2.822)\\ 2.173\\ (1.638 = 2.822)\\ 2.17\\ (1.638 = 2.822)\\ 2.17\\ (1.640 = 2.881)\\ \end{array}$ | $\begin{array}{c} 1.131\\ (1.408\ m-1.211)\\ 1.131\\ (1.408\ m-1.21)\\ 1.291\\ (1.60\ m-1.59)\\ 1.291\\ (1.60\ m-1.591)\\ 1.297\\ (1.60\ m-1.561)\\ 1.294\\ (1.659\ m-1.569)\\ 1.294\\ (1.659\ m-1.562)\\ 1.294\\ (1.681\ m-1.$ | $\begin{array}{c} 1.11\\ (1.086 n - 1.211)\\ (1.086 n - 1.211)\\ 1.231\\ (1.046 n - 1.211)\\ 1.231\\ (1.046 n - 1.539)\\ 1.237\\ (1.066 n - 1.561)\\ 1.237\\ (1.066 n - 1.561)\\ 1.234\\ (1.051 n - 1.562)\\ 1.234\\ (1.061 n - 1.548)\\ 1.541\\
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2715 9-100 | $\begin{array}{c} 2.546\\ (.2312 \pm 2.724)\\ (.2312 \pm 2.724)\\ (.2312 \pm 2.724)\\ (.2312 \pm 2.724)\\ (.2544 \pm 7.224)\\ (.2544 \pm $ | $\begin{array}{c} 2.584\\ 2.584\\ 2.584\\ 2.584\\ 2.584\\ 2.584\\ 3.$ | $\begin{array}{c} 1.849\\ (1.712-1.924)\\ 1.849\\ (1.712-1.924)\\ 1.5756-1.524\\ (1.712-1.924)\\ 1.5756-1.5249\\ (1.5756-1.524)\\ 1.5756-1.5249\\ (1.5756-1.524)\\ 1.556-1.524\\ (1.576-1.524)\\ 1.556\\ 1.556-1.524\\ 1.556\\ 1$
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1.22 \\ 1.$ | $\begin{array}{c} 1.13\\ (1.68 = 1.21)\\ 1.291\\ (1.68 = 1.59)\\ (1.68 = 1.59)\\ 1.291\\ (1.68 = 1.59)\\ 1.291\\ (1.68 = 1.59)\\ 1.294\\ (1.69 = 1.59)\\ 1.294\\ (1.69 = 1.59)\\ 1.294\\ (1.69 = 1.59)\\ 1.294\\ (1.69 = 1.59)\\ 1.294\\ (1.69 = 1.59)\\ 1.294\\ (1.69 = 1.59)\\ 1.294\\ (1.69 = 1.59)\\ 1.294\\ (1.69 = 1.59)\\ 1.294\\ (1.69 = 1.59)\\ 1.294\\ (1.69 = 1.59)\\ 1.294\\ 1.$ |
| Mediated years in the market plant of the Mediated Hermiter (Mediated Hermiter) and Mediated Hermiter (Mediated Hermiter) (Mediated Hermit | big harpsone, -50-88 big harpsone, -50-88 fig harpsone, -50-88 big harpsone, -50-88

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(1.378 n 1.529)
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1. | $\begin{array}{c} 1.11\\ (1.681\pm1.21)\\ (1.681\pm1.21)\\ (1.681\pm1.21)\\ (1.681\pm1.21)\\ (1.681\pm1.23)\\ (1.681\pm1.23)\\ (1.681\pm1.23)\\ (1.681\pm1.23)\\ (1.681\pm1.53)\\ (1.$ | $\begin{array}{c} 1.13\\ (1.08) = 1210\\ (1.08) = 1210\\ (1.08) = 1210\\ (1.08) = 1210\\ (1.08) = 1210\\ (1.08) = 120$ | 1.13
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(| $\begin{array}{c} (.68 \pm11) \\ (.1.88 \pm11) \\ (.1.88 \pm11) \\29 \\ (.1.8 \pm12) \\29 \\ (.1.8 \pm13) \\29 $ | $\begin{array}{c} 1.13\\ (1.086 = 1.21)\\ 1.291\\ (1.66 = 1.59)\\ 1.291\\ (1.66 = 1.59)\\ 1.291\\ (1.66 = 1.59)\\ 1.291\\ (1.66 = 1.59)\\ 1.294\\ (1.879 = 1.59)\\ 1.294\\ (1.879 = 1.59)\\ 1.294\\ (1.879 = 1.59)\\ 1.294\\ (1.819 = 1.59)\\ 1.294\\ (1.819 = 1.59)\\ 1.294\\ (1.819 = 1.59)\\ 1.294\\ (1.819 = 1.59)\\ 1.294\\ (1.819 = 1.59)\\ 1.29\\ 1.29\\ (1.819 = 1.59)\\ 1.29\\ 1.29\\ 1.29\\ 1.29\\ 1.29\\ 1.29\\ 1.29\\ 1.29\\ 1.29\\
1.29\\ 1$ |
| Mediated years in the | table sparsers, >>>0.88 table sparsers, >>>0.88 table sparsers, >>>>0.88 table sparsers, >>>>0.88 table sparsers, >>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>>

 | Mabahary in para
Matshafay in para | | | | | | | | $\begin{array}{c} 7,07\\ (5,89 \mbox{ with 123)}\\ (6,89 \mbox{ with 123)}\\ (6,89 \mbox{ with 123)}\\ (6,80 \mbox{ with 133)}\\ (6,80 \m$ | $\begin{array}{c} (4.26 \pm 0.056) \\ 7.42 \\ 1.42 \\ 1.41 \\ 4.13 \\ 4.14 \\$ | $\begin{array}{c} (23)^{-100}$ | (3) 300 400 400 400 400 400 400 400 400 400
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2715 - 271 | $\begin{array}{c} 1.546\\ (.2312 \pm 2.754)\\ (.2312 \pm 7.254)\\ (.2312 \pm $ | 2.554
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(1.614 = 2.217)
 | $\begin{array}{c} 1.11\\ (.168 + 1.21)\\ (.168 + 1.21)\\ 1.23\\ (.168 + 1.23)\\ 1.23\\ (.268 + 1.23)\\ 1.23\\ (.268 + 1.23)\\ 1.24\\ (.268 + 1.23)\\ 1.24\\ (.268 + 1.23)\\ 1.24\\ (.268 + 1.23)\\ 1.24\\ (.268 + 1.24)\\ 1.24\\ (.268 + 1.24)\\ 1.24\\ (.268 + 1.24)\\ 1.24\\ (.268 + 1.24)\\ 1.24\\ (.268 + 1.24)\\ 1.24\\ 1.$ | $\begin{array}{c} 1.01\\ (1.086 = 1.21)\\ (1.086 = 1.21)\\ (1.086 = 1.21)\\ (1.086 = 1.21)\\ (1.086 = 1.21)\\ (1.086 = 1.09)\\ (1.097 = 1.09)\\ ($ | 1.131
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appendix rable of Reality isk	s used by age and sex for each outco	1010 00 001 1155 HECO12	s except fo	or ambient air pol	llution alcohol, a	and smoking.									А	iges											
Risk - Outcome	Category / Units	Morbidity / Mortality	Sex	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Complete hearing loss due to ag related and other hearing loss	p- Low exposure, 85-90dB	Morbidity	Both								3.004 (1.775 to 4.913)	2.986 (1.804 to 4.882)	3.482 (2.473 to 4.715)	3.478 (2.449 to 4.702)	3.858 (2.702 to 5.284)	3.813 (2.745 to 5.341)	3.917 (2.805 to 5.383)	3.985 (2.826 to 5.477)	2.701 (1.874 to 3.658)	2.695 (1.885 to 3.731)	1.812 (1.376 to 2.371)	1.824 (1.359 to 2.407)	1.222 (1.015 to 1.475)				
Complete hearing loss with ring due to age-related and other hear	ing Low exposure, 85-90dB	Morbidity	Both								3.004 (1.775 to 4.913)	2.986 (1.804 to 4.882)	3.482 (2.473 to 4.715)	3.478 (2.449 to 4.702)	3.858 (2.702 to 5.284)	3.813 (2.745 to 5.341)	3.917 (2.805 to 5.383)	3.985 (2.826 to 5.477)	2.701 (1.874 to 3.658)	2.695 (1.885 to 3.731)	1.812 (1.376 to 2.371)	1.824 (1.359 to 2.407)	1.222 (1.015 to 1.475)				
Mild hearing loss due to age-reli and other hearing loss	ated No exposure	Morbidity	Both								1.0 (1.0 to 1.0)																
Mild hearing loss with ringing d to ane-related and other hearing	lae No esposure	Morbidity	Both								1.0 (1.0 to 1.0)																
Moderate hearing loss due to age	P- No exposure	Morbidity	Both								1.0 (1.0 m 1.0)	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
Moderate hearing loss with ringi due to age-related and other hear	ing ring No exposure	Morbidity	Both								1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
Moderately severe hearing loss of	hae No exposure	Morbidity	Both								(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0) 1.0
to age-related and other hearing Moderately severe hearing loss v ringing due to age-related and et	aith her. No execute	Marbidity	Brath								(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0) 1.0
Severe hearing loss with ringing	dae No empoure	Marbidity	Bath								(1.0 to 1.0) 1.0																
to age-related and other hearing Severe hearing loss due to age-	loss No equalit	Mathing .	D. d								(1.0 to 1.0) 1.0																
related and other hearing loss Profound hearing loss date to an	No exposure	Morbidity	Both								(1.0 to 1.0) 1.0																
related and other hearing loss Profound hearing loss with ringi	" No exposure	Morbidity	Both								(1.0 to 1.0)																
due to age-related and other heat	ring No exposure	Morbidity	Both								(1.0 to 1.0)																
Complete hearing loss did to ag related and other hearing loss Complete hearing loss with rine	e- No exposure	Morbidity	Both								(1.0 to 1.0)																
due to age-related and other heat	ring No exposure	Morbidity	Both					r r			(1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	(1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	(1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Occupational ergonomic fact	ors																										
Low back pain	Professional, technical and related workers	Morbidity	Both								1.173 (1.066 to 1.282)	1.172 (1.062 to 1.283)	1.169 (1.065 to 1.283)	1.17 (1.062 to 1.284)	1.17 (1.062 to 1.285)	1.172 (1.062 to 1.283)	1.171 (1.071 to 1.27)	1.169 (1.063 to 1.288)	1.171 (1.059 to 1.281)	1.17 (1.058 to 1.286)	1.17 (1.07 to 1.279)	1.172 (1.065 to 1.287)	1.172 (1.07 to 1.283)	1.172 (1.07 to 1.283)			
Low back pain	Administrative and managerial workers	Morbidity	Both								1.211 (0.964 to 1.508)	1.21 (0.964 to 1.492)	1.209 (0.965 to 1.487)	1.209 (0.963 to 1.524)	1.207 (0.976 to 1.496)	1.207 (0.965 to 1.5)	1.205 (0.946 to 1.489)	1.205 (0.967 to 1.472)	1.205 (0.961 to 1.509)	1.203 (0.948 to 1.515)	1.209 (0.976 to 1.479)	1.21 (0.964 to 1.49)	1.203 (0.961 to 1.501)	1.203 (0.961 to 1.501)			
Low back pain	Clerical and related workers	Morbidity	Both								1.0 (1.0 to 1.0)																
Low back pain	Sales workers	Morbidity	Both								1.22 (1.029 to 1.434)	1.21 (1.018 to 1.418)	1.213 (1.028 to 1.434)	1.214 (1.004 to 1.448)	1.207 (1.017 to 1.445)	1.218 (1.016 to 1.455)	1.212 (1.012 to 1.444)	1.216 (1.01 to 1.448)	1.219 (1.019 to 1.45)	1.211 (1.014 to 1.444)	1.213 (1.015 to 1.455)	1.21 (1.007 to 1.423)	1.214 (1.017 to 1.446)	1.214 (1.017 to 1.446)			
Low back pain	Service workers	Morbidity	Both								1.472 (1.385 to 1.568)	1.472 (1.383 to 1.569)	1.471 (1.372 to 1.563)	1.472 (1.382 to 1.571)	1.469 (1.378 to 1.567)	1.472 (1.377 to 1.57)	1.469 (1.375 to 1.568)	1.47 (1.378 to 1.57)	1.472 (1.379 to 1.575)	1.472 (1.381 to 1.572)	1.474 (1.386 to 1.571)	1.47 (1.377 to 1.568)	1.472 (1.38 to 1.571)	1.472 (1.38 to 1.571)			
Low back pain	Agriculture, animal husbandry and forestry workers, fishermen and hunters	Morbidity	Both								3.789 (2.58 to 5.376)	3.762 (2.621 to 5.284)	3.869 (2.642 to 5.486)	3.775 (2.569 to 5.369)	3.774 (2.606 to 5.314)	3.771 (2.532 to 5.317)	3.793 (2.632 to 5.361)	3.785 (2.556 to 5.333)	3.776 (2.645 to 5.157)	3.792 (2.536 to 5.421)	3.802 (2.684 to 5.428)	3.746 (2.609 to 5.175)	3.77 (2.635 to 5.151)	3.77 (2.635 to 5.151)			
Low back pain	Production and related workers, transport equipment operators and labourers	Morbidity	Both								1.543 (1.409 to 1.679)	1.54 (1.406 to 1.676)	1.542 (1.415 to 1.677)	1.543 (1.413 to 1.695)	1.542 (1.416 to 1.685)	1.543 (1.418 to 1.685)	1.541 (1.402 to 1.684)	1.542 (1.41 to 1.684)	1.541 (1.404 to 1.683)	1.54 (1.414 to 1.677)	1.54 (1.408 to 1.683)	1.538 (1.408 to 1.673)	1.541 (1.41 to 1.677)	1.541 (1.41 to 1.677)			
Low back pain	Background	Morbidity	Both								1.0 (1.0 to 1.0)																
Non-exclusive breastfeeding																											
Lower respiratory infections	None	Both	Both			1.739	1.739	1		1		1		1	1	1				1	1			1	1		
Lower respiratory infections	Partial	Both	Both			1.483	1.483																				
Lower respiratory infections	Predominant	Both	Both			(1.206 to 1.792) 1.369	(1.206 to 1.792) 1.369																				
Lower rearistory infections	Furbraise	Breth	Brath			(1.055 to 1.8)	(1.055 to 1.8)																				
Disabased Server	Nome	Buch	Rush			(1.0 to 1.0) 3.605	(1.0 to 1.0) 3.605																				
Product Process	b. d.t.					(2.716 to 4.703) 2.633	(2.716 to 4.703) 2.633																				
Diamocal discuss	Parta	nom	Dom			(1.942 to 3.481) 2.346	(1.942 to 3.481) 2.346																				
Duarthocal docuses	Predominant	Both	Both			(1.667 to 3.234) 1.0	(1.667 to 3.234) 1.0																				
Duarhoeal docases	Exclusive	Both	Both			(1.0 to 1.0)	(1.0 to 1.0)																				
Discontinued breastreeding							1313	1313																			
Diarthoeal diseases	Continued	Both	Both				(1.111 to 1.549)	(1.111 to 1.549)																			
Diarrhoeal diseases	Not continued	Both	Both				(1.0 to 1.0)	(1.0 to 1.0)																			
Child underweight																											
Lower respiratory infections	ke 6->	Both	Both				2.593 (1.908 to 4.39)	(1.908 to 4.39)																			
Lower respiratory infections	<-2 sil	Both	Both				1.365 (1.215 to 1.755)	1.365 (1.215 to 1.755)																			
Lower respiratory infections	<-1 sal	Both	Both				1.145 (1.044 to 1.364)	1.145 (1.044 to 1.364)																			
Lower respiratory infections	-1 sd and above	Both	Both				1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																			
Diarthocal diseases	<-3 sal	Both	Both				2.332 (2.076 to 2.802)	2.332 (2.076 to 2.802)																			
Diarrhoeal diseases	<-2 sd	Both	Both				1.23 (1.163 to 1.314)	1.23 (1.163 to 1.314)																			
Diarrhoeal diseases	<-1 sal	Both	Both				1.088 (1.046 to 1.134)	1.088 (1.046 to 1.134)																			
Diarrhoeal diseases	-1 sd and above	Both	Both				1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																			
Measles	<-3 sal	Both	Both				5.668 (1.767 to 12.414)	5.668 (1.767 to 12.414)																			
Measles	<-2 sil	Both	Both				2.458 (1.26 to 5.118)	2.458 (1.26 to 5.118)																			
Measles	<-1 sd	Both	Both				0.995 (0.5 to 1.726)	0.995 (0.5 to 1.726)																			
Measles	-1 sd and above	Both	Both				1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																			
Child wasting																											
Lower respiratory infections	<-3 sal	Both	Both				47.67 (15.923 jn 94.87.6)	47.67 (15.923 p. 94 874)																			
Lower respiratory infections	<-2 sd	Both	Both				20.455	20.455																			
Lower respiratory infections	<-1 ad	Both	Both				5.941	(1064 to 37.929) 5.941																			
Lower respiratory infections	-1 sd and above	Both	Both				(1.972 to 11.992)	(1.972 to 11.992)																			
Diambocal diseases	<-3 ml	Both	Both				(1.0 to 1.0) 105.759	(1.0 to 1.0) 105.759																			
Disarboard Jonana	<-2 wl	Bente	Berk				(42.198 to 157.813) 23.261	(42.198 to 157.813) 23.261																			
Diamon a constant	e tud	Doin	Dom.				(9.02 to 35.845) 6.601	(9.02 to 35.845) 6.601																			
Diarrhoeal diseases	<-1 10	Both	Both				(2.158 to 11.243) 1.0	(2.158 to 11.243) 1.0																			
Diarrhoeal diseases	-1 sd and above	Both	Both				(1.0 to 1.0)	(1.0 to 1.0)																			

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Risk - Outcome	Category / Units	Morbidity / Mortali	ty Sex	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Measles	<-3 ad	Both	Both				37.936 (5.088 to 199.126)	37.936 (5.088 to 199.126)																			
Measles	<-2 ad	Both	Both				8.477 (1.33 to 42.777)	8.477 (1.33 to 42.777)																			
Measles	<-1 sd	Both	Both				1.833 (0.569 to 8.985)	1.833 (0.569 to 8.985)																			
Measles	-1 sd and above	Both	Both				1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																			
Child stunting																											
Lower respiratory infections	<-3 ad	Both	Both				2.355 (1.15 to 5.114)	2.355 (1.15 to 5.114)																			
Lower respiratory infections	<-2 ad	Both	Both				1.318 (1.014 to 2.165)	1.318 (1.014 to 2.165)																			
Lower respiratory infections	<-1 ad	Both	Both				1.158 (0.998 to 1.655)	1.158 (0.998 to 1.655)																			
Lower respiratory infections	-1 sd and above	Both	Both				1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																			
Diarrhoeal diseases	<-3 ad	Both	Both				1.851 (1.28 to 2.699)	1.851 (1.28 to 2.699)																			
Diarrhoeal diseases	<-2 ad	Both	Both				1.222 (1.067 to 1.5)	1.222 (1.067 to 1.5)																			
Diarrhocal diseases	<-1 ad	Both	Both				1.111 (1.023 to 1.273)	1.111 (1.023 to 1.273)																			
Diarrhoeal diseases	-1 sd and above	Both	Both				1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																			
Measles	<-3 ad	Both	Both				2.487 (1.129 to 6.528)	2.487 (1.129 to 6.528)																			
Measles	<-2 ad	Both	Both				1.54 (1.029 to 3.222)	1.54 (1.029 to 3.222)																			
Measles	<-1 ad	Both	Both				1.103 (0.861 to 1.719)	1.103 (0.861 to 1.719)																			
Measles	-1 sd and above	Both	Both				1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																			
Low birth weight and short ge	station																										
Lower respiratory infections	Birth prevalence - $\left[0,24\right)$ wks, $\left[0,500\right)$ g	Mortality	Males		1564.792 (1056.542 to 2116.062)	618.595 (458.842 to 812.921)																					
Lower respiratory infections	Birth prevalence - [0, 24) wks, [0, 500) g	Mortality	Females		(1050.664 to 2311.977)	713.571 (526.178 to 921.018)																					
Lower respiratory infections	Birth prevalence - [0, 24) wks, [500, 1000) g	Mortality	Males	0	1155.815 (825.412 to 1506.837)	457.5 (352.552 to 573.483)																					
Lower respiratory infections	Birth prevalence - [0, 24) wks, [500, 1000) g	Mortality	Females	0	1169.123 (802.003 to 1617.979)	515.406 (396.713 to 641.541)																					
Lower respiratory infections	Birth prevalence - [24, 26) wks, [500, 1000) 8	Mortality	Males	0	955.583 (723.748 to 1244.265)	443.357 (363.03 to 534.695)																					
Lower respiratory infections	Birth prevalence - [24, 26) wks, [500, 1000) g	Mortality	Females		947.143 (702.662 to 1237.093)	487.549 (387.307 to 603.498)																					
Lower respiratory infections	Birth prevalence - [26, 28) wks, [500, 1000) g	Mortality	Males		497.817 (377.617 to 648.547)	330.886 (261.438 to 401.709)																					
Lower respiratory infections	Birth prevalence - [26, 28) wks, [500, 1000) g	Mortality	Females		483.682 (354.946 to 629.517)	344.618 (274.427 to 419.864)																					
Lower respiratory infections	Birth prevalence - [30, 32) wks, [500, 1000) g	Mortality	Males		236.614 (163.821 to 324.502)	149.995 (117.866 to 188.368)																					
Lower respiratory infections	Birth prevalence - [30, 32) wks, [500, 1000) g	Mortality	Females		229.197 (157.606 to 317.194)	152.117 (120.779 to 190.583)																					
Lower respiratory infections	Birth prevalence - [28, 30) wks, [500, 1000) g	Mortality	Males		297.629 (214.953 to 396.586)	216.995 (173.321 to 271.466)																					
Lower respiratory infections	Birth prevalence - [28, 30) wks, [500, 1000) g	Mortality	Females		281.056 (198.176 to 386.635)	219.884 (174.264 to 272.704)																					
Lower respiratory infections	Birth prevalence - [26, 28) wks, [1000, 1500) g	Mortality	Males		267.91 (210.177 to 332.92)	164.167 (132.898 to 200.569)																					
Lower respiratory infections	Birth prevalence - [26, 28) wks, [1000, 1500) g	Mortality	Females		266.509 (197.461 to 346.932)	174.222 (137.431 to 217.349)																					
Lower respiratory infections	Birth prevalence - [34, 36) wks, [1000, 1500) g	Mortality	Males		142.056 (98.086 to 197.774)	52.86 (42.914 to 64.617)																					
Lower respiratory infections	Birth prevalence - [34, 36) wks, [1000, 1500) g	Mortality	Females		141.899 (95.864 to 197.656)	57.421 (46.452 to 71.339)																					
Lower respiratory infections	Birth prevalence - [28, 30) wks, [1500, 2000) g	Mortality	Males		127.966 (97.178 to 167.026)	50.018 (40.539 to 61.919)																					
Lower respiratory infections	Birth prevalence - [28, 30) wks, [1500, 2000) g	Mortality	Females		130.924 (96.513 to 172.188)	57.275 (46.36 to 70.038)																					
Lower respiratory infections	Birth prevalence - [28, 30) wks, [1000, 1500) g	Mortality	Males		158.563 (120.99 to 204.947)	103.32 (83.486 to 127.144)																					
Lower respiratory infections	Birth prevalence - [28, 30) wks, [1000, 1500) g	Mortality	Females		153.905 (112.327 to 200.786)	107.529 (86.954 to 131.78)																					
Lower respiratory infections	Birth prevalence - [32, 34) wks, [1000, 1500) g	Mortality	Males		117.142 (81.354 to 161.101)	53.185 (43.049 to 66.274)																					
Lower respiratory infections	Birth prevalence - [32, 34) wks, [1000, 1500) g	Mortality	Females		115.171 (79.363 to 159.206)	56.034 (45.982 to 68.36)																					
Lower respiratory infections	Birth prevalence - [30, 32) wks, [1000, 1500) g	Mortality	Males		119.308 (87.769 to 160.885)	67.163 (54.863 to 82.638)																					
Lower respiratory infections	Birth prevalence - [30, 32) wks, [1000, 1500) g	Mortality	Females		115.448 (84.272 to 156.425)	69.14 (55.873 to 85.012)																					
Lower respiratory infections	Birth prevalence - [37, 38) wks, [1500, 2000) g	Mortality	Males		62.972 (46.159 to 83.484)	24.148 (20.066 to 29.406)																					
Lower respiratory infections	Birth prevalence - [37, 38) wks, [1500, 2000) g	Mortality	Females		59.988 (43.974 to 79.053)	26.719 (21.746 to 32.816)																					
Lower respiratory infections	Birth prevalence - [36, 37] wks, [1500, 2000) g	Mortality	Males		60.218 (43.669 to 82.48)	23.031 (18.793 to 28.483)																					
Lower respiratory infections	Birth prevalence - [36, 37) wks, [1500, 2000) g	Mortality	Females		58.527 (42.172 to 80.557)	25.143 (20.331 to 30.566)																					
Lower respiratory infections	Birth prevalence - [30, 32) wks, [2000, 2500) g	Mortality	Males		67.971 (50.354 to 88.935)	18.03 (14.621 to 22.103)																					
Lower respiratory infections	Birth prevalence - [30, 32) wks, [2000, 2500) g	Mortality	Females		69.383 (49.108 to 94.583)	22.069 (17.836 to 27.163)																					
Lower respiratory infections	Birth prevalence - [30, 32) wks, [1500, 2000) g	Mortality	Males		77.369 (59.702 to 99.232)	31.079 (25.786 to 36.724)																					
Lower respiratory infections	Birth prevalence - [30, 32) wks, [1500, 2000) g	Mortality	Females		76.134 (56.885 to 100.996)	34.756 (28.764 to 41.849)																					
Lower respiratory infections	Birth prevalence - [34, 36) wks, [1500, 2000) g	Mortality	Males		55.555 (39.553 to 75.104)	21.346 (17.677 to 26.143)																					
Lower respiratory infections	Birth prevalence - [34, 36) wks, [1500, 2000) g	Mortality	Females		54.335 (38.617 to 75.24)	23.046 (18.743 to 28.287)																					
Lower respiratory infections	Birth prevalence - [32, 34) wks, [1500, 2000) g	Mortality	Males		57.155 (42.484 to 73.651)	23.114 (19.028 to 27.915)																					
Lower respiratory infections	Birth prevalence - [32, 34) wks, [1500, 2000) g	Mortality	Females		56.101 (39.794 to 76.295)	25.149 (20.615 to 30.388)																					
Lower respiratory infections	Birth prevalence - [32, 34) wks, [2000, 2500) g	Mortality	Males		37.444 (29.026 to 48.227)	12.233 (10.252 to 14.477)																					
Lower respiratory infections	Birth prevalence - [32, 34) wks, [2000, 2500) g	Mortality	Females		36.874 (26.658 to 49.653)	14.384 (12.095 to 17.03)																					
Lower respiratory infections	Birth prevalence - [40, 42) wks, [2000, 2500) g	Mortality	Males		18.092 (13.292 to 23.719)	9.23 (7.037 to 11.454)																					

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Risk - Outcome	Category / Units	Morbidity / Mortalit	v Sex	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Lower respiratory infections	Birth prevalence - [40, 42) wks, [2000, 26000 -	Mortality	Females		15.574	9.975																					
Lower respiratory infections	Birth prevalence - [38, 40) wks, [2000,	Mortality	Males		13.104	8.198																					
Lower restinatory infections	2500) g Birth prevalence - [38, 40) wks, [2000,	Manulity	Females		(9.829 to 16.99) 11.308	(6.786 to 9.959) 8.577																					
	2500) g Birth prevalence - [32, 34) wks, [2500,				(8.389 to 14.38) 33.063	(7.04 to 10.449) 8.441																					
Lower respiratory intections	3000) g Birth menalence - [32-34] wite [2500	stortany	Manos		(24.393 to 43.503) 32.812	(6.822 to 10.431) 10.398																					
Lower respiratory infections	3000) g	Mortality	Females		(23.439 to 45.567)	(8.227 to 13.042)																					
Lower respiratory infections	2500) g	Mortality	Males		(16.305 to 29.433)	(7.859 to 11.112)																					
Lower respiratory infections	Birth prevalence - [34, 36) wks, [2000, 2500) g	Mortality	Females		21.297 (15.657 to 28.761)	10.295 (8.548 to 12.273)																					
Lower respiratory infections	Birth prevalence - [37, 38) wks, [2000, 2500) g	Mortality	Males		13.0 (10.102 to 16.456)	8.096 (6.724 to 9.676)																					
Lower respiratory infections	Birth prevalence - [37, 38) wks, [2000, 2500) g	Mortality	Females		11.563 (8.805 to 15.11)	8.467 (6.994 to 10.342)																					
Lower respiratory infections	Birth prevalence - [36, 37) wks, [2000, 2500) g	Mortality	Males		14.401 (10.789 to 18.654)	8.221 (6.917 to 9.923)																					
Lower respiratory infections	Birth prevalence - [36, 37) wks, [2000, 2500) #	Mortality	Females		13.513 (9.817 to 17.947)	8.654 (7.215 to 10.369)																					
Lower respiratory infections	Birth prevalence - [34, 36) wks, [2500, 2000) -	Mortality	Males		13.419	5.562																					
Lower respiratory infections	Birth prevalence - [34, 36) wks, [2500,	Monality	Females		13.266	6.395																					
Lower restiratory infections	3000) g Birth prevalence - [34, 36) wks, [4000,	Manulity	Malex		(9.666 to 17.689) 23.096	(5.292 to 7.606) 2.895																					
i i i i i i i i i i i i i i i i i i i	4500) g Birth meyalence - [34, 36) wks. [4000.				(14.708 to 35.098) 25.038	(2.245 to 3.716) 3.778																					
Lower respiratory infections	4500) g Rightmansharar (24, 36) mbs (2000	Mortality	Females		(15.763 to 37.255)	(2.855 to 4.925)																					
Lower respiratory infections	3500) g	Mortality	Males		(10.222 to 18.478)	(3.449 to 5.338)																					
Lower respiratory infections	Barth prevalence - [34, 36) wks, [3000, 3500) g	Mortality	Females		14.375 (10.269 to 20.114)	5.265 (4.145 to 6.564)																					
Lower respiratory infections	Birth prevalence - [36, 37) wks, [2500, 3000) g	Mortality	Males		4.874 (4.014 to 5.713)	3.699 (3.134 to 4.374)																					
Lower respiratory infections	Birth prevalence - [36, 37) wks, [2500, 3000) g	Mortality	Females		4.609 (3.731 to 5.61)	3.898 (3.235 to 4.67)																					
Lower respiratory infections	Birth prevalence - [34, 36) wks, [3500, 4000) g	Mortality	Males		18.024 (12.279 to 25.547)	3.657 (2.838 to 4.675)																					
Lower respiratory infections	Birth prevalence - [34, 36) wks, [3500, 4000) #	Mortality	Females		19.263 (12.567 to 27.924)	4.634 (3.576 to 5.989)																					
Lower respiratory infections	Birth prevalence - [37, 38) wks, [2500,	Mortality	Males		3.306	3.194																					
Lower respiratory infections	S000) g Birth prevalence - [37, 38) wks, [2500,	Manulity	Females		(2.8210 3.843) 2.991	3.242																					
I information	3000) g Birth prevalence - [40, 42) wks, [2500,	Matelia	Malar		(2.496 to 3.61) 3.771	(2.745 to 3.817) 3.175																					
Lower respectively includes	3000) g Birth menalence - (40, 47) wise (2500	Juntany	ALL A		(3.002 to 4.693) 3.244	(2.56 to 3.923) 3.228																					
Lower respiratory infections	3000) g	Mortality	Females		(2.486 to 4.159)	(2.605 to 3.985)																					
Lower respiratory infections	3000) g	Mortality	Males		(2.274 to 3.309)	(2.44 to 3.548)																					
Lower respiratory infections	Barth prevalence - [38, 40) wks, [2500, 3000) g	Mortality	Females		2.376 (1.91 to 2.886)	2.938 (2.434 to 3.503)																					
Lower respiratory infections	Birth prevalence - [36, 37) wks, [3000, 3500) g	Mortality	Males		3.774 (3.094 to 4.497)	2.466 (2.058 to 2.929)																					
Lower respiratory infections	Birth prevalence - [36, 37) wks, [3000, 3500) g	Mortality	Females		3.73 (2.981 to 4.646)	2.715 (2.277 to 3.218)																					
Lower respiratory infections	Birth prevalence - [36, 37) wks, [4000, 4500) g	Mortality	Males		6.826 (5.212 to 9.045)	1.77 (1.491 to 2.082)																					
Lower respiratory infections	Birth prevalence - [36, 37) wks, [4000, 45000 m	Mortality	Females		7.269 (5.144 to 9.821)	2.177 (1.786 to 2.607)																					
Lower respiratory infections	Birth prevalence - [36, 37) wks, [3500,	Mortality	Males		4.544	2.057																					
Lower respiratory infections	Birth prevalence - [36, 37) wks, [3500,	Monality	Females		4.662	2.398																					
Lower restinatory infections	4000) g Birth prevalence - [37, 38) wks, [3000,	Manulity	Malex		2.007	1.888																					
	3500) g Birth prevalence - [37, 38) wks, [3000,				(1.759 to 2.293) 1.925	(1.613 to 2.224) 1.972																					
Lower respiratory intections	3500) g Birth menalence - [37-38) wise [4000	stortaity	Penales		(1.582 to 2.328) 3.28	(1.669 to 2.313)																					
Lower respiratory infections	4500) g Risk smallerer (27, 38) adv (4000	Mortality	Males		(2.596 to 4.133)	(1.171 to 1.532)																					
Lower respiratory infections	4500) g	Mortality	Females		(2.649 to 4.5)	(1.333 to 1.835)																					
Lower respiratory infections	Barth prevalence - [37, 38) wks, [3500, 4000) g	Mortality	Males		2.128 (1.833 to 2.466)	(1.299 to 1.76)																					
Lower respiratory infections	Birth prevalence - [37, 38) wks, [3500, 4000) g	Mortality	Females		2.142 (1.694 to 2.67)	1.661 (1.411 to 1.961)																					
Lower respiratory infections	Birth prevalence - [40, 42) wks, [3000, 3500) g	Mortality	Males		1.436 (1.245 to 1.65)	1.47 (1.199 to 1.8)																					
Lower respiratory infections	Birth prevalence - [40, 42) wks, [3000, 3500) g	Mortality	Females		1.326 (1.069 to 1.614)	1.465 (1.188 to 1.775)																					
Lower respiratory infections	Birth prevalence - [38, 40) wks, [3000, 3500) g	Mortality	Males		1.33 (1.155 to 1.53)	1.559 (1.305 to 1.851)																					
Lower respiratory infections	Birth prevalence - [38, 40) wks, [3000, 3500) g	Mortality	Females		1.224 (1.0 to 1.492)	1.564 (1.304 to 1.847)																					
Lower respiratory infections	- Birth prevalence - [38, 40) wks, [4000, 4500) r	Mortality	Males		1.787 (1.453 to 2.182)	1.175 (1.005 to 1.371)																					
Lower respiratory infections	Birth prevalence - [38, 40) wks, [4000,	Mortality	Females		1.877	1.224																					
Lower restinatory infections	4500) g Birth prevalence - [38, 40) wks, [3500,	Manulity	Malex		(1.46/10.2.388)	1.173																					
	4000) g Birth prevalence - [38, 40) wks, [3500,				(1.478 to 2.147) 1.892	(1.0 to 1.377) 1.23																					
Lower respiratory intections	4000) g Birth menalence - [40, 471 wise 13490	Mortainy	Centares		(1.481 to 2.352) 1.0	(1.03 to 1.46) 1.003																					
Lower respiratory infections	4000) g	Mortality	Males		(1.0 to 1.0)	(1.0 to 1.046)																					
Lower respiratory infections	4000) g	Mortality	Females		(1.0 to 1.013)	(1.0 to 1.006)																					
Lower respiratory infections	Bath prevalence - [40, 42) wks, [4000, 4500) g	Mortality	Males		1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																					
Lower respiratory infections	Birth prevalence - [40, 42) wks, [4000, 4500) g	Mortality	Females		1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																					
Lower respiratory infections	Birth prevalence - [28, 30) wks, [2000, 2500) g	Mortality	Males		117.172 (83.895 to 158.056)	27.726 (21.877 to 34.972)																					
Lower respiratory infections	Birth prevalence - [28, 30) wks, [2000, 2500) g	Mortality	Females		121.682 (84.349 to 171.375)	33.983 (26.101 to 43.404)																					
Lower respiratory infections	Birth prevalence - [28, 30) wks, [2500, 3000) g	Mortality	Males		77.948 (54.687 to 105.047)	16.608 (12.653 to 21.188)																					
Lower respiratory infections	Birth prevalence - [28, 30] wks, [2500, 3000) p	Mortality	Females		79.193 (54.099 to 112.234)	20.387 (15.089 to 26.434)																					
Lower respiratory infections	Birth prevalence - [28, 30) wks, [3000,	Mortality	Males		42.199	10.082																					
Lower respiratory infectives	Birth prevalence - [28, 30) wks, [3000,	Mortality	Females		(29.891 to 57.227) 42.551	(7.777 to 13.056) 11.989																					
	3000) g		1		(28.25 to 63.209)	(9.084 to 15.544)	I																				

opendix Table 6a. Relative risk	s used by age and sex for each outcor	ne for all risk factors ex	ept for ambient a	r pollution alcohol,	and smosting.									А	Ages											
Risk - Outcome	Category / Units	Morbidity / Mortality	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Lower respiratory infections	Birth prevalence - [30, 32) wks, [2500, 2000) -	Mortality N	ales	58.722	12.115													I								
Lower respiratory infections	Birth prevalence - [30, 32) wks, [2500,	Mortality Fe	nales	59.522	15.364																					
I anno anniatan infastiana	3000) g Birth prevalence - [30, 32) wks, [3000,	Mastelite A		(42.058 to 82.793) 45.67	(11.936 to 19.581) 8.381																					
iowi napizwa mitina	3500) g Birth menulence - (30, 32) wks, (3000,	sectory a		(32.014 to 65.531) 46.104	(6.362 to 10.842) 10.506																					
Lower respiratory infections	3500) g	Mortality Fe	nalex	(30.243 to 66.207)	(8.041 to 13.513)																					
Lower respiratory infections	Both prevalence - [30, 32) wks, [3500, 4000) g	Mortality N	iales	36.334 (21.558 to 54.813)	5.698 (4.293 to 7.349)																					
Lower respiratory infections	Birth prevalence - [30, 32) wks, [3500, 4000) g	Mortality Fe	nales	37.931 (22.692 to 61.276)	6.892 (5.175 to 9.058)																					
Lower respiratory infections	Birth prevalence - [32, 34) wks, [3000, 3500) g	Mortality N	iales	34.016 (23.515 to 48.37)	6.577 (4.934 to 8.436)																					
Lower respiratory infections	Birth prevalence - [32, 34) wks, [3000, 3500) g	Mortality Fe	nales	34.585 (22.909 to 49.754)	8.314 (6.362 to 10.789)																					
Lower respiratory infections	Birth prevalence - [32, 34) wks, [3500,	Mortality M	iales	36.248	5.068																					
Lower respiratory infections	Birth prevalence - [32, 34) wks, [3500,	Mortality Fe	nales	38.098	6.476																					
	4000) g Birth prevalence - [36, 37) wks, [1000,			(23.301 to 59.429) 166.686	(4.666 to 8.689) 57.535																					
Lower respiratory intections	1500) g Birth menalence - [36-37) wks [1000	Montality A	105	(118.487 to 222.581 169.725	(45.999 to 71.742) 63.564																					
Lower respiratory infections	1500) g	Mortality Fe	nalex	(119.017 to 229.008	(50.068 to 80.703)																					
Lower respiratory infections	Bath prevalence - [38, 40) wks, [1000, 1500) g	Mortality N	iales	174.066 (125.125 to 232.507	57.966 (44.393 to 73.241)																					
Lower respiratory infections	Birth prevalence - [38, 40) wks, [1000, 1500) g	Mortality Fe	nales	171.557 (121.585 to 237.047	65.208 (48.821 to 84.308)																					
Lower respiratory infections	Birth prevalence - [38, 40) wks, [1500, 2000) g	Mortality M	iales	67.302 (49.547 to 89.055)	25.206 (20.365 to 31.168)																					
Lower respiratory infections	Birth prevalence - [38, 40) wks, [1500, 2000) #	Mortality Fe	nales	62.19 (45 884 to 83 445)	28.05 (22.625 in 35.139)																					
Lower respiratory infections	Birth prevalence - [40, 42) wks, [1500,	Mortality N	iales	76.673	25.785																					
Lower restinators infections	Birth prevalence - [40, 42) wks, [1500,	Marolity Fr	makes	70.411	29.113																					
iowi napizwa mitina	2000) g	socially re		(49.221 to 97.952) 1564.792	(21.355 to 38.272) 618.595																					
Upper respiratory infections	Barth prevalence - [0, 24) wks, [0, 500) g	Mortality N	ales	(1056.542 to 2116.062) 1600.122	(458.842 to 812.921)																					
Upper respiratory infections	Birth prevalence - [0, 24) wkx, [0, 500) g	Mortality Fe	nales	(1050.664 to 2011 977)	(526.178 to 921.018)																					
Upper respiratory infections	Birth prevalence - [0, 24) wks, [500, 1000) 8	Mortality N	ales	1155.815 (825.412 to 1506.83)	457.5 (352.552 to 573.483)																					
Upper respiratory infections	Birth prevalence - [0, 24) wkx, [500, 1000) g	Mortality Fe	nales	1169.123 (802.003 to 1617.97)	515.406 (396.713 to 641.541)																					
Upper respiratory infections	Birth prevalence - [24, 26) wks, [500, 1000) 8	Mortality N	ales	955.583 (723.748 to 1244.265	443.357 (363.03 to 534.695)																					
Upper respiratory infections	Birth prevalence - [24, 26) wks, [500, 1000)	Mortality Fe	nales	947.143 (702.462 to 1227.00)	487.549																					
Upper respiratory infections	8 Birth prevalence - [26, 28) wks, [500, 1000)	Mortality N	ales	497.817	330.886																					
Unite material and infections	8 Birth prevalence - [26, 28) wks, [500, 1000)	Marolity Fr	makes	483.682	344.618																					
	8 Birth meyalence - (30, 32) wks, (500, 1000)			(354.946 to 629.517 236.614	(274.427 to 419.864) 149.995																					
Upper respiratory intections	8	stortaity N	105	(163.821 to 324.502	(117.866 to 188.368)																					
Upper respiratory infections	8	Mortality Fe	nales	(157.606 to 317.194	(120.779 to 190.583)																					
Upper respiratory infections	Birth prevalence - [28, 30) wks, [500, 1000) 8	Mortality N	ales	297.629 (214.953 to 396.586	216.995 (173.321 to 271.466)																					
Upper respiratory infections	Birth prevalence - [28, 30) wks, [500, 1000) 8	Mortality Fe	nales	281.056 (198.176 to 386.635	219.884 (174.264 to 272.704)																					
Upper respiratory infections	Birth prevalence - [26, 28) wks, [1000, 1500) g	Mortality N	ales	267.91 (210.177 to 332.92)	164.167 (132.898 to 200.569)																					
Upper respiratory infections	Birth prevalence - [26, 28) wks, [1000, 1500) r	Mortality Fe	nales	266.509 (197.461 to 346.932	174.222 (137.431 to 217.349)																					
Upper respiratory infections	Birth prevalence - [34, 36) wks, [1000, 1500) -	Mortality N	iales	142.056	52.86 (42.014 in 61.617)																					
Upper respiratory infections	Birth prevalence - [34, 36) wks, [1000,	Mortality Fe	nales	141.899	57.421																					
	1500) g Birth prevalence - [28, 30) wks, [1500,			(95.864 to 197.656) 127.966	(46.452 to 71.339) 50.018																					
Upper respiratory intections	2000) g Birth menalence - [28, 30) wks. [1500	Montality A	105	(97.178 to 167.026) 130.924	(40.539 to 61.919) 57.275																					
Upper respiratory infections	2000) g	Mortality Fe	nales	(96.513 to 172.188)	(46.36 to 70.038)																					
Upper respiratory infections	1500) g	Mortality N	ales	(120.99 to 204.947)	(83.486 to 127.144)																					
Upper respiratory infections	Birth prevalence - [28, 30) wks, [1000, 1500) g	Mortality Fe	nales	153.905 (112.327 to 200.786	107.529 (86.954 to 131.78)																					
Upper respiratory infections	Birth prevalence - [32, 34) wks, [1000, 1500) g	Mortality N	ales	117.142 (81.354 to 161.101)	53.185 (43.049 to 66.274)																					
Upper respiratory infections	Birth prevalence - [32, 34) wks, [1000, 1500) g	Mortality Fe	nales	115.171 (79.363 to 159.206)	56.034 (45.982 to 68.36)																					
Upper respiratory infections	Birth prevalence - [30, 32) wks, [1000, 1500) r	Mortality M	iales	119.308 (87.769 to 160 885)	67.163 (54.863 to 82.638)																					
Upper respiratory infections	Bith prevalence - [30, 32) wks, [1000, 1500) a	Mortality Fe	nales	115.448	(9).14 (55.873 95.013-																					
Upper respiratory infections	Birth prevalence - [37, 38) wks, [1500, 20000 -	Mortality N	ales	62.972	24.148																					
Upper prepiratory infection-	Bith prevalence - [37, 38] wks, [1500,	Mortality E.	nales	(+0.159 to 83.484) 59.988	(20,000 to 29,406) 26.719																					
	2000) g Birth prevalence - [36. 37) wks. [1500	Mar Pe		(43.974 to 79.053) 60.218	(21.746 to 32.816) 23.031																					
Upper respiratory infections	2000) g	Mortality N	ales	(43.669 to 82.48)	(18.793 to 28.483)																					
Upper respiratory infections	Bath prevalence - [36, 37) wks, [1500, 2000) g	Mortality Fe	nales	58.527 (42.172 to 80.557)	25.143 (20.331 to 30.566)																					
Upper respiratory infections	Birth prevalence - [30, 32) wks, [2000, 2500) g	Mortality N	iales	67.971 (50.354 to 88.935)	18.03 (14.621 to 22.103)																					
Upper respiratory infections	Birth prevalence - [30, 32) wks, [2000, 2500) g	Mortality Fe	nales	69.383 (49.108 to 94.583)	22.069 (17.836 to 27.163)																					
Upper respiratory infections	Birth prevalence - [30, 32) wks, [1500, 2000) g	Mortality N	ales	77.369 (59.702 to 99.232)	31.079 (25.786 to 36.724)																					
Upper respiratory infections	- Bitth prevalence - [30, 32) wks, [1500, 2000) g	Mortality Fe	nales	76.134 (56.885 to 100 page)	34.756 (28.764 in 41 8.440																					
Upper respiratory infections	Birth prevalence - [34, 36) wks, [1500,	Mortality N	ales	55.555	21.346																					
Linear maximum infenti	2000) g Birth prevalence - [34, 36) wks, [1500,	Mandin 1	makes	(39.553 to 75.104) 54.335	(17.677 to 26.143) 23.046																					
oppa neprawy mechani	2000) g Birth menulence - (32, 34) wks, (1 500	sectarity PE		(38.617 to 75.24) 57.155	(18.743 to 28.287) 23.114																					
Upper respiratory infections	2000) g Bith membros - [37 24) when [1600]	Mortality N	ales	(42.484 to 73.651)	(19.028 to 27.915) 25.140																					
Upper respiratory infections	2000) g	Mortality Fe	nales	56.101 (39.794 to 76.295)	25.149 (20.615 to 30.388)																					
Upper respiratory infections	Birth prevalence - [32, 34) wks, [2000, 2500) g	Mortality N	iales	37.444 (29.026 to 48.227)	12.233 (10.252 to 14.477)																					
Upper respiratory infections	Birth prevalence - [32, 34) wks, [2000, 2500) g	Mortality Fe	nales	36.874 (26.658 to 49.653)	14.384 (12.095 to 17.03)																					
Upper respiratory infections	Birth prevalence - [40, 42) wks, [2000, 2500) g	Mortality N	ales	18.092 (13.292 to 23.719)	9.23 (7.037 to 11.454)																					
			•		•	•																				

				, and the particular p		ind an and a line.									A	ges											
Risk - Outcome	Category / Units	Morbidity / Mortalit	v Sex	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Upper respiratory infections	Birth prevalence - [40, 42) wks, [2000, 26000 -	Mortality	Females		15.574	9.975																					
Upper respiratory infections	Birth prevalence - [38, 40) wks, [2000,	Mortality	Males		13.104	8.198																					
line mainten infection	2500) g Birth prevalence - [38, 40) wks, [2000,	Manulity	Females		(9.829 to 16.99) 11.308	(6.786 to 9.959) 8.577																					
	2500) g Birth prevalence - [32, 34) wks, [2500,				(8.389 to 14.38) 33.063	(7.04 to 10.449) 8.441																					
Upper respiratory intections	3000) g Birth menalence - [32-34] wise [2500	stortany	NIBO		(24.393 to 43.503) 32.812	(6.822 to 10.431) 10.398																					
Upper respiratory infections	3000) g	Mortality	Females		(23.439 to 45.567)	(8.227 to 13.042)																					
Upper respiratory infections	2500) g	Mortality	Males		(16.305 to 29.433)	(7.859 to 11.112)																					
Upper respiratory infections	Birth prevalence - [34, 36) wks, [2000, 2500) g	Mortality	Females		21.297 (15.657 to 28.761)	10.295 (8.548 to 12.273)																					
Upper respiratory infections	Birth prevalence - [37, 38) wks, [2000, 2500) g	Mortality	Males		13.0 (10.102 to 16.456)	8.096 (6.724 to 9.676)																					
Upper respiratory infections	Birth prevalence - [37, 38) wks, [2000, 2500) g	Mortality	Females		11.563 (8.805 to 15.11)	8.467 (6.994 to 10.342)																					
Upper respiratory infections	Birth prevalence - [36, 37) wks, [2000, 2500) g	Mortality	Males		14.401 (10.789 to 18.654)	8.221 (6.917 to 9.923)																					
Upper respiratory infections	Birth prevalence - [36, 37) wks, [2000, 2500) #	Mortality	Females		13.513 (9.817 to 17.947)	8.654 (7.215 to 10.369)																					
Upper respiratory infections	Birth prevalence - [34, 36) wks, [2500, 2000) -	Mortality	Males		13.419	5.562																					
Unner respiratory infections	Birth prevalence - [34, 36) wks, [2500,	Monality	Females		13.266	6.395																					
Uniter respiratory infections	3000) g Birth prevalence - [34, 36) wks, [4000,	Manulity	Malex		(9.666 to 17.689) 23.096	(5.292 to 7.606) 2.895																					
cipa najnavi incluar	4500) g Birth meyalence - [34, 36) wks, [4000,				(14.708 to 35.098) 25.038	(2.245 to 3.716) 3.778																					
Upper respiratory infections	4500) g Rightmansharar (24, 36) mbs (2000	Mortality	Females		(15.763 to 37.255)	(2.855 to 4.925)																					
Upper respiratory infections	3500) g	Mortality	Males		(10.222 to 18.478)	(3.449 to 5.338)																					
Upper respiratory infections	Barth prevalence - [34, 36) wks, [3000, 3500) g	Mortality	Females		14.375 (10.269 to 20.114)	5.265 (4.145 to 6.564)																					
Upper respiratory infections	Birth prevalence - [36, 37) wks, [2500, 3000) g	Mortality	Males		4.874 (4.014 to 5.713)	3.699 (3.134 to 4.374)																					
Upper respiratory infections	Birth prevalence - [36, 37) wks, [2500, 3000) g	Mortality	Females		4.609 (3.731 to 5.61)	3.898 (3.235 to 4.67)																					
Upper respiratory infections	Birth prevalence - [34, 36) wks, [3500, 4000) g	Mortality	Males		18.024 (12.279 to 25.547)	3.657 (2.838 to 4.675)																					
Upper respiratory infections	Birth prevalence - [34, 36) wks, [3500,	Mortality	Females		19.263	4.634																					
Upper respiratory infections	Birth prevalence - [37, 38) wks, [2500,	Mortality	Males		3.306	3.194																					
University infections	Su00) g Birth prevalence - [37, 38) wks, [2500,	Manulity	Females		(2.8210 3.843) 2.991	3.242																					
United and interview.	3000) g Birth prevalence - [40, 42) wks, [2500,	Matelia	Mala		(2.496 to 3.61) 3.771	(2.745 to 3.817) 3.175																					
cipa napizory method	3000) g Birth menalence - [40, 47) wise [2500	Juntany	ALLA		(3.002 to 4.693) 3.244	(2.56 to 3.923) 3.228																					
Upper respiratory infections	3000) g	Mortality	Females		(2.486 to 4.159)	(2.605 to 3.985)																					
Upper respiratory infections	3000) g	Mortality	Males		(2.274 to 3.309)	(2.44 to 3.548)																					
Upper respiratory infections	Barth prevalence - [38, 40) wks, [2500, 3000) g	Mortality	Females		2.376 (1.91 to 2.886)	2.938 (2.434 to 3.503)																					
Upper respiratory infections	Birth prevalence - [36, 37) wks, [3000, 3500) g	Mortality	Males		3.774 (3.094 to 4.497)	2.466 (2.058 to 2.929)																					
Upper respiratory infections	Birth prevalence - [36, 37) wks, [3000, 3500) g	Mortality	Females		3.73 (2.981 to 4.646)	2.715 (2.277 to 3.218)																					
Upper respiratory infections	Birth prevalence - [36, 37) wks, [4000, 4500) g	Mortality	Males		6.826 (5.212 to 9.045)	1.77 (1.491 to 2.082)																					
Upper respiratory infections	Birth prevalence - [36, 37) wks, [4000, 4500) g	Mortality	Females		7.269 (5.144 to 9.821)	2.177 (1.786 to 2.607)																					
Upper respiratory infections	Birth prevalence - [36, 37) wks, [3500, 40000 -	Mortality	Males		4.544	2.057																					
Upper respiratory infections	Birth prevalence - [36, 37) wks, [3500,	Mortality	Females		4.662	2.398																					
Unite restitutory infections	4000) g Birth prevalence - [37, 38) wks, [3000,	Manulity	Malex		2.007	1.888																					
	3500) g Birth prevalence - [37, 38) wks, [3000,				(1.759 to 2.293) 1.925	(1.613 to 2.224) 1.972																					
Opper respiratory intections	3500) g Birth menulence - (37-38) wise (4000	stortaity	remates		(1.582 to 2.328) 3.28	(1.669 to 2.313)																					
Upper respiratory infections	4500) g Rightmannham (27, 28) who (4000)	Mortality	Males		(2.596 to 4.133)	(1.171 to 1.532)																					
Upper respiratory infections	4500) g	Mortality	Females		(2.649 to 4.5)	(1.333 to 1.835)																					
Upper respiratory infections	Barth prevalence - [37, 38) wks, [3500, 4000) g	Mortality	Males		2.128 (1.833 to 2.466)	(1.299 to 1.76)																					
Upper respiratory infections	Birth prevalence - [37, 38) wks, [3500, 4000) g	Mortality	Females		2.142 (1.694 to 2.67)	1.661 (1.411 to 1.961)																					
Upper respiratory infections	Birth prevalence - [40, 42) wks, [3000, 3500) g	Mortality	Males		1.436 (1.245 to 1.65)	1.47 (1.199 to 1.8)																					
Upper respiratory infections	Birth prevalence - [40, 42) wks, [3000, 3500) g	Mortality	Females		1.326 (1.069 to 1.614)	1.465 (1.188 to 1.775)																					
Upper respiratory infections	Birth prevalence - [38, 40) wks, [3000, 3500) g	Mortality	Males		1.33 (1.155 to 1.53)	1.559 (1.305 to 1.851)																					
Upper respiratory infections	Birth prevalence - [38, 40) wks, [3000, 3500) g	Mortality	Females		1.224 (1.0 to 1.492)	1.564 (1.304 to 1.847)																					
Upper respiratory infections	- Birth prevalence - [38, 40) wks, [4000, 45000 m	Mortality	Males		1.787	1.175																					
Upper respiratory infections	Birth prevalence - [38, 40) wks, [4000,	Mortality	Females		1.877	1.224																					
Unite restitutory infections	4500) g Birth prevalence - [38, 40) wks, [3500,	Manulity	Malex		(1.46/10.2.388)	1.173																					
	4000) g Birth prevalence - [38, 40) wks, [3500,				(1.478 to 2.147) 1.892	(1.0 to 1.377) 1.23																					
upper respiratory intections	4000) g Birth menalence - [40-47) why 13600	Mortainy	remates		(1.481 to 2.352) 1.0	(1.03 to 1.46) 1.003																					
Upper respiratory infections	4000) g Rightmandara (40, 42) who (2000)	Mortality	Males		(1.0 to 1.0)	(1.0 to 1.046)																					
Upper respiratory infections	4000) g	Mortality	Females		(1.0 to 1.013)	(1.0 to 1.006)																					
Upper respiratory infections	Bath prevalence - [40, 42) wks, [4000, 4500) g	Mortality	Males		1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																					
Upper respiratory infections	Birth prevalence - [40, 42) wks, [4000, 4500) g	Mortality	Females		1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																					
Upper respiratory infections	Birth prevalence - [28, 30) wks, [2000, 2500) g	Mortality	Males		117.172 (83.895 to 158.056)	27.726 (21.877 to 34.972)																					
Upper respiratory infections	Birth prevalence - [28, 30) wks, [2000, 2500) g	Mortality	Females		121.682 (84.349 to 171.375)	33.983 (26.101 to 43.404)																					
Upper respiratory infections	Birth prevalence - [28, 30) wks, [2500, 3000) g	Mortality	Males		77.948 (54.687 to 105.047)	16.608 (12.653 to 21.188)																					
Upper respiratory infections	Birth prevalence - [28, 30) wks, [2500, 3000) g	Mortality	Females		79.193 (54.099 to 112.234)	20.387 (15.089 to 26.434)																					
Upper respiratory infections	Birth prevalence - [28, 30) wks, [3000,	Mortality	Males		42.199	10.082																					
Upper respiratory infectives	Birth prevalence - [28, 30) wks, [3000,	Mortality	Females		(29.891 to 57.227) 42.551	(7.777 to 13.056) 11.989																					
	3900) g		1		(28.25 to 63.209)	(9.084 to 15.544)	I																				

opendix Table 6a. Kelative risk	s used by age and sex for each outcon	ne for all risk factors (except for ambient a	r pollution alcohol,	and smoking.								Ages											
Risk - Outcome	Category / Units	Morbidity / Mortality	Sex All-age	0-6 days	7-27 days	28-364 days 1-4 years	5-9 years	10-14 years	15-19 years	20-24 years 25-	19 years 30-3	4 years 35-39 yea	ars 40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Upper respiratory infections	Birth prevalence - [30, 32) wks, [2500, 3000) e	Mortality	Males	58.722 (42.419 to 78.873)	12.115 (9.518 to 15.521)		1 1								I	I								
Upper respiratory infections	Bith prevalence - [30, 32) wks, [2500,	Mortality	Females	59.522	15.364																			
Unner respiratory infections	Birth prevalence - [30, 32) wks, [3000,	Mortality	Males	(42.058.05.82.793) 45.67	8.381																			
Name empiristent infections	3500) g Birth prevalence - [30, 32) wks, [3000,	Mastelite	Employ	(32.014 to 65.531) 46.104	(6.362 to 10.842) 10.506																			
Copie inspiratory interiories	3500) g Birth merudence - (30, 32) wks, (3500,			(30.243 to 66.207) 36.334	(8.041 to 13.513) 5.698																			
Upper respiratory infections	4000) g Rish semalaran (20, 22) ada (2600	Mortality	Males	(21.558 to 54.813) 27.021	(4.293 to 7.349)																			
Upper respiratory infections	4000) g	Mortality	Females	(22.692 to 61.276)	(5.175 to 9.058)																			
Upper respiratory infections	Both prevalence - [32, 34) wks, [3000, 3500) g	Mortality	Males	34.016 (23.515 to 48.37)	6.577 (4.934 to 8.436)																			
Upper respiratory infections	Birth prevalence - [32, 34) wks, [3000, 3500) g	Mortality	Females	34.585 (22.909 to 49.754)	8.314 (6.362 to 10.789)																			
Upper respiratory infections	Birth prevalence - [32, 34) wks, [3500, 4000) g	Mortality	Males	36.248 (23.158 to 54.67)	5.068 (3.761 to 6.741)																			
Upper respiratory infections	Birth prevalence - [32, 34) wks, [3500, 4000) g	Mortality	Females	38.098 (23.301 to 59.429)	6.476 (4.666 to 8.689)																			
Upper respiratory infections	Birth prevalence - [36, 37) wks, [1000, 1500) g	Mortality	Males	166.686 (118.487 to 222.581)	57.535 (45.999 to 71.742)																			
Upper respiratory infections	Birth prevalence - [36, 37) wks, [1000, 1500) g	Mortality	Females	169.725 (119.017 to 229.008)	63.564 (50.068 to 80.703)																			
Upper respiratory infections	Birth prevalence - [38, 40) wks, [1000, 1500) a	Mortality	Males	174.066 (125.125 to 232.507)	57.966 (44.393 to 73.241)																			
Upper respiratory infections	Birth prevalence - [38, 40) wks, [1000,	Mortality	Females	171.557	65.208																			
Unner respiratory infections	Birth prevalence - [38, 40) wks, [1500,	Mortality	Males	67.302	(48.82110 84.308) 25.206																			
Unter respiratory infections	2000) g Birth prevalence - [38, 40) wks, [1500,	Manulin	Females	(49.547 to 89.055) 62.19	(20.365 to 31.168) 28.05																			
	2000) g Birth prevalence - [40, 42] wks, [1500,			(45.884 to 83.445) 76.673	(22.625 to 35.139) 25.785																			
Opper respiratory intections	2000) g Birth remains - [40, 47) wis: [1500	stortany	MING	(56.177 to 102.468) 70.411	(19.387 to 34.168) 29.113																			
Upper respiratory infections	2000) g	Mortality	Females	(49.221 to 97.952) 1564.792	(21.355 to 38.272)																			
Otitis media	Birth prevalence - [0, 24) wks, [0, 500) g	Mortality	Males	(1056.542 to 2116.062) 1600.122	(458.842 to 812.921																			
Otitis media	Birth prevalence - [0, 24) wks, [0, 500) g	Mortality	Females	(1050.664 to 2111.977)	713.571 (526.178 to 921.018	>																		
Otitis media	Birth prevalence - [0, 24) wks, [500, 1000) 8	Mortality	Males	1155.815 (825.412 to 1506.837	457.5 (352.552 to 573.483																			
Otitis media	Birth prevalence - [0, 24) wks, [500, 1000) 8	Mortality	Females	1169.123 (802.003 to 1617.979	515.406 (396.713 to 641.541)																		
Otitis media	Birth prevalence - [24, 26) wks, [500, 1000) g	Mortality	Males	955.583 (723.748 to 1244.265	443.357 (363.03 to 534.695)																			
Otitis media	Birth prevalence - [24, 26) wks, [500, 1000)	Mortality	Females	947.143 (702.662 to 1237.093	487.549 (387.307 to 603.498	>																		
Otitis media	Birth prevalence - [26, 28) wks, [500, 1000) 8	Mortality	Males	497.817 (377.617 to 648.547)	330.886 (261.438 to 401.709	, ,																		
Otitis media	Birth prevalence - [26, 28) wks, [500, 1000)	Mortality	Females	483.682 (354.946 to 629.517)	344.618 (274.427 to 419.864																			
Otitis media	o Birth prevalence - [30, 32) wks, [500, 1000)	Mortality	Males	236.614	149.995																			
Otitis media	8 Birth prevalence - [30, 32) wks, [500, 1000)	Mortality	Females	229.197	152.117																			
Otitis media	8 Birth prevalence - [28, 30) wks, [500, 1000)	Mortality	Males	297.629	216.995	2																		
Orizia madia	g Birth prevalence - [28, 30) wks, [500, 1000)	Manulin	Females	(214.953 to 396.586) 281.056	(173.321 to 271.466 219.884	9																		
Origina analas	8 Birth prevalence - [26, 28) wks, [1000,	Matelia	Maler	(198.176 to 386.635) 267.91	(174.264 to 272.704 164.167																			
Contra andrea	1500) g Birth prevalence - [26, 28) wks, [1000,	Mariany		(210.177 to 332.92) 266.509	(132.898 to 200.569 174.222	>																		
Onits media	1500) g Birth menulence - [34, 36) wise [1000]	stortany	remaies	(197.461 to 346.932) 142.056	(137.431 to 217.349 52.86																			
Otitis media	1500) g Rish muselman (24, 26) who (1000	Mortality	Males	(98.086 to 197.774)	(42.914 to 64.617)																			
Otitis media	1500) g	Mortality	Females	(95.864 to 197.656)	(46.452 to 71.339)																			
Otitis media	2000) g	Mortality	Males	(97.178 to 167.026)	(40.539 to 61.919)																			
Otitis media	Both prevalence - [28, 30) wks, [1500, 2000) g	Mortality	Females	(96.513 to 172.188)	57.275 (46.36 to 70.038)																			
Otitis media	Birth prevalence - [28, 30) wks, [1000, 1500) g	Mortality	Males	158.563 (120.99 to 204.947)	103.32 (83.486 to 127.144)																			
Otitis media	Birth prevalence - [28, 30) wks, [1000, 1500) g	Mortality	Females	153.905 (112.327 to 200.786)	107.529 (86.954 to 131.78)																			
Otitis media	Birth prevalence - [32, 34) wks, [1000, 1500) g	Mortality	Males	117.142 (81.354 to 161.101)	53.185 (43.049 to 66.274)																			
Otitis media	Birth prevalence - [32, 34) wks, [1000, 1500) g	Mortality	Females	115.171 (79.363 to 159.206)	56.034 (45.982 to 68.36)																			
Otitis media	Birth prevalence - [30, 32) wks, [1000, 1500) g	Mortality	Males	119.308 (87.769 to 160.885)	67.163 (54.863 to 82.638)																			
Otitis media	Birth prevalence - [30, 32) wks, [1000, 1500) g	Mortality	Females	115.448 (84.272 to 156.425)	69.14 (55.873 to 85.012)																			
Otitis media	Birth prevalence - [37, 38) wks, [1500, 2000) a	Mortality	Males	62.972 (46.159 to 83.484)	24.148 (20.066 to 29.406)																			
Otitis media	- Birth prevalence - [37, 38) wks, [1500, 2000) g	Mortality	Females	59.988 (43.974 to 79.052)	26.719 (21.746 in 32.814)																			
Otitis media	Birth prevalence - [36, 37) wks, [1500,	Mortality	Males	60.218	23.031																			
Otitis media	Birth prevalence - [36, 37) wks, [1500,	Mortality	Females	(43.889 83.48) 58.527	(18.1931028.483) 25.143																			
Orizin media	2000) g Birth prevalence - [30, 32) wks, [2000,	Manulin	Malex	(42.172 to 80.557) 67.971	(20.331 to 30.566) 18.03																			
Origina and a	2500) g Birth prevalence - (30, 32) wks, (2000,	Matelia	Emerica	(50.354 to 88.935) 69.383	(14.621 to 22.103) 22.069																			
Ories means	2500) g Birth prevalence - [30, 32) wks, [1500.	Mortany	e samaniti	(49.108 to 94.583) 77.369	(17.836 to 27.163) 31.079																			
Otitis media	2000) g Birth mercelence - [30, 37) why [1600	Mortality	MINES	(59.702 to 99.232) 76.134	(25.786 to 36.724) 34.756																			
Otitis media	2000) g Rich mandama (24, 26) ada (1700)	Mortality	Females	(56.885 to 100.996)	(28.764 to 41.849) 21.246																			
Otitis media	2000) g	Mortality	Males	(39.553 to 75.104)	(17.677 to 26.143)																			
Otitis media	1500, 2000) g	Mortality	Females	54.335 (38.617 to 75.24)	25.046 (18.743 to 28.287)																			
Otitis media	Birth prevalence - [32, 34) wks, [1500, 2000) g	Mortality	Males	57.155 (42.484 to 73.651)	23.114 (19.028 to 27.915)																			
Otitis media	Birth prevalence - [32, 34) wks, [1500, 2000) g	Mortality	Females	56.101 (39.794 to 76.295)	25.149 (20.615 to 30.388)																			
Otitis media	Birth prevalence - [32, 34) wks, [2000, 2500) g	Mortality	Males	37.444 (29.026 to 48.227)	12.233 (10.252 to 14.477)																			
Otitis media	Birth prevalence - [32, 34) wks, [2000, 2500) g	Mortality	Females	36.874 (26.658 to 49.653)	14.384 (12.095 to 17.03)																			
Otitis media	Birth prevalence - [40, 42) wks, [2000, 2500) g	Mortality	Males	18.092 (13.292 to 23.719)	9.23 (7.037 to 11.454)																			

															A	ges											
NI 0 .	6			All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Otitis media	Category / Units Birth prevalence - [40, 42) wks, [2000,	Morbidity / Morta	Females		15.574	9.975		I				I					1					1					
	2500) g Birth prevalence - [38, 40) wks, [2000,	M			(11.516 to 20.778) 13.104	(7.82 to 12.46) 8.198																					
Othis mean	2500) g	storcarty	MIRO		(9.829 to 16.99)	(6.786 to 9.959)																					
Otitis media	2500) g	Mortality	Females		(8.389 to 14.38)	(7.04 to 10.449)																					
Otitis media	Birth prevalence - [32, 34) wks, [2500, 3000) g	Mortality	Males		33.063 (24.393 to 43.503)	8.441 (6.822 to 10.431)																					
Otitis media	Birth prevalence - [32, 34) wks, [2500, 3000) g	Mortality	Females		32.812 (23.439 to 45.567)	10.398 (8.227 to 13.042)																					
Otitis media	Birth prevalence - [34, 36) wks, [2000, 25000 -	Mortality	Males		21.925 (16.305 to 20.422)	9.367																					
Otitis media	Bith prevalence - [34, 36) wks, [2000,	Mortality	Females		21.297	10.295																					
	2500) g Birth prevalence - [37, 38) wks, [2000,	M - P			(15.657 to 28.761) 13.0	(8.548 to 12.273) 8.096																					
Othis mean	2500) g Birth manufacture (27, 28) with (2000)	storcarty	MIRO		(10.102 to 16.456)	(6.724 to 9.676)																					
Otitis media	2500) g	Mortality	Females		(8.805 to 15.11)	(6.994 to 10.342)																					
Otitis media	Birth prevalence - [36, 37] wks, [2000, 2500) g	Mortality	Males		14.401 (10.789 to 18.654)	8.221 (6.917 to 9.923)																					
Otitis media	Birth prevalence - [36, 37) wks, [2000, 2500) g	Mortality	Females		13.513 (9.817 to 17.942)	8.654 (7.215 to 10.369)																					
Otitis media	Birth prevalence - [34, 36) wks, [2500, 3000) g	Mortality	Males		13.419 (10.387 to 16.819)	5.562 (4.646 to 6.696)																					
Otitis media	Birth prevalence - [34, 36) wks, [2500, 2000) -	Mortality	Females		13.266	6.395																					
Otitis media	Bith prevalence - [34, 36) wks, [4000,	Mortality	Males		23.096	2.895																					
	4500) g Birth prevalence - (34, 36) wks. (4000.				(14.708 to 35.098) 25.038	(2.245 to 3.716) 3.778																					
Othis mean	4500) g Birth menulation (24, 26) solar (2000	storcarty	Pemases		(15.763 to 37.255)	(2.855 to 4.925)																					
Otitis media	3500) g	Mortality	Males		(10.222 to 18.478)	(3.449 to 5.338)																					
Otitis media	Bath prevalence - [34, 36) wks, [3000, 3500) g	Mortality	Females		(10.269 to 20.114)	5.265 (4.145 to 6.564)																					
Otitis media	Birth prevalence - [36, 37) wks, [2500, 3000) g	Mortality	Males		4.874 (4.014 to 5.713)	3.699 (3.134 to 4.374)																					
Otitis media	Birth prevalence - [36, 37) wks, [2500, 3000) g	Mortality	Females		4.609 (3.731 to 5.61)	3.898 (3.235 to 4.67)																					
Otitis media	Birth prevalence - [34, 36) wks, [3500, 4000) r	Mortality	Males		18.024 (12.279 to 25.547)	3.657 (2.838 to 4.675)																					
Otitis media	Birth prevalence - [34, 36) wks, [3500,	Mortality	Females		19.263	4.634																					
Oticia analia	4000) g Birth prevalence - [37, 38) wks, [2500,	Mantalian	Malar		(12.567 to 27.924) 3.306	(3.576 to 5.989) 3.194																					
Control and and	3000) g Birth menulance - [37-38) wks [2500	Janaany	74454		(2.82 to 3.843) 2.991	(2.691 to 3.803) 3.242																					
Otitis media	3000) g	Mortality	Females		(2.496 to 3.61)	(2.745 to 3.817)																					
Otitis media	Bath prevalence - [40, 42] wks, [2500, 3000) g	Mortality	Males		(3.002 to 4.693)	3.175 (2.56 to 3.923)																					
Otitis media	Birth prevalence - [40, 42) wks, [2500, 3000) g	Mortality	Females		3.244 (2.486 to 4.159)	3.228 (2.605 to 3.985)																					
Otitis media	Birth prevalence - [38, 40) wks, [2500, 3000) g	Mortality	Males		2.755 (2.274 to 3.309)	2.944 (2.44 to 3.548)																					
Otitis media	Birth prevalence - [38, 40) wks, [2500, 3000) g	Mortality	Females		2.376 (1.91 to 2.886)	2.938 (2.434 to 3.503)																					
Otitis media	Birth prevalence - [36, 37) wks, [3000, 35000 r	Mortality	Males		3.774 (3.094 to 4.497)	2.466 (2.058 to 2.929)																					
Otitis media	Birth prevalence - [36, 37] wks, [3000,	Mortality	Females		3.73	2.715																					
Orizin modia	3300) g Birth prevalence - [36, 37) wks, [4000,	Montality	Malex		6.826	1.77																					
Origina analis	4500) g Birth prevalence - [36, 37) wks, [4000,	Mantalian	Emole		(5.212 to 9.045) 7.269	(1.491 to 2.082) 2.177																					
Only and a	4500) g Birth prevalence - [36, 37) wks, [3500,	Martin			(5.144 to 9.821) 4.544	(1.786 to 2.607) 2.057																					
Othis mean	4000) g Risk menskanse (36, 37) solo (3600	stortany	MIRO		(3.64 to 5.622)	(1.74 to 2.44)																					
Otitis media	4000) g	Mortality	Females		(3.577 to 6.014)	(1.98 to 2.864)																					
Otitis media	3500) g	Mortality	Males		(1.759 to 2.293)	(1.613 to 2.224)																					
Otitis media	Birth prevalence - [37, 38) wks, [3000, 3500) g	Mortality	Females		1.925 (1.582 to 2.328)	1.972 (1.669 to 2.313)																					
Otitis media	Birth prevalence - [37, 38) wks, [4000, 4500) g	Mortality	Males		3.28 (2.596 to 4.133)	1.335 (1.171 to 1.532)																					
Otitis media	Birth prevalence - [37, 38) wks, [4000, 4500) g	Mortality	Females		3.521 (2.649 to 4.5)	1.559 (1.333 to 1.835)																					
Otitis media	Birth prevalence - [37, 38) wks, [3500, 4000) g	Mortality	Males		2.128 (1.833 to 2.466)	1.505 (1.299 to 1.76)																					
Otitis media	Birth prevalence - [37, 38) wks, [3500, 4000) g	Mortality	Females		2.142 (1.694 to 2.67)	1.661 (1.411 to 1.961)																					
Otitis media	Birth prevalence - [40, 42) wks, [3000, 35000 r	Mortality	Males		1.436 (1.245 to 1.65)	1.47 (1.199 m 1.8)																					
Otitis media	Birth prevalence - [40, 42) wks, [3000,	Mortality	Females		1.326	1.465																					
Otitis media	Birth prevalence - [38, 40) wks, [3000,	Mortality	Males		(1.009 to 1.014)	(1.166 to 1.775) 1.559																					
Otitis molia	5500) g Birth prevalence - [38, 40) wks, [3000,	Matulity	Females		(1.155 to 1.53) 1.224	(4.305 to 1.851) 1.564																					
0000 - 5	3500) g Birth prevalence - [38, 40) wks, [4000.	y	. consett		(1.0 to 1.492) 1.787	(1.304 to 1.847) 1.175																					
Ornix media	4500) g Birth merculance, 128, 400 mile, 14000	stortatity	Males		(1.453 to 2.182)	(1.005 to 1.371)																					
Otitis media	4500) g	Mortality	Females		(1.467 to 2.388)	(1.022 to 1.465)																					
Otitis media	4000) g	Mortality	Males		(1.478 to 2.147)	(1.0 to 1.377)																					
Otitis media	Birth prevalence - [38, 40) wks, [3500, 4000) g	Mortality	Females		1.892 (1.481 to 2.352)	1.23 (1.03 to 1.46)																					
Otitis media	Birth prevalence - [40, 42) wks, [3500, 4000) g	Mortality	Males		1.0 (1.0 to 1.0)	1.003 (1.0 to 1.046)																					
Otitis media	Birth prevalence - [40, 42) wks, [3500, 4000) g	Mortality	Females		1.002 (1.0 to 1.013)	1.001 (1.0 to 1.006)																					
Otitis media	Birth prevalence - [40, 42) wks, [4000, 4500) g	Mortality	Males		1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																					
Otitis media	Birth prevalence - [40, 42] wks, [4000, 4500) a	Mortality	Females		1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																					
Otitis media	Birth prevalence - [28, 30] wks, [2000, 2500) #	Mortality	Males		117.172 (83.895 br 159.07 ···	27.726																					
Otitis media	Birth prevalence - [28, 30) wks, [2000,	Mortality	Females		121.682	33.983																					
Otitis modes	2300) g Birth prevalence - [28, 30) wks, [2500,	Manulin	Maler		(84.34910 171.375) 77.948	(28.101 to 43.404) 16.608																					
	3000) g Birth prevalence - [28, 30) wks. [2500.	y			(54.687 to 105.047) 79.193	(12.653 to 21.188) 20.387																					
Otitis media	3000) g Birth prevalence - 128-300 wise 13000	Mortabity	Females		(54.099 to 112.236) 42.199	(15.089 to 26.434) 10.082																					
Otitis media	3500) g Birth merculance, 128, 300 min 12000	Mortality	Males		(29.891 to 57.227)	(7.777 to 13.056)																					
Otitis media	3500) g	Mortality	Females		(28.25 to 63.209)	(9.084 to 15.544)	I																				

ppendix Table 6a. Kelative risi	ks used by age and sex for each outo	come for all risk factor:	s except t	or ambient air pollution	aconor	and Su2021112							Ages												
Risk - Outcome	Category / Units	Morbidity / Mortality	v Sex	All-age 0-6	days	7-27 days	28-364 days 1-4 years	5-9 years 10-14 years	urs 15-19 ye	ears 20-24 ye	ars 25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Otitis media	Birth prevalence - [30, 32) wks, [2500, 20000 -	Mortality	Males	55	(722	12.115																			
Otitis media	Bith prevalence - [30, 32) wks, [2500,	Mortality	Females	(42.419	(522	(9.5181015.521)																			
Origina and the	3000) g Birth prevalence - [30, 32) wks, [3000,	Manufite	Malar	(42.058	to 82.793) 5.67	(11.936 to 19.581) 8.381																			
China and a	3500) g Birth mevalence - (30, 32) wks. (3000,	Juniary	ALLA.	(32.014	to 65.531)	(6.362 to 10.842) 10.506																			
Otitis media	3500) g	Mortality	Females	(30.243	to 66.207)	(8.041 to 13.513)																			
Otitis media	Both prevalence - [30, 32) wks, [3500, 4000) g	Mortality	Males	(21.558	to 54.813)	5.698 (4.293 to 7.349)																			
Otitis media	Birth prevalence - [30, 32) wks, [3500, 4000) g	Mortality	Females	37 (22.692	.931 to 61.276)	6.892 (5.175 to 9.058)																			
Otitis media	Birth prevalence - [32, 34) wks, [3000, 3500) g	Mortality	Males	34 (23.515	.016 to 48.37)	6.577 (4.934 to 8.436)																			
Otitis media	Birth prevalence - [32, 34) wks, [3000, 3500) g	Mortality	Females	34 (22.909	.585 to 49.754)	8.314 (6.362 to 10.789)																			
Otitis media	Birth prevalence - [32, 34) wks, [3500,	Mortality	Males	36	248	5.068																			
Otitis media	Birth prevalence - [32, 34) wks, [3500,	Mortality	Females	38	.098	6.476																			
Origina mention	4000) g Birth prevalence - [36, 37) wks, [1000,	Manufitz	Malar	(23.301	10 59,429) 5.686	(4.666 to 8.689) 57.535																			
China and a	1500) g Birth mevalence - [36, 37) wks. [1000.	Juniary	ALLA.	(118.487	to 222.581) 0.725	(45.999 to 71.742) 63.564																			
Otitis media	1500) g	Mortality	Females	(119.017	to 229.008)	(50.068 to 80.703)																			
Otitis media	1500) g	Mortality	Males	(125.125	to 232.507)	(44.393 to 73.241)																			
Otitis media	Birth prevalence - [38, 40) wks, [1000, 1500) g	Mortality	Females	(121.585	1.557 to 237.047)	65.208 (48.821 to 84.308)																			
Otitis media	Birth prevalence - [38, 40) wks, [1500, 2000) g	Mortality	Males	67 (49.547	.302 to 89.055)	25.206 (20.365 to 31.168)																			
Otitis media	Birth prevalence - [38, 40) wks, [1500, 2000) #	Mortality	Females	60	2.19	28.05 (22.625 to 35.139)																			
Otitis media	Birth prevalence - [40, 42) wks, [1500,	Mortality	Males	76	.673	25.785																			
Otitis molia	Birth prevalence - [40, 42) wks, [1500,	Manufity	Females	(56.177	6 102.468) (411	29.113																			
No. 1	2000) g			(49.221 156	to 97.952) 4.792	(21.355 to 38.272) 618.595																			
Duarthoeal diseases	Both prevalence - [0, 24) wks, [0, 500) g	Mortality	Males	(105) 211- 160	6.542 to 6.0671 0.122	(458.842 to 812.921																			
Diarrhoeal diseases	Birth prevalence - [0, 24) wks, [0, 500) g	Mortality	Females	(105)	0.664 to	(526.178 to 921.018)																			
Diarrhoeal diseases	Birth prevalence - [0, 24) wks, [500, 1000) 8	Mortality	Males	115 (825.412 t	5.815 o 1506.837)	457.5 (352.552 to 573.483)																			
Diarrhoral diseases	Birth prevalence - [0, 24) wks, [500, 1000) g	Mortality	Females	(802.003 s	9.123 o 1617.979	515.406 (396.713 to 641.541																			
Diarrhoeal diseases	Birth prevalence - [24, 26) wks, [500, 1000 g	0 Mortality	Males	95 (723.748 t	5.583 o 1244.265)	443.357 (363.03 to 534.695)																			
Diarrhoeal diseases	Birth prevalence - [24, 26) wks, [500, 1000	0 Mortality	Females	94	7.143	487.549																			
Diarrhoeal diseases	o Birth prevalence - [26, 28) wks, [500, 1000	0 Mortality	Males	49	7.817	330.886																			
Distributed discours	8 Birth prevalence - [26, 28) wks, [500, 1000	0 Manufity	Females	(377.617	8.682	344.618																			
No. 1	8 Birth prevalence - [30, 32) wks, [500, 1000	0		(354.946	to 629.517) 5.614	(274.427 to 419.864 149.995																			
Diamoral diseases	8 Richaramatana (20.22) ada (600.1000	MORTHIN	MIDO	(163.821	to 324.502)	(117.866 to 188.368																			
Diarrhoeal diseases	8	Mortality	Females	(157.606	to 317.194)	(120.779 to 190.583)																			
Diarrhoeal diseases	Both prevalence - [28, 30) wks, [500, 1000 8	9 Mortality	Males	(214.953	(.629 to 396.586)	216.995 (173.321 to 271.466																			
Diarrhoeal diseases	Birth prevalence - [28, 30) wks, [500, 1000 8	0 Mortality	Females	28 (198.176	1.056 to 386.635)	219.884 (174.264 to 272.704																			
Diarrhoeal diseases	Birth prevalence - [26, 28) wks, [1000, 1500) g	Mortality	Males	26 (210.177	7.91 to 332.92)	164.167 (132.898 to 200.569																			
Diarrhoeal diseases	Birth prevalence - [26, 28) wks, [1000, 1500) #	Mortality	Females	26 (197.461	5.509 to 346.932)	174.222 (137.431 to 217.349																			
Diarrhoeal diseases	Birth prevalence - [34, 36) wks, [1000, 1500) -	Mortality	Males	14	2.056	52.86 (42.014 in 61.617)																			
Diamhoral diseases	Birth prevalence - [34, 36) wks, [1000,	Mortality	Females	14	1.899	57.421																			
Disastered Surgers	1500) g Birth prevalence - [28, 30) wks, [1500,	Manufite	Malar	(95.864)	o 197.636) 7.966	(46.452 to 71.539) 50.018																			
Diamonal Concess	2000) g Birth mevalence - (28, 30) wks. (1500.	Juniary	ALLA.	(97.178)	o 167.026)).924	(40.539 to 61.919) 57.275																			
Duarthoeal diseases	2000) g Rishmundama 128, 200 mln (1000	Mortality	Females	(96.513)	o 172.188)	(46.36 to 70.038)																			
Diarrhoeal diseases	1500) g	Mortality	Males	(120.99)	o 204.947)	(83.486 to 127.144)																			
Diarrhoeal diseases	Both prevalence - [28, 30) wks, [1000, 1500) g	Mortality	Females	(112.327	5.905 to 200.786)	107.529 (86.954 to 131.78)																			
Diarrhoeal diseases	Birth prevalence - [32, 34) wks, [1000, 1500) g	Mortality	Males	(81.354)	7.142 o 161.101)	53.185 (43.049 to 66.274)																			
Diarrhoeal diseases	Birth prevalence - [32, 34) wks, [1000, 1500) g	Mortality	Females	(79.363)	5.171 o 159.206)	56.034 (45.982 to 68.36)																			
Diarrhoeal diseases	Birth prevalence - [30, 32) wks, [1000, 1500) g	Mortality	Males	(87.769)	0.308 o 160.885)	67.163 (54.863 to 82.638)																			
Diarrhoeal diseases	Birth prevalence - [30, 32) wks, [1000, 1500) #	Mortality	Females	11:	5.448	69.14 (55.873 in 85.012)																			
Diarrhoeal diseases	Birth prevalence - [37, 38) wks, [1500, 2000) -	Mortality	Males	62	.972	24.148																			
Diambogal discours	2000) g Birth prevalence - [37, 38) wks, [1500,	Montality	Fermine	(46.159		(20.000 to 29.406) 26.719																			
The second	2000) g Birth prevalence - (36, 37) wks. (1500			(43.974	to 79.053) 218	(21.746 to 32.816) 23.031																			
Duarthoeal diseases	2000) g Riskersenheum (26, 27) sels (1800	Mortality	Males	(43.669	to 82.48)	(18.793 to 28.483) 26.142																			
Diarrhoeal diseases	2000) g	Mortality	Females	(42.172	to 80.557)	(20.331 to 30.566)																			
Diarrhoeal diseases	Both prevalence - [30, 32) wks, [2000, 2500) g	Mortality	Males	(50.354	.971 to 88.935)	18.03 (14.621 to 22.103)																			
Diarrhoeal diseases	Birth prevalence - [30, 32) wks, [2000, 2500) g	Mortality	Females	69 (49.108	.383 to 94.583)	22.069 (17.836 to 27.163)																			
Diarrhoeal diseases	Birth prevalence - [30, 32) wks, [1500, 2000) g	Mortality	Males	77 (59.702	.369 to 99.232)	31.079 (25.786 to 36.724)																			
Diarrhoeal diseases	Birth prevalence - [30, 32) wks, [1500, 2000) #	Mortality	Females	76	.134 o 100.9961	34.756 (28.764 to 41 849)																			
Diarrhoeal diseases	Birth prevalence - [34, 36) wks, [1500,	Mortality	Males	55	.555	21.346																			
Diambogal discours	2000) g Birth prevalence - [34, 36) wks, [1500,	Montality	Fermine	(39.553 54	10 75.104) .335	(17.677 to 26.143) 23.046																			
The second	2000) g Birth prevalence - [32, 34) wks. [1500			(38.617	to 75.24)	(18.743 to 28.287) 23.114																			
Diamboral diseases	2000) g Birth menalence - (32, 34) wky (1600	Mortality	Males	(42.484	to 73.651)	(19.028 to 27.915) 25.149																			
Diarrhoeal diseases	2000) g	Mortality	Females	(39.794	10 76.295)	(20.615 to 30.388)																			
Diarrhoeal diseases	Both prevalence - [32, 34) wks, [2000, 2500) g	Mortality	Males	37 (29.026	.444 to 48.227)	12.233 (10.252 to 14.477)																			
Diamboral diseases	Birth prevalence - [32, 34) wks, [2000, 2500) g	Mortality	Females	36 (26.658	.874 to 49.653)	14.384 (12.095 to 17.03)																			
Diarrhoeal diseases	Birth prevalence - [40, 42) wks, [2000, 2500) g	Mortality	Males	18 (13.292	:092 to 23.719)	9.23 (7.037 to 11.454)																			

															A	ges											
Rick - Outcome	Category / Units	Morbidity / Mortality	Ser	All-age 0-6	6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Diamboral diseases	Birth prevalence - [40, 42) wks, [2000,	Mortality I	Females	19	5.574	9.975												1									
Diambogal diseases	2500) g Birth prevalence - [38, 40) wks, [2000,	Mortality	Males	(11.516	3.104	8.198																					
No. 1	2500) g Birth prevalence - [38, 40) wks, [2000,	M . P.		(9.829	9 to 16.99) 1.308	(6.786 to 9.959) 8.577																					
Diamonal diseases	2500) g Birth prevalence - (32, 34) wks, (2500,			(8.389	9 to 14.38) (3.063	(7.04 to 10.449) 8.441																					
Duarthoeal diseases	3000) g Risk-susselses (22, 24) who (2600	Mortality	Males	(24.393	3 to 43.503)	(6.822 to 10.431)																					
Diarrhoeal diseases	3000) g	Mortality 1	Females	(23.439	9 to 45.567)	(8.227 to 13.042)																					
Diarrhoeal diseases	Birth prevalence - [34, 36) wks, [2000, 2500) g	Mortality	Males	(16.305	(1.925 5 to 29.433)	9.367 (7.859 to 11.112)																					
Diarrhoeal diseases	Birth prevalence - [34, 36) wks, [2000, 2500) g	Mortality 1	Females	(15.657	1.297 7 to 28.761)	10.295 (8.548 to 12.273)																					
Diarrhoeal diseases	Birth prevalence - [37, 38) wks, [2000, 2500) g	Mortality	Males	(10.102	13.0 2 to 16.456)	8.096 (6.724 to 9.676)																					
Diamboeal diseases	Birth prevalence - [37, 38) wks, [2000, 2500) #	Mortality I	Females	(8.805	1.563 5 to 15.11)	8.467 (6.994 to 10.342)																					
Diamboeal diseases	Birth prevalence - [36, 37) wks, [2000, 25000 -	Mortality	Males	(10.789	4.401	8.221																					
Diambogal diseases	Birth prevalence - [36, 37) wks, [2000,	Mortality	Females	13	3.513	8.654																					
Distributed diseases	2500) g Birth prevalence - [34, 36) wks, [2500,	Mertility	Malex	(9.817)	to 17.942) 3.419	(7.215 to 10.369) 5.562																					
	3000) g Birth prevalence - (34, 36) wks, (2500,			(10.387	7 to 16.819) 3.266	(4.646 to 6.696) 6.395																					
Duarthoeal diseases	3000) g Birth musclerup (24, 36) who (4000	Mortality	Females	(9.666	to 17.689)	(5.292 to 7.606)																					
Diarrhoeal diseases	4500) g	Mortality	Males	(14.708	8 to 35.098)	(2.245 to 3.716)																					
Diarrhoeal diseases	Birth prevalence - [34, 36) wks, [4000, 4500) g	Mortality I	Females	(15.763	5.038 3 to 37.255)	3.778 (2.855 to 4.925)																					
Diarrhoeal diseases	Birth prevalence - [34, 36) wks, [3000, 3500) g	Mortality	Males	(10.222	4.006 2 to 18.478)	4.322 (3.449 to 5.338)																					
Diarrhoeal diseases	Birth prevalence - [34, 36) wks, [3000, 3500) g	Mortality I	Females	(10.269	4.375 9 to 20.114)	5.265 (4.145 to 6.564)																					
Diarrhoeal diseases	Birth prevalence - [36, 37) wks, [2500, 3000) g	Mortality	Males	4 (4.014	4.874 4 to 5.713)	3.699 (3.134 to 4.374)																					
Diarrhoeal diseases	Birth prevalence - [36, 37) wks, [2500, 2000) -	Mortality	Females	4	4.609	3.898																					
Diambogal diseases	Birth prevalence - [34, 36) wks, [3500,	Mortality	Males	18	8.024	3.657																					
Disabased Susses	4000) g Birth prevalence - [34, 36) wks, [3500,	Matchin	Frenchas	(12.279	91025.547) 9.263	(2.838 to 4.675) 4.634																					
Diamonal diseases	4000) g Birth prevalence - (37, 38) wks, (2500,			(12.567	7 to 27.924) 3.306	(3.576 to 5.989) 3.194																					
Duarthoeal diseases	3000) g	Mortality	Males	(2.82	to 3.843)	(2.691 to 3.803)																					
Diarrhoeal diseases	3000) g	Mortality	Females	(2.496	6 to 3.61)	(2.745 to 3.817)																					
Diarrhoeal diseases	Birth prevalence - [40, 42) wks, [2500, 3000) g	Mortality	Males	3 (3.002	3.771 2 to 4.693)	3.175 (2.56 to 3.923)																					
Diarrhoeal diseases	Birth prevalence - [40, 42) wks, [2500, 3000) g	Mortality I	Females	3 (2.486	3.244 6 to 4.159)	3.228 (2.605 to 3.985)																					
Diarrhoeal diseases	Birth prevalence - [38, 40) wks, [2500, 3000) g	Mortality	Males	2 (2.274	2.755 4 to 3.309)	2.944 (2.44 to 3.548)																					
Diarrhoeal diseases	Birth prevalence - [38, 40) wks, [2500, 3000) g	Mortality I	Females	2 (1.91	2.376 to 2.886)	2.938 (2.434 to 3.503)																					
Diarrhoeal diseases	Birth prevalence - [36, 37) wks, [3000, 35000 r	Mortality	Males	3 094	3.774	2.466 (2.058 to 2.929)																					
Diarrhoeal diseases	Birth prevalence - [36, 37) wks, [3000,	Mortality I	Females		3.73	2.715																					
Disabased Susses	3500) g Birth prevalence - [36, 37) wks, [4000,	Mastelite	Maler	(2.981	1 to 4.646) 6.826	(2.277 to 3.218) 1.77																					
Product and an	4500) g Birth prevalence - [36, 37) wks, [4000,	Mariany	Tank I	(5.212	2 to 9.045) 7.269	(1.491 to 2.082) 2.177																					
Dumiseli diselses	4500) g Birth menulence - [36: 37) wise [3500	stortany	remates	(5.144	4 to 9.821) 4 544	(1.786 to 2.607) 2.057																					
Diarrhoeal diseases	4000) g	Mortality	Males	(3.64	to 5.622)	(1.74 to 2.44)																					
Diarrhoeal diseases	4000) g	Mortality	Females	(3.577	4.002 7 to 6.014)	(1.98 to 2.864)																					
Diarrhoeal diseases	Birth prevalence - [37, 38) wks, [3000, 3500) g	Mortality	Males	2 (1.759	2.007 9 to 2.293)	1.888 (1.613 to 2.224)																					
Diamboeal diseases	Birth prevalence - [37, 38) wks, [3000, 3500) g	Mortality I	Females	(1.582	1.925 2 to 2.328)	1.972 (1.669 to 2.313)																					
Diarrhoeal diseases	Birth prevalence - [37, 38) wks, [4000, 4500) g	Mortality	Males	(2.596	3.28 6 to 4.133)	1.335 (1.171 to 1.532)																					
Diarrhoeal diseases	Birth prevalence - [37, 38) wks, [4000, 4500) g	Mortality I	Females	3 (2.64	3.521 49 to 4.5)	1.559 (1.333 to 1.835)																					
Diarrhoeal diseases	Birth prevalence - [37, 38) wks, [3500, 4000) r	Mortality	Males	(1.833	2.128 3 to 2.466)	1.505 (1.299 to 1.76)																					
Diamboeal diseases	Birth prevalence - [37, 38) wks, [3500,	Mortality I	Females	2	2.142	1.661																					
Diambocal diseases	Birth prevalence - [40, 42) wks, [3000,	Mortality	Males	(1.094	1.436	1.47																					
Diambocal disease*	5500) g Birth prevalence - [40, 42) wks, [3000,	Mortality	Females	(1.245	1.326	(1.199 to 1.8) 1.465																					
Distributed discour-	3500) g Birth prevalence - [38, 40) wks, [3000,	Mertility	Malex	(1.069	9 to 1.614) 1.33	(1.188 to 1.775) 1.559																					
The second second	3500) g Birth prevalence - [38, 40) wks, 13000.			(1.155	5 to 1.53) 1.224	(1.305 to 1.851) 1.564																					
Duarthoeal diseases	3500) g Birth menulence - 138, 400 wdw 14000	Mortality 1	remates	(1.0	to 1.492) 1.787	(1.304 to 1.847)																					
Diarrhoeal diseases	4500) g	Mortality	Males	(1.453	3 to 2.182)	(1.005 to 1.371)																					
Diamboeal diseases	Bath prevalence - [38, 40) wks, [4000, 4500) g	Mortality I	Females	(1.467	1.877 7 to 2.388)	(1.022 to 1.465)																					
Diarrhoeal diseases	Birth prevalence - [38, 40) wks, [3500, 4000) g	Mortality	Males	1 (1.478	1.785 8 to 2.147)	1.173 (1.0 to 1.377)																					
Diarrhoeal diseases	Birth prevalence - [38, 40) wks, [3500, 4000) g	Mortality I	Females	1 (1.481	1.892 1 to 2.352)	1.23 (1.03 to 1.46)																					
Diarrhoeal diseases	Birth prevalence - [40, 42) wks, [3500, 4000) g	Mortality	Males	(1.0	1.0 0 to 1.0)	1.003 (1.0 to 1.046)																					
Diarrhoeal diseases	Birth prevalence - [40, 42) wks, [3500, 4000) g	Mortality I	Females	1 (1.0)	1.002 to 1.013)	1.001 (1.0 to 1.006)																					
Diarrhoeal diseases	- Birth prevalence - [40, 42) wks, [4000, 4500) n	Mortality	Males		1.0 0 to 1.0)	1.0																					
Diarrhoeal diseases	Birth prevalence - [40, 42) wks, [4000,	Mortality	Females	(1.0	1.0	1.0																					
Diambocal disease*	4500) g Birth prevalence - [28, 30) wks, [2000,	Mortality	Males	(1.0	17.172	(1.0 to 1.0) 27.726																					
Disadar 1.5	2500) g Birth prevalence - [28, 30) wks, [2000.	Mark	Ferral	(83.895)	to 158.056) 21.682	(21.877 to 34.972) 33.983																					
Luarnocal diseases	2500) g Birth menulence - 128, 300 wdw 12500	successfully 1	- unates	(84.349	to 171.375) 7.948	(26.101 to 43.404) 16.408																					
Diarrhoeal diseases	3000) g	Mortality	Males	(54.687	to 105.047)	(12.653 to 21.188)																					
Diarrhoeal diseases	nomh prevalence - [28, 30) wks, [2500, 3000) g	Mortality I	Females	(54.099)	9.193 to 112.236)	20.387 (15.089 to 26.434)																					
Diarrhoeal diseases	Birth prevalence - [28, 30) wks, [3000, 3500) g	Mortality	Males	42 (29.891	(2.199 1 to 57.227)	10.082 (7.777 to 13.056)																					
Diarrhoeal diseases	Birth prevalence - [28, 30) wks, [3000, 3500) g	Mortality I	Females	42 (28.25)	2.551 to 63.209)	11.989 (9.084 to 15.544)																					

spendix rable ba. Relative rist	is used by age and sex for each outcom	101 all FISK lactors e	exceptions	amplent all pollution al	conton and	smosting.									A	ges											
Rick - Ontcome	Category / Unite M	arhidity / Mortality	Ser	All-age 0-6 d	ays	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Diamboral diseases	Birth prevalence - [30, 32] wks, [2500,	Mortality	Males	58.7	22	12.115																					
Diambogal diseases	Birth prevalence - [30, 32) wks, [2500,	Mortality	Females	(42.41910 59.5)	22	15.364																					
Disabased James	3000) g Birth prevalence - [30, 32) wks, [3000,	Mantalian	Maler	(42.058 to 45.6	82.793) (7	11.936 to 19.581) 8.381																					
	3500) g Birth menulence - (30, 32) wks, (3000,	January .	mana	(32.014 to 46.1)	65.531) ()4	(6.362 to 10.842) 10.506																					
Durrhoeal diseases	3500) g Risk susselses (20, 22) who (2000)	Mortality	Females	(30.243 to	66.207) ((8.041 to 13.513)																					
Diarrhoeal diseases	4000) g	Mortality	Males	(21.558 to	54.813)	(4.293 to 7.349)																					
Diarrhoeal diseases	Both prevalence - [30, 32) wks, [3500, 4000) g	Mortality	Females	(22.692 to	61.276)	6.892 (5.175 to 9.058)																					
Diarrhoeal diseases	Birth prevalence - [32, 34) wks, [3000, 3500) g	Mortality	Males	34.0 (23.515 to	48.37)	6.577 (4.934 to 8.436)																					
Diarrhoeal diseases	Birth prevalence - [32, 34) wks, [3000, 3500) g	Mortality	Females	34.5 (22.909 to	85 49.754) (8.314 (6.362 to 10.789)																					
Diarrhoeal diseases	Birth prevalence - [32, 34) wks, [3500, 4000) g	Mortality	Males	36.2- (23.158 to	18 54.67)	5.068 (3.761 to 6.741)																					
Diarrhocal diseases	Birth prevalence - [32, 34) wks, [3500, 4000) g	Mortality	Females	38.0 (23.301 to	98 59.429)	6.476 (4.666 to 8.689)																					
Diarrhoeal diseases	Birth prevalence - [36, 37) wks, [1000, 15000 -	Mortality	Males	166.6	86	57.535																					
Diambogal diseases	Birth prevalence - [36, 37) wks, [1000,	Mortality	Females	169.7	25	63.564																					
Distributed diseases	1500) g Birth prevalence - [38, 40) wks, [1000,	Martality	Malex	174.0	66 (:	57.966																					
No. 1	1500) g Birth prevalence - [38, 40) wks, [1000,		To a da	(125.125 to 171.5	232.507) (· 57	44.393 to 73.241) 65.208																					
Diamiseal diseases	1500) g Birth merculance - [38, 40) wise (1500	storcarty	remates	(121.585 to 67.3)	237.047) (·	48.821 to 84.308) 25.206																					
Diarrhoeal diseases	2000) g	Mortality	Males	(49.547 to	89.055) (J	20.365 to 31.168)																					
Diarrhoeal diseases	2000) g	Mortality	Females	(45.884 to	9 83.445) (J	22.625 to 35.139)																					
Diarrhoeal diseases	Birth prevalence - [40, 42) wks, [1500, 2000) g	Mortality	Males	76.6 (56.177 to	73 102.468) (25.785 19.387 to 34.168)																					
Diarrhoeal diseases	Birth prevalence - [40, 42) wks, [1500, 2000) g	Mortality	Females	70.4 (49.221 to	11 97.952) (3	29.113 21.355 to 38.272)																					
Pneumococcal meningitis	Birth prevalence - [0, 24) wks, [0, 500) g	Mortality	Males	1564. (1056.5 2116.0	792 42 to (4:	618.595 58.842 to 812.921)																					
Pneumococcal meningitis	Birth prevalence - [0, 24) wks, [0, 500) g	Mortality	Females	1600 (1050.6	122 64 to (5	713.571 26.178 to 921.018)																					
Pneumococcal meningitis	Birth prevalence - [0, 24) wks, [500, 1000)	Mortality	Males	1155.1	815	457.5																					
Pneumococcal meningitis	o Birth prevalence - [0, 24) wks, [500, 1000)	Mortality	Females	1169.	123	515.406																					
Paramacrocal menineitia	8 Birth prevalence - [24, 26) wks, [500, 1000)	Martality	Malex	(802.003.05	83	443.357																					
Provincial manipula	8 Birth prevalence - [24, 26) wks, [500, 1000)	Mantalian	Females	(723.748 to) 947.1	1244.265) (3 43	487.549																					
	8 Birth prevalence - (26, 28) wks, (500, 1000)	January .	- conserve	(702.662 to 497.8	1237.093) (3: 17	87.307 to 603.498) 330.886																					
Pneumococcal menungitis	8 Birth merculance - (26-28) wise (500-1000)	Mortality	Males	(377.617 to	648.547) (2) 82	61.438 to 401.709) 344.618																					
Pneumococcal meningitis	8	Mortality	Females	(354.946 to	629.517) (2	74.427 to 419.864)																					
Pneumococcal meningitis	8	Mortality	Males	(163.821 to	324.502) (1	149.995 17.866 to 188.368)																					
Pneumococcal meningitis	Birth prevalence - [30, 32) wks, [500, 1000) 8	Mortality	Females	229.1 (157.606 to	97 317.194) (1:	152.117 20.779 to 190.583)																					
Pneumococcal meningitis	Birth prevalence - [28, 30] wks, [500, 1000) 8	Mortality	Males	297.6 (214.953 to	29 396.586) (1	216.995 73.321 to 271.466)																					
Pneumococcal meningitis	Birth prevalence - [28, 30) wks, [500, 1000) g	Mortality	Females	281.0 (198.176 to	56 386.635) (1	219.884 74.264 to 272.704)																					
Pneumococcal meningitis	Birth prevalence - [26, 28) wks, [1000, 1500) g	Mortality	Males	267.5 (210.177 to	332.92) (1:	164.167 32.898 to 200.569)																					
Pneumococcal meningitis	Birth prevalence - [26, 28) wks, [1000, 1500) a	Mortality	Females	266.5 (197.461 to	09 346.932) (1	174.222 37.431 to 217.349)																					
Pneumococcal meningitis	Birth prevalence - [34, 36) wks, [1000, 1500) a	Mortality	Males	142.0	56 197 774)	52.86 42.914 in 64.617)																					
Pneumococcal meningitis	Birth prevalence - [34, 36) wks, [1000,	Mortality	Females	141.8	99	57.421																					
Pneumococcal menineitis	Birth prevalence - [28, 30) wks, [1500,	Mortality	Males	(95.86410	66	50.018																					
Procession and and and and a	2000) g Birth prevalence - [28, 30) wks, [1500,	Mantalian	Females	(97.178 to 130.9	24 (·	40.539 to 61.919) 57.275																					
hanneed a stampto	2000) g Birth prevalence - [28, 30) wks, [1000,	Marin	No.	(96.513 to 158.5	172.188) (63	(46.36 to 70.038) 103.32																					
Preunococca menugins	1500) g Birth merculance - [28, 30) wise [1000]	storcarty	Males	(120.99 to 1	204.947) (8 05	13.486 to 127.144)																					
Pneumococcal meningitis	1500) g	Mortality	Females	(112.327 to	200.786) (1	86.954 to 131.78)																					
Pneumococcal meningitis	1500) g	Mortality	Males	(81.354 to	(61.101)	53.185 43.049 to 66.274)																					
Pneumococcal meningitis	Both prevalence - [32, 34) wks, [1000, 1500) g	Mortality	Females	(79.363 to	71 159.206) (56.034 (45.982 to 68.36)																					
Pneumococcal meningitis	Birth prevalence - [30, 32) wks, [1000, 1500) g	Mortality	Males	119.3 (87.769 to	08 160.885) (1	67.163 54.863 to 82.638)																					
Pneumococcal meningitis	Birth prevalence - [30, 32) wks, [1000, 1500) g	Mortality	Females	115.4 (84.272 to	48 156.425) (1	69.14 55.873 to 85.012)																					
Pneumococcal meningitis	Birth prevalence - [37, 38) wks, [1500, 2000) g	Mortality	Males	62.9 (46.159 to	72 83.484) (J	24.148 20.066 to 29.406)																					
Pneumococcal meningitis	Birth prevalence - [37, 38) wks, [1500, 2000) g	Mortality	Females	59.90 (43.974 to	88 79.053) (J	26.719 21.746 to 32.816)																					
Pneumococcal meningitis	Birth prevalence - [36, 37] wks, [1500, 2000) g	Mortality	Males	60.2 (43.669 to	18 (82.48)	23.031 18.793 to 28.483)																					
Pneumococcal meningitis	Birth prevalence - [36, 37) wks, [1500, 2000) -	Mortality	Females	58.5	27	25.143																					
Pneumococcal meningitis	Birth prevalence - [30, 32) wks, [2000, 2500) -	Mortality	Males	67.9	71	18.03																					
Pneumococcal menineiti+	2.500) g Birth prevalence - [30, 32) wks, [2000, 2.500)	Mortality	Females	(50.354 to 69.3	83.933) (22.069																					
Pneumocreved manimiti-	2:500) g Birth prevalence - [30, 32) wks, [1500,	Mortality	Males	(49.108 to 77.3)	99 (99	17.4536 to 27.163) 31.079																					
numerous manipus	2000) g Birth prevalence - [30, 32) wks. [1500.	Markany Markany		(59.702 to 76 1	99.232) (. 34	25.786 to 36.724) 34.756																					
Pneumococcal menungitis	2000) g Birth mercalence - [34-36) wisy [1500	Mortality	remates	(56.885 to	100.996) (J	28.764 to 41.849) 21.346																					
Pneumococcal meningitis	2000) g Birth recordence - [34, 26) where [1,600]	Mortality	Males	(39.553 to	75.104) (17.677 to 26.143)																					
Pneumococcal meningitis	2000) g	Mortality	Females	54.3 (38.617 to	(75.24)	18.743 to 28.287)																					
Pneumococcal meningitis	north prevalence - [32, 34) wks, [1500, 2000) g	Mortality	Males	57.1 (42.484 to	73.651) (25.114 19.028 to 27.915)																					
Pneumococcal meningitis	Birth prevalence - [32, 34) wks, [1500, 2000) g	Mortality	Females	56.1 (39.794 to	01 76.295) (25.149 20.615 to 30.388)																					
Pneumococcal meningitis	Birth prevalence - [32, 34) wks, [2000, 2500) g	Mortality	Males	37.4 (29.026 to	44 48.227) (12.233 10.252 to 14.477)																					
Pneumococcal meningitis	Birth prevalence - [32, 34) wks, [2000, 2500) g	Mortality	Females	36.8' (26.658 to	74 49.653) (14.384 (12.095 to 17.03)																					
Pneumococcal meningitis	Birth prevalence - [40, 42) wks, [2000, 2500) g	Mortality	Males	18.0 (13.292 to	23.719)	9.23 (7.037 to 11.454)																					
				1																							

	counce by age and sex for each outer			e annoc ne an p		nu anoxing.									А	iges											
NI G				All-age	0-6 days	7-27 days	28-364 days 1	-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Preumococcal menineitis	Birth prevalence - [40, 42) wks, [2000,	Mortality / Mortali	Females		15.574	9.975		I				1	I		1	1 1	1	1					1 1				
	2500) g Birth prevalence - [38, 40) wks, [2000,				(11.516 to 20.778) 13.104	(7.82 to 12.46) 8.198																					
Pheumococca menugins	2500) g	stortany	MIRO		(9.829 to 16.99)	(6.786 to 9.959)																					
Pneumococcal meningitis	2500) g	Mortality	Females		(8.389 to 14.38)	(7.04 to 10.449)																					
Pneumococcal meningitis	Birth prevalence - [32, 34) wks, [2500, 3000) g	Mortality	Males		33.063 (24.393 to 43.503)	8.441 (6.822 to 10.431)																					
Pneumococcal meningitis	Birth prevalence - [32, 34) wks, [2500, 3000) g	Mortality	Females		32.812 (23.439 to 45.567)	10.398 (8.227 to 13.042)																					
Pneumococcal meningitis	Birth prevalence - [34, 36) wks, [2000, 2500) -	Mortality	Males		21.925	9.367																					
Pneumococcal menineitis	Bith prevalence - [34, 36) wks, [2000,	Mortality	Females		21.297	10.295																					
	2500) g Birth mevalence - (37, 38) wks. (2000,				(15.657 to 28.761) 13.0	(8.548 to 12.273) 8.096																					
Pheumococca menugins	2500) g Risk muschener (27, 28) sole (2000	stortany	MIRO		(10.102 to 16.456)	(6.724 to 9.676)																					
Pneumococcal meningitis	2500) g	Mortality	Females		(8.805 to 15.11)	(6.994 to 10.342)																					
Pneumococcal meningitis	Birth prevalence - [36, 37) wks, [2000, 2500) g	Mortality	Males		14.401 (10.789 to 18.654)	8.221 (6.917 to 9.923)																					
Pneumococcal meningitis	Birth prevalence - [36, 37) wks, [2000, 2500) g	Mortality	Females		13.513 (9.817 to 17.942)	8.654 (7.215 to 10.369)																					
Pneumococcal meningitis	Birth prevalence - [34, 36) wks, [2500, 3000) g	Mortality	Males		13.419 (10.387 to 16.819)	5.562 (4.646 to 6.696)																					
Pneumococcal meningitis	Birth prevalence - [34, 36) wks, [2500, 2000) -	Mortality	Females		13.266	6.395																					
Presence and meminating	Birth prevalence - [34, 36) wks, [4000,	Martility	Malex		23.096	2.895																					
	4500) g Birth menulence - (34, 36) wks. (4000.				(14.708 to 35.098) 25.038	(2.245 to 3.716) 3.778																					
Pneumococcal meningitis	4500) g	Mortality	Females		(15.763 to 37.255)	(2.855 to 4.925)																					
Pneumococcal meningitis	13500) g	Mortality	Males		(10.222 to 18.478)	4.322 (3.449 to 5.338)																					
Pneumococcal meningitis	Birth prevalence - [34, 36) wks, [3000, 3500) g	Mortality	Females		14.375 (10.269 to 20.114)	5.265 (4.145 to 6.564)																					
Pneumococcal meningitis	Birth prevalence - [36, 37) wks, [2500, 3000) g	Mortality	Males		4.874 (4.014 to 5.713)	3.699 (3.134 to 4.374)																					
Pneumococcal meningitis	Birth prevalence - [36, 37) wks, [2500, 3000) #	Mortality	Females		4.609 (3.731 to 5.61)	3.898 (3.235 to 4.67)																					
Pneumococcal menineitis	Birth prevalence - [34, 36) wks, [3500,	Mortality	Malex		18.024	3.657																					
Prov. 1	4000) g Birth prevalence - [34, 36) wks, [3500,				(12.279 to 25.547) 19.263	(2.838 to 4.675) 4.634																					
Pheumococca menugins	4000) g Risk muschener (27, 28) min (2600	stortany	Pemases		(12.567 to 27.924) 2.206	(3.576 to 5.989)																					
Pneumococcal meningitis	3000) g	Mortality	Males		(2.82 to 3.843)	(2.691 to 3.803)																					
Pneumococcal meningitis	Birth prevalence - [37, 38) wks, [2500, 3000) g	Mortality	Females		2.991 (2.496 to 3.61)	3.242 (2.745 to 3.817)																					
Pneumococcal meningitis	Birth prevalence - [40, 42) wks, [2500, 3000) g	Mortality	Males		3.771 (3.002 to 4.693)	3.175 (2.56 to 3.923)																					
Pneumococcal meningitis	Birth prevalence - [40, 42) wks, [2500, 3000) g	Mortality	Females		3.244 (2.486 to 4.159)	3.228 (2.605 to 3.985)																					
Pneumococcal meningitis	Birth prevalence - [38, 40) wks, [2500, 2000) -	Mortality	Males		2.755	2.944																					
Pneumococcal menineitis	Bith prevalence - [38, 40) wks, [2500,	Mortality	Females		2.376	2.938																					
Personal environition	3000) g Birth prevalence - [36, 37) wks, [3000,	Mantalian	Malar		(1.91 to 2.886) 3.774	(2.434 to 3.503) 2.466																					
Pheumococca menugins	3500) g Birth menulance - [36-37) wks [3000	storcarty	MINO		(3.094 to 4.497) 3.73	(2.058 to 2.929) 2.715																					
Pneumococcal meningitis	3500) g	Mortality	Females		(2.981 to 4.646)	(2.277 to 3.218)																					
Pneumococcal meningitis	Birth prevalence - [36, 37) wks, [4000, 4500) g	Mortality	Males		6.826 (5.212 to 9.045)	1.77 (1.491 to 2.082)																					
Pneumococcal meningitis	Birth prevalence - [36, 37) wks, [4000, 4500) g	Mortality	Females		7.269 (5.144 to 9.821)	2.177 (1.786 to 2.607)																					
Pneumococcal meningitis	Birth prevalence - [36, 37) wks, [3500, 4000) g	Mortality	Males		4.544 (3.64 to 5.622)	2.057 (1.74 to 2.44)																					
Pneumococcal meningitis	Birth prevalence - [36, 37) wks, [3500, 4000) g	Mortality	Females		4.662 (3.577 to 6.014)	2.398 (1.98 to 2.864)																					
Pneumococcal meningitis	Birth prevalence - [37, 38) wks, [3000, 25000 -	Mortality	Males		2.007	1.888																					
Pneumococcal menineitis	Birth prevalence - [37, 38) wks, [3000,	Mortality	Females		1.925	1.972																					
Personal maximitie	3500) g Birth prevalence - [37, 38) wks, [4000,	Mantality	Malar		3.28	1.335																					
-	4500) g Birth merculence - (37, 38) wks. (4000.				(2.596 to 4.133) 3.521	(1.171 to 1.532) 1.559																					
Pneumococcal meningitis	4500) g	Mortality	Females		(2.649 to 4.5)	(1.333 to 1.835)																					
Pneumococcal meningitis	Both prevalence - [37, 38) wks, [3500, 4000) g	Mortality	Males		2.128 (1.833 to 2.466)	(1.299 to 1.76)																					
Pneumococcal meningitis	Birth prevalence - [37, 38) wks, [3500, 4000) g	Mortality	Females		2.142 (1.694 to 2.67)	1.661 (1.411 to 1.961)																					
Pneumococcal meningitis	Birth prevalence - [40, 42) wks, [3000, 3500) g	Mortality	Males		1.436 (1.245 to 1.65)	1.47 (1.199 to 1.8)																					
Pneumococcal meningitis	Birth prevalence - [40, 42) wks, [3000, 3500) g	Mortality	Females		1.326 (1.069 to 1.614)	1.465 (1.188 to 1.775)																					
Pneumococcal meningitis	Birth prevalence - [38, 40) wks, [3000, 3500) #	Mortality	Males		1.33	1.559																					
Presence and meminating	Birth prevalence - [38, 40) wks, [3000,	Martility	Females		1.224	1.564																					
Bernard and and and a second	3500) g Birth prevalence - [38, 40) wks, [4000,	Mantalian	Malar		(1.0 to 1.492) 1.787	(1.304 to 1.847) 1.175																					
Pheumococca menugins	4500) g Risk musclasse (28, 40) sole (4000	storcarty	MINO		(1.453 to 2.182)	(1.005 to 1.371)																					
Pneumococcal meningitis	4500) g	Mortality	Females		(1.467 to 2.388)	(1.022 to 1.465)																					
Pneumococcal meningitis	Both prevalence - [38, 40) wks, [3500, 4000) g	Mortality	Males		1.785 (1.478 to 2.147)	1.173 (1.0 to 1.377)																					
Pneumococcal meningitis	Birth prevalence - [38, 40) wks, [3500, 4000) g	Mortality	Females		1.892 (1.481 to 2.352)	1.23 (1.03 to 1.46)																					
Pneumococcal meningitis	Birth prevalence - [40, 42) wks, [3500, 4000) g	Mortality	Males		1.0 (1.0 to 1.0)	1.003 (1.0 to 1.046)																					
Pneumococcal meningitis	Birth prevalence - [40, 42) wks, [3500, 4000) g	Mortality	Females		1.002 (1.0 to 1.013)	1.001 (1.0 to 1.006)																					
Pneumococcal meningitis	Birth prevalence - [40, 42) wks, [4000, 4500) g	Mortality	Males		1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																					
Pneumococcal meningitis	Birth prevalence - [40, 42) wks, [4000,	Mortality	Females		1.0	1.0																					
Parameter and manipulti-	+500) g Birth prevalence - [28, 30) wks, [2000,	Martality	Malex		(1.0 to 1.0) 117.172	(1.0 to 1.0) 27.726																					
- management	2500) g Bith mevalence - 128-300 wise 12000	ana any	-		(83.895 to 158.056) 121.682	(21.877 to 34.972) 33.983																					
Pneumococcal meningitis	2500) g Rich musclama (28, 30) ada (2000)	Mortality	Pemales		(84.349 to 171.375)	(26.101 to 43.404)																					
Pneumococcal meningitis	3000) g	Mortality	Males		(54.687 to 105.047)	(12.653 to 21.188)																					
Pneumococcal meningitis	Birth prevalence - [28, 30) wks, [2500, 3000) g	Mortality	Females		79.193 (54.099 to 112.236)	20.387 (15.089 to 26.434)																					
Pneumococcal meningitis	Birth prevalence - [28, 30) wks, [3000, 3500) g	Mortality	Males		42.199 (29.891 to 57.227)	10.082 (7.777 to 13.056)																					
Pneumococcal meningitis	Birth prevalence - [28, 30) wks, [3000, 3500) g	Mortality	Females		42.551 (28.25 to 63.209)	11.989 (9.084 to 15.544)																					

ppendix Table 6a. Relative risks	s used by age and sex for each oute	ome for all risk factors	except for ambient a	r pollution alcohol,	and smoking.								Ages										
Birk Outerma	Cotoner / Unite	Manhidita / Mantalita	All-age	0-6 days	7-27 days	28-364 days 1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years 30-3	34 years 35-39	years 40-44 years	45-49 years	50-54 years	55-59 years 6	0-64 years 65-	69 years 70-7	14 years 75-7	79 years 80-84 y	ars 85-89 year	s 90-94 years	95+ years
Pneumococcal meningitis	Birth prevalence - [30, 32) wks, [2500,	Mortality	Males	58.722	12.115																		
Preumocoved menineitis	3000) g Birth prevalence - [30, 32) wks, [2500,	Manufity	Females	(42.419 to 78.873) 59.522	(9.518 to 15.521) 15.364																		
	3000) g Birth prevalence - [30, 32) wks, [3000,	M		(42.058 to 82.793) 45.67	(11.936 to 19.581) 8.381																		
Pheumococca meningins	3500) g Birth menalence - (30-32) wise (3000	storcarry	MIRS	(32.014 to 65.531) 46.104	(6.362 to 10.842) 10.506																		
Pneumococcal meningitis	3500) g	Mortality	Females	(30.243 to 66.207)	(8.041 to 13.513)																		
Pneumococcal meningitis	Birth prevalence - [30, 32) wks, [3500, 4000) g	Mortality	Males	36.334 (21.558 to 54.813)	5.698 (4.293 to 7.349)																		
Pneumococcal meningitis	Birth prevalence - [30, 32) wks, [3500, 4000) g	Mortality	Females	37.931 (22.692 to 61.276)	6.892 (5.175 to 9.058)																		
Pneumococcal meningitis	Birth prevalence - [32, 34) wks, [3000, 3500) g	Mortality	Males	34.016 (23.515 to 48.37)	6.577 (4.934 to 8.436)																		
Pneumococcal meningitis	Birth prevalence - [32, 34) wks, [3000, 3500) #	Mortality	Females	34.585 (22.909 to 49.754)	8.314 (6.362 to 10.789)																		
Pneumococcal menineitis	Birth prevalence - [32, 34) wks, [3500,	Mortality	Males	36.248	5.068																		
December 201	4000) g Birth prevalence - [32, 34) wks, [3500,	Manufilite	Ferrela	(23.158 to 54.67) 38.098	(3.761 to 6.741) 6.476																		
	4000) g Birth mevalence - (36, 37) wks, (1000,			(23.301 to 59.429) 166.686	(4.666 to 8.689) 57.535																		
Pneumococcal meningitis	1500) g Rishermahara (36, 27) ada (1000	Mortality	Males	(118.487 to 222.581)	(45.999 to 71.742)																		
Pneumococcal meningitis	1500) g	Mortality	Females	(119.017 to 229.008)	(50.068 to 80.703)																		
Pneumococcal meningitis	Birth prevalence - [38, 40) wks, [1000, 1500) g	Mortality	Males	174.066 (125.125 to 232.507)	57.966 (44.393 to 73.241)																		
Pneumococcal meningitis	Birth prevalence - [38, 40) wks, [1000, 1500) g	Mortality	Females	171.557 (121.585 to 237.047)	65.208 (48.821 to 84.308)																		
Pneumococcal meningitis	Birth prevalence - [38, 40) wks, [1500, 2000) g	Mortality	Males	67.302 (49.547 to 89.055)	25.206 (20.365 to 31.168)																		
Pneumococcal meningitis	Birth prevalence - [38, 40) wks, [1500, 2000) #	Mortality	Females	62.19 (45 884 to 83 445)	28.05 (22.625 to 35.139)																		
Pneumococcal meningitis	Birth prevalence - [40, 42) wks, [1500,	Mortality	Males	76.673	25.785																		
Preumocoved menineitis	Birth prevalence - [40, 42) wks, [1500,	Mandity	Females	(56.17716-102.468) 70.411	29.113																		
	2000) g	M F.		(49.221 to 97.952) 1564.792	(21.355 to 38.272) 618.595																		
ri initienzae type is meninguis	isanii prevazence - [0, 24) wici, [0, 500) g	storcarry	MINS	(1056.542 to 2116.062) 1600.122	(458.842 to 812.921)																		
H influenzae type B meningitis	Birth prevalence - [0, 24) wks, [0, 500) g	Mortality	Females	(1050.664 to 2011.027)	(526.178 to 921.018)																		
H influenzae type B meningitis	Birth prevalence - [0, 24) wks, [500, 1000) g	Mortality	Males	(825.412 to 1506.837)	457.5 (352.552 to 573.483)																		
H influenzae type B meningitis	Birth prevalence - [0, 24) wks, [500, 1000) 8	Mortality	Females	1169.123 (802.003 to 1617.979)	515.406 (396.713 to 641.541)																		
H influenzae type B meningitis	Birth prevalence - [24, 26) wks, [500, 1000 g) Mortality	Males	955.583 (723.748 to 1244.265)	443.357 (363.03 to 534.695)																		
H influenzae type B meningitis	Birth prevalence - [24, 26) wks, [500, 1000) Mortality	Females	947.143 (702.662 to 1237.093)	487.549 (387.307 in 603.498)																		
H influenzae type B meningitis	o Birth prevalence - [26, 28) wks, [500, 1000) Mortality	Males	497.817	330.886																		
H influences tone B menimities	8 Birth prevalence - [26, 28) wks, [500, 1000	Mandin	Females	483.682	344.618																		
	8 Birth prevalence - [30, 32) wks, [500, 1000) M P.		(354.946 to 629.517) 236.614	(274.427 to 419.864) 149.995																		
ri initiettise type is meningitis	8 Richmanham (20.22) who (500.1000	MORTANY	MIRS	(163.821 to 324.502) 220.107	(117.866 to 188.368)																		
H influenzae type B meningitis	8	Mortality	Females	(157.606 to 317.194)	(120.779 to 190.583)																		
H influenzae type B meningitis	Barth prevalence - [28, 30] wks, [500, 1000 8) Mortality	Males	297.629 (214.953 to 396.586)	216.995 (173.321 to 271.466)																		
H influenzae type B meningitis	Birth prevalence - [28, 30) wks, [500, 1000 8) Mortality	Females	281.056 (198.176 to 386.635)	219.884 (174.264 to 272.704)																		
H influenzae type B meningitis	Birth prevalence - [26, 28) wks, [1000, 1500) g	Mortality	Males	267.91 (210.177 to 332.92)	164.167 (132.898 to 200.569)																		
H influenzae type B meningitis	Birth prevalence - [26, 28) wks, [1000, 1500) g	Mortality	Females	266.509 (197.461 to 346.932)	174.222 (137.431 to 217.349)																		
H influenzae type B meningitis	Birth prevalence - [34, 36) wks, [1000, 1500) #	Mortality	Males	142.056 (98.086 to 197.774)	52.86 (42.914 to 64.617)																		
H influenzae type B meningitis	Birth prevalence - [34, 36) wks, [1000,	Mortality	Females	141.899	57.421																		
Hinformus trav Prominsitio	1500) g Birth prevalence - [28, 30) wks, [1500,	Manufilite	Mile	(95.864 to 197.656) 127.966	(46.452 to 71.539) 50.018																		
in annual type in annughts	2000) g Birth meyalence - (28, 30) wks, (1500,			(97.178 to 167.026) 130.924	(40.539 to 61.919) 57.275																		
H influenzae type B meningriss	2000) g Rishermahara (28, 20) ada (1000	Mortality	Females	(96.513 to 172.188)	(46.36 to 70.038)																		
H influenzae type B meningitis	1500) g	Mortality	Males	(120.99 to 204.947)	(83.486 to 127.144)																		
H influenzae type B meningitis	Barth prevalence - [28, 30) wks, [1000, 1500) g	Mortality	Females	(112.327 to 200.786)	107.529 (86.954 to 131.78)																		
H influenzae type B meningitis	Birth prevalence - [32, 34) wks, [1000, 1500) g	Mortality	Males	117.142 (81.354 to 161.101)	53.185 (43.049 to 66.274)																		
H influenzae type B meningitis	Birth prevalence - [32, 34) wks, [1000, 1500) g	Mortality	Females	115.171 (79.363 to 159.206)	56.034 (45.982 to 68.36)																		
H influenzae type B meningitis	Birth prevalence - [30, 32) wks, [1000, 1500) g	Mortality	Males	119.308 (87.769 to 160.885)	67.163 (54.863 to 82.638)																		
H influenzae type B meningitis	Birth prevalence - [30, 32) wks, [1000, 1500) g	Mortality	Females	115.448 (84.272 to 156.475)	69.14 (55.873 to 85.012)																		
H influenzae type B meningitis	Birth prevalence - [37, 38) wks, [1500, 2000) #	Mortality	Males	62.972	24.148																		
H influenzae type B menineitis	Birth prevalence - [37, 38) wks, [1500,	Mortality	Females	59.988	26.719																		
Hardsman has R	2000) g Birth prevalence - [36, 37) wks, [1500,	Mantalita	Malar	(43.974 to 79.053) 60.218	(21.746 to 32.816) 23.031																		
ri initiettise type is meningitis	2000) g Birth menalence - [36-37) wks [1500	storcarry	MIRS	(43.669 to 82.48) 58.527	(18.793 to 28.483) 25.143																		
H influenzae type B meningitis	2000) g	Mortality	Females	(42.172 to 80.557)	(20.331 to 30.566)																		
H influenzae type B meningitis	Both prevalence - [30, 32] wks, [2000, 2500) g	Mortality	Males	67.971 (50.354 to 88.935)	18.03 (14.621 to 22.103)																		
H influenzae type B meningitis	Birth prevalence - [30, 32) wks, [2000, 2500) g	Mortality	Females	69.383 (49.108 to 94.583)	22.069 (17.836 to 27.163)																		
H influenzae type B meningitis	Birth prevalence - [30, 32) wks, [1500, 2000) g	Mortality	Males	77.369 (59.702 to 99.232)	31.079 (25.786 to 36.724)																		
H influenzae type B meningitis	Birth prevalence - [30, 32) wks, [1500, 2000) g	Mortality	Females	76.134 (56.885 to 100.996)	34.756 (28.764 to 41.849)																		
H influenzae type B meningitis	Birth prevalence - [34, 36) wks, [1500, 2000) -	Mortality	Males	55.555	21.346																		
H influenzae type B menjoaritis	Birth prevalence - [34, 36) wks, [1500,	Mortality	Females	(39.353 to 75.104) 54.335	(17.0771026.143) 23.046																		
History	2000) g Birth prevalence - [32, 34) wks, [1500.	M	Malar	(38.617 to 75.24) 57.155	(18.743 to 28.287) 23.114																		
ri intractorae type B meningitis	2000) g Birth prevalence - (32, 34) wks (1500	asortality	militis	(42.484 to 73.651) 56.101	(19.028 to 27.915) 25.149																		
H influenzae type B meningitis	2000) g	Mortality	Females	(39.794 to 76.295)	(20.615 to 30.388)																		
H influenzae type B meningitis	asini prevaance - [32, 34) wks, [2000, 2500) g	Mortality	Males	57.444 (29.026 to 48.227)	(10.252 to 14.477)																		
H influenzae type B meningitis	Birth prevalence - [32, 34) wks, [2000, 2500) g	Mortality	Females	36.874 (26.658 to 49.653)	14.384 (12.095 to 17.03)																		
H influenzae type B meningitis	Birth prevalence - [40, 42) wks, [2000, 2500) g	Mortality	Males	18.092 (13.292 to 23.719)	9.23 (7.037 to 11.454)																		

opendix rable ba. Relative risks	used by age and sex tor each outco	ne for an risk factor	ALC: US	r anoren ar p	ontition second, s	ind shidsing.									A	ges											
Risk - Outcome	Category / Units	Morbidity / Mortalit	v Sex	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
H influenzae type B meningitis	Birth prevalence - [40, 42) wks, [2000, 2500) a	Mortality	Females		15.574 (11.516 to 20.778)	9.975 (7.82 to 12.46)																					
H influenzae type B meningitis	Birth prevalence - [38, 40) wks, [2000, 2500) #	Mortality	Males		13.104 (9.879 to 16.99)	8.198 (6.786 to 9.959)																					
H influenzae type B meningitis	Birth prevalence - [38, 40) wks, [2000, 2500) =	Mortality	Females		11.308 (8.389 to 14.38)	8.577 (7.04 to 10.449)																					
H influenzae type B meningitis	Birth prevalence - [32, 34) wks, [2500, 20000 -	Mortality	Males		33.063	8.441																					
H influenzae type B meningitis	Bith prevalence - [32, 34) wks, [2500,	Mortality	Females		32.812	10.398																					
Hinfluenzae vere B menineitis	5000) g Birth prevalence - [34, 36) wks, [2000,	Mortality	Males		(23.43910.43.367) 21.925	9.367																					
Hinfluenzae type B menineitis	2500) g Birth prevalence - [34, 36) wks, [2000,	Mortality	Females		(16.305 to 29.433) 21.297	(7.859 to 11.112) 10.295																					
Hinfluence tone B menineitia	2500) g Birth prevalence - [37, 38) wks, [2000,	Manulity	Malex		(15.657 to 28.761) 13.0	(8.548 to 12.273) 8.096																					
Hinfluence type B meninging	2500) g Birth prevalence - [37, 38) wks, [2000,	Manulity	Females		(10.102 to 16.456) 11.563	(6.724 to 9.676) 8.467																					
II and the providence of the second second	2500) g Birth prevalence - [36, 37) wks, [2000,	Mariany	No.		(8.805 to 15.11) 14.401	(6.994 to 10.342) 8.221																					
H intractive type is meninging	2500) g Birth prevalence - [36, 37) wks, [2000,	Mortany	Males -		(10.789 to 18.654) 13.513	(6.917 to 9.923) 8.654																					
H intractive type is meninging	2500) g Birth prevalence - [34, 36) wks, [2500,	Mortany	remates		(9.817 to 17.942) 13.419	(7.215 to 10.369) 5.562																					
H influenzae type B meningitis	3000) g Birth merculance - [34, 36) why [2500	Moriality	Males		(10.387 to 16.819)	(4.646 to 6.696) 6 395																					
H influenzae type B meningitis	3000) g Birth menulation (24, 36) solio (4000	Mortality	Females		(9.666 to 17.689)	(5.292 to 7.606)																					
H influenzae type B meningitis	4500) g	Mortality	Males		(14.708 to 35.098)	(2.245 to 3.716)																					
H influenzae type B meningitis	4500) g	Mortality	Females		(15.763 to 37.255)	(2.855 to 4.925)																					
H influenzae type B meningitis	1500) g	Mortality	Males		(10.222 to 18.478)	4.322 (3.449 to 5.338)																					
H influenzae type B meningitis	Barth prevalence - [34, 36) wks, [3000, 3500) g	Mortality	Females		14.575 (10.269 to 20.114)	5.265 (4.145 to 6.564)																					
H influenzae type B meningitis	Birth prevalence - [36, 37] wks, [2500, 3000) g	Mortality	Males		4.874 (4.014 to 5.713)	3.699 (3.134 to 4.374)																					
H influenzae type B meningitis	Birth prevalence - [36, 37) wks, [2500, 3000) g	Mortality	Females		4.609 (3.731 to 5.61)	3.898 (3.235 to 4.67)																					
H influenzae type B meningitis	Birth prevalence - [34, 36) wks, [3500, 4000) g	Mortality	Males		18.024 (12.279 to 25.547)	3.657 (2.838 to 4.675)																					
H influenzae type B meningitis	Birth prevalence - [34, 36) wks, [3500, 4000) g	Mortality	Females		19.263 (12.567 to 27.924)	4.634 (3.576 to 5.989)																					
H influenzae type B meningitis	Birth prevalence - [37, 38) wks, [2500, 3000) g	Mortality	Males		3.306 (2.82 to 3.843)	3.194 (2.691 to 3.803)																					
H influenzae type B meningitis	Birth prevalence - [37, 38) wks, [2500, 3000) g	Mortality	Females		2.991 (2.496 to 3.61)	3.242 (2.745 to 3.817)																					
H influenzae type B meningitis	Birth prevalence - [40, 42) wks, [2500, 3000) g	Mortality	Males		3.771 (3.002 to 4.693)	3.175 (2.56 to 3.923)																					
H influenzae type B meningitis	Birth prevalence - [40, 42) wks, [2500, 3000) g	Mortality	Females		3.244 (2.486 to 4.159)	3.228 (2.605 to 3.985)																					
H influenzae type B meningitis	Birth prevalence - [38, 40) wks, [2500, 3000) g	Mortality	Males		2.755 (2.274 to 3.309)	2.944 (2.44 to 3.548)																					
H influenzae type B meningitis	Birth prevalence - [38, 40) wks, [2500, 3000) g	Mortality	Females		2.376 (1.91 to 2.886)	2.938 (2.434 to 3.503)																					
H influenzae type B meningitis	Birth prevalence - [36, 37) wks, [3000, 3500) g	Mortality	Males		3.774 (3.094 to 4.497)	2.466 (2.058 to 2.929)																					
H influenzae type B meningitis	Birth prevalence - [36, 37) wks, [3000, 3500) #	Mortality	Females		3.73	2.715 (2.277 in 3.218)																					
H influenzae type B meningitis	Binh prevalence - [36, 37] wks, [4000, 4500) #	Mortality	Males		6.826 (5.212 to 9.045)	1.77																					
H influenzae type B meningitis	Birth prevalence - [36, 37] wks, [4000,	Mortality	Females		7.269	2.177																					
H influenzae type B meningitis	4300) g Birth prevalence - [36, 37) wks, [3500, 4000) =	Mortality	Males		4544	2.057																					
H influenzae type B meningitis	Birth prevalence - [36, 37] wks, [3500,	Mortality	Females		4.662	2.398																					
H influenzae type B meningitis	Birth prevalence - [37, 38) wks, [3000,	Mortality	Males		2.007	1.888																					
H influenzae type B meningitis	Bith prevalence - [37, 38] wks, [3000,	Mortality	Females		1.925	(1.81318.2.2.24)																					
Hinfluenzae type B menineitis	3500) g Birth prevalence - [37, 38) wks, [4000,	Mortality	Males		(1.582 to 2.328) 3.28	(1.669 to 2.313) 1.335																					
H influenzae type B meningitis	4500) g Birth prevalence - [37, 38) wks, [4000,	Mortality	Females		(2.596 to 4.133) 3.521	(1.171 to 1.532) 1.559																					
Hinfluenzae two B menineitis	4500) g Birth prevalence - [37, 38) wks, [3500,	Mortality	Males		(2.649 to 4.5) 2.128	(1.333 to 1.835) 1.505																					
Hinfluence type B meninging	4000) g Birth prevalence - [37, 38) wks, [3500,	Manulity	Females		(1.833 to 2.466) 2.142	(1.299 to 1.76) 1.661																					
H influence have R maximitie	4000) g Birth prevalence - [40, 42) wks, [3000,	Manufity	Malar		(1.694 to 2.67) 1.436	(1.411 to 1.961) 1.47																					
H influence type B managers	3500) g Birth prevalence - [40, 42) wks, [3000,	Mastality	Employ		(1.245 to 1.65) 1.326	(1.199 to 1.8) 1.465																					
Hinfluence type B meninging	3500) g Birth prevalence - [38, 40) wks, [3000,	Manulity	Malex		(1.069 to 1.614) 1.33	(1.188 to 1.775) 1.559																					
H influence have R maximitie	3500) g Birth prevalence - [38, 40) wks, [3000,	Manufity	Employ		(1.155 to 1.53) 1.224	(1.305 to 1.851) 1.564																					
H influence type B managers	3500) g Birth prevalence - [38, 40) wks, [4000,	Mastality	Malar		(1.0 to 1.492) 1.787	(1.304 to 1.847) 1.175																					
II and the providence of the second second	4500) g Birth prevalence - [38, 40) wks, [4000,	Mariany	To a day		(1.453 to 2.182) 1.877	(1.005 to 1.371) 1.224																					
H influenzae type B meningitis	4500) g Birth prevalence - [38, 40) wks, [3500,	Moriality	Females		(1.467 to 2.388) 1.785	(1.022 to 1.465) 1.173																					
H intractive type is meninging	4000) g Birth prevalence - [38, 40) wks, [3500,	Mortany	Males -		(1.478 to 2.147) 1.892	(1.0 to 1.377) 1.23																					
H influenzae type B meningitis	4000) g Birth merculance - [40, 47) who [3500	Moriality	Females		(1.481 to 2.352)	(1.03 to 1.46)																					
H influenzae type B meningitis	4000) g Birth recentlence - [40, 47) who [3500	Mortality	Males		(1.0 to 1.0)	(1.0 to 1.046)																					
H influenzae type B meningitis	4000) g Birth merculance - [40, 47) why (4000	Mortality	Females		(1.0 to 1.013)	(1.0 to 1.006)																					
H influenzae type B meningitis	4500) g Rith revolutor - [40, 42] W0, [4000]	Mortality	Males		(1.0 to 1.0)	(1.0 to 1.0)																					
H influenzae type B meningitis	4500) g Rith revolutor - [28, 200 min 12000	Mortality	Females		(1.0 to 1.0)	(1.0 to 1.0) 27 224																					
H influenzae type B meningitis	2500) g Risk muselson (20, 20, 1, 2000	Mortality	Males		(83.895 to 158.056)	(21.877 to 34.972)																					
H influenzae type B meningitis	2500) g Risk sussiance - (28, 30) wks, (2000, 2500) g	Mortality	Females		(84.349 to 171.375)	33.383 (26.101 to 43.404)																					
H influenzae type B meningitis	aroon prevasence - (28, 30) wks, (2500, 3000) g	Mortality	Males		(54.687 to 105.047)	(12.653 to 21.188)																					
H influenzae type B meningitis	Bath prevalence - [28, 30] wks, [2500, 3000) g	Mortality	Females		79.193 (54.099 to 112.236)	20.387 (15.089 to 26.434)																					
H influenzae type B meningitis	Bath prevalence - [28, 30) wks, [3000, 3500) g	Mortality	Males		42.199 (29.891 to 57.227)	10.082 (7.777 to 13.056)																					
H influenzae type B meningitis	Bath prevalence - [28, 30) wks, [3000, 3500) g	Mortality	Females		42.551 (28.25 to 63.209)	11.989 (9.084 to 15.544)																					

pendix Table 6a. Relative risks	used by age and sex for each outco	me for all risk factors exc	epi (or ambient an	1201111001202000	and smoxing.									Age	es											
Risk - Outcome	Category / Units	Morbidity / Mortality S	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
H influenzae type B meningitis	Birth prevalence - [30, 32) wks, [2500, 2000) -	Mortality M	iles	58.722	12.115						11															
H influenzae type B meningitis	S000) g Birth prevalence - [30, 32) wks, [2500,	Mortality Fer	ules	(42,41916 78,873) 59,522	(9.3181013.321) 15.364																					
Hindunan tan Remainin	3000) g Birth prevalence - [30, 32) wks, [3000,	Mastelite M		(42.058 to 82.793) 45.67	(11.936 to 19.581) 8.381																					
in annual type is included	3500) g Birth menulence - (30, 32) wks. (3000.	January A		(32.014 to 65.531) 46.104	(6.362 to 10.842) 10.506																					
H influenzae type B menungsias	3500) g Birth menulation (20, 22) with (2600	Mortality Fer	ules	(30.243 to 66.207) 26.324	(8.041 to 13.513)																					
H influenzae type B meningitis	4000) g	Mortality M	iles	(21.558 to 54.813)	(4.293 to 7.349)																					
H influenzae type B meningitis	Birth prevalence - (30, 32) wks, (3500, 4000) g	Mortality Fer	ules	37.931 (22.692 to 61.276)	6.892 (5.175 to 9.058)																					
H influenzae type B meningitis	Birth prevalence - [32, 34) wks, [3000, 3500) g	Mortality M	iles	34.016 (23.515 to 48.37)	6.577 (4.934 to 8.436)																					
H influenzae type B meningitis	Birth prevalence - [32, 34) wks, [3000, 3500) g	Mortality Fer	ules	34.585 (22.909 to 49.754)	8.314 (6.362 to 10.789)																					
H influenzae type B meningitis	Birth prevalence - [32, 34) wks, [3500, 4000) g	Mortality M	ıles	36.248 (23.158 to 54.67)	5.068 (3.761 to 6.741)																					
H influenzae type B meningitis	Birth prevalence - [32, 34) wks, [3500, 4000) p	Mortality Fer	ules	38.098 (23.301 to 59.429)	6.476 (4.666 to 8.689)																					
H influenzae type B meningitis	Birth prevalence - [36, 37) wks, [1000,	Mortality M	des	166.686	57.535																					
H influenzae type B menineitis	Birth prevalence - [36, 37) wks, [1000,	Mortality Fe	ules	169.725	63.564																					
Hindunan tan Remainin	1500) g Birth prevalence - [38, 40) wks, [1000,	Mastelite M		174.066	57.966 (50.068 to 80.705)																					
in annual type is included	1500) g Birth menulence - (38, 40) wks, (1000,	January A		(125.125 to 232.507) 171.557) (44.393 to 73.241) 65.208																					
H influenzae type B menungsias	1500) g Birth menulation (28, 40) with (1800)	Mortality Fer	ules	(121.585 to 237.047	(48.821 to 84.308) 25.206																					
H influenzae type B meningitis	2000) g	Mortality M	iles	(49.547 to 89.055)	(20.365 to 31.168)																					
H influenzae type B meningitis	Birth prevalence - [38, 40) wks, [1500, 2000) g	Mortality Fer	ules	62.19 (45.884 to 83.445)	28.05 (22.625 to 35.139)																					
H influenzae type B meningitis	Birth prevalence - [40, 42) wks, [1500, 2000) g	Mortality M	iles	76.673 (56.177 to 102.468)	25.785 (19.387 to 34.168)																					
H influenzae type B meningitis	Birth prevalence - [40, 42) wks, [1500, 2000) g	Mortality Fer	ules	70.411 (49.221 to 97.952)	29.113 (21.355 to 38.272)																					
Meningococcal infection	Birth prevalence - [0, 24) wks, [0, 500) g	Mortality M	ıles	1564.792 (1056.542 to	618.595 (458.842 to 812.921	0																				
Meningococcal infection	Birth prevalence - [0, 24) wks, [0, 500) g	Mortality Fer	ules	1600.122 (1050.664 to	713.571 (526.178 to 921.018	0																				
Meningococcal infection	Birth prevalence - [0, 24) wks, [500, 1000)	Mortality M	des	1155.815	457.5																					
Menineococcal infection	8 Birth prevalence - [0, 24) wks, [500, 1000)	Mortality Fe	ules	(825.412 to 1506.83) 1169.123	515.406	5)																				
M	8 Birth prevalence - [24, 26) wks, [500, 1000)			(802.003 to 1617.979 955.583	(396.713 to 641.541) 443.357	0																				
Meningococca interior	8 Birth menulance - (24, 26) with (500, 1000)	Montany M	ues.	(723.748 to 1244.265 947 143	i) (363.03 to 534.695) 487 549)																				
Meningococcal infection	8	Mortality Fer	ules	(702.662 to 1237.093	(387.307 to 603.498	8)																				
Meningococcal infection	Bath prevalence - [26, 28) wks, [500, 1000) 8	Mortality M	ıles	497.817 (377.617 to 648.547)	3:30.886) (261.438 to 401.709)	0																				
Meningrococcal infection	Birth prevalence - (26, 28) wks, (500, 1000) 8	Mortality Fer	ules	483.682 (354.946 to 629.517)	344.618) (274.427 to 419.864	ŧ)																				
Meningococcal infection	Birth prevalence - [30, 32) wks, [500, 1000) 8	Mortality M	iles	236.614 (163.821 to 324.502	149.995) (117.866 to 188.368	š)																				
Meningococcal infection	Birth prevalence - [30, 32) wks, [500, 1000) g	Mortality Fer	ules	229.197 (157.606 to 317.194	152.117 (120.779 to 190.583	3)																				
Meningococcal infection	Birth prevalence - [28, 30) wks, [500, 1000)	Mortality M	iles	297.629 (214.953 to 396.586	216.995 (173.321 to 271.466																					
Meningococcal infection	Birth prevalence - [28, 30) wks, [500, 1000)	Mortality Fer	ules	281.056 (198.176 to 386.635	219.884 (174.264 to 272.204																					
Meningococcal infection	Birth prevalence - [26, 28) wks, [1000,	Mortality M	iles	267.91	164.167																					
Menineococcal infection	Birth prevalence - [26, 28) wks, [1000,	Mortality Fe	ules	266.509	174.222	n																				
Maximum linfaction	1500) g Birth prevalence - [34, 36) wks, [1000,	Mastelite M		(197.461 to 346.932 142.056) (137.431 to 217.349 52.86	<i>n</i>																				
Manageroran marcan	1500) g Birth prevalence - [34, 36) wks, [1000,	Markany A		(98.086 to 197.774) 141.899	(42.914 to 64.617) 57.421	ł.																				
Meningococca intection	1500) g Birth menulence - [28, 30) why [1500	storiality Per	11124	(95.864 to 197.656) 127.966	(46.452 to 71.339) 50.018	•																				
Meningococcal infection	2000) g	Mortality M	des	(97.178 to 167.026)	(40.539 to 61.919)																					
Meningococcal infection	2000) g	Mortality Fer	ules	(96.513 to 172.188)	(46.36 to 70.038)																					
Meningococcal infection	Bath prevalence - [28, 30) wks, [1000, 1500) g	Mortality M	ıles	158.563 (120.99 to 204.947)	103.32 (83.486 to 127.144))																				
Meningococcal infection	Birth prevalence - [28, 30) wks, [1000, 1500) g	Mortality Fer	ules	153.905 (112.327 to 200.786)	107.529 (86.954 to 131.78)																					
Meningococcal infection	Birth prevalence - [32, 34) wks, [1000, 1500) g	Mortality M	iles	117.142 (81.354 to 161.101)	53.185 (43.049 to 66.274)																					
Meningococcal infection	Birth prevalence - [32, 34) wks, [1000, 1500) g	Mortality Fer	ules	115.171 (79.363 to 159.206)	56.034 (45.982 to 68.36)																					
Meningococcal infection	Birth prevalence - [30, 32) wks, [1000, 1500) g	Mortality M	iles	119.308 (87.769 to 160.885)	67.163 (54.863 to 82.638)																					
Meningococcal infection	Birth prevalence - [30, 32) wks, [1000, 1500) g	Mortality Fer	ules	115.448 (84.272 to 156.425)	69.14 (55.873 in 85.012)																					
Meningococcal infection	Birth prevalence - [37, 38) wks, [1500, 2000) -	Mortality M	iles	62.972	24.148																					
Meningpeoceal infection	Birth prevalence - [37, 38) wks, [1500,	Mortality Fer	ules	(%4.1.7910.83.484) 59.988	26.719																					
-	2000) g Birth prevalence - [36, 37) wks, [1500,	Mandity M	der.	(43.974 to 79.053) 60.218	(21.746 to 32.816) 23.031																					
mangototta antion	2000) g Birth menulence - (36, 37) wks, (1500,	January A		(43.669 to 82.48) 58.527	(18.793 to 28.483) 25.143																					
Menungococcal infection	2000) g Birth menalence - (30, 32) wite (2000	Mortality Fer	ules	(42.172 to 80.557) 67.971	(20.331 to 30.566) 18.03																					
Meningrococcal infection	2500) g	Mortality M	iles	(50.354 to 88.935)	(14.621 to 22.103)																					
Meningococcal infection	north prevalence - [30, 32) wks, [2000, 2500) g	Mortality Fer	ules	69.383 (49.108 to 94.583)	22.069 (17.836 to 27.163)																					
Meningococcal infection	Batth prevalence - [30, 32) wks, [1500, 2000) g	Mortality M	iles	77.369 (59.702 to 99.232)	31.079 (25.786 to 36.724)																					
Meningococcal infection	Birth prevalence - [30, 32) wks, [1500, 2000) g	Mortality Fer	ules	76.134 (56.885 to 100.996)	34.756 (28.764 to 41.849)																					
Meningococcal infection	Birth prevalence - [34, 36) wks, [1500, 2000) g	Mortality M	iles	55.555 (39.553 to 75.104)	21.346 (17.677 to 26.143)																					
Meningococcal infection	Birth prevalence - [34, 36) wks, [1500, 2000) g	Mortality Fer	ules	54.335 (38.617 to 75.24)	23.046 (18.743 to 28.287)																					
Meningococcal infection	- Birth prevalence - [32, 34) wks, [1500, 2000) #	Mortality M	ıles	57.155 (42.484 to 73.651)	23.114 (19.028 in 27.915)																					
Meningococcal infection	Birth prevalence - [32, 34) wks, [1500,	Mortality Fe	ules	56.101	25.149																					
Meningrococcal infectives	Bith prevalence - [32, 34) wks, [2000,	Mortality M	iles	(39.794 to 76.295) 37.444	(20.015 to 30.388) 12.233																					
Maximum and infesti	2500) g Birth prevalence - [32, 34) wks, [2000,	Mandin T	-	(29.026 to 48.227) 36.874	(10.252 to 14.477) 14.384	1																				
Meningscocca intection	2500) g Birth prevalence - [40, 42) wks, [2000	Mortany Fe		(26.658 to 49.653) 18.092	(12.095 to 17.03) 9.23																					
meningscoccal infection	2500) g	suoraiity M	ana.	(13.292 to 23.719)	(7.037 to 11.454)	1																				

															Aş	ges											
Birk Outrons	Catagory / Units	Manhidita / Mantali	ia. 6	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Meningococcal infection	Birth prevalence - [40, 42) wks, [2000,	Mortality	Females		15.574	9.975																					
Maximum and infection	2500) g Birth prevalence - [38, 40) wks, [2000,	Mantality	Malar		(11.516 to 20.778) 13.104	(7.82 to 12.46) 8.198																					
	2500) g Birth menulence - (38, 40) wks, (2000,				(9.829 to 16.99) 11.308	(6.786 to 9.959) 8.577																					
Meningococcal infection	2500) g Risk musikana (22, 24) ada (2500)	Mortality	Females		(8.389 to 14.38)	(7.04 to 10.449)																					
Meningococcal infection	3000) g	Mortality	Males		(24.393 to 43.503)	(6.822 to 10.431)																					
Meningococcal infection	Birth prevalence - [32, 34) wks, [2500, 3000) g	Mortality	Females		32.812 (23.439 to 45.567)	10.398 (8.227 to 13.042)																					
Meningococcal infection	Birth prevalence - [34, 36) wks, [2000, 2500) g	Mortality	Males		21.925 (16.305 to 29.433)	9.367 (7.859 to 11.112)																					
Meningococcal infection	Birth prevalence - [34, 36) wks, [2000, 2500) g	Mortality	Females		21.297 (15.657 to 28.761)	10.295 (8.548 to 12.273)																					
Meningococcal infection	Birth prevalence - [37, 38) wks, [2000, 2500) g	Mortality	Males		13.0 (10.102 to 16.456)	8.096 (6.724 to 9.676)																					
Meningococcal infection	Birth prevalence - [37, 38) wks, [2000, 2500) #	Mortality	Females		11.563 (8.805 to 15.11)	8.467 (6.994 to 10.347)																					
Meningococcal infection	Birth prevalence - [36, 37] wks, [2000,	Mortality	Males		14.401	8.221																					
Meximum-corceal infections	2500) g Birth prevalence - [36, 37) wks, [2000,	Martality	Females		(10.78910 18.054)	8.654																					
Maximum distantion	2500) g Birth prevalence - [34, 36) wks, [2500,	Mantalian	Malar		(9.817 to 17.942) 13.419	(7.215 to 10.369) 5.562																					
mangeot and and	3000) g Birth menulance - [34, 36) wire [2500	January	ALL A		(10.387 to 16.819)	(4.646 to 6.696) 6 395																					
Meningococcal infection	3000) g	Mortality	Females		(9.666 to 17.689)	(5.292 to 7.606)																					
Meningococcal infection	4500) g	Mortality	Males		(14.708 to 35.098)	(2.245 to 3.716)																					
Meningococcal infection	Birth prevalence - [34, 36) wks, [4000, 4500) g	Mortality	Females		25.038 (15.763 to 37.255)	3.778 (2.855 to 4.925)																					
Meningococcal infection	Birth prevalence - [34, 36) wks, [3000, 3500) g	Mortality	Males		14.006 (10.222 to 18.478)	4.322 (3.449 to 5.338)																					
Meningococcal infection	Birth prevalence - [34, 36) wks, [3000, 3500) g	Mortality	Females		14.375 (10.269 to 20.114)	5.265 (4.145 to 6.564)																					
Meningococcal infection	Birth prevalence - [36, 37) wks, [2500, 3000) a	Mortality	Males		4.874 (4.014 to 5.713)	3.699 (3.134 to 4.374)																					
Meningococcal infection	Birth prevalence - [36, 37] wks, [2500,	Mortality	Females		4.609	3.898																					
Meninger-coreal infection	Birth prevalence - [34, 36) wks, [3500,	Martality	Malex		(3.731 83 3.81) 18.024	(3.235 65 4.67) 3.657																					
N	4000) g Birth prevalence - [34, 36) wks, [3500,				(12.279 to 25.547) 19.263	(2.838 to 4.675) 4.634																					
Menngococcai intection	4000) g Birth menulance - [37-38) wise [2500	storcarty	Pemales		(12.567 to 27.924) 3 306	(3.576 to 5.989) 3.194																					
Meningococcal infection	3000) g	Mortality	Males		(2.82 to 3.843)	(2.691 to 3.803)																					
Meningococcal infection	Both prevalence - [37, 38) wks, [2500, 3000) g	Mortality	Females		(2.496 to 3.61)	3.242 (2.745 to 3.817)																					
Meningococcal infection	Birth prevalence - [40, 42) wks, [2500, 3000) g	Mortality	Males		3.771 (3.002 to 4.693)	3.175 (2.56 to 3.923)																					
Meningococcal infection	Birth prevalence - [40, 42) wks, [2500, 3000) g	Mortality	Females		3.244 (2.486 to 4.159)	3.228 (2.605 to 3.985)																					
Meningococcal infection	Birth prevalence - [38, 40) wks, [2500, 3000) g	Mortality	Males		2.755 (2.274 to 3.309)	2.944 (2.44 to 3.548)																					
Meningococcal infection	Birth prevalence - [38, 40) wks, [2500, 3000) a	Mortality	Females		2.376 (1.91 to 2.886)	2.938 (2.434 to 3.503)																					
Meningococcal infection	Birth prevalence - [36, 37] wks, [3000, 25000 -	Mortality	Males		3.774	2.466																					
Meningococcal infection	Birth prevalence - [36, 37) wks, [3000,	Mortality	Females		3.73	2.715																					
Maximum distantion	3500) g Birth prevalence - [36, 37) wks, [4000,	Mantalian	Malar		(2.981 to 4.646) 6.826	(2.277 to 3.218) 1.77																					
Managerouta and the	4500) g Birth prevalence - [36, 37) wks, [4000,	Mariany	Trank.		(5.212 to 9.045) 7.269	(1.491 to 2.082) 2.177																					
Menngococcai intection	4500) g Birth menulance - [36-37) wise [3500	storcarty	Pemales		(5.144 to 9.821) 4 544	(1.786 to 2.607) 2.057																					
Meningococcal infection	4000) g	Mortality	Males		(3.64 to 5.622)	(1.74 to 2.44)																					
Meningococcal infection	4000) g	Mortality	Females		4.002 (3.577 to 6.014)	(1.98 to 2.864)																					
Meningococcal infection	Birth prevalence - [37, 38) wks, [3000, 3500) g	Mortality	Males		2.007 (1.759 to 2.293)	1.888 (1.613 to 2.224)																					
Meningococcal infection	Birth prevalence - [37, 38) wks, [3000, 3500) g	Mortality	Females		1.925 (1.582 to 2.328)	1.972 (1.669 to 2.313)																					
Meningococcal infection	Birth prevalence - [37, 38) wks, [4000, 4500) g	Mortality	Males		3.28 (2.596 to 4.133)	1.335 (1.171 to 1.532)																					
Meningococcal infection	Birth prevalence - [37, 38) wks, [4000, 4500) g	Mortality	Females		3.521 (2.649 to 4.5)	1.559 (1.333 to 1.835)																					
Meningococcal infection	Birth prevalence - [37, 38) wks, [3500, 4000) a	Mortality	Males		2.128 (1.833 to 2.466)	1.505 (1.299 to 1.76)																					
Meningococcal infection	Birth prevalence - [37, 38) wks, [3500,	Mortality	Females		2.142	1.661																					
Meningococcal infections	-000) g Birth prevalence - [40, 42) wks, [3000,	Mortality	Maler		(1.094 to 2.67) 1.436	(1.411 to 1.961) 1.47																					
Maximum and infection	3500) g Birth prevalence - [40, 42) wks, [3000,	Mantalian	Employ		(1.245 to 1.65) 1.326	(1.199 to 1.8) 1.465																					
Maximum 12.5	3500) g Birth prevalence - [38, 40) wks, [3000.	M	A COMMON		(1.069 to 1.614) 1.33	(1.188 to 1.775) 1.559																					
encompositional internation	3500) g Birth mevalence - [38, 40) wise [3000	somany	ALLICK		(1.155 to 1.53) 1.224	(1.305 to 1.851) 1.564																					
Meningococcal infection	3500) g Rink muselsma (28, 40) who (2000)	Mortality	Females		(1.0 to 1.492)	(1.304 to 1.847)																					
Meningococcal infection	4500) g	Mortality	Males		(1.453 to 2.182)	(1.005 to 1.371)																					
Meningococcal infection	Birth prevalence - [38, 40) wks, [4000, 4500) g	Mortality	Females		1.877 (1.467 to 2.388)	1.224 (1.022 to 1.465)																					
Meningococcal infection	Birth prevalence - [38, 40) wks, [3500, 4000) g	Mortality	Males		1.785 (1.478 to 2.147)	1.173 (1.0 to 1.377)																					
Meningococcal infection	Birth prevalence - [38, 40) wks, [3500, 4000) g	Mortality	Females		1.892 (1.481 to 2.352)	1.23 (1.03 to 1.46)																					
Meningococcal infection	Birth prevalence - [40, 42) wks, [3500, 4000) g	Mortality	Males		1.0 (1.0 to 1.0)	1.003 (1.0 to 1.046)																					
Meningococcal infection	Birth prevalence - [40, 42) wks, [3500, 4000) g	Mortality	Females		1.002 (1.0 to 1.013)	1.001 (1.0 to 1.006)																					
Meningococcal infection	Birth prevalence - [40, 42) wks, [4000, 4500) =	Mortality	Males		1.0	1.0																					
Meningococcal infection	Birth prevalence - [40, 42) wks, [4000,	Mortality	Females		1.0	1.0																					
Meningspeeced infection	4500) g Birth prevalence - [28, 30) wks, [2000,	Montality	Malex		(1.0 to 1.0) 117.172	(1.0 to 1.0) 27.726																					
Marine and Aria	2500) g Birth prevalence - [28, 30) wks, [2000.				(83.895 to 158.056) 121.682	(21.877 to 34.972) 33.983																					
Meningococcal infection	2500) g Birth mevalence - (28, 30) wks, (2500	Mortality	Females		(84.349 to 171.375) 77.948	(26.101 to 43.404) 16.608																					
Meningococcal infection	3000) g Rink muslame (20. 200 b. 2000)	Mortality	Males		(54.687 to 105.047)	(12.653 to 21.188)																					
Meningococcal infection	amn prevaaence - (28, 30) wiss, (2500, 3000) g	Mortality	Females		(54.099 to 112.236)	20.587 (15.089 to 26.434)																					
Meningococcal infection	Birth prevalence - [28, 30) wks, [3000, 3500) g	Mortality	Males		42.199 (29.891 to 57.227)	10.082 (7.777 to 13.056)																					
Meningococcal infection	Birth prevalence - [28, 30) wks, [3000, 3500) g	Mortality	Females		42.551 (28.25 to 63.209)	11.989 (9.084 to 15.544)																					

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Rick - Ontcome	Catanory / Units	Morbidity / Mortality	Sar	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Meningococcal infection	Birth prevalence - [30, 32) wks, [2500,	Mortality	Males		58.722	12.115																					
Meninerococcal infection	Solid) g Birth prevalence - [30, 32) wks, [2500,	Mortality	Females		(42.41930 78.873) 59.522	(9.5181015.521)																					
Maximum and infection	3000) g Birth prevalence - [30, 32) wks, [3000,	Mantality	Malar		(42.058 to 82.793) 45.67	(11.936 to 19.581) 8.381																					
Mangood and and	3500) g Birth mevalence - (30, 32) wks, (3000,	January			(32.014 to 65.531) 46.104	(6.362 to 10.842) 10.506																					
Meningscoccal infection	3500) g Rick merchanis (20, 22) who (2500	Mortality	Females		(30.243 to 66.207) 26.234	(8.041 to 13.513)																					
Meningococcal infection	4000) g	Mortality	Males		(21.558 to 54.813)	(4.293 to 7.349)																					
Meningococcal infection	Birth prevalence - [30, 32] wks, [3500, 4000) g	Mortality	Females		37.931 (22.692 to 61.276)	6.892 (5.175 to 9.058)																					
Meningococcal infection	Birth prevalence - [32, 34) wks, [3000, 3500) g	Mortality	Males		34.016 (23.515 to 48.37)	6.577 (4.934 to 8.436)																					
Meningococcal infection	Birth prevalence - [32, 34) wks, [3000, 3500) g	Mortality	Females		34.585 (22.909 to 49.754)	8.314 (6.362 to 10.789)																					
Meningococcal infection	Birth prevalence - [32, 34) wks, [3500, 4000) g	Mortality	Males		36.248 (23.158 to 54.67)	5.068 (3.761 to 6.741)																					
Meningococcal infection	Birth prevalence - [32, 34) wks, [3500, 4000) g	Mortality	Females		38.098 (23.301 to 59.429)	6.476 (4.666 to 8.689)																					
Meningococcal infection	Birth prevalence - [36, 37) wks, [1000, 1500) #	Mortality	Males		166.686 (118.487 to 222.581)	57.535 (45.999 to 71.747)																					
Meningococcal infection	Birth prevalence - [36, 37) wks, [1000, 1500) #	Mortality	Females		169.725 (119.017 to 229.008)	63.564 (50.068 to 80.703)																					
Meningococcal infection	Birth prevalence - [38, 40) wks, [1000,	Mortality	Males		174.066	57.966																					
Meninerococcal infection	1500) g Birth prevalence - [38, 40) wks, [1000,	Mortality	Females		171.557	(44.3931873.241) 65.208																					
	1500) g Birth prevalence - [38, 40) wks, [1500,	M - P-			(121.585 to 237.047) 67.302	(48.821 to 84.308) 25.206																					
Meningscoccai intection	2000) g Birth mevalence - (38, 40) wks, (1500,	storcarry	Males		(49.547 to 89.055) 62.19	(20.365 to 31.168) 28.05																					
Meningscoccal infection	2000) g Rick merchanic (40, 42) who (1500)	Mortality	Females		(45.884 to 83.445) 76.673	(22.625 to 35.139)																					
Meningococcal infection	2000) g	Mortality	Males		(56.177 to 102.468)	(19.387 to 34.168)																					
Meningococcal infection	Birth prevalence - [40, 42) wks, [1500, 2000) g	Mortality	Females		70.411 (49.221 to 97.952) 1564 792	29.113 (21.355 to 38.272)																					
Other meningitis	Birth prevalence - [0, 24) wks, [0, 500) g	Mortality	Males		(1056.542 to 2116.062)	618.595 (458.842 to 812.921)																					
Other meningitis	Birth prevalence - [0, 24) wks, [0, 500) g	Mortality	Females		(1050.664 to 2011 927)	713.571 (526.178 to 921.018)																					
Other meningitis	Birth prevalence - [0, 24) wks, [500, 1000) g	Mortality	Males		1155.815 (825.412 to 1506.837)	457.5 (352.552 to 573.483)																					
Other meningitis	Birth prevalence - [0, 24) wks, [500, 1000) 8	Mortality	Females		1169.123 (802.003 to 1617.979)	515.406 (396.713 to 641.541)																					
Other meningitis	Birth prevalence - [24, 26) wks, [500, 1000)	Mortality	Males		955.583 (723.748 to 1244.265)	443.357 (363.03 to 534.695)																					
Other meningitis	Birth prevalence - [24, 26) wks, [500, 1000)	Mortality	Females		947.143	487.549																					
Other menineitis	8 Birth prevalence - [26, 28) wks, [500, 1000)	Mortality	Males		497.817	330.886																					
Other menimuitia	8 Birth prevalence - [26, 28] wks, [500, 1000)	Martility	Females		(377.617 to 648.547) 483.682	(261.438 to 401.709) 344.618																					
	8 Birth prevalence - [30, 32) wks, [500, 1000)	M			(354.946 to 629.517) 236.614	(274.427 to 419.864) 149.995																					
Cial analysis	8 Birth mevalence - (30, 32) wks, (500, 1000)	January .			(163.821 to 324.502) 229.197	(117.866 to 188.368) 152.117																					
Other menungitis	8 Richmanham 128 20 min 1600 1000	Mortality	Females		(157.606 to 317.194) 207.620	(120.779 to 190.583) 216 005																					
Other meningitis	8	Mortality	Males		(214.953 to 396.586)	(173.321 to 271.466)																					
Other meningitis	Birth prevalence - [28, 30) wks, [500, 1000) 8	Mortality	Females		281.056 (198.176 to 386.635)	219.884 (174.264 to 272.704)																					
Other meningitis	Birth prevalence - [26, 28) wks, [1000, 1500) g	Mortality	Males		267.91 (210.177 to 332.92)	164.167 (132.898 to 200.569)																					
Other meningitis	Birth prevalence - [26, 28) wks, [1000, 1500) g	Mortality	Females		266.509 (197.461 to 346.932)	174.222 (137.431 to 217.349)																					
Other meningitis	Birth prevalence - [34, 36) wks, [1000, 1500) g	Mortality	Males		142.056 (98.086 to 197.774)	52.86 (42.914 to 64.617)																					
Other meningitis	Birth prevalence - [34, 36) wks, [1000, 1500) g	Mortality	Females		141.899 (95.864 to 197.656)	57.421 (46.452 to 71.339)																					
Other meningitis	Birth prevalence - [28, 30) wks, [1500, 2000) g	Mortality	Males		127.966 (97.178 to 167.026)	50.018 (40.539 to 61.919)																					
Other meningitis	Birth prevalence - [28, 30) wks, [1500, 2000) g	Mortality	Females		130.924 (96.513 to 172.188)	57.275 (46.36 to 70.038)																					
Other meningitis	Birth prevalence - [28, 30) wks, [1000, 1500) #	Mortality	Males		158.563 (120.99 to 204.947)	103.32 (83.486 to 127.144)																					
Other meningitis	Birth prevalence - [28, 30) wks, [1000,	Mortality	Females		153.905	107.529																					
Other meningitis	Birth prevalence - [32, 34) wks, [1000,	Mortality	Males		117.142	53.185																					
Other menimuitia	1500) g Birth prevalence - [32, 34) wks, [1000,	Martility	Females		(81.354 to 161.101) 115.171	(43.049 to 66.274) 56.034																					
Other merinaitia	1 500) g Birth prevalence - [30, 32) wks, [1000,	Montalia	Male		(19.365 to 159.206) 119.308	(45.982 to 68.36) 67.163																					
Other marinetic	1500) g Birth prevalence - [30, 32) wks, [1000,	Marilin	Females		(87.769 to 160.885) 115.448	(54.863 to 82.638) 69.14																					
Other meanights	1500) g Birth menalence - [37 38) wire [1500	storcarry	remates		(84.272 to 156.425) 62.972	(55.873 to 85.012) 24.148																					
Other meningitis	2000) g Birth menalence - [37 38) wise [1500	Mortality	Males		(46.159 to 83.484) 59.988	(20.066 to 29.406) 26.719																					
Other meningitis	2000) g	Mortality	Females		(43.974 to 79.053)	(21.746 to 32.816)																					
Other meningitis	Birth prevalence - [36, 37] wks, [1500, 2000) g	Mortality	Males		60.218 (43.669 to 82.48)	25.051 (18.793 to 28.483)																					
Other meningitis	Buth prevalence - [36, 37) wks, [1500, 2000) g	Mortality	Females		58.527 (42.172 to 80.557)	25.143 (20.331 to 30.566)																					
Other meningitis	Birth prevalence - [30, 32) wks, [2000, 2500) g	Mortality	Males		67.971 (50.354 to 88.935)	18.03 (14.621 to 22.103)																					
Other meningitis	Birth prevalence - [30, 32) wks, [2000, 2500) g	Mortality	Females		69.383 (49.108 to 94.583)	22.069 (17.836 to 27.163)																					
Other meningitis	Birth prevalence - [30, 32) wks, [1500, 2000) g	Mortality	Males		77.369 (59.702 to 99.232)	31.079 (25.786 to 36.724)																					
Other meningitis	Birth prevalence - [30, 32) wks, [1500, 2000) g	Mortality	Females		76.134 (56.885 to 100.996)	34.756 (28.764 to 41.849)																					
Other meningitis	Birth prevalence - [34, 36) wks, [1500, 2000) #	Mortality	Males		55.555 (39.553 to 75.104)	21.346 (17.677 to 26.143)																					
Other meningitis	Birth prevalence - [34, 36) wks, [1500, 2000) p	Mortality	Females		54.335 (38.617 to 75.24)	23.046 (18.743 to 78.787)																					
Other meningitis	Birth prevalence - [32, 34) wks, [1500, 2000) #	Mortality	Males		57.155 (42.484 tr 73.661)	23.114																					
Other meningitis	Birth prevalence - [32, 34) wks, [1500,	Mortality	Females		56.101	25.149																					
Other menimities	2000) g Birth prevalence - [32, 34) wks, [2000,	Manulity	Malex		(39.794 to 76.295) 37.444	(20.615 to 30.388) 12.233																					
Other marinetic	2500) g Birth prevalence - [32, 34) wks, [2000,	Marilin	Females		(29.026 to 48.227) 36.874	(10.252 to 14.477) 14.384																					
our nenigus	2500) g Birth prevalence - [40, 42) wks, [2000.	Mortany	- emailes		(26.658 to 49.653) 18.092	(12.095 to 17.03) 9.23																					
Other meninguis	2500) g	stortany	ALLIES	I	(13.292 to 23.719)	(7.037 to 11.454)																					

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Risk - Outcome	Category / Units	Morbidity / Mortali	tv Sex	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Other meningitis	Birth prevalence - [40, 42) wks, [2000, 2500) e	Mortality	Females		15.574 (11 516 to 20 778)	9.975 (7.82 to 12.46)																					
Other meningitis	Birth prevalence - [38, 40) wks, [2000,	Mortality	Males		13.104	8.198																					
Other menineitis	2300) g Birth prevalence - [38, 40) wks, [2000,	Mandin	Females		(9.8291818.99)	8.577																					
Other environitio	2500) g Birth prevalence - [32, 34) wks, [2500,	Massilia	Malar		(8.389 to 14.38) 33.063	(7.04 to 10.449) 8.441																					
Contractinguit	3000) g Birth merculance , [32, 34) wise [2500	Juniary	Auto		(24.393 to 43.503) 32.812	(6.822 to 10.431) 10.398																					
Other meningitis	3000) g	Mortality	Females		(23.439 to 45.567)	(8.227 to 13.042)																					
Other meningitis	2500) g	Mortality	Males		(16.305 to 29.433)	(7.859 to 11.112)																					
Other meningitis	Barth prevalence - [34, 36) wks, [2000, 2500) g	Mortality	Females		(15.657 to 28.761)	(8.548 to 12.273)																					
Other meningitis	Birth prevalence - [37, 38) wks, [2000, 2500) g	Mortality	Males		13.0 (10.102 to 16.456)	8.096 (6.724 to 9.676)																					
Other meningitis	Birth prevalence - [37, 38) wks, [2000, 2500) g	Mortality	Females		11.563 (8.805 to 15.11)	8.467 (6.994 to 10.342)																					
Other meningitis	Birth prevalence - [36, 37) wks, [2000, 2500) g	Mortality	Males		14.401 (10.789 to 18.654)	8.221 (6.917 to 9.923)																					
Other meningitis	Birth prevalence - [36, 37) wks, [2000, 2500) #	Mortality	Females		13.513 (9.817 to 17.942)	8.654 (7.215 to 10.369)																					
Other meningitis	Birth prevalence - [34, 36) wks, [2500, 3000) e	Montality	Males		13.419 (10.387 to 16.819)	5.562 (4.646 to 6.696)																					
Other meningitis	Birth prevalence - [34, 36) wks, [2500, 2000) -	Mortality	Females		13.266	6.395																					
Other menineitis	Birth prevalence - [34, 36) wks, [4000,	Mortality	Males		23.096	2.895																					
Other environitio	4500) g Birth prevalence - [34, 36) wks, [4000,	Mastality	Emales		(14.708 to 35.098) 25.038	(2.245 to 3.716) 3.778																					
Contractinguit	4500) g Birth prevalence - (34, 36) wks. (3000,				(15.763 to 37.255) 14.006	(2.855 to 4.925) 4.322																					
Other meningins	3500) g Birth merculance - [34, 36) wise [3000	sommery	MBO		(10.222 to 18.478) 14.375	(3.449 to 5.338) 5.265																					
Other meningitis	3500) g	Mortality	Females		(10.269 to 20.114)	(4.145 to 6.564)																					
Other meningitis	3000) g	Mortality	Males		4.874 (4.014 to 5.713)	(3.134 to 4.374)																					
Other meningitis	Barth prevalence - [36, 37] wks, [2500, 3000) g	Mortality	Females		4.609 (3.731 to 5.61)	3.898 (3.235 to 4.67)																					
Other meningitis	Birth prevalence - [34, 36) wks, [3500, 4000) g	Mortality	Males		18.024 (12.279 to 25.547)	3.657 (2.838 to 4.675)																					
Other meningitis	Birth prevalence - [34, 36) wks, [3500, 4000) g	Mortality	Females		19.263 (12.567 to 27.924)	4.634 (3.576 to 5.989)																					
Other meningitis	Birth prevalence - [37, 38) wks, [2500, 3000) g	Mortality	Males		3.306 (2.82 to 3.843)	3.194 (2.691 to 3.803)																					
Other meningitis	Birth prevalence - [37, 38) wks, [2500, 3000) g	Mortality	Females		2.991 (2.496 to 3.61)	3.242 (2.745 to 3.817)																					
Other meningitis	Birth prevalence - [40, 42) wks, [2500, 3000 a	Montality	Males		3.771 (3.002 to 4.693)	3.175 (2.56 to 3.923)																					
Other meningitis	Birth prevalence - [40, 42) wks, [2500, 2000) -	Mortality	Females		3.244	3.228																					
Other meningitis	Birth prevalence - [38, 40) wks, [2500,	Mortality	Males		2.755	2.944																					
Other menineitis	3000) g Birth prevalence - [38, 40) wks, [2500,	Mortality	Females		(2.274 to 3.309) 2.376	(2.44 to 3.548) 2.938																					
Other environitio	3000) g Birth prevalence - [36, 37) wks, [3000,	Mastelite	Maler		(1.91 to 2.886) 3.774	(2.434 to 3.503) 2.466																					
Ola mingh	3500) g Birth prevalence - [36, 37) wks, [3000,	Martin	T and		(3.094 to 4.497) 3.73	(2.058 to 2.929) 2.715																					
Other meningins	3500) g Birth merculance - [36-37) wise [4000	sommery	remates		(2.981 to 4.646) 6.826	(2.277 to 3.218)																					
Other meningitis	4500) g Rishemmelana (26, 27) ada (4000	Mortality	Males		(5.212 to 9.045) 7.260	(1.491 to 2.082)																					
Other meningitis	4500) g	Mortality	Females		(5.144 to 9.821)	(1.786 to 2.607)																					
Other meningitis	4000) g	Mortality	Males		4.544 (3.64 to 5.622)	(1.74 to 2.44)																					
Other meningitis	Barth prevalence - [36, 37] wks, [3500, 4000) g	Mortality	Females		4.662 (3.577 to 6.014)	2.398 (1.98 to 2.864)																					
Other meningitis	Birth prevalence - [37, 38) wks, [3000, 3500) g	Mortality	Males		2.007 (1.759 to 2.293)	1.888 (1.613 to 2.224)																					
Other meningitis	Birth prevalence - [37, 38) wks, [3000, 3500) g	Mortality	Females		1.925 (1.582 to 2.328)	1.972 (1.669 to 2.313)																					
Other meningitis	Birth prevalence - [37, 38) wks, [4000, 4500) g	Mortality	Males		3.28 (2.596 to 4.133)	1.335 (1.171 to 1.532)																					
Other meningitis	Birth prevalence - [37, 38) wks, [4000, 4500) g	Mortality	Females		3.521 (2.649 to 4.5)	1.559 (1.333 to 1.835)																					
Other meningitis	Birth prevalence - [37, 38) wks, [3500, 4000) g	Mortality	Males		2.128 (1.833 to 2.466)	1.505 (1.299 to 1.76)																					
Other meningitis	Birth prevalence - [37, 38) wks, [3500, 4000) #	Montality	Females		2.142 (1.694 to 2.67)	1.661 (1.411 to 1.961)																					
Other meningitis	Birth prevalence - [40, 42) wks, [3000, 3500) n	Mortality	Males		1.436 (1.245 to 1.65)	1.47 (1.199 m 1.8)																					
Other meningitis	Birth prevalence - [40, 42] wks, [3000, 25000 -	Mortality	Females		1.326	1.465																					
Other meningitis	Birth prevalence - [38, 40) wks, [3000, 25000 -	Mortality	Males		1.33	1.559																					
Other menineitis	3300) g Birth prevalence - [38, 40) wks, [3000,	Mortality	Females		(1.135 to 1.53) 1.224	(1.305 to 1.851) 1.564																					
Other menineitis	3500) g Birth prevalence - [38, 40) wks, [4000,	Mortality	Male		(1.0 to 1.492) 1.787	(1.304 to 1.847) 1.175																					
Olive a straights	4500) g Birth prevalence - [38, 40) wks, [4000.				(1.453 to 2.182) 1.877	(1.005 to 1.371) 1.224																					
Oner mennights	4500) g Birth prevalence - [38, 40) wks, [3500,	Mortany	remates		(1.467 to 2.388) 1.785	(1.022 to 1.465) 1.173																					
Other meningitis	4000) g Rightmanlana (28, 40) min (2600	Mortality	Males		(1.478 to 2.147)	(1.0 to 1.377)																					
Other meningitis	4000) g Rich mandama (40, 42) adv (2000)	Mortality	Females		(1.481 to 2.352)	(1.03 to 1.46)																					
Other meningitis	ann prevaence - [40, 42] wks, [3500, 4000) g	Mortality	Males		1.0 (1.0 to 1.0)	(1.0 to 1.046)																					
Other meningitis	Bath prevalence - [40, 42] wks, [3500, 4000) g	Mortality	Females		1.002 (1.0 to 1.013)	1.001 (1.0 to 1.006)																					
Other meningitis	Birth prevalence - [40, 42) wks, [4000, 4500) g	Mortality	Males		1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																					
Other meningitis	Birth prevalence - [40, 42) wks, [4000, 4500) g	Mortality	Females		1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																					
Other meningitis	Birth prevalence - [28, 30) wks, [2000, 2500) g	Mortality	Males		117.172 (83.895 to 158.056)	27.726 (21.877 to 34.972)																					
Other meningitis	Birth prevalence - [28, 30) wks, [2000, 2500) g	Mortality	Females		121.682 (84.349 to 171.375)	33.983 (26.101 to 43.404)																					
Other meningitis	Birth prevalence - [28, 30) wks, [2500, 3000) g	Mortality	Males		77.948 (54.687 to 105.047)	16.608 (12.653 to 21.188)																					
Other meningitis	Birth prevalence - [28, 30) wks, [2500, 3000) g	Mortality	Females		79.193 (54.099 to 112.236)	20.387 (15.089 to 26.434)																					
Other meningitis	Birth prevalence - [28, 30) wks, [3000, 3500) =	Mortality	Males		42.199 (79.891 to 57.227)	10.082																					
Other meningitis	Birth prevalence - [28, 30] wks, [3000,	Mortality	Females		42.551	11.989																					
	2009 B			1	(48.47.40.03.209)	(2.009 10 13.394)	1																				

ppendix rable oa. Kelative i	isks used by age and sex for each outcon	ne tor an risk facto	a stexacion	0.0000000000000000000000000000000000000	0101000 8100000	nu snioking.									А	iges											
Pirk - Outcome	Catanory / Unite	Morbidity / Mortali	ity Say	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Other meningitis	Bith prevalence - [30, 32) wks, [2500, 2000 -	Mortality	Males		58.722	12.115						L															
Other meningitis	S000) g Birth prevalence - [30, 32) wks, [2500,	Mortality	Females		(42.41910 78.873) 59.522	(9.5181015.321)																					
Other menineitis	3000) g Birth prevalence - [30, 32) wks, [3000,	Martality	Malex		(42.058 to 82.793) 45.67	(11.936 to 19.581) 8.381																					
Other maniputio	3500) g Birth prevalence - [30, 32) wks, [3000,	Manufita	Emple		(32.014 to 65.531) 46.104	(6.362 to 10.842) 10.506																					
Charl Including	3500) g Birth mevalence - (30, 32) wks. (3500,	January			(30.243 to 66.207) 36.334	(8.041 to 13.513) 5.698																					
Other meningitis	4000) g Birth menalewer - (30, 32) wise (3500	Moriality	Males		(21.558 to 54.813) 37.931	(4.293 to 7.349) 6.892																					
Other meningitis	4000) g	Mortality	Females		(22.692 to 61.276)	(5.175 to 9.058)																					
Other meningitis	3500) g	Mortality	Males		(23.515 to 48.37)	(4.934 to 8.436)																					
Other meningitis	Birth prevalence - [32, 34) wks, [3000, 3500) g	Mortality	Females		34.585 (22.909 to 49.754)	8.314 (6.362 to 10.789)																					
Other meningitis	Birth prevalence - [32, 34) wks, [3500, 4000) g	Mortality	Males		36.248 (23.158 to 54.67)	5.068 (3.761 to 6.741)																					
Other meningitis	Birth prevalence - [32, 34) wks, [3500, 4000) g	Mortality	Females		38.098 (23.301 to 59.429)	6.476 (4.666 to 8.689)																					
Other meningitis	Birth prevalence - [36, 37) wks, [1000, 1500) g	Mortality	Males		166.686 (118.487 to 222.581)	57.535 (45.999 to 71.742)																					
Other meningitis	Birth prevalence - [36, 37) wks, [1000, 1500) g	Mortality	Females		169.725 (119.017 to 229.008)	63.564 (50.068 to 80.703)																					
Other meningitis	Birth prevalence - [38, 40) wks, [1000, 1500) g	Mortality	Males		174.066 (125.125 to 232.507)	57.966 (44.393 to 73.241)																					
Other meningitis	Birth prevalence - [38, 40) wks, [1000,	Mortality	Females		171.557	65.208																					
Other meningitis	Birth prevalence - [38, 40) wks, [1500,	Mortality	Males		67.302 (40.547 to 50.055)	25.206																					
Other menineitis	2000) g Birth prevalence - [38, 40) wks, [1500,	Mortality	Females		62.19	28.05																					
Other menineitis	2000) g Birth prevalence - [40, 42) wks, [1500,	Manulity	Malex		(45.884 to 83.445) 76.673	(22.625 to 35.139) 25.785																					
	2000) g Birth prevalence - [40, 42) wks, [1500,				(56.177 to 102.468) 70.411	(19.387 to 34.168) 29.113																					
Other meningins	2000) g	stortany	remates		(49.221 to 97.952) 1564.792	(21.355 to 38.272) 618 595																					
Encephalitis	Birth prevalence - [0, 24) wks, [0, 500) g	Mortality	Males		(1056.542 to 2116.062) 1600.122	(458.842 to 812.921)																					
Encephalitis	Birth prevalence - [0, 24) wks, [0, 500) g	Mortality	Females		(1050.664 to 2011 977)	(526.178 to 921.018)																					
Encephalitis	Birth prevalence - [0, 24) wks, [500, 1000) 8	Mortality	Males		(825.412 to 1506.837)	457.5 (352.552 to 573.483)																					
Encephalitis	Birth prevalence - [0, 24) wks, [500, 1000) 8	Mortality	Females		1169.123 (802.003 to 1617.979)	515.406 (396.713 to 641.541)																					
Encephalitis	Birth prevalence - [24, 26) wks, [500, 1000) 8	Mortality	Males		955.583 (723.748 to 1244.265)	443.357 (363.03 to 534.695)																					
Encephalitis	Birth prevalence - [24, 26) wks, [500, 1000) 8	Mortality	Females		947.143 (702.662 to 1237.093)	487.549 (387.307 to 603.498)																					
Encephalitis	Birth prevalence - [26, 28) wks, [500, 1000) 8	Mortality	Males		497.817 (377.617 to 648.547)	330.886 (261.438 to 401.709)																					
Encephalitis	Birth prevalence - [26, 28) wks, [500, 1000) 8	Mortality	Females		483.682 (354.946 to 629.517)	344.618 (274.427 to 419.864)																					
Encephalitis	Birth prevalence - [30, 32) wks, [500, 1000)	Mortality	Males		236.614 (163.821 to 324.502)	149.995 (117.866 to 188.368)																					
Encephalitis	Birth prevalence - [30, 32) wks, [500, 1000)	Mortality	Females		229.197 (157.606 to 317.194)	152.117 (120.779 to 190.583)																					
Encephalitis	o Birth prevalence - [28, 30) wks, [500, 1000)	Mortality	Males		297.629	216.995																					
Encertulitis	8 Birth prevalence - [28, 30) wks, [500, 1000)	Mortality	Females		281.056	219.884																					
Envertualitie	8 Birth prevalence - [26, 28) wks, [1000,	Manulity	Malex		(198.176 to 386.635) 267.91	(174.264 to 272.704) 164.167																					
Energhalitie	1500) g Birth prevalence - [26, 28) wks, [1000,	Manufita	Emple		(210.177 to 332.92) 266.509	(132.898 to 200.569) 174.222																					
	1500) g Birth prevalence - [34, 36) wks, [1000,				(197.461 to 346.932) 142.056	(137.431 to 217.349) 52.86																					
Enceptants	1500) g Birth mevalence - (34, 36) wks. [1000.	stortany	Males		(98.086 to 197.774) 141.899	(42.914 to 64.617) 57.421																					
Encephalitis	1500) g Rich muschang (28, 20) sels (1600	Moriality	Females		(95.864 to 197.656)	(46.452 to 71.339)																					
Encephalitis	2000) g	Mortality	Males		(97.178 to 167.026)	(40.539 to 61.919)																					
Encephalitis	2000) g	Mortality	Females		(96.513 to 172.188)	(46.36 to 70.038)																					
Encephalitis	Birth prevalence - [28, 30) wks, [1000, 1500) g	Mortality	Males		158.563 (120.99 to 204.947)	103.32 (83.486 to 127.144)																					
Encephalitis	Birth prevalence - [28, 30) wks, [1000, 1500) g	Mortality	Females		153.905 (112.327 to 200.786)	107.529 (86.954 to 131.78)																					
Encephalitis	Birth prevalence - [32, 34) wks, [1000, 1500) g	Mortality	Males		117.142 (81.354 to 161.101)	53.185 (43.049 to 66.274)																					
Encephalitis	Birth prevalence - [32, 34) wks, [1000, 1500) g	Mortality	Females		115.171 (79.363 to 159.206)	56.034 (45.982 to 68.36)																					
Encephalitis	Birth prevalence - [30, 32) wks, [1000, 1500) g	Mortality	Males		119.308 (87.769 to 160.885)	67.163 (54.863 to 82.638)																					
Encephalitis	Birth prevalence - [30, 32) wks, [1000, 1500) g	Mortality	Females		115.448 (84.272 to 156.425)	69.14 (55.873 to 85.012)																					
Encephalitis	Birth prevalence - [37, 38) wks, [1500, 2000) g	Mortality	Males		62.972 (46.159 to 83.484)	24.148 (20.066 to 29.406)																					
Encephalitis	Birth prevalence - [37, 38) wks, [1500, 2000) g	Mortality	Females		59.988 (43.974 to 79.053)	26.719 (21.746 to 32.816)																					
Encephalitis	Birth prevalence - [36, 37) wks, [1500, 2000) g	Mortality	Males		60.218 (43.669 to 82.48)	23.031 (18.793 to 28.483)																					
Encephalitis	Birth prevalence - [36, 37] wks, [1500, 2000) -	Mortality	Females		58.527	25.143																					
Encephalitis	2000) g Birth prevalence - [30, 32) wks, [2000, 2000)	Mortality	Males		(42.1726580.357) 67.971	18.03																					
Encephalitis	2500) g Birth prevalence - [30, 32) wks, [2000, 2000)	Mortality	Female-		(20.354 to 88.935) 69.383	(14.821 to 22.103)																					
Encentralisis	2500) g Birth prevalence - [30, 32) wks, [1500,	Montality	Maley		(49.108 to 94.583) 77.369	(17.836 to 27.163) 31.079																					
Encode V	2000) g Birth prevalence - [30, 32) wks, [1500.	M			(59.702 to 99.232) 76.134	(25.786 to 36.724) 34.756																					
inceptability	2000) g Birth prevalence - (34, 36) wks. (1500	autulity	remates		(56.885 to 100.996) 55.555	(28.764 to 41.849) 21.346																					
Encephalitis	2000) g Birth prevalence - 134 36) wise [1500	Mortality	Males		(39.553 to 75.104) 54.335	(17.677 to 26.143) 23.046																					
Encephalitis	2000) g	Mortality	Females		(38.617 to 75.24)	(18.743 to 28.287)																					
Encephalitis	2000) g	Mortality	Males		57.155 (42.484 to 73.651)	23.114 (19.028 to 27.915)																					
Encephalitis	Batth prevalence - [32, 34) wks, [1500, 2000) g	Mortality	Females		56.101 (39.794 to 76.295)	25.149 (20.615 to 30.388)																					
Encephalitis	Birth prevalence - [32, 34) wks, [2000, 2500) g	Mortality	Males		37.444 (29.026 to 48.227)	12.233 (10.252 to 14.477)																					
Encephalitis	Birth prevalence - [32, 34) wks, [2000, 2500) g	Mortality	Females		36.874 (26.658 to 49.653)	14.384 (12.095 to 17.03)																					
Encephalitis	Birth prevalence - [40, 42) wks, [2000, 2500) g	Mortality	Males		18.092 (13.292 to 23.719)	9.23 (7.037 to 11.454)																					

				tor annorene arr p											Aş	ges											
NI 0 .				All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Encertulitis	Category / Units Birth prevalence - [40, 42) wks, [2000,	Mortility / Mortal	Females		15.574	9.975		I				I	I			11		1					11				
	2500) g Birth prevalence - [38, 40) wks, [2000,				(11.516 to 20.778) 13.104	(7.82 to 12.46) 8.198																					
Enceptains	2500) g	storcarry	MIRO		(9.829 to 16.99)	(6.786 to 9.959)																					
Encephalitis	2500) g	Mortality	Females		(8.389 to 14.38)	(7.04 to 10.449)																					
Encephalitis	Birth prevalence - [32, 34) wks, [2500, 3000) g	Mortality	Males		33.063 (24.393 to 43.503)	8.441 (6.822 to 10.431)																					
Encephalitis	Birth prevalence - [32, 34) wks, [2500, 3000) g	Mortality	Females		32.812 (23.439 to 45.567)	10.398 (8.227 to 13.042)																					
Encephalitis	Birth prevalence - [34, 36) wks, [2000, 25000 -	Mortality	Males		21.925 (16.305 to 20.422)	9.367																					
Encephalitis	Bith prevalence - [34, 36) wks, [2000,	Mortality	Females		21.297	10.295																					
- -	2500) g Birth prevalence - [37, 38) wks, [2000,				(15.657 to 28.761) 13.0	(8.548 to 12.273) 8.096																					
Enceptains	2500) g Birth musclemen (27, 28) who (2000)	storcarry	MIRO		(10.102 to 16.456)	(6.724 to 9.676)																					
Encephalitis	2500) g	Mortality	Females		(8.805 to 15.11)	(6.994 to 10.342)																					
Encephalitis	Birth prevalence - [36, 37) wks, [2000, 2500) g	Mortality	Males		14.401 (10.789 to 18.654)	8.221 (6.917 to 9.923)																					
Encephalitis	Birth prevalence - [36, 37) wks, [2000, 2500) g	Mortality	Females		13.513 (9.817 to 17.942)	8.654 (7.215 to 10.369)																					
Encephalitis	Birth prevalence - [34, 36) wks, [2500, 3000) g	Mortality	Males		13.419 (10.387 to 16.819)	5.562 (4.646 to 6.696)																					
Encephalitis	Birth prevalence - [34, 36) wks, [2500, 3000) e	Mortality	Females		13.266 (2.665 to 17.682)	6.395 (5.292 to 7.606)																					
Encertulitis	Birth prevalence - [34, 36) wks, [4000,	Mortality	Males		23.096	2.895																					
- -	4500) g Birth prevalence - [34, 36) wks, [4000,		n		(14.708 to 35.098) 25.038	(2.245 to 3.716) 3.778																					
Enceptains	4500) g Risk muschener (24, 26) who (2000)	storcarry	Pemases		(15.763 to 37.255)	(2.855 to 4.925)																					
Encephalitis	3500) g	Mortality	Males		(10.222 to 18.478)	(3.449 to 5.338)																					
Encephalitis	15000, 3500) g	Mortality	Females		(10.269 to 20.114)	(4.145 to 6.564)																					
Encephalitis	Birth prevalence - [36, 37) wks, [2500, 3000) g	Mortality	Males		4.874 (4.014 to 5.713)	3.699 (3.134 to 4.374)																					
Encephalitis	Birth prevalence - [36, 37) wks, [2500, 3000) g	Mortality	Females		4.609 (3.731 to 5.61)	3.898 (3.235 to 4.67)																					
Encephalitis	Birth prevalence - [34, 36) wks, [3500, 4000) e	Mortality	Males		18.024 (12.279 to 25.547)	3.657 (2.838 to 4.675)																					
Encephalitis	Birth prevalence - [34, 36) wks, [3500,	Mortality	Females		19.263	4.634																					
Envertuititis	4000) g Birth prevalence - [37, 38) wks, [2500,	Mertility	Malex		(12.567 to 27.924) 3.306	(3.576 to 5.989) 3.194																					
	3000) g Birth menulance - [37-38) with [2500	,			(2.82 to 3.843) 2.991	(2.691 to 3.803) 3.242																					
Encephalitis	3000) g	Mortality	Females		(2.496 to 3.61)	(2.745 to 3.817)																					
Encephalitis	3000) g	Mortality	Males		(3.002 to 4.693)	(2.56 to 3.923)																					
Encephalitis	Birth prevalence - [40, 42) wks, [2500, 3000) g	Mortality	Females		3.244 (2.486 to 4.159)	3.228 (2.605 to 3.985)																					
Encephalitis	Birth prevalence - [38, 40) wks, [2500, 3000) g	Mortality	Males		2.755 (2.274 to 3.309)	2.944 (2.44 to 3.548)																					
Encephalitis	Birth prevalence - [38, 40) wks, [2500, 3000) g	Mortality	Females		2.376 (1.91 to 2.886)	2.938 (2.434 to 3.503)																					
Encephalitis	Birth prevalence - [36, 37) wks, [3000, 35000 e	Mortality	Males		3.774 (3.094 to 4.497)	2.466 (2.058 to 2.929)																					
Encephalitis	Birth prevalence - [36, 37) wks, [3000,	Mortality	Females		3.73	2.715																					
Example dist	3500) g Birth prevalence - [36, 37) wks, [4000,	Mantality	Malar		(2.981 to 4.646) 6.826	(2.277 to 3.218) 1.77																					
	4500) g Birth prevalence - [36, 37) wks, [4000,				(5.212 to 9.045) 7.269	(1.491 to 2.082) 2.177																					
	4500) g Birth menulence - (36, 37) wks. (3500,				(5.144 to 9.821) 4.544	(1.786 to 2.607) 2.057																					
Encephalotis	4000) g	Mortality	Males		(3.64 to 5.622)	(1.74 to 2.44)																					
Encephalitis	4000) g	Mortality	Females		(3.577 to 6.014)	(1.98 to 2.864)																					
Encephalitis	atini prevazitce - [37, 36) wisi, [3000, 3500) g	Mortality	Males		(1.759 to 2.293)	(1.613 to 2.224)																					
Encephalitis	Birth prevalence - [37, 38) wks, [3000, 3500) g	Mortality	Females		1.925 (1.582 to 2.328)	1.972 (1.669 to 2.313)																					
Encephalitis	Birth prevalence - [37, 38) wks, [4000, 4500) g	Mortality	Males		3.28 (2.596 to 4.133)	1.335 (1.171 to 1.532)																					
Encephalitis	Birth prevalence - [37, 38) wks, [4000, 4500) g	Mortality	Females		3.521 (2.649 to 4.5)	1.559 (1.333 to 1.835)																					
Encephalitis	Birth prevalence - [37, 38) wks, [3500, 4000) g	Mortality	Males		2.128 (1.833 to 2.466)	1.505 (1.299 to 1.76)																					
Encephalitis	Birth prevalence - [37, 38) wks, [3500, 40000 -	Mortality	Females		2.142	1.661																					
Encephalitis	Birth prevalence - [40, 42) wks, [3000,	Mortality	Males		1.436	1.47																					
Encertulitis	Birth prevalence - [40, 42) wks, [3000,	Mortality	Females		1.326	1.465																					
Encertulina	3500) g Birth prevalence - [38, 40) wks, [3000,	Mortality	Maler		(1.069 to 1.614) 1.33	(4.188 to 1.775) 1.559																					
	3500) g Birth menulence - (38, 40) wks, (3000,				(1.155 to 1.53) 1.224	(1.305 to 1.851) 1.564																					
Lange Bills	3500) g Birth mevalence - [38, 40) w/o: [4000	stortaity	. emailes		(1.0 to 1.492) 1.787	(1.304 to 1.847) 1.175																					
Encephalotis	4500) g	Mortality	Males		(1.453 to 2.182)	(1.005 to 1.371)																					
Encephalitis	4500) g	Mortality	Females		(1.467 to 2.388)	(1.022 to 1.465)																					
Encephalitis	Bath prevalence - [38, 40) wks, [3500, 4000) g	Mortality	Males		1.785 (1.478 to 2.147)	1.173 (1.0 to 1.377)																					
Encephalitis	Birth prevalence - [38, 40) wks, [3500, 4000) g	Mortality	Females		1.892 (1.481 to 2.352)	1.23 (1.03 to 1.46)																					
Encephalitis	Birth prevalence - [40, 42) wks, [3500, 4000) g	Mortality	Males		1.0 (1.0 to 1.0)	1.003 (1.0 to 1.046)																					
Encephalitis	Birth prevalence - [40, 42) wks, [3500, 4000) g	Mortality	Females		1.002 (1.0 to 1.013)	1.001 (1.0 to 1.006)																					
Encephalitis	Birth prevalence - [40, 42) wks, [4000, 4500) g	Mortality	Males		1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																					
Encephalitis	Birth prevalence - [40, 42) wks, [4000, 4500) #	Mortality	Females		1.0	1.0																					
Encephalitis	Birth prevalence - [28, 30) wks, [2000, 2600	Mortality	Males		117.172	27.726																					
Encertulisis	2300) g Birth prevalence - [28, 30) wks, [2000,	Marality	Female		(43.895 to 158.056) 121.682	(21.877 to 34.972) 33.983																					
Ennederic	2500) g Birth prevalence - [28, 30) wks, [2500,	Manadia	Mal		(84.349 to 171.375) 77.948	(26.101 to 43.404) 16.608																					
Exceptions	3000) g Birth mevalence - 128-30) wise 12600	Mortainy	MINES		(54.687 to 105.047) 79.193	(12.653 to 21.188) 20 %7																					
Encephalitis	3000) g Rich musclama (20. 200 - 1. 200-	Mortality	Females		(54.099 to 112.236)	(15.089 to 26.434)																					
Encephalitis	3500) g	Mortality	Males		(29.891 to 57.227)	(7.777 to 13.056)																					
Encephalitis	north prevalence - [28, 30) wks, [3000, 3500) g	Mortality	Females		42.551 (28.25 to 63.209)	(9.084 to 15.544)																					

ppendix Table 6a. Relative risi	is used by age and sex for each oute	0003101-801-058-08000	stexcept		on alconor	and showing						А	Ages											
Risk - Outcome	Category / Units	Morbidity / Mortalit	v Sex	All-age	0-6 days	7-27 days	28-364 days 1-4 years	5-9 years 10-14 years	15-19 years	20-24 years 2	25-29 years 30-34 yea	s 35-39 years	40-44 years	45-49 years	50-54 years	55-59 years 6	i0-64 years 65-6	years 70-7	4 years	75-79 years 80-8	4 years 85-8	9 years 90-94	years	95+ years
Encephalitis	Birth prevalence - [30, 32) wks, [2500, 2000) -	Mortality	Males	(1)	58.722 410 to 78 973)	12.115					1													
Encerebalitis	Birth prevalence - [30, 32) wks, [2500,	Mortality	Females	(42)	59.522	(9.5181015.521)																		
Enumberitie	3000) g Birth prevalence - [30, 32) wks, [3000,	Mantality	Malar	(42)	45.67	(11.936 to 19.581) 8.381																		
	3500) g Birth merculence - (30, 32) wks, (3000,	January .		(32)	.014 to 65.531) 46.104	(6.362 to 10.842) 10.506																		
Encephalitas	3500) g	Mortality	Females	(30.	243 to 66.207)	(8.041 to 13.513)																		
Encephalitis	Both prevalence - [30, 32) wks, [3500, 4000) g	Mortality	Males	(21:	36.334 .558 to 54.813)	5.698 (4.293 to 7.349)																		
Encephalitis	Birth prevalence - [30, 32) wks, [3500, 4000) g	Mortality	Females	(22)	37.931 .692 to 61.276)	6.892 (5.175 to 9.058)																		
Encephalitis	Birth prevalence - [32, 34) wks, [3000, 3500) g	Mortality	Males	(23	34.016 1.515 to 48.37)	6.577 (4.934 to 8.436)																		
Encephalitis	Birth prevalence - [32, 34) wks, [3000, 3500) a	Mortality	Females	(22)	34.585 909 to 49.754)	8.314 (6.362 to 10.789)																		
Encephalitis	Birth prevalence - [32, 34) wks, [3500,	Mortality	Males		36.248	5.068																		
Encertualitis	4000) g Birth prevalence - [32, 34) wks, [3500,	Mortality	Females	(25	38.098	6.476																		
Enumberia	4000) g Birth prevalence - [36, 37) wks, [1000,	Manufility	Malar	(23.	301 to 59.429) 166.686	(4.666 to 8.689) 57.535																		
	1500) g Birth merculence - (36, 37) wks, (1000,	January .		(118)	487 to 222.581 169.725) (45.999 to 71.742) 63.564																		
Encephalitas	1500) g	Mortality	Females	(119)	.017 to 229.008) (50.068 to 80.703)																		
Encephalitis	Both prevalence - [38, 40) wks, [1000, 1500) g	Mortality	Males	(125.	174.066 .125 to 232.507) (44.393 to 73.241)																		
Encephalitis	Birth prevalence - [38, 40) wks, [1000, 1500) g	Mortality	Females	(121:	171.557 .585 to 237.047	65.208 (48.821 to 84.308)																		
Encephalitis	Birth prevalence - [38, 40) wks, [1500, 2000) g	Mortality	Males	(49).	67.302 .547 to 89.055)	25.206 (20.365 to 31.168)																		
Encephalitis	Birth prevalence - [38, 40) wks, [1500, 2000) #	Mortality	Females	(45)	62.19 884 to 83.445)	28.05 (22.625 to 35.139)																		
Encephalitis	Birth prevalence - [40, 42) wks, [1500,	Mortality	Males	(76.673	25.785																		
Favordulitis	Birth prevalence - [40, 42) wks, [1500,	Monthline	Females	(56.1	70.411	29.113																		
N	2000) g			(49.	221 to 97.952) 1564.792	(21.355 to 38.272) 618.595																		
Neonatal preterm birth	Bath prevalence - [0, 24) wks, [0, 500) g	Mortality	Males	0	1056.542 to 2116.0621 1600.122	(458.842 to 812.921																		
Neonatal preterm birth	Birth prevalence - [0, 24) wks, [0, 500) g	Mortality	Females	0	1050.664 to	(526.178 to 921.018																		
Neomatal preterm birth	Birth prevalence - [0, 24) wks, [500, 1000) 8	Mortality	Males	(825.4	1155.815 412 to 1506.837	457.5 (352.552 to 573.483																		
Neomatal preterm birth	Birth prevalence - [0, 24) wks, [500, 1000) g	Mortality	Females	(802.0	1169.123 103 to 1617.979	515.406 (396.713 to 641.541																		
Neonatal preterm birth	Birth prevalence - [24, 26) wks, [500, 1000 8) Mortality	Males	(723.7	955.583 748 to 1244.265	443.357 (363.03 to 534.695)																		
Neonatal preterm birth	Birth prevalence - [24, 26) wks, [500, 1000) Mortality	Females	(70) 4	947.143	487.549																		
Neomatal preterm birth	o Birth prevalence - [26, 28) wks, [500, 1000) Mortality	Males		497.817	330.886																		
Noomial meterm birth	8 Birth prevalence - [26, 28] wks, [500, 1000	Mandin	Females	(and	483.682	344.618	2																	
N	8 Birth prevalence - [30, 32) wks, [500, 1000) M		(354)	946 to 629.517 236.614) (274.427 to 419.864 149.995	•																	
Neomana preterm serin	8 Rishmundana (20.22) ada (600.1000	stortany	Made	(163.	821 to 324.502) (117.866 to 188.368																		
Neomatal preterm birth	8	Mortality	Females	(157)	.606 to 317.194) (120.779 to 190.583)																	
Neomatal preterm birth	Bath prevalence - [28, 30) wks, [500, 1000 8) Mortality	Males	(214)	297.629 .953 to 396.586	216.995) (173.321 to 271.466	•																	
Neonatal preterm birth	Birth prevalence - [28, 30) wks, [500, 1000 8) Mortality	Females	(198.	281.056 176 to 386.635	219.884) (174.264 to 272.704																		
Neomatal preterm birth	Birth prevalence - [26, 28) wks, [1000, 1500) g	Mortality	Males	(210	267.91 1.177 to 332.92)	164.167 (132.898 to 200.569																		
Neomatal preterm birth	Birth prevalence - [26, 28) wks, [1000, 1500) #	Mortality	Females	0.97	266.509 461 to 346.932	174.222 (137.431 to 217.349																		
Neomatal preterm birth	Birth prevalence - [34, 36) wks, [1000, 1500) -	Mortality	Males		142.056	52.86	-																	
Neomatal preterm birth	Birth prevalence - [34, 36) wks, [1000,	Mortality	Females		141.899	57.421																		
Normal materia biath	1500) g Birth prevalence - [28, 30) wks, [1500,	Mantality	Malar	(95.3	127.966	(46.452 to 71.339) 50.018																		
	2000) g Birth merculence - (28, 30) wks, (1500,	January .		(97.1	178 to 167.026) 130.924	(40.539 to 61.919) 57.275																		
Neonatal preterm birth	2000) g Rish muscleme 128, 200 mln 11000	Mortality	Females	(96.5	513 to 172.188)	(46.36 to 70.038)																		
Neomatal preterm birth	1500) g	Mortality	Males	(120	1.99 to 204.947)	(83.486 to 127.144)																		
Neomatal preterm birth	Birth prevalence - [28, 30) wks, [1000, 1500) g	Mortality	Females	(112.	153.905 .327 to 200.786	107.529 (86.954 to 131.78)																		
Neonatal preterm birth	Birth prevalence - [32, 34) wks, [1000, 1500) g	Mortality	Males	(81.3	117.142 354 to 161.101)	53.185 (43.049 to 66.274)																		
Neomatal preterm birth	Birth prevalence - [32, 34) wks, [1000, 1500) g	Mortality	Females	(79.3	115.171 363 to 159.206)	56.034 (45.982 to 68.36)																		
Neonatal preterm birth	Birth prevalence - [30, 32) wks, [1000, 1500) g	Mortality	Males	(87.5	119.308 769 to 160.885)	67.163 (54.863 to 82.638)																		
Neomatal preterm birth	Birth prevalence - [30, 32) wks, [1000, 1500) #	Mortality	Females		115.448 272 m 156.425	69.14 (55.873 to 85.012)																		
Neomatal preterm birth	Birth prevalence - [37, 38) wks, [1500, 2000) -	Mortality	Males	(64.2	62.972	24.148																		
Neoenital workers hirth	2000) g Birth prevalence - [37, 38) wks, [1500,	Montality	Female	(46.	59.988	(20.006 to 29.406) 26.719																		
Marcal Article	2000) g Birth prevalence - (36, 37) wks. (1500			(43.	974 to 79.053) 60.218	(21.746 to 32.816) 23.031																		
Neonatal preterm birth	2000) g Risk manulana, 136, 27) ada, 11800	Mortality	Males	(43	669 to 82.48)	(18.793 to 28.483)																		
Neonatal preterm birth	2000) g	Mortality	Females	(42.	.172 to 80.557)	(20.331 to 30.566)																		
Neomatal preterm birth	Birth prevalence - [30, 32) wks, [2000, 2500) g	Mortality	Males	(50.	67.971 .354 to 88.935)	18.03 (14.621 to 22.103)																		
Neomatal preterm birth	Birth prevalence - [30, 32) wks, [2000, 2500) g	Mortality	Females	(49.	69.383 108 to 94.583)	22.069 (17.836 to 27.163)																		
Neonatal preterm birth	Birth prevalence - [30, 32) wks, [1500, 2000) g	Mortality	Males	(59).	77.369 .702 to 99.232)	31.079 (25.786 to 36.724)																		
Neomatal preterm birth	Birth prevalence - [30, 32) wks, [1500, 2000) g	Mortality	Females	156.5	76.134 885 to 100.9960	34.756 (28.764 to 41 849)																		
Neomatal preterm birth	Birth prevalence - [34, 36) wks, [1500,	Mortality	Males	(55.555	21.346																		
Neoenital workers hirth	2000) g Birth prevalence - [34, 36) wks, [1500,	Montality	Female	(39.	.353 to 75.104) 54.335	(17.677 to 26.143) 23.046																		
Marcal Article	2000) g Birth prevalence - (32, 34) wks. (1500			(38	57.155	(18.743 to 28.287) 23.114																		
Neonatal preterm beth	2000) g Birth menalence . (32-34) wky. (1600	Mortality	Males	(42)	484 to 73.651) 56 101	(19.028 to 27.915) 25.149																		
Neonatal preterm birth	2000) g	Mortality	Females	(39.	.794 to 76.295)	(20.615 to 30.388)																		
Neomatal preterm birth	Birth prevalence - [32, 34) wks, [2000, 2500) g	Mortality	Males	(29)	37.444 .026 to 48.227)	12.233 (10.252 to 14.477)																		
Neomatal preterm birth	Birth prevalence - [32, 34) wks, [2000, 2500) g	Mortality	Females	(26)	36.874 .658 to 49.653)	14.384 (12.095 to 17.03)																		
Neonatal preterm birth	Birth prevalence - [40, 42) wks, [2000, 2500) g	Mortality	Males	(13:	18.092 292 to 23.719)	9.23 (7.037 to 11.454)																		

				e annotene ant po											А	ıges											
Risk - Outcome	Category / Units	Morbidity / Mortality	Sex	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Neomatal preterm birth	Birth prevalence - [40, 42) wks, [2000,	Mortality	Females		15.574	9.975												1									
Neonatal preterm birth	2500) g Birth prevalence - [38, 40) wks, [2000,	Mortality	Males		13.104	8.198																					
	2500) g Birth prevalence - [38, 40) wks, [2000,	M			(9.829 to 16.99) 11.308	(6.786 to 9.959) 8.577																					
	2500) g Birth menulence - (32, 34) wks, (2500,	January .	- canades		(8.389 to 14.38) 33.063	(7.04 to 10.449) 8.441																					
Neonatal preterm birth	3000) g Birth manalanan (22, 24) ada (2000)	Mortality	Malex		(24.393 to 43.503)	(6.822 to 10.431)																					
Neomatal preterm birth	3000) g	Mortality	Females		(23.439 to 45.567)	(8.227 to 13.042)																					
Neomatal preterm birth	Bath prevalence - [34, 36) wiss, [2000, 2500) g	Mortality	Males		(16.305 to 29.433)	9.367 (7.859 to 11.112)																					
Neomatal preterm birth	Birth prevalence - [34, 36) wks, [2000, 2500) g	Mortality	Females		21.297 (15.657 to 28.761)	10.295 (8.548 to 12.273)																					
Neonatal preterm birth	Birth prevalence - [37, 38) wks, [2000, 2500) g	Mortality	Males		13.0 (10.102 to 16.456)	8.096 (6.724 to 9.676)																					
Neonatal preterm birth	Birth prevalence - [37, 38) wks, [2000, 2500) g	Mortality	Females		11.563 (8.805 to 15.11)	8.467 (6.994 to 10.342)																					
Neomatal preterm birth	Birth prevalence - [36, 37) wks, [2000, 2500) g	Mortality	Males		14.401 (10.789 to 18.654)	8.221 (6.917 to 9.923)																					
Neomatal preterm birth	Birth prevalence - [36, 37] wks, [2000, 2500) #	Mortality	Females		13.513 (9.817 to 17.947)	8.654 (7.215 to 10.369)																					
Neonatal preterm birth	Birth prevalence - [34, 36) wks, [2500,	Mortality	Males		13.419	5.562																					
Noomial meterm kith	5000) g Birth prevalence - [34, 36) wks, [2500,	Martality	Females		(10.387 10 16.819) 13.266	6.395																					
	3000) g Birth prevalence - [34, 36) wks, [4000,				(9.666 to 17.689) 23.096	(5.292 to 7.606) 2.895																					
reconaria preterm term	4500) g Birth menalewer - [34-36) wise [4000	stortany	MERCY		(14.708 to 35.098) 25.038	(2.245 to 3.716) 3.778																					
Neomatal preterm birth	4500) g	Mortality	Females		(15.763 to 37.255)	(2.855 to 4.925)																					
Neonatal preterm birth	ann prevaence - [54, 56) was, [5000, 3500) g	Mortality	Males		(10.222 to 18.478)	4.322 (3.449 to 5.338)																					
Neomatal preterm birth	Bath prevalence - [34, 36) wiss, [3000, 3500) g	Mortality	Females		(10.269 to 20.114)	5.265 (4.145 to 6.564)																					
Neonatal preterm birth	Birth prevalence - [36, 37) wks, [2500, 3000) g	Mortality	Males		4.874 (4.014 to 5.713)	3.699 (3.134 to 4.374)																					
Neomatal preterm birth	Birth prevalence - [36, 37) wks, [2500, 3000) g	Mortality	Females		4.609 (3.731 to 5.61)	3.898 (3.235 to 4.67)																					
Neonatal preterm birth	Birth prevalence - [34, 36) wks, [3500, 4000) g	Mortality	Males		18.024 (12.279 to 25.547)	3.657 (2.838 to 4.675)																					
Neomatal preterm birth	Birth prevalence - [34, 36) wks, [3500, 4000) g	Mortality	Females		19.263 (12.567 to 27.924)	4.634 (3.576 to 5.989)																					
Neonatal preterm birth	Bith prevalence - [37, 38) wks, [2500, 2000) -	Mortality	Males		3.306	3.194																					
Neonatal preterm birth	Birth prevalence - [37, 38) wks, [2500,	Mortality	Females		2.991	3.242																					
Noomial meterm birth	3000) g Birth prevalence - [40, 42) wks, [2500,	Martality	Malex		(2.496 to 3.61) 3.771	(2.745 to 3.817) 3.175																					
	3000) g Birth menulence - (40, 42) wks, (2500,				(3.002 to 4.693) 3.244	(2.56 to 3.923) 3.228																					
Neonatal preterm birth	3000) g Birth menalewer - [38, 40) wise [2500	Mortality	Females		(2.486 to 4.159) 2.755	(2.605 to 3.985) 2.944																					
Neonatal preterm birth	3000) g Birth manufanar (28, 40) min (2000)	Mortality	Males		(2.274 to 3.309)	(2.44 to 3.548)																					
Neoental preterm birth	3000) g	Mortality	Females		(1.91 to 2.886)	(2.434 to 3.503)																					
Neonatal preterm birth	Bath prevalence - [36, 37) wks, [3000, 3500) g	Mortality	Males		3.774 (3.094 to 4.497)	2.466 (2.058 to 2.929)																					
Neoental preterm birth	Birth prevalence - [36, 37) wks, [3000, 3500) g	Mortality	Females		3.73 (2.981 to 4.646)	2.715 (2.277 to 3.218)																					
Neonatal preterm birth	Birth prevalence - [36, 37) wks, [4000, 4500) g	Mortality	Males		6.826 (5.212 to 9.045)	1.77 (1.491 to 2.082)																					
Neonatal preterm birth	Birth prevalence - [36, 37) wks, [4000, 4500) g	Mortality	Females		7.269 (5.144 to 9.821)	2.177 (1.786 to 2.607)																					
Neonatal preterm birth	Birth prevalence - [36, 37) wks, [3500, 4000) g	Mortality	Males		4.544 (3.64 to 5.622)	2.057 (1.74 to 2.44)																					
Neomatal preterm birth	Birth prevalence - [36, 37) wks, [3500, 4000) g	Mortality	Females		4.662 (3.577 to 6.014)	2.398 (1.98 to 2.864)																					
Neonatal preterm birth	Birth prevalence - [37, 38) wks, [3000, 26000 -	Mortality	Males		2.007	1.888																					
Neonatal preterm birth	Birth prevalence - [37, 38) wks, [3000,	Mortality	Females		1.925	1.972																					
Neonatal meterm birth	3500) g Birth prevalence - [37, 38) wks, [4000,	Mortality	Males		3.28	1.335																					
Noomial meterm birth	4500) g Birth prevalence - [37, 38) wks, [4000,	Martality	Females		(2.596 to 4.133) 3.521	(1.171 to 1.532) 1.559																					
N	4500) g Birth prevalence - [37, 38) wks, [3500,				(2.649 to 4.5) 2.128	(1.333 to 1.835) 1.505																					
Neonara preterm term	4000) g Birth mevalence - (37, 38) wks. (3500,	Mortany	MERCY		(1.833 to 2.466) 2.142	(1.299 to 1.76) 1.661																					
Neonatal preterm birth	4000) g Birth menalence - (40, 42) who (2000)	Mortality	remates		(1.694 to 2.67)	(1.411 to 1.961)																					
Neoental preterm birth	3500) g Rick manifestary (at the to page 1	Mortality	Males		(1.245 to 1.65)	(1.199 to 1.8)																					
Neomatal preterm birth	3500) g Risk muchania (20, 40, 40, 1000)	Mortality	Females		(1.069 to 1.614)	(1.188 to 1.775)																					
Neonatal preterm birth	asmn prevaænce - [38, 40] wiss [3000, 3500) g	Mortality	Males		1.53 (1.155 to 1.53)	1.559 (1.305 to 1.851)																					
Neoeutal preterm birth	Birth prevalence - [38, 40) wks, [3000, 3500) g	Mortality	Females		1.224 (1.0 to 1.492)	1.564 (1.304 to 1.847)																					
Neonatal preterm birth	Birth prevalence - [38, 40) wks, [4000, 4500) g	Mortality	Males		1.787 (1.453 to 2.182)	1.175 (1.005 to 1.371)																					
Neomatal preterm birth	Birth prevalence - [38, 40) wks, [4000, 4500) g	Mortality	Females		1.877 (1.467 to 2.388)	1.224 (1.022 to 1.465)																					
Neomatal preterm birth	Birth prevalence - [38, 40) wks, [3500, 4000) g	Mortality	Males		1.785 (1.478 to 2.147)	1.173 (1.0 to 1.377)																					
Neonatal preterm birth	Birth prevalence - [38, 40) wks, [3500, 4000) g	Mortality	Females		1.892 (1.481 to 2.352)	1.23 (1.03 to 1.46)																					
Neonatal preterm birth	Birth prevalence - [40, 42) wks, [3500, 4000) e	Mortality	Males		1.0	1.003																					
Neonatal preterm birth	Birth prevalence - [40, 42) wks, [3500, 40000 r	Mortality	Females		1.002	1.001																					
Neonatal meterm birth	4000) g Birth prevalence - [40, 42) wks, [4000,	Mortality	Males		(1.0 to 1.013)	(LUID 1.006) 1.0																					
Neomatal workers birth	4500) g Birth prevalence - [40, 42) wks, [4000,	Mortalia	Females		(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0																					
Normal States	4500) g Birth prevalence - [28, 30) wks, [2000.	Mart	Mal		(1.0 to 1.0) 117.172	(1.0 to 1.0) 27.726																					
reconaria preterm peth	2500) g Bith prevalence - 128 30) wise 12000	мотапу	MINEX		(83.895 to 158.056) 121.682	(21.877 to 34.972) 33.983																					
Neoeatal preterm birth	2500) g Bith mendance - 128, 300 min - 12600	Mortality	Females		(84.349 to 171.375) 77 0.49	(26.101 to 43.404)																					
Neonatal preterm birth	3000) g	Mortality	Males		(54.687 to 105.047)	(12.653 to 21.188)																					
Neomatal preterm birth	Both prevalence - [28, 30) wks, [2500, 3000) g	Mortality	Females		79.193 (54.099 to 112.236)	20.387 (15.089 to 26.434)																					
Neonatal preterm birth	Birth prevalence - [28, 30) wks, [3000, 3500) g	Mortality	Males		42.199 (29.891 to 57.227)	10.082 (7.777 to 13.056)																					
Neomatal preterm birth	Birth prevalence - [28, 30) wks, [3000, 3500) g	Mortality	Females		42.551 (28.25 to 63.209)	11.989 (9.084 to 15.544)																					

opendix Table 6a. Relative ris	is used by age and sex for each outco	me for all risk factors ex	cept for ambient	air pollution alcohol.	and snowing.									1	Ages											
Risk - Outcome	Category / Units	Morbidity / Mortality	All-ag	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Neomatal preterm birth	Birth prevalence - [30, 32] wks, [2500, 2000) -	Mortality 1	hles	58.722	12.115	1	I										I									
Neomatal preterm birth	S000) g Birth prevalence - [30, 32) wks, [2500,	Mortality F	males	(42,41910 /8.873) 59.522	(9.5181015.521)																					
Normal materia kink	3000) g Birth prevalence - [30, 32) wks, [3000,	Manufility	li la	(42.058 to 82.793) 45.67	(11.936 to 19.581) 8.381																					
	3500) g Birth meyalence - [30, 32) wks, [3000,	Juniary .		(32.014 to 65.531) 46.104	(6.362 to 10.842) 10.506																					
Neonatal preterm birth	3500) g	Mortality F	males	(30.243 to 66.207)	(8.041 to 13.513)																					
Neonatal preterm birth	Bath prevalence - [30, 32) wks, [3500, 4000) g	Mortality	fales	36.334 (21.558 to 54.813)	(4.293 to 7.349)																					
Neonatal preterm birth	Birth prevalence - [30, 32) wks, [3500, 4000) g	Mortality F	males	37.931 (22.692 to 61.276)	6.892 (5.175 to 9.058)																					
Neonatal preterm birth	Birth prevalence - [32, 34) wks, [3000, 3500) g	Mortality	fales	34.016 (23.515 to 48.37)	6.577 (4.934 to 8.436)																					
Neonatal preterm birth	Birth prevalence - [32, 34) wks, [3000, 3500) #	Mortality F	males	34.585 (22.909 to 49.754)	8.314 (6.362 to 10.789)																					
Neomatal preterm birth	Birth prevalence - [32, 34) wks, [3500,	Mortality	fales	36.248	5.068																					
Normal materia kink	4000) g Birth prevalence - [32, 34) wks, [3500,	Mantalia, E		(23.158 to 54.67) 38.098	(3.761 to 6.741) 6.476																					
Neonata preterm term	4000) g Birth menalence - [36 37) wise [1000	Mortany P	masex	(23.301 to 59.429)	(4.666 to 8.689) 57 535																					
Neonatal preterm birth	1500) g Birth muscherer (26, 27) who (1000	Mortality	hiles	(118.487 to 222.581	 (45.999 to 71.742) 62.664 																					
Neonatal preterm birth	1500) g	Mortality F	males	(119.017 to 229.008	 (50.068 to 80.703) 																					
Neonatal preterm birth	Birth prevalence - [38, 40) wks, [1000, 1500) g	Mortality	fales	174.066 (125.125 to 232.507	57.966 (44.393 to 73.241)																					
Neonatal preterm birth	Birth prevalence - [38, 40) wks, [1000, 1500) g	Mortality F	males	171.557 (121.585 to 237.047	65.208 (48.821 to 84.308)																					
Neonatal preterm birth	Birth prevalence - [38, 40) wks, [1500, 2000) g	Mortality	fales	67.302 (49.547 to 89.055)	25.206 (20.365 to 31.168)																					
Neomatal preterm birth	Birth prevalence - [38, 40) wks, [1500,	Mortality F	males	62.19	28.05																					
Neomatal preterm birth	2000) g Birth prevalence - [40, 42) wks, [1500,	Mortality	lales	76.673	25.785																					
	2000) g Birth mevalence - [40, 42] wks, [1500,			(56.177 to 102.468 70.411) (19.387 to 34.168) 29.113																					
Neomatal preterm birth	2000) g	Mortality F	malex	(49.221 to 97.952) 1564.792	(21.355 to 38.272)																					
birth asphysia and trauma	Birth prevalence - [0, 24) wks, [0, 500) g	Mortality	fales	(1056.542 to 21.16.062) 1600.122	(458.842 to 812.921																					
Neomatal encephalopathy due to birth asphysia and trauma	Birth prevalence - [0, 24) wks, [0, 500) g	Mortality F	males	(1050.664 to 2111.977)	713.571 (526.178 to 921.018																					
Neonatal encephalopathy due to birth asphysia and trauma	 Birth prevalence - [0, 24) wks, [500, 1000) g 	Mortality	hiles	1155.815 (825.412 to 1506.83	457.5 7) (352.552 to 573.483	,																				
Neomatal encephalopathy due to birth asphysia and trauma	Birth prevalence - [0, 24) wks, [500, 1000)	Mortality F	males	1169.123 (802.003 to 1617.97	515.406 9) (396.713 to 641.541	,																				
Neomatal encephalopathy due to birth surboxis and trauma	Birth prevalence - [24, 26) wks, [500, 1000)	Mortality	hiles	955.583 (773.748 to 1244.26	443.357																					
Neonatal encephalopathy due to	8 Birth prevalence - [24, 26) wks, [500, 1000)	Mortality F	males	947.143	487.549																					
birth asphysia and trauma Neonatal encephalopathy due to	8 Birth prevalence - [26, 28) wks, [500, 1000)	N		(702.662 to 1237.09 497.817	3) (387.307 to 603.498) 330.886																					
birth asphysia and trauma	8 Richardone (26.28) adv. (500, 1000)	storcarty	ma	(377.617 to 648.547	7) (261.438 to 401.709) 244.618																					
birth asphysia and trauma	8 8	Mortality F	males	(354.946 to 629.517	7) (274.427 to 419.864																					
Neonatal encephalopathy due to birth asphysia and trauma	Bath prevalence - [30, 32] wks, [500, 1000) 8	Mortality	fales	256.614 (163.821 to 324.501	(117.866 to 188.368)	·																				
Neomatal encephalopathy due to birth asphysia and tratama	 Birth prevalence - [30, 32) wks, [500, 1000) 8 	Mortality F	males	229.197 (157.606 to 317.194	152.117 (120.779 to 190.583	, ,																				
Neonatal encephalopathy due to birth asphyxia and trauma	Birth prevalence - [28, 30] wks, [500, 1000) 8	Mortality	fales	297.629 (214.953 to 396.588	216.995 5) (173.321 to 271.466	,																				
Neomatal encephalopathy due to birth surboxis and trauma	Birth prevalence - [28, 30) wks, [500, 1000)	Mortality F	males	281.056 (198.176 to 386.639	219.884 (174.264 to 272.204																					
Neonatal encephalopathy due to	Birth prevalence - [26, 28) wks, [1000,	Mortality	fales	267.91	164.167																					
burth asphysica and trauma Neomatal encephalopathy due to	1500) g Birth prevalence - [26, 28) wks, [1000,	Metality F	males	(210.177 to 332.92 266.509) (132.898 to 200.569 174.222																					
birth asphysia and trauma Neonatal encephalopathy due to	1500) g Birth prevalence - [34, 36) wks, [1000,	N		(197.461 to 346.93) 142.056	 (137.431 to 217.349) 52.86 																					
birth asphysia and trauma Neomital oncombalarathy due to	1500) g Birth menalence - [34, 36) wise [1000	Mortany	ma	(98.086 to 197.774)	(42.914 to 64.617) 57.421																					
birth asphysia and trauma	1500) g	Mortality F	males	(95.864 to 197.656) (46.452 to 71.339)																					
Neomital encephalopathy due to birth asphysia and trauma	Bath prevalence - [28, 30] wks, [1500, 2000) g	Mortality	fales	(97.178 to 167.026	(40.539 to 61.919)																					
Neonatal encephalopathy due to birth asphysia and trazma	 Birth prevalence - [28, 30) wks, [1500, 2000) g 	Mortality F	males	130.924 (96.513 to 172.188	57.275 (46.36 to 70.038)																					
Neonatal encephalopathy due to birth asphysia and trauma	 Birth prevalence - [28, 30) wks, [1000, 1500) g 	Mortality	hiles	158.563 (120.99 to 204.947)	103.32 (83.486 to 127.144)																					
Neomatal encephalopathy due to birth assboxia and trauma	 Birth prevalence - [28, 30) wks, [1000, 1500) p 	Mortality F	males	153.905 (112.327 to 200.786	107.529 (86.954 to 131.78)																					
Neomatal encephalopathy due to	- Birth prevalence - [32, 34) wks, [1000, 1500) #	Mortality	fales	117.142	53.185																					
Neomaal encephalopathy doe to	Bith prevalence - [32, 34) wks, [1000,	Mortality F	males	115.171	56.034																					
rorm aspnysta and trauma Neomatal encephalopathy due to	 300) g Birth prevalence - [30, 32) wks, [1000, 	Mertality	him	(19.363 to 159.206) 119.308	, (+3.982 to 68.36) 67.163																					
birth asphysia and trauma Neomatal encephalopathy due to	1500) g Birth prevalence - [30, 32) wks, [1000.	Marti V		(87.769 to 160.885 115.448) (54.863 to 82.638) 69.14																					
birth asphysia and trauma	1500) g Birth menalence - (27, 28) when (1600)	Mortany F		(84.272 to 156.425) (55.873 to 85.012) 24.149																					
birth asphysia and traama	2000) g	Mortality	hiles	(46.159 to \$3.484)	(20.066 to 29.406)																					
Neomatal encephalopathy due to birth asphysia and traama	a north prevalence - [37, 38) wks; [1500, 2000) g	Mortality F	males	59.988 (43.974 to 79.053)	26.719 (21.746 to 32.816)																					
Neomatal encephalopathy due to birth asphysia and trauma	 Birth prevalence - [36, 37) wks, [1500, 2000) g 	Mortality	fales	60.218 (43.669 to 82.48)	23.031 (18.793 to 28.483)																					
Neonatal encephalopathy due to birth asphysia and traama	 Birth prevalence - [36, 37) wks, [1500, 2000) g 	Mortality F	males	58.527 (42.172 to 80.557)	25.143 (20.331 to 30.566)																					
Neomatal encephalopathy due to birth asphysia and transm	Birth prevalence - [30, 32) wks, [2000, 2500) #	Mortality	lales	67.971 (50.354 to \$8.935)	18.03 (14.621 to 22 103)																					
Neomital encephalopathy due to	Birth prevalence - [30, 32) wks, [2000, 2500) -	Mortality F	males	69.383	22.069																					
turni aopitysta ana trauma Neonatal encephalopathy due ti	Birth prevalence - [30, 32] wks, [1500,	Mortality	lales	(49.10810.94.583) 77.369	(17.85610.27.165) 31.079																					
birth asphysia and trauma Neonatal enceshalorathy due to	2000) g Birth prevalence - [30, 32) wks. [1500			(59.702 to 99.232) 76.134	(25.786 to 36.724) 34.756																					
birth asphysia and trauma	2000) g Birth menalence - (24, 26) who (1600	Mortality F	manex	(56.885 to 100.996) (28.764 to 41.849) 21.246																					
birth asphysia and trauma	2000) g	Mortality	lales	(39.553 to 75.104)	(17.677 to 26.143)																					
Neomatal encephalopathy due to birth asphysia and traama	b Both prevalence - [34, 36) wks, [1500, 2000) g	Mortality F	males	54.335 (38.617 to 75.24)	23.046 (18.743 to 28.287)																					
Neomatal encephalopathy due to birth asphysia and trauma	Birth prevalence - [32, 34) wks, [1500, 2000) g	Mortality	fales	57.155 (42.484 to 73.651)	23.114 (19.028 to 27.915)																					
Neomatal encephalopathy due to birth asphysia and trauma	 Birth prevalence - [32, 34) wks, [1500, 2000) g 	Mortality F	males	56.101 (39.794 to 76.295)	25.149 (20.615 to 30.388)																					
Neomatal encephalopathy due to	Birth prevalence - [32, 34) wks, [2000, 25000 a	Mortality	hiles	37.444	12.233 (10.252 to 14.422)																					
Neomatal encephalopathy due to	Bith prevalence - [32, 34) wks, [2000,	Mortality F	males	36.874	14.384																					
birth asphysia and traama Neomatal encephalopathy due to	2500) g Birth prevalence - [40, 42) wks, [2000,	Mastella	blar.	(26.658 to 49.653) 18.092	(12.095 to 17.03) 9.23																					
birth asphyxia and trauma	2500) g	secondity		(13.292 to 23.719)	(7.037 to 11.454)	1																				

																Aş	ges											
Index Index <t< th=""><th>Risk - Outcome</th><th>Category / Units</th><th>Morbidity / Mortality</th><th>v Sex</th><th>All-age</th><th>0-6 days</th><th>7-27 days</th><th>28-364 days</th><th>1-4 years</th><th>5-9 years</th><th>10-14 years</th><th>15-19 years</th><th>20-24 years</th><th>25-29 years</th><th>30-34 years</th><th>35-39 years</th><th>40-44 years</th><th>45-49 years</th><th>50-54 years</th><th>55-59 years</th><th>60-64 years</th><th>65-69 years</th><th>70-74 years</th><th>75-79 years</th><th>80-84 years</th><th>85-89 years</th><th>90-94 years</th><th>95+ years</th></t<>	Risk - Outcome	Category / Units	Morbidity / Mortality	v Sex	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Name <t< td=""><td>Neonatal encephalopathy due to</td><td>Birth prevalence - [40, 42) wks, [2000, 26000 -</td><td>Mortality</td><td>Females</td><td></td><td>15.574</td><td>9.975</td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td></t<>	Neonatal encephalopathy due to	Birth prevalence - [40, 42) wks, [2000, 26000 -	Mortality	Females		15.574	9.975																					
NumeNumN	Neomatal encephalopathy due to	Birth prevalence - [38, 40) wks, [2000,	Mortality	Males		13.104	8.198																					
Number Number Number Number Number Number Number Number Numbe	berth asphysia and trauma Neonatal encephalopathy due to	2500) g Birth prevalence - [38, 40) wks, [2000,	Mastality	Females		(9.829 to 16.99) 11.308	(6.786 to 9.959) 8.577																					
	birth asphyxia and trianna Neonatal encephalopathy due to	2500) g Birth prevalence - [32, 34) wks, [2500,				(8.389 to 14.38) 33.063	(7.04 to 10.449) 8.441																					
	birth asphysia and trauma	3000) g Rightmansharar (22, 24) mbr (2600	stortaity	Males		(24.393 to 43.503)	(6.822 to 10.431)																					
Number <td< td=""><td>birth asphysia and trauma</td><td>3000) g</td><td>Mortality</td><td>Females</td><td></td><td>(23.439 to 45.567)</td><td>(8.227 to 13.042)</td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td></td<>	birth asphysia and trauma	3000) g	Mortality	Females		(23.439 to 45.567)	(8.227 to 13.042)																					
Number Number Number Number Number Number Number Number Numbe	Neonatal encephalopathy due to birth asphysia and tratama	Barth prevalence - [34, 36) wks, [2000, 2500) g	Mortality	Males		21.925 (16.305 to 29.433)	9.367 (7.859 to 11.112)																					
Network <td>Neomatal encephalopathy due to birth asphysia and trauma</td> <td>Birth prevalence - [34, 36) wks, [2000, 2500) g</td> <td>Mortality</td> <td>Females</td> <td></td> <td>21.297 (15.657 to 28.761)</td> <td>10.295 (8.548 to 12.273)</td> <td></td>	Neomatal encephalopathy due to birth asphysia and trauma	Birth prevalence - [34, 36) wks, [2000, 2500) g	Mortality	Females		21.297 (15.657 to 28.761)	10.295 (8.548 to 12.273)																					
Name<	Neomatal encephalopathy due to birth asphyxia and trauma	Birth prevalence - [37, 38) wks, [2000, 2500) g	Mortality	Males		13.0 (10.102 to 16.456)	8.096 (6.724 to 9.676)																					
	Neonatal encephalopathy due to birth asobycia and tratama	Birth prevalence - [37, 38) wks, [2000, 2500) g	Mortality	Females		11.563 (8.805 to 15.11)	8.467 (6.994 to 10.342)																					
	Neonatal encephalopathy due to	Birth prevalence - [36, 37) wks, [2000, 2600) -	Mortality	Males		14.401	8.221																					
	Neonatal encephalopathy due to	2300) g Birth prevalence - [36, 37) wks, [2000,	Martility	Females		(10.78910 18.034) 13.513	8.654																					
	birth asphysia and trauma Neomatal encephalopathy due to	2500) g Birth prevalence - [34, 36) wks, [2500,				(9.817 to 17.942) 13.419	(7.215 to 10.369) 5.562																					
	birth asphysia and trauma Neomtol encombaloreathy due to	3000) g Birth menulence - [34, 36) wise [2500	January	mane		(10.387 to 16.819)	(4.646 to 6.696) 6 395																					
	birth asphysia and trauma	3000) g	Mortality	Females		(9.666 to 17.689)	(5.292 to 7.606)																					
	Neonatal encephalopathy due to birth asphysia and tratama	Barth prevalence - [34, 36) wks, [4000, 4500) g	Mortality	Males		23.096 (14.708 to 35.098)	(2.245 to 3.716)																					
Network Network <t< td=""><td>Neonatal encephalopathy due to birth asphyxia and trauma</td><td>Birth prevalence - [34, 36) wks, [4000, 4500) g</td><td>Mortality</td><td>Females</td><td></td><td>25.038 (15.763 to 37.255)</td><td>3.778 (2.855 to 4.925)</td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td></t<>	Neonatal encephalopathy due to birth asphyxia and trauma	Birth prevalence - [34, 36) wks, [4000, 4500) g	Mortality	Females		25.038 (15.763 to 37.255)	3.778 (2.855 to 4.925)																					
	Neomatal encephalopathy due to birth asphysia and trauma	Birth prevalence - [34, 36) wks, [3000, 3500) g	Mortality	Males		14.006 (10.222 to 18.478)	4.322 (3.449 to 5.338)																					
	Neonatal encephalopathy due to birth asobycia and tratama	Birth prevalence - [34, 36) wks, [3000, 3500) g	Mortality	Females		14.375 (10.269 to 20.114)	5.265 (4.145 to 6.564)																					
	Neomatal encephalopathy due to	Birth prevalence - [36, 37) wks, [2500,	Mortality	Males		4.874	3.699																					
	Neomatal encephalopathy due to	Bith prevalence - [36, 37) wks, [2500,	Mortality	Females		4.609	3.898																					
	birth asphysia and trauma Neonatal encershaloreathy due to	3000) g Birth meyalence - [34, 36) wks. [3500.				(3.731 to 5.61) 18.024	(3.235 to 4.67) 3.657																					
Name<	birth asphysia and trauma	4000) g Rightmansharar (24, 36) mbs (2600	Mortality	Males		(12.279 to 25.547)	(2.838 to 4.675)																					
	birth asphysia and trauma	4000) g	Mortality	Females		(12.567 to 27.924)	(3.576 to 5.989)																					
	Neomatal encephalopathy due to birth asphysia and trauma	Birth prevalence - [37, 38) wks, [2500, 3000) g	Mortality	Males		3.306 (2.82 to 3.843)	3.194 (2.691 to 3.803)																					
	Neonatal encephalopathy due to birth asphysia and trauma	Birth prevalence - [37, 38) wks, [2500, 3000) g	Mortality	Females		2.991 (2.496 to 3.61)	3.242 (2.745 to 3.817)																					
	Neonatal encephalopathy due to birth asphysia and trauma	Birth prevalence - [40, 42) wks, [2500, 3000) g	Mortality	Males		3.771 (3.002 to 4.693)	3.175 (2.56 to 3.923)																					
	Neomatal encephalopathy due to birth asobycia and trauma	Birth prevalence - [40, 42) wks, [2500, 3000) g	Mortality	Females		3.244 (2.486 to 4.159)	3.228 (2.605 to 3.985)																					
	Neonatal encephalopathy due to	Birth prevalence - [38, 40) wks, [2500,	Mortality	Males		2.755	2.944																					
	Neomatal encephalopathy due to	S000) g Birth prevalence - [38, 40) wks, [2500,	Mortality	Females		2.376	2.938																					
	birth asphyxia and trianna Neonatal encephalopathy due to	3000) g Birth prevalence - [36, 37) wks, [3000,	M			(1.91 to 2.886) 3.774	(2.434 to 3.503) 2.466																					
Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name <	birth asphysia and trauma Neomtal encombalorative due to	3500) g Birth menalence - [36-37) wise [3000	stortaity	Males		(3.094 to 4.497) 3.73	(2.058 to 2.929) 2.715																					
	birth asphysia and tratana	3500) g	Mortality	Females		(2.981 to 4.646)	(2.277 to 3.218)																					
Network <t< td=""><td>Neomatal encephalopathy due to birth asphysia and trauma</td><td>Birth prevalence - [36, 37) wks, [4000, 4500) g</td><td>Mortality</td><td>Males</td><td></td><td>6.826 (5.212 to 9.045)</td><td>1.77 (1.491 to 2.082)</td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td></t<>	Neomatal encephalopathy due to birth asphysia and trauma	Birth prevalence - [36, 37) wks, [4000, 4500) g	Mortality	Males		6.826 (5.212 to 9.045)	1.77 (1.491 to 2.082)																					
Network	Neomatal encephalopathy due to birth asphysia and trauma	Birth prevalence - [36, 37) wks, [4000, 4500) g	Mortality	Females		7.269 (5.144 to 9.821)	2.177 (1.786 to 2.607)																					
Network Network Network Network Network Network Network	Neomatal encephalopathy due to birth asphysia and trauma	Birth prevalence - [36, 37) wks, [3500, 4000) g	Mortality	Males		4.544 (3.64 to 5.622)	2.057 (1.74 to 2.44)																					
Network Network Network Network Network Network Network Network Network Network <t< td=""><td>Neonatal encephalopathy due to birth asphysia and trauma</td><td>Birth prevalence - [36, 37) wks, [3500, 4000) g</td><td>Mortality</td><td>Females</td><td></td><td>4.662 (3.577 to 6.014)</td><td>2.398 (1.98 to 2.864)</td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td></t<>	Neonatal encephalopathy due to birth asphysia and trauma	Birth prevalence - [36, 37) wks, [3500, 4000) g	Mortality	Females		4.662 (3.577 to 6.014)	2.398 (1.98 to 2.864)																					
Name Name <th< td=""><td>Neomatal encephalopathy due to birth authorits and trauma</td><td>Birth prevalence - [37, 38) wks, [3000, 35000 m</td><td>Mortality</td><td>Males</td><td></td><td>2.007 (1.759 to 2.793)</td><td>1.888 (1.613 to 2.224)</td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td></th<>	Neomatal encephalopathy due to birth authorits and trauma	Birth prevalence - [37, 38) wks, [3000, 35000 m	Mortality	Males		2.007 (1.759 to 2.793)	1.888 (1.613 to 2.224)																					
	Neomatal encephalopathy due to	Birth prevalence - [37, 38) wks, [3000,	Mortality	Females		1.925	1.972																					
Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network Network	Neonatal encephalopathy due to	3500) g Birth prevalence - [37, 38) wks, [4000,	Martility	Males		3.28	1.335																					
Nationality	birth asphysia and trauma Neomatal encephalopathy due to	4500) g Birth prevalence - [37, 38) wks, [4000,				(2.596 to 4.133) 3.521	(1.171 to 1.532) 1.559																					
Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Name Nam Nam Name	birth asphysia and trauma Neomtol encombaloreathy due to	4500) g Birth menulence - [37-38) wise [3500	January	- unart		(2.649 to 4.5) 2.128	(1.333 to 1.835)																					
Name Name Name Name Name Name Reserved Reserved Name Name Name	birth asphysia and trauma	4000) g	Mortality	Males		(1.833 to 2.466)	(1.299 to 1.76)																					
Name Note Note Note Note Reserved Reserved Reserved Reserved Reserved Reserved Reserved Reser	Neonatal encephalopathy due to birth asphysia and trauma	Barth prevalence - [37, 38) wks, [3500, 4000) g	Mortality	Females		2.142 (1.694 to 2.67)	(1.411 to 1.961)																					
Matrix Matrix Matrix Matrix Matrix Matrix Matrix Matrix Matrix	Neonatal encephalopathy due to birth asphysia and traama	Bath prevalence - [40, 42) wks, [3000, 3500) g	Mortality	Males		1.436 (1.245 to 1.65)	1.47 (1.199 to 1.8)																					
Index Mathematical Mathematical Mathematical Mathematical Name Mathematical Mathematical Mathemat	Neomatal encephalopathy due to birth asphyxia and trauma	Birth prevalence - [40, 42) wks, [3000, 3500) g	Mortality	Females		1.326 (1.069 to 1.614)	1.465 (1.188 to 1.775)																					
Subscription Subscription Subscription Subscription Subscription Subscription Subscription Subscription Subscription Subscripinon Subscription Subsc	Neomatal encephalopathy doe to birth asphysia and tratama	Birth prevalence - [38, 40) wks, [3000, 3500) g	Mortality	Males		1.33 (1.155 to 1.53)	1.559 (1.305 to 1.851)																					
Index Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide Import Provide <	Neonatal encephalopathy due to birth asphysia and trauma	Birth prevalence - [38, 40) wks, [3000, 3500) g	Mortality	Females		1.224 (1.0 to 1.492)	1.564 (1.304 to 1.847)																					
	Neonatal encephalopathy due to	Birth prevalence - [38, 40) wks, [4000, 46000 -	Mortality	Males		1.787	1.175																					
Name Ord Ord <td>Neomatal encephalopathy due to</td> <td>Birth prevalence - [38, 40) wks, [4000,</td> <td>Mortality</td> <td>Females</td> <td></td> <td>1.877</td> <td>1.224</td> <td></td>	Neomatal encephalopathy due to	Birth prevalence - [38, 40) wks, [4000,	Mortality	Females		1.877	1.224																					
Name Only (M) Org Org Org Org Name Second Second </td <td>berth asphysia and trauma Neonatal encephalopathy due to</td> <td>4500) g Birth prevalence - [38, 40) wks, [3500,</td> <td>Mastality</td> <td>Maler</td> <td></td> <td>(1.467 to 2.388) 1.785</td> <td>(1.022 to 1.465) 1.173</td> <td></td>	berth asphysia and trauma Neonatal encephalopathy due to	4500) g Birth prevalence - [38, 40) wks, [3500,	Mastality	Maler		(1.467 to 2.388) 1.785	(1.022 to 1.465) 1.173																					
Non-granual	birth asphysia and trauma Neonatal encershaloreathy due to	4000) g Birth mevalence - [38, 40) wks, [3500,				(1.478 to 2.147) 1.892	(1.0 to 1.377) 1.23																					
Name Main Main Main Main Main Main Name Name Name Name Name Name Name Name Name Name N	birth asphysia and trauma	4000) g	Mortality	Females		(1.481 to 2.352)	(1.03 to 1.46)																					
And conduction to specify and conduction to specify and conductionAnd conduction to specify and conductionAnd conduction to specify and conductionAnd conduction to specify and conductionName and conduction to specify and conduction to specify and conductionName and conduction to specify and conduction to specify and conductionName and conduction to specify and conductionName and conduction to specify and conductionName and conduction to specify and conduction to specify and conductionName and conduction to specify and conductionName and conduction to specify and conductionName and conduction to specify and conduction to specify and conductionName and conduction to specify and conductionName and conduction to specify and conductionName and conduction to specify and conduction to specify and conductionName and conduction to specify and conductionName and conduction to specify and conductionName and conduction to specify and conduction to specify and conductionName and conduction to specify and conductionName and conduction to specify and conductionName and conduction to specify and conduction to specify and conductionName and conduction to specify and conductionName and conduction to specify and conduction to specify and conductionName and conduction to specify and conduction to specify and conductionName and conduction to specify and conductionName and conduction to specify and conduction	birth asphysia and trauma	4000) g	Mortality	Males		(1.0 to 1.0)	(1.0 to 1.046)																					
Anotamelande and anotamelande anotamelan	Neonatat encephalopathy due to birth asphysia and traama	north prevalence - [40, 42) wks, [3500, 4000) g	Mortality	Females		1.002 (1.0 to 1.013)	1.001 (1.0 to 1.006)																					
And mandmand matrix And matrix And matrix And matrix And matrix Main and matrix Main and Ma	Neonatal encephalopathy due to birth asphyxia and trauma	Birth prevalence - [40, 42) wks, [4000, 4500) g	Mortality	Males		1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																					
And complexing free billing Subscription Subscription Subscription Subscription Name combining Subscription Subscription Subscripiiiiiiiiiiiiiiiii	Neomatal encephalopathy due to birth asphysia and tratama	Birth prevalence - [40, 42) wks, [4000, 4500) g	Mortality	Females		1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																					
August angle Aller	Neonatal encephalopathy due to birth asphysia and trauma	Birth prevalence - [28, 30) wks, [2000, 2500) g	Mortality	Males		117.172 (83.895 to 158.056)	27.726 (21.877 to 34.972)																					
Next and analysing of the strength of the stre	Neonatal encephalopathy due to	Birth prevalence - [28, 30) wks, [2000, 2500) #	Mortality	Females		121.682 (84.349 to 171.277)	33.983																					
Integration State	Neomatal encephalopathy due to	Birth prevalence - [28, 30) wks, [2500,	Mortality	Males		77.948	16.608																					
International methylinia and specific and speci	birth asphyxia and trauma Neonatal encephalopathy due to	3000) g Birth prevalence - [28, 30) wks, [2500,	Mastality	Females		(54.687 to 105.047) 79.193	(12.653 to 21.188) 20.387																					
Ising Sill Sill Manual Opposite	birth asphysia and trauma Neomatal on orbital orating the term	3000) g Birth menalence - [28-30) wise 13000	sucrtainty	remates		(54.099 to 112.236) 42.199	(15.089 to 26.434) 10.087																					
New York Company and Company a	birth asphysia and trauma	3500) g Bish mushem (28, 30) ada (2000)	Mortality	Males		(29.891 to 57.227)	(7.777 to 13.056)																					
	birth asphysia and traama	3500) g	Mortality	Females		(28.25 to 63.209)	(9.084 to 15.544)	l																				

			pontition involtor.										A	ges											
Risk - Outcome Category / Units M	orbidity / Mortality	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Neonatal encephalopathy due to Birth prevalence - [30, 32] wks, [2500, kink andward and another 2000 a	Mortality N	ales	58.722	12.115	1																				-
Neonatal encephalopathy due to Birth prevalence - (30, 32) wks, (2500,	Mortality Fe	nales	59.522	15.364																					
berth asphysia and tratama 3000) g Neonatal encephaloputhy due to Birth prevalence - (30, 32) wks, (3000,	Martality N	ala.	(42.058 to 82.793) 45.67	(11.936 to 19.581) 8.381																					
birth asphysia and trauma 3500) g Neomatal encephalopathy dae to Birth prevalence - [30, 32] wks, [3000,			(32.014 to 65.531) 46.104	(6.362 to 10.842) 10.506																					
birth asphysia and trauma 3500) g Necessital encembracements due to Birth merculence - (30-32) why (3500	sumarry Pe	12225	(30.243 to 66.207) 36.334	(8.041 to 13.513) 5.698																					
birth aphysia and trauma 4000) g	Mortality N	ales	(21.558 to 54.813)	(4.293 to 7.349)																					
Neonaial encephalopathy dae to Bath prevalence - [30, 32] wks, [5500, birth asphysia and tratama 4000) g	Mortality Fe	nales	37.931 (22.692 to 61.276)	6.892 (5.175 to 9.058)																					
Neoenital encephalopathy due to Birth prevalence - (32, 34) wks, (3000, birth asphysia and traama 3500) g	Mortality N	ales	34.016 (23.515 to 48.37)	6.577 (4.934 to 8.436)																					
Neonatal encephalopathy dae to Birth prevalence - [32, 34) wks, [3000, birth asphysia and trauma 3500) g	Mortality Fe	nales	34.585 (22.909 to 49.754)	8.314 (6.362 to 10.789)																					
Necental encephalopathy due to Birth prevalence - [32, 34] wks, [3500, birth asphysia and trauma 4000) g	Mortality N	ales	36.248 (23.158 to 54.67)	5.068 (3.761 to 6.741)																					
Neomatal encephalopathy due to Birth prevalence - [32, 34] wks, [3500, birth authorits and tracers 40000 p	Mortality Fe	nales	38.098 (23.301 to 59.429)	6.476 (4.666 to 8.689)																					
Neonatal encephalopathy due to Birth prevalence - (36, 37) wks, [1000,	Mortality N	ales	166.686	57.535																					
Neonatal encephalopathy due to Birth prevalence - [36, 37] wks, [1000,	Mortality Fe	nales	169.725	63.564																					
birth asphysia and trauma 1500) g Neomatal encephalopathy dae to Birth prevalence - [38, 40) wks, [1000,	M		(119.017 to 229.008) 174.066	(50.068 to 80.703) 57.966																					
birth asphysia and trauma 1500) g Neomini on-ombulements due to Birth meruslence - 138–400 with 11000	subrainy N	105	(125.125 to 232.507) 171.557	(44.393 to 73.241) 65 208																					
birth asphysia and trauma 1500) g	Mortality Fe	nales	(121.585 to 237.047)	(48.821 to 84.308)																					
Neonaial encephalopathy dae to Bath prevalence - [38, 40] wio, [1500, birth asphysia and traama 2000) g	Mortality N	ales	67.302 (49.547 to 89.055)	25.206 (20.365 to 31.168)																					
Neonatal encephalopathy due to Birth prevalence - [38, 40) wks, [1500, birth apphysia and trauma 2000) g	Mortality Fe	nales	62.19 (45.884 to 83.445)	28.05 (22.625 to 35.139)																					
Neonatal encephalopathy dae to Birth prevalence - [40, 42] wks, [1500, birth asphysia and trauma 2000) g	Mortality N	iales	76.673 (56.177 to 102.468)	25.785 (19.387 to 34.168)																					
Neomatal encephalopathy due to Birth prevalence - [40, 42] wks, [1500, birth asphysia and trauma 2000) g	Mortality Fe	nales	70.411 (49.221 to 97.952)	29.113 (21.355 to 38.272)																					
Necessatal sepsis and other necessatal Birth prevalence - [0, 24] wks, [0, 500) g	Mortality N	ales	1564.792 (1056.542 to	618.595																					
Neonatal sepsis and other neonatal Birth merculence - 10, 24) wks. 10, 500) a	Mortality Fe	nales	2116.062) 1600.122 (1050.664 to	713.571	,																				
infections Neonatal sepsis and other neonatal Birth prevalence - [0, 24) wks, [500, 1000)	Manufata N		1155.815	(526.178 to 921.018) 457.5)																				
infections g Neoental servis and other neonatal Birth prevalence - 10, 24) wks. (500, 1000)	analany a		(825.412 to 1506.837 1169.123) (352.552 to 573.483) 515.406)																				
infections g	Mortality Fe	nalex	(802.003 to 1617.979	(396.713 to 641.541)																				
reconsta separa and other neonatal institu prevalence - [24, 26] wes, [500, 1000] infections g	Mortality N	ales	(723.748 to 1244.265	(363.03 to 534.695)																					
Neonital sepsis and other neonatal Birth prevalence - (24, 26) wks, (500, 1000) infections g	Mortality Fe	nales	947.143 (702.662 to 1237.093	487.549 (387.307 to 603.498))																				
Neonatal sepsis and other neonatal Birth prevalence - [26, 28) wks, [500, 1000) infections g	Mortality N	ales	497.817 (377.617 to 648.547)	330.886 (261.438 to 401.709	, ,																				
Neomatal sepsis and other neonatal Birth prevalence - [26, 28) wks, [500, 1000) infections g	Mortality Fe	nales	483.682 (354.946 to 629.517)	344.618 (274.427 to 419.864	,																				
Neonatal sepsis and other neonatal Birth prevalence - [30, 32] wks, [500, 1000) infections	Mortality N	ales	236.614 (163.821 to 324.502)	149.995 (117.866 to 188.368	,																				
Neonatal sepsis and other neonatal Birth prevalence - [30, 32] wks, [500, 1000)	Mortality Fe	nales	229.197 (157.606 to 317.194)	152.117 (120.779 to 190.583																					
Necential sepsis and other neconatal Birth prevalence - [28, 30] wko, [500, 1000]	Mortality N	ales	297.629	216.995	<i>.</i>																				
infections g Neonatal sepsis and other neonatal Birth prevalence - [28, 30] wks, [500, 1000)	Manufito En		(214.953 to 396.586) 281.056	219.884	*																				
infections g Neonatal sepsis and other neonatal Birth prevalence - [26, 28] wks, [1000,	Marine N		(198.176 to 386.635) 267.91	(174.264 to 272.704) 164.167)																				
infections 1500) g Neomini and other neominal Brith menulonce - 126-283 why (1000	Mortany N	105	(210.177 to 332.92) 266 509	(132.898 to 200.569) 174.222)																				
infections 1500) g	Mortality Fe	nales	(197.461 to 346.932)	(137.431 to 217.349)																				
Neonatal sepors and other neonatal infections 1500) g	Mortality N	ales	142.056 (98.086 to 197.774)	52.86 (42.914 to 64.617)																					
Neoental sepsis and other neonatal Birth prevalence - (34, 36) wks, [1000, infections 1500) g	Mortality Fe	nales	141.899 (95.864 to 197.656)	57.421 (46.452 to 71.339)																					
Neonatal sepsis and other neonatal infections Birth prevalence - [28, 30) wks, [1500, 2000) g	Mortality N	ales	127.966 (97.178 to 167.026)	50.018 (40.539 to 61.919)																					
Neomatal sepsis and other neonatal Birth prevalence - [28, 30] wks, [1500, infections 2000) g	Mortality Fe	nales	130.924 (96.513 to 172.188)	57.275 (46.36 to 70.038)																					
Neonatal sepsis and other neonatal Birth prevalence - [28, 30) wks, [1000, infections 1500) r	Mortality N	ales	158.563 (120.99 to 204.947)	103.32 (83.486 to 127.144)																					
Neonatal sepsis and other neonatal Birth prevalence - [28, 30] wks, [1000, infections	Mortality Fe	nales	153.905 (112.327 to 200.786)	107.529 (86.954 to 131.78)																					
Neonatal sepsis and other neonatal Birth prevalence - (32, 34) wks, [1000,	Mortality N	ales	117.142	53.185																					
infections 1500) g Neonatal sepsis and other neonatal Birth prevalence - [32, 34) wks, [1000,	Manufator En		(81.354 to 161.101) 115.171	(43.049 to 66.274) 56.034																					
infections 1500) g Neonatal sepsis and other neonatal Birth prevalence - [30, 32) wks, [1000,	Mandan N		(79.363 to 159.206) 119.308	(45.982 to 68.36) 67.163																					
infections 1500) g Neonatal sepsis and other neonatal Birth prevalence - [30, 32) wks. [1000.	Marin -		(87.769 to 160.885) 115.448	(54.863 to 82.638) 69.14																					
infections 1500) g	Mortality Fe	nalex	(84.272 to 156.425)	(55.873 to 85.012)																					
infections and on an one and an one an on one an	Mortality N	intes	(46.159 to \$3.484)	(20.066 to 29.406)																					
Neonatal sepors and other neonatal Barth prevalence - [37, 38] wks, [1500, infections 2000) g	Mortality Fe	nales	59.988 (43.974 to 79.053)	26.719 (21.746 to 32.816)																					
Neonatal sepsis and other neonatal infections Birth prevalence - [36, 37] wks, [1500, 2000) g	Mortality N	ales	60.218 (43.669 to 82.48)	23.031 (18.793 to 28.483)																					
Neoental sepsis and other neonatal Birth prevalence - [36, 37] wks, [1500, infections 2000) g	Mortality Fe	nales	58.527 (42.172 to 80.557)	25.143 (20.331 to 30.566)																					
Neonatal sepsis and other neonatal Birth prevalence - [30, 32] wks, [2000, infections 2500) r	Mortality N	ales	67.971 (50.354 to 88.935)	18.03 (14.621 to 22.103)																					
Neomatal sepsis and other neomatal Birth prevalence - [30, 32] wks, [2000, 2600 p	Mortality Fe	nales	69.383 (40.109 to 04.593)	22.069																					
Neonatal sepsis and other neonatal Birth prevalence - [30, 32] wks, [1500,	Mortality N	ales	77.369	31.079																					
meetions 2000) g Neonatal sepsis and other neonatal Birth prevalence - [30, 32) wks, [1500,	Marality	maler	(59.702 to 99.232) 76.134	(25.786 to 36.724) 34.756																					
infections 2000) g Neonatal sepsis and other neonatal Birth prevalence - 134, 36) wks. /1 500	watany Pe		(56.885 to 100.996) 55.555	(28.764 to 41.849) 21.346																					
infections 2000) g Neomial series and other neomatal Brith menalence - [34, 36, 11000	Mortality N	ases	(39.553 to 75.104) \$4.335	(17.677 to 26.143) 23.046																					
infections 2000) g	Mortality Fe	nales	(38.617 to 75.24)	(18.743 to 28.287)	1																				
reconstant seriors and other neonatal Borth prevalence - [32, 34] wks, [1500, infections 2000) g	Mortality N	iales	57.155 (42.484 to 73.651)	25.114 (19.028 to 27.915)	1																				
Necental sepsis and other neonatal Birth prevalence - [32, 34] wks, [1500, infections 2000) g	Mortality Fe	nales	56.101 (39.794 to 76.295)	25.149 (20.615 to 30.388)																					
Neonatal sepsis and other neonatal infections 2500) g	Mortality N	inles	37.444 (29.026 to 48.227)	12.233 (10.252 to 14.477)																					
Neonatal sepsis and other neonatal infections 2500) g	Mortality Fe	nales	36.874 (26.658 to 49.653)	14.384 (12.095 to 17.03)																					
Neonatal sepsis and other neonatal infections 2500 n	Mortality N	iales	18.092 (13.292 to 23.719)	9.23 (7.037 to 11.454)																					
		I	(1 (1																				

											Aş			iges												
Risk - Outcome Category / Units	Morbidity / Mortality Sex	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years	
Neonatal sepsis and other neonatal infections 2500) g	Mortality Femal	DS	15.574 (11.516 to 20.778)	9.975 (7.82 to 12.46)																						
Neonatal sepsis and other neonatal Birth prevalence - [38, 40) wks, [2000, infections 2500) r	Mortality Male	ĸ	13.104 (9.829 to 16.99)	8.198 (6.786 to 9.959)																						
Neomatal serpsis and other neonatal Birth prevalence - [38, 40) wks, [2000, 2000] -	Mortality Female	D	11.308	8.577																						
Neonatal sepsis and other neonatal Birth prevalence - [32, 34] wks, [2500,	Mortality Male		33.063	8.441																						
infections 5000) g Neonatal sepsis and other neonatal Birth prevalence - [32, 34) wks, [2500,	Mantality Email		(24.393 to 43.503) 32.812	(6.822 to 10.431) 10.398																						
infections 3000) g Necessital service and other reconstal Birth recondence - [34, 36) who [2000	storiany rema-	5	(23.439 to 45.567) 21.925	(8.227 to 13.042) 9.367																						
infections 2500 g	Mortality Male	K.	(16.305 to 29.433)	(7.859 to 11.112)																						
infections 2500) g	Mortality Female	D)	(15.657 to 28.761)	(8.548 to 12.273)																						
Necential sepsis and other neconatal infections 2500) g	Mortality Male	,	13.0 (10.102 to 16.456)	8.096 (6.724 to 9.676)																						
Neonatal sepsis and other neonatal Birth prevalence - [37, 38) wks, [2000, infections 2500) g	Mortality Female	DS	11.563 (8.805 to 15.11)	8.467 (6.994 to 10.342)																						
Neonatal sepsis and other neonatal infections 2500) g	Mortality Male		14.401 (10.789 to 18.654)	8.221 (6.917 to 9.923)																						
Networked Service and other networked Birth prevalence - [36, 37] wks, [2000, infections 2500) g	Mortality Femal	DN	13.513 (9.817 to 17.942)	8.654 (7.215 to 10.369)																						
Neonatal sepsis and other neonatal Birth prevalence - [34, 36] wks, [2500, infections 3000 n	Mortality Male	ĸ	13.419 (10.387 to 16.819)	5.562 (4.646 to 6.696)																						
Neomatal servis and other neonatal Birth prevalence - [34, 36] wks, [2500, 2000 -	Mortality Female	D	13.266	6.395																						
Neonatal sepsis and other neonatal Birth prevalence - [34, 36) wks, [4000,	Mortality Male		23.096	2.895																						
infections 4500) g Neonatal sepsis and other neonatal Birth prevalence - [34, 36) wks, [4000,	Mantality Email		(14.708 to 35.098) 25.038	(2.245 to 3.716) 3.778																						
infections 4500) g Neonatal sensis and other neonatal Birth prevalence - [34, 36) wks. [3000.	Section 2		(15.763 to 37.255) 14.006	(2.855 to 4.925) 4.322																						
infections 3500) g	Mortality Male	·	(10.222 to 18.478)	(3.449 to 5.338)																						
infections aport and outer monantal and prevalence - (199, 50) was, (2000,	Mortality Female	D)	(10.269 to 20.114)	(4.145 to 6.564)																						
Necessial separs and other necessial infections 3000) g	Mortality Male	K.	4.8/4 (4.014 to 5.713)	3.699 (3.134 to 4.374)																						
Neonatal sepsis and other neonatal Birth prevalence - [36, 37] wks, [2500, infections 3000) g	Mortality Female	DS	4.609 (3.731 to 5.61)	3.898 (3.235 to 4.67)																						
Neomatal sepsis and other neonatal Birth prevalence - [34, 36) wks, [3500, infections 4000) g	Mortality Male		18.024 (12.279 to 25.547)	3.657 (2.838 to 4.675)																						
Necentral serpsis and other necentral Birth prevalence - [34, 36) wks, [3500, 4000) r	Mortality Female	5	19.263 (12.567 to 27.924)	4.634 (3.576 to 5.989)																						
Neomatal sepsis and other neomatal Birth prevalence - [37, 38) wks, [2500, infections 30000 s	Mortality Male		3.306 (2.82 to 3.843)	3.194 (2.691 to 3.893)																						
Network and other neonatal Birth prevalence - [37, 38) wks, [2500,	Mortality Femal	DN	2.991	3.242																						
Neonatal sepsis and other neonatal Birth prevalence - [40, 42] wks, [2500,	Mortality Male		3.771	3.175																						
infections 3000) g Neonatal sepsis and other neonatal Birth prevalence - [40, 42] wks, [2500,			(3.002 to 4.693) 3.244	(2.56 to 3.923) 3.228																						
infections 3000) g Neonatal servis and other neonatal Birth prevalence - (38, 40) wks. (2500,	Mortany Pena	24	(2.486 to 4.159) 2.755	(2.605 to 3.985) 2.944																						
infections 3000) g	Mortality Male	·	(2.274 to 3.309)	(2.44 to 3.548)																						
infections aport and outer monantal and prevalence - [.36, 40] was, [2.500, 3000) g	Mortality Female	D)	(1.91 to 2.886)	(2.434 to 3.503)																						
Neonatal sepsis and other neonatal infections 3500) g	Mortality Male		3.774 (3.094 to 4.497)	2.466 (2.058 to 2.929)																						
Neonatal sepsis and other neonatal Birth prevalence - [36, 37) wks, [3000, infections 3500) g	Mortality Female	DS	3.73 (2.981 to 4.646)	2.715 (2.277 to 3.218)																						
Neonatal sepsis and other neonatal infections Birth prevalence - [36, 37] wks, [4000, 4500) g	Mortality Male		6.826 (5.212 to 9.045)	1.77 (1.491 to 2.082)																						
Networked Service and other networked Birth prevalence - [36, 37] wks, [4000, 4500) g	Mortality Femal	DN	7.269 (5.144 to 9.821)	2.177 (1.786 to 2.607)																						
Neomatal sepsis and other neomatal Birth prevalence - [36, 37] wks, [3500, infections 40000 s	Mortality Male		4.544	2.057 (1.74 to 2.44)																						
Neomatal servis and other neonatal Birth prevalence - [36, 37] wks, [3500, infection:	Mortality Female	D	4.662	2.398																						
Networking Service and other networking Birth prevalence - [37, 38) wks, [3000,	Mortality Male		2.007	1.888																						
infections 3500) g Neonatal sepsis and other neonatal Birth prevalence - [37, 38) wks, [3000,	Mantality Email		(1.759 to 2.293) 1.925	(1.613 to 2.224) 1.972																						
infections 3500) g Neonatal sepsis and other neonatal Birth prevalence - [37, 38) wks, [4000,			(1.582 to 2.328) 3.28	(1.669 to 2.313) 1.335																						
infections 4500) g Neonatal servis and other neonatal Birth prevalence - (37, 38) wks. (4000.	Mortany Make	K.	(2.596 to 4.133) 3.521	(1.171 to 1.532) 1.559																						
infections 4500) g	Mortality Female	DA	(2.649 to 4.5)	(1.333 to 1.835)																						
vectorial separation other neonatal in the prevalence - (37, 38) woo, (5500, infections 4000) g	Mortality Male		(1.833 to 2.466)	(1.299 to 1.76)																						
Neonatal separs and other neonatal Barth prevalence - [37, 38] wio, [5200, infections 4000) g	Mortality Female	DS	2.142 (1.694 to 2.67)	1.661 (1.411 to 1.961)																						
Neonatal sepsis and other neonatal Birth prevalence - [40, 42] wks, [3000, infections 3500) g	Mortality Male	ĸ	1.436 (1.245 to 1.65)	1.47 (1.199 to 1.8)																						
Neonatal sepsis and other neonatal Birth prevalence - [40, 42] wks, [3000, infections 3500) g	Mortality Female	25	1.326 (1.069 to 1.614)	1.465 (1.188 to 1.775)																						
Neonatal sepsis and other neonatal infections 3500) g	Mortality Male	,	1.33 (1.155 to 1.53)	1.559 (1.305 to 1.851)																						
Neonatal sepsis and other neonatal Birth prevalence - [38, 40] wks, [3000, infections 3500) g	Mortality Femal	25	1.224 (1.0 to 1.492)	1.564 (1.304 to 1.847)																						
Neonatal sepsis and other neonatal Birth prevalence - [38, 40) wks, [4000, infections 4500) g	Mortality Male	,	1.787 (1.453 to 2.182)	1.175 (1.005 to 1.371)																						
Neonatal sepsis and other neonatal informations 4000 a	Mortality Femal	5	1.877	1.224 (1.022 to 1.465)																						
Networking and other networking Birth prevalence - [38, 40] wks, [3500,	Mortality Male		1.785	1.173																						
infections 4000) g Neonatal sepsis and other neonatal Birth prevalence - [38, 40) wks, [3500,	Montality Formal		(1.478 to 2.147) 1.892	(1.0 to 1.377) 1.23																						
infections 4000) g Neonatal sepsis and other neonatal Birth prevalence - [40, 42] wks. 13500.	March		(1.481 to 2.352) 1.0	(1.03 to 1.46) 1.003																						
infections 4000) g Neomini and other neominal Birth menulence - (40, 47) who (3500)	Mortality Male		(1.0 to 1.0)	(1.0 to 1.046)																						
infections 4000) g	Mortainty Female	25	(1.0 to 1.013)	(1.0 to 1.006)																						
www.neura report and orner neonatal Birth prevalence - [40, 42] wks, [4000, infections 4500) g	Mortality Male	K.	1.0 (1.0 to 1.0)	(1.0 to 1.0)																						
Necentral sepsis and other neonatal Birth prevalence - [40, 42] wks, [4000, infections 4500) g	Mortality Female	2N	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																						
Neonatal sepsis and other neonatal Birth prevalence - [28, 30] wks, [2000, infections 2500) g	Mortality Male	ĸ	117.172 (83.895 to 158.056)	27.726 (21.877 to 34.972)																						
Neonatal sepsis and other neonatal infections 2500) g	Mortality Femal	DS	121.682 (84.349 to 171.375)	33.983 (26.101 to 43.404)																						
Neonatal sepsis and other neonatal Birth prevalence - [28, 30) wks, [2500, infections 3000) g	Mortality Male		77.948 (54.687 to 105.047)	16.608 (12.653 to 21.188)																						
Neonatal sepsis and other neonatal Birth prevalence - [28, 30) wks, [2500, infections 3000) g.	Mortality Femal	DS	79.193 (54.099 to 112.236)	20.387 (15.089 to 26.434)																						
Neonatal sepsis and other neonatal informations 2000 a	Mortality Male		42.199	10.082																						
Neonatal sepsis and other neonatal Neonatal sepsis and other neonatal	Mortality Female	zs.	42.551	11.989																						
intections 3500) g		1	(28.25 to 63.209)	(v.084 to 15.544)	1																					
	and by age and set to card outcom				a and substang.									A	lges											
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Risk - Outcome	Category / Units	Morbidity / Mortality	All-a	ige 0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Neonatal sepsis and other neonat	al Birth prevalence - [30, 32) wks, [2500,	Mortality M	ales .	58.722	12.115																					
infections Neonatal sepsis and other neonat	3000) g al Birth prevalence - [30, 32) wks, [2500,	Martality Fr	makes	(42.419 to 78.8) 59.522	5) (9.518 to 15.521) 15.364																					
infections Noomtal service and other records	3000) g al. Birth menalence - [30, 37) wks [3000	,		(42.058 to 82.7) 45.67	3) (11.936 to 19.581) 8 381	0																				
infections	3500) g	Mortality N	ales	(32.014 to 65.5)	 (6.362 to 10.842) 																					
infections	a Binn previaence - [50, 52) wis, [5000, 3500) g	Mortality Fe	nales	(30.243 to 66.2)	7) (8.041 to 13.513)																					
Neonatal sepsis and other neonat- infections	al Birth prevalence - [30, 32) wks, [3500, 4000) g	Mortality N	ales	36.334 (21.558 to 54.8	5.698 (4.293 to 7.349)																					
Neomatal sepsis and other neonat- infections	al Birth prevalence - [30, 32) wks, [3500, 4000) n	Mortality Fe	nales	37.931 (22.692 to 61.2)	6.892 (5.175 to 9.058)																					
Neomatal sepsis and other neonat	al Birth prevalence - [32, 34) wks, [3000,	Mortality N	ales	34.016	6.577																					
infections Neomatal sepsis and other neonat	3500) g al Birth prevalence - [32, 34) wks, [3000,	Mastelia E-	-	(23.515 to 48.3 34.585) (4.934 to 8.436) 8.314																					
infections Noomtal service and other records	3500) g al. Birth menalence - [32: 34) wks [3500	January 14		(22.909 to 49.7) 36.248	4) (6.362 to 10.789) 5.068																					
infections	4000) g	Mortality N	ales	(23.158 to 54.6) (3.761 to 6.741)																					
Neonatal sepsos and other neonati infections	al Batth prevalence - [32, 34) wks, [3500, 4000) g	Mortality Fe	nales	(23.301 to 59.4)	6.476 (4.666 to 8.689)																					
Neonatal sepsis and other neonat- infections	al Birth prevalence - [36, 37) wks, [1000, 1500) g	Mortality N	ales	166.686 (118.487 to 222.5	57.535 (45.999 to 71.742)																					
Neonatal sepsis and other neonat- infections	al Birth prevalence - [36, 37) wks, [1000, 1500) n	Mortality Fe	nales	169.725 (119.017 to 229.0	63.564 (50.068 to 80.703)																					
Neomatal sepsis and other neonat-	al Birth prevalence - [38, 40) wks, [1000, 15000 -	Mortality N	ales	174.066	57.966																					
Neonatal sepsis and other neonat	al Birth prevalence - [38, 40) wks, [1000,	Martality Fr	males	171.557	65.208	,																				
infections Noomtal service and other records	1500) g al. Birth menalence - [38, 40) wise [1500]	,		(121.585 to 237.0 67.302	47) (48.821 to 84.308) 25.206	0																				
infections	2000) g	Mortality N	ales	(49.547 to 89.0)	5) (20.365 to 31.168)	0																				
Neonatal sepsos and other neonati infections	al Barth prevalence - [38, 40) wks, [1500, 2000) g	Mortality Fe	nales	62.19 (45.884 to 83.4-	28.05 (22.625 to 35.139)																					
Neonatal sepsis and other neonat- infections	al Birth prevalence - [40, 42) wks, [1500, 2000) g	Mortality N	ales	76.673 (56.177 to 102.4	25.785 (19.387 to 34.168)																					
Neonatal sepsis and other neonat- infections	al Birth prevalence - [40, 42) wks, [1500, 2000) n	Mortality Fe	nales	70.411 (49.221 to 97.9)	29.113 (21.355 to 38.272																					
Hemolytic disease and other	Birth prevalence - [0, 24) wks, [0, 500) g	Mortality M	ales	1564.792 (1056.542 to	618.595																					
neonatal jaundace Hemolytic disease and other	N. 4			2116.062) 1600.122	(458.842 to 812.92 713.571	1)																				
neonatal jaundice	Birth previaence - [0, 24) wick, [0, 500) g	Mortany Pe	nases.	1050.884 50	(526.178 to 921.01)	8)																				
neonatal jaundice	8	Mortality N	ales	(825.412 to 1506.	37) (352.552 to 573.48)	3)																				
Hemolytic disease and other neonatal jaundice	Birth prevalence - [0, 24) wks, [500, 1000) 8	Mortality Fe	nales	1169.123 (802.003 to 1617.	515.406 (396.713 to 641.54	1)																				
Hemolytic disease and other neonatal jaundice	Birth prevalence - [24, 26) wks, [500, 1000) 8	Mortality M	ales	955.583 (723.748 to 1244.	443.357 (363.03 to 534.695	i)																				
Hemolytic disease and other	Birth prevalence - [24, 26) wks, [500, 1000)	Mortality Fe	nales	947.143	487.549	o.																				
Hemolytic disease and other	o Birth prevalence - [26, 28) wks, [500, 1000)	Mortality M	ales	497.817	330.886																					
neonatal jaundace Hemolytic disease and other	8 Birth prevalence - [26, 28) wks, [500, 1000)	M		483.682	(261.438 to 401.70 344.618	9)																				
neonatal jaundice	8 Rishmanlana (20.22) ada (500.1000)	Mortany Pe	nases.	(354.946 to 629.) 236.614	17) (274.427 to 419.86-	4)																				
neonatal jaundice	8	Mortality N	ales	(163.821 to 324.)	02) (117.866 to 188.36)	8)																				
Hemolytic disease and other neonatal jaundice	Birth prevalence - [30, 32) wks, [500, 1000) 8	Mortality Fe	nales	229.197 (157.606 to 317.)	152.117 (120.779 to 190.58)	3)																				
Hemolytic disease and other neonatal jaundice	Birth prevalence - [28, 30) wks, [500, 1000) g	Mortality M	ales	297.629 (214.953 to 396.5	216.995 (173.321 to 271.46	6)																				
Hemolytic disease and other recorded immediate	Birth prevalence - [28, 30) wks, [500, 1000)	Mortality Fe	nales	281.056 (198.176 to 386 t	219.884 (174.264 to 272.20)	a																				
Hemolytic disease and other	o Birth prevalence - [26, 28) wks, [1000,	Mortality M	iales	267.91	164.167	-																				
neonatal jaundice Hemolytic disease and other	1500) g Birth prevalence - [26, 28) wks, [1000,	Mastalia E-		(210.177 to 332. 266.509	 (132.898 to 200.56) 174.222 	9)																				
neonatal jaundice	1500) g Rish muschen: (24, 26) who (1000)	January 14		(197.461 to 346.5	32) (137.431 to 217.34) \$2.96	9)																				
neonatal jaundice	1500) g	Mortality N	ales	(98.086 to 197.7	(42.914 to 64.617))																				
Hemolytic disease and other neonatal jaundice	Birth prevalence - [34, 36) wks, [1000, 1500) g	Mortality Fe	nales	141.899 (95.864 to 197.6	57.421 (46.452 to 71.339)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [28, 30) wks, [1500, 2000) g	Mortality M	iales	127.966 (97.178 to 167.0	50.018 (40.539 to 61.919)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [28, 30) wks, [1500, 2000) #	Mortality Fe	nales	130.924 (96.513 to 172.1	57.275 8) (46.36 to 70.038)																					
Hemolytic disease and other	Birth prevalence - [28, 30) wks, [1000, 1500) -	Mortality M	iales	158.563	103.32																					
Hemolytic disease and other	Birth prevalence - [28, 30) wks, [1000,	Matulity Fr	makes	153.905	107.529	.)																				
neonatal jaundice Hemolytic disease and other	1500) g Birth prevalence - (32, 34) wks. (1000,			(112.327 to 200.) 117.142	36) (86.954 to 131.78) 53.185)																				
neonatal jaundice	1500) g	Mortality N	ales	(81.354 to 161.1	(43.049 to 66.274	0																				
Hemolytic disease and other neonatal jaundice	Barth prevalence - [32, 34) wks, [1000, 1500) g	Mortality Fe	nales	(79.363 to 159.2	56.034 (45.982 to 68.36)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [30, 32) wks, [1000, 1500) g	Mortality N	iales	119.308 (87.769 to 160.8	67.163 (54.863 to 82.638))																				
Hemolytic disease and other neonatal jaundice	Birth prevalence - [30, 32) wks, [1000, 1500) g	Mortality Fe	nales	115.448 (84.272 to 156.4	69.14 5) (55.873 to 85.012																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [37, 38) wks, [1500, 2000) n	Mortality M	ales	62.972 (46.159 to 83.4)	24.148 (20.066 to 29.406)																					
Hemolytic disease and other	Birth prevalence - [37, 38) wks, [1500,	Mortality Fe	nales	59.988	26.719																					
neonatal jaundace Hemolytic disease and other	2000) g Birth prevalence - [36, 37) wks, [1500,	Mastelia N	-	(43.974 to 79.0) 60.218	3) (21.746 to 32.816) 23.031)																				
neonatal jaundice	2000)g Rishmundara (36, 27)mln (1600	stortany s	100	(43.669 to 82.4	(18.793 to 28.483) 26.142																					
neonatal jaundice	2000) g	Mortality Fe	nales	(42.172 to 80.5)	7) (20.331 to 30.566)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [30, 32) wks, [2000, 2500) g	Mortality M	ales	67.971 (50.354 to 88.9)	18.03 (14.621 to 22.103)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [30, 32) wks, [2000, 2500) g	Mortality Fe	nales	69.383 (49.108 to 94.51	22.069 (17.836 to 27.163)																					
Hemolytic disease and other recorded issuedice	Birth prevalence - [30, 32) wks, [1500, 2000) #	Mortality M	ales	77.369 (59.702 to 99.2)	31.079 (25.786 to 36.724)																					
Hemolytic disease and other	Birth prevalence - [30, 32) wks, [1500, 2000) -	Mortality Fe	nales	76.134	34.756																					
neonatal jaundice Hemolytic disease and other	2000) g Birth prevalence - [34, 36) wks, [1500,	Martality	ala:	(56.885 to 100.9 55.555	 (28.764 to 41.849) 21.346 	, I																				
neonatal jaundice Hemolytic disease and other	2000) g Birth prevalence - [34, 36) wise [1500	Mortany N		(39.553 to 75.10 64 376	 (17.677 to 26.143) 23.046 																					
neonatal jaundice	2000) g	Mortality Fe	nates	(38.617 to 75.2	(18.743 to 28.287																					
Hemolytic disease and other neonatal jaundice	Both prevalence - [32, 34) wks, [1500, 2000) g	Mortality M	iales	57.155 (42.484 to 73.65	23.114 (19.028 to 27.915)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [32, 34) wks, [1500, 2000) g	Mortality Fe	nales	56.101 (39.794 to 76.2	25.149 (20.615 to 30.388)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [32, 34) wks, [2000, 2500) g	Mortality M	iales	37.444 (29.026 to 48 ?	12:233 (10:252 to 14:477	,																				
Hemolytic disease and other	Birth prevalence - [32, 34) wks, [2000,	Mortality Fe	nales	36.874	14.384																					
neonatal jaundice Hemolytic disease and other	2500) g Birth prevalence - [40, 42) wks, [2000,	Mortality	ales	(26.658 to 49.6) 18.092	9.23 (12.095 to 17.03)	1																				
neonatal jaundice	2500) g		1	(13.292 to 23.7	(7.037 to 11.454)	1																				

	rused by age and see for each onte			an ponution acono	, and smoking,									А	iges											
Risk - Outcome	Category / Units	Morbidity / Mortality	Sex All-ag	e 0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Hemolytic disease and other	Birth prevalence - [40, 42) wks, [2000, 26000 -	Mortality	Females	15.574	9.975																					
Hemolytic disease and other	Birth prevalence - [38, 40) wks, [2000,	Mortality	Males	13.104	8.198																					
neonatal jaundace Hemolytic disease and other	2500) g Birth prevalence - [38, 40) wks, [2000,	Mantalia	Employ	(9.829 to 16.99) 11.308	(6.786 to 9.959) 8.577																					
neonatal jaundice Hemolytic disease and other	2500) g Birth prevalence - [32, 34) wks, [2500,			(8.389 to 14.38) 33.063	(7.04 to 10.449) 8.441																					
neonatal jaundice Hemolotic diarane and other	3000) g Birth merculance - [32-34) wise [2500	stortany	ALLIS	(24.393 to 43.503 32.812) (6.822 to 10.431) 10.398																					
neonatal jaundice	3000) g	Mortality	Females	(23.439 to 45.567) (8.227 to 13.042)																					
Hemolytic disease and other neonatal jaundice	Both prevalence - [34, 36) wks, [2000, 2500) g	Mortality	Males	21.925 (16.305 to 29.433	9.367) (7.859 to 11.112)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [34, 36) wks, [2000, 2500) g	Mortality	Females	21.297 (15.657 to 28.76)	10.295 (8.548 to 12.273)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [37, 38) wks, [2000, 2500) g	Mortality	Males	13.0 (10.102 to 16.456	8.096 (6.724 to 9.676)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [37, 38) wks, [2000, 2500) g	Mortality	Females	11.563 (8.805 to 15.11)	8.467 (6.994 to 10.342)																					
Hemolytic disease and other	Birth prevalence - [36, 37) wks, [2000, 2500) n	Mortality	Males	14.401 (10.789 to 18.654	8.221																					
Hemolytic disease and other	Birth prevalence - [36, 37) wks, [2000,	Mortality	Females	13.513	8.654																					
neonatal jaundace Hemolytic disease and other	2500) g Birth prevalence - [34, 36) wks, [2500,	Mortality	Males	(9.817 to 17.942 13.419	(7.215 to 10.369) 5.562																					
neonatal jaundice Hemolytic disease and other	3000) g Birth prevalence - [34, 36) wks, [2500,	M	F - 1	(10.387 to 16.819 13.266) (4.646 to 6.696) 6.395																					
neonatal jaundice Hemolotic diarane and other	3000) g Birth merculance - [34, 36) wise [4000	storcarry	remates	(9.666 to 17.689 23.096	(5.292 to 7.606) 2.895																					
neonatal jaundice	4500) g	Mortality	Males	(14.708 to 35.098) (2.245 to 3.716)																					
Hemolytic disease and other neonatal jaundice	Barth prevalence - [34, 36) wks, [4000, 4500) g	Mortality	Females	25.038 (15.763 to 37.255) (2.855 to 4.925)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [34, 36) wks, [3000, 3500) g	Mortality	Males	14.006 (10.222 to 18.478	4.322) (3.449 to 5.338)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [34, 36) wks, [3000, 3500) g	Mortality	Females	14.375 (10.269 to 20.114	5.265) (4.145 to 6.564)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [36, 37) wks, [2500, 3000) g	Mortality	Males	4.874 (4.014 to 5.713)	3.699 (3.134 to 4.374)																					
Hemolytic disease and other	Birth prevalence - [36, 37) wks, [2500, 2000) -	Mortality	Females	4.609	3.898 (2.225 to 4.67)																					
Hemolytic disease and other	Birth prevalence - [34, 36) wks, [3500,	Mortality	Males	18.024	3.657																					
neonatal jaundace Hemolytic disease and other	4000) g Birth prevalence - [34, 36) wks, [3500,	Mantality	Employ	(12.279 to 25.54) 19.263) (2.838 to 4.675) 4.634																					
neonatal jaundice Hemolytic disease and other	4000) g Birth meyalence - [37, 38) wks, [2500,	January .		(12.567 to 27.924 3.306) (3.576 to 5.989) 3.194																					
neonatal jaundice	3000) g	Mortality	Males	(2.82 to 3.843)	(2.691 to 3.803)																					
neonatal jaundice	3000) g	Mortality	Females	(2.496 to 3.61)	(2.745 to 3.817)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [40, 42) wks, [2500, 3000) g	Mortality	Males	3.771 (3.002 to 4.693)	3.175 (2.56 to 3.923)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [40, 42) wks, [2500, 3000) g	Mortality	Females	3.244 (2.486 to 4.159)	3.228 (2.605 to 3.985)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [38, 40) wks, [2500, 3000) g	Mortality	Males	2.755 (2.274 to 3.309)	2.944 (2.44 to 3.548)																					
Hemolytic disease and other neonatal jaurdice	Birth prevalence - [38, 40) wks, [2500, 3000) g	Mortality	Females	2.376 (1.91 to 2.886)	2.938 (2.434 to 3.503)																					
Hemolytic disease and other	Birth prevalence - [36, 37) wks, [3000, 2600) -	Mortality	Males	3.774	2.466																					
Hemolytic disease and other	Birth prevalence - [36, 37) wks, [3000,	Mortality	Females	3.73	2.715																					
neonatal jaundace Hemolytic disease and other	3500) g Birth prevalence - [36, 37) wks, [4000,	Mantalia	Malar	(2.981 to 4.646) 6.826	(2.277 to 3.218)																					
neonatal jaundice Hemolytic disease and other	4500) g Birth meyalence - [36, 37) wks, [4000,	January .		(5.212 to 9.045) 7.269	(1.491 to 2.082) 2.177																					
neonatal jaundice Hemolotic diarane and other	4500) g Birth merculance - [36, 37) wise [3500	Mortality	Females	(5.144 to 9.821) 4 544	(1.786 to 2.607) 2.057																					
neonatal jaundice	4000) g	Mortality	Males	(3.64 to 5.622)	(1.74 to 2.44)																					
neonatal jaundice	4000) g	Mortality	Females	4.002 (3.577 to 6.014)	(1.98 to 2.864)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [37, 38) wks, [3000, 3500) g	Mortality	Males	2.007 (1.759 to 2.293)	1.888 (1.613 to 2.224)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [37, 38) wks, [3000, 3500) g	Mortality	Females	1.925 (1.582 to 2.328)	1.972 (1.669 to 2.313)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [37, 38) wks, [4000, 4500) g	Mortality	Males	3.28 (2.596 to 4.133)	1.335 (1.171 to 1.532)																					
Hemolytic disease and other neonatal jaurdice	Birth prevalence - [37, 38) wks, [4000, 4500) g	Mortality	Females	3.521 (2.649 to 4.5)	1.559 (1.333 to 1.835)																					
Hemolytic disease and other	Birth prevalence - [37, 38) wks, [3500,	Mortality	Males	2.128	1.505																					
Hemolytic disease and other	Birth prevalence - [37, 38) wks, [3500,	Mortality	Females	2.142	1.661																					
neonatal jaundace Hemolytic disease and other	4000) g Birth prevalence - [40, 42) wks, [3000,	Mantalia	Malar	(1.694 to 2.67) 1.436	(1.411 to 1.961) 1.47																					
neonatal jaundice Hemolytic disease and other	3500) g Birth prevalence - [40, 42) wks, [3000.	Marchine	Emerica	(1.245 to 1.65) 1.326	(1.199 to 1.8) 1.465																					
neonatal jaundice Hemolytic disease and other	3500) g Birth prevalence - [38, 40) wks, [3000,	Mortany	remaies	(1.069 to 1.614) 1.33	(1.188 to 1.775) 1.559																					
neonatal jaundice	3500)g Risk mersken: (28, 40) ode 10000	Mortality	auto	(1.155 to 1.53)	(1.305 to 1.851)																					
neronatal jaundice	annun prevatence - (38, 40) was, (3000, 3500) g	Mortality	Females	(1.0 to 1.492)	(1.304 to 1.847)																					
Hemolytic disease and other neonatal jaundice	north prevalence - [38, 40) wks, [4000, 4500) g	Mortality	Males	1.787 (1.453 to 2.182)	1.175 (1.005 to 1.371)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [38, 40) wks, [4000, 4500) g	Mortality	Females	1.877 (1.467 to 2.388)	1.224 (1.022 to 1.465)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [38, 40) wks, [3500, 4000) g	Mortality	Males	1.785 (1.478 to 2.147)	1.173 (1.0 to 1.377)																					
Hemolytic disease and other neonatal jaurdice	Birth prevalence - [38, 40) wks, [3500, 4000) g	Mortality	Females	1.892 (1.481 to 2.352)	1.23 (1.03 to 1.46)																					
Hemolytic disease and other	Birth prevalence - [40, 42) wks, [3500, 4000) #	Mortality	Males	1.0	1.003																					
Hemolytic disease and other	Birth prevalence - [40, 42) wks, [3500,	Mortality	Females	1.002	1.001																					
neonatal jaundice Hemolytic disease and other	-4000) g Birth prevalence - [40, 42) wks, [4000,	Manufin	Malex	(1.0 to 1.013) 1.0	(1.0 to 1.006) 1.0																					
neonatal jaundice Hemolytic disease and other	4500) g Birth prevalence - [40, 42) wks. (4000	Mortany	manX	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0																					
neonatal jaundice Hemolotic discuss and ad	4500) g Birth menalence - (20, 20) and - (2000)	Mortality	remaies	(1.0 to 1.0)	(1.0 to 1.0)																					
neonatal jaundice	2500) g	Mortality	Males	(83.895 to 158.05	5) (21.877 to 34.972)																					
Hemolytic disease and other neonatal jaundice	north prevalence - [28, 30) wks, [2000, 2500) g	Mortality	Females	121.682 (84.349 to 171.37	33.983 (26.101 to 43.404)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [28, 30) wks, [2500, 3000) g	Mortality	Males	77.948 (54.687 to 105.04	16.608 (12.653 to 21.188)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [28, 30) wks, [2500, 3000) g	Mortality	Females	79.193 (54.099 to 112.23	20.387 (15.089 to 26.434)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [28, 30) wks, [3000, 3500) g	Mortality	Males	42.199 (29.891 to 57.22)	10.082 (7.777 to 13.056)																					
Hemolytic disease and other neonatal jaundice	- Birth prevalence - [28, 30) wks, [3000, 3500) #	Mortality	Females	42.551 (28.25 to 63.209	11.989 (9.084 to 15.544)																					
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															Ag	ges											
Birk Outroom	Catagory / Unite M	andridites / Mantalites	S	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Hemolytic disease and other	Bith prevalence - [30, 32) wks, [2500,	Mortality	Males		58.722	12.115																					
neonatal jaundice Hemolytic disease and other	3000) g Birth prevalence - [30, 32) wks, [2500,	Marality	Females		(42.419 to 78.873) 59.522	(9.518 to 15.521) 15.364																					
neonatal jaundice Hemolytic disease and other	3000) g Birth menulence - (30, 32) wks, (3000,				(42.058 to 82.793) 45.67	(11.936 to 19.581) 8.381																					
neonatal jaundice Hemolotic diarane and other	3500) g Birth merculance - (30-32) wite (3000	stortany	MING		(32.014 to 65.531) 46.104	(6.362 to 10.842) 10 506																					
neonatal jaundice	3500) g	Mortality	Females		(30.243 to 66.207)	(8.041 to 13.513)																					
Hemolytic disease and other neonatal jaundice	Bath prevalence - [30, 32] wks, [3500, 4000) g	Mortality	Males		36.334 (21.558 to 54.813)	5.698 (4.293 to 7.349)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [30, 32) wks, [3500, 4000) g	Mortality	Females		37.931 (22.692 to 61.276)	6.892 (5.175 to 9.058)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [32, 34) wks, [3000, 3500) g	Mortality	Males		34.016 (23.515 to 48.37)	6.577 (4.934 to 8.436)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [32, 34) wks, [3000, 3500) a	Mortality	Females		34.585 (22.909 to 49.754)	8.314 (6.362 to 10.789)																					
Hemolytic disease and other	Birth prevalence - [32, 34) wks, [3500,	Mortality	Males		36.248	5.068																					
Hemolytic disease and other	4000) g Birth prevalence - [32, 34) wks, [3500,	Marality	Females		38.098	6.476																					
neonatal jaundice Hemolytic disease and other	4000) g Birth prevalence - [36, 37) wks, [1000,				(23.301 to 59.429) 166.686	(4.666 to 8.689) 57.535																					
neonatal jaundice Hemolotic diarane and other	1500) g Birth merculance - [36, 37) why [1000	stortany	MING		(118.487 to 222.581) 169.725	(45.999 to 71.742) 63.564																					
neonatal jaundice	1500) g	Mortality	Females		(119.017 to 229.008)	(50.068 to 80.703)																					
Hemolytic disease and other neonatal jaundice	Bath prevalence - [38, 40) wio, [1000, 1500) g	Mortality	Males		(125.125 to 232.507)	57.966 (44.393 to 73.241)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [38, 40) wks, [1000, 1500) g	Mortality	Females		171.557 (121.585 to 237.047)	65.208 (48.821 to 84.308)																					
Hemolytic disease and other neonatal jaundice	Birth prevalence - [38, 40) wks, [1500, 2000) g	Mortality	Males		67.302 (49.547 to 89.055)	25.206 (20.365 to 31.168)																					
Hemolytic disease and other neonatal jaurdice	Birth prevalence - [38, 40) wks, [1500, 2000) #	Mortality	Females		62.19 (45.884 to 83.445)	28.05 (22.625 to 35.139)																					
Hemolytic disease and other	Birth prevalence - [40, 42) wks, [1500, 2000) -	Mortality	Males		76.673	25.785																					
Hemolytic disease and other	Birth prevalence - [40, 42) wks, [1500,	Mortality	Females		70.411	29.113																					
neonatal jaundace	2000) g	Mantality	Male		(49.221 to 97.952) 1564.792	(21.355 to 38.272) 618.595																					
Cristi accessi accorda	anna prevanace - (0, 24) was, (0, 500) g	sectary			2116 062) 1600.122	(458.842 to 812.921) 712.671																					
Other neonatal disorders	Birth prevalence - [0, 24) wkx, [0, 500) g	Mortality	Females		(1050.664 to 2011.077)	(526.178 to 921.018)																					
Other neomatal disorders	Bath prevalence - [0, 24) wks, [500, 1000) 8	Mortality	Males	c	1155.815 825.412 to 1506.837)	457.5 (352.552 to 573.483)																					
Other neonatal disorders	Birth prevalence - [0, 24) wks, [500, 1000) 8	Mortality	Females	c	1169.123 802.003 to 1617.979)	515.406 (396.713 to 641.541)																					
Other neonatal disorders	Birth prevalence - [24, 26) wks, [500, 1000) g	Mortality	Males	c	955.583 723.748 to 1244.265)	443.357 (363.03 to 534.695)																					
Other neonatal disorders	Birth prevalence - [24, 26) wks, [500, 1000)	Mortality	Females		947.143 702.662 to 1237.093)	487.549 (387.307 to 603.498)																					
Other neomatal disorders	Birth prevalence - [26, 28) wks, [500, 1000)	Mortality	Males		497.817	330.886																					
Other neonatal disorders	8 Birth prevalence - [26, 28) wks, [500, 1000)	Mortality	Females	ľ	483.682	344.618																					
Other excepted disorders	8 Birth prevalence - [30, 32) wks, [500, 1000)	Mantalian	Male	1	(354.946 to 629.517) 236.614	(274.427 to 419.864) 149.995																					
Cristi accura accura	8 Birth merculance - [30, 32] wire (500, 1000)	analany			(163.821 to 324.502) 229.197	(117.866 to 188.368) 152.117																					
Other neonatal disorders	8 Birth muscleum (28, 20) who (500, 1000)	Mortality	Females		(157.606 to 317.194) 207.620	(120.779 to 190.583) 216.005																					
Other neonatal disorders	8 8 8 8 9 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	Mortality	Males		(214.953 to 396.586)	(173.321 to 271.466)																					
Other neonatal disorders	Birth prevalence - [28, 30) wks, [500, 1000) 8	Mortality	Females		281.056 (198.176 to 386.635)	219.884 (174.264 to 272.704)																					
Other neomatal disorders	Birth prevalence - [26, 28) wks, [1000, 1500) g	Mortality	Males		267.91 (210.177 to 332.92)	164.167 (132.898 to 200.569)																					
Other neonatal disorders	Birth prevalence - [26, 28) wks, [1000, 1500) g	Mortality	Females		266.509 (197.461 to 346.932)	174.222 (137.431 to 217.349)																					
Other neonatal disorders	Birth prevalence - [34, 36) wks, [1000, 1500) a	Mortality	Males		142.056 (98.086 to 197.774)	52.86 (42.914 to 64.617)																					
Other neonatal disorders	Birth prevalence - [34, 36) wks, [1000, 1600) -	Mortality	Females		141.899	57.421																					
Other neonatal disorders	Birth prevalence - [28, 30) wks, [1500,	Mortality	Males		127.966	50.018																					
Other excepted disorders	2000) g Birth prevalence - [28, 30) wks, [1500,	Mantality	Employ		(97.178 to 167.026) 130.924	(40.539 to 61.919) 57.275																					
	2000) g Birth merulence - (28, 30) wks, (1000,		- under		(96.513 to 172.188) 158.563	(46.36 to 70.038) 103.32																					
Other neonaria disorders	1500) g Birth manufana (28, 20) who (1000	stortany	MING		(120.99 to 204.947) 152.005	(83.486 to 127.144)																					
Other neonatal disorders	1500) g	Mortality	Females		(112.327 to 200.786)	(86.954 to 131.78)																					
Other neonatal disorders	namn prevalence - [32, 34) wks, [1000, 1500) g	Mortality	Males		(81.354 to 161.101)	53.185 (43.049 to 66.274)																					
Other neonatal disorders	Birth prevalence - [32, 34) wks, [1000, 1500) g	Mortality	Females		115.171 (79.363 to 159.206)	56.034 (45.982 to 68.36)																					
Other neomatal disorders	Birth prevalence - [30, 32) wks, [1000, 1500) g	Mortality	Males		119.308 (87.769 to 160.885)	67.163 (54.863 to 82.638)																					
Other neonatal disorders	Birth prevalence - [30, 32) wks, [1000, 1500) g	Mortality	Females		115.448 (84.272 to 156.425)	69.14 (55.873 to 85.012)																					
Other neomatal disorders	Birth prevalence - [37, 38) wks, [1500, 2000) g	Mortality	Males		62.972 (46.159 to 83.484)	24.148 (20.066 to 29.406)																					
Other neonatal disorders	Birth prevalence - [37, 38) wks, [1500, 2000) #	Mortality	Females		59.988 (43.974 to 79.053)	26.719 (21.746 to 32.816)																					
Other neomatal disorders	Birth prevalence - [36, 37] wks, [1500, 2000) -	Mortality	Males		60.218	23.031																					
Other excepted disorders	2000) g Birth prevalence - [36, 37) wks, [1500,	Mantalian	Employ		(43.669 to 82.48) 58.527	(18.193 to 28.485) 25.143																					
Oder and 1 Sector	2000) g Birth prevalence - [30, 32) wks, [2000,	Mastelli	Male		(42.172 to 80.557) 67.971	(20.331 to 30.566) 18.03																					
Other neonatal desortary	2500) g Birth menalence - (30, 32) wise (2000	somany	will be		(50.354 to 88.935) 69.383	(14.621 to 22.103) 22.069																					
Other neonatal disorders	2500) g	Mortality	Females		(49.108 to 94.583)	(17.836 to 27.163)																					
Other neonatal disorders	2000) g	Mortality	Males		(59.702 to 99.232)	(25.786 to 36.724)																					
Other neomatal disorders	Birth prevalence - [30, 32) wks, [1500, 2000) g	Mortality	Females		76.134 (56.885 to 100.996)	34.756 (28.764 to 41.849)																					
Other neonatal disorders	Birth prevalence - [34, 36) wks, [1500, 2000) g	Mortality	Males		55.555 (39.553 to 75.104)	21.346 (17.677 to 26.143)																					
Other neonatal disorders	Birth prevalence - [34, 36) wks, [1500, 2000) g	Mortality	Females		54.335 (38.617 to 75.24)	23.046 (18.743 to 28.287)																					
Other neonatal disorders	- Birth prevalence - [32, 34) wks, [1500, 2000) #	Mortality	Males		57.155 (42.484 to 73.651)	23.114 (19.028 to 27.915)																					
Other neomatal disorders	Birth prevalence - [32, 34) wks, [1500, 2000) -	Mortality	Females		56.101	25.149																					
Other assessial discarder-	2000) g Birth prevalence - [32, 34) wks, [2000,	Montality	Malex		(39.794 to 76.295) 37.444	(20.615 to 30.388) 12.233																					
Course an ordered descriptions	2500) g Birth prevalence - [32, 34) wks, [2000.	ana udity	-mana		(29.026 to 48.227) 36.874	(10.252 to 14.477) 14.384																					
Other neomital disorders	2500) g Birth mevalence - (40 47) wise (2000	Mortality	remates		(26.658 to 49.653) 18.097	(12.095 to 17.03) 9.23																					
Other neonatal disorders	2500) g	Mortality	Males		(13.292 to 23.719)	(7.037 to 11.454)																					

															A	ges											
Risk - Outcome	Category / Units	Morbidity / Mortality	Sex	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Other neonatal disorders	Birth prevalence - [40, 42] wks, [2000, 25000 -	Mortality	Females	au	15.574	9.975 (7.82) in 12.46)												11									
Other neonatal disorders	Birth prevalence - [38, 40) wks, [2000,	Mortality	Males	(11.	13.104	8.198																					
Other recented disorders	2500) g Birth prevalence - [38, 40) wks, [2000,	Manulity	Females	(9.	11.308	(6.786 to 9.959) 8.577																					
Other excepted formulae	2500) g Birth prevalence - [32, 34) wks, [2500,	Manufity	Mala	(8-	1.389 to 14.38) 33.063	(7.04 to 10.449) 8.441																					
	3000) g Birth menulance - [32-34) wise [2500	Juniany	ALL A	(24.	32 812	(6.822 to 10.431) 10.398																					
Other neonatal disorders	3000) g	Mortality	Females	(23)	1.439 to 45.567)	(8.227 to 13.042)																					
Other neonatal disorders	2500) g	Mortality	Males	(16.	i.305 to 29.433)	(7.859 to 11.112)																					
Other neomatal disorders	Bath prevalence - [34, 36) wks, [2000, 2500) g	Mortality	Females	(15)	21.297 i.657 to 28.761)	(8.548 to 12.273)																					
Other neoratal disorders	Birth prevalence - [37, 38) wks, [2000, 2500) g	Mortality	Males	(10.	13.0 1.102 to 16.456)	8.096 (6.724 to 9.676)																					
Other neonatal disorders	Birth prevalence - [37, 38) wks, [2000, 2500) g	Mortality	Females	(8.	11.563 1.805 to 15.11)	8.467 (6.994 to 10.342)																					
Other neonatal disorders	Birth prevalence - [36, 37) wks, [2000, 2500) g	Mortality	Males	(10:	14.401 1.789 to 18.654)	8.221 (6.917 to 9.923)																					
Other neonatal disorders	Birth prevalence - [36, 37) wks, [2000, 2500) g	Mortality	Females	(9.8	13.513 817 to 17.942)	8.654 (7.215 to 10.369)																					
Other neonatal disorders	Birth prevalence - [34, 36) wks, [2500, 3000) g	Mortality	Males	(10.	13.419 1.387 to 16.819)	5.562 (4.646 to 6.696)																					
Other neonatal disorders	Birth prevalence - [34, 36) wks, [2500, 3000) p	Mortality	Females	(9.6	13.266 666 to 17.68%	6.395 (5.292 to 7.606)																					
Other neonatal disorders	Birth prevalence - [34, 36) wks, [4000, 45000 -	Mortality	Males	au	23.096	2,895																					
Other neonatal disorders	Birth prevalence - [34, 36) wks, [4000,	Mortality	Females		25.038	3.778																					
Other recorded disorders	4500) g Birth prevalence - [34, 36) wks, [3000,	Manulity	Malex	(15.	14.006	(2.855 to 4.925) 4.322																					
Other excepted formulae	3500) g Birth prevalence - [34, 36) wks, [3000,	Manufity	Employ	(10.	14.375	(3.449 to 5.338) 5.265																					
	3500) g Birth menulence - (36, 37) wks, (2500,	Juniary .		(10.	4.874	(4.145 to 6.564) 3.699																					
Other neonatal disorders	3000) g Birth manufaran (36, 37) mbu (2000	Mortality	Males	(4.)	1.014 to 5.713)	(3.134 to 4.374)																					
Other neonatal disorders	3000) g	Mortality	Females	(3	3.731 to 5.61)	(3.235 to 4.67)																					
Other neomatal disorders	Both prevalence - [34, 36) wks, [3500, 4000) g	Mortality	Males	(12.	18.024 1.279 to 25.547)	3.657 (2.838 to 4.675)																					
Other neonatal disorders	Birth prevalence - [34, 36) wks, [3500, 4000) g	Mortality	Females	(12.	19.263 1.567 to 27.924)	4.634 (3.576 to 5.989)																					
Other neoratal disorders	Birth prevalence - [37, 38) wks, [2500, 3000) g	Mortality	Males	(2	3.306 2.82 to 3.843)	3.194 (2.691 to 3.803)																					
Other neonatal disorders	Birth prevalence - [37, 38) wks, [2500, 3000) g	Mortality	Females	(2	2.991 2.496 to 3.61)	3.242 (2.745 to 3.817)																					
Other neonatal disorders	Birth prevalence - [40, 42) wks, [2500, 3000) g	Mortality	Males	(3)	3.771 1.002 to 4.693)	3.175 (2.56 to 3.923)																					
Other neonatal disorders	Birth prevalence - [40, 42) wks, [2500, 3000) g	Mortality	Females	2	3.244 1.486 to 4.159)	3.228 (2.605 to 3.985)																					
Other neomatal disorders	Birth prevalence - [38, 40) wks, [2500, 3000) g	Mortality	Males	(2:	2.755 2.274 to 3.309)	2.944 (2.44 to 3.548)																					
Other neonatal disorders	Birth prevalence - [38, 40) wks, [2500, 3000) #	Mortality	Females		2.376	2.938 (2.434 to 3.503)																					
Other neonatal disorders	Birth prevalence - [36, 37) wks, [3000, 2500) -	Mortality	Males		3.774	2.466																					
Other neonatal disorders	Birth prevalence - [36, 37) wks, [3000,	Mortality	Females		3.73	2.715																					
Other recented disorders	3500) g Birth prevalence - [36, 37) wks, [4000,	Manulity	Malex	(2.)	6.826	(2.277 to 3.218) 1.77																					
Other recorded disorders	4500) g Birth prevalence - [36, 37) wks, [4000,	Manulity	Females	0.	7.269	(1.491 to 2.082) 2.177																					
Other excepted formulae	4500) g Birth prevalence - [36, 37) wks, [3500,	Manufity	Mala	(5.	4.544 to 9.821)	(1.786 to 2.607) 2.057																					
	4000) g Birth menulence - (36, 37) wks, (3500,	Juniany		(3	3.64 to 5.622) 4.662	(1.74 to 2.44) 2.398																					
Other neomatal disorders	4000) g Birth menalence - [37, 38) wise [3000	Mortality	Females	(3.	1.577 to 6.014)	(1.98 to 2.864)																					
Other neonatal disorders	3500) g	Mortality	Males	(I.:	.759 to 2.293)	(1.613 to 2.224)																					
Other neonatal disorders	3500) g	Mortality	Females	(1.	.582 to 2.328)	(1.669 to 2.313)																					
Other neonatal disorders	4500) g	Mortality	Males	(2.	3.28 1.596 to 4.133)	(1.171 to 1.532)																					
Other neonatal disorders	4500) g	Mortality	Females	đ	(2.649 to 4.5)	(1.333 to 1.835)																					
Other neoratal disorders	Birth prevalence - [37, 38) wks, [3500, 4000) g	Mortality	Males	(1.	2.128 .833 to 2.466)	1.505 (1.299 to 1.76)																					
Other neonatal disorders	Birth prevalence - [37, 38) wks, [3500, 4000) g	Mortality	Females	(1	2.142 1.694 to 2.67)	1.661 (1.411 to 1.961)																					
Other neonatal disorders	Birth prevalence - [40, 42) wks, [3000, 3500) g	Mortality	Males	(1	1.436 1.245 to 1.65)	1.47 (1.199 to 1.8)																					
Other neonatal disorders	Birth prevalence - [40, 42) wks, [3000, 3500) g	Mortality	Females	(L)	1.326 .069 to 1.614)	1.465 (1.188 to 1.775)																					
Other neonatal disorders	Birth prevalence - [38, 40) wks, [3000, 3500) g	Mortality	Males	(1	1.33 1.155 to 1.53)	1.559 (1.305 to 1.851)																					
Other neonatal disorders	Birth prevalence - [38, 40) wks, [3000, 3500) g	Mortality	Females	a	1.224 (1.0 to 1.492)	1.564 (1.304 to 1.847)																					
Other neonatal disorders	Birth prevalence - [38, 40) wks, [4000, 4500) g	Mortality	Males	a	1.787 .453 to 2.182)	1.175 (1.005 to 1.371)																					
Other neoratal disorders	Birth prevalence - [38, 40) wks, [4000, 4500) g	Mortality	Females	a.	1.877 .467 to 2.388)	1.224 (1.022 to 1.465)																					
Other neonatal disorders	Birth prevalence - [38, 40) wks, [3500,	Mortality	Males		1.785	1.173																					
Other neonatal disorders	Bith prevalence - [38, 40) wks, [3500,	Mortality	Females	(L) (L)	1.892	1.23																					
Other neoratal disorders	000) g Birth prevalence - [40, 42) wks, [3500,	Mortality	Males	0.	1.0	1.003																					
Other necessarial disconter-	+000) g Birth prevalence - [40, 42) wks, [3500,	Montaline	Females	'	(1.0 to 1.0) 1.002	(1.0 to 1.046) 1.001																					
Other any second second	4000) g Birth prevalence - [40, 42] wks, [4000.	Mark	Mala	a	(1.0 to 1.013) 1.0	(1.0 to 1.006) 1.0																					
Other neonatal desorders	4500) g Birth prevalence - [40, 42) wks. [4000	successfully	-ALLIER	·	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0																					
Other neonatal disorders	4500) g Bith prevalence - [28, 30) wdw (2000	Mortality	remaies		(1.0 to 1.0)	(1.0 to 1.0) 27,726																					
Other neoratal disorders	2500) g Risk sensions (20, 20, 1, 2007	Mortality	Males	(83.8	895 to 158.056)	(21.877 to 34.972)																					
Other neonatal disorders	2500) g	Mortality	Females	(84.3	349 to 171.375)	(26.101 to 43.404)																					
Other neonatal disorders	nonn prevalence - [28, 30) wks, [2500, 3000) g	Mortality	Males	(54.6	/1.948 .687 to 105.047)	16.608 (12.653 to 21.188)																					
Other neoratal disorders	Birth prevalence - [28, 30) wks, [2500, 3000) g	Mortality	Females	(\$4.0	79.193 .099 to 112.236)	20.387 (15.089 to 26.434)																					
Other neonatal disorders	Birth prevalence - [28, 30) wks, [3000, 3500) g	Mortality	Males	(29)	42.199 9.891 to 57.227)	10.082 (7.777 to 13.056)																					
Other neonatal disorders	Birth prevalence - [28, 30) wks, [3000, 3500) g	Mortality	Females	(28	42.551 8.25 to 63.209)	11.989 (9.084 to 15.544)																					

opendix Table 6a. Relative risk	s used by age and sex for each outco	me for all risk factors ex	cept for ambient	air pollution alcohol.	and snoxing.									А	Ages											
Risk - Outcome	Category / Units	Morbidity / Mortality	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Other neomatal disorders	Birth prevalence - [30, 32) wks, [2500, 3000) g	Mortality	dales	58.722 (42.419 to 78.873)	12.115 (9.518 to 15.521)																					
Other neonatal disorders	Birth prevalence - [30, 32) wks, [2500, 2000) -	Mortality F	males	59.522 (#2.058 to \$2.202)	15.364																					
Other neonatal disorders	Birth prevalence - [30, 32) wks, [3000,	Mortality	dales	45.67	8.381																					
Other neormatal disorders	Bith prevalence - [30, 32) wks, [3000,	Mortality F	males	46.104	(6.3621810.842)																					
	3500) g Birth prevalence - [30, 32) wks, [3500,			(30.243 to 66.207) 36.334	(8.041 to 13.513) 5.698																					
Other neonatal disorders	4000) g Birth menalewer - (30-32) wire (3500	Mortality	dates	(21.558 to 54.813) 37.931	(4.293 to 7.349) 6.892																					
Other neonatal disorders	4000) g	Mortality F	males	(22.692 to 61.276)	(5.175 to 9.058)																					
Other neoental disorders	3500) g	Mortality	dales	(23.515 to 48.37)	(4.934 to 8.436)																					
Other neonatal disorders	Birth prevalence - [32, 34) wks, [3000, 3500) g	Mortality F	males	34.585 (22.909 to 49.754)	8.314 (6.362 to 10.789)																					
Other neonatal disorders	Birth prevalence - [32, 34) wks, [3500, 4000) g	Mortality	dales	36.248 (23.158 to 54.67)	5.068 (3.761 to 6.741)																					
Other neonatal disorders	Birth prevalence - [32, 34) wks, [3500, 4000) g	Mortality F	males	38.098 (23.301 to 59.429)	6.476 (4.666 to 8.689)																					
Other neonatal disorders	Birth prevalence - [36, 37) wks, [1000, 1500) g	Mortality	dales	166.686 (118.487 to 222.581	57.535) (45.999 to 71.742)																					
Other neonatal disorders	Birth prevalence - [36, 37) wks, [1000, 1500) g	Mortality F	males	169.725 (119.017 to 229.008	63.564 (50.068 to 80.703)																					
Other neonatal disorders	Birth prevalence - [38, 40) wks, [1000, 1500) #	Mortality	dales	174.066 (125.125 to 232.507	57.966 (44.393 to 73.241)																					
Other neonatal disorders	Birth prevalence - [38, 40) wks, [1000,	Mortality F	males	171.557	65.208																					
Other excepted disorders	1500) g Birth prevalence - [38, 40) wks, [1500,	Manufility	41) m	(121.585 to 237.047 67.302	(48.821 to 84.308) 25.206																					
Odar and a distant	2000) g Birth prevalence - [38, 40) wks, [1500,	Marking 7		(49.547 to 89.055) 62.19	(20.365 to 31.168) 28.05																					
Other neonatal disorders	2000) g Birth menalewer - [40, 47) wire (1500	Mortality F	malex	(45.884 to 83.445) 76.673	(22.625 to 35.139) 25.785																					
Other neonatal disorders	2000) g	Mortality	dates	(56.177 to 102.468	(19.387 to 34.168)																					
Other neonatal disorders	Birth prevalence - [40, 42) wio, [1500, 2000) g	Mortality F	males	70.411 (49.221 to 97.952) 1564 792	29.113 (21.355 to 38.272)																					
Sudden infant death syndrome	Birth prevalence - [0, 24) wks, [0, 500) g	Mortality	dales	(1056.542 to 2116.062)	618.595 (458.842 to 812.921)																				
Sudden infant death syndrome	Birth prevalence - [0, 24) wks, [0, 500) g	Mortality F	males	(1050.664 to 2011.977)	713.571 (526.178 to 921.018	, ,																				
Sudden infant death syndrome	Birth prevalence - [0, 24) wks, [500, 1000) g	Mortality	dales	1155.815 (825.412 to 1506.83	457.5 7) (352.552 to 573.483)																				
Sudden infant death syndrome	Birth prevalence - [0, 24) wks, [500, 1000) g	Mortality F	males	1169.123 (802.003 to 1617.97	515.406 9) (396.713 to 641.541	, ,																				
Sudden infant death syndrome	Birth prevalence - [24, 26) wks, [500, 1000)	Mortality	dales	955.583 (723.748 to 1244.26)	443.357 5) (363.03 to 534.695)																					
Sudden infant death syndrome	Birth prevalence - [24, 26) wks, [500, 1000)	Mortality F	males	947.143	487.549																					
Sudden infant death syndrome	8 Birth prevalence - [26, 28) wks, [500, 1000)	Mortality	dales	497.817	330.886	,																				
Sudden infant death conductive	8 Birth prevalence - [26, 28) wks, [500, 1000)	Mandity F	male	483.682	344.618	,																				
Seeking infest doub contemp	8 Birth prevalence - [30, 32) wks, [500, 1000)	Manufito	61) m	(354.946 to 629.517 236.614	(274.427 to 419.864 149.995)																				
States man state spectrum	8 Birth mevalence - (30, 32) wks, (500, 1000)	January .		(163.821 to 324.502 229.197	(117.866 to 188.368) 152.117)																				
Sudden infant death syndrome	8 Birth menalewer - (28, 30) wire (500, 1000)	Mortality F	malex	(157.606 to 317.194 297.629	(120.779 to 190.583) 216.995)																				
Sudden infant death syndrome	8	Mortality	dates	(214.953 to 396.58t	i) (173.321 to 271.466	•																				
Sudden infant death syndrome	8 8 8 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9	Mortality F	males	(198.176 to 386.635	(174.264 to 272.704)																				
Sudden infant death syndrome	Bath prevalence - [26, 28) wis, [1000, 1500) g	Mortality 1	dales	(210.177 to 332.92	(132.898 to 200.569)	>																				
Sudden infant death syndrome	Birth prevalence - [26, 28) wks, [1000, 1500) g	Mortality F	males	266.509 (197.461 to 346.931	174.222 (137.431 to 217.349))																				
Sudden infant death syndrome	Birth prevalence - [34, 36) wks, [1000, 1500) g	Mortality	dales	142.056 (98.086 to 197.774	52.86 (42.914 to 64.617)																					
Sudden infant death syndrome	Birth prevalence - [34, 36) wks, [1000, 1500) g	Mortality F	males	141.899 (95.864 to 197.656	57.421 (46.452 to 71.339)																					
Sudden infant death syndrome	Birth prevalence - [28, 30) wks, [1500, 2000) g	Mortality	dales	127.966 (97.178 to 167.026	50.018 (40.539 to 61.919)																					
Sudden infant death syndrome	Birth prevalence - [28, 30) wks, [1500, 2000) g	Mortality F	males	130.924 (96.513 to 172.188	57.275 (46.36 to 70.038)																					
Sudden infant death syndrome	Birth prevalence - [28, 30) wks, [1000, 1500) #	Mortality	dales	158.563 (120.99 to 204.947	103.32 (83.486 to 127.144)																					
Sudden infant death syndrome	Birth prevalence - [28, 30) wks, [1000, 1600) -	Mortality F	males	153.905	107.529																					
Sudden infant death syndrome	Birth prevalence - [32, 34) wks, [1000,	Mortality	dales	117.142	53.185																					
Sudden infant death syndrome	Birth prevalence - [32, 34) wks, [1000,	Mortality F	males	115.171	(43.04910.06.274) 56.034																					
Sudden infort death cond	1 500) g Birth prevalence - [30, 32) wks, [1000,	Mortality	dales.	(79.363 to 159.206) 119.308	(45.982 to 68.36) 67.163																					
Surbley infort doubt could	1500) g Birth prevalence - [30, 32) wks, [1000,	Maralin .	make	(87.769 to 160.885 115.448	(54.863 to 82.638) 69.14																					
STRAKET INGER GREEN SYREPOTE	1500) g Birth prevalence - [37, 38) wks, [1500	secondity P		(84.272 to 156.425) 62.972	(55.873 to 85.012) 24.148																					
Sudden infant death syndrome	2000) g Birth menalence - [37-38) wise [1500	Mortality	anas .	(46.159 to 83.484) 50 088	(20.066 to 29.406) 26.719																					
Sudden infant death syndrome	2000) g	Mortality F	males	(43.974 to 79.053)	(21.746 to 32.816)																					
Sudden infant death syndrome	north prevalence - [36, 37] wks, [1500, 2000) g	Mortality	dales	60.218 (43.669 to 82.48)	23.031 (18.793 to 28.483)																					
Sudden infant death syndrome	Bath prevalence - [36, 37) wks, [1500, 2000) g	Mortality F	males	58.527 (42.172 to 80.557)	25.143 (20.331 to 30.566)																					
Sudden infant death syndrome	Birth prevalence - [30, 32) wks, [2000, 2500) g	Mortality	dales	67.971 (50.354 to 88.935)	18.03 (14.621 to 22.103)																					
Sudden infant death syndrome	Birth prevalence - [30, 32) wks, [2000, 2500) g	Mortality F	males	69.383 (49.108 to 94.583)	22.069 (17.836 to 27.163)																					
Sudden infant death syndrome	Birth prevalence - [30, 32) wks, [1500, 2000) g	Mortality	dales	77.369 (59.702 to 99.232)	31.079 (25.786 to 36.724)																					
Sudden infant death syndrome	Birth prevalence - [30, 32) wks, [1500, 2000) g	Mortality F	makes	76.134 (56.885 to 100.996	34.756 (28.764 to 41.849)																					
Sudden infant death syndrome	Birth prevalence - [34, 36) wks, [1500, 2000) r	Mortality	dales	55.555 (39.553 to 75 104)	21.346 (17.677 to 26 143)																					
Sudden infant death syndrome	Birth prevalence - [34, 36) wks, [1500, 2000) #	Mortality F	males	54.335 (38.617 to 75.24)	23.046 (18.743 to 28.397)																					
Sudden infant death syndrome	Birth prevalence - [32, 34) wks, [1500, 2000) -	Mortality	dales	(Jacuar 1 at 75,24) 57,155 (42,484 to 72 (21)	23.114																					
Sudden infant death syndrome	Birth prevalence - [32, 34) wks, [1500,	Mortality F	males	(42.464 10 /3.651) 56.101	25.149																					
Sudden infant death your	2000) g Birth prevalence - [32, 34) wks, [2000,	Martility	dalar.	(39.794 to 76.295) 37.444	(20.615 to 30.388) 12.233																					
Souther information	2500) g Birth prevalence - [32, 34) wks, [2000.	Manut'		(29.026 to 48.227) 36.874	(10.252 to 14.477) 14.384																					
suuaen intant death syndrome	2500) g Birth prevalence - [40, 42) wks, (2000	suoradity F		(26.658 to 49.653) 18.092	(12.095 to 17.03) 9.23																					
Sussen intant death syndrome	2500) g	secretatiy	marek	(13.292 to 23.719)	(7.037 to 11.454)	1																				

	o lace of age and sea for each one.														A	ges											
Risk - Outcome	Category / Units	Morbidity / Mortality	Sex	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Sudden infant death syndrome	Birth prevalence - [40, 42) wks, [2000, 26000 -	Mortality	Females		15.574	9.975																					
Sudden infant death syndrome	Birth prevalence - [38, 40) wks, [2000,	Mortality	Males		13.104	8.198																					
Sudden infant death conducate	2500) g Birth prevalence - [38, 40) wks, [2000,	Monthliny	Females		(9.8291018.99)	(6.786109.939) 8.577																					
Seeking inferst dusts combinen	2500) g Birth prevalence - [32, 34) wks, [2500,	Manuliu	Malar		(8.389 to 14.38) 33.063	(7.04 to 10.449) 8.441																					
Shake man wan yarouk	3000) g Birth menulance - [32-34) wise [2500	Sectory	ALL A		(24.393 to 43.503) 32.812	(6.822 to 10.431) 10.398																					
Sudden infant death syndrome	3000) g	Mortality	Females		(23.439 to 45.567)	(8.227 to 13.042)																					
Sudden infant death syndrome	2500) g	Mortality	Males		(16.305 to 29.433)	9.367 (7.859 to 11.112)																					
Sudden infant death syndrome	Birth prevalence - [34, 36) wks, [2000, 2500) g	Mortality	Females		21.297 (15.657 to 28.761)	10.295 (8.548 to 12.273)																					
Sudden infant death syndrome	Birth prevalence - [37, 38) wks, [2000, 2500) g	Mortality	Males		13.0 (10.102 to 16.456)	8.096 (6.724 to 9.676)																					
Sudden infant death syndrome	Birth prevalence - [37, 38) wks, [2000, 2500) g	Mortality	Females		11.563 (8.805 to 15.11)	8.467 (6.994 to 10.342)																					
Sudden infant death syndrome	Birth prevalence - [36, 37) wks, [2000, 2500) g	Mortality	Males		14.401 (10.789 to 18.654)	8.221 (6.917 to 9.923)																					
Sudden infant death syndrome	Birth prevalence - [36, 37) wks, [2000, 2500) #	Mortality	Females		13.513 (9.817 to 17.947)	8.654 (7.215 to 10.369)																					
Sudden infant death syndrome	Birth prevalence - [34, 36) wks, [2500, 2000) -	Mortality	Males		13.419	5.562																					
Sudden infant death syndrome	Birth prevalence - [34, 36) wks, [2500,	Mortality	Females		13.266	6.395																					
Sudden infant denth sondrome	3000) g Birth prevalence - [34, 36) wks, [4000,	Monthline	Malex		(9.666 to 17.689) 23.096	(5.292 to 7.606) 2.895																					
	4500) g Birth meyalence - [34, 36) wks, [4000,				(14.708 to 35.098) 25.038	(2.245 to 3.716) 3.778																					
Sudden mlant death syndrome	4500) g Birth menalence - [34, 36) wise [3000	Mortality	Females		(15.763 to 37.255) 14.006	(2.855 to 4.925) 4 322																					
Sudden infant death syndrome	3500) g	Mortality	Males		(10.222 to 18.478)	(3.449 to 5.338)																					
Sudden infant death syndrome	3500) g	Mortality	Females		(10.269 to 20.114)	(4.145 to 6.564)																					
Sudden infant death syndrome	Birth prevalence - [36, 37) wks, [2500, 3000) g	Mortality	Males		4.874 (4.014 to 5.713)	3.699 (3.134 to 4.374)																					
Sudden infant death syndrome	Birth prevalence - [36, 37) wks, [2500, 3000) g	Mortality	Females		4.609 (3.731 to 5.61)	3.898 (3.235 to 4.67)																					
Sudden infant death syndrome	Birth prevalence - [34, 36) wks, [3500, 4000) g	Mortality	Males		18.024 (12.279 to 25.547)	3.657 (2.838 to 4.675)																					
Sudden infant death syndrome	Birth prevalence - [34, 36) wks, [3500, 4000) g	Mortality	Females		19.263 (12.567 to 27.924)	4.634 (3.576 to 5.989)																					
Sudden infant death syndrome	Birth prevalence - [37, 38) wks, [2500, 3000) g	Mortality	Males		3.306 (2.82 to 3.843)	3.194 (2.691 to 3.803)																					
Sudden infant death syndrome	Birth prevalence - [37, 38) wks, [2500, 3000) #	Mortality	Females		2.991 (2.496 to 3.61)	3.242 (2.745 to 3.817)																					
Sudden infant death syndrome	Birth prevalence - [40, 42) wks, [2500,	Mortality	Males		3.771	3.175																					
Sudden infant denth conductor	3000) g Birth prevalence - [40, 42) wks, [2500,	Monthline	Females		(3.002 to 4.693) 3.244	(2.56 to 3.923) 3.228																					
Seeking inferst dustic symbolic	3000) g Birth prevalence - [38, 40) wks, [2500,	Manuliu	Malar		(2.486 to 4.159) 2.755	(2.605 to 3.985) 2.944																					
	3000) g Birth prevalence - [38, 40) wks, [2500,	Mariany	E		(2.274 to 3.309) 2.376	(2.44 to 3.548) 2.938																					
Staten mun acan synatone	3000) g Birth menalence - [36-37) wise [3000	stortany	Pemases		(1.91 to 2.886) 3.774	(2.434 to 3.503) 2.466																					
Sudden infant death syndrome	3500) g	Mortality	Males		(3.094 to 4.497)	(2.058 to 2.929)																					
Sudden infant death syndrome	3500) g	Mortality	Females		(2.981 to 4.646)	(2.277 to 3.218)																					
Sudden infant death syndrome	Barth prevalence - [36, 37) wks, [4000, 4500) g	Mortality	Males		6.826 (5.212 to 9.045)	1.77 (1.491 to 2.082)																					
Sudden infant death syndrome	Birth prevalence - [36, 37] wks, [4000, 4500) g	Mortality	Females		7.269 (5.144 to 9.821)	2.177 (1.786 to 2.607)																					
Sudden infant death syndrome	Birth prevalence - [36, 37) wks, [3500, 4000) g	Mortality	Males		4.544 (3.64 to 5.622)	2.057 (1.74 to 2.44)																					
Sudden infant death syndrome	Birth prevalence - [36, 37) wks, [3500, 4000) g	Mortality	Females		4.662 (3.577 to 6.014)	2.398 (1.98 to 2.864)																					
Sudden infant death syndrome	Birth prevalence - [37, 38) wks, [3000, 3500) g	Mortality	Males		2.007 (1.759 to 2.293)	1.888 (1.613 to 2.224)																					
Sudden infant death syndrome	Birth prevalence - [37, 38) wks, [3000, 3500) g	Mortality	Females		1.925 (1.582 to 2.328)	1.972 (1.669 to 2.313)																					
Sudden infant death syndrome	Birth prevalence - [37, 38) wks, [4000, 45000 m	Mortality	Males		3.28 (2.595 to 4.133)	1.335 (1.171 to 1.532)																					
Sudden infant death syndrome	Birth prevalence - [37, 38) wks, [4000, 46000 -	Mortality	Females		3.521	1.559																					
Sudden infant death syndrome	Birth prevalence - [37, 38) wks, [3500,	Mortality	Males		2.128	1.505																					
Sudden infant death syndrome	4000) g Birth prevalence - [37, 38) wks, [3500,	Mortality	Females		2.142	1.661																					
Sudden infort death conde	4000) g Birth prevalence - [40, 42) wks, [3000,	Magnality	Male		(1.694 to 2.67) 1.436	(1.411 to 1.961) 1.47																					
	3500) g Birth prevalence - [40, 42) wks, [3000,				(1.245 to 1.65) 1.326	(1.199 to 1.8) 1.465																					
Souther information	3500) g Birth prevalence - [38, 40) wks, [3000.	Mar 11	Mal		(1.069 to 1.614) 1.33	(1.188 to 1.775) 1.559																					
suusaen intiati death syndrome	3500) g Birth menalence - [38, 40) wise 13000	soutidity	MIRES		(1.155 to 1.53) 1.224	(1.305 to 1.851) 1.564																					
Sudden infant death syndrome	3500)g Bishamahan (28.40)ala (200	Mortality	Females		(1.0 to 1.492)	(1.304 to 1.847)																					
Sudden infant death syndrome	4500) g	Mortality	Males		(1.453 to 2.182)	(1.005 to 1.371)																					
Sudden infant death syndrome	north prevalence - [38, 40) wks, [4000, 4500) g	Mortality	Females		1.877 (1.467 to 2.388)	1.224 (1.022 to 1.465)																					
Sudden infant death syndrome	Birth prevalence - [38, 40) wks, [3500, 4000) g	Mortality	Males		1.785 (1.478 to 2.147)	1.173 (1.0 to 1.377)																					
Sudden infant death syndrome	Birth prevalence - [38, 40) wks, [3500, 4000) g	Mortality	Females		1.892 (1.481 to 2.352)	1.23 (1.03 to 1.46)																					
Sudden infant death syndrome	Birth prevalence - [40, 42) wks, [3500, 4000) g	Mortality	Males		1.0 (1.0 to 1.0)	1.003 (1.0 to 1.046)																					
Sudden infant death syndrome	Birth prevalence - [40, 42) wks, [3500, 4000) g	Mortality	Females		1.002 (1.0 to 1.013)	1.001 (1.0 to 1.006)																					
Sudden infant death syndrome	Birth prevalence - [40, 42) wks, [4000, 4500) g	Mortality	Males		1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																					
Sudden infant death syndrome	- Birth prevalence - [40, 42) wks, [4000, 45000 m	Mortality	Females		1.0	1.0																					
Sudden infant death syndrome	Birth prevalence - [28, 30] wks, [2000, 2500) -	Mortality	Males		117.172	27.726																					
Sudden infant death syndrome	Birth prevalence - [28, 30] wks, [2000,	Mortality	Females		(87.695 10 156.056) 121.682	33.983																					
Sudden infant death sources	2500) g Birth prevalence - [28, 30) wks, [2500,	Mortality	Males		(84.549 to 171.375) 77.948	(20.101 to 43.404) 16.608																					
Souther infort doubles	3000) g Birth prevalence - [28, 30) wks, [2500,	Mastality	Employ		(54.687 to 105.047) 79.193	(12.653 to 21.188) 20.387																					
Subarn intant arath syndrome	3000) g Birth prevalence - [28, 30) wks. [3000	Morany	. emases		(54.099 to 112.236) 42.199	(15.089 to 26.434) 10.082																					
Sudden infant death syndrome	3500) g Birth menalence - [28, 30) wise 13000	Mortality	Males		(29.891 to 57.227) 42.551	(7.777 to 13.056) 11.989																					
Sudden infant death syndrome	3500) g	Mortality	Females	ļ	(28.25 to 63.209)	(9.084 to 15.544)	I																				

Appendix Table 6a. Relative risk	s used by age and sex for each ou	come for all risk factors ex	ept for ambient ai	r pollution alcohol,	and smoking.									А	iges											
Risk - Outcome	Category / Units	Morbidity / Mortality	ex All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Sudden infant death syndrome	Birth prevalence - [30, 32) wks, [2500, 3000) g	Mortality N	ales	58.722 (42.419 to 78.873)	12.115 (9.518 to 15.521)																					
Sudden infant death syndrome	Birth prevalence - [30, 32) wks, [2500, 3000) g	Mortality Fe	nales	59.522 (42.058 to 82.793)	15.364 (11.936 to 19.581)																					
Sudden infant death syndrome	Birth prevalence - [30, 32) wks, [3000, 3500) g	Mortality N	ales	45.67 (32.014 to 65.531)	8.381 (6.362 to 10.842)																					
Sudden infant death syndrome	Birth prevalence - [30, 32) wks, [3000, 3500) g	Mortality Fe	nales	46.104 (30.243 to 66.207)	10.506 (8.041 to 13.513)																					
Sudden infant death syndrome	Birth prevalence - [30, 32) wks, [3500, 4000) g	Mortality N	ales	36.334 (21.558 to 54.813)	5.698 (4.293 to 7.349)																					
Sudden infant death syndrome	Birth prevalence - [30, 32) wks, [3500, 4000) g	Mortality Fe	nales	37.931 (22.692 to 61.276)	6.892 (5.175 to 9.058)																					
Sudden infant death syndrome	Birth prevalence - [32, 34) wks, [3000, 3500) g	Mortality N	ales	34.016 (23.515 to 48.37)	6.577 (4.934 to 8.436)																					
Sudden infant death syndrome	Birth prevalence - [32, 34) wks, [3000, 3500) g	Mortality Fe	nales	34.585 (22.909 to 49.754)	8.314 (6.362 to 10.789)																					
Sudden infant death syndrome	Birth prevalence - [32, 34) wks, [3500, 4000) g	Mortality N	ales	36.248 (23.158 to 54.67)	5.068 (3.761 to 6.741)																					
Sudden infant death syndrome	Birth prevalence - [32, 34) wks, [3500, 4000) g	Mortality Fe	nales	38.098 (23.301 to 59.429)	6.476 (4.666 to 8.689)																					
Sudden infant death syndrome	Birth prevalence - [36, 37) wks, [1000, 1500) g	Mortality N	ales	166.686 (118.487 to 222.581)	57.535 (45.999 to 71.742)																					
Sudden infant death syndrome	Birth prevalence - [36, 37) wks, [1000, 1500) g	Mortality Fe	nales	169.725 (119.017 to 229.008)	63.564) (50.068 to 80.703)																					
Sudden infant death syndrome	Birth prevalence - [38, 40) wks, [1000, 1500) g	Mortality N	ales	174.066 (125.125 to 232.507)	57.966 (44.393 to 73.241)																					
Sudden infant death syndrome	Birth prevalence - [38, 40) wks, [1000, 1500) g	Mortality Fe	nales	171.557 (121.585 to 237.047)	65.208 (48.821 to 84.308)																					
Sudden infant death syndrome	Birth prevalence - [38, 40) wks, [1500, 2000) g	Mortality N	ales	67.302 (49.547 to 89.055)	25.206 (20.365 to 31.168)																					
Sudden infant death syndrome	Birth prevalence - [38, 40) wks, [1500, 2000) g	Mortality Fe	nales	62.19 (45.884 to 83.445)	28.05 (22.625 to 35.139)																					
Sudden infant death syndrome	Birth prevalence - [40, 42) wks, [1500, 2000) g	Mortality N	ales	76.673 (56.177 to 102.468)	25.785 (19.387 to 34.168)																					
Sudden infant death syndrome	Birth prevalence - [40, 42) wks, [1500, 2000) g	Mortality Fe	nales	70.411 (49.221 to 97.952)	29.113 (21.355 to 38.272)																					
Iron deficiency																										
Maternal haemorrhage	1 g'dL	Both F	oth						1.252 (1.087 to 1.425)																	
Maternal sepsis and other pregna related infections	mcy 1 g dL	Both F	oth						1.252 (1.087 to 1.425)																	
Maternal hypertensive disorders	1 g/dL	Both F	oth						1.252 (1.087 to 1.425)																	
Maternal obstructed labour and uterine rupture	1 g'dL	Both E	oth						1.252 (1.087 to 1.425)																	
Maternal abortive outcome	1 g'dL	Both E	oth						1.252 (1.087 to 1.425)																	
Ectopic prognancy	1 gidL	Both F	oth						1.252 (1.087 to 1.425)																	
Indirect maternal deaths	1 gidL	Both F	oth						1.252 (1.087 to 1.425)																	
Late maternal deaths	1 gidL	Both F	oth						1.252 (1.087 to 1.425)																	
Maternal deaths aggravated by HIV/AIDS	1 gidL	Both F	oth						1.252 (1.087 to 1.425)																	
Other maternal disorders	1 g'dL	Both F	oth						1.252 (1.087 to 1.425)																	
Vitamin A deficiency																										
Lower respiratory infections	Vitamin A deficient	Both F	oth			1.33 (1.114 to 1.585)	1.33 (1.114 to 1.585)																			
Lower respiratory infections	Not deficient	Both F	oth			1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																			
Diarrhoeal diseases	Vitamin A deficient	Both E	oth			2.444 (2.268 to 2.626)	2.444 (2.268 to 2.626)																			
Diarrhoeal diseases	Not deficient	Both E	oth			1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																			
Measles	Vitamin A deficient	Both F	oth			3.515 (2.526 to 4.671)	3.515 (2.526 to 4.671)																			
Measles	Not deficient	Both E	oth			1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																			
Zinc deficiency																										
Lower respiratory infections	Zinc deficient	Morbidity E	oth				1.837 (1.275 to 2.523)																			
Lower respiratory infections	Zinc deficient	Mortality E	oth				1.672 (0.458 to 4.135)																			
Lower respiratory infections	Not deficient	Both F	oth				1.0 (1.0 to 1.0)																			
Diarrhoeal diseases	Zinc deficient	Morbidity E	oth				1.903 (1.517 to 2.335)																			
Diarrhoeal diseases	Zinc deficient	Mortality E	oth				1.951 (0.905 to 3.909)																			
Diarrhoeal diseases	Not deficient	Both F	oth				1.0 (1.0 to 1.0)																			
Chewing tobacco																										
Lip and oral cavity cancer	Exposed	Both M	ales										3.021 (2.015 to 4.47)													
Lip and oral cavity cancer	Exposed	Both Fe	nales										8.09 (5.463 to 11.767)													
Oesophageal cancer	Exposed	Both F	oth	-									2.571 (1.854 to 3.416)													
Second-hand smoke																										
Otitis media	Exposed	Morbidity E	oth	1.37 (1.238 to 1.493)	1.37 (1.238 to 1.493)	1.37 (1.238 to 1.493)	1.37 (1.238 to 1.493)	1.37 (1.238 to 1.493)	1.37 (1.238 to 1.493)																	
Otitis media	Exposed	Mortality E	oth	1.37 (1.238 to 1.493)	1.37 (1.238 to 1.493)	1.37 (1.238 to 1.493)	1.37 (1.238 to 1.493)	1.37 (1.238 to 1.493)	1.37 (1.238 to 1.493)																	
Otitis media	Not exposed	Morbidity E	oth	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)																	
Otitis media	Not exposed	Mortality E	oth	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)						1	1			i.							
Breast cancer	Exposed	Morbidity E	oth									1.072 (1.017 to 1.126)														
Breast cancer	Exposed	Mortality F	oth									1.072 (1.017 to 1.126)														
Breast cancer	Not exposed	Morbidity E	oth									1.0 (1.0 to 1.0)														
Breast cancer	Not exposed	Mortality E	oth									1.0 (1.0 to 1.0)														
Dict low in fruits																										

Appendix Table 6a. Relative risk	is used by age and sex for each oute	come for all risk factors exce	pt for ambient air p	ollution alcohol, a	nd smoking.									A	ges											
Rick - Ontroma	Catanory / Unite	Marbidity / Martality Se	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Lip and oral cavity cancer	100 giday	Both Bo	ú.			I	I			1		1.042 (0.994 to 1.091)	1.042 (0.994 to 1.091)	1.042 (0.994 to 1.091)	1.042 (0.994 to 1.091)	1.042 (0.994 to 1.091)	1.042 (0.994 to 1.091)	1.042 (0.994 to 1.091)	1.042 (0.994 to 1.091)	1.042 (0.994 to 1.091)	1.042 (0.994 to 1.091)	1.042 (0.994 to 1.091)	1.042 (0.994 to 1.091)	1.042 (0.994 to 1.091)	1.042 (0.994 to 1.091)	1.042 (0.994 to 1.091)
Nasopharynx cancer	100 giday	Both Bo	th									1.043 (0.992 to 1.092)	1.043 (0.992 to 1.092)	1.043 (0.992 to 1.092)	1.043 (0.992 to 1.092)	1.043 (0.992 to 1.092)	1.043 (0.992 to 1.092)	1.043 (0.992 to 1.092)	1.043 (0.992 to 1.092)	1.043 (0.992 to 1.092)	1.043 (0.992 to 1.092)	1.043 (0.992 to 1.092)	1.043 (0.992 to 1.092)	1.043 (0.992 to 1.092)	1.043 (0.992 to 1.092)	1.043 (0.992 to 1.092)
Other pharynx cancer	100 gʻday	Both Bo	th.									1.042 (0.996 to 1.095)	1.042 (0.996 to 1.095)	1.042 (0.996 to 1.095)	1.042 (0.996 to 1.095)	1.042 (0.996 to 1.095)	1.042 (0.996 to 1.095)	1.042 (0.996 to 1.095)	1.042 (0.996 to 1.095)	1.042 (0.996 to 1.095)	1.042 (0.996 to 1.095)	1.042 (0.996 to 1.095)	1.042 (0.996 to 1.095)	1.042 (0.996 to 1.095)	1.042 (0.996 to 1.095)	1.042 (0.996 to 1.095)
Oesophageal cancer	100 giday	Both Bo	th									1.153 (1.033 to 1.288)	1.153 (1.033 to 1.288)	1.153 (1.033 to 1.288)	1.153 (1.033 to 1.288)	1.153 (1.033 to 1.288)	1.153 (1.033 to 1.288)	1.153 (1.033 to 1.288)	1.153 (1.033 to 1.288)	1.153 (1.033 to 1.288)	1.153 (1.033 to 1.288)	1.153 (1.033 to 1.288)	1.153 (1.033 to 1.288)	1.153 (1.033 to 1.288)	1.153 (1.033 to 1.288)	1.153 (1.033 to 1.288)
Larynx cancer	100 gʻday	Both Bo	th.									1.042 (0.995 to 1.095)	1.042 (0.995 to 1.095)	1.042 (0.995 to 1.095)	1.042 (0.995 to 1.095)	1.042 (0.995 to 1.095)	1.042 (0.995 to 1.095)	1.042 (0.995 to 1.095)	1.042 (0.995 to 1.095)	1.042 (0.995 to 1.095)	1.042 (0.995 to 1.095)	1.042 (0.995 to 1.095)	1.042 (0.995 to 1.095)	1.042 (0.995 to 1.095)	1.042 (0.995 to 1.095)	1.042 (0.995 to 1.095)
Tracheal, bronchus, and lung ca	ncer 100 g/day	Both Bo	th.									1.076 (1.031 to 1.123)	1.076 (1.031 to 1.123)	1.076 (1.031 to 1.123)	1.076 (1.031 to 1.123)	1.076 (1.031 to 1.123)	1.076 (1.031 to 1.123)	1.076 (1.031 to 1.123)	1.076 (1.031 to 1.123)	1.076 (1.031 to 1.123)	1.076 (1.031 to 1.123)	1.076 (1.031 to 1.123)	1.076 (1.031 to 1.123)	1.076 (1.031 to 1.123)	1.076 (1.031 to 1.123)	1.076 (1.031 to 1.123)
Ischaemic heart disease	100 giday	Both Bo	th									1.254 (1.083 to 1.442)	1.209 (1.07 to 1.361)	1.159 (1.054 to 1.271)	1.131 (1.045 to 1.221)	1.125 (1.043 to 1.211)	1.114 (1.039 to 1.193)	1.099 (1.034 to 1.167)	1.087 (1.03 to 1.146)	1.078 (1.027 to 1.13)	1.07 (1.025 to 1.117)	1.064 (1.022 to 1.106)	1.057 (1.02 to 1.095)	1.057 (1.02 to 1.095)	1.057 (1.02 to 1.095)	1.057 (1.02 to 1.095)
Ischaemic stroke	100 giday	Both Bo	th									2.024 (1.465 to 2.818)	1.834 (1.39 to 2.444)	1.621 (1.301 to 2.043)	1.48 (1.239 to 1.787)	1.403 (1.204 to 1.653)	1.333 (1.171 to 1.533)	1.272 (1.142 to 1.432)	1.222 (1.116 to 1.348)	1.181 (1.096 to 1.283)	1.145 (1.078 to 1.225)	1.114 (1.061 to 1.175)	1.054 (1.029 to 1.082)	1.054 (1.029 to 1.082)	1.054 (1.029 to 1.082)	1.054 (1.029 to 1.082)
Intracerebral hemorrhage	100 giday	Both Bo	th									1.688 (1.319 to 2.182)	1.576 (1.273 to 1.972)	1.444 (1.215 to 1.732)	1.365 (1.18 to 1.595)	1.336 (1.167 to 1.544)	1.3 (1.15 to 1.483)	1.26 (1.131 to 1.415)	1.226 (1.115 to 1.358)	1.193 (1.099 to 1.305)	1.164 (1.084 to 1.256)	1.133 (1.069 to 1.207)	1.065 (1.034 to 1.1)	1.065 (1.034 to 1.1)	1.065 (1.034 to 1.1)	1.065 (1.034 to 1.1)
Suburachnoid hemoerhage	100 gʻday	Both Bo	th.									1.688 (1.319 to 2.182)	1.576 (1.273 to 1.972)	1.444 (1.215 to 1.732)	1.365 (1.18 to 1.595)	1.336 (1.167 to 1.544)	1.3 (1.15 to 1.483)	1.26 (1.131 to 1.415)	1.226 (1.115 to 1.358)	1.193 (1.099 to 1.305)	1.164 (1.084 to 1.256)	1.133 (1.069 to 1.207)	1.065 (1.034 to 1.1)	1.065 (1.034 to 1.1)	1.065 (1.034 to 1.1)	1.065 (1.034 to 1.1)
Diabetes mellitus type 2	100 giday	Both Bo	th.									1.125 (1.027 to 1.238)	1.122 (1.026 to 1.232)	1.119 (1.026 to 1.226)	1.113 (1.024 to 1.214)	1.102 (1.022 to 1.194)	1.093 (1.02 to 1.176)	1.085 (1.019 to 1.16)	1.076 (1.017 to 1.143)	1.068 (1.015 to 1.128)	1.061 (1.014 to 1.114)	1.052 (1.012 to 1.098)	1.036 (1.008 to 1.066)	1.036 (1.008 to 1.066)	1.036 (1.008 to 1.066)	1.036 (1.008 to 1.066)
Diet low in vegetables																										
Ischaemic heart disease	100 giday	Both Bo	th.									1.249 (1.089 to 1.446)	1.205 (1.074 to 1.362)	1.154 (1.057 to 1.269)	1.126 (1.047 to 1.219)	1.121 (1.045 to 1.21)	1.111 (1.042 to 1.193)	1.098 (1.037 to 1.168)	1.086 (1.032 to 1.148)	1.077 (1.029 to 1.133)	1.07 (1.027 to 1.12)	1.064 (1.024 to 1.109)	1.057 (1.022 to 1.097)	1.057 (1.022 to 1.097)	1.057 (1.022 to 1.097)	1.057 (1.022 to 1.097)
Ischaemic stroke	100 giday	Both Bo	th.									1.249 (1.049 to 1.463)	1.211 (1.042 to 1.388)	1.165 (1.033 to 1.3)	1.132 (1.027 to 1.238)	1.113 (1.023 to 1.203)	1.095 (1.02 to 1.17)	1.079 (1.017 to 1.141)	1.065 (1.014 to 1.116)	1.054 (1.012 to 1.096)	1.044 (1.009 to 1.077)	1.035 (1.007 to 1.061)	1.017 (1.004 to 1.029)	1.017 (1.004 to 1.029)	1.017 (1.004 to 1.029)	1.017 (1.004 to 1.029)
Intracerebral hemorrhage	100 gʻday	Both Bo	th.									1.177 (1.046 to 1.326)	1.153 (1.04 to 1.278)	1.122 (1.032 to 1.22)	1.102 (1.027 to 1.184)	1.095 (1.025 to 1.17)	1.086 (1.023 to 1.153)	1.075 (1.02 to 1.134)	1.066 (1.018 to 1.117)	1.057 (1.015 to 1.101)	1.049 (1.013 to 1.086)	1.04 (1.011 to 1.071)	1.02 (1.005 to 1.035)	1.02 (1.005 to 1.035)	1.02 (1.005 to 1.035)	1.02 (1.005 to 1.035)
Suburachnoid hemoerhage	100 gʻday	Both Bo	th.									1.177 (1.046 to 1.326)	1.153 (1.04 to 1.278)	1.122 (1.032 to 1.22)	1.102 (1.027 to 1.184)	1.095 (1.025 to 1.17)	1.086 (1.023 to 1.153)	1.075 (1.02 to 1.134)	1.066 (1.018 to 1.117)	1.057 (1.015 to 1.101)	1.049 (1.013 to 1.086)	1.04 (1.011 to 1.071)	1.02 (1.005 to 1.035)	1.02 (1.005 to 1.035)	1.02 (1.005 to 1.035)	1.02 (1.005 to 1.035)
Diet low in legumes																										
Ischaemic heart disease	50 g/day	Both Bo	th									1.499 (1.18 to 1.89)	1.453 (1.166 to 1.801)	1.388 (1.144 to 1.677)	1.332 (1.125 to 1.573)	1.287 (1.11 to 1.49)	1.237 (1.092 to 1.401)	1.181 (1.071 to 1.303)	1.139 (1.055 to 1.23)	1.111 (1.045 to 1.183)	1.089 (1.036 to 1.146)	1.074 (1.03 to 1.12)	1.101 (1.041 to 1.165)	1.101 (1.041 to 1.165)	1.101 (1.041 to 1.165)	1.101 (1.041 to 1.165)
Diet low in whole grains																										
Ischaemic heart disease	50 g/day	Both Bo	th									1.478 (1.274 to 1.722)	1.387 (1.225 to 1.578)	1.285 (1.168 to 1.418)	1.228 (1.136 to 1.333)	1.216 (1.129 to 1.313)	1.194 (1.117 to 1.281)	1.165 (1.1 to 1.238)	1.141 (1.086 to 1.203)	1.125 (1.076 to 1.179)	1.112 (1.068 to 1.16)	1.102 (1.062 to 1.145)	1.097 (1.059 to 1.138)	1.097 (1.059 to 1.138)	1.097 (1.059 to 1.138)	1.097 (1.059 to 1.138)
Ischaemic stroke	50 g/day	Both Bo	th									2.075 (1.669 to 2.517)	1.863 (1.548 to 2.199)	1.624 (1.406 to 1.849)	1.466 (1.309 to 1.625)	1.38 (1.255 to 1.505)	1.304 (1.206 to 1.401)	1.241 (1.165 to 1.316)	1.189 (1.13 to 1.247)	1.15 (1.104 to 1.195)	1.117 (1.081 to 1.151)	1.09 (1.063 to 1.116)	1.041 (1.029 to 1.053)	1.041 (1.029 to 1.053)	1.041 (1.029 to 1.053)	1.041 (1.029 to 1.053)
Intracerebral hemorrhage	50 g/day	Both Bo	th									1.596 (1.406 to 1.825)	1.484 (1.333 to 1.662)	1.349 (1.244 to 1.471)	1.276 (1.194 to 1.369)	1.258 (1.182 to 1.344)	1.232 (1.165 to 1.309)	1.201 (1.143 to 1.267)	1.176 (1.126 to 1.233)	1.15 (1.108 to 1.198)	1.128 (1.092 to 1.169)	1.106 (1.076 to 1.139)	1.05 (1.036 to 1.065)	1.05 (1.036 to 1.065)	1.05 (1.036 to 1.065)	1.05 (1.036 to 1.065)
Subarachaoid hemoerhage	50 g/day	Both Bo	th.									1.596 (1.406 to 1.825)	1.484 (1.333 to 1.662)	1.349 (1.244 to 1.471)	1.276 (1.194 to 1.369)	1.258 (1.182 to 1.344)	1.232 (1.165 to 1.309)	1.201 (1.143 to 1.267)	1.176 (1.126 to 1.233)	1.15 (1.108 to 1.198)	1.128 (1.092 to 1.169)	1.106 (1.076 to 1.139)	1.05 (1.036 to 1.065)	1.05 (1.036 to 1.065)	1.05 (1.036 to 1.065)	1.05 (1.036 to 1.065)
Diabetes mellitus type 2	50 g/day	Both Bo	th.									1.231 (1.125 to 1.349)	1.226 (1.122 to 1.341)	1.22 (1.119 to 1.331)	1.208 (1.113 to 1.313)	1.189 (1.103 to 1.283)	1.172 (1.094 to 1.256)	1.156 (1.085 to 1.232)	1.139 (1.077 to 1.207)	1.125 (1.069 to 1.184)	1.111 (1.061 to 1.163)	1.095 (1.053 to 1.14)	1.064 (1.036 to 1.094)	1.064 (1.036 to 1.094)	1.064 (1.036 to 1.094)	1.064 (1.036 to 1.094)
Diet low in nuts and seeds																										
Ischaemic heart disease	4.05 g/day	Morbidity Bo	th									1.176 (1.055 to 1.322)	1.143 (1.045 to 1.259)	1.105 (1.033 to 1.188)	1.084 (1.027 to 1.15)	1.081 (1.026 to 1.144)	1.074 (1.024 to 1.132)	1.064 (1.021 to 1.114)	1.056 (1.018 to 1.099)	1.05 (1.016 to 1.089)	1.046 (1.015 to 1.081)	1.042 (1.014 to 1.075)	1.039 (1.013 to 1.069)	1.039 (1.013 to 1.069)	1.039 (1.013 to 1.069)	1.039 (1.013 to 1.069)
Ischaemic heart disease	4.05 g/day	Mortality Bo	th.									1.209 (1.128 to 1.296)	1.169 (1.105 to 1.239)	1.124 (1.077 to 1.174)	1.099 (1.062 to 1.138)	1.095 (1.06 to 1.133)	1.088 (1.055 to 1.122)	1.076 (1.048 to 1.105)	1.066 (1.042 to 1.092)	1.059 (1.037 to 1.082)	1.054 (1.034 to 1.075)	1.05 (1.032 to 1.069)	1.046 (1.029 to 1.064)	1.046 (1.029 to 1.064)	1.046 (1.029 to 1.064)	1.046 (1.029 to 1.064)
Diabetes mellitus type 2	4.05 g/day	Both Bo	th.									1.05 (1.025 to 1.075)	1.049 (1.025 to 1.073)	1.048 (1.024 to 1.071)	1.045 (1.023 to 1.068)	1.041 (1.021 to 1.062)	1.038 (1.019 to 1.056)	1.035 (1.018 to 1.052)	1.031 (1.016 to 1.046)	1.028 (1.014 to 1.042)	1.025 (1.013 to 1.037)	1.022 (1.011 to 1.032)	1.015 (1.007 to 1.022)	1.015 (1.007 to 1.022)	1.015 (1.007 to 1.022)	1.015 (1.007 to 1.022)
Diet low in milk																										
Colon and rectum cancer	226.8 g/day	Both Bo	th									1.113 (1.039 to 1.202)	1.113 (1.039 to 1.202)	1.113 (1.039 to 1.202)	1.113 (1.039 to 1.202)	1.113 (1.039 to 1.202)	1.113 (1.039 to 1.202)	1.113 (1.039 to 1.202)	1.113 (1.039 to 1.202)	1.113 (1.039 to 1.202)	1.113 (1.039 to 1.202)	1.113 (1.039 to 1.202)	1.113 (1.039 to 1.202)	1.113 (1.039 to 1.202)	1.113 (1.039 to 1.202)	1.113 (1.039 to 1.202)
Diet high in red meat																										
Colon and rectum cancer	100 giday	Both Bo	th									1.167 (1.033 to 1.309)	1.167 (1.033 to 1.309)	1.167 (1.033 to 1.309)	1.167 (1.033 to 1.309)	1.167 (1.033 to 1.309)	1.167 (1.033 to 1.309)	1.167 (1.033 to 1.309)	1.167 (1.033 to 1.309)	1.167 (1.033 to 1.309)	1.167 (1.033 to 1.309)	1.167 (1.033 to 1.309)	1.167 (1.033 to 1.309)	1.167 (1.033 to 1.309)	1.167 (1.033 to 1.309)	1.167 (1.033 to 1.309)
Diabetes mellitus type 2	100 gʻday	Both Bo	th.									1.322 (1.037 to 1.603)	1.314 (1.036 to 1.588)	1.305 (1.035 to 1.569)	1.288 (1.034 to 1.536)	1.26 (1.031 to 1.48)	1.236 (1.028 to 1.433)	1.213 (1.026 to 1.389)	1.19 (1.023 to 1.345)	1.169 (1.021 to 1.306)	1.15 (1.019 to 1.269)	1.128 (1.016 to 1.229)	1.086 (1.011 to 1.152)	1.086 (1.011 to 1.152)	1.086 (1.011 to 1.152)	1.086 (1.011 to 1.152)
Diet high in processed meat																										
Colon and rectum cancer	50 g/day	Both Bo	th									1.179 (1.093 to 1.267)	1.179 (1.093 to 1.267)	1.179 (1.093 to 1.267)	1.179 (1.093 to 1.267)	1.179 (1.093 to 1.267)	1.179 (1.093 to 1.267)	1.179 (1.093 to 1.267)	1.179 (1.093 to 1.267)	1.179 (1.093 to 1.267)	1.179 (1.093 to 1.267)	1.179 (1.093 to 1.267)	1.179 (1.093 to 1.267)	1.179 (1.093 to 1.267)	1.179 (1.093 to 1.267)	1.179 (1.093 to 1.267)
Ischaemic heart disease	50 g/day	Both Bo	th									2.568 (1.047 to 4.657)	2.124 (1.038 to 3.478)	1.72 (1.028 to 2.489)	1.545 (1.022 to 2.093)	1.547 (1.022 to 2.097)	1.52 (1.022 to 2.037)	1.467 (1.02 to 1.922)	1.422 (1.018 to 1.826)	1.386 (1.017 to 1.75)	1.354 (1.016 to 1.683)	1.325 (1.015 to 1.622)	1.252 (1.012 to 1.475)	1.252 (1.012 to 1.475)	1.252 (1.012 to 1.475)	1.252 (1.012 to 1.475)
Diabetes mellitus type 2	50 g/day	Both Bo	sh									1.94 (1.395 to 2.545)	1.913 (1.386 to 2.496)	1.881 (1.375 to 2.439)	1.824 (1.354 to 2.337)	1.731 (1.319 to 2.173)	1.653 (1.289 to 2.038)	1.583 (1.261 to 1.918)	1.512 (1.233 to 1.798)	1.45 (1.207 to 1.696)	1.393 (1.183 to 1.603)	1.332 (1.157 to 1.505)	1.216 (1.105 to 1.323)	1.216 (1.105 to 1.323)	1.216 (1.105 to 1.323)	1.216 (1.105 to 1.323)
Diet high in sugar-sweetened	beverages																									
Ischaemic heart disease	2.5 giday	Both Bo	th									1.377 (0.933 to 1.883)	1.311 (0.943 to 1.717)	1.232 (0.955 to 1.521)	1.195 (0.961 to 1.436)	1.186 (0.963 to 1.413)	1.172 (0.965 to 1.381)	1.156 (0.968 to 1.343)	1.14 (0.971 to 1.306)	1.124 (0.974 to 1.27)	1.11 (0.977 to 1.238)	1.095 (0.98 to 1.205)	1.067 (0.985 to 1.143)	1.067 (0.985 to 1.143)	1.067 (0.985 to 1.143)	1.067 (0.985 to 1.143)
Diabetes mellitus type 2	2.5 giday	Both Bo	th									1.263 (1.129 to 1.4)	1.257 (1.126 to 1.39)	1.25 (1.123 to 1.379)	1.237 (1.117 to 1.358)	1.214 (1.106 to 1.322)	1.195 (1.097 to 1.292)	1.177 (1.088 to 1.264)	1.158 (1.079 to 1.235)	1.141 (1.071 to 1.209)	1.125 (1.063 to 1.185)	1.107 (1.055 to 1.158)	1.08 (1.041 to 1.118)	1.08 (1.041 to 1.118)	1.08 (1.041 to 1.118)	1.08 (1.041 to 1.118)
Diet low in fibre																										
Colon and rectum cancer	20 g/day	Both Bo	th									1.236 (1.133 to 1.35)	1.236 (1.133 to 1.35)	1.236 (1.133 to 1.35)	1.236 (1.133 to 1.35)	1.236 (1.133 to 1.35)	1.236 (1.133 to 1.35)	1.236 (1.133 to 1.35)	1.236 (1.133 to 1.35)	1.236 (1.133 to 1.35)	1.236 (1.133 to 1.35)	1.236 (1.133 to 1.35)	1.236 (1.133 to 1.35)	1.236 (1.133 to 1.35)	1.236 (1.133 to 1.35)	1.236 (1.133 to 1.35)
Ischaemic heart disease	20 g/day	Both Bo	th									1.688 (1.415 to 2.028)	1.622 (1.379 to 1.922)	1.529 (1.326 to 1.776)	1.45 (1.28 to 1.654)	1.387 (1.243 to 1.558)	1.318 (1.202 to 1.455)	1.242 (1.156 to 1.342)	1.184 (1.119 to 1.258)	1.147 (1.096 to 1.205)	1.118 (1.077 to 1.163)	1.097 (1.064 to 1.135)	1.133 (1.087 to 1.185)	1.133 (1.087 to 1.185)	1.133 (1.087 to 1.185)	1.133 (1.087 to 1.185)
Diet low in calcium																										
Colon and rectum cancer	1 giday	Both Bo	th									1.372 (1.268 to 1.485)	1.372 (1.268 to 1.485)	1.372 (1.268 to 1.485)	1.372 (1.268 to 1.485)	1.372 (1.268 to 1.485)	1.372 (1.268 to 1.485)	1.372 (1.268 to 1.485)	1.372 (1.268 to 1.485)	1.372 (1.268 to 1.485)	1.372 (1.268 to 1.485)	1.372 (1.268 to 1.485)	1.372 (1.268 to 1.485)	1.372 (1.268 to 1.485)	1.372 (1.268 to 1.485)	1.372 (1.268 to 1.485)
Diet low in seafood omega-3	fatty acids																									
Ischaemic heart disease	100 mg/day	Morbidity Bo	th									1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Ischaemic heart disease	100 mg/day	Mortality Bo	th									1.291 (1.109 to 1.505)	1.249 (1.094 to 1.428)	1.199 (1.077 to 1.338)	1.173 (1.067 to 1.293)	1.165 (1.064 to 1.279)	1.154 (1.06 to 1.26)	1.14 (1.055 to 1.235)	1.126 (1.05 to 1.211)	1.113 (1.045 to 1.189)	1.101 (1.04 to 1.167)	1.088 (1.035 to 1.145)	1.062 (1.025 to 1.102)	1.062 (1.025 to 1.102)	1.062 (1.025 to 1.102)	1.062 (1.025 to 1.102)
Diet low in polyunsaturated	fatty acids																									
Ischaemic heart disease	5% energy/day	Both Bo	û.									1.267 (1.098 to 1.452)	1.211 (1.079 to 1.352)	1.148 (1.056 to 1.244)	1.114 (1.044 to 1.186)	1.111 (1.043 to 1.181)	1.101 (1.039 to 1.165)	1.086 (1.033 to 1.14)	1.075 (1.029 to 1.121)	1.068 (1.026 to 1.11)	1.063 (1.025 to 1.102)	1.06 (1.024 to 1.097)	1.063 (1.025 to 1.102)	1.063 (1.025 to 1.102)	1.063 (1.025 to 1.102)	1.063 (1.025 to 1.102)
Diet high in trans fatty acids																										
Ischaemic heart disease	2% energy/day	Both Bo	th.					-				1.901 (1.591 to 2.275)	1.775 (1.514 to 2.085)	1.615 (1.415 to 1.848)	1.517 (1.352 to 1.707)	1.461 (1.316 to 1.627)	1.396 (1.274 to 1.535)	1.323 (1.225 to 1.433)	1.264 (1.186 to 1.352)	1.222 (1.157 to 1.294)	1.186 (1.132 to 1.246)	1.158 (1.112 to 1.207)	1.15 (1.107 to 1.197)	1.15 (1.107 to 1.197)	1.15 (1.107 to 1.197)	1.15 (1.107 to 1.197)
Dict high in sodium **																										
	Non-Black, Non-Hypertensive	Both Bo	th.								(-1.366 -1.937 to -0.795)	-1.882 (-2.434 to -1.330)	-2.397 (-2.967 to -1.828)	-2.913 (-3.533 to -2.292)	-3.428 (-4.126 to -2.730)	-3.944 (-4.738 to -3.150)	-4.459 (-5.362 to -3.556)	-4.975 (-5.995 to -3.954)	-5.490 (-6.634 to -4.347)						
]	Non-Black, Hypertensive	Both Bo	th.									-3.300 -4.147 to -2.454)	-3.816 (-4.547 to -3.085)	-4.331 (-4.959 to -3.704)	-4.847 (-5.389 to -4.305)	-5.363 (-5.848 to -4.877)	-5.878 (-6.346 to -5.411)	-6.394 (-6.886 to -5.901)	-6.909 (-7.464 to -6.354)	-7.425 (-8.069 to -6.781)						
	Black, Non-Hypertensive	Both Bo	th								6	-3.910 -5.065 to -2.755)	-4.426 (-5.564 to -3.287)	-4.941 (-6.081 to -3.802)	-5.457 (-6.616 to -4.298)	-5.972 (-7.168 to -4.777)	-6.488 (-7.735 to -5.241)	-7.004 (-8.316 to -5.691)	-7.519 (-8.909 to -6.129)	-8.035 (-9.512 to -6.557)						
	Black, Hypertensive	Both Bo	th.									-5.844 -7.222 to -4.467)	-6.360 (-7.663 to -5.057)	-6.876 (-8.117 to -5.635)	-7.391 (-8.584 to -6.198)	-7.907 (-9.068 to -6.745)	-8.422 (-9.569 to -7.275)	-8.938 (-10.088 to -7.788)	-9.453 (-10.624 to -8.282)	-9.969 (-11.178 to -8.760)						

	e used by age and sex for each of		onalex (equil	tor ambient an p	onution aconor, a	and shipsing									A	ges											
Risk - Outcome	Category / Units	Morbidity / Mortali	ity Sex	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Stornach cancer	1 giday	Both	Both										1.199 (0.987 to 1.443)	1.205 (1.008 to 1.428)	1.205 (0.989 to 1.459)	1.202 (0.996 to 1.443)	1.209 (0.996 to 1.448)	1.198 (0.984 to 1.429)	1.204 (1.008 to 1.428)	1.2 (0.982 to 1.457)	1.206 (1.003 to 1.431)	1.21 (0.995 to 1.444)	1.203 (0.987 to 1.433)	1.205 (0.984 to 1.459)	1.205 (0.984 to 1.459)	1.205 (0.984 to 1.459)	1.205 (0.984 to 1.459)
Intimate partner violence (H	IV PAF approach)																										
HIV/AIDS - Drug-susceptible Tuberculosis	Exposed	morbidity	Both	1.59 (1.3 to 1.94)																							
HIV/AIDS - Drug-susceptible Tuberculosis	Exposed	mortality	Both	1.59 (1.3 to 1.94)																							
HIV/AIDS - Drug-susceptible Tuberculosis	Not exposed	morbidity	Both	1.0 (1.0 to 1.0)																							
HIV/AIDS - Drug-susceptible Tabaeculouis	Not exposed	mortality	Both	1.0																							
HIV/AIDS - Multidrug-resistant Tuberculosis without extensive	t drug Exposed	morbidity	Both	1.59																							
HIV/AIDS - Maltidrug-resistant Tuberculosis without extensive	t drug Exposed	mortality	Both	1.59																							
HIV/AIDS - Multidrug-resistant Tuberculosis without extensive	t drug Not exposed	morbidity	Both	1.0																							
HIV/AIDS - Multidrug-resistant Tuberculosis without extensive	t drue Not exposed	mortality	Both	(1.0 to 1.0)																							
HIV/AIDS - Extensively drug-	Ermond	murhistic	Both	(1.0 to 1.0) 1.59																							
resistant Tuberculosis HIV/AIDS - Extensively drug-	Ermond	martality	Brath	(1.3 to 1.94) 1.59																							
resistant Tuberculosis HIV/AIDS - Extensively drug-	No			(1.3 to 1.94) 1.0																							
resistant Tuberculosis HIV/AIDS - Extensively drug-	Not exposed	moreany	Boin	(1.0 to 1.0) 1.0																							
resistant Tuberculosis HIV/AIDS resulting in other	Not exposed	mortany	Boin	(1.0 to 1.0)																							
diseases HIV/AIDS resulting in other	Exposed	morbidity	Both	(1.3 to 1.94)																							
diseases HIV/ADS resulting in other	Exposed	mortality	Both	(1.3 to 1.94)																							
diseases	Not exposed	morbidity	Both	(1.0 to 1.0)																							
HIV/AIDS resulting in other diseases	Not exposed	mortality	Both	1.0 (1.0 to 1.0)			1	-		-					-			-				-					
Intimate partner violence (ex	xposure approach)						1.000	1077	14.7	10.7	1017	1077	1/17	1000	1011	1-	1017	1.5	1.1.1	1.000	1077	1.000	167	1000	14.7	14-1	1071
Maternal abortive outcome	Exposed	Both	Both		(1.154 to 3.098)	1.96 (1.145 to 3.167)	(1.169 to 3.094)	(1.151 to 3.204)	(1.137 to 3.211)	(1.144 to 3.144)	(1.168 to 3.019)	(1.124 to 3.269)	(1.161 to 3.187)	(1.154 to 3.119)	(1.128 to 3.04)	2.0 (1.175 to 3.159)	(1.175 to 3.202)	(1.156 to 3.107)	(1.132 to 3.217)	(1.164 to 3.09)	(1.146 to 3.147)	(1.154 to 3.189)	(1.188 to 3.11)	(1.176 to 3.148)	(1.192 to 3.065)	(1.149 to 3.087)	(1.136 to 3.152)
Maternal abortive outcome	Not exposed	Both	Both		1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)				
Major depressive disorder	Exposed	Morbidity	Both		1.474 (1.099 to 1.949)	1.456 (1.051 to 1.898)	1.464 (1.099 to 1.914)	1.454 (1.103 to 1.922)	1.457 (1.088 to 1.917)	1.46 (1.106 to 1.92)	1.454 (1.094 to 1.872)	1.455 (1.098 to 1.92)	1.459 (1.091 to 1.917)	1.447 (1.079 to 1.885)	1.458 (1.099 to 1.921)	1.442 (1.068 to 1.924)	1.449 (1.061 to 1.942)	1.448 (1.104 to 1.875)	1.458 (1.111 to 1.905)	1.461 (1.083 to 1.92)	1.458 (1.094 to 1.928)	1.456 (1.09 to 1.912)	1.462 (1.097 to 1.914)	1.47 (1.116 to 1.892)	1.447 (1.061 to 1.898)	1.448 (1.064 to 1.879)	1.45 (1.081 to 1.915)
Major depressive disorder	Not exposed	Morbidity	Both		1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)				
Childhood sexual abuse																											
Major depressive disorder	Exposed	Morbidity	Both		1.631 (1.405 to 1.877)	1.631 (1.422 to 1.881)	1.633 (1.409 to 1.893)	1.63 (1.404 to 1.896)	1.627 (1.396 to 1.863)	1.63 (1.407 to 1.881)	1.635 (1.401 to 1.894)	1.637 (1.41 to 1.879)	1.638 (1.42 to 1.888)	1.633 (1.416 to 1.876)	1.633 (1.416 to 1.874)	1.634 (1.417 to 1.882)	1.632 (1.411 to 1.91)	1.631 (1.416 to 1.889)	1.634 (1.402 to 1.883)	1.634 (1.408 to 1.866)	1.634 (1.422 to 1.886)	1.634 (1.412 to 1.878)	1.633 (1.406 to 1.886)	1.634 (1.419 to 1.899)	1.637 (1.412 to 1.856)	1.636 (1.401 to 1.877)	1.635 (1.402 to 1.903)
Major depressive disorder	Not exposed	Morbidity	Both		1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)				
Alcohol use disorders	Exposed	Both	Both		1.55 (1.186 to 1.952)	1.549 (1.194 to 2.024)	1.551 (1.208 to 1.983)	1.55 (1.184 to 1.941)	1.551 (1.172 to 2.013)	1.55 (1.202 to 1.965)	1.553 (1.182 to 2.008)	1.545 (1.194 to 1.939)	1.557 (1.202 to 1.994)	1.54 (1.167 to 2.047)	1.541 (1.184 to 1.953)	1.554 (1.187 to 1.97)	1.552 (1.196 to 1.997)	1.549 (1.188 to 2.0)	1.55 (1.183 to 1.971)	1.562 (1.206 to 1.992)	1.546 (1.206 to 1.973)	1.556 (1.19 to 1.971)	1.562 (1.179 to 1.982)	1.545 (1.187 to 1.961)	1.549 (1.2 to 1.967)	1.545 (1.211 to 1.967)	1.549 (1.19 to 1.944)
Alcohol use disorders	Not exposed	Both	Both		1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)				
Bullying victimization																											
Major depressive disorder	First exposed to bullying in current year	morbidity	Both	2.016 (1.488 to 2.731)																							
Major depressive disorder	First exposed to bullying 1 year prior	morbidity	Both	1.955 (1.481 to 2.579)																							
Major depressive disorder	First exposed to bullying 2 years prior	morbidity	Both	1.895 (1.471 to 2.441)																							
Major depressive disorder	First exposed to bullying 3 years prior	morbidity	Both	1.838 (1.457 to 2.318)																							
Major depressive disorder	First exposed to bullying 4 years prior	morbidity	Both	1.782 (1.437 to 2.21)																							
Major depressive disorder	First exposed to bullying 5 years prior	morbidity	Both	1.728 (1.41 to 2.117)																							
Major depressive disorder	First exposed to bullying 6 years prior	morbidity	Both	1.675 (1.376 to 2.04)																							
Major depressive disorder	First exposed to ballying 7 years prior	morbidity	Both	1.624 (1.333 to 1.979)																							
Major depressive disorder	First exposed to bullying 8 years prior	morbidity	Both	1.575 (1.284 to 1.932)																							
Major depressive disorder	First exposed to bullying 9 years prior	morbidity	Both	1.527 (1.229 to 1.898)																							
Major depressive disorder	First exposed to bullying 10 years prior	morbidity	Both	1.481 (1.171 to 1.873)																							
Major depressive disorder	First exposed to bullying 11 years prior	morbidity	Both	1.436 (1.112 to 1.855)																							
Major depressive disorder	First exposed to bullying 12 years prior	morbidity	Both	1.392 (1.052 to 1.843)																							
Major depressive disorder	First exposed to ballying 13 years prior	morbidity	Both	1.35 (0.993 to 1.836)																							
Major depressive disorder	First exposed to ballying 14 years prior	morbidity	Both	1.309 (0.936 to 1.832)																							
Major depressive disorder	First exposed to bullying 15 years prior	morbidity	Both	1.269 (0.881 to 1.83)																							
Major depressive disorder	First exposed to bullying 16 years prior	morbidity	Both	1.231																							
Major depressive disorder	First exposed to bullying 17 years prior	morbidity	Both	1.193																							
Major depressive disorder	First exposed to bullying 18 years prior	morbidity	Both	(0.770 40 1.831) 1.157 40.73 to 1.921																							
Major depressive disorder	First exposed to bullying 19 years prior	morbidity	Both	1.122																							
Major dereessive disorder	First exposed to bullvine 20 years wine	morbidity	Both	(1.088 at 1.0837)																							
Major dereessive downlar	First exceed to bullying 21 wars wine	morbidi~	Both	(0.643 to 1.842) 1.055																							
Major demonstration disconter	First exceed to bellvine 22 war min-	peubidity	Reath	(0.603 to 1.847) 1.023																							
Maine de contra de contra	Never exposed to bullying or first expose	ad	noni n	(0.565 to 1.852) 1.0																							
major depressive disorder	over 22 years prior	morbukiy	Both	(1.0 to 1.0) 2.016																							
Anxiety disorders	 init exposed to bullying in current year 	morbukiy	Both	(1.488 to 2.731) 1.955																							
Anxiety disorders	First exposed to bullying 1 year prior	morbidity	Both	(1.481 to 2.579)																							
Anxiety disorders	First exposed to bullying 2 years prior	morbidity	Both	(1.471 to 2.441)																							
Anxiety disorders	First exposed to bullying 3 years prior	morbidity	Both	(1.457 to 2.318)	l I																						

															А	lges											
Risk - Outcome	Category / Units	Morbidity / Mortality	Sex All-1	ge 0-	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Anxiety disorders	First exposed to bullying 4 years prior	morbidity	Both 1.78 (1.437 tr	t 2.21)																							
Anxiety disorders	First exposed to bullying 5 years prior	morbidity	Both (1.41 to)	i .117)																							
Anxiety disorders	First exposed to bullying 6 years prior	morbidity	Both (1.376 to	i 2.04)																							
Anxiety disorders	First exposed to bullying 7 years prior	morbidity	Both (1.333 to	i 1.979)																							
Anxiety disorders	First exposed to bullying 8 years prior	morbūdity	Both (1.284 to	i 1.932)																							
Anxiety disorders	First exposed to bullying 9 years prior	morbidity	Both (1.229 to	r 1.898)																							
Anxiety disorders	First exposed to bullying 10 years prior	morbidity	Both (1.171 to	i 1.873)																							
Anxiety disorders	First exposed to bullying 11 years prior	morbidity	Both (1.112 to	i 1.855)																							
Anxiety disorders	First exposed to bullying 12 years prior	morbidity	Both (1.052 to	t 1.843)																							
Anxiety disorders	First exposed to ballying 13 years prior	morbidity	Both (0.993 to	1.836)																							
Anxiety disorders	First exposed to bullying 14 years prior	morbūdity	Both (0.936 to) 1.832)																							
Anxiety disorders	First exposed to bullying 15 years prior	morbidity	Both (0.881 tr) 1.83)																							
Anxiety disorders	First exposed to bullying 16 years prior	morbūdity	Both (0.828 to	1 1.83)																							
Anxiety disorders	First exposed to bullying 17 years prior	morbidity	Both (0.778 to	i 1.831)																							
Anxiety disorders	First exposed to ballying 18 years prior	morbidity	Both (0.73 to	(.834)																							
Anxiety disorders	First exposed to bullying 19 years prior	morbūdity	Both (0.685 to	t 1.837)																							
Anxiety disorders	First exposed to bullying 20 years prior	morbūdity	Both (0.643 to	i 1.842)																							
Anxiety disorders	First exposed to ballying 21 years prior	morbūlity	Both (0.603 to	i 1.847)																							
Anxiety disorders	First exposed to ballying 22 years prior	morbidity	Both (0.565 to	l.852)																							
Anxiety disorders	Never exposed to bullying or first exposed over 22 years prior	morbūlity	Both (1.0 to	1.0)																							
Low physical activity																											
Colon and rectum cancer	0 METs	Both	Both (1.0 to	1.0)																							
Colon and rectum cancer	10200 METs	Both	Both 0.81 (0.758 to	l 1.864)																							
Colon and rectum cancer	10800 METs	Both	Both 0.80 (0.757 to)).862)																							
Colon and rectum cancer	11400 METs	Both	Both (0.756 to	1 0.86)																							
Colon and rectum cancer	1200 METx	Both	Both 0.95 (0.88 to	i .032)																							
Colon and rectum cancer	12000 METs	Both	Both 0.80 (0.754 to	i 1.858)																							
Colon and rectum cancer	12600 METs	Both	Both (0.753 to	i 1.856)																							
Colon and rectum cancer	13200 METs	Both	Both 0.80 (0.752 to	t 0.855)																							
Colon and rectum cancer	13800 METs	Both	Both (0.75 to)	.854)																							
Colon and rectum cancer	14400 METs	Both	Both 0.75	1 (851)																							
Colon and rectum cancer	15000 METs	Both	Both 0.75	5 0.85)																							
Colon and rectum cancer	15600 METa	Both	Both (0.744 to	i 1.849)																							
Colon and rectum cancer	16200 METs	Both	Both (0.75	849)																							
Colon and rectum cancer	16800 METa	Both	Both (0.739 to	1																							
Colon and rectum cancer	17400 METs	Both	Both (0.78	0.847)																							
Colon and rectum cancer	1800 METs	Both	Both (0.821 to	1.048)																							
Colon and rectum cancer	18000 METs	Both	Both (0.733 to	1																							
Colon and rectum cancer	18600 METs	Both	Both 0.78	5																							
Colon and rectum cancer	19200 METs	Both	Both 0.78	1																							
Colon and rectum cancer	19800 METs	Both	Both 0.72	1.845)																							
Colon and rectum cancer	20400 METs	Both	Both 0.7	845)																							
Colon and rectum cancer	21000 METs	Both	Both (0.717 to	1.845)																							
Colon and rectum cancer	21600 METs	Both	Both 0.77	1846)																							
Colon and rectum cancer	22200 METs	Both	Both 0.77	i 1.847)																							
Colon and rectum cancer	22800 METs	Both	Both 0.77	1																							
Colon and rectum cancer	23400 METs	Both	(0.708 is Both 0.77	1.0.40)																							
Colon and rectum cancer	2400 METx	Both	0.702 to 0.88 Both 0.88	(1958)																							
Colon and rectum cancer	24000 METs	Both	0.76 Both 0.76																								
Colon and rectum cancer	24600 METs	Both	(0.698 to Both 0.7t																								
Colon and rectum cancer	25200 METs	Both	(0.694 in Both 0.76	6 0000																							
Colon and rectum cancer	25800 METs	Both	(0.689 to Both 0.76	u.a5) k																							
Colon and rectum cancer	26400 METs	Both	(0.685 to Both 0.76	1.8-49/) E																							
Colon and rectum cancer	27000 METs	Both	(0.679 to Both 0.7	.631)																							
Colon and rectum cancer	27600 METs	Both	(0.675 to Both 0.75	1855) k																							
Colon and rectum cancer	28200 METs	Both	(0.671 to 0.75	1855) 1																							
Colon and rectum cancer	28800 METa	Both	(0.668 to 0.75	3.856) i																							
COURSE AND PRODUCT OFFICE		10.000	(0.664 to	1.856)																							

oppendix rable oa. Kelative i si	is used by age and sex for	each official for all fisk facto		tor amorent are p	onution arconol.	and anosting.									A	iges											
Risk - Outcome	Category / Ur	its Morbidity / Mortali	ty Sex	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Colon and rectum cancer	29400 METs	Both	Both	0.753 (0.659 to 0.856)																							
Colon and rectum cancer	3000 METs	Both	Both	0.833 (0.76 to 0.907)																							
Colon and rectum cancer	30000 METs	Both	Both	0.751 (0.654 to 0.856)																							
Colon and rectum cancer	30600 METs	Both	Both	0.749																							
Colon and rectum cancer	31200 METs	Both	Both	0.748																							
Colon and rectum cancer	31800 METs	Both	Both	0.746																							
Colon and rectum cancer	32400 METs	Both	Both	(0.64 to 0.859) 0.744																							
Colon and rectam curver	33010 META	Breth	Brath	(0.635 to 0.86) 0.743																							
Colon and rectam canver	3600 MFTv	Breth	Beath	(0.632 to 0.86) 0.831																							
Color relation areas	4200 MET-	Prob	Rest	(0.761 to 0.903) 0.829																							
Colored and television	1000 1077	Dola Dola	D.d.	(0.761 to 0.898) 0.827																							
Color and rectain career	\$400 MET-	Post	Post	(0.761 to 0.893) 0.825																							
			Liona -	(0.763 to 0.89) 0.978																							
Colon and rectum cancer	600 METa	Both	Both	(0.94 to 1.016) 0.824																							
Colon and rectum cancer	6000 METs	Both	Both	(0.763 to 0.887) 0.822																							
Colon and rectum cancer	6600 METx	Both	Both	(0.762 to 0.884)																							
Colon and rectum cancer	7200 METs	Both	Both	(0.762 to 0.88)																							
Colon and rectum cancer	7800 METs	Both	Both	(0.761 to 0.877)																							
Colon and rectum cancer	8400 METs	Both	Both	(0.76 to 0.872)																							
Colon and rectum cancer	9000 METs	Both	Both	(0.758 to 0.871)																							
Colon and rectum cancer	9600 METs	Both	Both	0.813 (0.758 to 0.868)																							
Breast cancer	0 METs	Both	Both	1.0 (1.0 to 1.0)																							
Breast cancer	10200 METs	Both	Both	0.908 (0.861 to 0.961)																							
Breast cancer	10800 METs	Both	Both	0.904 (0.859 to 0.952)																							
Breast cancer	11400 METs	Both	Both	0.9 (0.855 to 0.948)																							
Breast cancer	1200 METs	Both	Both	0.974 (0.942 to 1.005)																							
Breast cancer	12000 METs	Both	Both	0.895 (0.85 to 0.946)																							
Breast cancer	12600 METs	Both	Both	0.891 (0.842 to 0.946)																							
Breast cancer	13200 METs	Both	Both	0.887 (0.832 to 0.948)																							
Breast cancer	13800 METs	Both	Both	0.883 (0.819 to 0.949)																							
Breast cancer	14400 METs	Both	Both	0.879 (0.807 to 0.952)																							
Breast cancer	15010 METs	Both	Both	0.874 (0.796 to 0.957)																							
Breast cancer	15600 METs	Both	Both	0.872 (0.797 to 0.951)																							
Breast cancer	16200 METs	Both	Both	0.87 (0.799 to 0.946)																							
Breast cancer	16800 METs	Both	Both	0.868 (0.801 to 0.941)																							
Breast cancer	17400 METs	Both	Both	0.865 (0.804 to 0.933)																							
Breast cancer	1800 METs	Both	Both	0.96 (0.914 to 1.008)																							
Breast cancer	18000 METs	Both	Both	0.863 (0.807 to 0.927)																							
Breast cancer	18600 METs	Both	Both	0.861 (0.807 to 0.919)																							
Breast cancer	19200 METs	Both	Both	0.859 (0.808 to 0.912)																							
Breast cancer	19800 METs	Both	Both	0.856 (0.809 to 0.909)																							
Breast cancer	20400 METs	Both	Both	0.854 (0.809 to 0.906)																							
Breast cancer	21010 METs	Both	Both	0.852 (0.809 to 0.9)																							
Breast cancer	21600 METs	Both	Both	0.85 (0.808 to 0.896)																							
Breast cancer	22200 METs	Both	Both	0.847 (0.805 to 0.893)																							
Breast cancer	22800 METs	Both	Both	0.845 (0.804 to 0.891)																							
Breast cancer	23400 METs	Both	Both	0.843 (0.801 to 0.888)																							
Breast cancer	2400 METs	Both	Both	0.957 (0.914 to 1.001)																							
Breast cancer	24000 METs	Both	Both	0.841 (0.797 to 0.888)																							
Breast cancer	24600 METa	Both	Both	0.839 (0.793 to 0.889)																							
Breast cancer	25200 METs	Both	Both	0.836 (0.789 to 0.891)																							
Breast cancer	25800 METs	Both	Both	0.834																							
Breast cancer	26400 METs	Both	Both	0.832 (0.779 to 0.897)																							
Breast cancer	27000 METs	Both	Both	0.83																							
Breast cancer	27600 METs	Both	Both	0.827																							
Breast cancer	28200 METs	Both	Both	0.825																							
Breast cancer	28800 METa	Both	Both	0.823																							
Breast cancer	29400 METs	Both	Both	(0.759 to 0.3) 0.821 (0.752 to 0.0021																							
				(1																						I

Appendix Table 6a. Relative ris	iks used by age and sex for each o	autcome for all risk factors exce	pt for ambient air	pollution alcohol.	and snoosing.									Ages												
Risk - Outcome	Category / Units	Morbidity / Mortality Se:	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years 15	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years 40-4	44 years 4	5-49 years 5	0-54 years	5-59 years	0-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Breast cancer	3000 METs	Both Bot	h 0.953 (0.913 to 0.994)																							
Breast cancer	30000 METa	Both Bot	h 0.818 (0.745 to 0.904)																							
Breast cancer	30600 METs	Both Bot	b 0.816 (0.738 to 0.906)																							
Breast cancer	31200 METs	Both Bot	h 0.814 (0.731 to 0.907)																							
Breast cancer	31800 METs	Both Bot	h 0.812 (0.724 to 0.909)																							
Breast cancer	32400 METs	Both Bot	h 0.809 (0.717 to 0.911)																							
Breast cancer	33010 METs	Both Bot	h 0.808 (0.713 to 0.912)																							
Breast cancer	3600 METs	Both Bot	0.949 (0.912 to 0.99)																							
Breast cancer	4200 METs	Both Bot	h 0.946 (0.909 to 0.985)																							
Breast cancer	4800 METs	Both Bot	h 0.942 (0.907 to 0.982)																							
Breast cancer	5400 METs	Both Bot	h 0.938 (0.903 to 0.979)																							
Breast cancer	600 METs	Both Bot	h 0.987 (0.971 to 1.003)																							
Breast cancer	6000 METs	Both Bot	h 0.935 (0.897 to 0.979)																							
Breast cancer	6600 METs	Both Bot	h 0.931 (0.889 to 0.978)																							
Breast cancer	7200 METs	Both Bot	h 0.928 (0.883 to 0.978)																							
Breast cancer	7800 METs	Both Bot	h 0.924 (0.875 to 0.979)																							
Breast cancer	8400 METs	Both Bot	h 0.92 (0.867 to 0.979)																							
Breast cancer	9000 METs	Both Bot	0.917 (0.858 to 0.979)																							
Breast cancer	9600 METs	Both Bot	h 0.912 (0.862 to 0.97)																							
Ischaemic heart disease	0 METs	Both Bot	h (1.0 to 1.0)																							
Ischaemic heart disease	10200 METs	Both Bot	0.73 (0.668 to 0.793)																							
Ischaemic heart disease	10800 METa	Both Bot	h 0.73 (0.669 to 0.791)																							
Ischaemic heart disease	11400 METs	Both Bot	h 0.729 (0.671 to 0.789)																							
Ischaemic heart disease	1200 METs	Both Bot	h 0.819 (0.714 to 0.928)																							
Ischaemic heart disease	12000 METs	Both Bot	h 0.728 (0.672 to 0.787)																							
Ischaemic heart disease	12600 METs	Both Bot	h 0.728 (0.672 to 0.786)																							
Ischaemic heart disease	13200 METs	Both Bot	h 0.727 (0.671 to 0.785)																							
Ischaemic heart disease	13800 METs	Both Bot	h 0.727 (0.67 to 0.784)																							
Ischaemic heart disease	14400 METs	Both Bot	h 0.726 (0.671 to 0.782)																							
Ischaemic heart disease	15010 METs	Both Bot	h 0.725 (0.67 to 0.781)																							
Ischaemic heart disease	15600 METs	Both Bot	h 0.725 (0.671 to 0.781)																							
Ischaemic heart disease	16200 METs	Both Bot	h 0.724 (0.671 to 0.779)																							
Ischaemic heart disease	16800 METs	Both Bot	h 0.724 (0.669 to 0.779)																							
Ischaemic heart disease	17400 METs	Both Bot	h (0.669 to 0.78)																							
Ischaemic heart disease	1800 METs	Both Bot	h 0.808 (0.739 to 0.886)																							
Ischaemic heart disease	18000 METs	Both Bot	h (0.668 to 0.78)																							
Ischaemic heart disease	18600 METs	Both Bot	h (0.667 to 0.781)																							
Ischaemic heart disease	19200 METs	Both Bot	h (0.666 to 0.781)																							
Ischaemic heart disease	19800 METs	Both Bot	h (0.665 to 0.781)																							
Ischaemic heart disease	20400 METs	Both Bot	h (0.664 to 0.781)																							
Ischaemic heart disease	21000 METs	Both Bot	h (0.663 to 0.782) 0.719																							
Ischaemic heart disease	21000 METs	Both Bot	(0.662 to 0.782) 0.718																							
Ischaeme heart disease	22200 METs	Both Bot	(0.66 to 0.783) 0.718																							
Inclusive a second second	22400 ME IS	nom Bot	(0.659 to 0.784) 0.717																							
Inclusive locat disease	23400 MET-	Both Bot	(0.657 to 0.786) 0.796																							
Jucharmir-heart disease	24010 METs	Both Bot	(0.711 to 0.895) 0.717																							
Ischaerreic heart disease	24600 METs	Bosh Rev	(0.655 to 0.787) b 0.716																							
Ischaemic heart disease	25200 METs	Both Bot	(U.653 to 0.788) b 0.715																							
Ischaemic heart disease	25800 METs	Both Bot	(0.65 to 0.789) b 0.715																							
Ischaemic heart disease	26400 METs	Both Bot	(0.648 to 0.79) h 0.714																							
Ischaerric heart disease	27000 METs	Both Bot	0.714 (0.643 to 0.704)																							
Ischaerric heart disease	27600 METs	Both Bot	b 0.713 (0.64 to 0.796)																							
Ischaemic heart disease	28200 METs	Both Bot	h 0.712 (0.636 to 0.798)																							
Ischaemic heart disease	28800 METs	Both Bot	h 0.712 (0.633 to 0.8)																							
Ischaemic heart disease	29400 METs	Both Bot	h 0.711 (0.63 to 0.801)																							
Ischaemic heart disease	3000 METs	Both Bot	h 0.776 (0.713 to 0.844)																							
			•																							

ppendix range of Relative ris	as used by age and sex i	O PACA DIRCOME IO AIL HSK DEIDTS EXC		politition accords.	and smosting.									A	ges											
Risk - Outcome	Category / 1	Units Morbidity/Mortality S	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Ischaemic heart disease	30000 META	Both B	oth 0.711 (0.626 to 0.801)		1	1			11								II					1				
Ischaemic heart disease	30600 METs	Both B	oth 0.71																							
Ischaemic heart disease	31200 METs	Both B	0.709																							
Ischaemic heart disease	31800 METs	Both B	(0.82 55 0.804) 0.709																							
Indonesia baset disease	22400 MET.	P-sh P	(0.616 to 0.806) 0.708																							
Inclusion in the second	32400 ME IX	Boin B	018 (0.615 to 0.808) . 0.708																							
Ischaemic heart disease	33000 METs	Both B	(0.613 to 0.81)																							
Ischaemic heart disease	3600 METx	Both B	(0.692 to 0.816)																							
Ischaemic heart disease	4200 METs	Both B	0.756 (0.652 to 0.821)																							
Ischaemic heart disease	4800 METx	Both B	0.736 (0.653 to 0.818)																							
Ischaemic heart disease	5400 METs	Both B	0.735 (0.654 to 0.815)																							
Ischaemic heart disease	600 METs	Both B	0.909 (0.857 to 0.964)																							
Ischaemic heart disease	6000 METs	Both B	0.734 (0.656 to 0.811)																							
Ischaemic heart disease	6600 METs	Both B	0.734 (0.657 to 0.808)																							
Ischaemic heart disease	7200 METs	Both B	0.733 (0.659 to 0.806)																							
Ischaemic heart disease	7800 METs	Both B	0.733																							
Ischaemic heart disease	8400 METs	Both B	(0.06 10 0.803) 0.732																							
Industria beart disease	9000 MFTv	Roth B	(0.662 to 0.801) 0.731																							
			(0.664 to 0.799) . 0.731																							
Ischiachtic neart disease	9600 ME18	DOIN D	(0.666 to 0.796)																							
Ischaemic stroke	0 METs	Both B	oth (1.0 to 1.0)																							
Ischaemic stroke	10200 METs	Both B	oth (0.653 to 0.864)																							
Ischaemic stroke	10800 METs	Both B	0.75 (0.653 to 0.858)																							
Ischaemic stroke	11400 METs	Both B	0.747 (0.653 to 0.851)																							
Ischaemic stroke	1200 METs	Both B	0.819 (0.662 to 0.999)																							
Ischaemic stroke	12000 METs	Both B	0.744 (0.651 to 0.845)																							
Ischaemic stroke	12600 METs	Both B	0.741 (0.65 to 0.84)																							
Ischaemic stroke	13200 METs	Both B	0.738 (0.65 to 0.833)																							
Ischaemic stroke	13800 METs	Both B	0.735																							
Ischaemic stroke	14400 METs	Both B	0.732																							
Ischaemic stroke	15000 METs	Both B	0.729																							
Industrie stroke	15600 METs	Roth B	(0.644 to 0.82) 0.727																							
hadronnia stanka	16200 MET-	Post P	(0.642 to 0.813) 0.724																							
I have been	10000 1077	Doin D	(0.639 to 0.809) 0.721																							
Defizence stroke	16800 342 18	DOIN D	018 (0.639 to 0.806) 0.718																							
Ischaemic stroke	17400 METs	Both B	oth (0.636 to 0.802)																							
Ischaemic stroke	1800 METx	Both B	(0.702 to 0.915)																							
Ischaemic stroke	18000 METs	Both B	oth (0.633 to 0.797)																							
Ischaemic stroke	18600 METs	Both B	oth (0.63 to 0.793)																							
Ischaemic stroke	19200 METs	Both B	0.709 (0.626 to 0.791)																							
Ischaemic stroke	19800 METs	Both B	0.706 (0.623 to 0.789)																							
Ischaemic stroke	20400 METs	Both B	0.703 (0.62 to 0.79)																							
Ischaemic stroke	21000 METs	Both B	0.7 (0.615 to 0.787)																							
Ischaemic stroke	21600 METs	Both B	0.697 (0.611 to 0.787)																							
Ischaemic stroke	22200 METs	Both B	0.694 (0.606 to 0.787)																							
Ischaemic stroke	22800 METs	Both B	0.691 (0.602 to 0.785)																							
Ischaemic stroke	23400 METs	Both B	0.688																							
Ischaemic stroke	2400 METs	Both B	(0.595 is 0.785) oth 0.785																							
Inclusive stroke	24000 METs	Roth B	(0.643 to 0.943) 0.685																							
hadronnia stanka	24600 MET.	Post P	(0.593 to 0.78) 0.682																							
Parameter Anone	2000 10-0		(0.586 to 0.779) 0.679																							
Ischaemic stroke	25200 METs	Both B	018 (0.579 to 0.778)																							
Ischaemic stroke	25800 METs	Both B	(0.572 to 0.776)																							
Ischaemic stroke	26400 METs	Both B	oth (0.566 to 0.777)																							
Ischaemic stroke	27000 METs	Both B	(0.56 to 0.779)																							
Ischaemic stroke	27600 METs	Both B	0.668 (0.554 to 0.781)																							
Ischaemic stroke	28200 METs	Both B	0.665 (0.547 to 0.781)																							
Ischaemic stroke	28800 METs	Both B	0.662 (0.541 to 0.783)																							
Ischaemic stroke	29400 METs	Both B	0.659 (0.534 to 0.785)																							
Ischaemic stroke	3000 METs	Both B	0.784 (0.701 to 0.888)																							
Ischaemic stroke	30000 METs	Both B	0.656 (0.527 to 0.787)																							

Appendix Table 6a. Relative ris	iks used by age and sex for each o	sutcome for all risk factors exo	ept for ambient air	pollution alcohol.	and smoxing.									Ages											
Risk - Outcome	Category / Units	Morbidity / Mortality S	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years 15	5-19 years 2	0-24 years 1	25-29 years	30-34 years	35-39 years 40-44	4 years 45-4	9 years 50-54	rears 55-59 yea	rs 60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Ischaemic stroke	30600 METs	Both Be	oth 0.653 (0.521 to 0.789)																						
Ischaemic stroke	31200 METs	Both Bo	0.65 (0.514 to 0.791)																						
Ischaemic stroke	31800 METa	Both Bo	0.647 (0.506 to 0.792)																						
Ischaemic stroke	32400 METa	Both Bo	0.644 (0.498 to 0.792)																						
Ischaemic stroke	33010 METs	Both Be	0.643 (0.493 to 0.792)																						
Ischaemic stroke	3600 METs	Both Bo	oth 0.783 (0.695 to 0.883)																						
Ischaemic stroke	4200 METs	Both Be	oth 0.782 (0.644 to 0.934)																						
Ischaemic stroke	4800 METs	Both Bo	oth 0.779 (0.647 to 0.926)																						
Ischaemic stroke	5400 METs	Both Bo	0.776 (0.648 to 0.918)																						
Ischaemic stroke	600 METs	Both Bo	0.91 (0.831 to 0.999)																						
Ischaemic stroke	6000 METs	Both Be	oth 0.774 (0.649 to 0.913)																						
Ischaemic stroke	6600 METs	Both Bo	oth 0.771 (0.65 to 0.905)																						
Ischaemic stroke	7200 METs	Both Bo	0.768 (0.652 to 0.898)																						
Ischaemic stroke	7800 METs	Both Be	0.765 (0.65 to 0.891)																						
Ischaemic stroke	8400 METs	Both Bo	0.762 (0.651 to 0.884)																						
Ischaemic stroke	9000 METs	Both Bo	0.759 (0.652 to 0.878)																						
Ischaemic stroke	9600 METs	Both Bo	oth 0.756 (0.652 to 0.87)																						
Diabetes mellitus type 2	0 METs	Both Bo	oth 1.0 (1.0 to 1.0)																						
Diabetes mellitus type 2	10200 METs	Both Be	oth 0.727 (0.684 to 0.774)																						
Diabetes mellitus type 2	10800 METs	Both Be	oth 0.726 (0.683 to 0.772)																						
Diabetes mellitus type 2	11400 METs	Both Bo	oth 0.725 (0.682 to 0.77)																						
Diabetes mellitus type 2	1200 METs	Both Bo	oth 0.961 (0.933 to 0.992)																						
Diabetes mellitus type 2	12000 METs	Both Bo	oth 0.723 (0.682 to 0.771)																						
Diabetes mellitus type 2	12600 METs	Both Be	oth 0.722 (0.682 to 0.768)																						
Diabetes mellitus type 2	13200 METs	Both Bo	oth 0.721 (0.681 to 0.766)																						
Diabetes mellitus type 2	13800 METs	Both Bo	oth 0.72 (0.68 to 0.765)																						
Diabetes mellitus type 2	14400 METs	Both Bo	0.719 (0.678 to 0.764)																						
Diabetes mellitus type 2	15010 METs	Both Bo	0.718 (0.678 to 0.762)																						
Diabetes mellitus type 2	15600 METa	Both Bo	oth 0.717 (0.677 to 0.761)																						
Diabetes mellitus type 2	16200 METs	Both Bo	oth (0.676 to 0.76)																						
Diabetes mellitus type 2	16800 METs	Both Be	oth 0.715 (0.674 to 0.759)																						
Diabetes mellitus type 2	17400 METs	Both Bo	oth 0.714 (0.673 to 0.759)																						
Diabetes mellitus type 2	1800 METs	Both Bo	oth 0.921 (0.867 to 0.985)																						
Diabetes mellitus type 2	18000 METs	Both Be	oth (0.67 to 0.759)																						
Diabetes mellitus type 2	18600 METs	Both Bo	oth (0.668 to 0.759)																						
Diabetes mellitus type 2	19200 METs	Both Bo	oth (0.665 to 0.758) 0.709																						
Dubetes mellitus type 2	19800 METs	Both Bo	(0.663 to 0.758) 0.708																						
Dubetes mellitus type 2	20400 METs	Both Bo	(0.66 to 0.759)																						
Durenes metanas type 2	21000 ME IN	boin b	(0.658 to 0.759) . 0.706																						
Dianenes metinus type 2	21000 META	Boll D	(0.656 to 0.759) 0.705																						
Diabetes mellitus type 2	22800 METa	Brok D.	(0.654 to 0.759) 0.704																						
Diabetes mellitus type 2	23400 METs	Both Re	(0.651 to 0.759) 0.703																						
Diabetes mellitus type 2	2400 METs	Both Be	(0.65 to 0.759) 0.882																						
Diabetes mellitus type 2	24000 METs	Both Be	(0.8 to 0.977) 0.702																						
Diabetes mellitus type 2	24600 METs	Both Be	(0.848 IS 0.76) oth 0.701																						
Diabetes mellitus type 2	25200 METs	Both B	(0.646 to 0.76) 0.7																						
Diabetes mellitus type 2	25800 METs	Both B	(0.043 to 0.76) 0.699 ah (0.64 to 0.77)																						
Diabetes mellitus type 2	26400 METs	Both B	(0.64 to 0.76)																						
Diabetes mellitus type 2	27000 METs	Both Bo	0.697 (0.635 to 0.76)																						
Diabetes mellitus type 2	27600 METs	Both Bo	oth 0.696 (0.631 to 0.761)																						
Diabetes mellitus type 2	28200 METs	Both Bo	oth 0.694 (0.628 to 0.762)																						
Diabetes mellitus type 2	28800 METs	Both Bo	oth 0.693 (0.624 to 0.762)																						
Diabetes mellitus type 2	29400 METs	Both Bo	oth 0.692 (0.621 to 0.763)																						
Diabetes mellitus type 2	3000 METs	Both Bo	0.834 (0.773 to 0.904)																						
Diabetes mellitus type 2	30000 METs	Both Bo	0.691 (0.618 to 0.764)																						
Diabetes mellitus type 2	30600 METs	Both Bo	0.69 (0.615 to 0.765)																						
																									,

Appendix Table 6a. Relative FISR	used by age and see	a tor reach on come	TOT SH HISK HEROTS IS	100000000000000000000000000000000000000	r ponution arconol, r	ind smosing.									Aş	ges											
Risk - Outcome	Category	/ Units Mo	orbidity / Mortality	Sex All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Diabetes mellitus type 2	31200 METs		Both	Both 0.689 (0.612 to 0.766																							
Diabetes mellitus type 2	31800 METs		Both	Both 0.688 (0.609 to 0.768																							
Diabetes mellitus type 2	32400 METs		Both	Both 0.687 (0.607 to 0.768																							
Diabetes mellitus type 2	33000 METs		Both	Both 0.686 (0.605 to 0.769																							
Diabetes mellitus type 2	3600 METx		Both	Both 0.786 (0.735 to 0.841																							
Diabetes mellitus type 2	4200 METs		Both	Both 0.737																							
Diabetes mellitus type 2	4800 METs		Both	Both 0.736 (0.685 to 0.791																							
Diabetes mellitus type 2	5400 METs		Both	Both 0.735 (0.684 to 0.789																							
Diabetes mellitus type 2	600 METs		Both	Both 0.98																							
Diabetes mellitus type 2	6000 METs		Both	Both 0.734																							
Diabetes mellitus type 2	6600 METx		Both	Both 0.733																							
Diabetes mellitus type 2	7200 METs		Both	Both 0.732																							
Diabetes mellitus type 2	7800 METs		Both	Both 0.731																							
Diabetes mellitus type 2	8400 METx		Both	Both 0.73																							
Diabetes mellitus type 2	9000 METx		Both	0.729 Both 0.729	,																						
Diabetes mellitus type 2	9600 METx		Both	0.554 iS 0.775 Both 0.728	,																						
High fasting plasma glucose (continuous)			(0.684 to 0.776)																						
lschaemic heart disease	mmol/L.		Both	Both									1.471	1.373	1.274	1.22	1.211	1.201	1.192	1.182	1.173	1.168	1.168	1.169	1.169	1.169	1.169
Ischaemic stroke	mmol/L.		Both	Both									(1.147 to 2.099) 1.526	(1.133 to 1.741) 1.4	(1.15 to 1.451) 1.275	(1.085 to 1.365) 1.21	(1.111 to 1.327) 1.204	(1.121 to 1.297) 1.199	(1.123 to 1.269)	(1.111 to 1.267) 1.188	(1.085 to 1.271) 1.183	(1.08 to 1.273)	(1.097 to 1.264) 1.162	1.133	(1.071 to 1.316)	(1.0/1 to 1.316)	(1.0/1 to 1.316)
Intracerebral hemorthane	mmol/L.		Both	Both									(1.11 to 2.227) 1.506	(1.101 to 1.85) 1.382	(1.081 to 1.561) 1.258	(1.044 to 1.441) 1.196	(1.075 to 1.383) 1.193	(1.095 to 1.334) 1.191	(1.108 to 1.302) 1.189	(1.101 to 1.295) 1.187	(1.069 to 1.308) 1.184	(1.056 to 1.309) 1.175	(1.075 to 1.295) 1.158	(1.055 to 1.332) 1.116	(1.055 to 1.332) 1.116	(1.055 to 1.332)	(1.055 to 1.332) 1.116
Subarachnoid hemorrhave	mmol/L.		Both	Both									(1.112 to 2.221) 1.506	(1.111 to 1.843) 1.382	(1.085 to 1.487) 1.258	(1.053 to 1.371) 1.196	(1.082 to 1.333)	(1.106 to 1.298) 1.191	(1.115 to 1.262) 1.189	(1.115 to 1.252) 1.187	(1.097 to 1.264) 1.184	(1.087 to 1.263) 1.175	(1.09 to 1.253) 1.158	(1.056 to 1.25) 1.116	(1.056 to 1.25) 1.116	(1.056 to 1.25) 1.116	(1.056 to 1.25)
Chronic kidney disease due to	mmol/L.		Both	Both									(1.112 to 2.221) 1.388	(1.111 to 1.843) 1.388	(1.085 to 1.487) 1.388	(1.053 to 1.371) 1.388	(1.082 to 1.333) 1.388	(1.106 to 1.298) 1.388	(1.115 to 1.262) 1.388	(1.115 to 1.252) 1.388	(1.097 to 1.264) 1.388	(1.087 to 1.263) 1.388	(1.09 to 1.233) 1.388	(1.056 to 1.25) 1.388	(1.056 to 1.25) 1.388	(1.056 to 1.25) 1.388	(1.056 to 1.25) 1.388
hypertension Chronic kidney disease due to	mmol/I		Brath	Broth									(1.272 to 1.512) 1.388														
glomerulonephritis Chronic kidney disease due to otl	ar meal/l		Brath	Beth									(1.272 to 1.512) 1.388														
and unspecified causes High fasting plasma glucose (categorical)							1					(1.272 to 1.512)														
Latent tuberculosis infection	Diabetic		Both	Both									2.73	2.801	2.871	2.798	2.581	2.364	2.147	1.93	1.713	1.598	1.587	1.559	1.559	1.559	1.559
Latent tuberculosis infection	Not diabetic		Both	Both									(1.973 to 3.602) 1.0	(2.056 to 3.658) 1.0	(2.047 to 3.698) 1.0	(1.97 to 3.63) 1.0	(1.91 to 3.268) 1.0	(1.817 to 2.943) 1.0	(1.692 to 2.673) 1.0	(1.485 to 2.44) 1.0	(1.231 to 2.317) 1.0	(1.125 to 2.239) 1.0	(1.184 to 2.115) 1.0	(1.182 to 2.174) 1.0	(1.182 to 2.174) 1.0	(1.182 to 2.174) 1.0	(1.182 to 2.174) 1.0
Drur-susceptible tubercalosis	Diabetic		Both	Both									(1.0 to 1.0) 2.73	(1.0 to 1.0) 2.801	(1.0 to 1.0) 2.871	(1.0 to 1.0) 2.798	(1.0 to 1.0) 2.581	(1.0 to 1.0) 2.364	(1.0 to 1.0) 2.147	(1.0 to 1.0) 1.93	(1.0 to 1.0) 1.713	(1.0 to 1.0) 1.598	(1.0 to 1.0) 1.587	(1.0 to 1.0) 1.559	(1.0 to 1.0) 1.559	(1.0 to 1.0) 1.559	(1.0 to 1.0) 1.559
Drur-susceptible tuberculosis	Not diabetic		Both	Both									(1.973 to 3.802)	(2.056 to 3.658) 1.0	(2.047 to 3.698) 1.0	(1.97 to 3.63) 1.0	(1.91 to 3.268)	(1.817 to 2.943)	(1.692 to 2.673)	(1.485 to 2.44) 1.0	(1.231 to 2.317) 1.0	(1.125 to 2.239)	(1.184 to 2.115) 1.0	(1.182 to 2.174) 1.0	(1.182 to 2.174)	(1.182 to 2.174)	(1.182 to 2.174) 1.0
Multideug-resistant tuberculosis	Disbais		Ruth	Reals									(1.0 to 1.0) 2.73	(1.0 to 1.0) 2.801	(1.0 to 1.0) 2.871	(1.0 to 1.0) 2.798	(1.0 to 1.0) 2.581	(1.0 to 1.0) 2.364	(1.0 to 1.0) 2.147	(1.0 to 1.0) 1.93	(1.0 to 1.0) 1.713	(1.0 to 1.0) 1.598	(1.0 to 1.0) 1.587	(1.0 to 1.0) 1.559	(1.0 to 1.0) 1.559	(1.0 to 1.0) 1.559	(1.0 to 1.0) 1.559
without extensive drug resistance Multidrug-resistant tuberculosis	Not Asherin		Ruth	Reals									(1.973 to 3.602) 1.0	(2.056 to 3.658) 1.0	(2.047 to 3.698) 1.0	(1.97 to 3.63) 1.0	(1.91 to 3.268) 1.0	(1.817 to 2.943) 1.0	(1.692 to 2.673) 1.0	(1.485 to 2.44) 1.0	(1.231 to 2.317) 1.0	(1.125 to 2.239) 1.0	(1.184 to 2.115) 1.0	(1.182 to 2.174) 1.0	(1.182 to 2.174) 1.0	(1.182 to 2.174) 1.0	(1.182 to 2.174) 1.0
without extensive drug resistance Extensively drug-resistant	Disbais		Ruth	Reals									(1.0 to 1.0) 2.73	(1.0 to 1.0) 2.801	(1.0 to 1.0) 2.871	(1.0 to 1.0) 2.798	(1.0 to 1.0) 2.581	(1.0 to 1.0) 2.364	(1.0 to 1.0) 2.147	(1.0 to 1.0) 1.93	(1.0 to 1.0) 1.713	(1.0 to 1.0) 1.598	(1.0 to 1.0) 1.587	(1.0 to 1.0) 1.559	(1.0 to 1.0) 1.559	(1.0 to 1.0) 1.559	(1.0 to 1.0) 1.559
taberculosis Extensively drug-resistant	Net Arbeite		Ruth	Reals									(1.973 to 3.602) 1.0	(2.056 to 3.658) 1.0	(2.047 to 3.698) 1.0	(1.97 to 3.63) 1.0	(1.91 to 3.268) 1.0	(1.817 to 2.943) 1.0	(1.692 to 2.673) 1.0	(1.485 to 2.44) 1.0	(1.231 to 2.317) 1.0	(1.125 to 2.239) 1.0	(1.184 to 2.115) 1.0	(1.182 to 2.174) 1.0	(1.182 to 2.174) 1.0	(1.182 to 2.174) 1.0	(1.182 to 2.174) 1.0
tuberculosis	Disbais		Ruth	Mala									(1.0 to 1.0) 1.527														
Color and restore server	Disbatis		Purk I										(1.081 to 2.304) 1.527														
Color and restore server	Not Asherin		Ruth	Mala									(1.086 to 2.315) 1.0														
Color and rectum cancer	Not diabene		Doin 1	ALLES									(1.0 to 1.0) 1.0														
													(1.0 to 1.0) 1.523														
Liver cancer due to wash	Durene		Doin 1	ALLES									(1.093 to 2.3) 1.512														
Liver cancer due to wash	Durenc		nom r	emaex									(1.083 to 2.293) 1.0														
Laver cancer due to NASH	root daabetisc		Both	MINES									(1.0 to 1.0) 1.0														
Liver cancer due to NASH	over maneric		DOIN P	And and a second s									(1.0 to 1.0) 1.523														
Laver cancer due to other causes	D-40000C		DOIN .	-manne									(1.093 to 2.3) 1.512														
Laver cancer due to other causes	Loubetic		Both F	emaars									(1.083 to 2.293) 1.0														
Laver cancer due to other causes	Not diabetic		Both	MING									(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)									
Liver cancer due to other causes	Not diabetic		Both F	emales									(1.0 to 1.0) 1.517	(1.0 to 1.0)	(1.0 to 1.0) 1.517	(1.0 to 1.0)	(1.0 to 1.0) 1.517	(1.0 to 1.0) 1.517									
Pancreatic cancer	Diabetic		Both	Males									(1.085 to 2.314)														
Panereatic cancer	Dubetic		Both F	emales									(1.075 to 2.311)														
Pancreatic cancer	Not diabetic		Both	Males									(1.0 to 1.0)	(0.1 ot 0.1)	(1.0 to 1.0)												
Pancreatic cancer	Not diabetic		Both F	emakes									(1.0 to 1.0)														
Tracheal, bronchas, and lung can	cer Diabetic		Both	Males									1.515 (1.076 to 2.309)	1.515 (1.076 to 2.309)	(1.076 to 2.309)	(1.076 to 2.309)	1.515 (1.076 to 2.309)	1.515 (1.076 to 2.309)	(1.076 to 2.309)	(1.076 to 2.309)	1.515 (1.076 to 2.309)	(1.076 to 2.309)	(1.076 to 2.309)	(1.076 to 2.309)	1.515 (1.076 to 2.309)	1.515 (1.076 to 2.309)	1.515 (1.076 to 2.309)
Tracheal, bronchus, and lung can	cer Diabetic		Both F	lemakes									1.512 (1.087 to 2.299)	1.512 (1.087 to 2.299)	(1.087 to 2.299)	1.512 (1.087 to 2.299)	1.512 (1.087 to 2.299)	1.512 (1.087 to 2.299)	1.512 (1.087 to 2.299)	1.512 (1.087 to 2.299)	1.512 (1.087 to 2.299)	1.512 (1.087 to 2.299)	1.512 (1.087 to 2.299)	1.512 (1.087 to 2.299)	1.512 (1.087 to 2.299)	1.512 (1.087 to 2.299)	1.512 (1.087 to 2.299)
Tracheal, bronchus, and lung can	cer Not diabetic		Both	Males									1.0 (1.0 to 1.0)														
Tracheal, bronchus, and lung can	cer Not diabetic		Both F	emales									1.0 (1.0 to 1.0)														
Breast cancer	Diabetic		Both	Both									1.513 (1.087 to 2.206)														
Breast cancer	Not diabetic		Both	Both									1.0 (1.0 to 1.0)														
Ovarian cancer	Diabetic		Both	Both									1.522 (1.092 to 2.32)														
Ovarian cancer	Not diabetic		Both	Both									1.0 (1.0 to 1.0)														

Appendix rable oa. Relative risks t	used by age	and sex for each outcome for all risk factors			on acconor, and	a smoosing.									A	iges											
Pick - Outcome	0	tanory / Unite Marbidity / Martality	All-a	ge O	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Bladder cancer	Diabetic	Both	Males										1.514 (1.076 to 2.256)	1.514 (1.076 to 2.256)	1.514 (1.076 to 2.256)	1.514 (1.076 tp 2.256)	1.514 (1.076 to 2.256)	1.514	1.514	1.514 (1.076 to 2.256)	1.514 (1.076 m 2.256)	1.514	1.514 (1.076 to 2.256)				
Bladder cancer	Diabetic	Both	Females										1.511	1.511	1.511	1.511	1.511	1.511	1.511	1.511	1.511	1.511	1.511	1.511	1.511	1.511	1.511
Bludder cancer	Not diabetic	Both	Males										1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
Blackler concorr	Not dishetic	Brith	Females										1.0	1.0	(1.0 10 1.0)	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	(1.018 1.0)	1.0	1.0	(1.0 10 1.0)
													(1.0 to 1.0) 8.264	(1.0 to 1.0) 6.651	(1.0 to 1.0) 5.039	(1.0 to 1.0) 4.138	(1.0 to 1.0) 3.947	(1.0 to 1.0) 3.756	(1.0 to 1.0) 3.565	(1.0 to 1.0) 3.374	(1.0 to 1.0) 3.183	(1.0 to 1.0) 2.992	(1.0 to 1.0) 2.801	(1.0 to 1.0) 2.324	(1.0 to 1.0) 2.324	(1.0 to 1.0) 2.324	(1.0 to 1.0) 2.324
Peripheral vascular disease	Dubetse	Both	Both										(6.044 to 9.303)	(5.375 to 7.452)	(4.436 to 5.673)	(3.561 to 4.76)	(3.447 to 4.504)	(3.347 to 4.232)	(3.214 to 3.972)	(3.079 to 3.714)	(2.92 to 3.466)	(2.76 to 3.25)	(2.558 to 3.05)	(1.977 to 2.671)	(1.977 to 2.671)	(1.977 to 2.671)	(1.977 to 2.671)
Peripheral vascular disease	Not diabetic	Both	Both										(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)									
Alzheimer's disease and other dementias	Diabetic	Both	Males										1.516 (1.084 to 2.295)	1.516 (1.084 to 2.295)	1.516 (1.084 to 2.295)	1.516 (1.084 to 2.295)	1.516 (1.084 to 2.295)	1.516 (1.084 to 2.295)									
Alzheimer's disease and other dementias	Diabetic	Both	Females										1.52 (1.08 to 2.301)	1.52 (1.08 to 2.301)	1.52 (1.08 to 2.301)	1.52 (1.08 to 2.301)	1.52 (1.08 to 2.301)	1.52 (1.08 to 2.301)									
Alzheimer's disease and other dementias	Not diabetic	Both	Males										1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)									
Alzheimer's disease and other dementias	Not diabetic	Both	Females										1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)									
Glaucoma	Diabetic	Both	Males										1.52	1.52	1.52	1.52	1.52	1.52	1.52	1.52	1.52	1.52	1.52	1.52	1.52	1.52	1.52
Glaucoma	Diabetic	Both	Females										1.516	1.516	1.516	1.516	1.516	1.516	1.516	1.516	1.516	1.516	1.516	1.516	1.516	1.516	1.516
	N												(1.08 to 2.328) 1.0	(1.08 to 2.328) 1.0	(1.08 to 2.328) 1.0	(1.08 to 2.328) 1.0	(1.08 to 2.328) 1.0	(1.08 to 2.328) 1.0									
Gincom	NOT diabetic	BOIR	MINO										(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)									
Glaucoma	Not diabetic	Both	Females										(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)									
Cataract	Diabetic	Both	Males										(1.09 to 2.259)	(1.09 to 2.259)	1.52 (1.09 to 2.259)	(1.09 to 2.259)	(1.09 to 2.259)	(1.09 to 2.259)									
Cataract	Diabetic	Both	Females										1.522 (1.094 to 2.289)	1.522 (1.094 to 2.289)	1.522 (1.094 to 2.289)	1.522 (1.094 to 2.289)	1.522 (1.094 to 2.289)	1.522 (1.094 to 2.289)									
Cataract	Not diabetic	Both	Males										1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)									
Cataract	Not diabetic	Both	Females										1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)									
High LDL cholesterol																											
Ischaernic heart disease	mmol/L	Both	Both					1		11			2.016	2.027	2.038	1.971	1.828	1.685	1.541	1.398	1.254	1.193	1.213	1.262	1.262	1262	1.262
Inclusionic stroke	mmal/I	Breh	Beath										(1.684 to 2.544) 1.67	(1.768 to 2.354) 1.626	(1.831 to 2.273) 1.583	(1.775 to 2.191) 1.518	(1.676 to 2.004) 1.434	(1.561 to 1.815)	(1.446 to 1.648) 1.265	(1.306 to 1.494) 1.181	(1.141 to 1.372) 1.096	(1.088 to 1.312) 1.062	(1.124 to 1.321) 1.077	(1.11 to 1.465)	(1.11 to 1.465)	(1.11 to 1.465) 1.116	(1.11 to 1.465) 1.116
High systelic blood pressure													(1.334 to 2.339)	(1.352 to 2.041)	(1.342 to 1.849)	(1.287 to 1.76)	(1.242 to 1.636)	(1.212 to 1.514)	(1.164 to 1.391)	(1.109 to 1.299)	(1.043 to 1.223)	(1.008 to 1.193)	(1.012 to 1.216)	(1.014 to 1.344)	(1.014 to 1.344)	(1.014 to 1.344)	(1.014 to 1.344)
righ systone blood pressure													1631	1.474	1317	1.229	1211	1.193	1.175	1157	1.139	1.127	112	1.104	1.104	1.104	1.104
Rheurratic heart disease	10 mmHg	Both	Both										(1.174 to 2.306)	(1.17 to 1.898)	(1.144 to 1.575)	(1.089 to 1.422)	(1.101 to 1.367)	(1.107 to 1.328)	(1.101 to 1.287)	(1.086 to 1.265)	(1.055 to 1.248)	(1.048 to 1.241)	(1.06 to 1.238)	(1.04 to 1.28)	(1.04 to 1.28)	(1.04 to 1.28)	(1.04 to 1.28)
Ischaemic heart disease	10 mmHg	Both	Both										(1.44 to 2.596)	(1.458 to 2.207)	(1.461 to 1.911)	(1.398 to 1.799)	(1.393 to 1.705)	(1.385 to 1.619)	(1.368 to 1.535)	(1.332 to 1.488)	(1.257 to 1.456)	(1.224 to 1.424)	(1.225 to 1.404)	(1.134 to 1.437)	(1.134 to 1.437)	(1.134 to 1.437)	(1.134 to 1.437)
Ischaemic stroke	10 mmHg	Both	Both										1.854 (1.395 to 2.588)	1.774 (1.427 to 2.252)	1.694 (1.404 to 2.035)	1.628 (1.354 to 1.95)	1.574 (1.36 to 1.823)	1.521 (1.361 to 1.698)	1.468 (1.344 to 1.596)	1.414 (1.302 to 1.524)	1.361 (1.214 to 1.49)	1.318 (1.168 to 1.451)	1.284 (1.179 to 1.389)	1.201 (1.109 to 1.37)	1.201 (1.109 to 1.37)	1.201 (1.109 to 1.37)	1.201 (1.109 to 1.37)
Intracerebral hemorrhage	10 mmHg	Both	Both										2.134 (1.555 to 2.919)	2.05 (1.593 to 2.648)	1.966 (1.589 to 2.465)	1.874 (1.492 to 2.302)	1.775 (1.484 to 2.114)	1.676 (1.446 to 1.932)	1.577 (1.402 to 1.754)	1.478 (1.331 to 1.619)	1.379 (1.207 to 1.54)	1.323 (1.162 to 1.495)	1.311 (1.193 to 1.45)	1.279 (1.126 to 1.515)	1.279 (1.126 to 1.515)	1.279 (1.126 to 1.515)	1.279 (1.126 to 1.515)
Subarachnoid hemorrhage	10 mmHg	Both	Both										2.134 (1.555 to 2.919)	2.05 (1.593 to 2.648)	1.966 (1.589 to 2.465)	1.874 (1.492 to 2.302)	1.775 (1.484 to 2.114)	1.676 (1.446 to 1.932)	1.577 (1.402 to 1.754)	1.478 (1.331 to 1.619)	1.379 (1.207 to 1.54)	1.323 (1.162 to 1.495)	1.311 (1.193 to 1.45)	1.279 (1.126 to 1.515)	1.279 (1.126 to 1.515)	1.279 (1.126 to 1.515)	1.279 (1.126 to 1.515)
Hypertensive heart disease	10 mmHg	Both	Both										2.862 (1.879 to 4.108)	2.838 (1.857 to 4.187)	2.814 (1.802 to 4.337)	2.703 (1.762 tp.4.186)	2.504 (1.804 to 3.758)	2.304 (1.722 to 3.464)	2.105 (1.645 to 3.336)	1.905 (1.447 to 3.171)	1.706 (1.188 to 3.215)	1.619 (1.053 to 3.136)	1.644 (1.078 to 3.091)	1.708 (1.103 to 3.258)	1.708 (1.103 m 3.258)	1.708 (1.103 to 3.258)	1.708 (1.103 to 3.258)
Non-rheumatic calcific acetic valve	e 10 mmHg	Both	Both										1.755	1.605	1.455	1.365	1.335	1.306	1.276	1.247	1.217	1.193	1.175	1.128	1.128	1.128	1.128
anene	10 mmHz	Park	Resh										(1.20510.2.423)	1.605	(1.278101.042)	1.365	1.335	1.306	(1.212101.342) 1.276	(1.183 to 1.303) 1.247	1.217	(1.116101.263)	1.175	1.128	(1.071 10 1.235) 1.128	1.128	1.128
			P. 4										(1.266 to 2.423) 1.76	(1.293 to 2.011) 1.631	(1.278 to 1.642) 1.503	(1.232 to 1.51) 1.423	(1.222 to 1.449) 1.392	(1.219 to 1.394) 1.361	(1.212 to 1.342) 1.33	(1.183 to 1.303) 1.299	(1.131 to 1.284) 1.268	(1.116 to 1.263) 1.237	(1.12 to 1.237) 1.208	(1.071 to 1.235) 1.134	(1.071 to 1.235) 1.134	(1.071 to 1.235) 1.134	(1.071 to 1.235) 1.134
Artis needaton and numer	10 mmrg	BOIR	DOIN										(1.336 to 2.43)	(1.379 to 2.026)	(1.396 to 1.644)	(1.34 to 1.505) 1.345	(1.328 to 1.457)	(1.313 to 1.411) 1.296	(1.293 to 1.369) 1.272	(1.265 to 1.333) 1.248	(1.233 to 1.308)	(1.202 to 1.277)	(1.177 to 1.238)	(1.092 to 1.185)	(1.092 to 1.185)	(1.092 to 1.185)	(1.092 to 1.185)
Aortic aneurysm	10 mmHg	Both	Both										(1.259 to 2.164)	(1.29 to 1.816)	(1.3 to 1.535)	(1.227 to 1.451)	(1.233 to 1.405)	(1.229 to 1.362)	(1.218 to 1.327)	(1.191 to 1.299)	(1.16 to 1.286)	(1.137 to 1.261)	(1.126 to 1.229)	(1.071 to 1.184)	(1.071 to 1.184)	(1.071 to 1.184)	(1.071 to 1.184)
Peripheral vascular disease	10 mmHg	Both	Both										1.728 (1.203 to 2.428)	(1.206 to 1.87)	(1.182 to 1.329)	(1.019 to 1.263)	(1.047 to 1.243)	1.146 (1.071 to 1.224)	1.15 (1.094 to 1.208)	(1.11 to 1.199)	(1.113 to 1.207)	(1.152 (1.104 to 1.201)	(1.098 to 1.176)	(1.054 to 1.154)	(1.054 to 1.154)	(1.095 (1.054 to 1.154)	1.095 (1.054 to 1.154)
Endocarditis	10 mmHg	Both	Both										1.755 (1.266 to 2.423)	1.605 (1.293 to 2.011)	1.455 (1.278 to 1.642)	1.365 (1.232 to 1.51)	1.335 (1.222 to 1.449)	1.306 (1.219 to 1.394)	1.276 (1.212 to 1.342)	1.247 (1.183 to 1.303)	1.217 (1.131 to 1.284)	1.193 (1.116 to 1.263)	1.175 (1.12 to 1.237)	1.128 (1.071 to 1.235)	1.128 (1.071 to 1.235)	1.128 (1.071 to 1.235)	1.128 (1.071 to 1.235)
Other cardiovascular and circulator diseases	^{ry} 10 mmHg	Both	Both										1.744 (1.339 to 2.396)	1.624 (1.384 to 2.006)	1.504 (1.405 to 1.626)	1.427 (1.354 to 1.498)	1.395 (1.336 to 1.452)	1.363 (1.318 to 1.406)	1.33 (1.296 to 1.365)	1.298 (1.266 to 1.332)	1.265 (1.231 to 1.303)	1.235 (1.201 to 1.27)	1.207 (1.177 to 1.235)	1.137 (1.095 to 1.187)	1.137 (1.095 to 1.187)	1.137 (1.095 to 1.187)	1.137 (1.095 to 1.187)
Chronic kidney disease due to diabetes mellitus type 1	10 mmHg	Both	Both										1.283 (1.186 to 1.397)	1.283 (1.186 to 1.397)	1.283 (1.186 to 1.397)	1.283 (1.186 to 1.397)	1.283 (1.186 to 1.397)	1.283 (1.186 to 1.397)									
Chronic kidney disease due to diabetes mellitus tone ?	10 mmHg	Both	Both										1.283 (1.186 to 1.397)	1.283 (1.186 to 1.397)	1.283 (1.186 to 1.397)	1.283 (1.186 m 1.397)	1.283 (1.186 to 1.397)	1.283 (1.186 to 1.397)									
Chronic kidney disease due to	10 mmHg	Both	Both										1.281	1.281	1.281	1.281	1.281	1.281	1.281	1.281	1.281	1.281	1.281	1.281	1.281	1.281	1.281
hypertension Chronic kidney disease due to	10 mmHz	Brith	Beath										(1.1810 1.383)	1.281	1.281	(1.18 to 1.585) 1.281	(1.1810 1.383)	(1.18 10 1.585)	1.281	1.281	(1.18101.383)	1.281	1.281	(1.18 10 1.383)	(1.18 10 1.383)	1.281	1.281
glomerulonephritis Chronic kidney disease due to other	- 10 - 11		Reals										(1.182 to 1.383) 1.282	(1.182 to 1.383) 1.282	(1.182 to 1.383) 1.282	(1.182 to 1.383) 1.282	(1.182 to 1.383) 1.282	(1.182 to 1.383) 1.282									
and unspecified causes	10 mmrg	BOIR	Boin		1								(1.181 to 1.395)	(1.181 to 1.395)	(1.181 to 1.395)	(1.181 to 1.395)	(1.181 to 1.395)	(1.181 to 1.395)									
righ body-mass muex in addres												1.201	1 201	1.20.1	1.201	1.201	1.201	1.20.1	1 201	1.201	1 301	1 201	1 201	1.201	1.201	1 201	1.201
Oesophageal cancer	5 kg/m2	Both	Males									(1.077 to 1.754)	(1.077 to 1.754)	(1.077 to 1.754)	(1.077 to 1.754)	(1.077 to 1.754)	(1.077 to 1.754)										
Oesophageal cancer	5 kg/m2	Both	Females									1.351 (1.012 to 1.745)	1.351 (1.012 to 1.745)	1.351 (1.012 to 1.745)	1.351 (1.012 to 1.745)	1.351 (1.012 to 1.745)	1.351 (1.012 to 1.745)										
Colon and rectum cancer	5 kg/m2	Both	Males									1.177 (1.145 to 1.208)	1.177 (1.145 to 1.208)	1.177 (1.145 to 1.208)	1.177 (1.145 to 1.208)	1.177 (1.145 to 1.208)	1.177 (1.145 to 1.208)										
Colon and rectum cancer	5 kg/m2	Both	Females									1.059 (1.031 to 1.083)	1.059 (1.031 to 1.083)	1.059 (1.031 to 1.083)	1.059 (1.031 to 1.083)	1.059 (1.031 to 1.083)	1.059 (1.031 to 1.083)										
Liver cancer due to hepatitis B	5 kg/m2	Both	Males									1.289 (1.109 to 1.491)	1.289 (1.109 to 1.491)	1.289 (1.109 to 1.491)	1.289 (1.109 to 1.491)	1.289 (1.109 to 1.497)	1.289 (1.109 to 1.491)	1.289 (1.109 to 1.491)	1.289 (1.109 to 1.491)	1.289 (1.109 to 1.491)	1.289 (1.109 to 1.491)	1.289 (1.109 to 1.491)					
Liver cancer due to hepatitis B	5 kg/m2	Both	Females									1.176	1.176	1.176	1.176	1.176	1.176	1.176	1.176	1.176	1.176	1.176	1.176	1.176 (1.03 m 1.224)	1.176	1.176	1.176
Liver cancer due to heratitis C	5 kg/m2	Both	Males									(1.00 10 1.334)	1.289	1.289	1.289	1.289	1.289	1.289	1.289	1.289	1.289	1289	1289	1.289	1.289	1289	1.289
Lines cancer due to honorali ?	5 ku()	Durik	Females									(1.109 to 1.491) 1.176	(1.109 to 1.491) 1.176	(1.109 to 1.491) 1.176	(1.109 to 1.491) 1.176	(1.109 to 1.491) 1.176	(1.109 to 1.491) 1.176										
Lavor Cancer due to neparatis C	- ng 412	DOIL										(1.03 to 1.334)	(1.03 to 1.334) 1,280	(1.03 to 1.334) 1.289	(1.03 to 1.334) 1.289	(1.03 to 1.334) 1.289	(1.03 to 1.334)	(1.03 to 1.334) 1.780	(1.03 to 1.334) 1.789	(1.03 to 1.334) 1.289	(1.03 to 1.334) 1,289	(1.03 to 1.334) 1,289	(1.03 to 1.334) 1,289	(1.03 to 1.334) 1.289	(1.03 to 1.334) 1.289	(1.03 to 1.334) 1,289	(1.03 to 1.334) 1.289
Liver cancer due to alcohol use	5 kg/m2	Both	Males									(1.109 to 1.491)	(1.109 to 1.491)	(1.109 to 1.491)	(1.109 to 1.491)	(1.109 to 1.491)	(1.109 to 1.491)										
Liver cancer due to alcohol use	5 kg/m2	Both	Females									(1.03 to 1.334)	1.176 (1.03 to 1.334)	1.176 (1.03 to 1.334)	(1.03 to 1.334)	(1.03 to 1.334)	1.176 (1.03 to 1.334)	1.176 (1.03 to 1.334)	(1.03 to 1.334)	1.176 (1.03 to 1.334)	1.176 (1.03 to 1.334)	1.176 (1.03 to 1.334)	(1.03 to 1.334)	1.176 (1.03 to 1.334)	(1.03 to 1.334)	1.176 (1.03 to 1.334)	(1.03 to 1.334)
Liver cancer due to other causes	5 kg/m2	Both	Males									1.289 (1.109 to 1.491)	1.289 (1.109 to 1.491)	1.289 (1.109 to 1.491)	1.289 (1.109 to 1.491)	1.289 (1.109 to 1.491)	1.289 (1.109 to 1.491)										
Liver cancer due to other causes	5 kg/m2	Both	Females									1.176 (1.03 to 1.334)	1.176 (1.03 to 1.334)	1.176 (1.03 to 1.334)	1.176 (1.03 to 1.334)	1.176 (1.03 to 1.334)	1.176 (1.03 to 1.334)										
Gallbladder and biliary tract cancer	5 kg/m2	Both	Males									1.155 (1.033 to 1.281)	1.155 (1.033 to 1.281)	1.155 (1.033 to 1.281)	1.155 (1.033 to 1.281)	1.155 (1.033 to 1.281)	1.155 (1.033 to 1.281)										
Gallbladder and biliary tract cancer	5 kg/m2	Both	Females									1.344 (1.223 to 1.477)	1.344 (1.223 to 1.477)	1.344 (1.223 to 1.477)	1.344 (1.223 to 1.477)	1.344 (1.223 to 1.477)	1.344 (1.223 to 1.477)										
Pancreatic cancer	5 kg/m2	Both	Males									1.071	1.071	1.071	1.071	1.071	1.071	1.071	1.071	1.071	1.071	1.071	1.071	1.071	1.071	1.071	1.071
Parents	5 ku - 1	Burk	Females									(0.999 to 1.153) 1.092	(0.999 to 1.153) 1.092	(0.999 to 1.153) 1.092	(0.999 to 1.153) 1.092	(0.999 to 1.153) 1.092	(0.999 to 1.153) 1.092										
												(1.037 to 1.144)	(1.037 to 1.144)	(1.037 to 1.144)	(1.037 to 1.144)	(1.037 to 1.144)	(1.037 to 1.144)										

Appendix Table 6a. Relative risks	used by age and sex for each outcon	ne for all risk factors exce		201111104 11100401	and Shittaning									Aş	ges											
Risk - Outcome	Category / Units	Morbidity / Mortality Sea	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Breast cancer (premenopausal)	5 kg/m2	Both Bot	-								0.89 (0.869 to 0.914)															
Breast cancer (postmenopausal)	5 kg/m2	Both Bot	h														1.089 (1.037 to 1.14)									
Uterine cancer	5 kg/m2	Both Bot	h								1.613 (1.543 to 1.681)															
Ovarian cancer	5 kg/m2	Both Bot	h								1.038 (0.999 to 1.077)															
Kidney cancer	5 kg/m2	Both Male	×								1.24 (1.171 to 1.313)															
Kidney cancer	5 kg/m2	Both Fema	les								1.32 (1.254 to 1.394)															
Thyroid cancer	5 kg/m2	Both Male	×								1.221 (1.068 to 1.381)															
Thyroid cancer	5 kg/m2	Both Fema	les								1.136 (1.094 to 1.178)															
Non-Hodgkin's lymphoma	5 kg/m2	Both Male	x								1.089 (1.038 to 1.143)															
Non-Hodgkin's lymphoma	5 kg/m2	Both Fema	les								1.068 (1.01 to 1.125)															
Maltiple myeloma	5 kg/m2	Both Male	s								1.089 (1.027 to 1.153)															
Multiple myeloma	5 kg/m2	Both Fema	les								1.092 (1.034 to 1.157)															
Acute lymphoid leukaemia	5 kg/m2	Both Male	x								1.086 (1.053 to 1.119)															
Acute lymphoid leakaemia	5 kg/m2	Both Fema	les								1.131 (1.061 to 1.208)															
Chronic lymphoid leukaemia	5 kg/m2	Both Male	×								1.086 (1.053 to 1.119)															
Chronic lymphoid leukaemia	5 kg/m2	Both Fema	les								1.131 (1.061 to 1.208)															
Acute myeloid leukaemia	5 kg/m2	Both Male	s								1.086 (1.053 to 1.119)															
Acute myeloid leukaemia	5 kg/m2	Both Fema	les								1.131 (1.061 to 1.208)															
Chronic myeloid leukaemia	5 kg/m2	Both Male	s								1.086 (1.053 to 1.119)															
Chronic myeloid leukaemia	5 kg/m2	Both Fema	les								1.131 (1.061 to 1.208)															
Other leukaemia	5 kg/m2	Both Male	×								1.086 (1.053 to 1.119)															
Other leukaemia	5 kg/m2	Both Fema	les								1.131 (1.061 to 1.208)															
Ischaemic heart disease	5 kg/m2	Both Bot	h								2.274 (1.259 to 3.683)	2.274 (1.259 to 3.683)	2.018 (1.3 to 3.099)	1.724 (1.533 to 1.93)	1.599 (1.418 to 1.784)	1.567 (1.458 to 1.68)	1.52 (1.417 to 1.631)	1.466 (1.372 to 1.557)	1.414 (1.325 to 1.504)	1.364 (1.287 to 1.448)	1.319 (1.242 to 1.4)	1.274 (1.187 to 1.365)	1.17 (1.091 to 1.252)	1.17 (1.091 to 1.252)	1.17 (1.091 to 1.252)	1.17 (1.091 to 1.252)
Ischaemic stroke	5 kg/m2	Both Bot	h								2.472 (1.4 to 3.975)	2.472 (1.4 to 3.975)	2.235 (1.457 to 3.329)	1.979 (1.699 to 2.313)	1.826 (1.6 to 2.075)	1.733 (1.581 to 1.898)	1.635 (1.479 to 1.795)	1.543 (1.441 to 1.653)	1.455 (1.345 to 1.566)	1.38 (1.31 to 1.458)	1.304 (1.234 to 1.376)	1.228 (1.16 to 1.304)	1.068 (0.992 to 1.143)	1.068 (0.992 to 1.143)	1.068 (0.992 to 1.143)	1.068 (0.992 to 1.143)
Intracerebral hemorrhage	5 kg/m2	Both Bot	h								3.066 (1.751 to 5.334)	3.066 (1.751 to 5.334)	2.913 (1.862 to 4.395)	2.597 (1.974 to 3.375)	2.389 (1.87 to 2.998)	2.199 (1.822 to 2.672)	1.996 (1.626 to 2.416)	1.805 (1.574 to 2.06)	1.665 (1.438 to 1.932)	1.523 (1.377 to 1.683)	1.41 (1.266 to 1.57)	1.295 (1.162 to 1.438)	1.07 (0.928 to 1.219)	1.07 (0.928 to 1.219)	1.07 (0.928 to 1.219)	1.07 (0.928 to 1.219)
Suburachnoid hemoerhage	5 kg/m2	Both Bot	h								3.066 (1.751 to 5.334)	3.066 (1.751 to 5.334)	2.913 (1.862 to 4.395)	2.597 (1.974 to 3.375)	2.389 (1.87 to 2.998)	2.199 (1.822 to 2.672)	1.996 (1.626 to 2.416)	1.805 (1.574 to 2.06)	1.665 (1.438 to 1.932)	1.523 (1.377 to 1.683)	1.41 (1.266 to 1.57)	1.295 (1.162 to 1.438)	1.07 (0.928 to 1.219)	1.07 (0.928 to 1.219)	1.07 (0.928 to 1.219)	1.07 (0.928 to 1.219)
Hypertensive heart disease	5 kg/m2	Both Bot	h								3.122 (1.588 to 5.498)	3.122 (1.588 to 5.498)	3.0 (1.783 to 4.902)	2.769 (1.816 to 4.211)	2.573 (1.743 to 3.636)	2.407 (1.717 to 3.293)	2.281 (1.602 to 3.188)	2.159 (1.505 to 3.036)	2.035 (1.452 to 2.822)	1.955 (1.343 to 2.698)	1.86 (1.296 to 2.617)	1.792 (1.17 to 2.553)	1.697 (1.069 to 2.618)	1.697 (1.069 to 2.618)	1.697 (1.069 to 2.618)	1.697 (1.069 to 2.618)
Atrial fibrillation and flatter	5 kg/m2	Both Male	×								1.344 (1.231 to 1.473)															
Atrial fibrillation and flutter	5 kg/m2	Both Fema	les								1.346 (1.22 to 1.475)															
Asthma	5 kg/m2	Both Male	×								1.409 (1.29 to 1.545)															
Asthma	5 kg/m2	Both Fema	les								1.402 (1.275 to 1.532)															
Gallbladder and biliary diseases	5 kg/m2	Both Male	×								1.464 (1.291 to 1.64)															
Gallbladder and biliary diseases	5 kg/m2	Both Fema	les								1.729 (1.571 to 1.893)															
Alzheimer's disease and other dementias	5 kg/m2	Both Male	×								1.218 (1.054 to 1.409)															
Alzheimer's disease and other dementias	5 kg/m2	Both Fema	les								1.214 (1.047 to 1.404)															
Diabetes mellitus type 2	5 kg/m2	Both Bot	h								3.547 (2.314 to 5.219)	3.547 (2.314 to 5.219)	3.455 (2.516 to 4.691)	3.349 (2.803 to 3.918)	3.16 (2.697 to 3.699)	2.864 (2.453 to 3.312)	2.624 (2.224 to 3.038)	2.417 (2.089 to 2.779)	2.215 (1.868 to 2.606)	2.046 (1.724 to 2.379)	1.896 (1.597 to 2.228)	1.74 (1.446 to 2.074)	1.461 (1.207 to 1.758)	1.461 (1.207 to 1.758)	1.461 (1.207 to 1.758)	1.461 (1.207 to 1.758)
Chronic kidney disease due to diabetes mellitus type 2	5 kg/m2	Both Bot	h											1.746 (1.054 to 2.746)	2.036 (1.298 to 3.044)	2.036 (1.298 to 3.044)	1.621 (1.063 to 2.378)	1.621 (1.063 to 2.378)	1.431 (0.802 to 2.396)	1.431 (0.802 to 2.396)	1.431 (0.802 to 2.396)	1.431 (0.802 to 2.396)				
Chronic kidney disease due to hypertension	5 kg/m2	Both Bot	h											1.763 (1.09 to 2.755)	2.044 (1.305 to 3.082)	2.044 (1.305 to 3.082)	1.605 (1.067 to 2.31)	1.605 (1.067 to 2.31)	1.437 (0.829 to 2.415)	1.437 (0.829 to 2.415)	1.437 (0.829 to 2.415)	1.437 (0.829 to 2.415)				
Chronic kidney disease due to glomerulonephritis	5 kg/m2	Both Bot	h											1.742 (1.021 to 2.775)	2.044 (1.254 to 3.154)	2.044 (1.254 to 3.154)	1.604 (1.109 to 2.254)	1.604 (1.109 to 2.254)	1.452 (0.851 to 2.345)	1.452 (0.851 to 2.345)	1.452 (0.851 to 2.345)	1.452 (0.851 to 2.345)				
Chronic kidney disease due to oth and unspecified causes	er 5 kg/m2	Both Bot	h											1.732 (1.052 to 2.681)	2.032 (1.216 to 3.101)	2.032 (1.216 to 3.101)	1.625 (1.068 to 2.365)	1.625 (1.068 to 2.365)	1.433 (0.778 to 2.344)	1.433 (0.778 to 2.344)	1.433 (0.778 to 2.344)	1.433 (0.778 to 2.344)				
Cataract	5 kg/m2	Both Male									1.104 (1.052 to 1.157)															
Cataract	5 kg/m2	Both Fema	les								1.104 (1.051 to 1.156)															
Low back pain	5 kg/m2	Morbidity Bot	h								1.1 (1.073 to 1.126)	1.1 (1.073 to 1.126)	1.1 (1.073 to 1.127)	1.101 (1.076 to 1.128)	1.1 (1.074 to 1.126)	1.099 (1.075 to 1.123)	1.1 (1.075 to 1.128)	1.1 (1.075 to 1.126)	1.101 (1.077 to 1.126)	1.1 (1.075 to 1.126)	1.1 (1.076 to 1.124)	1.1 (1.075 to 1.124)	1.1 (1.074 to 1.125)	1.1 (1.074 to 1.125)	1.1 (1.074 to 1.125)	1.1 (1.074 to 1.125)
Gout	5 kg/m2	Both Male	x								1.628 (1.34 to 1.964)															
Gout	5 kg/m2	Both Fema	les								1.493 (1.322 to 1.677)															
Osteoarthritis of the hip	5 kg/m2	Morbidity Male	×								1.11 (1.06 to 1.157)															
Osteoarthritis of the knee	5 kg/m2	Morbidity Male	×								1.37 (1.201 to 1.538)															
Osteoarthritis of the hip	5 kg/m2	Morbidity Fema	les								1.112 (1.062 to 1.16)															
Osteoarthritis of the knee	5 kg/m2	Morbidity Fema	kes								1.375 (1.188 to 1.559)															
High body-mass index in child	Iren																									
Asthma	Obese	Both Male	×				1.315 (1.098 to 1.515)	1.315 (1.095 to 1.513)	1.316 (1.096 to 1.522)	1.321 (1.103 to 1.531)																
Asthma	Obese	Both Fema	les				1.313 (1.098 to 1.522)	1.316 (1.102 to 1.524)	1.317 (1.091 to 1.522)	1.316 (1.102 to 1.516)																
Asthma	Overweight	Both Male	×				1.312 (1.1 to 1.512)	1.316 (1.105 to 1.508)	1.316 (1.094 to 1.525)	1.319 (1.11 to 1.526)																
Asthma	Overweight	Both Fema	les				1.315 (1.0% to 1.518)	1.318 (1.1 to 1.519)	1.316 (1.106 to 1.519)	1.316 (1.091 to 1.521)																
Low bone mineral density																										
Non-hip fractures	0.1 gicm2	Both Male	×												1.077 (1.074 to 1.08)	1.114 (1.112 to 1.115)	1.151 (1.057 to 1.258)	1.182 (1.1 to 1.264)	1.214 (1.148 to 1.284)	1.247 (1.186 to 1.309)	1.297 (1.241 to 1.353)	1.339 (1.278 to 1.399)	1.37 (1.297 to 1.448)	1.37 (1.297 to 1.448)	1.37 (1.297 to 1.448)	1.37 (1.297 to 1.448)
Hip fractures	0.1 g/cm2	Both Male													2.945 (2.121 to 3.906)	2.85 (2.133 to 3.819)	2.614 (2.019 to 3.326)	2.438 (2.002 to 2.966)	2.286 (1.964 to 2.662)	2.184 (1.912 to 2.473)	2.102 (1.888 to 2.322)	1.921 (1.786 to 2.084)	1.732 (1.631 to 1.84)	1.732 (1.631 to 1.84)	1.732 (1.631 to 1.84)	1.732 (1.631 to 1.84)

Appendix Table 6a. Relative risk	is used by age and sex for each o	autcome for all risk factors	except f	or ambient air p	ollution alcohol,	and smoking.									1	ges											-
Risk - Outcome	Category / Units	Morbidity / Mortality	Sex	All-age	0-6 days	7-27 days	28-364 days	1-4 years	5-9 years	10-14 years	15-19 years	20-24 years	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Non-hip fractures	0.1 g/cm2	Both	Females													1.083 (1.08 to 1.087)	1.118 (1.116 to 1.12)	1.163 (1.065 to 1.273)	1.203 (1.119 to 1.294)	1.239 (1.161 to 1.317)	1.287 (1.216 to 1.361)	1.343 (1.273 to 1.418)	1.401 (1.33 to 1.479)	1.437 (1.352 to 1.524)	1.437 (1.352 to 1.524)	1.437 (1.352 to 1.524)	1.437 (1.352 to 1.524)
Hip fractures	0.1 g/cm2	Both	Females													3.255 (2.261 to 4.486)	2.94 (2.146 to 3.899)	2.713 (2.074 to 3.427)	2.642 (2.094 to 3.271)	2.474 (2.061 to 2.95)	2.412 (2.058 to 2.768)	2.32 (2.077 to 2.571)	2.118 (1.938 to 2.299)	1.876 (1.748 to 2.001)	1.876 (1.748 to 2.001)	1.876 (1.748 to 2.001)	1.876 (1.748 to 2.001)
Impaired kidney function																											
Ischaemic heart disease	Stage 5 CKD	Both	Both										6.403 (1.632 to 17.373)	6.579 (1.642 to 17.922)	6.222 (1.812 to 16.832)	6.454 (1.486 to 18.425)	6.576 (1.712 to 19.892)	6.638 (1.619 to 18.398)	6.447 (1.575 to 17.801)	3.682 (2.162 to 6.067)	3.674 (2.083 to 6.132)	3.058 (1.883 to 4.917)	3.066 (1.895 to 4.796)	2.611 (1.17 to 5.088)	2.546 (1.085 to 4.94)	2.593 (1.177 to 5.083)	2.545 (1.026 to 4.924)
Ischaemic heart disease	Stage 4 CKD	Both	Both										4.187 (1.636 to 8.618)	4.138 (1.626 to 9.02)	4.158 (1.576 to 9.128)	4.096 (1.662 to 8.682)	4.106 (1.554 to 8.24)	4.15 (1.564 to 9.113)	4.15 (1.579 to 9.015)	2.596 (1.79 to 3.639)	2.54 (1.751 to 3.555)	2.263 (1.621 to 3.112)	2.272 (1.624 to 3.038)	2.01 (1.167 to 3.381)	2.01 (1.18 to 3.267)	2.041 (1.192 to 3.214)	2.029 (1.172 to 3.309)
Ischaemic heart disease	Stage 3 CKD	Both	Both										1.508 (1.187 to 1.905)	1.51 (1.183 to 1.902)	1.509 (1.195 to 1.905)	1.509 (1.195 to 1.893)	1.519 (1.201 to 1.94)	1.506 (1.169 to 1.856)	1.507 (1.182 to 1.887)	1.406 (1.233 to 1.597)	1.403 (1.227 to 1.615)	1.375 (1.205 to 1.564)	1.381 (1.23 to 1.544)	1.316 (1.12 to 1.534)	1.321 (1.146 to 1.532)	1.312 (1.129 to 1.525)	1.312 (1.125 to 1.522)
Ischaemic heart disease	Albuminira	Both	Both										1.627 (1.249 to 2.107)	1.64 (1.252 to 2.097)	1.63 (1.216 to 2.103)	1.638 (1.26 to 2.081)	1.636 (1.255 to 2.08)	1.632 (1.246 to 2.074)	1.628 (1.253 to 2.071)	1.172 (1.072 to 1.272)	1.173 (1.067 to 1.287)	1.041 (0.918 to 1.178)	1.041 (0.914 to 1.173)	0.923 (0.749 to 1.107)	0.927 (0.762 to 1.115)	0.923 (0.754 to 1.125)	0.921 (0.75 to 1.122)
Ischaemic heart disease	None	Both	Both										1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Ischaemic stroke	Stage 5 CKD	Both	Both										6.665 (0.633 to 28.506)	6.631 (0.586 to 26.898)	6.571 (0.671 to 25.958)	6.66 (0.529 to 28.335)	6.568 (0.674 to 27.459)	7.071 (0.699 to 27.677)	6.859 (0.656 to 28.559)	3.305 (1.133 to 7.402)	3.34 (1.183 to 7.005)	2.175 (1.085 to 4.053)	2.154 (1.054 to 4.058)	1.608 (0.478 to 3.837)	1.567 (0.488 to 3.819)	1.539 (0.499 to 3.799)	1.544 (0.454 to 3.893)
Ischaemic stroke	Stage 4 CKD	Both	Both										3.642 (0.769 to 10.195)	3.665 (0.716 to 11.442)	3.493 (0.721 to 10.629)	3.634 (0.744 to 10.727)	3.518 (0.743 to 9.943)	3.579 (0.832 to 10.234)	3.576 (0.774 to 10.582)	2.291 (1.209 to 3.935)	2.287 (1.13 to 4.048)	1.708 (1.023 to 2.625)	1.67 (1.026 to 2.52)	1.298 (0.628 to 2.387)	1.331 (0.65 to 2.537)	1.323 (0.61 to 2.458)	1.301 (0.624 to 2.507)
Ischaemic stroke	Stage 3 CKD	Both	Both										1.063 (0.699 to 1.513)	1.073 (0.733 to 1.534)	1.071 (0.73 to 1.524)	1.072 (0.717 to 1.565)	1.081 (0.721 to 1.571)	1.077 (0.725 to 1.55)	1.077 (0.724 to 1.542)	1.324 (1.072 to 1.619)	1.323 (1.069 to 1.629)	1.255 (1.064 to 1.462)	1.254 (1.08 to 1.46)	1.155 (0.925 to 1.405)	1.148 (0.936 to 1.393)	1.147 (0.931 to 1.385)	1.156 (0.932 to 1.41)
Ischaemic stroke	Albuminira	Both	Both										2.09 (1.406 to 3.11)	2.077 (1.425 to 2.946)	2.074 (1.405 to 2.996)	2.093 (1.413 to 3.032)	2.092 (1.365 to 3.032)	2.085 (1.391 to 2.983)	2.079 (1.4 to 2.988)	1.478 (1.289 to 1.683)	1.487 (1.319 to 1.692)	1.272 (1.093 to 1.49)	1.27 (1.076 to 1.497)	1.11 (0.854 to 1.433)	1.102 (0.834 to 1.419)	1.107 (0.835 to 1.446)	1.109 (0.822 to 1.429)
Ischaemic stroke	None	Both	Both										1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Intracerebral hemorrhage	Stage 5 CKD	Both	Both										6.665 (0.633 to 28.506)	6.631 (0.586 to 26.898)	6.571 (0.671 to 25.958)	6.66 (0.529 to 28.335)	6.568 (0.674 to 27.459)	7.071 (0.699 to 27.677)	6.859 (0.656 to 28.559)	3.305 (1.133 to 7.402)	3.34 (1.183 to 7.005)	2.175 (1.085 to 4.053)	2.154 (1.054 to 4.058)	1.608 (0.478 to 3.837)	1.567 (0.488 to 3.819)	1.539 (0.499 to 3.799)	1.544 (0.454 to 3.893)
Intracerebral hemorrhage	Stage 4 CKD	Both	Both										3.642 (0.769 to 10.195)	3.665 (0.716 to 11.442)	3.493 (0.721 to 10.629)	3.634 (0.744 to 10.727)	3.518 (0.743 to 9.943)	3.579 (0.832 to 10.234)	3.576 (0.774 to 10.582)	2.291 (1.209 to 3.935)	2.287 (1.13 to 4.048)	1.708 (1.023 to 2.625)	1.67 (1.026 to 2.52)	1.298 (0.628 to 2.387)	1.331 (0.65 to 2.537)	1.323 (0.61 to 2.458)	1.301 (0.624 to 2.507)
Intracerebral hemorrhage	Stage 3 CKD	Both	Both										1.063 (0.699 to 1.513)	1.073 (0.733 to 1.534)	1.071 (0.73 to 1.524)	1.072 (0.717 to 1.565)	1.081 (0.721 to 1.571)	1.077 (0.725 to 1.55)	1.077 (0.724 to 1.542)	1.324 (1.072 to 1.619)	1.323 (1.069 to 1.629)	1.255 (1.064 to 1.462)	1.254 (1.08 to 1.46)	1.155 (0.925 to 1.405)	1.148 (0.936 to 1.393)	1.147 (0.931 to 1.385)	1.156 (0.932 to 1.41)
Intracerebral hemorrhage	Albuminira	Both	Both										2.09 (1.406 to 3.11)	2.077 (1.425 to 2.946)	2.074 (1.405 to 2.996)	2.093 (1.413 to 3.032)	2.092 (1.365 to 3.032)	2.085 (1.391 to 2.983)	2.079 (1.4 to 2.988)	1.478 (1.289 to 1.683)	1.487 (1.319 to 1.692)	1.272 (1.093 to 1.49)	1.27 (1.076 to 1.497)	1.11 (0.854 to 1.433)	1.102 (0.834 to 1.419)	1.107 (0.835 to 1.446)	1.109 (0.822 to 1.429)
Intracerebral hemorrhage	None	Both	Both										1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Peripheral vascular disease	Stage 5 CKD	Both	Both										207.753 (30.856 to 704.06)	200.617 (34.114 to 695.696)	211.239 (32.055 to 800.634)	205.926 (31.084 to 751.759)	200.856 (33.786 to 732.241)	207.342 (30.718 to 650.978)	202.164 (32.222 to 701.663)	17.647 (7.939 to 32.318)	17.353 (8.097 to 31.55)	5.347 (2.334 to 10.628)	5.427 (2.272 to 10.749)	1.895 (0.429 to 5.39)	1.852 (0.444 to 5.291)	1.856 (0.437 to 5.035)	1.925 (0.427 to 5.377)
Peripheral vascular disease	Stage 4 CKD	Both	Both										58.362 (15.568 to 149.001)	54.062 (15.177 to 136.039)	54.638 (14.697 to 139.549)	55.201 (16.279 to 139.398)	54.066 (14.955 to 142.728)	55.805 (15.506 to 137.121)	55.87 (15.838 to 139.768)	7.713 (4.647 to 11.89)	7.845 (4.686 to 12.511)	3.141 (1.836 to 5.218)	3.213 (1.87 to 5.196)	1.464 (0.558 to 3.037)	1.47 (0.601 to 3.006)	1.462 (0.599 to 3.073)	1.488 (0.593 to 3.038)
Peripheral vascular disease	Stage 3 CKD	Both	Both										3.083 (1.826 to 4.774)	3.102 (1.839 to 4.864)	3.045 (1.792 to 4.748)	3.114 (1.906 to 4.86)	3.039 (1.817 to 4.807)	3.117 (1.883 to 4.881)	3.113 (1.839 to 4.781)	2.055 (1.57 to 2.665)	2.05 (1.584 to 2.635)	1.525 (1.216 to 1.891)	1.531 (1.221 to 1.92)	1.142 (0.868 to 1.461)	1.158 (0.875 to 1.513)	1.147 (0.871 to 1.476)	1.158 (0.879 to 1.484)
Peripheral vascular disease	Albuminira	Both	Both										3.542 (1.986 to 5.721)	3.572 (1.98 to 5.768)	3.57 (1.977 to 5.944)	3.569 (1.991 to 5.91)	3.543 (1.934 to 5.923)	3.607 (1.98 to 6.203)	3.552 (1.902 to 5.879)	1.758 (1.442 to 2.135)	1.759 (1.451 to 2.101)	1.336 (1.033 to 1.695)	1.327 (1.018 to 1.727)	1.037 (0.664 to 1.542)	1.029 (0.687 to 1.478)	1.033 (0.646 to 1.496)	1.038 (0.68 to 1.548)
Peripheral vascular disease	None	Both	Both										1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Goat	Stage 5 CKD	Both	Both										2.749 (2.491 to 3.017)	2.747 (2.47 to 3.035)	2.747 (2.482 to 3.035)	2.736 (2.466 to 3.014)	2.741 (2.48 to 3.053)	2.729 (2.47 to 3.007)	2.74 (2.479 to 3.038)	2.747 (2.485 to 3.049)	2.742 (2.479 to 3.021)	2.742 (2.496 to 3.039)	2.74 (2.48 to 3.042)	2.736 (2.468 to 3.013)	2.736 (2.468 to 3.013)	2.736 (2.468 to 3.013)	2.736 (2.468 to 3.013)
Gout	Stage 4 CKD	Both	Both										2.745 (2.473 to 3.032)	2.743 (2.484 to 3.014)	2.742 (2.473 to 3.034)	2.742 (2.442 to 3.029)	2.746 (2.494 to 3.045)	2.748 (2.483 to 3.045)	2.744 (2.462 to 3.029)	2.749 (2.477 to 3.018)	2.755 (2.495 to 3.038)	2.747 (2.473 to 3.034)	2.738 (2.463 to 3.041)	2.744 (2.478 to 3.021)	2.744 (2.478 to 3.021)	2.744 (2.478 to 3.021)	2.744 (2.478 to 3.021)
Gout	Stage 3 CKD	Both	Both										2.748 (2.457 to 3.035)	2.753 (2.48 to 3.038)	2.743 (2.465 to 3.024)	2.748 (2.491 to 3.029)	2.743 (2.467 to 3.051)	2.749 (2.473 to 3.042)	2.744 (2.475 to 3.017)	2.741 (2.464 to 3.034)	2.745 (2.478 to 3.037)	2.739 (2.465 to 3.006)	2.745 (2.473 to 3.029)	2.747 (2.475 to 3.028)	2.747 (2.475 to 3.028)	2.747 (2.475 to 3.028)	2.747 (2.475 to 3.028)
Gout	Albaninira	Both	Both										1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Gout	None	Both	Both										1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
* Occupational noise rela ** Shifts are reported for	ative risk values are for sequela r diet hich in sodium as the estimatics	n is based on mediation through	oh hioh sy	stolic blood pressu	D ^e																						

** Shifts are reported for diet high in sodium as the estimation is based on mediation through high systolic blood pressure.

Risk-outcome pairs with 100% attribution		
Alcohol use	High fasting plasma glucose	Occupational particulate matter, gases, and fumes
Liver cancer due to alcohol use	Diabetes mellitus	Coal workers pneumoconiosis
Cirrhosis due to alcohol use	Chronic kidney disease due to diabetes mellitus	Unsafe sex
Alcohol use disorders	High systolic blood pressure	Syphilis
Childhood underweight	Hypertensive heart disease	Chlamydial infection
Protein-energy malnutrition	Chronic kidney disease due to hypertension	Gonococcal infection
Childhood wasting	Iron deficiency	Trichomoniasis
Protein-energy malnutrition	Iron-deficiency anaemia	Genital herpes
Drug use	Low glomerular filtration rate	Other sexually transmitted diseases
Opiod use disorders	Chronic kidney disease due to diabetes mellitus	Cervical cancer
Cocaine use disorders	Chronic kidney disease due to hypertension	Sexually transmitted diseases excluding HIV
Amphetamine use disorders	Chronic kidney disease due to glomerulonephritis	Vitamin A deficiency
Cannabis use disorders	Chronic kidney disease due to other causes	Vitamin A deficiency
Other drug use disorders		

Appendix Table 6b. Relative risks use	d by age and	sex for each ou	itcome for	the particulat	e matter integ	rated exposur	e response cu	rve.											
	Category /	Morbidity /	~					L		L	A	.ge	L	I					
Risk - Outcome	Units	Mortality	Sex	All ages	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Ambient particulate matter pollution (FM2.5)				2.29															
Lower respiratory infections	$600 \ \mu\text{g/m}^3$	Both	Both	(1.968 to 2.776)															
Lower respiratory infections	$500 \ \mu g/m^3$	Both	Both	2.347 (1.936 to 2.735)															
Lower respiratory infections	$400 \ \mu g/m^3$	Both	Both	2.297 (1.883 to 2.687)															
Lower respiratory infections	$300 \; \mu g/m^3$	Both	Both	2.213 (1.809 to 2.615)															
Lower respiratory infections	$200 \; \mu g/m^3$	Both	Both	2.062 (1.702 to 2.44)															
Lower respiratory infections	$150\;\mu g/m^3$	Both	Both	1.938 (1.629 to 2.281)															
Lower respiratory infections	135 µg/m³	Both	Both	1.891 (1.6 to 2.209)															
Lower respiratory infections	120 µg/m ³	Both	Both	1.838 (1.571 to 2.129)															
Lower respiratory infections	105 µg/m ³	Both	Both	1.778 (1.54 to 2.05)															
Lower respiratory infections	90 µg/m³	Both	Both	1.711 (1.505 to 1.945)															
Lower respiratory infections	75 µg/m³	Both	Both	1.634 (1.455 to 1.827)															
Lower respiratory infections	60 µg/m ³	Both	Both	1.546 (1.4 to 1.711)															
Lower respiratory infections	45 µg/m³	Both	Both	1.443 (1.323 to 1.576)															
Lower respiratory infections	30 µg/m ³	Both	Both	1.322 (1.225 to 1.428)															
Lower respiratory infections	25 µg/m³	Both	Both	1.276 (1.184 to 1.379)															
Lower respiratory infections	20 µg/m ³	Both	Both	1.226 (1.14 to 1.335)															
Lower respiratory infections	15 µg/m³	Both	Both	1.171 (1.093 to 1.282)															
Lower respiratory infections	10 µg/m ³	Both	Both	1.108 (1.046 to 1.219)															
Lower respiratory infections	5 µg/m ³	Both	Both	1.025 (1.0 to 1.119)															
Lower respiratory infections	$0 \ \mu g/m^3$	Both	Both	1.0 (1.0 to 1.0)															
Tracheal, bronchus, and lung cancer	600 µg/m ³	Both	Both	2.541 (2.222 to 2.868)															
Tracheal, bronchus, and lung cancer	$500 \ \mu g/m^3$	Both	Both	2.369 (2.074 to 2.675)															
Tracheal, bronchus, and lung cancer	$400 \ \mu g/m^3$	Both	Both	2.185 (1.919 to 2.469)															
Tracheal, bronchus, and lung cancer	300 µg/m ³	Both	Both	1.982 (1.748 to 2.229)															
Tracheal, bronchus, and lung cancer	$200 \; \mu g/m^3$	Both	Both	1.753 (1.56 to 1.958)															
Tracheal, bronchus, and lung cancer	$150\;\mu g/m^3$	Both	Both	1.622 (1.454 to 1.802)															
Tracheal, bronchus, and lung cancer	$135\;\mu g/m^3$	Both	Both	1.58 (1.421 to 1.754)															
Tracheal, bronchus, and lung cancer	$120\;\mu g/m^3$	Both	Both	1.536 (1.385 to 1.701)															
Tracheal, bronchus, and lung cancer	105 µg/m ³	Both	Both	1.49 (1.348 to 1.648)															
I				1 7	I														

Appendix Table 6b. Relative risks us	ed by age and	sex for each or	utcome for	the particulat	e matter integ	grated exposur	e response cu	irve.											
	Category /	Morbidity /	0								A	lige	1 1 1 1						
Kisk - Outcome	Units	Mortality	Sex	All ages	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Tracheal, bronchus, and lung cancer	90 μg/m ³	Both	Both	1.442															
To deal book and have a set	75	Date	Deth	(1.309 to 1.588) 1.39															
I racheai, bronchus, and lung cancer	/5 μg/m²	Both	Both	(1.269 to 1.526)															
Tracheal, bronchus, and lung cancer	60 μg/m ³	Both	Both	(1.227 to 1.458)															
Tracheal, bronchus, and lung cancer	45 μg/m ³	Both	Both	(1.18 to 1.383)															
Tracheal, bronchus, and lung cancer	$30 \ \mu g/m^3$	Both	Both	1.203 (1.128 to 1.295)															
Tracheal, bronchus, and lung cancer	$25 \ \mu g/m^3$	Both	Both	1.177 (1.109 to 1.261)															
Tracheal, bronchus, and lung cancer	$20 \ \mu\text{g/m}^{\scriptscriptstyle 3}$	Both	Both	1.148 (1.089 to 1.223)															
Tracheal, bronchus, and lung cancer	$15 \ \mu g/m^3$	Both	Both	1.116 (1.067 to 1.178)															
Tracheal, bronchus, and lung cancer	$10 \ \mu g/m^3$	Both	Both	1.077 (1.041 to 1.126)															
Tracheal, bronchus, and lung cancer	$5 \ \mu g/m^3$	Both	Both	1.02 (1.0 to 1.056)															
Tracheal, bronchus, and lung cancer	$0\ \mu g/m^3$	Both	Both	1.0 (1.0 to 1.0)															
Ischaemic heart disease	600 µg/m³	Both	Both		2.244 (1.922 to 2.608)	2.142 (1.848 to 2.464)	2.033 (1.78 to 2.343)	1.917 (1.671 to 2.195)	1.831 (1.624 to 2.079)	1.743 (1.557 to 1.943)	1.661 (1.5 to 1.837)	1.583 (1.437 to 1.739)	1.506 (1.384 to 1.639)	1.434 (1.329 to 1.545)	1.371 (1.287 to 1.462)	1.307 (1.231 to 1.384)	1.249 (1.191 to 1.309)	1.193 (1.151 to 1.241)	1.141 (1.108 to 1.174)
Ischaemic heart disease	500 µg/m3	Both	Both		2.166 (1.852 to 2.523)	2.073 (1.784 to 2.395)	1.971 (1.726 to 2.279)	1.863 (1.618 to 2.133)	1.783 (1.581 to 2.031)	1.701 (1.52 to 1.898)	1.625 (1.465 to 1.8)	1.551 (1.406 to 1.705)	1.479 (1.358 to 1.608)	1.412 (1.307 to 1.52)	1.352 (1.269 to 1.442)	1.292 (1.216 to 1.368)	1.237 (1.179 to 1.297)	1.184 (1.142 to 1.232)	1.134 (1.101 to 1.168)
Ischaemic heart disease	400 µg/m ³	Both	Both		2.076 (1.77 to 2.427)	1.993 (1.708 to 2.317)	1.901 (1.663 to 2.201)	1.8 (1.559 to 2.069)	1.727 (1.53 to 1.977)	1.652 (1.475 to 1.843)	1.582 (1.425 to 1.755)	1.515 (1.371 to 1.665)	1.448 (1.329 to 1.576)	1.386 (1.282 to 1.496)	1.331 (1.249 to 1.419)	1.274 (1.199 to 1.351)	1.223 (1.165 to 1.282)	1.174 (1.131 to 1.222)	1.127 (1.094 to 1.161)
Ischaemic heart disease	300 µg/m ³	Both	Both		1.971 (1.678 to 2.305)	1.898 (1.622 to 2.217)	1.816 (1.585 to 2.11)	1.725 (1.492 to 1.981)	1.662 (1.469 to 1.903)	1.594 (1.42 to 1.782)	1.532 (1.38 to 1.703)	1.471 (1.332 to 1.618)	1.41 (1.295 to 1.535)	1.354 (1.251 to 1.463)	1.304 (1.224 to 1.394)	1.253 (1.179 to 1.329)	1.206 (1.149 to 1.264)	1.161 (1.119 to 1.208)	1.117 (1.085 to 1.152)
Ischaemic heart disease	200 µg/m3	Both	Both		1.838 (1.562 to 2.163)	1.779 (1.515 to 2.08)	1.71 (1.493 to 1.989)	1.631 (1.413 to 1.879)	1.579 (1.393 to 1.809)	1.521 (1.356 to 1.704)	1.467 (1.325 to 1.634)	1.416 (1.282 to 1.561)	1.363 (1.253 to 1.488)	1.314 (1.215 to 1.423)	1.271 (1.192 to 1.362)	1.225 (1.154 to 1.302)	1.184 (1.128 to 1.241)	1.144 (1.103 to 1.19)	1.106 (1.073 to 1.139)
Ischaemic heart disease	150 µg/m ³	Both	Both		1.754 (1.492 to 2.069)	1.703 (1.452 to 1.992)	1.643 (1.433 to 1.914)	1.572 (1.362 to 1.807)	1.526 (1.346 to 1.75)	1.474 (1.313 to 1.652)	1.426 (1.29 to 1.591)	1.38 (1.253 to 1.52)	1.332 (1.226 to 1.455)	1.288 (1.192 to 1.397)	1.249 (1.172 to 1.339)	1.207 (1.138 to 1.283)	1.17 (1.115 to 1.226)	1.133 (1.094 to 1.179)	1.098 (1.066 to 1.132)
Ischaemic heart disease	135 µg/m³	Both	Both		1.726 (1.466 to 2.036)	1.677 (1.43 to 1.965)	1.62 (1.413 to 1.889)	1.551 (1.345 to 1.782)	1.507 (1.33 to 1.732)	1.457 (1.299 to 1.634)	1.412 (1.278 to 1.576)	1.367 (1.243 to 1.507)	1.321 (1.217 to 1.443)	1.279 (1.184 to 1.387)	1.241 (1.165 to 1.331)	1.201 (1.132 to 1.277)	1.165 (1.11 to 1.221)	1.129 (1.09 to 1.176)	1.095 (1.064 to 1.129)
Ischaemic heart disease	120 µg/m ³	Both	Both		1.695 (1.439 to 2.0)	1.649 (1.408 to 1.935)	1.595 (1.394 to 1.86)	1.529 (1.326 to 1.759)	1.487 (1.313 to 1.71)	1.44 (1.284 to 1.614)	1.396 (1.266 to 1.559)	1.354 (1.232 to 1.491)	1.31 (1.207 to 1.431)	1.269 (1.176 to 1.376)	1.233 (1.158 to 1.321)	1.194 (1.127 to 1.27)	1.16 (1.106 to 1.215)	1.125 (1.086 to 1.171)	1.092 (1.061 to 1.126)
Ischaemic heart disease	105 µg/m ³	Both	Both		1.661 (1.413 to 1.963)	1.619 (1.383 to 1.904)	1.567 (1.373 to 1.829)	1.505 (1.306 to 1.731)	1.466 (1.295 to 1.686)	1.421 (1.268 to 1.593)	1.38 (1.253 to 1.542)	1.339 (1.22 to 1.474)	1.297 (1.196 to 1.418)	1.258 (1.167 to 1.363)	1.224 (1.15 to 1.312)	1.187 (1.121 to 1.262)	1.154 (1.101 to 1.209)	1.121 (1.082 to 1.167)	1.089 (1.058 to 1.122)
Ischaemic heart disease	$90\;\mu g/m^3$	Both	Both		1.624 (1.382 to 1.921)	1.585 (1.356 to 1.86)	1.537 (1.347 to 1.794)	1.478 (1.284 to 1.7)	1.442 (1.276 to 1.66)	1.399 (1.251 to 1.568)	1.361 (1.238 to 1.518)	1.323 (1.206 to 1.457)	1.283 (1.185 to 1.402)	1.246 (1.157 to 1.35)	1.214 (1.141 to 1.3)	1.179 (1.113 to 1.252)	1.147 (1.094 to 1.202)	1.115 (1.078 to 1.161)	1.085 (1.055 to 1.119)
Ischaemic heart disease	$75 \ \mu g/m^3$	Both	Both		1.583 (1.345 to 1.871)	1.547 (1.325 to 1.813)	1.503 (1.32 to 1.755)	1.448 (1.259 to 1.665)	1.415 (1.254 to 1.628)	1.376 (1.232 to 1.542)	1.34 (1.22 to 1.495)	1.305 (1.19 to 1.437)	1.267 (1.171 to 1.383)	1.233 (1.146 to 1.335)	1.202 (1.131 to 1.287)	1.169 (1.106 to 1.241)	1.14 (1.088 to 1.193)	1.11 (1.073 to 1.155)	1.081 (1.051 to 1.114)
Ischaemic heart disease	$60\;\mu g/m^3$	Both	Both		1.535 (1.308 to 1.813)	1.504 (1.291 to 1.761)	1.464 (1.287 to 1.709)	1.413 (1.232 to 1.621)	1.384 (1.229 to 1.591)	1.348 (1.21 to 1.511)	1.315 (1.2 to 1.463)	1.283 (1.172 to 1.412)	1.249 (1.156 to 1.361)	1.217 (1.134 to 1.317)	1.189 (1.12 to 1.272)	1.158 (1.096 to 1.229)	1.131 (1.081 to 1.183)	1.103 (1.067 to 1.148)	1.076 (1.047 to 1.109)
Ischaemic heart disease	$45\;\mu g/m^3$	Both	Both		1.479 (1.268 to 1.741)	1.452 (1.251 to 1.695)	1.417 (1.25 to 1.652)	1.372 (1.2 to 1.573)	1.347 (1.202 to 1.546)	1.314 (1.184 to 1.473)	1.286 (1.176 to 1.431)	1.257 (1.152 to 1.381)	1.226 (1.139 to 1.337)	1.198 (1.118 to 1.295)	1.173 (1.107 to 1.253)	1.145 (1.086 to 1.214)	1.12 (1.072 to 1.171)	1.095 (1.06 to 1.138)	1.07 (1.042 to 1.102)
Ischaemic heart disease	$30 \ \mu g/m^3$	Both	Both		1.406 (1.219 to 1.651)	1.385 (1.205 to 1.609)	1.357 (1.205 to 1.578)	1.318 (1.163 to 1.506)	1.299 (1.167 to 1.492)	1.271 (1.15 to 1.423)	1.247 (1.146 to 1.387)	1.224 (1.126 to 1.342)	1.197 (1.116 to 1.303)	1.173 (1.098 to 1.265)	1.152 (1.09 to 1.229)	1.128 (1.071 to 1.193)	1.106 (1.061 to 1.156)	1.084 (1.05 to 1.126)	1.062 (1.036 to 1.093)
Ischaemic heart disease	$25\;\mu g/m^3$	Both	Both		1.376 (1.198 to 1.614)	1.357 (1.187 to 1.574)	1.332 (1.187 to 1.546)	1.296 (1.149 to 1.479)	1.278 (1.152 to 1.464)	1.253 (1.137 to 1.402)	1.231 (1.133 to 1.365)	1.209 (1.115 to 1.323)	1.185 (1.105 to 1.289)	1.163 (1.09 to 1.253)	1.143 (1.082 to 1.219)	1.12 (1.066 to 1.184)	1.1 (1.056 to 1.149)	1.079 (1.047 to 1.121)	1.059 (1.033 to 1.089)
Ischaemic heart disease	$20\;\mu g/m^3$	Both	Both		1.341 (1.175 to 1.566)	1.324 (1.166 to 1.533)	1.303 (1.166 to 1.506)	1.27 (1.132 to 1.446)	1.254 (1.134 to 1.43)	1.232 (1.122 to 1.374)	1.212 (1.119 to 1.342)	1.192 (1.103 to 1.302)	1.17 (1.095 to 1.272)	1.15 (1.081 to 1.237)	1.132 (1.074 to 1.206)	1.111 (1.06 to 1.173)	1.093 (1.051 to 1.141)	1.073 (1.042 to 1.114)	1.055 (1.03 to 1.084)
Ischaemic heart disease	15 µg/m ³	Both	Both		1.298 (1.143 to 1.508)	1.284 (1.139 to 1.476)	1.266 (1.14 to 1.458)	1.237 (1.112 to 1.404)	1.225 (1.113 to 1.386)	1.205 (1.103 to 1.341)	1.188 (1.102 to 1.315)	1.171 (1.088 to 1.276)	1.152 (1.081 to 1.249)	1.134 (1.07 to 1.219)	1.118 (1.064 to 1.188)	1.1 (1.052 to 1.159)	1.083 (1.044 to 1.13)	1.066 (1.037 to 1.107)	1.05 (1.026 to 1.078)
Ischaemic heart disease	$10\;\mu g/m^3$	Both	Both		1.238 (1.105 to 1.425)	1.229 (1.101 to 1.402)	1.215 (1.102 to 1.385)	1.192 (1.083 to 1.339)	1.183 (1.086 to 1.325)	1.168 (1.077 to 1.293)	1.155 (1.078 to 1.268)	1.141 (1.068 to 1.241)	1.125 (1.062 to 1.217)	1.112 (1.053 to 1.19)	1.099 (1.049 to 1.162)	1.084 (1.041 to 1.139)	1.07 (1.036 to 1.113)	1.056 (1.03 to 1.095)	1.042 (1.021 to 1.069)

Appendix Table ob. Relative risks us	Category /	Morbidity /	iteome for	the particulat	te matter integ	rated exposu	re response et	irve.			A	sge							
Risk - Outcome	Units	Mortality	Sex	All ages	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Ambient particulate matter pollution (PM2.5)																		
Ischaemic heart disease	$5\ \mu g/m^3$	Both	Both		1.097 (1.0 to 1.284)	1.095 (1.0 to 1.266)	1.088 (1.0 to 1.253)	1.079 (1.0 to 1.228)	1.077 (1.0 to 1.209)	1.071 (1.0 to 1.188)	1.067 (1.0 to 1.181)	1.062 (1.0 to 1.166)	1.055 (1.0 to 1.145)	1.049 (1.0 to 1.126)	1.045 (1.0 to 1.114)	1.038 (1.0 to 1.098)	1.032 (1.0 to 1.081)	1.026 (1.0 to 1.067)	1.02 (1.0 to 1.051)
Ischaemic heart disease	$0 \ \mu g/m^3$	Both	Both		1.0 (1.0 to 1.0)														
Stroke	$600 \ \mu g/m^3$	Both	Both		1.704 (1.392 to 1.966)	1.641 (1.36 to 1.902)	1.587 (1.303 to 1.823)	1.535 (1.297 to 1.749)	1.481 (1.269 to 1.674)	1.436 (1.245 to 1.606)	1.39 (1.225 to 1.552)	1.347 (1.201 to 1.489)	1.306 (1.186 to 1.427)	1.267 (1.159 to 1.367)	1.228 (1.136 to 1.313)	1.191 (1.117 to 1.256)	1.158 (1.097 to 1.215)	1.123 (1.074 to 1.167)	1.091 (1.058 to 1.127)
Stroke	$500 \ \mu g/m^3$	Both	Both		1.664 (1.355 to 1.925)	1.606 (1.328 to 1.862)	1.555 (1.275 to 1.791)	1.506 (1.271 to 1.719)	1.455 (1.244 to 1.651)	1.413 (1.223 to 1.585)	1.37 (1.207 to 1.529)	1.329 (1.184 to 1.473)	1.291 (1.17 to 1.412)	1.254 (1.146 to 1.354)	1.216 (1.125 to 1.302)	1.181 (1.109 to 1.247)	1.15 (1.089 to 1.208)	1.117 (1.068 to 1.161)	1.087 (1.053 to 1.123)
Stroke	$400 \; \mu g/m^3$	Both	Both		1.619 (1.314 to 1.881)	1.565 (1.293 to 1.822)	1.518 (1.244 to 1.755)	1.472 (1.242 to 1.68)	1.426 (1.217 to 1.623)	1.387 (1.199 to 1.559)	1.346 (1.186 to 1.505)	1.308 (1.165 to 1.452)	1.273 (1.153 to 1.394)	1.238 (1.131 to 1.34)	1.203 (1.113 to 1.29)	1.171 (1.098 to 1.237)	1.141 (1.081 to 1.2)	1.11 (1.061 to 1.155)	1.082 (1.049 to 1.117)
Stroke	$300 \ \mu g/m^3$	Both	Both		1.565 (1.27 to 1.825)	1.516 (1.249 to 1.773)	1.474 (1.208 to 1.709)	1.432 (1.21 to 1.641)	1.39 (1.185 to 1.587)	1.355 (1.174 to 1.525)	1.318 (1.162 to 1.479)	1.283 (1.143 to 1.425)	1.251 (1.133 to 1.373)	1.22 (1.114 to 1.322)	1.187 (1.098 to 1.275)	1.157 (1.086 to 1.224)	1.131 (1.072 to 1.189)	1.102 (1.054 to 1.147)	1.076 (1.043 to 1.111)
Stroke	$200 \; \mu g/m^3$	Both	Both		1.496 (1.219 to 1.757)	1.454 (1.203 to 1.712)	1.418 (1.166 to 1.646)	1.381 (1.17 to 1.591)	1.345 (1.148 to 1.542)	1.314 (1.142 to 1.483)	1.282 (1.132 to 1.445)	1.251 (1.117 to 1.394)	1.223 (1.108 to 1.343)	1.196 (1.094 to 1.299)	1.167 (1.081 to 1.255)	1.14 (1.072 to 1.207)	1.117 (1.06 to 1.175)	1.092 (1.046 to 1.137)	1.068 (1.036 to 1.104)
Stroke	$150 \; \mu g/m^3$	Both	Both		1.453 (1.189 to 1.705)	1.415 (1.175 to 1.667)	1.382 (1.141 to 1.605)	1.348 (1.146 to 1.558)	1.315 (1.125 to 1.509)	1.288 (1.123 to 1.453)	1.259 (1.115 to 1.42)	1.231 (1.102 to 1.375)	1.205 (1.094 to 1.325)	1.181 (1.081 to 1.283)	1.154 (1.07 to 1.24)	1.13 (1.063 to 1.195)	1.108 (1.053 to 1.166)	1.085 (1.04 to 1.13)	1.063 (1.032 to 1.098)
Stroke	$135 \; \mu g/m^3$	Both	Both		1.438 (1.179 to 1.69)	1.401 (1.166 to 1.653)	1.37 (1.133 to 1.591)	1.337 (1.138 to 1.545)	1.305 (1.118 to 1.498)	1.279 (1.117 to 1.443)	1.251 (1.109 to 1.412)	1.224 (1.096 to 1.366)	1.199 (1.089 to 1.318)	1.175 (1.077 to 1.277)	1.15 (1.067 to 1.236)	1.126 (1.06 to 1.191)	1.105 (1.05 to 1.163)	1.082 (1.038 to 1.127)	1.062 (1.031 to 1.096)
Stroke	$120\;\mu\text{g/m}^{3}$	Both	Both		1.421 (1.168 to 1.672)	1.386 (1.156 to 1.637)	1.357 (1.125 to 1.576)	1.325 (1.13 to 1.531)	1.295 (1.111 to 1.486)	1.269 (1.111 to 1.433)	1.243 (1.103 to 1.403)	1.216 (1.091 to 1.355)	1.192 (1.083 to 1.311)	1.169 (1.073 to 1.271)	1.145 (1.063 to 1.231)	1.122 (1.057 to 1.186)	1.102 (1.047 to 1.16)	1.08 (1.036 to 1.125)	1.06 (1.029 to 1.094)
Stroke	$105 \ \mu g/m^3$	Both	Both		1.404 (1.157 to 1.65)	1.37 (1.146 to 1.619)	1.342 (1.115 to 1.559)	1.312 (1.121 to 1.517)	1.283 (1.104 to 1.472)	1.258 (1.103 to 1.421)	1.233 (1.097 to 1.393)	1.208 (1.085 to 1.342)	1.185 (1.078 to 1.304)	1.163 (1.068 to 1.265)	1.139 (1.059 to 1.226)	1.117 (1.053 to 1.181)	1.098 (1.044 to 1.156)	1.077 (1.034 to 1.122)	1.058 (1.027 to 1.093)
Stroke	$90 \; \mu g/m^3$	Both	Both		1.384 (1.144 to 1.632)	1.352 (1.135 to 1.599)	1.326 (1.105 to 1.539)	1.297 (1.112 to 1.501)	1.269 (1.096 to 1.457)	1.246 (1.095 to 1.41)	1.222 (1.09 to 1.381)	1.198 (1.078 to 1.332)	1.177 (1.072 to 1.293)	1.156 (1.062 to 1.258)	1.133 (1.055 to 1.219)	1.112 (1.049 to 1.176)	1.094 (1.041 to 1.151)	1.074 (1.032 to 1.118)	1.055 (1.026 to 1.09)
Stroke	$75 \ \mu g/m^3$	Both	Both		1.362 (1.131 to 1.608)	1.332 (1.122 to 1.574)	1.307 (1.094 to 1.52)	1.28 (1.102 to 1.482)	1.254 (1.086 to 1.439)	1.233 (1.086 to 1.398)	1.211 (1.082 to 1.368)	1.188 (1.071 to 1.321)	1.167 (1.065 to 1.283)	1.148 (1.056 to 1.249)	1.126 (1.05 to 1.212)	1.106 (1.045 to 1.17)	1.089 (1.038 to 1.146)	1.07 (1.029 to 1.114)	1.053 (1.024 to 1.087)
Stroke	$60 \; \mu g/m^3$	Both	Both		1.336 (1.116 to 1.574)	1.309 (1.108 to 1.542)	1.286 (1.083 to 1.492)	1.261 (1.09 to 1.461)	1.237 (1.076 to 1.418)	1.217 (1.074 to 1.377)	1.197 (1.073 to 1.352)	1.175 (1.063 to 1.306)	1.157 (1.059 to 1.27)	1.138 (1.05 to 1.239)	1.118 (1.045 to 1.203)	1.1 (1.041 to 1.163)	1.084 (1.035 to 1.14)	1.066 (1.026 to 1.109)	1.049 (1.021 to 1.083)
Stroke	$45\;\mu\text{g/m}^{3}$	Both	Both		1.305 (1.099 to 1.532)	1.281 (1.093 to 1.502)	1.261 (1.069 to 1.458)	1.237 (1.077 to 1.433)	1.216 (1.064 to 1.392)	1.198 (1.062 to 1.354)	1.18 (1.063 to 1.331)	1.161 (1.054 to 1.288)	1.144 (1.051 to 1.254)	1.127 (1.044 to 1.226)	1.109 (1.039 to 1.192)	1.091 (1.036 to 1.153)	1.077 (1.03 to 1.132)	1.061 (1.023 to 1.103)	1.046 (1.019 to 1.079)
Stroke	$30 \; \mu g/m^3$	Both	Both		1.265 (1.077 to 1.48)	1.244 (1.073 to 1.455)	1.227 (1.054 to 1.415)	1.207 (1.061 to 1.396)	1.189 (1.05 to 1.357)	1.174 (1.05 to 1.326)	1.158 (1.05 to 1.303)	1.141 (1.043 to 1.264)	1.126 (1.041 to 1.233)	1.112 (1.035 to 1.208)	1.096 (1.031 to 1.175)	1.081 (1.029 to 1.141)	1.069 (1.025 to 1.122)	1.054 (1.019 to 1.095)	1.041 (1.016 to 1.073)
Stroke	$25\;\mu\text{g/m}^{\scriptscriptstyle 3}$	Both	Both		1.248 (1.069 to 1.46)	1.229 (1.066 to 1.436)	1.213 (1.047 to 1.396)	1.194 (1.054 to 1.38)	1.177 (1.045 to 1.342)	1.163 (1.045 to 1.312)	1.149 (1.045 to 1.291)	1.133 (1.039 to 1.253)	1.119 (1.037 to 1.223)	1.106 (1.031 to 1.2)	1.091 (1.028 to 1.168)	1.076 (1.027 to 1.136)	1.065 (1.022 to 1.117)	1.051 (1.018 to 1.091)	1.038 (1.014 to 1.07)
Stroke	$20\;\mu\text{g/m}^{3}$	Both	Both		1.227 (1.06 to 1.43)	1.211 (1.056 to 1.409)	1.196 (1.041 to 1.373)	1.179 (1.047 to 1.359)	1.164 (1.039 to 1.32)	1.151 (1.039 to 1.297)	1.137 (1.04 to 1.275)	1.123 (1.034 to 1.24)	1.11 (1.032 to 1.213)	1.098 (1.027 to 1.189)	1.084 (1.024 to 1.16)	1.071 (1.024 to 1.129)	1.06 (1.02 to 1.112)	1.047 (1.016 to 1.087)	1.036 (1.012 to 1.067)
Stroke	15 µg/m³	Both	Both		1.202 (1.05 to 1.396)	1.188 (1.046 to 1.381)	1.176 (1.033 to 1.346)	1.16 (1.039 to 1.334)	1.146 (1.033 to 1.295)	1.135 (1.031 to 1.277)	1.123 (1.033 to 1.256)	1.11 (1.028 to 1.222)	1.099 (1.027 to 1.196)	1.088 (1.023 to 1.175)	1.076 (1.02 to 1.148)	1.064 (1.02 to 1.12)	1.055 (1.017 to 1.104)	1.043 (1.013 to 1.081)	1.033 (1.01 to 1.064)
Stroke	$10\;\mu\text{g/m}^{3}$	Both	Both		1.167 (1.036 to 1.347)	1.155 (1.032 to 1.333)	1.146 (1.024 to 1.303)	1.133 (1.027 to 1.297)	1.122 (1.025 to 1.261)	1.113 (1.023 to 1.248)	1.104 (1.024 to 1.228)	1.093 (1.021 to 1.196)	1.083 (1.019 to 1.174)	1.075 (1.017 to 1.157)	1.064 (1.015 to 1.133)	1.054 (1.015 to 1.107)	1.047 (1.012 to 1.094)	1.037 (1.01 to 1.073)	1.028 (1.008 to 1.057)
Stroke	$5 \; \mu g/m^3$	Both	Both		1.073 (1.0 to 1.236)	1.069 (1.0 to 1.224)	1.066 (1.0 to 1.211)	1.059 (1.0 to 1.193)	1.054 (1.0 to 1.184)	1.051 (1.0 to 1.171)	1.046 (1.0 to 1.15)	1.043 (1.0 to 1.135)	1.039 (1.0 to 1.123)	1.035 (1.0 to 1.11)	1.03 (1.0 to 1.093)	1.025 (1.0 to 1.075)	1.022 (1.0 to 1.07)	1.017 (1.0 to 1.051)	1.013 (1.0 to 1.042)
Stroke	$0\;\mu g/m^3$	Both	Both		1.0 (1.0 to 1.0)														
Chronic obstructive pulmonary disease	$600 \ \mu g/m^3$	Both	Both	2.335 (1.956 to 2.701)															
Chronic obstructive pulmonary disease	$500 \ \mu g/m^3$	Both	Both	2.231 (1.864 to 2.585)															
Chronic obstructive pulmonary disease	$400 \; \mu g/m^3$	Both	Both	2.114 (1.767 to 2.447)															
Chronic obstructive pulmonary disease	$300 \ \mu g/m^3$	Both	Both	1.979 (1.653 to 2.292)															
Chronic obstructive pulmonary disease	$200 \ \mu g/m^3$	Both	Both	1.815 (1.52 to 2.099)															
Chronic obstructive pulmonary disease	150 µg/m ³	Both	Both	1.714 (1.441 to 1.976)															
Chronic obstructive pulmonary disease	$135\;\mu\text{g/m}^3$	Both	Both	1.681 (1.415 to 1.94)															

Appendix Table 6b. Relative risks us	ed by age and	sex for each ou	tcome for	the particulat	e matter integ	rated exposur	e response cu	rve.											
	Category /	Morbidity /						•	•		A	ge	•	•	•				
Risk - Outcome	Units	Mortality	Sex	All ages	25-29 years	30-34 years	35-39 years	40-44 years	45-49 years	50-54 years	55-59 years	60-64 years	65-69 years	70-74 years	75-79 years	80-84 years	85-89 years	90-94 years	95+ years
Ambient particulate matter pollution (PM2.5)																			
Chronic obstructive pulmonary disease	$120\;\mu\text{g/m}^{3}$	Both	Both	1.645 (1.388 to 1.899)															
Chronic obstructive pulmonary disease	$105\;\mu\text{g/m}^3$	Both	Both	1.606 (1.359 to 1.854)															
Chronic obstructive pulmonary disease	90 µg/m ³	Both	Both	1.564 (1.328 to 1.804)															
Chronic obstructive pulmonary disease	$75 \; \mu g/m^3$	Both	Both	1.518 (1.294 to 1.745)															
Chronic obstructive pulmonary disease	$60 \ \mu g/m^3$	Both	Both	1.466 (1.258 to 1.68)															
Chronic obstructive pulmonary disease	45 µg/m ³	Both	Both	1.405 (1.217 to 1.607)															
Chronic obstructive pulmonary disease	$30 \ \mu g/m^3$	Both	Both	1.33 (1.168 to 1.512)															
Chronic obstructive pulmonary disease	25 µg/m³	Both	Both	1.3 (1.149 to 1.473)															
Chronic obstructive pulmonary disease	20 µg/m³	Both	Both	1.266 (1.126 to 1.426)															
Chronic obstructive pulmonary disease	15 µg/m³	Both	Both	1.224 (1.101 to 1.371)															
Chronic obstructive pulmonary disease	10 µg/m ³	Both	Both	1.17 (1.07 to 1.296)															
Chronic obstructive pulmonary disease	$5\ \mu g/m^3$	Both	Both	1.06 (1.0 to 1.17)															
Chronic obstructive pulmonary disease	$0\ \mu g/m^3$	Both	Both	1.0 (1.0 to 1.0)															
Diabetes mellitus type 2	$600 \ \mu g/m^3$	Both	Both	1.448 (1.33 to 1.545)															
Diabetes mellitus type 2	$500 \ \mu g/m^3$	Both	Both	1.447 (1.324 to 1.545)															
Diabetes mellitus type 2	$400 \ \mu g/m^3$	Both	Both	1.446 (1.317 to 1.545)															
Diabetes mellitus type 2	$300 \ \mu g/m^3$	Both	Both	1.445 (1.306 to 1.545)															
Diabetes mellitus type 2	200 µg/m3	Both	Both	1.443 (1.292 to 1.545)															
Diabetes mellitus type 2	150 µg/m3	Both	Both	1.441 (1.281 to 1.545)															
Diabetes mellitus type 2	135 µg/m3	Both	Both	1.44 (1.276 to 1.545)															
Diabetes mellitus type 2	120 µg/m ³	Both	Both	1.439															
Diabetes mellitus type 2	105 µg/m ³	Both	Both	(1.272 to 1.544) 1.438															
Diabetes mellitus type 2	90 µg/m ³	Both	Both	(1.266 to 1.538) 1.436															
Diabetes mellitus type 2	75 µg/m	Both	Both	(1.26 to 1.533) 1.433															
Diabetes mellitus type 2	/5 μg/m	D-th	Deth	(1.253 to 1.53) 1.429															
Diabetes methics type 2	60 μg/m-	Boun	Both	(1.245 to 1.526) 1.422															
Diabetes mellitus type 2	45 μg/m ²	Both	Both	(1.234 to 1.52) 1.405															
Diabetes mellitus type 2	30 µg/m ³	Both	Both	(1.219 to 1.504)															
Diabetes mellitus type 2	25 µg/m3	Both	Both	(1.213 to 1.501)															
Diabetes mellitus type 2	$20 \ \mu\text{g/m}^3$	Both	Both	1.375 (1.197 to 1.494)															
Diabetes mellitus type 2	$15 \ \mu g/m^3$	Both	Both	1.345 (1.177 to 1.489)															
Diabetes mellitus type 2	$10 \ \mu g/m^3$	Both	Both	1.282 (1.116 to 1.466)															
Diabetes mellitus type 2	$5 \ \mu g/m^3$	Both	Both	1.089 (1.0 to 1.346)															
Diabetes mellitus type 2	$0\ \mu g/m^3$	Both	Both	1.0 (1.0 to 1.0)															

Risk - Outcome	Category / Units	Morbidity / Mortality	Sex All ages
Alcohol use			ž
Tukarsulasis	72 g/day	Poth Pot	3.507
Tuberculosis	72 g/uay	BOUIL BOU	2.994
Tuberculosis	60 g/day	Both Bot	h (1.972 to 4.204
Tuberculosis	48 g/day	Both Bot	h (1.701 to 3.51)
Tuberculosis	36 g/day	Both Bot	2.058 h (1.485 to 2.795
Tuboroulogia	24 g/day	Poth Pot	1.531 (1.165 to 1.08)
Tuberculosis	24 g/uay	BOUII BOU	1.101
Tuberculosis	12 g/day	Both Bot	h (0.815 to 1.425
Tuberculosis	0 g/day	Both Bot	h (1.0 to 1.0)
Lower respiratory infections	72 g/day	Both Bot	1.357 h (1.113 to 1.648
Lower recritatory infections	60 g/day	Poth Pot	1.226
	00 g/ uay	Both Bot	1.127
Lower respiratory infections	48 g/day	Both Bot	h (0.936 to 1.327
Lower respiratory infections	36 g/day	Both Bot	h (0.928 to 1.219
Lower respiratory infections	24 g/day	Both Bot	1.026 h (0.901 to 1.167
Lower recritatory infections	12 g/day	Poth Pot	1.013
Lower respiratory intections	IZ B/ngà	BOUII BOU	1.0
Lower respiratory infections	0 g/day	Both Bot	h (1.0 to 1.0)
Oesophageal cancer	72 g/day	Both Bot	h (2.074 to 3.348
Oesophageal cancer	60 g/day	Both Bot	2.452 h (1.905 to 3.094
Oesophageal cancer	48 g/dav	Both Bot	2.202 h (1.73 to 2.703)
	40 b) ddy		1.815
Oesophageal cancer	36 g/day	Both Bot	h (1.468 to 2.222 1.466
Oesophageal cancer	24 g/day	Both Bot	h (1.209 to 1.764
Oesophageal cancer	12 g/day	Both Bot	h (1.031 to 1.439
Oesophageal cancer	0g/dav	Both Bot	1.0 h (1.0 to 1.0)
	- ()		1.424
Liver cancer due to alcohol use	72 g/day	Both Bot	h (1.088 to 1.855 1.372
Liver cancer due to alcohol use	60 g/day	Both Bot	h (1.093 to 1.692
Liver cancer due to alcohol use	48 g/day	Both Bot	h (1.036 to 1.639
Liver cancer due to alcohol use	36 g/day	Both Bot	1.225 h (1.009 to 1.455
			1.14
Liver cancer due to alconol use	24 g/day	Both Bot	n (0.934 to 1.359 1.067
Liver cancer due to alcohol use	12 g/day	Both Bot	h (0.936 to 1.207
Liver cancer due to alcohol use	0 g/day	Both Bot	h (1.0 to 1.0)
Larynx cancer	72 g/day	Both Bot	2.461 h (1.758 to 3.228
	co «/d	Doth Do	2.144
Larynx cancer	60 g/day	воtn Bot	n (1.46 to 2.935) 1.813
Larynx cancer	48 g/day	Both Bot	h (1.3 to 2.421)

Bisk Outcome	Catagomy / Units	Manhidity / Mantality	
Kisk - Outcome	Category / Units	Morbidity / Mortality	Sex All ages
			1 521
Larynx cancer	36 g/day	Both Bot	h (1.126 to 2.061)
			1.304
Larynx cancer	24 g/day	Both Bot	h (1.006 to 1.659)
Larynx cancer	12 g/day	Both Bot	h (0.903 to 1.386)
			1.0
Larynx cancer	0 g/day	Both Bot	h (1.0 to 1.0) 1 476
Breast cancer	72 g/day	Both Bot	h (1.282 to 1.691)
	aa ()		1.452
Breast cancer	60 g/day	Both Bot	n (1.312 to 1.599) 1 443
Breast cancer	48 g/day	Both Bot	h (1.348 to 1.542)
D		D-th D-t	1.433
Breast cancer	36 g/day	Both Bot	n (1.311 to 1.551) 1.329
Breast cancer	24 g/day	Both Bot	h (1.237 to 1.419)
Proact concor	12 g/day	Roth Rot	1.17 (1.081 to 1.265)
Diedst Califer	12 g/uay	ВОШ ВОГ	1.0
Breast cancer	0 g/day	Both Bot	h (1.0 to 1.0)
Colon and rectum cancer	72 g/day	Both Bot	1.616 h (1.38 to 1.861)
	, 2 6, ady		1.468
Colon and rectum cancer	60 g/day	Both Bot	h (1.329 to 1.615)
Colon and rectum cancer	48 g/day	Both Bot	1.323 h (1.156 to 1.501)
			1.237
Colon and rectum cancer	36 g/day	Both Bot	h (1.148 to 1.336)
Colon and rectum cancer	24 g/day	Both Bot	h (1.067 to 1.248)
			1.078
Colon and rectum cancer	12 g/day	Both Bot	h (1.034 to 1.124)
Colon and rectum cancer	0 g/day	Both Bot	h (1.0 to 1.0)
	70 ()		4.858
Lip and oral cavity cancer	/2 g/day	Both Bot	n (3.74 to 6.076) 3.766
Lip and oral cavity cancer	60 g/day	Both Bot	h (2.839 to 4.9)
Lin and eral cavity cancer	48 g/day	Poth Pot	2.991 (2.282 to 2.896)
	40 g/uay	Both Bot	2.311
Lip and oral cavity cancer	36 g/day	Both Bot	h (1.757 to 2.929)
Lip and oral cavity cancer	24 g/day	Both Bot	1.738 h (1.383 to 2.161)
	246/009		1.293
Lip and oral cavity cancer	12 g/day	Both Bot	h (1.076 to 1.551)
Lip and oral cavity cancer	0 g/day	Both Bot	h (1.0 to 1.0)
	0. 7		4.545
Nasopharynx cancer	72 g/day	Both Bot	h (4.1 to 4.982)
Nasopharynx cancer	60 g/day	Both Bot	h (3.509 to 4.102)
			3.062
Nasopharynx cancer	48 g/day	Both Bot	h (2.873 to 3.258) געב ג
Nasopharynx cancer	36 g/day	Both Bot	h (2.25 to 2.552)
Nacarbarran	24 - /	Dath - ·	1.839
Nasopharynx cancer	24 g/day	вотп Bot	n (1.// to 1.907) 1.371
Nasopharynx cancer	12 g/day	Both Bot	h (1.341 to 1.398)

Risk - Outcome	Category / Units	Morbidity / Mortality	Sex	All ages
Alcohol use	Cutegory / Cints	Norblang / Nortanty		Ann uges
				1.0
Nasopharynx cancer	0 g/day	Both	Both	(1.0 to 1.0)
Other pharvny cancer	72 g/day	Both	Both	4.764 (3 315 to 6 576)
	72 B/ 00 y	both	both	3.972
Other pharynx cancer	60 g/day	Both	Both	(2.813 to 5.354)
Other pharynx cancer	48 g/day	Both	Both	3.199 (2.202 to 4.407)
		2011	2000	2.519
Other pharynx cancer	36 g/day	Both	Both	(1.843 to 3.299)
Other pharynx cancer	24 g/day	Both	Both	1.943 (1.467 to 2.484)
				1.472
Other pharynx cancer	12 g/day	Both	Both	(1.234 to 1.742)
Other pharynx cancer	0 g/day	Both	Both	(1.0 to 1.0)
	72 - / -!	Deth	N 4 - 1 -	1.091
ischaemic heart disease	72 g/day	Both	Male	0.993
Ischaemic heart disease	60 g/day	Both	Male	(0.883 to 1.105)
Ischaomic boart discase	48 g/day	Poth	Malo	0.906 (0.797 to 1.035)
	40 g/ uay	both	Wate	0.871
Ischaemic heart disease	36 g/day	Both	Male	(0.788 to 0.964)
lschaemic heart disease	24 g/day	Both	Male	(0.779 to 0.943)
				0.865
Ischaemic heart disease	12 g/day	Both	Male	(0.79 to 0.948) 1 0
Ischaemic heart disease	0 g/day	Both	Male	(1.0 to 1.0)
	72 a/day	Dath	Famala	1.107
	72 g/uay	BOUI	remaie	(0.894 (0 1.541) 1.012
Ischaemic heart disease	60 g/day	Both	Female	(0.869 to 1.174)
Ischaemic heart disease	48 g/day	Both	Female	0.932 (0.786 to 1.113)
				0.882
Ischaemic heart disease	36 g/day	Both	Female	(0.781 to 0.997)
Ischaemic heart disease	24 g/day	Both	Female	(0.749 to 0.948)
	42 //			0.823
ischaemic heart disease	12 g/day	Both	Female	(0.733 to 0.926) 1.0
Ischaemic heart disease	0 g/day	Both	Female	(1.0 to 1.0)
Ischaemic stroke	72 g/day	Both	Male	1.451 (1.228 to 1.69)
	, 2 8, 000	2011		1.312
Ischaemic stroke	60 g/day	Both	Male	(1.167 to 1.471)
Ischaemic stroke	48 g/day	Both	Male	(0.98 to 1.353)
	aa ()			1.057
Ischaemic stroke	36 g/day	Both	Male	(0.931 to 1.192) 0.97
Ischaemic stroke	24 g/day	Both	Male	(0.862 to 1.088)
Ischaemic stroke	12 g/day	Both	Male	0.938 (0.83 to 1.054)
	12 5/009	both	Whate	1.0
Ischaemic stroke	0 g/day	Both	Male	(1.0 to 1.0)
lschaemic stroke	72 g/day	Both	Female	1.43 (1.147 to 1.771)
				1.3
Ischaemic stroke	60 g/day	Both	Female	(1.121 to 1.496)

Risk - Outcome	Category / Units	Morbidity / Mortality	Sex	All ages
Alcohol use	Cuttgory / Chits	inter blancy / inter tailing		7 in ages
				1.145
Ischaemic stroke	48 g/day	Both	Female	(0.946 to 1.359)
Ischaemic stroke	36 g/day	Both	Female	(0.834 to 1.149)
Ischaemic stroke	24 g/day	Both	Female	0.85 (0.726 to 0.985)
	24 6/ 00 y	both	Ternale	0.824
Ischaemic stroke	12 g/day	Both	Female	(0.718 to 0.939)
Ischaemic stroke	0 g/day	Both	Female	(1.0 to 1.0)
Intracorobral homorrhage	72 g/day	Path	Mala	1.971 (1.662 to 2.216)
intracerebra nemornage	72 g/uay	both	Wale	1.705
Intracerebral hemorrhage	60 g/day	Both	Male	(1.45 to 1.991)
Intracerebral hemorrhage	48 g/day	Both	Male	(1.182 to 1.768)
		Dath	Mala	1.31 (1.105 to 1.530)
intracerebrai nemormage	36 g/uay	BOLI	Male	1.162
Intracerebral hemorrhage	24 g/day	Both	Male	(0.973 to 1.385)
Intracerebral hemorrhage	12 g/day	Both	Male	1.068 (0.945 to 1.214)
				1.0
Intracerebral hemorrhage	0 g/day	Both	Male	(1.0 to 1.0) 2.276
Intracerebral hemorrhage	72 g/day	Both	Female	(1.701 to 2.934)
Intracerebral hemorrhage	60 g/dav	Both	Female	1.964 (1.536 to 2.464)
				1.614
Intracerebral hemorrhage	48 g/day	Both	Female	(1.245 to 2.048) 1 337
Intracerebral hemorrhage	36 g/day	Both	Female	(1.065 to 1.664)
Intracerebral bemorrhage	24 g/day	Both	Female	1.11 (0.884 to 1.367)
	246/009	both	- Cillaic	1.031
Intracerebral hemorrhage	12 g/day	Both	Female	(0.897 to 1.18)
Intracerebral hemorrhage	0 g/day	Both	Female	(1.0 to 1.0)
Hypertensive beart disease	72 g/day	Both	Both	1.86 (1.445 to 2.358)
Hypertensive heart disease	72 g/udy	both	both	1.705
Hypertensive heart disease	60 g/day	Both	Both	(1.297 to 2.175)
Hypertensive heart disease	48 g/day	Both	Both	(1.25 to 2.049)
Hypertoneive boart disease	26 g/day	Path	Poth	1.479 (1.222 to 1.759)
Hypertensive neart disease	36 g/uay	BOLI	Both	1.315
Hypertensive heart disease	24 g/day	Both	Both	(1.136 to 1.526)
Hypertensive heart disease	12 g/day	Both	Both	(0.913 to 1.198)
	o. / I			1.0
Hypertensive heart disease	U g/day	Both	Both	(1.0 to 1.0) 1.535
Atrial fibrillation and flutter	72 g/day	Both	Both	(1.348 to 1.728)
Atrial fibrillation and flutter	60 g/day	Both	Both	1.411 (1.26 to 1.569)
				1.312
Atrial fibrillation and flutter	48 g/day	Both	Both	(1.218 to 1.407) 1.214
Atrial fibrillation and flutter	36 g/day	Both	Both	(1.145 to 1.29)
Atrial fibrillation and flutter	24 g/dav	Both	Both	1.131 (1.067 to 1.204)

Risk - Outcome	Category / Units	Morbidity / Mortality	Sex	All ages
lcohol use				
Atrial fibrillation and flutter	12 g/day	Both	Both	1.066 (1.034 to 1.102)
Atrial fibrillation and flutter	0 g/day	Both	Both	1.0 (1.0 to 1.0)
Cirrhosis and other chronic liver diseases due to alcohol use	72 g/day	Both	Both	9.427 (6.131 to 13.804
Cirrhosis and other chronic liver diseases due to alcohol use	60 g/day	Both	Both	6.274 (3.958 to 9.319)
Cirrhosis and other chronic liver diseases due to alcohol use	48 g/day	Both	Both	4.673 (3.25 to 6.717)
Cirrhosis and other chronic liver diseases due to alcohol use	36 g/day	Both	Both	(2.309 to 4.485)
Cirrhosis and other chronic liver diseases due to alcohol use	24 g/day	Both	Both	(1.521 to 2.688)
Cirrhosis and other chronic liver diseases due to alcohol use	12 g/day	Both	Both	(0.943 to 1.611)
Cirrhosis and other chronic liver diseases due to alcohol use	0 g/day	Both	Both	(1.0 to 1.0)
Pancreatitis	72 g/day	Both	Both	(2.473 to 4.458)
Pancreatitis	60 g/day	Both	Both	(1.415 to 3.389)
Pancreatitis	48 g/day	Both	Both	(1.199 to 2.477)
Pancreatitis	36 g/day	Both	Both	(1.062 to 2.021)
Pancreatitis	24 g/day	Both	Both	(0.874 to 1.67)
Pancreatitis	12 g/day	Both	Both	(0.791 to 1.481
Pancreatitis	0 g/day	Both	Both	(1.0 to 1.0) 2 48
Epilepsy	72 g/day	Both	Both	(1.929 to 3.144 2 186
Epilepsy	60 g/day	Both	Both	(1.781 to 2.622
Epilepsy	48 g/day	Both	Both	(1.438 to 2.369
Epilepsy	36 g/day	Both	Both	(1.303 to 1.898 1.353
Epilepsy	24 g/day	Both	Both	(1.118 to 1.633 1.177
Epilepsy	12 g/day	Both	Both	(1.059 to 1.316 1.0
Epilepsy	0 g/day	Both	Both	(1.0 to 1.0) 1.198
Diabetes mellitus	72 g/day	Both	Male	(1.065 to 1.337 1.165
Diabetes mellitus	60 g/day	Both	Male	(0.998 to 1.342 1.084
Diabetes mellitus	48 g/day	Both	Male	(0.933 to 1.239 1.0
Diabetes mellitus	36 g/day	Both	Male	(0.891 to 1.119 0.932
Diabetes mellitus	24 g/day	Both	Male	(0.841 to 1.03) 0.921
Diabetes mellitus	12 g/day	Both	Male	(0.833 to 1.015 1.0
Diabetes mellitus	0 g/day	Both	Male	(1.0 to 1.0) 1.172
Diabetes mellitus	72 g/day	Both	Female	(0.81 to 1.652)

Risk - Outcome	Category / Units	Morbidity / Mortality	Sex	All ages
Alcohol use	Cutegory / Chits	inor blatty / inor talley		in uges
				1.074
Diabetes mellitus	60 g/day	Both F	emale	(0.765 to 1.443)
Diabetes mellitus	48 g/day	Both F	emale	0.945 (0.737 to 1.173)
				0.836
Diabetes mellitus	36 g/day	Both F	emale	(0.702 to 0.981) 0.76
Diabetes mellitus	24 g/day	Both F	emale	(0.66 to 0.872)
Diabetes mellitus	12 g/day	Both F	emale	0.733 (0.658 to 0.826)
	22.8/ 004			1.0
Diabetes mellitus	0 g/day	Both F	emale	(1.0 to 1.0)
Transport injuries	72 g/day	Both B	oth	(1.201 to 2.032)
				1.456
Transport injuries	60 g/day	Both B	oth	(1.186 to 1.818) 1.366
Transport injuries	48 g/day	Both B	oth	(1.101 to 1.692)
Transport injuries	36 g/dav	Both B	oth	1.288 (1.089 to 1.534)
	00 B/ 44 P	5000		1.22
Transport injuries	24 g/day	Both B	oth	(1.062 to 1.4)
Transport injuries	12 g/day	Both B	oth	(1.021 to 1.346)
Transportisiuris		Dath D		1.0 (1.0 to 1.0)
i ransport injuries	0 g/day	BOLU B	oth	(1.0 to 1.0) 1.266
Unintentional injuries	72 g/day	Both B	oth	(1.063 to 1.555)
Unintentional injuries	60 g/dav	Both B	oth	1.221 (1.059 to 1.46)
	00 8/ 44/	5000		1.182
Unintentional injuries	48 g/day	Both B	oth	(1.024 to 1.428)
Unintentional injuries	36 g/day	Both B	oth	(1.054 to 1.347)
	24 - / /	D-th D	- 4 4	1.154
Unintentional injuries	24 g/day	BOLU B	oth	(1.046 (0 1.319)
Unintentional injuries	12 g/day	Both B	oth	(1.016 to 1.187)
Unintentional iniuries	0 g/dav	Both B	oth	1.0 (1.0 to 1.0)
	- 8, ,			1.927
Self-harm	72 g/day	Both B	oth	(1.398 to 2.665) 1 734
Self-harm	60 g/day	Both B	oth	(1.29 to 2.308)
Calf have	40 a /day	Dath D		1.545
Seit-narm	48 g/day	BOLU B	oth	(1.132 to 2.048) 1.376
Self-harm	36 g/day	Both B	oth	(1.05 to 1.751)
Self-harm	24 g/dav	Both B	oth	1.23 (0.972 to 1.533)
	0. 7			1.107
Self-harm	12 g/day	Both B	oth	(0.908 to 1.343)
Self-harm	0 g/day	Both B	oth	(1.0 to 1.0)
	72 a /day	Dath D		1.516
Interpersonal violence	72 g/uay	BOLII B	oth	1.452
Interpersonal violence	60 g/day	Both B	oth	(1.215 to 1.719)
Interpersonal violence	48 g/dav	Both B	oth	1.396 (1.118 to 1.739)
	- 0/ /	_		1.345
Interpersonal violence	36 g/day	Both B	oth	(1.14 to 1.585)

Appendix Table 6c. Relative risks used by age and sex for each outcome for alcohol use globally. **Risk - Outcome** Category / Units Morbidity / Mortality Sex All ages Alcohol use 1.256 (1.055 to 1.46) Interpersonal violence 24 g/day Both Both 1.129 (0.963 to 1.317) Interpersonal violence 12 g/day Both Both 1.0 Interpersonal violence 0 g/day Both Both (1.0 to 1.0)

Appendix Table 6d. Relative risks used by age and sex for each of																	
Risk - Outcome	Category / Units	Morbidity / Mortality	Sex	30 to 34	35 to 39	40 to 44	45 to 49	50 to 54	55 to 59	A 60 to 64	.ge 65 to 69	70 to 74	75 to 79	80 to 84	95 to 90	90 to 94	95 plur
Smoking				30 10 34	351039	401044	45 10 49	50 to 54	5510 59	60 10 64	05 10 09	70 to 74	751079	80 10 84	85 10 89	50 (0 54	95 pius
				1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
Latent tuberculosis infection	0 Cigarettes Per Smoker Per Day	Both	Male	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)						
Latent tuberculosis infection	0 Cigarettes Per Smoker Per Day	Both	Female	(1.0 to 1.0) 2.073	(1.0 to 1.0) 2.073	(1.0 to 1.0) 2.073	(1.0 to 1.0) 2.073	(1.0 to 1.0) 2.073	(1.0 to 1.0) 2.073	(1.0 to 1.0) 2.073	(1.0 to 1.0) 2.073						
Latent tuberculosis infection	9 Cigarettes Per Smoker Per Day	Both	Male	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)						
Latent tuberculosis infection	9 Cigarettes Per Smoker Per Day	Both	Female	(1.605 to 2.549)	(1.605 to 2.549) 2 311	(1.605 to 2.549)	(1.605 to 2.549) 2 311	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549) 2 311	(1.605 to 2.549)	(1.605 to 2.549) 2 311	(1.605 to 2.549)				
Latent tuberculosis infection	18 Cigarettes Per Smoker Per Day	Both	Male	(1.808 to 2.921) 2 311	(1.808 to 2.921) 2 311	(1.808 to 2.921)	(1.808 to 2.921) 2 311	(1.808 to 2.921) 2 311	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921) 2 311	(1.808 to 2.921)	(1.808 to 2.921) 2 311	(1.808 to 2.921) 2 311
Latent tuberculosis infection	18 Cigarettes Per Smoker Per Day	Both	Female	(1.808 to 2.921) 3.029	(1.808 to 2.921) 3.029	(1.808 to 2.921)	(1.808 to 2.921) 3 029	(1.808 to 2.921) 3.029	(1.808 to 2.921)) (1.808 to 2.921) 3 029	(1.808 to 2.921) 3.029	(1.808 to 2.921) 3.029	(1.808 to 2.921) 3.029	(1.808 to 2.921) 3 029	(1.808 to 2.921) 3.029	(1.808 to 2.921) 3.029	(1.808 to 2.921) 3.029
Latent tuberculosis infection	27 Cigarettes Per Smoker Per Day	Both	Male	(2.315 to 3.969) 3.029	(2.315 to 3.969) 3.029	(2.315 to 3.969) 3.029	(2.315 to 3.969) 3.029	(2.315 to 3.969) 3.029	(2.315 to 3.969) 3.029	(2.315 to 3.969) 3.029	(2.315 to 3.969) 3.029						
Latent tuberculosis infection	27 Cigarettes Per Smoker Per Day	Both	Female	(2.315 to 3.969) 4.013	(2.315 to 3.969) 4.013	(2.315 to 3.969) 4.013	(2.315 to 3.969) 4.013	(2.315 to 3.969) 4.013	(2.315 to 3.969) 4.013	(2.315 to 3.969) 4.013	(2.315 to 3.969) 4.013						
Latent tuberculosis infection	36 Cigarettes Per Smoker Per Day	Both	Male	(2.49 to 5.991) 4.013	(2.49 to 5.991) 4.013	(2.49 to 5.991) 4.013	(2.49 to 5.991) 4.013	(2.49 to 5.991) 4.013	(2.49 to 5.991) 4.013	(2.49 to 5.991) 4.013	(2.49 to 5.991) 4.013						
Latent tuberculosis infection	36 Cigarettes Per Smoker Per Day	Both	Female	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)						
Drug-susceptible tuberculosis	0 Cigarettes Per Smoker Per Day	Both	Male	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)						
Drug-susceptible tuberculosis	0 Cigarettes Per Smoker Per Day	Both	Female	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)						
Drug-susceptible tuberculosis	9 Cigarettes Per Smoker Per Day	Both	Male	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)						
Drug-susceptible tuberculosis	9 Cigarettes Per Smoker Per Day	Both	Female	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)						
Drug-susceptible tuberculosis	18 Cigarettes Per Smoker Per Day	Both	Male	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)						
Drug-susceptible tuberculosis	18 Cigarettes Per Smoker Per Day	Both	Female	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)						
Drug-susceptible tuberculosis	27 Cigarettes Per Smoker Per Day	Both	Male	(2.315 to 3.969)) (2.315 to 3.969)	(2.315 to 3.969)											
Drug-susceptible tuberculosis	27 Cigarettes Per Smoker Per Day	Both	Female	(2.315 to 3.969)	(2.315 to 3.969)	(2.315 to 3.969)	(2.315 to 3.969)	(2.315 to 3.969)	(2.315 to 3.969)	(2.315 to 3.969)	(2.315 to 3.969)						
Drug-susceptible tuberculosis	36 Cigarettes Per Smoker Per Day	Both	Male	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	4.015 (2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)
Drug-susceptible tuberculosis	36 Cigarettes Per Smoker Per Day	Both	Female	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)						
Multidrug-resistant tuberculosis without extensive drug resistance	0 Cigarettes Per Smoker Per Day	Both	Male	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)						
Multidrug-resistant tuberculosis without extensive drug resistance	0 Cigarettes Per Smoker Per Day	Both	Female	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)						
Multidrug-resistant tuberculosis without extensive drug resistance	9 Cigarettes Per Smoker Per Day	Both	Male	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)						
Multidrug-resistant tuberculosis without extensive drug resistance	9 Cigarettes Per Smoker Per Day	Both	Female	(1.605 to 2.549)	2.073) (1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)					
Multidrug-resistant tuberculosis without extensive drug resistance	18 Cigarettes Per Smoker Per Day	Both	Male	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)						
Multidrug-resistant tuberculosis without extensive drug resistance	18 Cigarettes Per Smoker Per Day	Both	Female	(1.808 to 2.921)) (1.808 to 2.921)	(1.808 to 2.921)											
Multidrug-resistant tuberculosis without extensive drug resistance	27 Cigarettes Per Smoker Per Day	Both	Male	(2.315 to 3.969)) (2.315 to 3.969)	(2.315 to 3.969)											
Multidrug-resistant tuberculosis without extensive drug resistance	27 Cigarettes Per Smoker Per Day	Both	Female	(2.315 to 3.969)	(2.315 to 3.969)	(2.315 to 3.969)	(2.315 to 3.969)	(2.315 to 3.969)	(2.315 to 3.969)	(2.315 to 3.969)	(2.315 to 3.969)						
Multidrug-resistant tuberculosis without extensive drug resistance	36 Cigarettes Per Smoker Per Day	Both	Male	(2.49 to 5.991) 4 013	(2.49 to 5.991)	(2.49 to 5.991) 4 013											
Multidrug-resistant tuberculosis without extensive drug resistance	36 Cigarettes Per Smoker Per Day	Both	Female	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)	(2.49 to 5.991)						
Extensively drug-resistant tuberculosis	0 Cigarettes Per Smoker Per Day	Both	Male	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)						
Extensively drug-resistant tuberculosis	0 Cigarettes Per Smoker Per Day	Both	Female	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)						
Extensively drug-resistant tuberculosis	9 Cigarettes Per Smoker Per Day	Both	Male	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)						
Extensively drug-resistant tuberculosis	9 Cigarettes Per Smoker Per Day	Both	Female	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)	(1.605 to 2.549)						
Extensively drug-resistant tuberculosis	18 Cigarettes Per Smoker Per Day	Both	Male	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)						
Extensively drug-resistant tuberculosis	18 Cigarettes Per Smoker Per Day	Both	Female	(1.808 to 2.921)	(1.808 to 2.921) 3 029	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921) 3.029	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921) 3 029	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921)	(1.808 to 2.921) 3 029
Extensively drug-resistant tuberculosis	27 Cigarettes Per Smoker Per Day	Both	Male	(2.315 to 3.969)) (2.315 to 3.969)	(2.315 to 3.969)											

Appendix Table 6d. Relative risks used by age and sex for each ou	tcome for smoking globally.																
Risk - Outcome	Category / Units	Morbidity / Mortality	Sex	30 to 34	35 to 39	40 to 44	45 to 49	50 to 54	55 to 59	A:	ge 65 to 69	70 to 74	75 to 79	80 to 84	85 to 89	90 to 94	95 plus
Smoking																	
Extensively drug-resistant tuberculosis	27 Cigarettes Per Smoker Per Day	Both	Female	3.029 (2.315 to 3.969)													
Extensively drug-resistant tuberculosis	36 Cigarettes Per Smoker Per Day	Both	Male	4.013 (2.49 to 5.991)													
Extensively drug-resistant tuberculosis	36 Cigarettes Per Smoker Per Day	Both	Female	4.013 (2.49 to 5.991)													
Lower respiratory infections	0 Cigarettes Per Smoker Per Day	Both	Male	1.0 (1.0 to 1.0)													
Lower respiratory infections	0 Cigarettes Per Smoker Per Day	Both	Female	(1.0 to 1.0)													
Lower respiratory infections	1.5 Cigarettes Per Smoker Per Day	Both	Male	(1.0 to 1.785)													
Lower respiratory infections	1.5 Cigarettes Per Smoker Per Day	Both	Female	(1.0 to 1.785)													
Lower respiratory infections	10 Cigarettes Per Smoker Per Day	Both	Male	(1.356 to 2.731) 1.958													
Lower respiratory infections	10 Cigarettes Per Smoker Per Day	Both	Female	(1.356 to 2.731) 2.434													
Lower respiratory infections	19.5 Cigarettes Per Smoker Per Day	Both	Male	(1.738 to 3.307) 2.434													
Lower respiratory infections	19.5 Cigarettes Per Smoker Per Day	Both	Female	(1.738 to 3.307) 3.158													
Lower respiratory infections	31.2 Cigarettes Per Smoker Per Day	Both	Male	(2.286 to 4.195) 3.158													
Lower respiratory infections	31.2 Cigarettes Per Smoker Per Day	Both	Female	(2.286 to 4.195) 1.0													
Lip and oral cavity cancer	0 Pack-Years	Both	Male	(1.0 to 1.0) 1.0													
Lip and oral cavity cancer	0 Pack-Years	Both	Female	(1.0 to 1.0) 3.19													
Lip and oral cavity cancer	17.7 Pack-Years	Both	Male	(2.162 to 4.362) 3.19													
Lip and oral cavity cancer	17.7 Pack-Years	Both	Female	(2.162 to 4.362) 3.754													
Lip and oral cavity cancer	35.4 Pack-Years	Both	Male	(2.335 to 5.44) 3.754													
Lip and oral cavity cancer	35.4 Pack-Years	Both	Female	(2.335 to 5.44) 4.017													
Lip and oral cavity cancer	53.1 Pack-Years	Both	Male	(2.376 to 6.004) 4.017													
Lip and oral cavity cancer	53.1 Pack-Years	Both	Female	(2.376 to 6.004) 5.517													
Lip and oral cavity cancer	70.8 Pack-Years	Both	Male	(2.574 to 9.804) 5.517													
Lip and oral cavity cancer	70.8 Pack-Years	Both	Female	(2.574 to 9.804) 7.27													
Lip and oral cavity cancer	88.5 Pack-Years	Both	Male	(2.599 to 14.741)													
				7.27 (2.599 to													
Lip and oral cavity cancer	88.5 Pack-Years	Both	Female	14.741) 1.0													
Nasopharynx cancer	0 Pack-Years	Both	Male	(1.0 to 1.0) 1.0													
Nasopharynx cancer	0 Pack-Years	Both	Female	(1.0 to 1.0) 1.879													
Nasopharynx cancer	10.7 Pack-Years	Both	Male	(1.34 to 2.522) 1.879													
Nasopharynx cancer	10.7 Pack-Years	Both	Female	(1.34 to 2.522) 1.992													
Nasopharynx cancer	21.4 Pack-Years	Both	Male	(1.322 to 2.865) 1.992													
Nasopharynx cancer	21.4 Pack-Years	Both	Female	(1.322 to 2.865) 2.297													
Nasopharynx cancer	32.1 Pack-Years	Both	Male	(1.448 to 3.32) 2.297													
Nasopharynx cancer	32.1 Pack-Years	Both	Female	(1.448 to 3.32) 2.662													
Nasopharynx cancer	42.9 Pack-Years	Both	Male	(1.646 to 4.007) 2.662													
Nasopharynx cancer	42.9 Pack-Years	Both	Female	(1.646 to 4.007) 3.377													
Nasopharynx cancer	53.6 Pack-Years	Both	Male	(2.01 to 5.041)													

Appendix Table 6d. Relative risks used by age and sex for each ou	tcome for smoking globally.																
Risk - Outcome	Category / Units	Morbidity / Mortality	Sex	30 to 34	35 to 39	40 to 44	45 to 49	50 to 54	55 to 59	A 60 to 64	.ge 65 to 69	70 to 74	75 to 79	80 to 84	85 to 89	90 to 94	95 plus
Smoking																	
Nasopharynx cancer	53.6 Pack-Years	Both	Female	3.377 (2.01 to 5.041)													
Nasopharynx cancer	64.3 Pack-Years	Both	Male	4.466 (2.643 to 6.831)													
Nasopharynx cancer	64.3 Pack-Years	Both	Female	4.466 (2.643 to 6.831)													
Other pharynx cancer	0 Pack-Years	Both	Male	1.0 (1.0 to 1.0)													
Other pharvnx cancer	0 Pack-Years	Both	Female	1.0 (1.0 to 1.0)													
Other pharvny cancer	17 7 Pack-Years	Both	Male	4.792 (3.267 to 6.498)													
	17.7 Pack Years	Both	Fomalo	4.792	4.792	4.792	4.792	4.792	4.792	4.792	4.792	4.792	4.792	4.792	4.792	4.792	4.792
		Buth	remaie	5.255	5.255	5.255	5.255	5.255	5.255	5.255	5.255	5.255	5.255	5.255	5.255	5.255	5.255
Other pharynx cancer	35.4 Pack-Years	Both	Male	(3.541 to 7.384) 5.255													
Other pharynx cancer	35.4 Pack-Years	Both	Female	(3.541 to 7.384) 6.588													
Other pharynx cancer	53.1 Pack-Years	Both	Male	(4.406 to 8.892) 6.588													
Other pharynx cancer	53.1 Pack-Years	Both	Female	(4.406 to 8.892) 7.21													
Other pharynx cancer	70.8 Pack-Years	Both	Male	(3.973 to 11.61) 7.21													
Other pharynx cancer	70.8 Pack-Years	Both	Female	(3.973 to 11.61) 11.149													
Other pharynx cancer	88.5 Pack-Years	Both	Male	(5.54 to 18.571) 11.149													
Other pharynx cancer	88.5 Pack-Years	Both	Female	(5.54 to 18.571)													
Oesophageal cancer	0 Pack-Years	Both	Male	(1.0 to 1.0)													
Oesophageal cancer	0 Pack-Years	Both	Female	(1.0 to 1.0)													
Oesophageal cancer	15 Pack-Years	Both	Male	(2.318 to 3.817)	3.038 (2.318 to 3.817)	3.038 (2.318 to 3.817)	(2.318 to 3.817)	3.038 (2.318 to 3.817)	3.038 (2.318 to 3.817)	3.038 (2.318 to 3.817)	3.038 (2.318 to 3.817)	3.038 (2.318 to 3.817)	(2.318 to 3.817)	3.038 (2.318 to 3.817)	3.038 (2.318 to 3.817)	3.038 (2.318 to 3.817)	(2.318 to 3.817)
Oesophageal cancer	15 Pack-Years	Both	Female	3.038 (2.318 to 3.817)													
Oesophageal cancer	30 Pack-Years	Both	Male	2.778 (1.884 to 3.796)													
Oesophageal cancer	30 Pack-Years	Both	Female	2.778 (1.884 to 3.796)													
Oesophageal cancer	45 Pack-Years	Both	Male	4.471 (3.099 to 5.83)													
Oesophageal cancer	45 Pack-Years	Both	Female	4.471 (3.099 to 5.83)													
Oesophageal cancer	60 Pack-Years	Both	Male	5.327 (2.865 to 8.227)													
Qesophageal cancer	60 Pack-Years	Both	Female	5.327 (2.865 to 8.227)													
				7.326 (3.276 to													
Oesophageal cancer	75 Pack-Years	Both	Male	12.451)	12.451)	12.451)	12.451)	12.451)	12.451)	12.451)	12.451)	12.451)	12.451)	12.451)	12.451)	12.451)	12.451)
0	75 Deals Verse	Deth	Famela	(3.276 to													
	75 Pack-rears	Both	Female	9.593	9.593	9.593	9.593	9.593	9.593	9.593	9.593	9.593	9.593	9.593	9.593	9.593	9.593
Oesopnageal cancer	90 Pack-Years	Both	Male .	(4.866 to 15.62) 9.593	9.593	(4.866 to 15.62) 9.593											
Oesophageal cancer	90 Pack-Years	Both	Female	(4.866 to 15.62) 1.0													
Stomach cancer	0 Pack-Years	Both	Male	(1.0 to 1.0) 1.0													
Stomach cancer	0 Pack-Years	Both	Female	(1.0 to 1.0) 1.427													
Stomach cancer	12.5 Pack-Years	Both	Male	(1.184 to 1.713) 1.427													
Stomach cancer	12.5 Pack-Years	Both	Female	(1.184 to 1.713) 1.607													
Stomach cancer	25 Pack-Years	Both	Male	(1.286 to 1.991) 1.607													
Stomach cancer	25 Pack-Years	Both	Female	(1.286 to 1.991) 1.907	(1.286 to 1.991)	(1.286 to 1.991) 1.907	(1.286 to 1.991)	(1.286 to 1.991)	(1.286 to 1.991) 1.907								
Stomach cancer	37.5 Pack-Years	Both	Male	(1.47 to 2.407)													

Appendix Table 6d. Relative risks used by age and sex for each ou																	
Risk - Outcome	Category / Units	Morbidity / Mortality	Sex	20 4- 24	25 4- 20	40.4- 44	45 4- 40	50 4- 54	55 4- 50	A	ge	70 4- 74	75 4- 70	80.4- 84	95 4- 90	00.4- 0.4	05 alua
Smoking				30 10 34	351039	401044	45 10 49	50 10 54	3510 59	6010 64	05 10 09	70 10 74	/510/9	80 10 84	83 10 87	901094	95 pius
•				1.907	1.907	1.907	1.907	1.907	1.907	1.907	1.907	1.907	1.907	1.907	1.907	1.907	1.907
Stomach cancer	37.5 Pack-Years	Both	Female	(1.47 to 2.407) 2.055													
Stomach cancer	50 Pack-Years	Both	Male	(1.558 to 2.544) 2.055													
Stomach cancer	50 Pack-Years	Both	Female	(1.558 to 2.544)													
Stomach cancer	75 Pack-Years	Both	Male	(1.547 to 2.928)													
Stomach cancer	75 Pack-Years	Both	Female	(1.547 to 2.928) 1.0													
Colon and rectum cancer	0 Pack-Years	Both	Male	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0
Colon and rectum cancer	0 Pack-Years	Both	Female	(1.0 to 1.0) 1.505													
Colon and rectum cancer	18.8 Pack-Years	Both	Male	(1.144 to 1.852) 1.505													
Colon and rectum cancer	18.8 Pack-Years	Both	Female	(1.144 to 1.852) 1.607													
Colon and rectum cancer	37.5 Pack-Years	Both	Male	(1.235 to 2.055) 1.607													
Colon and rectum cancer	37.5 Pack-Years	Both	Female	(1.235 to 2.055) 1.583													
Colon and rectum cancer	56.2 Pack-Years	Both	Male	(1.155 to 2.103) 1.583													
Colon and rectum cancer	56.2 Pack-Years	Both	Female	(1.155 to 2.103) 1.0													
Liver cancer due to hepatitis B	0 Pack-Years	Both	Male	(1.0 to 1.0) 1.0													
Liver cancer due to hepatitis B	0 Pack-Years	Both	Female	(1.0 to 1.0) 1.512													
Liver cancer due to hepatitis B	14.6 Pack-Years	Both	Male	(1.04 to 2.038) 1.512													
Liver cancer due to hepatitis B	14.6 Pack-Years	Both	Female	(1.04 to 2.038) 1.638													
Liver cancer due to hepatitis B	29.2 Pack-Years	Both	Male	(1.099 to 2.256) 1.638	(1.099 to 2.256)	(1.099 to 2.256) 1.638	(1.099 to 2.256) 1.638	(1.099 to 2.256) 1.638									
Liver cancer due to hepatitis B	29.2 Pack-Years	Both	Female	(1.099 to 2.256) 1.83	(1.099 to 2.256) 1.83	(1.099 to 2.256)	(1.099 to 2.256) 1.83	(1.099 to 2.256) 1.83	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256) 1.83	(1.099 to 2.256)	(1.099 to 2.256) 1.83	(1.099 to 2.256) 1.83	(1.099 to 2.256) 1.83	(1.099 to 2.256)
Liver cancer due to hepatitis B	43.8 Pack-Years	Both	Male	(1.224 to 2.606) 1.83													
Liver cancer due to hepatitis B	43.8 Pack-Years	Both	Female	(1.224 to 2.606) 1.786	(1.224 to 2.606)	(1.224 to 2.606) 1.786	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606) 1.786	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606) 1.786	(1.224 to 2.606)	(1.224 to 2.606) 1.786	(1.224 to 2.606)	(1.224 to 2.606)
Liver cancer due to hepatitis B	58.3 Pack-Years	Both	Male	(1.139 to 2.722) 1.786	(1.139 to 2.722) 1.786	(1.139 to 2.722) 1.786	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722) 1.786	(1.139 to 2.722)	(1.139 to 2.722) 1.786	(1.139 to 2.722) 1.786	(1.139 to 2.722)	(1.139 to 2.722) 1.786	(1.139 to 2.722)	(1.139 to 2.722) 1.786
Liver cancer due to hepatitis B	58.3 Pack-Years	Both	Female	(1.139 to 2.722) 1.937													
Liver cancer due to hepatitis B	72.9 Pack-Years	Both	Male	(1.147 to 3.012) 1.937													
Liver cancer due to hepatitis B	72.9 Pack-Years	Both	Female	(1.147 to 3.012) 1.0													
Liver cancer due to hepatitis C	0 Pack-Years	Both	Male	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0) 1.0
Liver cancer due to hepatitis C	0 Pack-Years	Both	Female	(1.0 to 1.0) 1.512	(1.0 to 1.0) 1.512	(1.0 to 1.0) 1.512	(1.0 to 1.0)	(1.0 to 1.0) 1.512	(1.0 to 1.0)	(1.0 to 1.0) 1.512	(1.0 to 1.0) 1.512	(1.0 to 1.0)	(1.0 to 1.0) 1.512	(1.0 to 1.0) 1.512	(1.0 to 1.0) 1.512	(1.0 to 1.0)	(1.0 to 1.0) 1.512
Liver cancer due to hepatitis C	14.6 Pack-Years	Both	Male	(1.04 to 2.038) 1.512													
Liver cancer due to hepatitis C	14.6 Pack-Years	Both	Female	(1.04 to 2.038) 1.638													
Liver cancer due to hepatitis C	29.2 Pack-Years	Both	Male	(1.099 to 2.256) 1.638													
Liver cancer due to hepatitis C	29.2 Pack-Years	Both	Female	(1.099 to 2.256) 1.83													
Liver cancer due to hepatitis C	43.8 Pack-Years	Both	Male	(1.224 to 2.606) 1.83													
Liver cancer due to hepatitis C	43.8 Pack-Years	Both	Female	(1.224 to 2.606) 1.786													
Liver cancer due to hepatitis C	58.3 Pack-Years	Both	Male	(1.139 to 2.722) 1.786													
Liver cancer due to hepatitis C	58.3 Pack-Years	Both	Female	(1.139 to 2.722) 1.937													
Liver cancer due to hepatitis C	72.9 Pack-Years	Both	Male	(1.147 to 3.012) 1.937													
Liver cancer due to hepatitis C	72.9 Pack-Years	Both	Female	(1.147 to 3.012)													

Appendix Table 6d. Relative risks used by age and sex for each ou	tcome for smoking globally.																
Risk - Outcome	Category / Units	Morbidity / Mortality	Sex	20 4- 24	25 4- 20	10 +- 11	45 4- 40	50 4- 54	55 4- 50	A:	ge (5 to (0	70 4- 74	75 4- 70	80.4- 84	95 4- 90	00 4- 0.1	05 -1
Smoking				30 10 34	351039	40 (0 44	43 10 49	50 to 54	5510 59	601064	05 10 09	701074	/5 10 /9	80 10 84	85 10 87	901094	95 pius
				1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
Liver cancer due to alcohol use	0 Pack-Years	Both	Male	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)
Liver cancer due to alcohol use	0 Pack-Years	Both	Female	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)
Liver cancer due to alcohol use	14.6 Pack-Years	Both	Male	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)
Liver cancer due to alcohol use	14.6 Pack-Years	Both	Female	1.512 (1.04 to 2.038)	1.512 (1.04 to 2.038)	1.512 (1.04 to 2.038)	1.512 (1.04 to 2.038)	1.512 (1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	1.512 (1.04 to 2.038)	1.512 (1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	1.512 (1.04 to 2.038)	1.512 (1.04 to 2.038)
Liver cancer due to alcohol use	29.2 Pack-Years	Both	Male	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)
Liver cancer due to alcohol use	29.2 Pack-Years	Both	Female	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)	1.638 (1.099 to 2.256)
Liver cancer due to alcohol use	43.8 Pack-Years	Both	Male	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)
Liver cancer due to alcohol use	43.8 Pack-Years	Both	Female	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)
Liver cancer due to alcohol use	58.3 Pack-Years	Both	Male	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)
Liver cancer due to alcohol use	58.3 Pack-Years	Both	Female	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)
				1.937	1.937	1.937	1.937	1.937	1.937	1.937	1.937	1.937	1.937	1.937	1.937	1.937	1.937
Liver cancer due to alcohol use	72.9 Pack-Years	Both	Male	(1.14/ to 3.012) 1.937	(1.14/ to 3.012) 1.937	(1.14/ to 3.012) 1.937	(1.14 / to 3.012) 1.937	(1.14/ to 3.012) 1.937	(1.14/ to 3.012) 1.937	(1.14/ to 3.012) 1.937	(1.14 / to 3.012) 1.937	(1.147 to 3.012) 1.937	(1.14/ to 3.012) 1.937	(1.14/ to 3.012) 1.937	(1.14 / to 3.012) 1.937	(1.147 to 3.012) 1.937	(1.14/ to 3.012) 1.937
Liver cancer due to alcohol use	72.9 Pack-Years	Both	Female	(1.147 to 3.012) 1.0	(1.147 to 3.012) 1.0	(1.147 to 3.012) 1.0	(1.147 to 3.012) 1.0	(1.147 to 3.012) 1.0	(1.147 to 3.012) 1.0	(1.147 to 3.012) 1.0	(1.147 to 3.012) 1.0	(1.147 to 3.012) 1.0	(1.147 to 3.012) 1.0	(1.147 to 3.012) 1.0	(1.147 to 3.012) 1.0	(1.147 to 3.012) 1.0	(1.147 to 3.012) 1.0
Liver cancer due to NASH	0 Pack-Years	Both	Male	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0
Liver cancer due to NASH	0 Pack-Years	Both	Female	(1.0 to 1.0) 1.512	(1.0 to 1.0) 1.512	(1.0 to 1.0) 1.512	(1.0 to 1.0) 1.512	(1.0 to 1.0) 1.512	(1.0 to 1.0) 1.512	(1.0 to 1.0) 1.512	(1.0 to 1.0) 1.512	(1.0 to 1.0) 1.512	(1.0 to 1.0) 1.512	(1.0 to 1.0) 1.512	(1.0 to 1.0) 1.512	(1.0 to 1.0) 1.512	(1.0 to 1.0) 1.512
Liver cancer due to NASH	14.6 Pack-Years	Both	Male	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)
Liver cancer due to NASH	14.6 Pack-Years	Both	Female	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)	(1.04 to 2.038)
Liver cancer due to NASH	29.2 Pack-Years	Both	Male	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)
Liver cancer due to NASH	29.2 Pack-Years	Both	Female	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)
Liver cancer due to NASH	43.8 Pack-Years	Both	Male	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)
Liver cancer due to NASH	43.8 Pack-Years	Both	Female	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)	1.83 (1.224 to 2.606)
Liver cancer due to NASH	58.3 Pack-Years	Both	Male	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)
Liver cancer due to NASH	58.3 Pack-Years	Both	Female	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)	1.786 (1.139 to 2.722)
Liver cancer due to NASH	72.9 Pack-Years	Both	Male	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)
Liver cancer due to NASH	72.9 Pack-Years	Both	Female	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)
Liver cancer due to other causes	0 Pack-Years	Both	Male	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
liver concer due to other courses	0 Pack Vears	Roth	Fomalo	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
	U Fack-reals	BUUI	remate	1.512	1.512	1.512	1.512	1.512	1.512	1.512	1.512	1.512	1.512	1.512	1.512	1.512	1.512
Liver cancer due to other causes	14.6 Pack-Years	Both	Male	(1.04 to 2.038) 1.512	(1.04 to 2.038) 1.512	(1.04 to 2.038) 1.512	(1.04 to 2.038) 1.512	(1.04 to 2.038) 1.512	(1.04 to 2.038) 1.512	(1.04 to 2.038) 1.512	(1.04 to 2.038) 1.512	(1.04 to 2.038) 1.512	(1.04 to 2.038) 1.512	(1.04 to 2.038) 1.512	(1.04 to 2.038) 1.512	(1.04 to 2.038) 1.512	(1.04 to 2.038) 1.512
Liver cancer due to other causes	14.6 Pack-Years	Both	Female	(1.04 to 2.038) 1.638	(1.04 to 2.038) 1.638	(1.04 to 2.038) 1.638	(1.04 to 2.038) 1.638	(1.04 to 2.038) 1.638	(1.04 to 2.038) 1.638	(1.04 to 2.038) 1.638	(1.04 to 2.038) 1.638	(1.04 to 2.038) 1.638	(1.04 to 2.038) 1.638	(1.04 to 2.038) 1.638	(1.04 to 2.038) 1.638	(1.04 to 2.038) 1.638	(1.04 to 2.038) 1.638
Liver cancer due to other causes	29.2 Pack-Years	Both	Male	(1.099 to 2.256) 1.638	(1.099 to 2.256) 1.638	(1.099 to 2.256) 1.638	(1.099 to 2.256) 1.638	(1.099 to 2.256) 1.638	(1.099 to 2.256) 1.638	(1.099 to 2.256) 1.638	(1.099 to 2.256) 1.638	(1.099 to 2.256) 1.638	(1.099 to 2.256) 1.638	(1.099 to 2.256) 1.638	(1.099 to 2.256) 1.638	(1.099 to 2.256) 1.638	(1.099 to 2.256) 1.638
Liver cancer due to other causes	29.2 Pack-Years	Both	Female	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256)	(1.099 to 2.256) 1.83	(1.099 to 2.256)
Liver cancer due to other causes	43.8 Pack-Years	Both	Male	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)
Liver cancer due to other causes	43.8 Pack-Years	Both	Female	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)	(1.224 to 2.606)
Liver cancer due to other causes	58.3 Pack-Years	Both	Male	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)
Liver cancer due to other causes	58.3 Pack-Years	Both	Female	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)	(1.139 to 2.722)
Liver cancer due to other causes	72.9 Pack-Years	Both	Male	(1.147 to 3.012)	1.937 (1.147 to 3.012)	(1.147 to 3.012)	(1.147 to 3.012)	(1.147 to 3.012)	(1.147 to 3.012)	(1.147 to 3.012)	(1.147 to 3.012)	(1.147 to 3.012)	(1.147 to 3.012)	(1.147 to 3.012)	(1.147 to 3.012)	(1.147 to 3.012)	(1.147 to 3.012)
Liver cancer due to other causes	72.9 Pack-Years	Both	Female	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.93/ (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.93/ (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)	1.937 (1.147 to 3.012)
Pancreatic cancer	0 Pack-Years	Both	Male	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)

Appendix Table 6d. Relative risks used by age and sex	x for each outcome for smoking globally.																
Risk - Outcome	Category / Units	Morbidity / Mortality	Sex	20 + 24	25 (20	101.11	15 - 10	50 + 54	55 - 50	A	Age	70 - 74	75 - 70	20 - 04	05 + 00	00 - 01	
Smoking				50 10 54	351039	40 10 44	45 to 49	50 10 54	5510 59	60 10 64	65 10 69	701074	751079	80 10 84	85 10 89	901094	95 pius
Pancreatic cancer	O Pack-Years	Both	Female	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
				1.653	1.653	1.653	1.653	1.653	1.653	1.653	1.653	1.653	1.653	1.653	1.653	1.653	1.653
Pancreatic cancer	15.8 Pack-Years	BOTH	Male	2.361	2.361	2.361	2.361	2.361	2.361	2.361	2.361	2.361	2.361	2.361	2.361	2.361	2.361
Pancreatic cancer	15.8 Pack-Years	Both	Female	(1.902 to 2.846) 1.799	(1.902 to 2.846) 1.799	(1.902 to 2.846) 1.799	(1.902 to 2.846) 1.799	(1.902 to 2.846) 1.799	(1.902 to 2.846) 1.799) (1.902 to 2.846) 1.799	(1.902 to 2.846) 1.799	(1.902 to 2.846) 1.799					
Pancreatic cancer	31.5 Pack-Years	Both	Male	(1.448 to 2.184)	(1.448 to 2.184 2 745	(1.448 to 2.184)	(1.448 to 2.184)	(1.448 to 2.184)	(1.448 to 2.184)) (1.448 to 2.184) 2 745	(1.448 to 2.184) 2 745	(1.448 to 2.184)	(1.448 to 2.184) 2 745	(1.448 to 2.184)	(1.448 to 2.184)	(1.448 to 2.184)	(1.448 to 2.184)
Pancreatic cancer	31.5 Pack-Years	Both	Female	(2.099 to 3.457)	(2.099 to 3.457	(2.099 to 3.457)	(2.099 to 3.457)	(2.099 to 3.457)	(2.099 to 3.457)) (2.099 to 3.457)	(2.099 to 3.457)	(2.099 to 3.457)					
Pancreatic cancer	47.3 Pack-Years	Both	Male	(1.75 to 2.605)	(1.75 to 2.605)	(1.75 to 2.605)	(1.75 to 2.605)	(1.75 to 2.605)	(1.75 to 2.605)	(1.75 to 2.605)	(1.75 to 2.605)	(1.75 to 2.605)	(1.75 to 2.605)	(1.75 to 2.605)	(1.75 to 2.605)	(1.75 to 2.605)	(1.75 to 2.605)
Pancreatic cancer	47.3 Pack-Years	Both	Female	3.281 (2.36 to 4.34)	3.281 (2.36 to 4.34)	3.281 (2.36 to 4.34)	3.281 (2.36 to 4.34)	3.281 (2.36 to 4.34)	3.281 (2.36 to 4.34)	3.281 (2.36 to 4.34)	3.281 (2.36 to 4.34)	3.281 (2.36 to 4.34)	3.281 (2.36 to 4.34)	3.281 (2.36 to 4.34)	3.281 (2.36 to 4.34)	3.281 (2.36 to 4.34)	3.281 (2.36 to 4.34)
Pancreatic cancer	63 Pack-Years	Both	Male	2.429 (1.793 to 3.211)	2.429 (1.793 to 3.211	2.429) (1.793 to 3.211)	2.429 (1.793 to 3.211)	2.429 (1.793 to 3.211)	2.429 (1.793 to 3.211)	2.429) (1.793 to 3.211)	2.429 (1.793 to 3.211	2.429) (1.793 to 3.211)	2.429 (1.793 to 3.211)				
Pancreatic cancer	63 Pack-Years	Both	Female	3.843 (2.059 to 6.055)	3.843 (2.059 to 6.055	3.843 (2.059 to 6.055)	3.843 (2.059 to 6.055)	3.843 (2.059 to 6.055)	3.843 (2.059 to 6.055)	3.843) (2.059 to 6.055)	3.843 (2.059 to 6.055	3.843) (2.059 to 6.055)	3.843 (2.059 to 6.055)				
	O Pack-Vears	Both	Male	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
	U Pack-Teals	Bour	iviale	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
Larynx cancer	0 Pack-Years	Both	Female	(1.0 to 1.0) 4.593	(1.0 to 1.0) 4.593	(1.0 to 1.0) 4.593	(1.0 to 1.0) 4.593	(1.0 to 1.0) 4.593	(1.0 to 1.0) 4.593	(1.0 to 1.0) 4.593	(1.0 to 1.0) 4.593	(1.0 to 1.0) 4.593	(1.0 to 1.0) 4.593	(1.0 to 1.0) 4.593	(1.0 to 1.0) 4.593	(1.0 to 1.0) 4.593	(1.0 to 1.0) 4.593
Larynx cancer	15 Pack-Years	Both	Male	(2.791 to 6.823) 4.593	(2.791 to 6.823 4.593	(2.791 to 6.823) 4.593	(2.791 to 6.823) 4.593	(2.791 to 6.823) 4.593	(2.791 to 6.823) 4.593) (2.791 to 6.823) 4.593	(2.791 to 6.823) 4.593	(2.791 to 6.823) 4.593					
Larynx cancer	15 Pack-Years	Both	Female	(2.791 to 6.823)	(2.791 to 6.823	(2.791 to 6.823)	(2.791 to 6.823)	(2.791 to 6.823)	(2.791 to 6.823)) (2.791 to 6.823)	(2.791 to 6.823	(2.791 to 6.823)	(2.791 to 6.823)				
				(5.061 to	(5.061 to	(5.061 to	(5.061 to	(5.061 to	(5.061 to	(5.061 to	(5.061 to	(5.061 to	(5.061 to	(5.061 to	(5.061 to	(5.061 to	(5.061 to
Larynx cancer	30 Pack-Years	Both	Male	12.894) 8.448	12.894) 8.448	12.894) 8.448	12.894) 8.448	12.894) 8.448	12.894) 8.448	12.894) 8.448	12.894) 8.448	12.894) 8.448	12.894) 8.448	12.894) 8.448	12.894) 8.448	12.894) 8.448	12.894) 8.448
				(5.061 to	(5.061 to	(5.061 to	(5.061 to	(5.061 to	(5.061 to	(5.061 to	(5.061 to	(5.061 to	(5.061 to	(5.061 to	(5.061 to	(5.061 to	(5.061 to
Larynx cancer	30 Pack-Years	Both	Female	12.894)	12.894)	12.894)	12.894)	12.894)	12.894)	12.894)	12.894)	12.894)	12.894)	12.894)	12.894)	12.894)	12.894)
				(9.234 to	(9.234 to	(9.234 to	(9.234 to	(9.234 to	(9.234 to	(9.234 to	(9.234 to	(9.234 to	(9.234 to	(9.234 to	(9.234 to	(9.234 to	(9.234 to
Larynx cancer	45 Pack-Years	Both	Male	25.714)	25.714)	25.714)	25.714)	25.714)	25.714)	25.714)	25.714)	25.714)	25.714)	25.714)	25.714)	25.714)	25.714)
				16.162	16.162	16.162	16.162 (0.334 to	16.162 (0.334 to	16.162 (0.224 to	16.162 (0.334 to	16.162	16.162 (0.224 to	16.162 (0.224 to	16.162	16.162 (0.224 to	16.162 (0.334 to	16.162
Larynx cancer	45 Pack-Years	Both	Female	25.714)	25.714)	25.714)	25.714)	25.714)	25.714)	25.714)	25.714)	25.714)	25.714)	25.714)	25.714)	25.714)	25.714)
				21.948	21.948	21.948	21.948	21.948	21.948	21.948	21.948	21.948	21.948	21.948	21.948	21.948	21.948
Lanuax cancor	60 Pack Voarc	Roth	Malo	(10.569 to	(10.569 to	(10.569 to	(10.569 to	(10.569 to	(10.569 to	(10.569 to	(10.569 to	(10.569 to	(10.569 to	(10.569 to	(10.569 to	(10.569 to	(10.569 to
Larynx cancer	bu Pack-rears	BOUN	wate	21.948	21.948	21.948	21.948	21.948	21.948	21.948	21.948	21.948	21.948	21.948	21.948	21.948	21.948
				(10.569 to	(10.569 to	(10.569 to	(10.569 to	(10.569 to	(10.569 to	(10.569 to	(10.569 to	(10.569 to	(10.569 to	(10.569 to	(10.569 to	(10.569 to	(10.569 to
Larynx cancer	60 Pack-Years	Both	Female	36.711)	36.711)	36.711)	36.711)	36.711)	36.711)	36.711)	36.711)	36.711)	36.711)	36.711)	36.711)	36.711)	36.711)
				(9.619 to	(9.619 to	(9.619 to	(9.619 to	(9.619 to	(9.619 to	(9.619 to	(9.619 to	(9.619 to	(9.619 to	(9.619 to	(9.619 to	(9.619 to	(9.619 to
Larynx cancer	75 Pack-Years	Both	Male	56.862)	56.862)	56.862)	56.862)	56.862)	56.862)	56.862)	56.862)	56.862)	56.862)	56.862)	56.862)	56.862)	56.862)
				26.147	26.147	26.147	26.147	26.147	26.147	26.147	26.147	26.147	26.147	26.147	26.147	26.147	26.147
Larvoy cancer	75 Pack-Years	Both	Female	(9.619 to 56.862)	(9.619 to 56.862)	(9.619 to 56.862)	(9.619 to 56.862)	(9.619 to 56.862)	(9.619 to 56.862)	(9.619 to 56.862)	(9.619 to 56.862)	(9.619 to 56.862)	(9.619 to 56.862)	(9.619 to 56.862)	(9.619 to 56.862)	(9.619 to 56.862)	(9.619 to 56.862)
	/or dex reals	bour	remare	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
Tracheal, bronchus, and lung cancer	0 Pack-Years	Both	Male	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)
Tracheal, bronchus, and lung cancer	0 Pack-Years	Both	Female	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Tracheal, bronchus, and lung cancer	5 Pack-Years	Both	Male	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)
Tracheal bronchus and lung cancer	5 Pack-Years	Both	Female	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)	1.758 (1.215 to 2.45)
	STack (cars			4.86	4.86	4.86	4.86	4.86	4.86	4.86	4.86	4.86	4.86	4.86	4.86	4.86	4.86
Tracheal, bronchus, and lung cancer	14.3 Pack-Years	Both	Male	(3.917 to 5.866) 4.86	4.86	4.86 4.86	(3.917 to 5.866) 4.86	(3.917 to 5.866) 4.86	(3.917 to 5.866) 4.86	4.86 (3.917 to 5.866)	(3.917 to 5.866) 4.86	(3.917 to 5.866) 4.86	4.86	(3.917 to 5.866) 4.86	(3.917 to 5.866) 4.86	(3.917 to 5.866) 4.86	(3.917 to 5.866) 4.86
Tracheal, bronchus, and lung cancer	14.3 Pack-Years	Both	Female	(3.917 to 5.866) 8.899	(3.917 to 5.866 8.899	(3.917 to 5.866) 8.899	(3.917 to 5.866) 8.899	(3.917 to 5.866) 8.899	(3.917 to 5.866) 8.899) (3.917 to 5.866) 8.899	(3.917 to 5.866 8.899	(3.917 to 5.866) 8.899	(3.917 to 5.866) 8.899				
				(7.377 to	(7.377 to	(7.377 to	(7.377 to	(7.377 to	(7.377 to	(7.377 to	(7.377 to	(7.377 to	(7.377 to	(7.377 to	(7.377 to	(7.377 to	(7.377 to
Tracheal, bronchus, and lung cancer	28.6 Pack-Years	Both	Male	10.562)	10.562)	10.562)	10.562)	10.562)	10.562)	10.562)	10.562)	10.562)	10.562)	10.562)	10.562)	10.562)	10.562)
				8.899 (7.377 to	8.899 (7.377 to	8.899 (7.377 to	8.899 (7.377 to	8.899 (7.377 to	8.899 (7.377 to	8.899 (7.377 to	8.899 (7.377 to	8.899 (7.377 to	8.899 (7.377 to	8.899 (7.377 to	8.899 (7.377 to	8.899 (7.377 to	8.899 (7.377 to
Tracheal, bronchus, and lung cancer	28.6 Pack-Years	Both	Female	10.562)	10.562)	10.562)	10.562)	10.562)	10.562)	10.562)	10.562)	10.562)	10.562)	10.562)	10.562)	10.562)	10.562)
· •				13.509	13.509	13.509	13.509	13.509	13.509	13.509	13.509	13.509	13.509	13.509	13.509	13.509	13.509
				(10.479 to	(10.479 to	(10.479 to	(10.479 to	(10.479 to	(10.479 to	(10.479 to	(10.479 to	(10.479 to	(10.479 to	(10.479 to	(10.479 to	(10.479 to	(10.479 to
Iracneal, bronchus, and lung cancer	42.9 Pack-Years	Both	Male	10.81)	13 500	15.81)	10.81)	13.500	10.81)	10.81)	18.81)	13.500	13.500	13 500	13 500	10.81)	10.81)
				(10.479 to	(10.479 to	(10.479 to	(10.479 to	(10.479 to	(10.479 to	(10.479 to	(10.479 to	(10.479 to	(10.479 to	(10.479 to	(10.479 to	(10.479 to	(10.479 to
Tracheal, bronchus, and lung cancer	42.9 Pack-Years	Both	Female	16.81)	16.81)	16.81)	16.81)	16.81)	16.81)	16.81)	16.81)	16.81)	16.81)	16.81)	16.81)	16.81)	16.81)

Appendix Table 6d. Relative risks used by age and sex for each ou	tcome for smoking globally.																
Risk - Outcome	Category / Units	Morbidity / Mortality	Sex		1	_		1	•	A	ge	-		1	1		
Smoking				30 to 34	35 to 39	40 to 44	45 to 49	50 to 54	55 to 59	60 to 64	65 to 69	70 to 74	75 to 79	80 to 84	85 to 89	90 to 94	95 plus
Shoking				14.83	14.83	14.83	14.83	14.83	14.83	14.83	14.83	14.83	14.83	14.83	14.83	14.83	14.83
				(10.64 to	(10.64 to	(10.64 to	(10.64 to	(10.64 to	(10.64 to	(10.64 to	(10.64 to	(10.64 to	(10.64 to	(10.64 to	(10.64 to	(10.64 to	(10.64 to
I racheal, bronchus, and lung cancer	57.1 Pack-Years	Both	Male	19.371) 14.83	19.371) 14.83	19.371) 14.83	19.3/1) 14.83	19.371) 14.83	19.371) 14.83	19.371) 14.83	19.371) 14.83	19.371) 14.83	19.371) 14.83	19.371) 14.83	19.371) 14.83	19.371) 14.83	19.371) 14.83
				(10.64 to	(10.64 to	(10.64 to	(10.64 to	(10.64 to	(10.64 to	(10.64 to	(10.64 to	(10.64 to	(10.64 to	(10.64 to	(10.64 to	(10.64 to	(10.64 to
Tracheal, bronchus, and lung cancer	57.1 Pack-Years	Both	Female	19.371) 18.644	19.371) 18.644	19.371) 18.644	19.3/1) 18.644	19.371) 18.644	19.371) 18.644	19.371) 18.644	19.371) 18.644	19.371) 18.644	19.371) 18.644	19.371) 18.644	19.371) 18.644	19.371) 18.644	19.3/1) 18.644
				(12.787 to	(12.787 to	(12.787 to	(12.787 to	(12.787 to	(12.787 to	(12.787 to	(12.787 to	(12.787 to	(12.787 to	(12.787 to	(12.787 to	(12.787 to	(12.787 to
I racheal, bronchus, and lung cancer	/1.4 Pack-Years	Both	Male	25.573) 18.644	25.573) 18.644	25.573) 18.644	25.573) 18.644	25.573) 18.644	25.573) 18.644	25.573) 18.644	25.573) 18.644	25.573) 18.644	25.573) 18.644	25.573) 18.644	25.573) 18.644	25.573) 18.644	25.573) 18.644
-	74 4 9 - 1 1	D ()	5	(12.787 to	(12.787 to	(12.787 to	(12.787 to	(12.787 to	(12.787 to	(12.787 to	(12.787 to	(12.787 to	(12.787 to	(12.787 to	(12.787 to	(12.787 to	(12.787 to
i racneai, bronchus, and lung cancer	/1.4 Pack-Years	BOTH	Female	25.573) 21.525	25.573) 21.525	25.573) 21.525	25.573) 21.525	25.573) 21.525	25.573) 21.525	25.573) 21.525	25.573) 21.525	25.573) 21.525	25.573) 21.525	25.573) 21.525	25.573) 21.525	25.573) 21.525	25.573) 21.525
				(13.899 to	(13.899 to	(13.899 to	(13.899 to	(13.899 to	(13.899 to	(13.899 to	(13.899 to	(13.899 to	(13.899 to	(13.899 to	(13.899 to	(13.899 to	(13.899 to
I racheal, bronchus, and lung cancer	85. / Pack-Years	Both	Male	30.415) 21.525	30.415) 21.525	21.525	30.415) 21.525	21.525	21.525	21.525	21.525	21.525	21.525	21.525	21.525	30.415) 21.525	21.525
				(13.899 to	(13.899 to	(13.899 to	(13.899 to	(13.899 to	(13.899 to	(13.899 to	(13.899 to	(13.899 to	(13.899 to	(13.899 to	(13.899 to	(13.899 to	(13.899 to
Tracheal, bronchus, and lung cancer	85.7 Pack-Years	Both	Female	30.415) 1.0	30.415) 1.0	30.415) 1.0	30.415) 1.0	30.415) 1.0	30.415) 1.0	30.415) 1.0	30.415) 1.0	30.415) 1.0	30.415) 1.0	30.415) 1.0	30.415) 1.0	30.415) 1.0	30.415) 1.0
Breast cancer	0 Pack-Years	Both	Female	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)
Breast cancer	12.8 Pack-Years	Both	Female	1.207 (1.094 to 1.322)	1.207 (1.094 to 1.32)	1.207 2) (1.094 to 1.322	1.207) (1.094 to 1.322)	1.207 (1.094 to 1.322)	1.207 (1.094 to 1.322)	1.207) (1.094 to 1.322)	1.207 (1.094 to 1.322	1.207 (1.094 to 1.322)					
				1.31	1.31	1.31	1.31	1.31	1.31	1.31	1.31	1.31	1.31	1.31	1.31	1.31	1.31
Breast cancer	25.5 Pack-Years	Both	Female	(1.205 to 1.431) 1.242	(1.205 to 1.43) 1.242	 (1.205 to 1.431 1.242) (1.205 to 1.431) 1.242	(1.205 to 1.431) 1.242	(1.205 to 1.431) 1.242) (1.205 to 1.431) 1.242	(1.205 to 1.431 1.242) (1.205 to 1.431) 1.242					
Breast cancer	38.2 Pack-Years	Both	Female	(1.095 to 1.383)	(1.095 to 1.38	3) (1.095 to 1.383) (1.095 to 1.383)	(1.095 to 1.383)	(1.095 to 1.383)) (1.095 to 1.383)	(1.095 to 1.383) (1.095 to 1.383)					
Breast cancer	63.8 Pack-Years	Both	Female	1.274 (1.083 to 1.492)	1.274 (1.083 to 1.49)	1.274 2) (1.083 to 1.492	1.274) (1.083 to 1.492)	1.274 (1.083 to 1.492)	1.274 (1.083 to 1.492)	1.274) (1.083 to 1.492)	1.274 (1.083 to 1.492	1.274 (1.083 to 1.492)					
				1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
Cervical cancer	0 Pack-Years	Both	Female	(1.0 to 1.0) 1.785	(1.0 to 1.0) 1.785	(1.0 to 1.0) 1.785	(1.0 to 1.0) 1.785	(1.0 to 1.0) 1.785	(1.0 to 1.0) 1.785	(1.0 to 1.0) 1.785	(1.0 to 1.0) 1.785	(1.0 to 1.0) 1.785	(1.0 to 1.0) 1.785	(1.0 to 1.0) 1.785	(1.0 to 1.0) 1.785	(1.0 to 1.0) 1.785	(1.0 to 1.0) 1.785
Cervical cancer	5 Pack-Years	Both	Female	(1.101 to 2.751)	(1.101 to 2.75	1) (1.101 to 2.751) (1.101 to 2.751)	(1.101 to 2.751)	(1.101 to 2.751)) (1.101 to 2.751)	(1.101 to 2.751) (1.101 to 2.751)					
Cervical cancer	7.4 Pack-Years	Both	Female	1.941 (1.17 to 2.981)	1.941 (1.17 to 2.981	1.941 (1.17 to 2.981)	1.941 (1.17 to 2.981)	1.941 (1.17 to 2.981)	1.941 (1.17 to 2.981)	1.941 (1.17 to 2.981)	1.941 (1.17 to 2.981)	1.941 (1.17 to 2.981)	1.941 (1.17 to 2.981)	1.941 (1.17 to 2.981)	1.941 (1.17 to 2.981)	1.941 (1.17 to 2.981)	1.941) (1.17 to 2.981)
				2.245	2.245	2.245	2.245	2.245	2.245	2.245	2.245	2.245	2.245	2.245	2.245	2.245	2.245
Cervical cancer	14.8 Pack-Years	Both	Female	(1.257 to 4.099) 3.355	(1.257 to 4.09) 3.355	9) (1.257 to 4.099 3.355) (1.257 to 4.099) 3.355	(1.257 to 4.099) 3.355	(1.257 to 4.099) 3.355) (1.257 to 4.099) 3.355) (1.257 to 4.099) 3.355						
Cervical cancer	22.1 Pack-Years	Both	Female	(1.72 to 6.124)	(1.72 to 6.124	(1.72 to 6.124)	(1.72 to 6.124)	(1.72 to 6.124)	(1.72 to 6.124)	(1.72 to 6.124)	(1.72 to 6.124)	(1.72 to 6.124)	(1.72 to 6.124)	(1.72 to 6.124)	(1.72 to 6.124)	(1.72 to 6.124)	(1.72 to 6.124)
Cervical cancer	29.5 Pack-Years	Both	Female	4.207 (1.745 to 8.534)	4.207 (1.745 to 8.53	4.207 4) (1.745 to 8.534	4.207) (1.745 to 8.534)	4.207 (1.745 to 8.534)	4.207 (1.745 to 8.534)	4.207) (1.745 to 8.534)	4.207 (1.745 to 8.534	4.207 (1.745 to 8.534)					
				1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
Prostate cancer	0 Cigarettes Per Smoker Per Day	Both	Male	(1.0 to 1.0) 1.192	(1.0 to 1.0) 1.192	(1.0 to 1.0) 1.192	(1.0 to 1.0) 1.192	(1.0 to 1.0) 1.192	(1.0 to 1.0) 1.192	(1.0 to 1.0) 1.192	(1.0 to 1.0) 1.192	(1.0 to 1.0) 1.192	(1.0 to 1.0) 1.192	(1.0 to 1.0) 1.192	(1.0 to 1.0) 1.192	(1.0 to 1.0) 1.192	(1.0 to 1.0) 1.192
Prostate cancer	7.5 Cigarettes Per Smoker Per Day	Both	Male	(1.008 to 1.382)	(1.008 to 1.38	2) (1.008 to 1.382) (1.008 to 1.382)	(1.008 to 1.382)	(1.008 to 1.382)) (1.008 to 1.382)	(1.008 to 1.382) (1.008 to 1.382)					
Prostate cancer	15 Cigarettes Per Smoker Per Day	Both	Male	1.173 (1.004 to 1.371)	1.173 (1.004 to 1.37)	1.173 1) (1.004 to 1.371	1.173) (1.004 to 1.371)	1.173 (1.004 to 1.371)	1.173 (1.004 to 1.371)	1.173 (1.004 to 1.371)	1.173 (1.004 to 1.371)	1.173 (1.004 to 1.371)	1.173 (1.004 to 1.371)	1.173 (1.004 to 1.371)	1.173 (1.004 to 1.371)	1.1/3 (1.004 to 1.371	1.173 .) (1.004 to 1.371)
				1.166	1.166	1.166	1.166	1.166	1.166	1.166	1.166	1.166	1.166	1.166	1.166	1.166	1.166
Prostate cancer	22.5 Cigarettes Per Smoker Per Day	Both	Male	(1.001 to 1.359) 1.235	1.235	9) (1.001 to 1.359 1.235	1.235 (1.001 to 1.359)	(1.001 to 1.359) 1.235	(1.001 to 1.359) 1.235	1.235 (1.001 to 1.359)	(1.001 to 1.359) 1.235	1.235) (1.001 to 1.359) 1.235				
Prostate cancer	30 Cigarettes Per Smoker Per Day	Both	Male	(1.006 to 1.511)	(1.006 to 1.51	1) (1.006 to 1.511) (1.006 to 1.511)	(1.006 to 1.511)	(1.006 to 1.511)) (1.006 to 1.511)	(1.006 to 1.511) (1.006 to 1.511)					
Prostate cancer	37.5 Cigarettes Per Smoker Per Day	Both	Male	1.33 (1.0 to 1.783)	1.33 (1.0 to 1.783)	1.33 (1.0 to 1.783)	1.33 (1.0 to 1.783)	1.33 (1.0 to 1.783)	1.33 (1.0 to 1.783)	1.33 (1.0 to 1.783)	1.33 (1.0 to 1.783)	1.33 (1.0 to 1.783)	1.33 (1.0 to 1.783)	1.33 (1.0 to 1.783)	1.33 (1.0 to 1.783)	1.33 (1.0 to 1.783)	1.33 (1.0 to 1.783)
V:	O Darle Veren	Deth	Mala	1.0 (1.0 to 1.0)	1.0	1.0 (1.0 to 1.0)	1.0	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0	1.0 (1.0 to 1.0)
kidney cancer	U Pack-Years	BOTH	wale	1.0	1.0	(1.0 10 1.0)	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
Kidney cancer	0 Pack-Years	Both	Female	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)
Kidney cancer	12.3 Pack-Years	Both	Male	(1.046 to 1.748)	(1.046 to 1.74	1.300 8) (1.046 to 1.748	1.300) (1.046 to 1.748)	(1.046 to 1.748)	(1.046 to 1.748)	1.300) (1.046 to 1.748)	(1.046 to 1.748)	(1.046 to 1.748)	(1.046 to 1.748)	(1.046 to 1.748)	(1.046 to 1.748)	(1.046 to 1.748	i) (1.046 to 1.748)
Kidooy cancor	12 2 Back Years	Both	Fomalo	1.366	1.366	1.366	1.366	1.366	1.366	1.366	1.366	1.366	1.366	1.366	1.366	1.366	1.366
	12.5 Fack-Teals	BOUT	remale	1.73	1.73	1.73	1.73	1.73	1.73	1.73	1.73	1.73	1.73	1.73	1.73	1.73	1.73
Kidney cancer	24.6 Pack-Years	Both	Male	(1.34 to 2.164)	(1.34 to 2.164	(1.34 to 2.164) 1 73	(1.34 to 2.164)	(1.34 to 2.164)	(1.34 to 2.164)	(1.34 to 2.164)	(1.34 to 2.164)	(1.34 to 2.164)	(1.34 to 2.164)	(1.34 to 2.164)	(1.34 to 2.164)	(1.34 to 2.164)	(1.34 to 2.164)
Kidney cancer	24.6 Pack-Years	Both	Female	(1.34 to 2.164)	(1.34 to 2.164	(1.34 to 2.164)	(1.34 to 2.164)	(1.34 to 2.164)	(1.34 to 2.164)	(1.34 to 2.164)	(1.34 to 2.164)	(1.34 to 2.164)	(1.34 to 2.164)	(1.34 to 2.164)	(1.34 to 2.164)	(1.34 to 2.164)	(1.34 to 2.164)
Kidney cancer	36 0 Pack-Vears	Both	Male	1.757 (1.296 to 2.309)	1.757	1.757 9) (1.296 to 2.309	1.757	1.757 (1.296 to 2.309)	1.757	1.757	1.757	1.757 (1.296 to 2.309)	1.757	1.757	1.757	1.757	1.757
nuncy canter	50.5 FOCK-FC013	5001	ware	1.757	1.757	1.757	1.757	1.757	1.757	1.757	1.757	1.757	1.757	1.757	1.757	1.757	1.757
Kidney cancer	36.9 Pack-Years	Both	Female	(1.296 to 2.309)	(1.296 to 2.30)	9) (1.296 to 2.309 1 897) (1.296 to 2.309) 1.892	(1.296 to 2.309) 1.892	(1.296 to 2.309) 1.897) (1.296 to 2.309) 1.892	(1.296 to 2.309) 1.892	(1.296 to 2.309) 1.892	(1.296 to 2.309) 1.892	(1.296 to 2.309)	(1.296 to 2.309) 1.892	(1.296 to 2.309)) (1.296 to 2.309) 1.892
Kidney cancer	49.2 Pack-Years	Both	Male	(1.429 to 2.44)	(1.429 to 2.44	(1.429 to 2.44)	(1.429 to 2.44)	(1.429 to 2.44)	(1.429 to 2.44)	(1.429 to 2.44)	(1.429 to 2.44)	(1.429 to 2.44)	(1.429 to 2.44)	(1.429 to 2.44)	(1.429 to 2.44)	(1.429 to 2.44)	(1.429 to 2.44)
Kidney cancer	49.2 Pack-Years	Both	Female	1.892 (1.429 to 2.44)	1.892 (1.429 to 2.44	1.892	1.892 (1.429 to 2.44)	1.892 (1.429 to 2.44)	1.892 (1.429 to 2.44)	1.892 (1.429 to 2.44)	1.892 (1.429 to 2.44)	1.892 (1.429 to 2.44)	1.892 (1.429 to 2.44)	1.892 (1.429 to 2.44)	1.892 (1.429 to 2.44)	1.892 (1.429 to 2.44)	1.892
			. emaie	2.006	2.006	2.006	2.006	2.006	2.006	2.006	2.006	2.006	2.006	2.006	2.006	2.006	2.006
Kidney cancer	61.5 Pack-Years	Both	Male	(1.254 to 3.032)	(1.254 to 3.03)	2) (1.254 to 3.032) (1.254 to 3.032)	(1.254 to 3.032)	(1.254 to 3.032)) (1.254 to 3.032)	(1.254 to 3.032) (1.254 to 3.032)					
Appendix Table 6d. Relative risks used by age and sex for each or	atcome for smoking globally.																
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Risk - Outcome	Category / Units	Morbidity / Mortality	Sex	30 to 34	35 to 39	40 to 44	45 to 49	50 to 54	55 to 59	60 to 64	.ge 65 to 69	70 to 74	75 to 79	80 to 84	85 to 89	90 to 94	95 plus
Smoking																	
Kidnev cancer	61.5 Pack-Years	Both	Female	2.006 (1.254 to 3.032)													
Bladder cancer	0 Pack-Years	Both	Male	1.0 (1.0 to 1.0)													
		Butt		1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0
Bladder Cancer	U Pack-Years	BOTH	Female	2.804	2.804	2.804	2.804	2.804	2.804	2.804	2.804	2.804	2.804	2.804	2.804	2.804	2.804
Bladder cancer	15 Pack-Years	Both	Male	(1.695 to 4.161) 2.804													
Bladder cancer	15 Pack-Years	Both	Female	(1.695 to 4.161) 3.328													
Bladder cancer	30 Pack-Years	Both	Male	(2.093 to 4.997) 3 328													
Bladder cancer	30 Pack-Years	Both	Female	(2.093 to 4.997)													
Bladder cancer	45 Pack-Years	Both	Male	(2.45 to 6.526)	(2.45 to 6.526)	(2.45 to 6.526)	(2.45 to 6.526)	4.235 (2.45 to 6.526)	(2.45 to 6.526)	(2.45 to 6.526)	(2.45 to 6.526)	(2.45 to 6.526)	(2.45 to 6.526)	(2.45 to 6.526)	(2.45 to 6.526)	(2.45 to 6.526)	(2.45 to 6.526)
Bladder cancer	45 Pack-Years	Both	Female	4.239 (2.45 to 6.526)													
Bladder cancer	60 Pack-Years	Both	Male	4.553 (2.544 to 7.643)													
Bladder cancer	60 Pack-Years	Both	Female	4.553 (2.544 to 7.643)													
Acute lymphoid leukaemia	0 Pack-Years	Both	Male	1.0 (1.0 to 1.0)													
Acute lymphoid leukaemia	0 Pack-Years	Both	Female	1.0 (1.0 to 1.0)													
	13 4 Back Voarc	Roth	Mala	2.011 (1.0 to 2.600)	2.011	2.011	2.011	2.011 (1.0 to 2.600)	2.011 (1.0 to 3.600)	2.011 (1.0 to 2.600)	2.011	2.011 (1.0 to 2.600)	2.011 (1.0 to 2.600)	2.011	2.011	2.011	2.011 (1.0 to 2.600)
	12.4 Pack-reals	Both	iviale	2.011	2.011	2.011	2.011	2.011	2.011	2.011	2.011	2.011	2.011	2.011	2.011	2.011	2.011
Acute lymphold leukaemia	12.4 Pack-Years	BOTH	Female	(1.0 to 3.609) 2.253													
Acute lymphoid leukaemia	24.8 Pack-Years	Both	Male	(1.071 to 3.961) 2.253													
Acute lymphoid leukaemia	24.8 Pack-Years	Both	Female	(1.071 to 3.961) 2.64													
Acute lymphoid leukaemia	37.1 Pack-Years	Both	Male	(1.483 to 4.386) 2.64													
Acute lymphoid leukaemia	37.1 Pack-Years	Both	Female	(1.483 to 4.386)													
Chronic lymphoid leukaemia	0 Pack-Years	Both	Male	(1.0 to 1.0)													
Chronic lymphoid leukaemia	0 Pack-Years	Both	Female	(1.0 to 1.0)													
Chronic lymphoid leukaemia	12.4 Pack-Years	Both	Male	2.011 (1.0 to 3.609)													
Chronic lymphoid leukaemia	12.4 Pack-Years	Both	Female	2.011 (1.0 to 3.609)													
Chronic lymphoid leukaemia	24.8 Pack-Years	Both	Male	2.253 (1.071 to 3.961)													
Chronic lymphoid leukaemia	24 8 Pack-Years	Both	Female	2.253 (1.071 to 3.961)													
	27.1 Bask Veers	Deth	Mala	2.64	2.64	2.64	2.64	2.64	2.64	2.64	2.64	2.64	2.64	2.64	2.64	2.64	2.64
	57.1 Pack-Years	BOLI	Male	2.64	2.64	2.64	2.64	2.64	2.64	2.64	2.64	2.64	2.64	2.64	2.64	2.64	2.64
Chronic lymphoid leukaemia	37.1 Pack-Years	Both	Female	(1.483 to 4.386) 1.0													
Acute myeloid leukaemia	0 Pack-Years	Both	Male	(1.0 to 1.0) 1.0													
Acute myeloid leukaemia Acute myeloid leukaemia	0 Pack-Years 12.4 Pack-Years	Both Both	Female Male	(1.0 to 1.0) 2.011													
Acute myeloid leukaemia	12.4 Pack-Years	Both	Female	2.011 (1.0 to 3.609)													
	24 8 Pack-Vears	Both	Male	2.253 (1.071 to 3.961)	2.253 (1.071 to 3.951)	2.253 (1.071 to 3.961)	2.253	2.253 (1.071 to 3.961)	2.253 (1.071 to 3.951)	2.253 (1.071 to 3.951)	2.253	2.253	2.253 (1.071 to 3.961)	2.253 (1.071 to 3.951)	2.253 (1.071 to 3.961)	2.253	2.253
		Buth		2.253	2.253	2.253	2.253	2.253	2.253	2.253	2.253	2.253	2.253	2.253	2.253	2.253	2.253
	24.8 Pack-Years	BOTH	Female	2.64	2.64	(1.071 to 3.961) 2.64	2.64	(1.0/1 to 3.961) 2.64	2.64	2.64	2.64	2.64	(1.0/1 to 3.961) 2.64	2.64	(1.0/1 to 3.961) 2.64	(1.071 to 3.961) 2.64	(1.0/1 to 3.961) 2.64
Acute myeloid leukaemia	37.1 Pack-Years	Both	Male	(1.483 to 4.386) 2.64													
Acute myeloid leukaemia	37.1 Pack-Years	Both	Female	(1.483 to 4.386) 1.0													
Chronic myeloid leukaemia	0 Pack-Years	Both	Male	(1.0 to 1.0) 1.0													
Chronic myeloid leukaemia	0 Pack-Years	Both	Female	(1.0 to 1.0) 2.011													
Chronic myeloid leukaemia	12.4 Pack-Years	Both	Male	(1.0 to 3.609)													

Appendix Table 6d. Relative risks used by age and sex for each ou	tcome for smoking globally.																
Risk - Outcome	Category / Units	Morbidity / Mortality	Sex	30 to 34	35 to 39	40 to 44	45 to 49	50 to 54	55 to 59	A 60 to 64	ge 65 to 69	70 to 74	75 to 79	80 to 84	85 to 89	90 to 94	95 plus
Smoking																	
Chronic myeloid leukaemia	12.4 Pack-Years	Both	Female	2.011 (1.0 to 3.609)	2.011 (1.0 to 3.609)	2.011 (1.0 to 3.609)	2.011 (1.0 to 3.609)	2.011 (1.0 to 3.609)	2.011 (1.0 to 3.609)	2.011 (1.0 to 3.609)	2.011 (1.0 to 3.609)	2.011 (1.0 to 3.609)	2.011 (1.0 to 3.609)	2.011 (1.0 to 3.609)	2.011 (1.0 to 3.609)	2.011 (1.0 to 3.609)	2.011 (1.0 to 3.609)
Chronic myeloid leukaemia	24.8 Pack-Years	Both	Male	2.253 (1.071 to 3.961)	2.253 (1.071 to 3.961	2.253 .) (1.071 to 3.961)	2.253 (1.071 to 3.961)	2.253 (1.071 to 3.961)	2.253 (1.071 to 3.961)	2.253 (1.071 to 3.961)	2.253 (1.071 to 3.961)	2.253 (1.071 to 3.961)	2.253 (1.071 to 3.961)	2.253 (1.071 to 3.961)	2.253 (1.071 to 3.961)	2.253 (1.071 to 3.961)	2.253 (1.071 to 3.961)
Chronic myeloid leukaemia	24.8 Pack-Years	Both	Female	2.253 (1.071 to 3.961)	2.253 (1.071 to 3.961	2.253 (1.071 to 3.961)	2.253) (1.071 to 3.961)	2.253 (1.071 to 3.961)									
Chronic myeloid leukaemia	37.1 Pack-Years	Both	Male	2.64 (1.483 to 4.386)	2.64 (1.483 to 4.386	2.64 6) (1.483 to 4.386)	2.64) (1.483 to 4.386)	2.64 (1.483 to 4.386)									
Chronic myeloid leukaemia	37.1 Pack-Years	Both	Female	2.64 (1.483 to 4.386)	2.64 (1.483 to 4.386	2.64 (1.483 to 4.386)	2.64) (1.483 to 4.386)	2.64 (1.483 to 4.386)									
Other leukaemia	0 Pack-Years	Both	Male	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Other leukaemia	0 Pack-Years	Both	Female	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	(1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Other leukaemia	12.4 Pack-Years	Both	Male	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)
Other leukaemia	12.4 Pack-Years	Both	Female	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)	(1.0 to 3.609)
Other leukaemia	24.8 Pack-Years	Both	Male	(1.071 to 3.961)	(1.071 to 3.961) (1.071 to 3.961) 2 253	(1.071 to 3.961)	(1.071 to 3.961)	(1.071 to 3.961)	(1.071 to 3.961)	(1.071 to 3.961)	(1.071 to 3.961)	(1.071 to 3.961)	(1.071 to 3.961)	(1.071 to 3.961)	(1.071 to 3.961)	(1.071 to 3.961)
Other leukaemia	24.8 Pack-Years	Both	Female	(1.071 to 3.961)	(1.071 to 3.961	.) (1.071 to 3.961)	(1.071 to 3.961)	(1.071 to 3.961)	(1.071 to 3.961)	(1.071 to 3.961)	(1.071 to 3.961)	(1.071 to 3.961)	(1.071 to 3.961)	(1.071 to 3.961)	(1.071 to 3.961)	(1.071 to 3.961)	(1.071 to 3.961)
Other leukaemia	37.1 Pack-Years	Both	Male	(1.483 to 4.386) 2.64	(1.483 to 4.386 2.64	i) (1.483 to 4.386) 2.64	(1.483 to 4.386) 2.64	(1.483 to 4.386) 2.64	(1.483 to 4.386) 2.64	(1.483 to 4.386) 2.64	(1.483 to 4.386) 2.64	(1.483 to 4.386) 2.64	(1.483 to 4.386) 2.64	(1.483 to 4.386) 2.64	(1.483 to 4.386) 2.64	(1.483 to 4.386) 2.64	(1.483 to 4.386) 2.64
Other leukaemia	37.1 Pack-Years	Both	Female	(1.483 to 4.386) 1.0	(1.483 to 4.386	i) (1.483 to 4.386)) (1.483 to 4.386) 1.0	(1.483 to 4.386) 1.0	(1.483 to 4.386) 1.0	(1.483 to 4.386) 1.0	(1.483 to 4.386)	(1.483 to 4.386) 1.0	(1.483 to 4.386)	(1.483 to 4.386) 1.0	(1.483 to 4.386) 1.0	(1.483 to 4.386) 1.0	(1.483 to 4.386)
Ischaemic heart disease	0 Cigarettes Per Smoker Per Day	Both	Male	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0
Ischaemic heart disease	0 Cigarettes Per Smoker Per Day	Both	Female	(1.0 to 1.0) 2.965	(1.0 to 1.0) 2.965	(1.0 to 1.0) 2.965	(1.0 to 1.0) 2.526	(1.0 to 1.0) 2.526	(1.0 to 1.0) 2.125	(1.0 to 1.0) 2.125	(1.0 to 1.0) 1.773	(1.0 to 1.0) 1.773	(1.0 to 1.0) 1.46	(1.0 to 1.0) 1.46	(1.0 to 1.0) 1.187	(1.0 to 1.0) 1.187	(1.0 to 1.0) 1.002
Ischaemic heart disease	11.2 Cigarettes Per Smoker Per Day	Both	Male	(2.571 to 3.372) 3.783	(2.571 to 3.372 3.783	(2.571 to 3.372) 3.783	(2.229 to 2.813) 3.183	(2.229 to 2.813) 3.183	(1.907 to 2.346) 2.608	(1.907 to 2.346) 2.608	(1.641 to 1.908) 2.131	(1.641 to 1.908) 2.131	(1.375 to 1.558) 1.68	(1.375 to 1.558) 1.68	(1.132 to 1.247) 1.277	(1.132 to 1.247) 1.277	(1.0 to 1.01) 1.004
Ischaemic heart disease	11.2 Cigarettes Per Smoker Per Day	Both	Female	(2.956 to 4.813) 3.29	(2.956 to 4.813 3.29	 (2.956 to 4.813) 3.29) (2.593 to 3.822) 2.671	(2.593 to 3.822) 2.671	(2.201 to 3.027) 2.188	(2.201 to 3.027) 2.188	(1.906 to 2.365) 1.784	(1.906 to 2.365) 1.784	(1.559 to 1.792) 1.457	(1.559 to 1.792) 1.457	(1.224 to 1.332) 1.188	(1.224 to 1.332) 1.188	(1.0 to 1.013) 1.001
Ischaemic heart disease	22.5 Cigarettes Per Smoker Per Day	Both	Male	(2.805 to 3.847) 6.684	(2.805 to 3.847 6.684	(2.805 to 3.847) 6.684) (2.327 to 3.056) 4.917	(2.327 to 3.056) 4.917	(1.945 to 2.427) 3.569	(1.945 to 2.427) 3.569	(1.626 to 1.964) 2.572	(1.626 to 1.964) 2.572	(1.369 to 1.557) 1.85	(1.369 to 1.557) 1.85	(1.147 to 1.234) 1.33	(1.147 to 1.234) 1.33	(1.0 to 1.007) 1.003
Ischaemic heart disease	22.5 Cigarettes Per Smoker Per Day	Both	Female	(5.11 to 8.413) 4.276	(5.11 to 8.413) 4.276	(5.11 to 8.413) 4.276	(3.923 to 5.931) 3.388	(3.923 to 5.931) 3.388	(3.012 to 4.161) 2.634	(3.012 to 4.161) 2.634	(2.258 to 2.875) 2.054	(2.258 to 2.875) 2.054	(1.709 to 1.999) 1.599	(1.709 to 1.999) 1.599	(1.279 to 1.386) 1.236	(1.279 to 1.386) 1.236	(1.0 to 1.012) 1.002
Ischaemic heart disease	33.8 Cigarettes Per Smoker Per Day	Both	Male	(3.407 to 5.103) 9.744	(3.407 to 5.103 9.744	(3.407 to 5.103) 9.744) (2.792 to 4.008)	(2.792 to 4.008)	(2.256 to 3.038)	(2.256 to 3.038)	(1.821 to 2.296)	(1.821 to 2.296)	(1.458 to 1.731)	(1.458 to 1.731)	(1.169 to 1.311)	(1.169 to 1.311)	(1.0 to 1.013)
lschaemic heart disease	33.8 Cigarettes Per Smoker Per Day	Both	Female	(7.026 to 12.903)	(7.026 to 12.903)	(7.026 to 12.903)	6.798 (5.18 to 8.755)	6.798 (5.18 to 8.755)	4.678 (3.713 to 5.73)	4.678 (3.713 to 5.73)	3.246 (2.742 to 3.794)	3.246 (2.742 to 3.794)	2.178 (1.938 to 2.425)	2.178 (1.938 to 2.425)	1.452 (1.369 to 1.536)	1.452 (1.369 to 1.536)	1.004 (1.0 to 1.014)
Ischaemic heart disease	45 Cigarettes Per Smoker Per Day	Both	Male	5.45 (3.718 to 7.541)	5.45 (3.718 to 7.541	5.45 .) (3.718 to 7.541)	4.139) (2.953 to 5.545)	4.139 (2.953 to 5.545)	3.075 (2.296 to 3.931)	3.075 (2.296 to 3.931)	2.325 (1.746 to 2.927)	2.325 (1.746 to 2.927)	1.737 (1.39 to 2.104)	1.737 (1.39 to 2.104)	1.303 (1.132 to 1.484)	1.303 (1.132 to 1.484)	(1.0 to 1.038)
	45 Cinemattee Des Caralise Des Deu	Deth	Formela	(7.222 to	(7.222 to	(7.222 to	7.729	7.729	5.097	5.097	3.414	3.414	2.239 (1.754 to 2.70)	2.239	1.436	1.436	1.006
ischaemic heart disease	45 Cigarettes Per Smoker Per Day	Both	remaie	5.574	5.574	5.574	4.182	4.182	3.016	3.016	2.227	2.227	1.647	1.647	1.229	1.229	1.008
ischaemic neart disease	56.2 Cigarettes Per Smoker Per Day	BOTH	ware	13.641	13.641	13.641	8.813	(2.877 to 5.614) 8.813	(2.218 (0 3.950)	(2.218 (0 3.956)	(1.65 (0 2.849)	(1.65 (0 2.849)	(1.280 to 2.018)	(1.280 to 2.018)	(1.022 (0 1.441)	(1.022 (0 1.441)	(1.0 to 1.05)
Ischaemic heart disease	56.2 Cigarettes Per Smoker Per Day	Both	Female	22.811)	22.811)	22.811)	14.032)	14.032)	(3.445 to 8.338)	(3.445 to 8.338)	(2.602 to 5.039)	(2.602 to 5.039)	(1.788 to 3.044)	(1.788 to 3.044)	(1.33 to 1.849)	(1.33 to 1.849)	(1.0 to 1.077)
lschaemic stroke	0 Cigarettes Per Smoker Per Day	Both	Male	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)
lschaemic stroke	0 Cigarettes Per Smoker Per Day	Both	Female	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0) 2 363	(1.0 to 1.0) 2 112	(1.0 to 1.0)	(1.0 to 1.0) 1 153	(1.0 to 1.0) 1 153	(1.0 to 1.0)						
Ischaemic stroke	10 Cigarettes Per Smoker Per Day	Both	Male	(1.976 to 2.74) 2.334	(1.976 to 2.74) 2.334	(1.976 to 2.74) 2.334	(1.84 to 2.402) 2.101	(1.84 to 2.402) 2.101	(1.639 to 2.061) 1.916	(1.639 to 2.061) 1.916	(1.461 to 1.755)	(1.461 to 1.755) 1.721	(1.289 to 1.48) 1.479	(1.289 to 1.48) 1.479	(1.101 to 1.204) 1.204	(1.101 to 1.204) 1.204	(1.0 to 1.041) 1.007
Ischaemic stroke	10 Cigarettes Per Smoker Per Day	Both	Female	(1.689 to 3.166) 2.319	(1.689 to 3.166 2.319	i) (1.689 to 3.166) 2.319) (1.605 to 2.723) 1.971	(1.605 to 2.723) 1.971	(1.509 to 2.33) 1.662	(1.509 to 2.33) 1.662	(1.435 to 2.014) 1.415	(1.435 to 2.014) 1.415	(1.317 to 1.639) 1.222	(1.317 to 1.639) 1.222	(1.116 to 1.301) 1.08	(1.116 to 1.301) 1.08	(1.0 to 1.027) 1.004
Ischaemic stroke	20 Cigarettes Per Smoker Per Day	Both	Male	(1.923 to 2.747) 3.737	(1.923 to 2.747 3.737	(1.923 to 2.747) 3.737) (1.703 to 2.277) 3.04	(1.703 to 2.277) 3.04	(1.454 to 1.884) 2.443	(1.454 to 1.884) 2.443	(1.285 to 1.56) 1.94	(1.285 to 1.56) 1.94	(1.143 to 1.307) 1.527	(1.143 to 1.307) 1.527	(1.045 to 1.119) 1.194	(1.045 to 1.119) 1.194	(1.0 to 1.016) 1.003
Ischaemic stroke	20 Cigarettes Per Smoker Per Day	Both	Female	(2.688 to 4.979) 3.178	(2.688 to 4.979 3.178	(2.688 to 4.979) 3.178) (2.299 to 3.938) 2.666	(2.299 to 3.938) 2.666	(1.876 to 3.059) 2.224	(1.876 to 3.059) 2.224	(1.598 to 2.277) 1.849	(1.598 to 2.277) 1.849	(1.326 to 1.723) 1.518	(1.326 to 1.723) 1.518	(1.127 to 1.274) 1.209	(1.127 to 1.274) 1.209	(1.0 to 1.014) 1.009
Ischaemic stroke	30 Cigarettes Per Smoker Per Day	Both	Male	(2.653 to 3.77) 5.491	(2.653 to 3.77) 5.491	(2.653 to 3.77) 5.491	(2.265 to 3.069) 4.206	(2.265 to 3.069) 4.206	(1.945 to 2.515) 3.136	(1.945 to 2.515) 3.136	(1.663 to 2.037) 2.3	(1.663 to 2.037) 2.3	(1.397 to 1.64) 1.721	(1.397 to 1.64) 1.721	(1.15 to 1.263) 1.292	(1.15 to 1.263) 1.292	(1.0 to 1.034) 1.007
Ischaemic stroke	30 Cigarettes Per Smoker Per Day	Both	Female	(3.813 to 7.703) 3.431	(3.813 to 7.703 3.431	(3.813 to 7.703) 3.431	(2.989 to 5.642) 2.734	(2.989 to 5.642) 2.734	(2.326 to 4.018) 2.117	(2.326 to 4.018) 2.117	(1.87 to 2.815) 1.659	(1.87 to 2.815) 1.659	(1.474 to 2.006) 1.337	(1.474 to 2.006) 1.337	(1.172 to 1.413) 1.119	(1.172 to 1.413) 1.119	(1.0 to 1.025) 1.011
Ischaemic stroke	40 Cigarettes Per Smoker Per Day	Both	Male	(2.454 to 4.606) 7.127	(2.454 to 4.606 7.127	i) (2.454 to 4.606) 7.127) (2.011 to 3.596)	(2.011 to 3.596)	(1.619 to 2.739)	(1.619 to 2.739)	(1.313 to 2.061)	(1.313 to 2.061)	(1.123 to 1.585)	(1.123 to 1.585)	(1.021 to 1.22)	(1.021 to 1.22)	(1.0 to 1.038)
Ischaemic stroke	40 Cigarettes Per Smoker Per Day	Both	Female	(4.104 to 11.541)	(4.104 to 11.541)	(4.104 to 11.541)	5.103 (3.045 to 7.724)	5.103 (3.045 to 7.724)	3.636 (2.299 to 5.219)	3.636 (2.299 to 5.219)	2.593 (1.808 to 3.589)	2.593 (1.808 to 3.589)	1.781 (1.325 to 2.376)	1.781 (1.325 to 2.376)	1.209 (1.0 to 1.43)	1.209 (1.0 to 1.43)	1.008 (1.0 to 1.036)

Appendix Table 6d. Relative risks used by age and sex for each ou	atcome for smoking globally.																
Risk - Outcome	Category / Units	Morbidity / Mortality	Sex	30 to 34	35 to 39	40 to 44	45 to 49	50 to 54	55 to 59	A 60 to 64	.ge 65 to 69	70 to 74	75 to 79	80 to 84	85 to 89	90 to 94	95 plus
Smoking																	
lschaemic stroke	50 Cigarettes Per Smoker Per Day	Both	Male	4.456 (2.943 to 6.299) 9.851	4.456 (2.943 to 6.299) 9.851	4.456 (2.943 to 6.299 9.851	3.491 (2.396 to 4.743) 6.922	3.491 (2.396 to 4.743) 6.922	2.659 (1.915 to 3.56)	2.659 (1.915 to 3.56)	2.048 (1.526 to 2.611)	2.048 (1.526 to 2.611)	1.579 (1.203 to 1.976)	1.579 (1.203 to 1.976)	1.211 (1.0 to 1.464)	1.211 (1.0 to 1.464)	1.035 (1.0 to 1.144)
lschaemic stroke	50 Cigarettes Per Smoker Per Day	Both	Female	(5.013 to 17.851) 5.317	(5.013 to 17.851) 5.317	(5.013 to 17.851) 5.317	(3.959 to 11.353) 4 155	(3.959 to 11.353) 4.155	5.0 (3.058 to 7.745) 3 175	5.0 (3.058 to 7.745) 3 175	3.569 (2.397 to 4.973) 2 395	3.569 (2.397 to 4.973) 2 395	2.352 (1.697 to 3.091) 1.844	2.352 (1.697 to 3.091) 1.844	1.453 (1.136 to 1.809) 1 385	1.453) (1.136 to 1.809) 1 385	1.026 (1.0 to 1.112) 1 101
Ischaemic stroke	60 Cigarettes Per Smoker Per Day	Both	Male	(2.793 to 8.673)	(2.793 to 8.673)	(2.793 to 8.673	(2.304 to 6.585)	(2.304 to 6.585)	(1.804 to 4.855)	(1.804 to 4.855)	(1.464 to 3.507)	(1.464 to 3.507)	(1.18 to 2.676)	(1.18 to 2.676)	(1.0 to 1.885)	(1.0 to 1.885)	(1.0 to 1.366)
Ischaemic stroke	60 Cigarettes Per Smoker Per Day	Both	Female	11.722 (4.845 to 23.87)	11.722 (4.845 to 23.87)	11.722 (4.845 to 23.87	(3.608 to) 16.396)	(3.608 to 16.396)	5.942 (3.01 to 11.062)	5.942 (3.01 to 11.062)	4.286 (2.319 to 7.079)	4.286 (2.319 to 7.079)	2.783 (1.643 to 4.284)	2.783 (1.643 to 4.284)	1.689 (1.093 to 2.378)	1.689) (1.093 to 2.378)	1.043 (1.0 to 1.234)
Intracerebral hemorrhage	0 Cigarettes Per Smoker Per Day	Both	Male	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)				
Intracerebral hemorrhage	0 Cigarettes Per Smoker Per Day	Both	Female	(1.0 to 1.0) 2.363	(1.0 to 1.0) 2.363	(1.0 to 1.0) 2.363	(1.0 to 1.0) 2.112	(1.0 to 1.0) 2.112	(1.0 to 1.0) 1.843	(1.0 to 1.0) 1.843	(1.0 to 1.0) 1.606	(1.0 to 1.0) 1.606	(1.0 to 1.0) 1.386	(1.0 to 1.0) 1.386	(1.0 to 1.0) 1.153	(1.0 to 1.0) 1.153	(1.0 to 1.0) 1.011
Intracerebral hemorrhage	10 Cigarettes Per Smoker Per Day	Both	Male	(1.976 to 2.74) 2.334	(1.976 to 2.74) 2.334	(1.976 to 2.74) 2.334	(1.84 to 2.402) 2.101	(1.84 to 2.402) 2.101	(1.639 to 2.061) 1.916	(1.639 to 2.061) 1.916	(1.461 to 1.755) 1.721	(1.461 to 1.755) 1.721	(1.289 to 1.48) 1.479	(1.289 to 1.48) 1.479	(1.101 to 1.204) 1.204	(1.101 to 1.204) 1.204	(1.0 to 1.041) 1.007
Intracerebral hemorrhage	10 Cigarettes Per Smoker Per Day	Both	Female	(1.689 to 3.166) 2.319	(1.689 to 3.166) 2.319	(1.689 to 3.166 2.319	(1.605 to 2.723) 1.971	(1.605 to 2.723) 1.971	(1.509 to 2.33) 1.662	(1.509 to 2.33) 1.662	(1.435 to 2.014) 1.415	(1.435 to 2.014) 1.415	(1.317 to 1.639) 1.222	(1.317 to 1.639) 1.222	(1.116 to 1.301) 1.08	(1.116 to 1.301) 1.08	(1.0 to 1.027) 1.004
Intracerebral hemorrhage	20 Cigarettes Per Smoker Per Day	Both	Male	(1.923 to 2.747) 3.737	(1.923 to 2.747) 3.737	(1.923 to 2.747 3.737	(1.703 to 2.277) 3.04	(1.703 to 2.277) 3.04	(1.454 to 1.884) 2.443	(1.454 to 1.884) 2.443	(1.285 to 1.56) 1.94	(1.285 to 1.56) 1.94	(1.143 to 1.307) 1.527	(1.143 to 1.307) 1.527	(1.045 to 1.119) 1.194	(1.045 to 1.119) 1.194	(1.0 to 1.016) 1.003
Intracerebral hemorrhage	20 Cigarettes Per Smoker Per Day	Both	Female	(2.688 to 4.979) 3.178	(2.688 to 4.979) 3.178	(2.688 to 4.979 3.178	(2.299 to 3.938) 2.666	(2.299 to 3.938) 2.666	(1.876 to 3.059) 2.224	(1.876 to 3.059) 2.224	(1.598 to 2.277) 1.849	(1.598 to 2.277) 1.849	(1.326 to 1.723)	(1.326 to 1.723)	(1.127 to 1.274) 1.209	(1.127 to 1.274) 1.209	(1.0 to 1.014) 1.009
Intracerebral hemorrhage	30 Cigarettes Per Smoker Per Day	Both	Male	(2.653 to 3.77) 5.491	(2.653 to 3.77) 5.491	(2.653 to 3.77) 5.491	(2.265 to 3.069) 4.206	(2.265 to 3.069) 4.206	(1.945 to 2.515) 3.136	(1.945 to 2.515) 3.136	(1.663 to 2.037) 2.3	(1.663 to 2.037) 2.3	(1.397 to 1.64) 1.721	(1.397 to 1.64) 1.721	(1.15 to 1.263) 1.292	(1.15 to 1.263) 1.292	(1.0 to 1.034) 1.007
Intracerebral hemorrhage	30 Cigarettes Per Smoker Per Day	Both	Female	(3.813 to 7.703) 3.431	(3.813 to 7.703) 3.431	(3.813 to 7.703 3.431	(2.989 to 5.642) 2.734	(2.989 to 5.642) 2.734	(2.326 to 4.018) 2.117	(2.326 to 4.018) 2.117	(1.87 to 2.815) 1.659	(1.87 to 2.815) 1.659	(1.474 to 2.006) 1.337	(1.474 to 2.006) 1.337	(1.172 to 1.413) 1.119	(1.172 to 1.413) 1.119	(1.0 to 1.025) 1.011
Intracerebral hemorrhage	40 Cigarettes Per Smoker Per Day	Both	Male	(2.454 to 4.606) 7.127	(2.454 to 4.606) 7.127	(2.454 to 4.606 7.127	(2.011 to 3.596)	(2.011 to 3.596)	(1.619 to 2.739)	(1.619 to 2.739)	(1.313 to 2.061)	(1.313 to 2.061)	(1.123 to 1.585)	(1.123 to 1.585)	(1.021 to 1.22)	(1.021 to 1.22)	(1.0 to 1.038)
Intracerebral hemorrhage	40 Cigarettes Per Smoker Per Day	Both	Female	(4.104 to 11.541)	(4.104 to 11.541)	(4.104 to 11.541)	5.103 (3.045 to 7.724)	5.103 (3.045 to 7.724)	3.636 (2.299 to 5.219)	3.636 (2.299 to 5.219)	2.593 (1.808 to 3.589)	2.593 (1.808 to 3.589)	1.781 (1.325 to 2.376)	1.781 (1.325 to 2.376)	1.209 (1.0 to 1.43)	1.209 (1.0 to 1.43)	1.008 (1.0 to 1.036)
Intracerebral hemorrhage	50 Cigarettes Per Smoker Per Day	Both	Male	4.456 (2.943 to 6.299) 9.851	4.456 (2.943 to 6.299) 9.851	4.456 (2.943 to 6.299) 9.851	3.491 (2.396 to 4.743) 6.922	3.491 (2.396 to 4.743) 6.922	2.659 (1.915 to 3.56)	2.659 (1.915 to 3.56)	2.048 (1.526 to 2.611)	2.048 (1.526 to 2.611)	1.579 (1.203 to 1.976)	1.579 (1.203 to 1.976)	1.211 (1.0 to 1.464)	1.211 (1.0 to 1.464)	1.035 (1.0 to 1.144)
Intracerebral hemorrhage	50 Cigarettes Per Smoker Per Day	Both	Female	(5.013 to 17.851) 5.317	(5.013 to 17.851) 5.317	(5.013 to 17.851) 5.317	(3.959 to 11.353) 4 155	(3.959 to 11.353) 4.155	5.0 (3.058 to 7.745) 3 175	5.0 (3.058 to 7.745) 3 175	3.569 (2.397 to 4.973) 2 395	3.569 (2.397 to 4.973) 2 395	2.352 (1.697 to 3.091) 1.844	2.352 (1.697 to 3.091) 1.844	1.453 (1.136 to 1.809) 1 385	1.453) (1.136 to 1.809) 1 385	1.026 (1.0 to 1.112) 1 101
Intracerebral hemorrhage	60 Cigarettes Per Smoker Per Day	Both	Male	(2.793 to 8.673)	(2.793 to 8.673)	(2.793 to 8.673	(2.304 to 6.585)	(2.304 to 6.585)	(1.804 to 4.855)	(1.804 to 4.855)	(1.464 to 3.507)	(1.464 to 3.507)	(1.18 to 2.676)	(1.18 to 2.676)	(1.0 to 1.885)	(1.0 to 1.885)	(1.0 to 1.366)
Intracerebral hemorrhage	60 Cigarettes Per Smoker Per Day	Both	Female	11.722 (4.845 to 23.87) 1.0	11.722 (4.845 to 23.87) 1.0	11.722 (4.845 to 23.87	(3.608 to 16.396)	(3.608 to 16.396)	5.942 (3.01 to 11.062)	5.942 (3.01 to 11.062)	4.286 (2.319 to 7.079) 1.0	4.286 (2.319 to 7.079)	2.783 (1.643 to 4.284)	2.783 (1.643 to 4.284)	1.689 (1.093 to 2.378) 1.0	1.689) (1.093 to 2.378) 1.0	1.043 (1.0 to 1.234)
Subarachnoid hemorrhage	0 Cigarettes Per Smoker Per Day	Both	Male	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)				
Subarachnoid hemorrhage	0 Cigarettes Per Smoker Per Day	Both	Female	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0) 1 153	(1.0 to 1.0)	(1.0 to 1.0)				
Subarachnoid hemorrhage	10 Cigarettes Per Smoker Per Day	Both	Male	(1.976 to 2.74) 2.334	(1.976 to 2.74) 2.334	(1.976 to 2.74) 2.334	(1.84 to 2.402) 2.101	(1.84 to 2.402) 2.101	(1.639 to 2.061) 1.916	(1.639 to 2.061) 1.916	(1.461 to 1.755) 1.721	(1.461 to 1.755) 1.721	(1.289 to 1.48) 1.479	(1.289 to 1.48) 1.479	(1.101 to 1.204) 1.204	(1.101 to 1.204) 1.204	(1.0 to 1.041) 1.007
Subarachnoid hemorrhage	10 Cigarettes Per Smoker Per Day	Both	Female	(1.689 to 3.166) 2.319	(1.689 to 3.166) 2.319	(1.689 to 3.166 2.319	(1.605 to 2.723) 1.971	(1.605 to 2.723) 1.971	(1.509 to 2.33) 1.662	(1.509 to 2.33) 1.662	(1.435 to 2.014) 1.415	(1.435 to 2.014) 1.415	(1.317 to 1.639) 1.222	(1.317 to 1.639) 1.222	(1.116 to 1.301) 1.08	(1.116 to 1.301) 1.08	(1.0 to 1.027) 1.004
Subarachnoid hemorrhage	20 Cigarettes Per Smoker Per Day	Both	Male	(1.923 to 2.747) 3.737	(1.923 to 2.747) 3.737	(1.923 to 2.747 3.737	(1.703 to 2.277) 3.04	(1.703 to 2.277) 3.04	(1.454 to 1.884) 2.443	(1.454 to 1.884) 2.443	(1.285 to 1.56) 1.94	(1.285 to 1.56) 1.94	(1.143 to 1.307) 1.527	(1.143 to 1.307) 1.527	(1.045 to 1.119) 1.194	(1.045 to 1.119) 1.194	(1.0 to 1.016) 1.003
Subarachnoid hemorrhage	20 Cigarettes Per Smoker Per Day	Both	Female	(2.688 to 4.979) 3.178	(2.688 to 4.979) 3.178	(2.688 to 4.979 3.178	(2.299 to 3.938) 2.666	(2.299 to 3.938) 2.666	(1.876 to 3.059) 2.224	(1.876 to 3.059) 2.224	(1.598 to 2.277) 1.849	(1.598 to 2.277) 1.849	(1.326 to 1.723) 1.518	(1.326 to 1.723) 1.518	(1.127 to 1.274) 1.209	(1.127 to 1.274) 1.209	(1.0 to 1.014) 1.009
Subarachnoid hemorrhage	30 Cigarettes Per Smoker Per Day	Both	Male	(2.653 to 3.77) 5.491	(2.653 to 3.77) 5.491	(2.653 to 3.77) 5.491	(2.265 to 3.069) 4.206	(2.265 to 3.069) 4.206	(1.945 to 2.515) 3.136	(1.945 to 2.515) 3.136	(1.663 to 2.037) 2.3	(1.663 to 2.037) 2.3	(1.397 to 1.64) 1.721	(1.397 to 1.64) 1.721	(1.15 to 1.263) 1.292	(1.15 to 1.263) 1.292	(1.0 to 1.034) 1.007
Subarachnoid hemorrhage	30 Cigarettes Per Smoker Per Day	Both	Female	(3.813 to 7.703) 3.431	(3.813 to 7.703) 3.431	(3.813 to 7.703) 3.431	(2.989 to 5.642) 2.734	(2.989 to 5.642) 2.734	(2.326 to 4.018) 2.117	(2.326 to 4.018) 2.117	(1.87 to 2.815) 1.659	(1.87 to 2.815) 1.659	(1.474 to 2.006) 1.337	(1.474 to 2.006) 1.337	(1.172 to 1.413) 1.119	(1.172 to 1.413) 1.119	(1.0 to 1.025) 1.011
Subarachnoid hemorrhage	40 Cigarettes Per Smoker Per Day	Both	Male	(2.454 to 4.606) 7.127	(2.454 to 4.606) 7.127	(2.454 to 4.606 7.127	(2.011 to 3.596)	(2.011 to 3.596)	(1.619 to 2.739)	(1.619 to 2.739)	(1.313 to 2.061)	(1.313 to 2.061)	(1.123 to 1.585)	(1.123 to 1.585)	(1.021 to 1.22)	(1.021 to 1.22)	(1.0 to 1.038)
Subarachnoid hemorrhage	40 Cigarettes Per Smoker Per Day	Both	Female	(4.104 to 11.541)	(4.104 to 11.541)	(4.104 to 11.541)	5.103 (3.045 to 7.724)	5.103 (3.045 to 7.724)	3.636 (2.299 to 5.219)	3.636 (2.299 to 5.219)	2.593 (1.808 to 3.589)	2.593 (1.808 to 3.589)	1.781 (1.325 to 2.376)	1.781 (1.325 to 2.376)	1.209 (1.0 to 1.43)	1.209 (1.0 to 1.43)	1.008 (1.0 to 1.036)
Subarachnoid hemorrhage	50 Cigarettes Per Smoker Per Day	Both	Male	4.456 (2.943 to 6.299) 9.851	4.456 (2.943 to 6.299) 9.851	4.456 (2.943 to 6.299) 9.851	3.491 (2.396 to 4.743) 6 922	3.491 (2.396 to 4.743) 6 922	(1.915 to 3.56)	(1.915 to 3.56)	2.048 (1.526 to 2.611)	(1.526 to 2.611)	(1.203 to 1.976)	(1.203 to 1.976)	1.211 (1.0 to 1.464)	(1.0 to 1.464)	(1.0 to 1.144)
Subarachnoid hemorrhage	50 Cigarettes Per Smoker Per Day	Both	Female	(5.013 to 17.851)	(5.013 to 17.851)	(5.013 to 17.851)	(3.959 to 11.353)	(3.959 to 11.353)	5.0 (3.058 to 7.745)	5.0 (3.058 to 7.745)	3.569 (2.397 to 4.973)	3.569 (2.397 to 4.973)	2.352 (1.697 to 3.091)	2.352 (1.697 to 3.091)	1.453 (1.136 to 1.809)	1.453) (1.136 to 1.809)	1.026 (1.0 to 1.112)
Subarachnoid hemorrhage	60 Cigarettes Per Smoker Per Day	Both	Male	5.317 (2.793 to 8.673)	5.317 (2.793 to 8.673)	5.317 (2.793 to 8.673	4.155 (2.304 to 6.585)	4.155 (2.304 to 6.585)	3.175 (1.804 to 4.855)	3.175 (1.804 to 4.855)	2.395 (1.464 to 3.507)	2.395 (1.464 to 3.507)	1.844 (1.18 to 2.676)	1.844 (1.18 to 2.676)	1.385 (1.0 to 1.885)	1.385 (1.0 to 1.885)	1.101 (1.0 to 1.366)
Subarachnoid hemorrhage	60 Cigarettes Per Smoker Per Day	Both	Female	11.722 (4.845 to 23.87)	11.722 (4.845 to 23.87)	11.722 (4.845 to 23.87	8.28 (3.608 to 16.396)	8.28 (3.608 to 16.396)	5.942 (3.01 to 11.062)	5.942 (3.01 to 11.062)	4.286 (2.319 to 7.079)	4.286 (2.319 to 7.079)	2.783 (1.643 to 4.284)	2.783 (1.643 to 4.284)	1.689 (1.093 to 2.378)	1.689) (1.093 to 2.378)	1.043 (1.0 to 1.234)
Atrial fibrillation and flutter	0 Cigarettes Per Smoker Per Day	Both	Male	(1.0 to 1.0)	(1.0 to 1.0)	1.0 (1.0 to 1.0)	(1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)

Appendix Table 6d. Relative risks used by age and sex for each out	tcome for smoking globally.																
Risk - Outcome	Category / Units	Morbidity / Mortality	Sex	30 to 34	35 to 39	40 to 44	45 to 49	50 to 54	55 to 59	A 60 to 64	e 65 to 69	70 to 74	75 to 79	80 to 84	85 to 89	90 to 94	95 plus
Smoking																	
Atrial fibrillation and flutter	0 Cigarettes Per Smoker Per Day	Both	Female	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Atrial fibrillation and flutter	6.3 Cigarettes Per Smoker Per Day	Both	Male	1.532 (1.044 to 2.151)	1.532 (1.044 to 2.151)	1.532 (1.044 to 2.151)	1.507 (1.048 to 2.051)	1.507 (1.048 to 2.051)	1.474 (1.035 to 1.935)	1.474 (1.035 to 1.935)	1.429 (1.039 to 1.787)	1.429 (1.039 to 1.787)	1.324 (1.052 to 1.59)	1.324 (1.052 to 1.59)	1.182 (1.0 to 1.358)	1.182 (1.0 to 1.358)	1.008 (1.0 to 1.047)
Atrial fibrillation and flutter	6.3 Cigarettes Per Smoker Per Day	Both	Female	1.532 (1.044 to 2.151)	1.532 (1.044 to 2.151)	1.532 (1.044 to 2.151)	1.507 (1.048 to 2.051)	1.507 (1.048 to 2.051)	1.474 (1.035 to 1.935)	1.474 (1.035 to 1.935)	1.429 (1.039 to 1.787)	1.429 (1.039 to 1.787)	1.324 (1.052 to 1.59)	1.324 (1.052 to 1.59)	1.182 (1.0 to 1.358)	1.182 (1.0 to 1.358)	1.008 (1.0 to 1.047)
Atrial fibrillation and flutter	12.6 Cigarettes Per Smoker Per Day	Both	Male	1.895 (1.239 to 2.794)	1.895 (1.239 to 2.794)	1.895 (1.239 to 2.794)	1.774 (1.21 to 2.55)	1.774 (1.21 to 2.55)	1.632 (1.154 to 2.241)	1.632 (1.154 to 2.241)	1.459 (1.101 to 1.896)	1.459 (1.101 to 1.896)	1.283 (1.024 to 1.607)	1.283 (1.024 to 1.607)	1.129 (1.0 to 1.32)	1.129 (1.0 to 1.32)	1.021 (1.0 to 1.111)
Atrial fibrillation and flutter	12.6 Cigarettes Per Smoker Per Day	Both	Female	1.895 (1.239 to 2.794)	1.895 (1.239 to 2.794)	1.895	1.774 (1.21 to 2.55)	1.774 (1.21 to 2.55)	1.632 (1.154 to 2.241)	1.632 (1.154 to 2.241)	1.459 (1.101 to 1.895)	1.459 (1.101 to 1.895)	1.283	1.283 (1.024 to 1.607)	1.129 (1.0 to 1.32)	1.129 (1.0 to 1.32)	1.021 (1.0 to 1.111)
Atrial fibrillation and fluttor	18.0 Cigarettes Per Smoker Per Day	Both	Mala	2.222	2.222 (1.42 to 2.234)	2.222	1.997	1.997	1.766	1.766	1.518	1.518 (1.122 to 2.048)	1.299 (1.002 to 1.660)	1.299 (1.002 to 1.660)	1.124	1.124	1.028
	10.5 Cigarettes Per Sinoker Per Day	Both	iviale	2.222	2.222	2.222	1.997	1.997	1.766	1.766	1.518	1.518	1.299	1.299	1.124	1.124	1.028
	18.9 Cigarettes Per Smoker Per Day	Both	remale	2.588	2.588	2.588	2.261	2.261	1.952	1.952	1.629	1.629	1.37	1.37	1.157	1.157	1.012
Atrial fibrillation and flutter	25.2 Cigarettes Per Smoker Per Day	Both	Male	(1.542 to 4.076) 2.588	(1.542 to 4.076) 2.588	(1.542 to 4.076) 2.588	(1.451 to 3.404) 2.261	(1.451 to 3.404) 2.261	(1.347 to 2.865) 1.952	(1.347 to 2.865) 1.952	(1.227 to 2.188) 1.629	(1.227 to 2.188) 1.629	(1.104 to 1.74) 1.37	(1.104 to 1.74) 1.37	(1.0 to 1.366) 1.157	(1.0 to 1.366) 1.157	(1.0 to 1.09) 1.012
Atrial fibrillation and flutter	25.2 Cigarettes Per Smoker Per Day	Both	Female	(1.542 to 4.076) 1.0	(1.542 to 4.076) 1.0	(1.542 to 4.076) 1.0	(1.451 to 3.404) 1.0	(1.451 to 3.404) 1.0	(1.347 to 2.865) 1.0	(1.347 to 2.865) 1.0	(1.227 to 2.188) 1.0	(1.227 to 2.188) 1.0	(1.104 to 1.74) 1.0	(1.104 to 1.74) 1.0	(1.0 to 1.366) 1.0	(1.0 to 1.366) 1.0	(1.0 to 1.09) 1.0
Aortic aneurysm	0 Cigarettes Per Smoker Per Day	Both	Male	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0
Aortic aneurysm	0 Cigarettes Per Smoker Per Day	Both	Female	(1.0 to 1.0) 5.046	(1.0 to 1.0) 5.046	(1.0 to 1.0) 5.046	(1.0 to 1.0) 4.201	(1.0 to 1.0) 4.201	(1.0 to 1.0) 3.404	(1.0 to 1.0) 3.404	(1.0 to 1.0) 2.669	(1.0 to 1.0) 2.669	(1.0 to 1.0) 2.014	(1.0 to 1.0) 2.014	(1.0 to 1.0) 1.441	(1.0 to 1.0) 1.441	(1.0 to 1.0) 1.005
Aortic aneurysm	12 Cigarettes Per Smoker Per Day	Both	Male	(3.685 to 6.516) 5.046	(3.685 to 6.516) 5.046	(3.685 to 6.516) 5.046	(3.207 to 5.298) 4.201	(3.207 to 5.298) 4.201	(2.713 to 4.162) 3.404	(2.713 to 4.162) 3.404	(2.263 to 3.044) 2.669	(2.263 to 3.044) 2.669	(1.79 to 2.242) 2.014	(1.79 to 2.242) 2.014	(1.327 to 1.563) 1.441	(1.327 to 1.563) 1.441	(1.0 to 1.03) 1.005
Aortic aneurysm	12 Cigarettes Per Smoker Per Day	Both	Female	(3.685 to 6.516) 9.532	(3.685 to 6.516) 9.532	(3.685 to 6.516) 9.532	(3.207 to 5.298) 6.722	(3.207 to 5.298) 6.722	(2.713 to 4.162) 4.62	(2.713 to 4.162) 4.62	(2.263 to 3.044) 3.116	(2.263 to 3.044) 3.116	(1.79 to 2.242) 2.076	(1.79 to 2.242) 2.076	(1.327 to 1.563)	(1.327 to 1.563) 1.387	(1.0 to 1.03) 1.004
Aortic aneurysm	24 Cigarettes Per Smoker Per Day	Both	Male	(6.977 to 12.48)	(6.977 to 12.48)	(6.977 to 12.48)	(5.101 to 8.596)	(5.101 to 8.596)	(3.71 to 5.762)	(3.71 to 5.762)	(2.577 to 3.724)	(2.577 to 3.724)	(1.797 to 2.395)	(1.797 to 2.395)	(1.242 to 1.547)	(1.242 to 1.547)	(1.0 to 1.021)
Aortic aneurysm	24 Cigarettes Per Smoker Per Day	Both	Female	(6.977 to 12.48)	(6.977 to 12.48)	(6.977 to 12.48)	(5.101 to 8.596)	(5.101 to 8.596)	(3.71 to 5.762)	(3.71 to 5.762)	(2.577 to 3.724)	(2.577 to 3.724)	(1.797 to 2.395)	(1.797 to 2.395)	(1.242 to 1.547)	(1.242 to 1.547)	(1.0 to 1.021)
				(7.774 to	(7.774 to	(7.774 to	(5.659 to	(5.659 to	5.626	5.626	3.697	3.697	2.455	2.455	1.593	1.593	1.006
Aortic aneurysm	36 Cigarettes Per Smoker Per Day	Both	Male	18.221) 12.382	18.221) 12.382	18.221) 12.382	11.932) 8.505	11.932) 8.505	(3.935 to 7.702)	(3.935 to 7.702)	(2.779 to 4.795)	(2.779 to 4.795)	(1.897 to 3.038)	(1.897 to 3.038)	(1.306 to 1.875)	(1.306 to 1.875)	(1.0 to 1.03)
Aortic aneurysm	36 Cigarettes Per Smoker Per Day	Both	Female	(7.774 to 18.221)	(7.774 to 18.221)	(7.774 to 18.221)	(5.659 to 11.932)	(5.659 to 11.932)	5.626 (3.935 to 7.702)	5.626 (3.935 to 7.702)	3.697 (2.779 to 4.795)	3.697 (2.779 to 4.795)	2.455 (1.897 to 3.038)	2.455 (1.897 to 3.038)	1.593 (1.306 to 1.875)	1.593 (1.306 to 1.875)	1.006 (1.0 to 1.03)
				14.479	14.479	14.479	9.77 (5.928 to	9.77 (5.928 to	6.258	6.258	3.954	3.954	2.503	2.503	1.562	1.562	1.034
Aortic aneurysm	48 Cigarettes Per Smoker Per Day	Both	Male	(7.446 to 24.32)	(7.446 to 24.32)	(7.446 to 24.32)	15.073) 9.77	15.073) 9.77	(4.093 to 9.24)	(4.093 to 9.24)	(2.756 to 5.443)	(2.756 to 5.443)	(1.879 to 3.295)	(1.879 to 3.295)	(1.216 to 1.946)	(1.216 to 1.946)	(1.0 to 1.156)
Aortic aneurysm	48 Cigarettes Per Smoker Per Day	Both	Female	14.479 (7.446 to 24.32)	14.479 (7.446 to 24.32)	14.479 (7.446 to 24.32)	(5.928 to 15.073)	(5.928 to 15.073)	6.258 (4.093 to 9.24)	6.258 (4.093 to 9.24)	3.954 (2.756 to 5.443)	3.954 (2.756 to 5.443)	2.503 (1.879 to 3.295)	2.503 (1.879 to 3.295)	1.562 (1.216 to 1.946)	1.562 (1.216 to 1.946)	1.034 (1.0 to 1.156)
Perinheral vascular disease	0 Cigarettes Per Smoker Per Day	Both	Male	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Perinheral vascular disease	O Cigarettes Per Smoker Per Dav	Both	Female	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Parinharal vascular disease	0 2 Circretter Per Smeker Per Day	Both	Mala	3.033 (1.979 to 4.598)	3.033 (1.979 to 4.598)	3.033 (1.979 to 4.598)	2.78 (1.756 to 4.24)	2.78 (1.756 to 4.24)	3.077 (1 721 to 4 737)	3.077 (1 721 to 4 737)	2.978 (2.177 to 3.749)	2.978	2.194 (1.724 to 2.64)	2.194 (1.724 to 2.64)	1.535 (1.238 to 1.832)	1.535	1.042
	9.5 Cigarettes Per Sinoker Per Day	Both	iviale	3.033	3.033	3.033	2.78	2.78	3.077	3.077	2.978	2.978	2.194	2.194	1.535	1.535	1.042
Peripheral vascular disease	9.3 Ligarettes Per Smoker Per Day	BOTH	Female	6.374	6.374	6.374	5.44	5.44	5.689	5.689	4.868	4.868	3.01	3.01	1.75	1.75	1.033
Peripheral vascular disease	18.6 Cigarettes Per Smoker Per Day	Both	Male	(3.686 to 10.26) 6.374	(3.686 to 10.26) 6.374	(3.686 to 10.26) 6.374	(3.032 to 8.936) 5.44	(3.032 to 8.936) 5.44	(2.725 to 8.816) 5.689	(2.725 to 8.816) 5.689	(3.556 to 6.076) 4.868	(3.556 to 6.076) 4.868	(2.439 to 3.56) 3.01	(2.439 to 3.56) 3.01	(1.446 to 2.077 1.75	1.75 1.75	(1.0 to 1.167) 1.033
Peripheral vascular disease	18.6 Cigarettes Per Smoker Per Day	Both	Female	(3.686 to 10.26)	(3.686 to 10.26)	(3.686 to 10.26)	(3.032 to 8.936) 7.43	(3.032 to 8.936) 7.43	(2.725 to 8.816) 7.069	(2.725 to 8.816) 7.069	(3.556 to 6.076)	(3.556 to 6.076)	(2.439 to 3.56)	(2.439 to 3.56)	(1.446 to 2.077)) (1.446 to 2.077)	(1.0 to 1.167)
Peripheral vascular disease	27.9 Cigarettes Per Smoker Per Day	Both	Male	8.997 (4.62 to 15.851)	8.997 (4.62 to 15.851)	8.997 (4.62 to 15.851)	(3.684 to 13.161)	(3.684 to 13.161)	(3.105 to 11.181)	(3.105 to 11.181)	5.123 (3.758 to 6.66)	5.123 (3.758 to 6.66)	2.887 (2.243 to 3.712)	2.887 (2.243 to 3.712)	1.651 (1.297 to 2.064)	1.651 (1.297 to 2.064)	1.047 (1.0 to 1.217)
				8.997	8.997	8.997	7.43 (3.684 to	7.43 (3.684 to	7.069 (3.105 to	7.069 (3.105 to	5.123	5.123	2.887	2.887	1.651	1.651	1.047
Peripheral vascular disease	27.9 Cigarettes Per Smoker Per Day	Both	Female	(4.62 to 15.851) 10.787	(4.62 to 15.851) 10.787	(4.62 to 15.851) 10.787	13.161) 8.838	13.161) 8.838	11.181) 8.201	11.181) 8.201	(3.758 to 6.66)	(3.758 to 6.66)	(2.243 to 3.712)	(2.243 to 3.712)	(1.297 to 2.064)	(1.297 to 2.064)	(1.0 to 1.217)
Perinheral vascular disease	37.2 Cigarettes Per Smoker Per Day	Both	Male	(4.702 to 20.954)	(4.702 to 20.954)	(4.702 to 20.954)	(3.845 to 18.062)	(3.845 to 18.062)	(3.609 to 14.687)	(3.609 to 14.687)	5.967 (3.758 to 9.089)	5.967 (3.758 to 9.089)	3.335 (2.184 to 4.946)	3.335 (2.184 to 4.946)	1.894 (1.247 to 2.825)	1.894 (1.247 to 2.825)	1.16 (1.0 to 1.68)
				10.787 (4.702 to	10.787	10.787	8.838 (3.845 to	8.838 (3.845 to	8.201 (3.609 to	8.201 (3.609 to	5 967	5 967	3 335	3 335	1 894	1 894	1 16
Peripheral vascular disease	37.2 Cigarettes Per Smoker Per Day	Both	Female	20.954)	20.954)	20.954)	18.062)	18.062)	14.687)	14.687)	(3.758 to 9.089)	(3.758 to 9.089)	(2.184 to 4.946)	(2.184 to 4.946)	(1.247 to 2.825)	(1.247 to 2.825)	(1.0 to 1.68)
		Deth	Mala	11.975	11.975	11.975	(3.819 to	(3.819 to	(3.611 to	(3.611 to	6.528	6.528	3.629	3.629	2.061	2.061	1.25
Peripheral vascular disease	46.5 Ligarettes Per Smoker Per Day	Both	Male	(4.06 to 25.2/3)	(4.00 to 25.2/3)	(4.66 to 25.2/3)	9.778	9.778	9.03	9.03	(5.542 to 11.14)	(3.542 to 11.14)	(1.986 to 6.135)	(1.986 to 6.135)	(1.118 to 3.541)	(1.118 to 3.541)	(1.0 to 2.038)
Peripheral vascular disease	46.5 Cigarettes Per Smoker Per Day	Both	Female	11.975 (4.66 to 25.273)	11.975 (4.66 to 25.273)	11.975 (4.66 to 25.273)	(3.819 to 20.981)	(3.819 to 20.981)	(3.611 to 18.958)	(3.611 to 18.958)	6.528 (3.542 to 11.14)	6.528 (3.542 to 11.14)	3.629 (1.986 to 6.135)	3.629 (1.986 to 6.135)	2.061 (1.118 to 3.541)	2.061 (1.118 to 3.541)	1.25 (1.0 to 2.038)
Chronic obstructive pulmonary disease	0 Pack-Years	Both	Male	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)

Appendix Table 6d. Relative risks used by age and sex for each out	come for smoking globally.																
Risk - Outcome	Category / Units	Morbidity / Mortality	Sex	30 to 34	35 to 39	40 to 44	45 to 49	50 to 54	55 to 59	A;	ge 65 to 69	70 to 74	75 to 79	80 to 84	85 to 89	90 to 94	95 plus
Smoking				501054	55 10 55	401044	401040	50 10 54	55 66 57	001004	0.5 10 05	101074	151015	001004	0.0 10 0.0	701074	50 pius
Chronic obstructive pulmonary disease	0 Pack-Years	Both	Female	1.0 (1.0 to 1.0)													
Chronic obstructive pulmonary disease	10 Pack-Years	Both	Male	2.512 (1.692 to 3.297)	2.512) (1.692 to 3.297)												
Chronic obstructive pulmonary disease	10 Pack-Years	Both	Female	2.512 (1.692 to 3.297)	2.512) (1.692 to 3.297)												
Chronic obstructive pulmonary disease	20 Pack-Years	Both	Male	3.942 (3.079 to 4.753)													
	20 Pack Years	Both	Fomalo	3.942 (2.070 to 4.752)	3.942	3.942	3.942	3.942	3.942	3.942 (2.070 to 4.752)	3.942 (2.070 to 4.752)	3.942 (2.070 to 4.752)	3.942 (2.070 to 4.752)	3.942	3.942	3.942 (2.070 to 4.752)	3.942
	20 Pack Years	Deth	Mala	4.619	4.619	4.619	4.619	4.619	4.619	4.619	4.619	4.619	4.619	4.619	4.619	4.619	4.619
chronic obstructive purmonary disease	30 Pack-rears	Both	iviale	4.619	4.619	4.619	4.619	4.619	4.619	4.619	4.619	4.619	4.619	4.619	4.619	4.619	4.619
Chronic obstructive pulmonary disease	30 Pack-Years	BOTH	Female	(3.761 to 5.569) 5.255	(3./61 to 5.569) 5.255	5.255	(3.761 to 5.569) 5.255	(3.761 to 5.569) 5.255	5.255	(3.761 to 5.569) 5.255	5.255	(3.761 to 5.569) 5.255					
Chronic obstructive pulmonary disease	40 Pack-Years	Both	Male	(4.289 to 6.601) 5.255													
Chronic obstructive pulmonary disease	40 Pack-Years	Both	Female	(4.289 to 6.601) 5.967													
Chronic obstructive pulmonary disease	50 Pack-Years	Both	Male	(4.774 to 7.471) 5.967													
Chronic obstructive pulmonary disease	50 Pack-Years	Both	Female	(4.774 to 7.471) 6.995													
Chronic obstructive pulmonary disease	60 Pack-Years	Both	Male	(5.423 to 9.578) 6.995													
Chronic obstructive pulmonary disease	60 Pack-Years	Both	Female	(5.423 to 9.578) 8 274													
Chronic obstructive pulmonary disease	70 Pack-Years	Both	Male	(6.17 to 11.785)													
Chronic obstructive pulmonary disease	70 Pack-Years	Both	Female	(6.17 to 11.785)													
		B. (1		(6.933 to													
chronic obstructive purnonary disease	80 Pack-rears	BOLH	wate	9.838	9.838	9.838	9.838	9.838	9.838	9.838	9.838	9.838	9.838	9.838	9.838	9.838	9.838
Chronic obstructive pulmonary disease	80 Pack-Years	Both	Female	(6.933 to 15.001)													
				12.905 (8.309 to													
Chronic obstructive pulmonary disease	90 Pack-Years	Both	Male	21.862) 12.905													
Chronic obstructive pulmonary disease	90 Pack-Years	Both	Female	(8.309 to 21.862)	(8.309 to 21.862)	(8.309 to 21.862)	(8.309 to 21.862)										
Asthma	0 Cigarettes Per Smoker Per Dav	Both	Male	1.0 (1.0 to 1.0)													
Asthma	O Cigarettes Per Smoker Per Day	Both	Female	1.0 (1.0 to 1.0)													
Actima	2.5. Cigarattes Per Smoker Per Day	Both	Male	1.236 (1.0 to 1.521)													
Asthma	2.5 Cigarettes Per Smoker Per Day	Both	Fomala	1.236 (1.0 to 1.521)	1.236	1.236 (1.0 to 1.521)											
Astrina	2.5 Cigarettes Per Silloker Per Day	Buth	remaie	1.659	1.659	1.659	1.659	1.659	1.659	1.659	1.659	1.659	1.659	1.659	1.659	1.659	1.659
Astrima	7.5 Cigarettes Per Smoker Per Day	BOTH	Male	1.659	1.659	1.659	1.659	1.659	1.659	1.659	1.659	1.659	1.659	1.659	1.659	1.659	1.659
Asthma	7.5 Cigarettes Per Smoker Per Day	Both	Female	(1.19 to 2.18) 1.858													
Asthma	15 Cigarettes Per Smoker Per Day	Both	Male	(1.323 to 2.404) 1.858													
Asthma	15 Cigarettes Per Smoker Per Day	Both	Female	(1.323 to 2.404) 1.987													
Asthma	22.5 Cigarettes Per Smoker Per Day	Both	Male	(1.373 to 2.875) 1.987													
Asthma	22.5 Cigarettes Per Smoker Per Day	Both	Female	(1.373 to 2.875) 2.366													
Asthma	30 Cigarettes Per Smoker Per Day	Both	Male	(1.453 to 3.847) 2.366													
Asthma	30 Cigarettes Per Smoker Per Day	Both	Female	(1.453 to 3.847) 1.0													
Peptic ulcer disease	0 Cigarettes Per Smoker Per Day	Both	Male	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0)						
Peptic ulcer disease	0 Cigarettes Per Smoker Per Day	Both	Female	(1.0 to 1.0)	(1.0 to 1.0) 1 824	(1.0 to 1.0)	(1.0 to 1.0) 1 824	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0) 1 824	(1.0 to 1.0)	(1.0 to 1.0) 1 824					
Peptic ulcer disease	6.5 Cigarettes Per Smoker Per Day	Both	Male	(1.46 to 2.2)													
Peptic ulcer disease	6.5 Cigarettes Per Smoker Per Day	Both	Female	(1.46 to 2.2)	1.824 (1.46 to 2.2)	1.824 (1.46 to 2.2)	1.824 (1.46 to 2.2)	1.824 (1.46 to 2.2)	1.824 (1.46 to 2.2)	1.824 (1.46 to 2.2)	1.824 (1.46 to 2.2)	1.824 (1.46 to 2.2)	1.824 (1.46 to 2.2)	1.824 (1.46 to 2.2)	1.824 (1.46 to 2.2)	1.824 (1.46 to 2.2)	(1.46 to 2.2)

Appendix Table 6d. Relative risks used by age and sex for each ou	tcome for smoking globally.																
Risk - Outcome	Category / Units	Morbidity / Mortality	Sex	30 to 34	35 to 39	40 to 44	45 to 49	50 to 54	55 to 59	A 60 to 64	ge 65 to 69	70 to 74	75 to 79	80 to 84	85 to 89	90 to 94	95 plus
Smoking																	
Pantic ulcar disease	13 Cigarettes Per Smoker Per Day	Both	Male	2.156 (1.764 to 2.568)	2.156 (1.764 to 2.568	2.156											
	15 Cigarettes Fel Silloker Fel Day	BUCH		2.156	2.156	2.156	2.156	2.156	2.156	2.156	2.156	2.156	2.156	2.156	2.156	2.156	2.156
Peptic ulcer disease	13 Cigarettes Per Smoker Per Day	Both	Female	(1.764 to 2.568) 2.519	2.519 (1.764 to 2.568	2.519 (1.764 to 2.568)											
Peptic ulcer disease	19.5 Cigarettes Per Smoker Per Day	Both	Male	(2.066 to 2.994) 2.519	2.066 to 2.994 (2.519) (2.066 to 2.994) 2.519											
Peptic ulcer disease	19.5 Cigarettes Per Smoker Per Day	Both	Female	(2.066 to 2.994) 2.552	(2.066 to 2.994 2.552) (2.066 to 2.994) 2.552											
Peptic ulcer disease	26 Cigarettes Per Smoker Per Day	Both	Male	(1.952 to 3.262)	(1.952 to 3.262) (1.952 to 3.262)											
Peptic ulcer disease	26 Cigarettes Per Smoker Per Day	Both	Female	(1.952 to 3.262)	(1.952 to 3.262) (1.952 to 3.262)											
Peptic ulcer disease	32.5 Cigarettes Per Smoker Per Day	Both	Male	(2.15 to 3.741)													
Peptic ulcer disease	32.5 Cigarettes Per Smoker Per Day	Both	Female	2.888 (2.15 to 3.741)													
Gallbladder and biliary diseases	0 Cigarettes Per Smoker Per Day	Both	Male	1.0 (1.0 to 1.0)													
Gallbladder and biliary diseases	0 Cigarettes Per Smoker Per Day	Both	Female	1.0 (1.0 to 1.0)													
Gallbladder and biliary diseases	8.8 Cigarettes Per Smoker Per Day	Both	Male	1.36 (1.197 to 1.53)													
Gallbladder and biliary diseases	8.8 Cigarettes Per Smoker Per Day	Both	Female	1.36 (1.197 to 1.53)													
Gallbladder and biliary diseases	17.5 Cigarettes Per Smoker Per Day	Both	Male	1.303 (1.178 to 1.429)	1.303 (1.178 to 1.429	1.303) (1.178 to 1.429)											
Gallbladder and biliary diseases	17.5 Cigarettes Per Smoker Per Day	Both	Female	1.303 (1.178 to 1.429)	1.303 (1.178 to 1.429	1.303) (1.178 to 1.429)											
Gallbladder and biliary diseases	26.2 Cigarettes Per Smoker Per Day	Both	Male	1.398 (1.238 to 1.56)													
Gallbladder and biliary diseases	26.2 Cigarettes Per Smoker Per Day	Both	Female	1.398 (1.238 to 1.56)													
Gallbladder and biliary diseases	35 Cigarettes Per Smoker Per Day	Both	Male	1.612 (1.295 to 1.96)													
Gallbladder and biliary diseases	35 Cigarettes Per Smoker Per Day	Both	Female	1.612 (1.295 to 1.96)													
Gallbladder and biliary diseases	43.8 Cigarettes Per Smoker Per Day	Both	Male	1.81 (1.451 to 2.164)	1.81 (1.451 to 2.164	1.81) (1.451 to 2.164)											
Gallbladder and biliary diseases	43.8 Cigarettes Per Smoker Per Day	Both	Female	1.81 (1.451 to 2.164)	1.81 (1.451 to 2.164	1.81) (1.451 to 2.164)											
Alzheimer's disease and other dementias	0 Cigarettes Per Smoker Per Dav	Both	Male	1.0 (1.0 to 1.0)													
Alzheimer's disease and other dementias	0 Cigarettes Per Smoker Per Day	Both	Female	1.0 (1.0 to 1.0)													
Alzheimer's disease and other dementias	12 Cigarettes Per Smoker Per Day	Both	Male	2.078 (1.396 to 2.844)	2.078 (1.396 to 2.844	2.078) (1.396 to 2.844)											
Alzheimer's disease and other dementias	12 Cigarettes Per Smoker Per Day	Both	Fomalo	2.078 (1.396 to 2.844)	2.078 (1.395 to 2.844)	2.078 (1.396 to 2.844)	2.078 (1.396 to 2.844)	2.078 (1.395 to 2.844)	2.078 (1.396 to 2.844)	2.078	2.078						
	12 Cigarettes Per Silloker Per Day	BUCH	remaie	2.936	2.936	2.936	2.936	2.936	2.936	2.936	2.936	2.936	2.936	2.936	2.936	2.936	2.936
Alzheimer's disease and other dementias	24 Cigarettes Per Smoker Per Day	Both	Male	(1.782 to 4.472) 2.936	2.936 (1.782 to 4.472	2.936 (1.782 to 4.472)											
Alzheimer's disease and other dementias	24 Cigarettes Per Smoker Per Day	Both	Female	(1.782 to 4.472) 3.737) (1.782 to 4.472) 3.737												
Alzheimer's disease and other dementias	36 Cigarettes Per Smoker Per Day	Both	Male	(1.972 to 6.019) 3.737	(1.972 to 6.019 3.737) (1.972 to 6.019) 3.737											
Alzheimer's disease and other dementias	36 Cigarettes Per Smoker Per Day	Both	Female	(1.972 to 6.019)	(1.972 to 6.019) (1.972 to 6.019)											
Alzheimer's disease and other dementias	48 Cigarettes Per Smoker Per Day	Both	Male	(2.075 to 7.232)	4.105 (2.075 to 7.232)	(2.075 to 7.232)	4.105 (2.075 to 7.232)	4.105 (2.075 to 7.232	4.103) (2.075 to 7.232)								
Alzheimer's disease and other dementias	48 Cigarettes Per Smoker Per Day	Both	Female	(2.075 to 7.232)	4.105 (2.075 to 7.232)	4.105 (2.075 to 7.232	4.103) (2.075 to 7.232)										
Parkinson's disease	0 Cigarettes Per Smoker Per Day	Both	Male	(1.0 to 1.0)													
Parkinson's disease	0 Cigarettes Per Smoker Per Day	Both	Female	(1.0 to 1.0)													
Parkinson's disease	7.5 Cigarettes Per Smoker Per Day	Both	Male	(0.644 to 0.989)	0.835 (0.644 to 0.989) (0.644 to 0.989)											
Parkinson's disease	7.5 Cigarettes Per Smoker Per Day	Both	Female	(0.644 to 0.989)	0.835 (0.644 to 0.989)	(0.644 to 0.989)	0.835 (0.644 to 0.989)	0.835 (0.644 to 0.989	0.835) (0.644 to 0.989)								
Parkinson's disease	15 Cigarettes Per Smoker Per Day	Both	Male	(0.531 to 0.869)	(0.531 to 0.869) (0.531 to 0.869)											
Parkinson's disease	15 Cigarettes Per Smoker Per Day	Both	Female	(0.531 to 0.869)	(0.531 to 0.869) (0.531 to 0.869)											
Parkinson's disease	22.5 Cigarettes Per Smoker Per Day	Both	Male	(0.44 to 0.789)													

Appendix Table 6d. Relative risks used by age and sex for each our	tcome for smoking globally.	Morbidity /									70						
Risk - Outcome	Category / Units	Mortality	Sex	30 to 34	35 to 39	40 to 44	45 to 49	50 to 54	55 to 59	60 to 64	65 to 69	70 to 74	75 to 79	80 to 84	85 to 89	90 to 94	95 plus
Smoking																	
Parkinson's disease	22.5 Cigarettes Per Smoker Per Day	Both	Female	0.604 (0.44 to 0.789)	0.604 (0.44 to 0.789)												
Parkinson's disease	30 Cigarettes Per Smoker Per Day	Both	Male	0.52 (0.346 to 0.71)	0.52 (0.346 to 0.71)												
Parkinson's disease	30 Cigarettes Per Smoker Per Day	Both	Female	0.52 (0.346 to 0.71)	0.52 (0.346 to 0.71)												
Parkinson's disease	37.5 Cigarettes Per Smoker Per Day	Both	Male	0.432 (0.238 to 0.644)	0.432 (0.238 to 0.644)												
Parkinson's disease	37.5 Cigarettes Per Smoker Per Day	Both	Female	0.432 (0.238 to 0.644)	0.432 (0.238 to 0.644)												
Parkinson's disease	45 Cigarettes Per Smoker Per Day	Both	Male	0.323 (0.127 to 0.561)	0.323 (0.127 to 0.561)												
Parkinson's disease	45 Cigarettes Per Smoker Per Day	Both	Female	(0.127 to 0.561)	0.323) (0.127 to 0.561)	(0.127 to 0.561)											
Multiple sclerosis	0 Cigarettes Per Smoker Per Day	Both	Male	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	(1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)
Multiple sclerosis	0 Cigarettes Per Smoker Per Day	Both	Female	1.0 (1.0 to 1.0)	1.0 (1.0 to 1.0)												
Multiple sclerosis	5.6 Cigarettes Per Smoker Per Day	Both	Male	(1.152 to 1.617)	(1.152 to 1.617)												
Multiple sclerosis	5.6 Cigarettes Per Smoker Per Day	Both	Female	(1.152 to 1.617)	(1.152 to 1.617)												
Multiple sclerosis	11.2 Cigarettes Per Smoker Per Day	Both	Male	(1.49 to 2.048)	(1.49 to 2.048)												
Multiple sclerosis	11.2 Cigarettes Per Smoker Per Day	Both	Female	(1.49 to 2.048)	(1.49 to 2.048)												
Multiple sclerosis	16.9 Cigarettes Per Smoker Per Day	Both	Male	(1.646 to 2.291) 1.965	(1.646 to 2.291) 1.965												
Multiple sclerosis	16.9 Cigarettes Per Smoker Per Day	Both	Female	(1.646 to 2.291) 2.04	(1.646 to 2.291) 2.04												
Multiple sclerosis	22.5 Cigarettes Per Smoker Per Day	Both	Male	(1.58 to 2.545)	(1.58 to 2.545)												
Multiple sclerosis	22.5 Cigarettes Per Smoker Per Day	Both	Female	(1.58 to 2.545)	(1.58 to 2.545)												
Diabetes mellitus type 2	0 Cigarettes Per Smoker Per Day	Both	Male	(1.0 to 1.0)	(1.0 to 1.0)												
Diabetes mellitus type 2	0 Cigarettes Per Smoker Per Day	Both	Female	(1.0 to 1.0)	(1.0 to 1.0)												
Diabetes mellitus type 2	6.5 Cigarettes Per Smoker Per Day	Both	Male	(1.213 to 1.653)	(1.213 to 1.653)												
Diabetes mellitus type 2	6.5 Cigarettes Per Smoker Per Day	Both	Female	(1.213 to 1.653)	(1.213 to 1.653)												
Diabetes mellitus type 2	12.9 Cigarettes Per Smoker Per Day	Both	Male	(1.131 to 1.756)	(1.131 to 1.756)												
Diabetes mellitus type 2	12.9 Cigarettes Per Smoker Per Day	Both	Female	(1.131 to 1.756) 1.637	(1.131 to 1.756) 1.637												
Diabetes mellitus type 2	19.4 Cigarettes Per Smoker Per Day	Both	Male	(1.25 to 2.031) 1.637	(1.25 to 2.031) 1.637												
Diabetes mellitus type 2	19.4 Cigarettes Per Smoker Per Day	Both	Female	(1.25 to 2.031)	(1.25 to 2.031)												
Diabetes mellitus type 2	25.8 Cigarettes Per Smoker Per Day	Both	Male	(1.355 to 1.992)	(1.355 to 1.992)												
Diabetes mellitus type 2	25.8 Cigarettes Per Smoker Per Day	Both	Female	(1.355 to 1.992)	(1.355 to 1.992)												
Diabetes mellitus type 2	32.3 Cigarettes Per Smoker Per Day	Both	Male	(1.221 to 2.458)	(1.221 to 2.458)												
Diabetes mellitus type 2	32.3 Cigarettes Per Smoker Per Day	Both	Female	(1.221 to 2.458)	(1.221 to 2.458)	(1.221 to 2.458) 2 163	(1.221 to 2.458)	(1.221 to 2.458) 2 163	(1.221 to 2.458) 2 163	(1.221 to 2.458)	(1.221 to 2.458) 2 163	(1.221 to 2.458)	(1.221 to 2.458) 2 163	(1.221 to 2.458)	(1.221 to 2.458)	(1.221 to 2.458) 2 163	(1.221 to 2.458) 2 163
Diabetes mellitus type 2	38.8 Cigarettes Per Smoker Per Day	Both	Male	(1.161 to 3.354)	(1.161 to 3.354)	(1.161 to 3.354) 2 163	(1.161 to 3.354) 2 163	(1.161 to 3.354)	(1.161 to 3.354)	(1.161 to 3.354) 2 163	(1.161 to 3.354)	(1.161 to 3.354)	(1.161 to 3.354) 2 163	(1.161 to 3.354) 2 163	(1.161 to 3.354)	(1.161 to 3.354)	(1.161 to 3.354)
Diabetes mellitus type 2	38.8 Cigarettes Per Smoker Per Day	Both	Female	(1.161 to 3.354) 1.0	(1.161 to 3.354) 1.0	(1.161 to 3.354) 1.0	(1.161 to 3.354)	(1.161 to 3.354) 1.0	(1.161 to 3.354) 1.0	(1.161 to 3.354)	(1.161 to 3.354)	(1.161 to 3.354) 1.0	(1.161 to 3.354)	(1.161 to 3.354) 1.0	(1.161 to 3.354) 1.0	(1.161 to 3.354) 1.0	(1.161 to 3.354)
Cataract	0 Cigarettes Per Smoker Per Day	Both	Male	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0)
Cataract	0 Cigarettes Per Smoker Per Day	Both	Female	(1.0 to 1.0) 1.438	(1.0 to 1.0) 1.438												
Cataract	9 Cigarettes Per Smoker Per Day	Both	Male	(1.288 to 1.608) 1.438	(1.288 to 1.608) 1.438												
Cataract	9 Cigarettes Per Smoker Per Day	Both	Female	(1.288 to 1.608)	(1.288 to 1.608)												
Cataract	18 Cigarettes Per Smoker Per Day	Both	Male	(1.288 to 1.694)	(1.288 to 1.694)												
Cataract	18 Cigarettes Per Smoker Per Day	Both	Female	(1.288 to 1.694)	(1.288 to 1.694)												

Appendix Table 6d. Relative risks used by age and sex for each ou	teome for smoking globally.	Morbidity /	0							А	ge						
Risk - Outcome	Category / Units	Mortality	Sex	30 to 34	35 to 39	40 to 44	45 to 49	50 to 54	55 to 59	60 to 64	65 to 69	70 to 74	75 to 79	80 to 84	85 to 89	90 to 94	95 plus
moking																	
Cataract	27 Cigarettes Per Smoker Per Day	Both	Male	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)	1.786) (1.356 to 2.266)	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)
Cataract	27 Cigarettes Per Smoker Per Day	Both	Female	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)	1.786 (1.356 to 2.266)
Cataract	36 Cigarettes Per Smoker Per Day	Both	Male	(1.21 to 2.734)	(1.21 to 2.734)	(1.21 to 2.734)	(1.21 to 2.734)	(1.21 to 2.734)	(1.21 to 2.734)	(1.21 to 2.734)	(1.21 to 2.734)	(1.21 to 2.734)	(1.21 to 2.734)	(1.21 to 2.734)	(1.21 to 2.734)	(1.21 to 2.734)	(1.21 to 2.734)
Cataract	36 Cigarettes Per Smoker Per Day	Both	Female	1.882 (1.21 to 2.734) 1.0	1.882 (1.21 to 2.734)	1.882 (1.21 to 2.734) 1.0	1.882 (1.21 to 2.734)	1.882 (1.21 to 2.734)	1.882 (1.21 to 2.734)	1.882 (1.21 to 2.734)	1.882 (1.21 to 2.734)	1.882 (1.21 to 2.734)	1.882 (1.21 to 2.734)	1.882 (1.21 to 2.734)	1.882 (1.21 to 2.734)	1.882 (1.21 to 2.734) 1.0	1.882 (1.21 to 2.734)
Age-related macular degeneration	0 Cigarettes Per Smoker Per Day	Both	Male	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0) 1.0	(1.0 to 1.0)	(1.0 to 1.0)	(1.0 to 1.0)
Age-related macular degeneration	0 Cigarettes Per Smoker Per Day	Both	Female	(1.0 to 1.0) 1.347	(1.0 to 1.0) 1.347	(1.0 to 1.0) 1.347	(1.0 to 1.0) 1.347	(1.0 to 1.0) 1.347	(1.0 to 1.0)	(1.0 to 1.0) 1.347	(1.0 to 1.0) 1.347	(1.0 to 1.0) 1.347	(1.0 to 1.0) 1.347	(1.0 to 1.0) 1.347	(1.0 to 1.0) 1.347	(1.0 to 1.0) 1.347	(1.0 to 1.0) 1.347
Age-related macular degeneration	7.5 Cigarettes Per Smoker Per Day	Both	Male	(1.0 to 1.769) 1.347	(1.0 to 1.769) 1.347	(1.0 to 1.769) 1.347	(1.0 to 1.769) 1.347	(1.0 to 1.769) 1.347	(1.0 to 1.769) 1.347	(1.0 to 1.769) 1.347	(1.0 to 1.769) 1.347	(1.0 to 1.769) 1.347	(1.0 to 1.769) 1.347	(1.0 to 1.769) 1.347	(1.0 to 1.769) 1.347	(1.0 to 1.769) 1.347	(1.0 to 1.769) 1.347
Age-related macular degeneration	7.5 Cigarettes Per Smoker Per Day	Both	Female	(1.0 to 1.769) 1.67	(1.0 to 1.769) 1.67	(1.0 to 1.769) 1.67	(1.0 to 1.769) 1.67	(1.0 to 1.769) 1.67	(1.0 to 1.769) 1.67	(1.0 to 1.769) 1.67	(1.0 to 1.769) 1.67	(1.0 to 1.769) 1.67	(1.0 to 1.769) 1.67	(1.0 to 1.769) 1.67	(1.0 to 1.769) 1.67	(1.0 to 1.769) 1.67	(1.0 to 1.769) 1.67
Age-related macular degeneration	15 Cigarettes Per Smoker Per Day	Both	Male	(1.152 to 2.349) 1.67	(1.152 to 2.349) 1.67) (1.152 to 2.349) 1.67	(1.152 to 2.349) 1.67	(1.152 to 2.349) 1.67	(1.152 to 2.349) 1.67	(1.152 to 2.349) 1.67	(1.152 to 2.349) 1.67	(1.152 to 2.349) 1.67	(1.152 to 2.349) 1.67	(1.152 to 2.349) 1.67	(1.152 to 2.349) 1.67	(1.152 to 2.349) 1.67	(1.152 to 2.349) 1.67
Age-related macular degeneration	15 Cigarettes Per Smoker Per Day	Both	Female	(1.152 to 2.349) 2.024	(1.152 to 2.349) 2.024) (1.152 to 2.349) 2.024	(1.152 to 2.349) 2.024	(1.152 to 2.349) 2.024	(1.152 to 2.349) 2.024	(1.152 to 2.349) 2.024	(1.152 to 2.349) 2.024	(1.152 to 2.349) 2.024	(1.152 to 2.349) 2.024	(1.152 to 2.349) 2.024	(1.152 to 2.349) 2.024	(1.152 to 2.349) 2.024	(1.152 to 2.349) 2.024
Age-related macular degeneration	22.5 Cigarettes Per Smoker Per Day	Both	Male	(1.375 to 2.775) 2.024	(1.375 to 2.775) 2.024) (1.375 to 2.775) 2.024	(1.375 to 2.775) 2.024	(1.375 to 2.775) 2.024	(1.375 to 2.775) 2.024	(1.375 to 2.775) 2.024	(1.375 to 2.775) 2.024	(1.375 to 2.775) 2.024	(1.375 to 2.775) 2.024	(1.375 to 2.775) 2.024	(1.375 to 2.775) 2.024	(1.375 to 2.775) 2.024	(1.375 to 2.775) 2.024
Age-related macular degeneration	22.5 Cigarettes Per Smoker Per Day	Both	Female	(1.375 to 2.775) 2.42	(1.375 to 2.775) 2.42) (1.375 to 2.775) 2.42	(1.375 to 2.775) 2.42	(1.375 to 2.775) 2.42	(1.375 to 2.775) 2.42	(1.375 to 2.775) 2.42	(1.375 to 2.775) 2.42	(1.375 to 2.775) 2.42	(1.375 to 2.775) 2.42	(1.375 to 2.775) 2.42	(1.375 to 2.775) 2.42	(1.375 to 2.775) 2.42	(1.375 to 2.775) 2.42
Age-related macular degeneration	30 Cigarettes Per Smoker Per Day	Both	Male	(1.624 to 3.354) 2.42	(1.624 to 3.354) 2.42) (1.624 to 3.354) 2.42	(1.624 to 3.354) 2.42	(1.624 to 3.354) 2.42	(1.624 to 3.354) 2.42	(1.624 to 3.354) 2.42	(1.624 to 3.354) 2.42	(1.624 to 3.354) 2.42	(1.624 to 3.354) 2.42	(1.624 to 3.354) 2.42	(1.624 to 3.354) 2.42	(1.624 to 3.354) 2.42	(1.624 to 3.354) 2.42
Age-related macular degeneration	30 Cigarettes Per Smoker Per Day	Both	Female	(1.624 to 3.354) 1.0	(1.624 to 3.354) 1.0) (1.624 to 3.354) 1.0	(1.624 to 3.354) 1.0	(1.624 to 3.354) 1.0	(1.624 to 3.354) 1.0	(1.624 to 3.354) 1.0	(1.624 to 3.354) 1.0	(1.624 to 3.354) 1.0	(1.624 to 3.354) 1.0	(1.624 to 3.354) 1.0	(1.624 to 3.354) 1.0	(1.624 to 3.354) 1.0	(1.624 to 3.354) 1.0
Reumatoid arthritis	0 Cigarettes Per Smoker Per Day	Both	Male	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0
Reumatoid arthritis	0 Cigarettes Per Smoker Per Day	Both	Female	(1.0 to 1.0) 1.406	(1.0 to 1.0) 1.406	(1.0 to 1.0) 1.406	(1.0 to 1.0) 1.406	(1.0 to 1.0) 1.406	(1.0 to 1.0) 1.406	(1.0 to 1.0) 1.406	(1.0 to 1.0) 1.406	(1.0 to 1.0) 1.406	(1.0 to 1.0) 1.406	(1.0 to 1.0) 1.406	(1.0 to 1.0) 1.406	(1.0 to 1.0) 1.406	(1.0 to 1.0) 1.406
Reumatoid arthritis	7.5 Cigarettes Per Smoker Per Day	Both	Male	(1.0 to 1.965) 1.406	(1.0 to 1.965) 1.406	(1.0 to 1.965) 1.406	(1.0 to 1.965) 1.406	(1.0 to 1.965) 1.406	(1.0 to 1.965) 1.406	(1.0 to 1.965) 1.406	(1.0 to 1.965) 1.406	(1.0 to 1.965) 1.406	(1.0 to 1.965) 1.406	(1.0 to 1.965) 1.406	(1.0 to 1.965) 1.406	(1.0 to 1.965) 1.406	(1.0 to 1.965) 1.406
Reumatoid arthritis	7.5 Cigarettes Per Smoker Per Day	Both	Female	(1.0 to 1.965) 1.742	(1.0 to 1.965) 1.742	(1.0 to 1.965) 1.742	(1.0 to 1.965) 1.742	(1.0 to 1.965) 1.742	(1.0 to 1.965) 1.742	(1.0 to 1.965) 1.742	(1.0 to 1.965) 1.742	(1.0 to 1.965) 1.742	(1.0 to 1.965) 1.742	(1.0 to 1.965) 1.742	(1.0 to 1.965) 1.742	(1.0 to 1.965) 1.742	(1.0 to 1.965) 1.742
Rheumatoid arthritis	15 Cigarettes Per Smoker Per Day	Both	Male	(1.101 to 2.633) 1.742	(1.101 to 2.633) 1.742) (1.101 to 2.633) 1.742	(1.101 to 2.633) 1.742	(1.101 to 2.633) 1.742	(1.101 to 2.633) 1.742	(1.101 to 2.633) 1.742	(1.101 to 2.633) 1.742	(1.101 to 2.633) 1.742	(1.101 to 2.633) 1.742	(1.101 to 2.633) 1.742	(1.101 to 2.633) 1.742	(1.101 to 2.633) 1.742	(1.101 to 2.633) 1.742
Reumatoid arthritis	15 Cigarettes Per Smoker Per Day	Both	Female	(1.101 to 2.633) 2.0	(1.101 to 2.633) 2.0) (1.101 to 2.633) 2.0	(1.101 to 2.633) 2.0	(1.101 to 2.633) 2.0	(1.101 to 2.633) 2.0	(1.101 to 2.633) 2.0	(1.101 to 2.633) 2.0	(1.101 to 2.633) 2.0	(1.101 to 2.633) 2.0	(1.101 to 2.633) 2.0	(1.101 to 2.633) 2.0	(1.101 to 2.633) 2.0	(1.101 to 2.633) 2.0
Reumatoid arthritis	22.5 Cigarettes Per Smoker Per Day	Both	Male	(1.167 to 3.187) 2.0	(1.167 to 3.187) 2.0	(1.167 to 3.187) 2.0	(1.167 to 3.187) 2.0	(1.167 to 3.187) 2.0	(1.167 to 3.187) 2.0	(1.167 to 3.187) 2.0	(1.167 to 3.187) 2.0	(1.167 to 3.187) 2.0	(1.167 to 3.187) 2.0	(1.167 to 3.187) 2.0	(1.167 to 3.187) 2.0	(1.167 to 3.187) 2.0	(1.167 to 3.187) 2.0
Reumatoid arthritis	22.5 Cigarettes Per Smoker Per Day	Both	Female	(1.167 to 3.187) 2.256	(1.167 to 3.187) 2.256	(1.167 to 3.187) 2.256	(1.167 to 3.187) 2.256	(1.167 to 3.187) 2.256	(1.167 to 3.187) 2.256	(1.167 to 3.187) 2.256	(1.167 to 3.187) 2.256	(1.167 to 3.187) 2.256	(1.167 to 3.187) 2.256	(1.167 to 3.187) 2.256	(1.167 to 3.187) 2.256	(1.167 to 3.187) 2.256	(1.167 to 3.187) 2.256
Reumatoid arthritis	30 Cigarettes Per Smoker Per Day	Both	Male	(1.147 to 3.913) 2.256	(1.147 to 3.913) 2.256	(1.147 to 3.913) 2.256	(1.147 to 3.913) 2.256	(1.147 to 3.913) 2.256	(1.147 to 3.913) 2.256	(1.147 to 3.913) 2.256	(1.147 to 3.913) 2.256	(1.147 to 3.913) 2.256	(1.147 to 3.913) 2.256	(1.147 to 3.913) 2.256	(1.147 to 3.913) 2.256	(1.147 to 3.913) 2.256	(1.147 to 3.913) 2.256
Reumatoid arthritis	30 Cigarettes Per Smoker Per Day	Both	Female	(1.147 to 3.913) 2.438	(1.147 to 3.913) 2.438	(1.147 to 3.913) 2.438	(1.147 to 3.913) 2.438	(1.147 to 3.913) 2.438	(1.147 to 3.913) 2.438	(1.147 to 3.913) 2.438	(1.147 to 3.913) 2.438	(1.147 to 3.913) 2.438	(1.147 to 3.913) 2.438	(1.147 to 3.913) 2.438	(1.147 to 3.913) 2.438	(1.147 to 3.913) 2.438	(1.147 to 3.913) 2.438
Reumatoid arthritis	37.5 Cigarettes Per Smoker Per Day	Both	Male	(1.15 to 4.713) 2.438	(1.15 to 4.713) 2.438	(1.15 to 4.713) 2.438	(1.15 to 4.713) 2.438	(1.15 to 4.713) 2.438	(1.15 to 4.713) 2.438	(1.15 to 4.713) 2.438	(1.15 to 4.713) 2.438	(1.15 to 4.713) 2.438	(1.15 to 4.713) 2.438	(1.15 to 4.713) 2.438	(1.15 to 4.713) 2.438	(1.15 to 4.713) 2.438	(1.15 to 4.713) 2.438
Reumatoid arthritis	37.5 Cigarettes Per Smoker Per Day	Both	Female	(1.15 to 4.713) 1.0	(1.15 to 4.713) 1.0	(1.15 to 4.713) 1.0	(1.15 to 4.713) 1.0	(1.15 to 4.713) 1.0	(1.15 to 4.713) 1.0	(1.15 to 4.713) 1.0	(1.15 to 4.713) 1.0	(1.15 to 4.713) 1.0	(1.15 to 4.713) 1.0	(1.15 to 4.713) 1.0	(1.15 to 4.713) 1.0	(1.15 to 4.713) 1.0	(1.15 to 4.713) 1.0
.ow back pain	0 Cigarettes Per Smoker Per Day	Both	Male	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0	(1.0 to 1.0) 1.0
ow back pain	0 Cigarettes Per Smoker Per Day	Both	Female	(1.0 to 1.0) 1.315	(1.0 to 1.0) 1.315	(1.0 to 1.0) 1.315	(1.0 to 1.0) 1.315	(1.0 to 1.0) 1.315	(1.0 to 1.0) 1.315	(1.0 to 1.0) 1.315	(1.0 to 1.0) 1.315	(1.0 to 1.0) 1.315	(1.0 to 1.0) 1.315	(1.0 to 1.0) 1.315	(1.0 to 1.0) 1.315	(1.0 to 1.0) 1.315	(1.0 to 1.0) 1.315
ow back pain	2.5 Cigarettes Per Smoker Per Day	Both	Male	(1.104 to 1.564) 1.315	(1.104 to 1.564) 1.315	(1.104 to 1.564) 1.315	(1.104 to 1.564) 1.315	(1.104 to 1.564) 1.315	(1.104 to 1.564) 1.315	(1.104 to 1.564) 1.315	(1.104 to 1.564) 1.315	(1.104 to 1.564) 1.315	(1.104 to 1.564) 1.315	(1.104 to 1.564) 1.315	(1.104 to 1.564) 1.315	(1.104 to 1.564) 1.315	(1.104 to 1.564) 1.315
ow back pain	2.5 Cigarettes Per Smoker Per Day	Both	Female	(1.104 to 1.564) 1.666	(1.104 to 1.564) 1.666	(1.104 to 1.564) 1.666	(1.104 to 1.564) 1.666	(1.104 to 1.564) 1.666	(1.104 to 1.564) 1.666	(1.104 to 1.564) 1.666	(1.104 to 1.564) 1.666	(1.104 to 1.564) 1.666	(1.104 to 1.564) 1.666	(1.104 to 1.564) 1.666	(1.104 to 1.564) 1.666	(1.104 to 1.564) 1.666	(1.104 to 1.564) 1.666
ow back pain	5 Cigarettes Per Smoker Per Day	Both	Male	(1.387 to 1.975) 1.666	(1.387 to 1.975) 1.666	(1.387 to 1.975) 1.666	(1.38/ to 1.9/5) 1.666	(1.387 to 1.975) 1.666	(1.38/ to 1.9/5) 1.666	(1.387 to 1.975) 1.666	(1.387 to 1.975) 1.666	(1.387 to 1.975) 1.666	(1.38/ to 1.9/5) 1.666 (4.207 to 4.075)	(1.387 to 1.975) 1.666	(1.38/ to 1.9/5) 1.666	(1.38/ to 1.9/5) 1.666	(1.387 to 1.975) 1.666
ow back pain	5 Cigarettes Per Smoker Per Day	Both	Female	(1.38/ to 1.9/5) 2.161	(1.38/ to 1.9/5) 2.161	2.161	(1.38/ to 1.9/5) 2.161	(1.387 to 1.975) 2.161	(1.38/ to 1.9/5) 2.161	(1.38/ to 1.9/5) 2.161	(1.38/ to 1.9/5) 2.161	(1.387 to 1.975) 2.161	(1.38/ to 1.9/5) 2.161	(1.387 to 1.975) 2.161	(1.38/ to 1.9/5) 2.161	(1.38/ to 1.9/5) 2.161	(1.38/ to 1.9/5) 2.161
ow back pain	10 Cigarettes Per Smoker Per Day	Both	iviale	(1.746 to 2.636) 2.161	2.161	2.161	(1.746 to 2.636) 2.161	(1.746 to 2.636) 2.161	(1.746 to 2.636) 2.161	(1.746 to 2.636) 2.161	(1.746 to 2.636) 2.161	(1.746 to 2.636) 2.161	(1.746 to 2.636) 2.161	(1.746 to 2.636) 2.161	(1.746 to 2.636) 2.161	(1.746 to 2.636) 2.161	(1.746 to 2.636) 2.161
ow back pain	15 Cigarettes Per Smoker Per Day	Both	remale	(1.740 (0 2.636) 2.244 (1.700 += 3.800)	2.244	2.244	(1.740 (0.2.036) 2.244 (1.700 to 2.005)	(1.740 t0 2.636) 2.244 (1.700 t= 2.800)	(1.740 t0 2.636) 2.244	(1.740 t0 2.636) 2.244	(1.740 (0 2.036) 2.244 (1.700 to 2.000)	2.244	(1.740 t0 2.036) 2.244 (1.700 t= 3.000)				
ow back pain	15 Ggarettes Per Smoker Per Day	Both	Fomale	(1.799 (0 2.806) 2.244 (1.700 += 2.800)	2.244	2.244	2.244	(1.799 t0 2.806) 2.244 (1.700 t- 2.806)	(1.799 t0 2.806) 2.244	(1.799 t0 2.806) 2.244	(1.799 to 2.806) 2.244	(1.799 to 2.806) 2.244	(1.799 t0 2.806) 2.244	(1.799 to 2.806) 2.244	(1.799 (0 2.806) 2.244 (1.700 to 2.805)	2.244	(1.799 t0 2.806) 2.244 (1.700 to 2.805)
	20 Gizerettes Per Smoker Per Day	BUTN	remaie	(1.799 to 2.806) 2.353	2.353	2.353	2.353	(1.799 to 2.806) 2.353	2.353	2.353	(1.799 to 2.806) 2.353	2.353	2.353	2.353	2.353	2.353	2.353
.ow back pain	20 Ligarettes Per Smoker Per Day	воth	wale	(1.614 to 3.307)	(1.614 to 3.307)) (1.614 to 3.307)	(1.614 to 3.307)	(1.614 to 3.307)	(1.614 to 3.307)	(1.614 to 3.307)	(1.614 to 3.307)	(1.614 to 3.307)	(1.614 to 3.307)	(1.614 to 3.307)	(1.614 to 3.307)	(1.614 to 3.307)	(1.614 to 3.307)

Appendix Table 6d. Relative risks used by age and sex for each o	utcome for smoking globally.																
Risk - Outcome	Category / Units	Morbidity / Mortality	Sex							А	ge						
		Mortanty		30 to 34	35 to 39	40 to 44	45 to 49	50 to 54	55 to 59	60 to 64	65 to 69	70 to 74	75 to 79	80 to 84	85 to 89	90 to 94	95 plus
Smoking																	
				2.353	2.353	2.353	2.353	2.353	2.353	2.353	2.353	2.353	2.353	2.353	2.353	2.353	2.353
Low back pain	20 Cigarettes Per Smoker Per Day	Both	Female	(1.614 to 3.307)													
				1.85	1.85	1.85	1.85	1.85	1.85	1.85	1.85	1.85	1.85	1.85	1.85	1.85	1.85
Hip fractures	1 Prevalence	Both	Male	(1.518 to 2.249)													
				1.85	1.85	1.85	1.85	1.85	1.85	1.85	1.85	1.85	1.85	1.85	1.85	1.85	1.85
Hip fractures	1 Prevalence	Both	Female	(1.518 to 2.249)													
				1.25	1.25	1.25	1.25	1.25	1.25	1.25	1.25	1.25	1.25	1.25	1.25	1.25	1.25
Non-hip fractures	1 Prevalence	Both	Male	(1.153 to 1.358)													
				1.25	1.25	1.25	1.25	1.25	1.25	1.25	1.25	1.25	1.25	1.25	1.25	1.25	1.25
Non-hip fractures	1 Prevalence	Both	Female	(1.153 to 1.358)													

Geography	Level
Global	0
Low SDI	1
Low-middle SDI	1
Middle SDI	1
High-middle SDI	1
High SDI	1
Central Europe, Eastern Europe, and Central Asia	1
Central Asia	2
Armenia	3
Azerbaijan	3
Georgia	3
Kazakhstan	3
Kyrgyzstan	3
Mongolia	3
Tajikistan	3
Turkmenistan	3
Uzbekistan	3
Central Europe	2
Albania	3
Bosnia and Herzegovina	3
Bulgaria	3
Croatia	3
Czech Republic	3
Hungary	3
Macedonia	3
Montenegro	3
Poland	3
Romania	3
Serbia	3
Slovakia	3
Slovenia	3
Eastern Europe	2
Belarus	3
Estonia	3
Latvia	3
Lithuania	3
Moldova	3
Russian Federation	3
Ukraine	3
High-income	1
Australasia	2
Australia	3
New Zealand	3
High-income Asia-Pacific	2
P i	3

Appendix Table 7. GBD location hierarc	thy with levels
Geography	Level
Japan	3
Aichi	4
Akita	4
Aomori	4
Chiba	4
Ehime	4
Fukui	4
Fukuoka	4
Fukushima	4
Gifu	4
Gunma	4
Hiroshima	4
Hokkaidō	4
Hyōgo	4
Ibaraki	4
Ishikawa	4
Iwate	4
Kagawa	4
Kagoshima	4
Kanagawa	4
Kōchi	4
Kumamoto	4
Kyōto	4
Mie	4
Miyagi	4
Miyazaki	4
Nagano	4
Nagasaki	4
Nara	4
Niigata	4
Ōita	4
Okayama	4
Okinawa	4
Ōsaka	4
Saga	4
Saitama	4
Shiga	4
Shimane	4
Shizuoka	4
Tochigi	4
Tokushima	4
Tōkyō	4
Tottori	4
Toyama	4
Wakayama	4

Geography	Level
Yamagata	4
Yamaguchi	4
Yamanashi	4
South Korea	3
Singapore	3
High-income North America	2
Canada	3
Greenland	3
USA	3
Alabama	4
Alaska	4
Arizona	4
Arkansas	4
California	4
Colorado	4
Connecticut	4
Delaware	4
Washington, DC	4
Florida	4
Georgia	4
Hawaii	4
Idaho	4
Illinois	4
Indiana	4
Iowa	4
Kansas	4
Kentucky	4
Louisiana	4
Maine	4
Maryland	4
Massachusetts	4
Michigan	4
Minnesota	4
Mississippi	4
Missouri	4
Montana	4
Nebraska	4
Nevada	4
New Hampshire	4
New Jersey	4
New Mexico	4
New York	4
North Carolina	4
North Dakota	4
Ohio	4
UIIU	4

Appendix Table 7. GBD location hierarchy with levels		
Geography	Level	
Oklahoma	4	
Oregon	4	
Pennsylvania	4	
Rhode Island	4	
South Carolina	4	
South Dakota	4	
Tennessee	4	
Texas	4	
Utah	4	
Vermont	4	
Virginia	4	
Washington	4	
West Virginia	4	
Wisconsin	4	
Wyoming	4	
Southern Latin America	2	
Argentina	3	
Chile	3	
Uruguay	3	
Western Europe	2	
Andorra	3	
Austria	3	
Belgium	3	
Cyprus	3	
Denmark	3	
Finland	3	
France	3	
Germany	3	
Greece	3	
Iceland	3	
Ireland	3	
Israel	3	
Italy	3	
Luxembourg	3	
Malta	3	
Netherlands	3	
Norway	3	
Portugal	3	
Spain	3	
Sweden	3	
Stockholm	4	
Sweden except Stockholm	4	
Switzerland	3	
United Kingdom	3	
England	4	

Appendix Table 7. GBD location hierarchy with levels		
Geography	Level	
East Midlands	5	
Derby	6	
Derbyshire	6	
Leicester	6	
Leicestershire	6	
Lincolnshire	6	
Northamptonshire	6	
Nottingham	6	
Nottinghamshire	6	
Rutland	6	
East of England	5	
Bedford	6	
Cambridgeshire	6	
Central Bedfordshire	6	
Essex	6	
Hertfordshire	6	
Luton	6	
Norfolk	6	
Peterborough	6	
Southend-on-Sea	6	
Suffolk	6	
Thurrock	6	
Greater London	5	
Barking and Dagenham	6	
Barnet	6	
Bexley	6	
Brent	6	
Bromley	6	
Camden	6	
Croydon	6	
Ealing	6	
Enfield	6	
Greenwich	6	
Hackney	6	
Hammersmith and Fulham	6	
Haringey	6	
Harrow	6	
Havering	6	
Hillingdon	6	
Hounslow	6	
Islington	6	
Kensington and Chelsea	6	
Kingston upon Thames	6	
Lambeth	6	
Lewisham	6	

Appendix Table 7. GBD location hierarchy with levels		
Geography	Level	
Merton	6	
Newham	6	
Redbridge	6	
Richmond upon Thames	6	
Southwark	6	
Sutton	6	
Tower Hamlets	6	
Waltham Forest	6	
Wandsworth	6	
Westminster	6	
North East England	5	
County Durham	6	
Darlington	6	
Gateshead	6	
Hartlepool	6	
Middlesbrough	6	
Newcastle upon Tyne	6	
North Tyneside	6	
Northumberland	6	
Redcar and Cleveland	6	
South Tyneside	6	
Stockton-on-Tees	6	
Sunderland	6	
North West England	5	
Blackburn with Darwen	6	
Blackpool	6	
Bolton	6	
Bury	6	
Cheshire East	6	
Cheshire West and Chester	6	
Cumbria	6	
Halton	6	
Knowsley	6	
Lancashire	6	
Liverpool	6	
Manchester	6	
Oldham	6	
Rochdale	6	
Salford	6	
Sefton	6	
St Helens	6	
Stockport	6	
Tameside	6	
Trafford	6	
Warrington	6	

Appendix Table 7. GBD location hierarchy with levels		
Geography	Level	
Wigan	6	
Wirral	6	
South East England	5	
Bracknell Forest	6	
Brighton and Hove	6	
Buckinghamshire	6	
East Sussex	6	
Hampshire	6	
Isle of Wight	6	
Kent	6	
Medway	6	
Milton Keynes	6	
Oxfordshire	6	
Portsmouth	6	
Reading	6	
Slough	6	
Southampton	6	
Surrey	6	
West Berkshire	6	
West Sussex	6	
Windsor and Maidenhead	6	
Wokingham	6	
South West England	5	
Bath and North East Somerset	6	
Bournemouth	6	
Bristol, City of	6	
Cornwall	6	
Devon	6	
Dorset	6	
Gloucestershire	6	
North Somerset	6	
Plymouth	6	
Poole	6	
Somerset	6	
South Gloucestershire	6	
Swindon	6	
Torbay	6	
Wiltshire	6	
West Midlands	5	
Birmingham	6	
Coventry	6	
Dudley	6	
Herefordshire, County of	6	
Sandwell	6	
Shropshire	6	

Appendix Table 7. GBD location hierarchy with levels		
Geography	Level	
Solihull	6	
Staffordshire	6	
Stoke-on-Trent	6	
Telford and Wrekin	6	
Walsall	6	
Warwickshire	6	
Wolverhampton	6	
Worcestershire	6	
Yorkshire and the Humber	5	
Barnsley	6	
Bradford	6	
Calderdale	6	
Doncaster	6	
East Riding of Yorkshire	6	
Kingston upon Hull, City of	6	
Kirklees	6	
Leeds	6	
North East Lincolnshire	6	
North Lincolnshire	6	
North Yorkshire	6	
Rotherham	6	
Sheffield	6	
Wakefield	6	
York	6	
Northern Ireland	4	
Scotland	4	
Wales	4	
Latin America and Caribbean	1	
Andean Latin America	2	
Bolivia	3	
Ecuador	3	
Peru	3	
Caribbean	2	
Antigua and Barbuda	3	
The Bahamas	3	
Barbados	3	
Belize	3	
Bermuda	3	
Cuba	3	
Dominica	3	
Dominican Republic	3	
Grenada	3	
Guyana	3	
Haiti	3	
Jamaica	3	

Appendix Table 7. GBD location hierarchy with levels		
Geography	Level	
Puerto Rico	3	
Saint Lucia	3	
Saint Vincent and the Grenadines	3	
Suriname	3	
Trinidad and Tobago	3	
Virgin Islands	3	
Central Latin America	2	
Colombia	3	
Costa Rica	3	
El Salvador	3	
Guatemala	3	
Honduras	3	
Mexico	3	
Aguascalientes	4	
Baja California	4	
Baja California Sur	4	
Campeche	4	
Chiapas	4	
Chihuahua	4	
Coahuila	4	
Colima	4	
Mexico City	4	
Durango	4	
Guanajuato	4	
Guerrero	4	
Hidalgo	4	
Jalisco	4	
México	4	
Michoacán de Ocampo	4	
Morelos	4	
Nayarit	4	
Nuevo León	4	
Oaxaca	4	
Puebla	4	
Querétaro	4	
Quintana Roo	4	
San Luis Potosí	4	
Sinaloa	4	
Sonora	4	
Tabasco	4	
Tamaulipas	4	
Tlaxcala	4	
Veracruz de Ignacio de la Llave	4	
Yucatán	4	
Zacatecas	4	

Appendix Table 7. GBD location hierarchy with levels			
Geography	Level		
Nicaragua	3		
Panama	3		
Venezuela	3		
Tropical Latin America	2		
Brazil	3		
Acre	4		
Alagoas	4		
Amapá	4		
Amazonas	4		
Bahia	4		
Ceará	4		
Distrito Federal	4		
Espírito Santo	4		
Goiás	4		
Maranhão	4		
Mato Grosso	4		
Mato Grosso do Sul	4		
Minas Gerais	4		
Pará	4		
Paraíba	4		
Paraná	4		
Pernambuco	4		
Piauí	4		
Rio de Janeiro	4		
Rio Grande do Norte	4		
Rio Grande do Sul	4		
Rondônia	4		
Roraima	4		
Santa Catarina	4		
São Paulo	4		
Sergipe	4		
Tocantins	4		
Paraguay	3		
North Africa and Middle East	1		
North Africa and Middle East	2		
Afghanistan	3		
Algeria	3		
Bahrain	3		
Egypt	3		
Iran	3		
Iraq	3		
Jordan	3		
Kuwait	3		
Lebanon	3		
Libya	3		

Geography	Level
Morocco	3
Palestine	3
Oman	3
Qatar	3
Saudi Arabia	3
Sudan	3
Syria	3
Tunisia	3
Turkey	3
United Arab Emirates	3
Yemen	3
South Asia	1
South Asia	2
Bangladesh	3
Bhutan	3
India	3
Andhra Pradesh	4
Arunachal Pradesh	4
Assam	4
Bihar	4
Chhattisgarh	4
Delhi	4
Goa	4
Gujarat	4
Haryana	4
Himachal Pradesh	4
Jammu and Kashmir	4
Jharkhand	4
Karnataka	4
Kerala	4
Madhya Pradesh	4
Maharashtra	4
Manipur	4
Meghalaya	4
Mizoram	4
Nagaland	4
Odisha	4
Punjab	4
Rajasthan	4
Sikkim	4
Tamil Nadu	4
Telangana	4
Tripura	4
Uttar Pradesh	4
Uttarakhand	4

Appendix Table 7. GBD location hierarchy with levels		
Geography	Level	
West Bengal	4	
Union Territories other than Delhi	4	
Nepal	3	
Pakistan	3	
Southeast Asia, East Asia, and Oceania	1	
East Asia	2	
China	3	
North Korea	3	
Taiwan (Province of China)	3	
Oceania	2	
American Samoa	3	
Federated States of Micronesia	3	
Fiji	3	
Guam	3	
Kiribati	3	
Marshall Islands	3	
Northern Mariana Islands	3	
Papua New Guinea	3	
Samoa	3	
Solomon Islands	3	
Tonga	3	
Vanuatu	3	
Southeast Asia	2	
Cambodia	3	
Indonesia	3	
Laos	3	
Malaysia	3	
Maldives	3	
Mauritius	3	
Myanmar	3	
Philippines	3	
Sri Lanka	3	
Seychelles	3	
Thailand	3	
Timor-Leste	3	
Vietnam	3	
Sub-Saharan Africa	1	
Central sub-Saharan Africa	2	
Angola	3	
Central African Republic	3	
Congo (Brazzaville)	3	
DR Congo	3	
Equatorial Guinea	3	
Gabon	3	
Eastern sub-Saharan Africa	2	

Appendix Table 7. GBD location hierarchy with levels		
Geography	Level	
Burundi	3	
Comoros	3	
Djibouti	3	
Eritrea	3	
Ethiopia	3	
Kenya	3	
Baringo	4	
Bomet	4	
Bungoma	4	
Busia	4	
Elgeyo Marakwet	4	
Embu	4	
Garissa	4	
Homa Bay	4	
Isiolo	4	
Kajiado	4	
Kakamega	4	
Kericho	4	
Kiambu	4	
Kilifi	4	
Kirinyaga	4	
Kisii	4	
Kisumu	4	
Kitui	4	
Kwale	4	
Laikipia	4	
Lamu	4	
Machakos	4	
Makueni	4	
Mandera	4	
Marsabit	4	
Meru	4	
Migori	4	
Mombasa	4	
Murang'a	4	
Nairobi	4	
Nakuru	4	
Nandi	4	
Narok	4	
Nyamira	4	
Nyandarua	4	
Nyeri	4	
Samburu	4	
Siaya	4	
Taita Taveta	4	
	I	

Appendix Table 7. GBD location hierarchy with levels		
Geography	Level	
Tana River	4	
Tharaka Nithi	4	
Trans Nzoia	4	
Turkana	4	
Uasin Gishu	4	
Vihiga	4	
Wajir	4	
West Pokot	4	
Madagascar	3	
Malawi	3	
Mozambique	3	
Rwanda	3	
Somalia	3	
South Sudan	3	
Tanzania	3	
Uganda	3	
Zambia	3	
Southern sub-Saharan Africa	2	
Botswana	3	
Lesotho	3	
Namibia	3	
South Africa	3	
Swaziland	3	
Zimbabwe	3	
Western sub-Saharan Africa	2	
Benin	3	
Burkina Faso	3	
Cameroon	3	
Cape Verde	3	
Chad	3	
Cote d'Ivoire	3	
The Gambia	3	
Ghana	3	
Guinea	3	
Guinea-Bissau	3	
Liberia	3	
Mali	3	
Mauritania	3	
Niger	3	
Nigeria	3	
Sao Tome and Principe	3	
Senegal	3	
Sierra Leone	3	
Togo	3	

of metabolic risk factors. We then co	mputed the excess attenuated r	sumated relative risks with and without adjustment acros isk for each mediation isk-cause set.	s all combinations
Risk Factor	Mediator	Cause	Mediation Factor
Lead exposure in bone	High systolic blood pressure	Rheumatic heart disease	(1 to 1)
Lead exposure in bone	High systolic blood pressure	Ischaemic heart disease	1 (1 to 1)
Lead exposure in bone	High systolic blood pressure	Ischaemic stroke	1 (1 to 1)
Lead exposure in bone	High systolic blood pressure	Intracerebral hemorrhage	1 (1 to 1)
Lead exposure in bone	High systolic blood pressure	Subarachnoid hemorrhage	1 (1 to 1)
Lead exposure in bone	High systolic blood pressure	Hypertensive heart disease	1 (1 to 1)
Lead exposure in bone	High systolic blood pressure	Other cardiomyopathy	1 (1 to 1)
Lead exposure in bone	High systolic blood pressure	Atrial fibrillation and flutter	1 (1 to 1)
Lead exposure in bone	High systolic blood pressure	Aortic aneurysm	1 (1 to 1)
Lead exposure in bone	High systolic blood pressure	Peripheral vascular disease	1 (1 to 1)
Lead exposure in bone	High systolic blood pressure	Other cardiovascular and circulatory diseases	1 (1 to 1)
Lead exposure in bone	High systolic blood pressure	Chronic kidney disease due to diabetes mellitus type 1	1 (1 to 1)
Lead exposure in bone	High systolic blood pressure	Chronic kidney disease due to diabetes mellitus type 2	1 (1 to 1)
Lead exposure in bone	High systolic blood pressure	Chronic kidney disease due to hypertension	1 (1 to 1)
Lead exposure in bone	High systolic blood pressure	Chronic kidney disease due to glomerulonephritis	1 (1 to 1)
Lead exposure in bone	High systolic blood pressure	Chronic kidney disease due to other and unspecified causes	1 (1 to 1)
Lead exposure in bone	Impaired kidney function	Chronic kidney disease due to diabetes mellitus type 1	1 (1 to 1)
Lead exposure in bone	Impaired kidney function	Chronic kidney disease due to diabetes mellitus type 2	1 (1 to 1)
Lead exposure in bone	Impaired kidney function	Chronic kidney disease due to hypertension	1 (1 to 1)
Lead exposure in bone	Impaired kidney function	Chronic kidney disease due to glomerulonephritis	1 (1 to 1)
Lead exposure in bone	Impaired kidney function	Chronic kidney disease due to other and unspecified causes	1 (1 to 1)
Smoking	High fasting plasma glucose	Diabetes mellitus type 2	1 (1 to 1)
Smoking	Low bone mineral density	Pedestrian road injuries	1 (1 to 1)
Smoking	Low bone mineral density	Cyclist road injuries	1 (1 to 1)
Smoking	Low bone mineral density	Motor vehicle road injuries	1 (1 to 1)
Smoking	Low bone mineral density	Other road injuries	1 (1 to 1)
Smoking	Low bone mineral density	Other transport injuries	1 (1 to 1)
Smoking	Low bone mineral density	Falls	1 (1 to 1)
Smoking	Low bone mineral density	Other exposure to mechanical forces	1 (1 to 1)
Smoking	Low bone mineral density	Non-venomous animal contact	1 (1 to 1)
Smoking	Low bone mineral density	Assault by other means	1 (1 to 1)
Diet low in fruits	High fasting plasma glucose	Ischaemic stroke	0.05 (0.04 to 0.06)

of metabolic risk factors. We then co	omputed the excess attenuated	risk for each mediation risks with and without adjustment acros	s an comomations
Risk Factor	Mediator	Cause	Mediation Factor
Diet low in fruits	High fasting plasma glucose	Diabetes mellitus type 2	0.99 (0.99 to 0.99)
Diet low in fruits	High LDL cholesterol	Ischaemic heart disease	0.06 (0.05 to 0.08)
Diet low in fruits	High LDL cholesterol	Ischaemic stroke	0.05 (0.04 to 0.06)
Diet low in fruits	High systolic blood pressure	Ischaemic heart disease	0.06 (0.05 to 0.08)
Diet low in fruits	High systolic blood pressure	Ischaemic stroke	0.05 (0.04 to 0.06)
Diet low in fruits	High systolic blood pressure	Intracerebral hemorrhage	0.02 (0.02 to 0.03)
Diet low in fruits	High systolic blood pressure	Subarachnoid hemorrhage	0.02 to 0.03)
Diet low in vegetables	High fasting plasma glucose	Ischaemic heart disease	$(0.02 \ 10 \ 0.03)$ 0.06 $(0.01 \ to 0.2)$
Diet low in vegetables	High fasting plasma glucose	Ischaemic stroke	0.08 (0.04 to 0.16)
Diet low in vegetables	High fasting plasma glucose	Intracerebral hemorrhage	0.08 0.04 to 0.16)
Diet low in vegetables	High fasting plasma glucose	Subarachnoid hemorrhage	0.08 (0.04 to 0.16)
Diet low in vegetables	High LDL cholesterol	Ischaemic heart disease	0.04 0.04 (0.02 to 0.05)
Diet low in vegetables	High LDL cholesterol	Ischaemic stroke	0.09
Diet low in vegetables	High systolic blood pressure	Ischaemic heart disease	0.04
Diet low in vegetables	High systolic blood pressure	Ischaemic stroke	0.03
Diet low in vegetables	High systolic blood pressure	Intracerebral hemorrhage	0.04
Diet low in vegetables	High systolic blood pressure	Subarachnoid hemorrhage	0.02 to 0.05)
Diet low in whole grains	High fasting plasma glucose	Diabetes mellitus type 2	(0.02100.03)
Diet low in whole grains	High LDL cholesterol	Ischaemic heart disease	0.39
Diet low in whole grains	High LDL cholesterol	Ischaemic stroke	0.16
Diet low in nuts and seeds	High fasting plasma glucose	Ischaemic heart disease	0.03
Diet low in nuts and seeds	High fasting plasma glucose	Diabetes mellitus type 2	0.99
Diet low in nuts and seeds	High LDL cholesterol	Ischaemic heart disease	0.28
Diet low in nuts and seeds	High systolic blood pressure	Ischaemic heart disease	(0.01 to 1.62) 0.34
Diet low in milk	Diet low in calcium	Colon and rectum cancer	(0.24 to 0.47)
Diet high in red meat	High fasting plasma glucose	Diabetes mellitus type 2	(1 to 1) 1
Diet high in processed meat	High fasting plasma glucose	Ischaemic heart disease	(1 to 1) 0.01
Diet high in processed meat	High fasting plasma glucose	Diabatec malitus type 2	(0.01 to 0.02) 1
Diat high in guess sweetened haven	Ligh fasting plasma glucose	Isabaamia haart dicassa	(1 to 1) 0.15
Diet hich in sugar-sweetened beverages	Itigh fasting plasma glucose	Dick store wellitere true 2	(0.1 to 0.2) 1
Diet nign in sugar-sweetened beverages	High fasting plasma glucose	Diabetes mellitus type 2	(1 to 1) 0.1
Diet high in sugar-sweetened beverages	High LDL cholesterol	Ischaemic heart disease	(0.05 to 0.15) 0.31
Diet high in sugar-sweetened beverages	High systolic blood pressure	Ischaemic heart disease	(0.28 to 0.34)

of metabolic risk factors. We then cor	nputed the excess attenuated ri	sk for each mediationrisk-cause set.	Mediation Factor
Diet high in sugar-sweetened beverages	High body-mass index	Ischaemic heart disease	
Diet high in sugar-sweetened beverages	High body-mass index	Diabetes mellitus type 2	
Diet low in fibre	Diet low in fruits	Ischaemic heart disease	(1 to 1)
Diet low in fibre	Diet low in vegetables	Ischaemic heart disease	(1 to 1) 1
Diet low in fibre	Diet low in whole grains	Ischaemic heart disease	(1 to 1) 1
Diet low in seafood omega-3 fatty acids	High systolic blood pressure	Ischaemic heart disease	(1 to 1) 0.01
Diet low in polyunsaturated fatty acids	High fasting plasma glucose	Ischaemic heart disease	(0 to 0.02) 0.57
Diet low in polyunsaturated fatty acids	High systolic blood pressure	Ischaemic heart disease	(0.39 to 0.77) 0.72
Diet high in trans fatty acids	High I DL cholesterol	Ischaemic heart disease	(0.57 to 0.89) 0.15
Diet high in trans fatty acids	High systelia blood prossure	Isohaomia haart disaasa	(0.02 to 0.24) 0.15
Diet high in radium	High systeme blood pressure		(0.02 to 0.24) 1
Diet nigh in sodium	High systolic blood pressure	Rheumatic heart disease	(1 to 1) 1
Diet high in sodium	High systolic blood pressure	Ischaemic heart disease	(1 to 1) 1
Diet high in sodium	High systolic blood pressure	Ischaemic stroke	(1 to 1)
Diet high in sodium	High systolic blood pressure	Intracerebral hemorrhage	(1 to 1)
Diet high in sodium	High systolic blood pressure	Subarachnoid hemorrhage	(1 to 1)
Diet high in sodium	High systolic blood pressure	Hypertensive heart disease	(1 to 1)
Diet high in sodium	High systolic blood pressure	Other cardiomyopathy	(1 to 1)
Diet high in sodium	High systolic blood pressure	Atrial fibrillation and flutter	(1 to 1)
Diet high in sodium	High systolic blood pressure	Aortic aneurysm	1 (1 to 1)
Diet high in sodium	High systolic blood pressure	Peripheral vascular disease	1 (1 to 1)
Diet high in sodium	High systolic blood pressure	Other cardiovascular and circulatory diseases	1 (1 to 1)
Diet high in sodium	High systolic blood pressure	Chronic kidney disease due to diabetes mellitus type 1	1 (1 to 1)
Diet high in sodium	High systolic blood pressure	Chronic kidney disease due to diabetes mellitus type 2	1 (1 to 1)
Diet high in sodium	High systolic blood pressure	Chronic kidney disease due to hypertension	1 (1 to 1)
Diet high in sodium	High systolic blood pressure	Chronic kidney disease due to glomerulonephritis	1 (1 to 1)
Diet high in sodium	High systolic blood pressure	Chronic kidney disease due to other and unspecified causes	1 (1 to 1)
Diet high in sodium	Impaired kidney function	Chronic kidney disease due to diabetes mellitus type 1	1 (1 to 1)
Diet high in sodium	Impaired kidney function	Chronic kidney disease due to diabetes mellitus type 2	(1 to 1) (1 to 1)
Diet high in sodium	Impaired kidney function	Chronic kidney disease due to hypertension	(1 to 1)
Diet high in sodium	Impaired kidney function	Chronic kidney disease due to glomerulonephritis	(1 (0 1))
Diet high in sodium	Impaired kidney function	Chronic kidney disease due to other and unspecified causes	$\begin{pmatrix} 1 & 0 & 1 \end{pmatrix}$
Childhood sexual abuse	Alcohol use	Alcohol use disorders	(1 to 1) 1

Risk Factor	Mediator	Cause	Mediation Factor
Low physical activity	High fasting plasma glucose	Ischaemic heart disease	0.14 (0.11 to 0.18)
Low physical activity	High fasting plasma glucose	Ischaemic stroke	0.08 (0.03 to 0.14)
Low physical activity	High fasting plasma glucose	Diabetes mellitus type 2	1
High fasting plasma glucose	High LDL cholesterol	Ischaemic heart disease	0.04
High fasting plasma glucose	High LDL cholesterol	Ischaemic stroke	0.04
High fasting plasma glucose	High systolic blood pressure	Ischaemic heart disease	(0.03 to 0.06) 0.1
High fasting plasma glucose	High systelic blood pressure	Ischaemic stroke	(0.08 to 0.11) 0.15
High fasting plasma glucose	Lligh systelia blood pressure		(0.14 to 0.17) 0.15
			(0.14 to 0.17) 0.15
High fasting plasma glucose	High systolic blood pressure	Subarachnoid hemorrhage	(0.14 to 0.17)
High fasting plasma glucose	Impaired kidney function	Chronic kidney disease due to diabetes mellitus type 1	(1 to 1)
High fasting plasma glucose	Impaired kidney function	Chronic kidney disease due to diabetes mellitus type 2	(1 to 1)
High fasting plasma glucose	Impaired kidney function	Chronic kidney disease due to hypertension	(1 to 1)
High fasting plasma glucose	Impaired kidney function	Chronic kidney disease due to glomerulonephritis	1 (1 to 1)
High fasting plasma glucose	Impaired kidney function	Chronic kidney disease due to other and unspecified causes	1 (1 to 1)
High LDL cholesterol	High systolic blood pressure	Ischaemic heart disease	0.09 (0.07 to 0.11)
High LDL cholesterol	High systolic blood pressure	Ischaemic stroke	0.16 (0.14 to 0.18)
High systolic blood pressure	Impaired kidney function	Chronic kidney disease due to diabetes mellitus type 1	1 (1 to 1)
High systolic blood pressure	Impaired kidney function	Chronic kidney disease due to diabetes mellitus type 2	1 (1 to 1)
High systolic blood pressure	Impaired kidney function	Chronic kidney disease due to hypertension	1 (1 to 1)
High systolic blood pressure	Impaired kidney function	Chronic kidney disease due to glomerulonephritis	(1 to 1)
High systolic blood pressure	Impaired kidney function	Chronic kidney disease due to other and unspecified causes	(1 to 1)
High body-mass index	High fasting plasma glucose	Ischaemic heart disease	0.15
High body-mass index	High fasting plasma glucose	Ischaemic stroke	0.22
High body-mass index	High fasting plasma glucose	Intracerebral hemorrhage	0.22
High body-mass index	High fasting plasma glucose	Subarachnoid hemorrhage	(0.13 to 0.32) 0.22
High body-mass index	High fasting plasma glucose	Diabetes mellitus type 2	(0.13 to 0.32) 1
High body mass index	High I DL abalastaral	Isohoomia haart digaga	(1 to 1) 0.1
			(0.05 to 0.15) 0.03
riigii oody-mass index		Ischaeinic stroke	(0 to 0.08) 0.31
High body-mass index	High systolic blood pressure	Ischaemic heart disease	(0.28 to 0.34)
High body-mass index	High systolic blood pressure	Ischaemic stroke	(0.57 to 0.72)
High body-mass index	High systolic blood pressure	Intracerebral hemorrhage	(0.58 to 0.73)
High body-mass index	High systolic blood pressure	Subarachnoid hemorrhage	0.65 (0.58 to 0.73)

For IHD, stroke, and diabetes we pooled all available cohorts and estimated relative risks with and without adjustment across all combinations						
of metabolic risk factors. We then com	of metabolic risk factors. We then computed the excess attenuated risk for each mediationrisk-cause set.					
Risk Factor	Mediator	Cause	Mediation Factor			
High body-mass index	High systolic blood pressure	Hypertensive heart disease	1 (1 to 1)			
High body-mass index	High systolic blood pressure	Atrial fibrillation and flutter	0.31 (0.28 to 0.34)			
High body-mass index	Impaired kidney function	Chronic kidney disease due to diabetes mellitus type 2	1 (1 to 1)			
High body-mass index	Impaired kidney function	Chronic kidney disease due to hypertension	1 (1 to 1)			
High body-mass index	Impaired kidney function	Chronic kidney disease due to glomerulonephritis	1 (1 to 1)			
High body-mass index	Impaired kidney function	Chronic kidney disease due to other and unspecified causes	1 (1 to 1)			

Geography	2017 SDI	SDI Quintile
Global	0.652	
Central Europe, Eastern Europe, and Central Asia	0.766	
Central Asia	0.673	
Armenia	0.702	High-middle SDI
Azerbaijan	0.701	High-middle SDI
Georgia	0.7	High-middle SDI
Kazakhstan	0.735	High-middle SDI
Kyrgyzstan	0.607	Low-middle SDI
Mongolia	0.662	Middle SDI
Tajikistan	0.523	Low-middle SDI
Turkmenistan	0.696	Middle SDI
Uzbekistan	0.63	Middle SDI
Central Europe	0.814	
Albania	0.685	Middle SDI
Bosnia and Herzegovina	0.713	High-middle SDI
Bulgaria	0.792	High-middle SDI
Croatia	0.825	High SDI
Czech Republic	0.851	High SDI
Hungary	0.817	High-middle SDI
Macedonia	0.754	High-middle SDI
Montenegro	0.788	High-middle SDI
Poland	0.844	High SDI
Romania	0.784	High-middle SDI
Serbia	0.752	High-middle SDI
Slovakia	0.842	High SDI
Slovenia	0.86	High SDI
Eastern Europe	0.785	
Belarus	0.773	High-middle SDI
Estonia	0.858	High SDI
Latvia	0.825	High SDI
Lithuania	0.841	High SDI
Moldova	0.676	Middle SDI
Russian Federation	0.792	High-middle SDI
Ukraine	0.74	High-middle SDI
High-income	0.854	
Australasia	0.869	
Australia	0.873	High SDI
New Zealand	0.842	High SDI
High-income Asia-Pacific	0.869	
Brunei	0.856	High SDI
Japan	0.865	High SDI
Aichi	0.875	High SDI
Akita	0.829	High SDI
Aomori	0.825	High SDI
Chiba	0.859	High SDI

Appendix Table 9. Socio-Demographic Index groupings by geography, based on 2017 values			
Geography	2017 SDI	SDI Quintile	
Ehime	0.838	High SDI	
Fukui	0.852	High SDI	
Fukuoka	0.855	High SDI	
Fukushima	0.831	High SDI	
Gifu	0.849	High SDI	
Gunma	0.851	High SDI	
Hiroshima	0.863	High SDI	
Hokkaidō	0.842	High SDI	
Hyōgo	0.86	High SDI	
Ibaraki	0.851	High SDI	
Ishikawa	0.856	High SDI	
Iwate	0.825	High SDI	
Kagawa	0.85	High SDI	
Kagoshima	0.83	High SDI	
Kanagawa	0.875	High SDI	
Kōchi	0.825	High SDI	
Kumamoto	0.832	High SDI	
Kyōto	0.873	High SDI	
Mie	0.854	High SDI	
Miyagi	0.85	High SDI	
Miyazaki	0.823	High SDI	
Nagano	0.851	High SDI	
Nagasaki	0.826	High SDI	
Nara	0.848	High SDI	
Niigata	0.843	High SDI	
Ōita	0.846	High SDI	
Okayama	0.856	High SDI	
Okinawa	0.818	High SDI	
Ōsaka	0.872	High SDI	
Saga	0.834	High SDI	
Saitama	0.852	High SDI	
Shiga	0.871	High SDI	
Shimane	0.831	High SDI	
Shizuoka	0.859	High SDI	
Tochigi	0.853	High SDI	
Tokushima	0.845	High SDI	
Tōkyō	0.924	High SDI	
Tottori	0.834	High SDI	
Toyama	0.86	High SDI	
Wakayama	0.84	High SDI	
Yamagata	0.832	High SDI	
Yamaguchi	0.849	High SDI	
Yamanashi	0.854	High SDI	
South Korea	0.872	High SDI	
Singapore	0.872	High SDI	

Geography	2017 SDI	SDI Quintile
High-income North America	0.868	
Canada	0.882	High SDI
Greenland	0.76	High-middle SDI
USA	0.867	High SDI
Alabama	0.837	High SDI
Alaska	0.861	High SDI
Arizona	0.845	High SDI
Arkansas	0.826	High SDI
California	0.872	High SDI
Colorado	0.882	High SDI
Connecticut	0.906	High SDI
Delaware	0.874	High SDI
Washington, DC	0.89	High SDI
Florida	0.864	High SDI
Georgia	0.848	High SDI
Hawaii	0.872	High SDI
Idaho	0.841	High SDI
Illinois	0.879	High SDI
Indiana	0.848	High SDI
Iowa	0.87	High SDI
Kansas	0.864	High SDI
Kentucky	0.831	High SDI
Louisiana	0.835	High SDI
Maine	0.872	High SDI
Maryland	0.896	High SDI
Massachusetts	0.913	High SDI
Michigan	0.868	High SDI
Minnesota	0.893	High SDI
Mississippi	0.819	High SDI
Missouri	0.853	High SDI
Montana	0.863	High SDI
Nebraska	0.873	High SDI
Nevada	0.847	High SDI
New Hampshire	0.904	High SDI
New Jersey	0.899	High SDI
New Mexico	0.835	High SDI
New York	0.893	High SDI
North Carolina	0.85	High SDI
North Dakota	0.88	High SDI
Ohio	0.858	High SDI
Oklahoma	0.838	High SDI
Oregon	0.871	High SDI
Pennsylvania	0.879	High SDI
Rhode Island	0.89	High SDI
South Carolina	0.846	High SDI

Appendix Table 9. Socio-Demographic Index groupings by geography, based on 2017 values			
Geography	2017 SDI	SDI Quintile	
South Dakota	0.86	High SDI	
Tennessee	0.837	High SDI	
Texas	0.838	High SDI	
Utah	0.856	High SDI	
Vermont	0.896	High SDI	
Virginia	0.885	High SDI	
Washington	0.884	High SDI	
West Virginia	0.825	High SDI	
Wisconsin	0.878	High SDI	
Wyoming	0.869	High SDI	
Southern Latin America	0.72		
Argentina	0.71	High-middle SDI	
Chile	0.748	High-middle SDI	
Uruguay	0.707	High-middle SDI	
Western Europe	0.857		
Andorra	0.902	High SDI	
Austria	0.866	High SDI	
Belgium	0.886	High SDI	
Cyprus	0.865	High SDI	
Denmark	0.918	High SDI	
Finland	0.893	High SDI	
France	0.865	High SDI	
Germany	0.87	High SDI	
Greece	0.817	High SDI	
Iceland	0.907	High SDI	
Ireland	0.882	High SDI	
Israel	0.816	High-middle SDI	
Italy	0.843	High SDI	
Luxembourg	0.916	High SDI	
Malta	0.836	High SDI	
Netherlands	0.912	High SDI	
Norway	0.911	High SDI	
Portugal	0.778	High-middle SDI	
Spain	0.825	High SDI	
Sweden	0.883	High SDI	
Stockholm	0.914	High SDI	
Sweden except Stockholm	0.873	High SDI	
Switzerland	0.889	High SDI	
United Kingdom	0.843	High SDI	
England	0.849	High SDI	
East Midlands	0.83	High SDI	
Derby	0.846	High SDI	
Derbyshire	0.817	High SDI	
Leicester	0.839	High SDI	
Leicestershire	0.846	High SDI	

Geography	2017 SDI	SDI Quintile
Lincolnshire	0.812	High SDI
Northamptonshire	0.829	High SDI
Nottingham	0.863	High SDI
Nottinghamshire	0.814	High SDI
Rutland	0.833	High SDI
East of England	0.84	High SDI
Bedford	0.838	High SDI
Cambridgeshire	0.871	High SDI
Central Bedfordshire	0.834	High SDI
Essex	0.832	High SDI
Hertfordshire	0.87	High SDI
Luton	0.833	High SDI
Norfolk	0.826	High SDI
Peterborough	0.818	High SDI
Southend-on-Sea	0.811	High SDI
Suffolk	0.821	High SDI
Thurrock	0.807	High SDI
Greater London	0.894	High SDI
Barking and Dagenham	0.802	High SDI
Barnet	0.865	High SDI
Bexley	0.826	High SDI
Brent	0.849	High SDI
Bromley	0.848	High SDI
Camden	0.93	High SDI
Croydon	0.833	High SDI
Ealing	0.865	High SDI
Enfield	0.839	High SDI
Greenwich	0.833	High SDI
Hackney	0.887	High SDI
Hammersmith and Fulham	0.927	High SDI
Haringey	0.854	High SDI
Harrow	0.848	High SDI
Havering	0.824	High SDI
Hillingdon	0.882	High SDI
Hounslow	0.879	High SDI
Islington	0.922	High SDI
Kensington and Chelsea	0.932	High SDI
Kingston upon Thames	0.89	High SDI
Lambeth	0.9	High SDI
Lewisham	0.843	High SDI
Merton	0.873	High SDI
Newham	0.838	High SDI
Redbridge	0.831	High SDI
Richmond upon Thames	0.902	High SDI
Southwark	0.912	High SDI

Appendix Table 9. Socio-Demographic Index groupings by geography, based on 2017 values			
Geography	2017 SDI	SDI Quintile	
Sutton	0.843	High SDI	
Tower Hamlets	0.905	High SDI	
Waltham Forest	0.819	High SDI	
Wandsworth	0.911	High SDI	
Westminster	0.927	High SDI	
North East England	0.821	High SDI	
County Durham	0.81	High SDI	
Darlington	0.825	High SDI	
Gateshead	0.826	High SDI	
Hartlepool	0.793	High SDI	
Middlesbrough	0.808	High SDI	
Newcastle upon Tyne	0.872	High SDI	
North Tyneside	0.825	High SDI	
Northumberland	0.808	High SDI	
Redcar and Cleveland	0.79	High SDI	
South Tyneside	0.794	High SDI	
Stockton-on-Tees	0.823	High SDI	
Sunderland	0.815	High SDI	
North West England	0.834	High SDI	
Blackburn with Darwen	0.802	High SDI	
Blackpool	0.781	High SDI	
Bolton	0.805	High SDI	
Bury	0.815	High SDI	
Cheshire East	0.864	High SDI	
Cheshire West and Chester	0.855	High SDI	
Cumbria	0.828	High SDI	
Halton	0.824	High SDI	
Knowsley	0.816	High SDI	
Lancashire	0.831	High SDI	
Liverpool	0.852	High SDI	
Manchester	0.885	High SDI	
Oldham	0.79	High SDI	
Rochdale	0.795	High SDI	
Salford	0.838	High SDI	
Sefton	0.812	High SDI	
St Helens	0.803	High SDI	
Stockport	0.843	High SDI	
Tameside	0.797	High SDI	
Trafford	0.873	High SDI	
Warrington	0.86	High SDI	
Wigan	0.798	High SDI	
Wirral	0.803	High SDI	
South East England	0.856	High SDI	
Bracknell Forest	0.869	High SDI	
Brighton and Hove	0.885	High SDI	

Buckinghamshire	0.865	High SDI
East Sussex	0.814	High SDI
Hampshire	0.85	High SDI
Isle of Wight	0.814	High SDI
Kent	0.828	High SDI
Medway	0.809	High SDI
Milton Keynes	0.86	High SDI
Oxfordshire	0.879	High SDI
Portsmouth	0.86	High SDI
Reading	0.895	High SDI
Slough	0.859	High SDI
Southampton	0.858	High SDI
Surrey	0.883	High SDI
West Berkshire	0.872	High SDI
West Sussex	0.843	High SDI
Windsor and Maidenhead	0.889	High SDI
Wokingham	0.885	High SDI
South West England	0.841	High SDI
Bath and North East Somerset	0.875	High SDI
Bournemouth	0.858	High SDI
Bristol, City of	0.884	High SDI
Cornwall	0.817	High SDI
Devon	0.837	High SDI
Dorset	0.825	High SDI
Gloucestershire	0.85	High SDI
North Somerset	0.832	High SDI
Plymouth	0.836	High SDI
Poole	0.842	High SDI
Somerset	0.816	High SDI
South Gloucestershire	0.867	High SDI
Swindon	0.847	High SDI
Torbay	0.79	High SDI
Wiltshire	0.829	High SDI
West Midlands	0.829	High SDI
Birmingham	0.84	High SDI
Coventry	0.848	High SDI
Dudley	0.799	High SDI
Herefordshire, County of	0.828	High SDI
Sandwell	0.797	High SDI
Shropshire	0.832	High SDI
Solihull	0.855	High SDI
Staffordshire	0.826	High SDI
Stoke-on-Trent	0.804	High SDI
Telford and Wrekin	0.822	High SDI

Appendix Table 9. Socio-Demographic Index groupings by geography, based on 2017 values			
Geography	2017 SDI	SDI Quintile	
Warwickshire	0.857	High SDI	
Wolverhampton	0.811	High SDI	
Worcestershire	0.833	High SDI	
Yorkshire and the Humber	0.83	High SDI	
Barnsley	0.787	High SDI	
Bradford	0.807	High SDI	
Calderdale	0.827	High SDI	
Doncaster	0.791	High SDI	
East Riding of Yorkshire	0.822	High SDI	
Kingston upon Hull, City of	0.813	High SDI	
Kirklees	0.816	High SDI	
Leeds	0.868	High SDI	
North East Lincolnshire	0.804	High SDI	
North Lincolnshire	0.811	High SDI	
North Yorkshire	0.839	High SDI	
Rotherham	0.796	High SDI	
Sheffield	0.853	High SDI	
Wakefield	0.806	High SDI	
York	0.879	High SDI	
Northern Ireland	0.835	High SDI	
Scotland	0.805	High SDI	
Wales	0.806	High SDI	
Latin America and Caribbean	0.64		
Andean Latin America	0.628		
Bolivia	0.587	Low-middle SDI	
Ecuador	0.636	Middle SDI	
Peru	0.636	Middle SDI	
Caribbean	0.638		
Antigua and Barbuda	0.715	High-middle SDI	
The Bahamas	0.756	High-middle SDI	
Barbados	0.739	High-middle SDI	
Belize	0.602	Low-middle SDI	
Bermuda	0.805	High-middle SDI	
Cuba	0.688	Middle SDI	
Dominica	0.687	Middle SDI	
Dominican Republic	0.593	Low-middle SDI	
Grenada	0.64	Middle SDI	
Guyana	0.584	Low-middle SDI	
Haiti	0.442	Low SDI	
Jamaica	0.679	Middle SDI	
Puerto Rico	0.813	High-middle SDI	
Saint Lucia	0.653	Middle SDI	
Saint Vincent and the Grenadines	0.608	Middle SDI	
Suriname	0.641	Middle SDI	
Trinidad and Tobago	0.698	Middle SDI	
Geography	2017 SDI	SDI Quintile	
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Virgin Islands	0.807	High-middle SDI	
Central Latin America	0.623		
Colombia	0.634	Middle SDI	
Costa Rica	0.662	Middle SDI	
El Salvador	0.593	Low-middle SDI	
Guatemala	0.524	Low-middle SDI	
Honduras	0.512	Low-middle SDI	
Mexico	0.628	Middle SDI	
Aguascalientes	0.659	Middle SDI	
Baja California	0.657	Middle SDI	
Baja California Sur	0.659	Middle SDI	
Campeche	0.616	Middle SDI	
Chiapas	0.533	Middle SDI	
Chihuahua	0.639	Middle SDI	
Coahuila	0.645	Middle SDI	
Colima	0.654	Middle SDI	
Mexico City	0.716	Middle SDI	
Durango	0.624	Middle SDI	
Guanajuato	0.621	Middle SDI	
Guerrero	0.562	Middle SDI	
Hidalgo	0.587	Middle SDI	
Jalisco	0.649	Middle SDI	
México	0.635	Middle SDI	
Michoacán de Ocampo	0.586	Middle SDI	
Morelos	0.635	Middle SDI	
Nayarit	0.62	Middle SDI	
Nuevo León	0.677	Middle SDI	
Oaxaca	0.561	Middle SDI	
Puebla	0.584	Middle SDI	
Querétaro	0.639	Middle SDI	
Quintana Roo	0.626	Middle SDI	
San Luis Potosí	0.621	Middle SDI	
Sinaloa	0.649	Middle SDI	
Sonora	0.65	Middle SDI	
Tabasco	0.611	Middle SDI	
Tamaulipas	0.647	Middle SDI	
Tlaxcala	0.604	Middle SDI	
Veracruz de Ignacio de la Llave	0.592	Middle SDI	
Yucatán	0.63	Middle SDI	
Zacatecas	0.608	Middle SDI	
Nicaragua	0.53	Low-middle SDI	
Panama	0.677	Middle SDI	
Venezuela	0.655	Middle SDI	
Tropical Latin America	0.662		
Brazil	0.663	Middle SDI	

Appendix Table 9. Socio-Demographic I	index groupings by geography, bas	ed on 2017 values
Geography	2017 SDI	SDI Quintile
Acre	0.602	Low-middle SDI
Alagoas	0.556	Low-middle SDI
Amapá	0.659	Middle SDI
Amazonas	0.629	Middle SDI
Bahia	0.591	Low-middle SDI
Ceará	0.6	Low-middle SDI
Distrito Federal	0.792	High-middle SDI
Espírito Santo	0.677	Middle SDI
Goiás	0.65	Middle SDI
Maranhão	0.507	Low-middle SDI
Mato Grosso	0.662	Middle SDI
Mato Grosso do Sul	0.65	Middle SDI
Minas Gerais	0.661	Middle SDI
Pará	0.579	Low-middle SDI
Paraíba	0.574	Low-middle SDI
Paraná	0.682	Middle SDI
Pernambuco	0.594	Low-middle SDI
Piauí	0.552	Low-middle SDI
Rio de Janeiro	0.709	High-middle SDI
Rio Grande do Norte	0.605	Low-middle SDI
Rio Grande do Sul	0.693	Middle SDI
Rondônia	0.622	Middle SDI
Roraima	0.646	Middle SDI
Santa Catarina	0.702	High-middle SDI
São Paulo	0.72	High-middle SDI
Sergipe	0.616	Middle SDI
Tocantins	0.611	Middle SDI
Paraguay	0.619	Middle SDI
North Africa and Middle East	0.639	
North Africa and Middle East	0.639	
Afghanistan	0.29	Low SDI
Algeria	0.696	Middle SDI
Bahrain	0.712	High-middle SDI
Egypt	0.604	Low-middle SDI
Iran	0.7	High-middle SDI
Iraq	0.585	Low-middle SDI
Jordan	0.697	Middle SDI
Kuwait	0.786	High-middle SDI
Lebanon	0.73	High-middle SDI
Libya	0.761	High-middle SDI
Morocco	0.579	Low-middle SDI
Palestine	0 541	Low-middle SDI
Oman	0.744	High-middle SDI
Qatar	0.766	High-middle SDI
Saudi Arabia	0.779	High-middle SDI

Geography	2017 SDI	SDI Quintile
Sudan	0.478	Low-middle SDI
Syria	0.611	Middle SDI
Tunisia	0.675	Middle SDI
Turkey	0.729	High-middle SDI
United Arab Emirates	0.795	High-middle SDI
Yemen	0.43	Low SDI
outh Asia	0.534	
South Asia	0.534	
Bangladesh	0.458	Low SDI
Bhutan	0.57	Low-middle SDI
India	0.55	Low-middle SDI
Andhra Pradesh	0.536	Low-middle SDI
Arunachal Pradesh	0.556	Low-middle SDI
Assam	0.53	Low-middle SDI
Bihar	0.433	Low SDI
Chhattisgarh	0.512	Low-middle SDI
Delhi	0.715	High-middle SDI
Goa	0.74	High-middle SDI
Gujarat	0.584	Low-middle SDI
Haryana	0.6	Low-middle SDI
Himachal Pradesh	0.633	Middle SDI
Jammu and Kashmir	0.59	Low-middle SDI
Jharkhand	0.487	Low-middle SDI
Karnataka	0.574	Low-middle SDI
Kerala	0.659	Middle SDI
Madhya Pradesh	0.487	Low-middle SDI
Maharashtra	0.618	Middle SDI
Manipur	0.59	Low-middle SDI
Meghalaya	0.565	Low-middle SDI
Mizoram	0.616	Middle SDI
Nagaland	0.633	Middle SDI
Odisha	0.524	Low-middle SDI
Punjab	0.622	Middle SDI
Rajasthan	0.492	Low-middle SDI
Sikkim	0.628	Middle SDI
Tamil Nadu	0.615	Middle SDI
Telangana	0.575	Low-middle SDI
Tripura	0.543	Low-middle SDI
Uttar Pradesh	0.488	Low-middle SDI
Uttarakhand	0.607	Middle SDI
West Bengal	0.538	Low-middle SDI
Union Territories other than Delhi	0.653	Middle SDI
Nepal	0.429	Low SDI
Pakistan	0.492	Low-middle SDI

Appendix Table 9. Socio-Demographic I	ndex groupings by geography, bas	ed on 2017 values
Geography	2017 SDI	SDI Quintile
East Asia	0.709	
China	0.707	High-middle SDI
North Korea	0.538	Low-middle SDI
Taiwan (Province of China)	0.864	High SDI
Oceania	0.471	
American Samoa	0.702	High-middle SDI
Federated States of Micronesia	0.575	Low-middle SDI
Fiji	0.641	Middle SDI
Guam	0.794	High-middle SDI
Kiribati	0.427	Low SDI
Marshall Islands	0.55	Low-middle SDI
Northern Mariana Islands	0.758	High-middle SDI
Papua New Guinea	0.419	Low SDI
Samoa	0.576	Low-middle SDI
Solomon Islands	0.425	Low SDI
Tonga	0.625	Middle SDI
Vanuatu	0.475	Low-middle SDI
Southeast Asia	0.641	
Cambodia	0.482	Low-middle SDI
Indonesia	0.648	Middle SDI
Laos	0.519	Low-middle SDI
Malavsia	0.759	High-middle SDI
Maldives	0.655	Middle SDI
Mauritius	0.72	High-middle SDI
Mvanmar	0.556	Low-middle SDI
Philippines	0.617	Middle SDI
Sri Lanka	0.68	Middle SDI
Seychelles	0.692	Middle SDI
Thailand	0.684	Middle SDI
Timor-Leste	0.505	Low-middle SDI
Vietnam	0.607	Middle SDI
Sub-Saharan Africa	0.446	
Central sub-Saharan Africa	0.457	
Angola	0.461	Low-middle SDI
Central African Republic	0.334	Low SDI
Congo (Brazzaville)	0.574	Low-middle SDI
DR Congo	0.364	Low SDI
Equatorial Guinea	0.625	Middle SDI
Gabon	0.651	Middle SDI
Eastern sub-Sabaran Africa	0.387	Wildle 5D1
Burundi	0.31	Low SDI
Comoros	0.434	Low SDI
Diibouti	0.495	Low-middle SDI
Eritro	0.400	
Ethiopia	0.224	
Europia	0.534	LOW SD1

Geography	2017 SDI	SDI Quintile
Kenya	0.499	Low-middle SDI
Baringo	0.444	Low-middle SDI
Bomet	0.496	Low-middle SDI
Bungoma	0.463	Low-middle SDI
Busia	0.438	Low-middle SDI
Elgeyo Marakwet	0.496	Low-middle SDI
Embu	0.533	Low-middle SDI
Garissa	0.334	Low-middle SDI
Homa Bay	0.425	Low-middle SDI
Isiolo	0.385	Low-middle SDI
Kajiado	0.534	Low-middle SDI
Kakamega	0.45	Low-middle SDI
Kericho	0.5	Low-middle SDI
Kiambu	0.58	Low-middle SDI
Kilifi	0.456	Low-middle SDI
Kirinyaga	0.533	Low-middle SDI
Kisii	0.522	Low-middle SDI
Kisumu	0.503	Low-middle SDI
Kitui	0.461	Low-middle SDI
Kwale	0.457	Low-middle SDI
Laikipia	0.556	Low-middle SDI
Lamu	0.453	Low-middle SDI
Machakos	0.518	Low-middle SDI
Makueni	0.469	Low-middle SDI
Mandera	0.295	Low-middle SDI
Marsabit	0.34	Low-middle SDI
Meru	0.508	Low-middle SDI
Migori	0.419	Low-middle SDI
Mombasa	0.568	Low-middle SDI
Murang'a	0.528	Low-middle SDI
Nairobi	0.674	Low-middle SDI
Nakuru	0.545	Low-middle SDI
Nandi	0.501	Low-middle SDI
Narok	0.402	Low-middle SDI
Nyamira	0.544	Low-middle SDI
Nyandarua	0.534	Low-middle SDI
Nyeri	0.554	Low-middle SDI
Samburu	0.308	Low-middle SDI
Siaya	0.46	Low-middle SDI
Taita Taveta	0.529	Low-middle SDI
Tana River	0.379	Low-middle SDI
Tharaka Nithi	0.528	Low-middle SDI
Trans Nzoia	0.496	Low-middle SDI
Turkana	0.295	Low-middle SDI
Uasin Gishu	0.545	Low-middle SDI

Appendix Table 9. Socio-Demographic I	Index groupings by geography, bas	ed on 2017 values
Geography	2017 SDI	SDI Quintile
Vihiga	0.477	Low-middle SDI
Wajir	0.243	Low-middle SDI
West Pokot	0.382	Low-middle SDI
Madagascar	0.331	Low SDI
Malawi	0.349	Low SDI
Mozambique	0.34	Low SDI
Rwanda	0.407	Low SDI
Somalia	0.235	Low SDI
South Sudan	0.275	Low SDI
Tanzania	0.412	Low SDI
Uganda	0.388	Low SDI
Zambia	0.472	Low-middle SDI
Southern sub-Saharan Africa	0.64	
Botswana	0.663	Middle SDI
Lesotho	0.493	Low-middle SDI
Namibia	0.616	Middle SDI
South Africa	0.677	Middle SDI
Swaziland	0.578	Low-middle SDI
Zimbabwe	0.463	Low-middle SDI
Western sub-Saharan Africa	0.441	
Benin	0.373	Low SDI
Burkina Faso	0.284	Low SDI
Cameroon	0.482	Low-middle SDI
Cape Verde	0.549	Low-middle SDI
Chad	0.253	Low SDI
Cote d'Ivoire	0.412	Low SDI
The Gambia	0.405	Low SDI
Ghana	0.537	Low-middle SDI
Guinea	0.325	Low SDI
Guinea-Bissau	0.349	Low SDI
Liberia	0.328	Low SDI
Mali	0.267	Low SDI
Mauritania	0.471	Low-middle SDI
Niger	0.191	Low SDI
Nigeria	0.493	Low-middle SDI
Sao Tome and Principe	0.488	Low-middle SDI
Senegal	0.373	Low SDI
Sierra Leone	0.357	Low SDI
Тодо	0.413	Low SDI

Appendix Table 10: Socio-Demographic	Index val	ues for all	estimated	GBD 201	7 locations	, 1990-201	7					-																
Location	1990	1991	1992	1993	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015	2016	2017
Global	0.523	0.529	0.534	0.539	0.543	0.548	0.553	0.557	0.561	0.566	0.571	0.576	0.581	0.585	0.59	0.595	0.601	0.606	0.611	0.616	0.62	0.624	0.628	0.633	0.639	0.644	0.647	0.652
Central Europe, Eastern Europe, and Central Asia	0.656	0.662	0.67	0.674	0.677	0.682	0.686	0.689	0.691	0.694	0.698	0.701	0.705	0.709	0.715	0.72	0.725	0.73	0.735	0.739	0.743	0.747	0.75	0.753	0.757	0.76	0.763	0.766
Central Asia	0.563	0.567	0.57	0.573	0.575	0.577	0.578	0.579	0.58	0.582	0.585	0.588	0.593	0.598	0.603	0.609	0.615	0.621	0.627	0.633	0.639	0.644	0.649	0.654	0.659	0.664	0.669	0.673
Armenia	0.555	0.559	0.56	0.562	0.565	0.567	0.57	0.573	0.577	0.581	0.586	0.592	0.6	0.61	0.619	0.629	0.639	0.65	0.66	0.667	0.673	0.678	0.683	0.687	0.691	0.695	0.699	0.702
Azerbaijan	0.611	0.614	0.616	0.617	0.610	0.615	0.61	0.607	0.604	0.601	0.6	0.6	0.602	0.605	0.608	0.615	0.625	0.635	0.645	0.654	0.650	0.672	0.671	0.684	0.689	0.694	0.698	0.701
Kazakhstan	0.613	0.615	0.619	0.625	0.632	0.638	0.643	0.645	0.646	0.623	0.651	0.656	0.622	0.624	0.628	0.633	0.683	0.689	0.696	0.034	0.039	0.707	0.708	0.878	0.082	0.723	0.093	0.735
Kurgyzstan	0.565	0.571	0.576	0.578	0.577	0.572	0.569	0.567	0.564	0.562	0.55	0.559	0.56	0.562	0.565	0.566	0.567	0.569	0.572	0.575	0.576	0.581	0.584	0.589	0.594	0.598	0.603	0.607
Mongolia	0.537	0.545	0.55	0.555	0.559	0.564	0.569	0.573	0.577	0.581	0.585	0.589	0.594	0.598	0.603	0.608	0.614	0.619	0.624	0.628	0.632	0.636	0.641	0.646	0.65	0.654	0.658	0.662
Tajikistan	0.474	0.481	0.485	0.487	0.486	0.481	0.474	0.468	0.463	0.459	0.455	0.454	0.456	0.462	0.465	0.466	0.472	0.479	0.483	0.488	0.494	0.501	0.506	0.51	0.514	0.517	0.52	0.523
Turkmenistan	0.588	0.592	0.594	0.599	0.602	0.604	0.606	0.606	0.606	0.607	0.61	0.613	0.617	0.622	0.628	0.635	0.638	0.641	0.644	0.647	0.651	0.657	0.663	0.669	0.678	0.685	0.691	0.696
Uzbekistan	0.481	0.484	0.487	0.493	0.497	0.502	0.508	0.513	0.52	0.526	0.532	0.537	0.543	0.549	0.555	0.56	0.565	0.57	0.575	0.581	0.587	0.592	0.598	0.604	0.611	0.618	0.624	0.63
Central Europe	0.665	0.671	0.677	0.683	0.69	0.698	0.705	0.711	0.717	0.723	0.731	0.738	0.745	0.751	0.757	0.762	0.767	0.772	0.776	0.782	0.788	0.793	0.797	0.802	0.805	0.808	0.811	0.814
Albania	0.548	0.545	0.542	0.541	0.542	0.546	0.552	0.558	0.566	0.577	0.584	0.593	0.602	0.611	0.619	0.627	0.635	0.642	0.648	0.653	0.658	0.661	0.665	0.668	0.672	0.676	0.681	0.685
Bosnia and Herzegovina	0.497	0.499	0.5	0.5	0.501	0.507	0.525	0.549	0.571	0.592	0.607	0.619	0.63	0.639	0.647	0.654	0.66	0.667	0.673	0.679	0.685	0.69	0.694	0.699	0.703	0.706	0.71	0.713
Bulgaria	0.658	0.668	0.676	0.684	0.693	0.699	0.705	0.706	0.704	0.703	0.708	0.715	0.721	0.726	0.731	0.736	0.741	0.746	0.751	0.757	0.765	0.771	0.775	0.778	0.781	0.784	0.788	0.792
Croatia	0.725	0.73	0.732	0.732	0.731	0.731	0.732	0.737	0.743	0.749	0.755	0.762	0.768	0.773	0.778	0.782	0.787	0.792	0.797	0.801	0.805	0.809	0.813	0.816	0.818	0.821	0.823	0.825
Czech Republic	0.711	0.717	0.726	0.74	0.757	0.769	0.777	0.783	0.788	0.794	0.799	0.804	0.809	0.814	0.819	0.823	0.827	0.83	0.833	0.836	0.84	0.843	0.846	0.847	0.848	0.848	0.849	0.851
Hungary	0.678	0.683	0.691	0.699	0.707	0.716	0.724	0.732	0.739	0.745	0.751	0.758	0.764	0.77	0.776	0.781	0.786	0.791	0.795	0.799	0.803	0.806	0.807	0.808	0.809	0.811	0.814	0.817
Martanagro	0.020	0.706	0.03	0.031	0.698	0.696	0.696	0.698	0.034	0.001	0.005	0.711	0.716	0.085	0.726	0.731	0.727	0.709	0.715	0.756	0.724	0.767	0.754	0.739	0.744	0.748	0.731	0.799
Poland	0.662	0.668	0.678	0.686	0.697	0.090	0.714	0.724	0.733	0.741	0.75	0.759	0.767	0.721	0.720	0.784	0.789	0.792	0.797	0.804	0.811	0.818	0.823	0.829	0.833	0.837	0.841	0.788
Romania	0.652	0.66	0.663	0.666	0.671	0.678	0.682	0.685	0.689	0.694	0.7	0.707	0.713	0.718	0.724	0.73	0.734	0.739	0.745	0.751	0.758	0.763	0.768	0.772	0.774	0.777	0.78	0.784
Serbia	0.632	0.638	0.643	0.642	0.641	0.641	0.643	0.648	0.653	0.655	0.661	0.665	0.669	0.675	0.684	0.692	0.699	0.705	0.709	0.713	0.718	0.723	0.729	0.736	0.742	0.747	0.75	0.752
Slovakia	0.684	0.69	0.699	0.71	0.722	0.732	0.74	0.748	0.756	0.764	0.772	0.779	0.784	0.788	0.793	0.798	0.804	0.809	0.814	0.818	0.823	0.828	0.832	0.834	0.836	0.838	0.839	0.842
Slovenia	0.741	0.747	0.753	0.759	0.764	0.769	0.775	0.781	0.788	0.794	0.801	0.808	0.814	0.819	0.824	0.828	0.833	0.837	0.841	0.843	0.846	0.848	0.85	0.852	0.854	0.856	0.858	0.86
Eastern Europe	0.678	0.685	0.694	0.698	0.7	0.704	0.708	0.708	0.71	0.711	0.712	0.713	0.715	0.72	0.727	0.734	0.739	0.745	0.751	0.756	0.761	0.764	0.767	0.772	0.776	0.779	0.783	0.785
Belarus	0.625	0.631	0.636	0.641	0.645	0.647	0.65	0.654	0.657	0.661	0.665	0.67	0.676	0.682	0.689	0.696	0.704	0.712	0.72	0.727	0.733	0.74	0.747	0.753	0.759	0.764	0.769	0.773
Estonia	0.711	0.719	0.728	0.736	0.742	0.746	0.75	0.755	0.761	0.766	0.772	0.778	0.783	0.788	0.794	0.799	0.806	0.813	0.82	0.826	0.832	0.838	0.843	0.847	0.851	0.854	0.856	0.858
Latvia	0.696	0.703	0.712	0.721	0.727	0.731	0.733	0.734	0.735	0.738	0.741	0.745	0.75	0.757	0.763	0.769	0.776	0.783	0.792	0.8	0.806	0.81	0.814	0.816	0.817	0.819	0.822	0.825
Lithuania	0.707	0.71	0.717	0.725	0.728	0.731	0.733	0.736	0.74	0.746	0.753	0.76	0.765	0.772	0.779	0.785	0.79	0.796	0.802	0.808	0.815	0.822	0.828	0.833	0.836	0.838	0.839	0.841
Moldova	0.575	0.578	0.58	0.582	0.583	0.584	0.584	0.582	0.58	0.577	0.574	0.574	0.577	0.582	0.588	0.595	0.602	0.61	0.618	0.624	0.632	0.64	0.647	0.654	0.66	0.666	0.671	0.676
Russian Federation	0.683	0.692	0.704	0.708	0.708	0.714	0.718	0.719	0.72	0.722	0.722	0.722	0.724	0.728	0.734	0.742	0.747	0.752	0.757	0.763	0.768	0.77	0.772	0.777	0.781	0.785	0.789	0.792
Ukraine	0.664	0.667	0.67	0.673	0.675	0.676	0.676	0.675	0.675	0.672	0.672	0.673	0.675	0.68	0.687	0.694	0.7	0.707	0.714	0.717	0.721	0.725	0.729	0.732	0.735	0.756	0.758	0.74
Australasia	0.783	0.774	0.779	0.783	0.787	0.792	0.796	0.798	0.813	0.804	0.807	0.875	0.814	0.817	0.835	0.822	0.825	0.820	0.829	0.832	0.830	0.851	0.855	0.859	0.862	0.851	0.855	0.854
Australia	0.786	0.79	0.793	0.797	0.801	0.805	0.81	0.814	0.818	0.822	0.825	0.829	0.833	0.837	0.84	0.843	0.844	0.845	0.848	0.851	0.854	0.856	0.86	0.864	0.867	0.869	0.871	0.873
New Zealand	0.765	0.768	0.771	0.774	0.777	0.78	0.783	0.786	0.79	0.794	0.798	0.802	0.805	0.807	0.809	0.811	0.811	0.811	0.813	0.816	0.819	0.823	0.828	0.832	0.835	0.838	0.84	0.842
High-income Asia-Pacific	0.783	0.789	0.794	0.799	0.804	0.809	0.813	0.817	0.82	0.823	0.826	0.83	0.833	0.836	0.839	0.842	0.844	0.846	0.849	0.851	0.853	0.856	0.858	0.861	0.863	0.865	0.867	0.869
Brunei	0.728	0.733	0.739	0.745	0.751	0.757	0.763	0.769	0.774	0.779	0.784	0.789	0.795	0.802	0.808	0.814	0.819	0.824	0.828	0.831	0.835	0.838	0.842	0.845	0.848	0.851	0.854	0.856
Japan	0.803	0.807	0.812	0.816	0.82	0.823	0.826	0.829	0.831	0.833	0.834	0.836	0.838	0.84	0.842	0.844	0.846	0.847	0.849	0.851	0.853	0.855	0.857	0.859	0.861	0.862	0.863	0.865
Aichi	0.812	0.816	0.821	0.825	0.829	0.833	0.836	0.839	0.841	0.843	0.844	0.846	0.847	0.85	0.852	0.854	0.855	0.856	0.858	0.86	0.862	0.864	0.866	0.869	0.871	0.872	0.873	0.875
Akita	0.766	0.77	0.775	0.778	0.781	0.785	0.788	0.791	0.793	0.795	0.796	0.798	0.8	0.802	0.805	0.806	0.808	0.81	0.812	0.814	0.816	0.818	0.821	0.823	0.825	0.826	0.827	0.829
Aomori	0.761	0.765	0.77	0.773	0.777	0.78	0.783	0.786	0.788	0.79	0.791	0.793	0.795	0.798	0.801	0.803	0.805	0.807	0.809	0.81	0.812	0.815	0.817	0.819	0.821	0.822	0.823	0.825
Chiba	0.803	0.807	0.812	0.816	0.82	0.823	0.827	0.829	0.831	0.832	0.833	0.834	0.836	0.837	0.839	0.841	0.842	0.843	0.845	0.846	0.848	0.85	0.852	0.854	0.856	0.857	0.858	0.859
Ehime	0.776	0.78	0.785	0.788	0.792	0.796	0.799	0.802	0.804	0.806	0.808	0.809	0.811	0.814	0.816	0.818	0.819	0.82	0.822	0.824	0.825	0.828	0.83	0.832	0.834	0.835	0.836	0.838
Fuku	0.784	0.789	0.794	0.798	0.802	0.806	0.81	0.813	0.815	0.818	0.819	0.821	0.823	0.826	0.828	0.83	0.832	0.834	0.836	0.837	0.839	0.842	0.844	0.840	0.848	0.849	0.85	0.852
Fukushima	0.769	0.773	0.778	0.781	0.784	0.788	0.791	0.793	0.794	0.795	0.797	0.798	0.8	0.803	0.807	0.809	0.811	0.813	0.814	0.816	0.819	0.821	0.823	0.825	0.827	0.828	0.829	0.831
Gifu	0.786	0.79	0.795	0.799	0.803	0.807	0.81	0.813	0.815	0.817	0.818	0.82	0.821	0.824	0.826	0.828	0.829	0.831	0.833	0.835	0.837	0.839	0.841	0.843	0.845	0.846	0.847	0.849
Gunma	0.787	0.791	0.796	0.8	0.803	0.807	0.81	0.813	0.815	0.816	0.818	0.819	0.821	0.824	0.827	0.829	0.831	0.833	0.835	0.837	0.839	0.841	0.843	0.845	0.847	0.848	0.849	0.851
Hiroshima	0.8	0.805	0.809	0.813	0.817	0.821	0.825	0.828	0.83	0.832	0.833	0.835	0.836	0.839	0.841	0.842	0.843	0.844	0.846	0.847	0.849	0.852	0.854	0.856	0.858	0.86	0.861	0.863
Hokkaidō	0.783	0.788	0.792	0.796	0.799	0.803	0.806	0.808	0.809	0.811	0.812	0.813	0.815	0.818	0.82	0.822	0.824	0.825	0.827	0.828	0.83	0.832	0.834	0.836	0.838	0.839	0.84	0.842
Hyōgo	0.798	0.803	0.807	0.811	0.815	0.819	0.822	0.825	0.827	0.829	0.831	0.833	0.835	0.837	0.839	0.841	0.842	0.843	0.845	0.846	0.848	0.85	0.852	0.854	0.856	0.857	0.858	0.86
Ibaraki	0.789	0.793	0.798	0.802	0.805	0.809	0.813	0.815	0.817	0.819	0.82	0.821	0.823	0.826	0.828	0.83	0.831	0.833	0.835	0.837	0.839	0.841	0.843	0.845	0.847	0.848	0.849	0.851
Ishikawa	0.789	0.795	0.8	0.804	0.808	0.812	0.815	0.818	0.82	0.822	0.824	0.826	0.828	0.831	0.833	0.835	0.837	0.839	0.84	0.842	0.844	0.846	0.848	0.85	0.852	0.853	0.854	0.856
Iwate	0.76	0.764	0.768	0.772	0.775	0.779	0.782	0.785	0.787	0.788	0.79	0.791	0.793	0.796	0.799	0.801	0.803	0.806	0.808	0.81	0.812	0.815	0.817	0.819	0.821	0.823	0.823	0.825
Kagawa	0.787	0.792	0.797	0.801	0.804	0.808	0.811	0.814	0.816	0.817	0.819	0.82	0.822	0.825	0.827	0.829	0.831	0.832	0.834	0.835	0.837	0.839	0.842	0.844	0.846	0.847	0.848	0.85
Kagoshima	0.768	0.772	0.777	0.781	0.784	0.788	0.791	0.793	0.795	0.797	0.798	0.799	0.801	0.804	0.806	0.808	0.809	0.81	0.812	0.814	0.816	0.818	0.82	0.823	0.825	0.827	0.828	0.83
Kanagawa Marki	0.818	0.823	0.827	0.831	0.835	0.838	0.842	0.844	0.846	0.848	0.849	0.85	0.852	0.854	0.856	0.857	0.859	0.86	0.862	0.863	0.865	0.866	0.868	0.87	0.871	0.873	0.873	0.875
Kumamoto	0.759	0.763	0.767	0.7/1	0.775	0.779	0.783	0.780	0.788	0.791	0.795	0.795	0.802	0.805	0.803	0.805	0.806	0.808	0.81	0.812	0.813	0.87	0.818	0.82	0.822	0.823	0.824	0.825
Kvöto	0.813	0.817	0.778	0.782	0.83	0.833	0.792	0.839	0.841	0.843	0.845	0.801	0.849	0.803	0.853	0.81	0.811	0.812	0.814	0.815	0.862	0.863	0.822	0.823	0.869	0.829	0.85	0.852
Mic	0.787	0.792	0.797	0.801	0.805	0.81	0.813	0.815	0.818	0.819	0.821	0.823	0.825	0.827	0.83	0.832	0.833	0.835	0.836	0.838	0.84	0.842	0.845	0.847	0.849	0.851	0.852	0.854
Miyagi	0.788	0.793	0.798	0.801	0.805	0.809	0.812	0.814	0.816	0.817	0.818	0.819	0.821	0.824	0.827	0.829	0.831	0.833	0.835	0.836	0.838	0.841	0.843	0.845	0.847	0.848	0.849	0.85
Miyazaki	0.764	0.768	0.772	0.775	0.778	0.781	0.784	0.786	0.788	0.79	0.791	0.793	0.795	0.797	0.8	0.801	0.802	0.804	0.805	0.807	0.809	0.812	0.814	0.816	0.819	0.82	0.821	0.823
Nagano	0.792	0.797	0.801	0.805	0.808	0.811	0.814	0.817	0.818	0.82	0.821	0.823	0.824	0.827	0.829	0.831	0.832	0.834	0.836	0.838	0.839	0.842	0.844	0.845	0.847	0.849	0.849	0.851
Nagasaki	0.766	0.77	0.774	0.777	0.781	0.784	0.787	0.79	0.792	0.794	0.795	0.797	0.799	0.801	0.803	0.805	0.806	0.808	0.809	0.811	0.813	0.815	0.817	0.82	0.822	0.823	0.824	0.826
Nara	0.789	0.794	0.799	0.803	0.807	0.811	0.814	0.817	0.819	0.821	0.822	0.824	0.825	0.827	0.829	0.831	0.832	0.834	0.835	0.836	0.838	0.839	0.841	0.843	0.844	0.845	0.846	0.848

Appendix Table 10: Socio-Demographic	Index val	ues for all	estimated	GBD 201	7 locations	, 1990-201	7					_	_									_		_				
Location	1990	1991	1992	1993	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015	2016	2017
Niigata	0.776	0.78	0.784	0.788	0.792	0.795	0.799	0.801	0.804	0.806	0.808	0.81	0.813	0.815	0.818	0.82	0.822	0.824	0.826	0.828	0.83	0.832	0.835	0.837	0.839	0.84	0.842	0.843
Oita	0.785	0.789	0.795	0.802	0.808	0.809	0.809	0.81	0.812	0.813	0.815	0.816	0.819	0.821	0.824	0.825	0.827	0.828	0.83	0.831	0.833	0.835	0.838	0.84	0.842	0.843	0.844	0.846
Okayama	0.79	0.794	0.799	0.803	0.807	0.811	0.814	0.817	0.819	0.821	0.823	0.824	0.82/	0.829	0.832	0.834	0.836	0.837	0.839	0.841	0.843	0.846	0.848	0.85	0.852	0.853	0.854	0.856
	0.735	0.737	0.823	0.827	0.778	0.834	0.837	0.84	0.842	0.78	0.781	0.785	0.780	0.789	0.793	0.795	0.855	0.799	0.858	0.802	0.861	0.807	0.809	0.812	0.868	0.815	0.810	0.818
Saga	0.773	0.777	0.782	0.785	0.788	0.792	0.795	0.797	0.799	0.8	0.801	0.803	0.805	0.808	0.81	0.812	0.814	0.815	0.817	0.819	0.821	0.823	0.825	0.827	0.83	0.831	0.832	0.834
Saitama	0.793	0.798	0.802	0.806	0.81	0.814	0.818	0.82	0.822	0.824	0.825	0.826	0.827	0.829	0.831	0.833	0.835	0.836	0.838	0.839	0.841	0.843	0.845	0.847	0.848	0.849	0.85	0.852
Shiga	0.804	0.809	0.814	0.819	0.823	0.827	0.83	0.833	0.836	0.838	0.84	0.841	0.843	0.846	0.848	0.85	0.852	0.853	0.855	0.857	0.858	0.861	0.863	0.865	0.867	0.868	0.869	0.871
Shimane	0.762	0.766	0.771	0.775	0.779	0.783	0.787	0.79	0.793	0.795	0.797	0.798	0.801	0.803	0.806	0.808	0.81	0.812	0.814	0.816	0.818	0.82	0.823	0.825	0.827	0.828	0.829	0.831
Shizuoka	0.798	0.802	0.807	0.81	0.814	0.818	0.821	0.823	0.825	0.826	0.827	0.828	0.83	0.833	0.835	0.837	0.838	0.84	0.842	0.843	0.846	0.848	0.85	0.853	0.855	0.856	0.857	0.859
Tochigi	0.787	0.792	0.797	0.801	0.805	0.809	0.813	0.815	0.817	0.819	0.82	0.821	0.823	0.826	0.829	0.831	0.832	0.834	0.836	0.838	0.84	0.842	0.845	0.847	0.849	0.85	0.851	0.853
Tokushima	0.776	0.781	0.786	0.79	0.794	0.798	0.802	0.805	0.807	0.809	0.812	0.814	0.816	0.819	0.822	0.824	0.826	0.828	0.83	0.831	0.833	0.835	0.837	0.839	0.841	0.842	0.843	0.845
Tokyo	0.87	0.875	0.879	0.883	0.887	0.892	0.896	0.899	0.901	0.903	0.905	0.906	0.908	0.91	0.911	0.913	0.914	0.915	0.916	0.917	0.918	0.919	0.92	0.921	0.922	0.923	0.923	0.924
Toyama	0.79	0.775	0.78	0.785	0.787	0.812	0.794	0.818	0.821	0.823	0.802	0.804	0.800	0.832	0.835	0.815	0.839	0.841	0.843	0.819	0.847	0.824	0.852	0.854	0.856	0.857	0.858	0.854
Wakayama	0.775	0.78	0.785	0.789	0.792	0.796	0.8	0.802	0.805	0.806	0.808	0.81	0.812	0.815	0.817	0.819	0.821	0.822	0.824	0.825	0.827	0.829	0.831	0.833	0.836	0.837	0.838	0.84
Yamagata	0.766	0.77	0.775	0.778	0.782	0.785	0.788	0.791	0.792	0.794	0.795	0.796	0.799	0.802	0.805	0.807	0.809	0.812	0.814	0.816	0.819	0.821	0.824	0.826	0.828	0.829	0.83	0.832
Yamaguchi	0.79	0.794	0.799	0.802	0.806	0.809	0.812	0.815	0.817	0.819	0.82	0.822	0.824	0.826	0.828	0.83	0.831	0.832	0.834	0.835	0.837	0.839	0.841	0.843	0.846	0.847	0.848	0.849
Yamanashi	0.791	0.796	0.8	0.804	0.808	0.812	0.815	0.818	0.82	0.822	0.823	0.825	0.827	0.829	0.832	0.834	0.836	0.837	0.839	0.841	0.842	0.844	0.846	0.848	0.85	0.851	0.853	0.854
South Korea	0.713	0.724	0.733	0.742	0.751	0.76	0.768	0.777	0.783	0.79	0.799	0.806	0.814	0.82	0.825	0.83	0.834	0.839	0.843	0.846	0.85	0.853	0.857	0.86	0.864	0.867	0.869	0.872
Singapore	0.736	0.744	0.75	0.758	0.765	0.772	0.78	0.786	0.79	0.79	0.797	0.808	0.815	0.82	0.824	0.827	0.832	0.838	0.844	0.849	0.854	0.857	0.86	0.863	0.865	0.868	0.87	0.872
High-income North America	0.784	0.786	0.789	0.793	0.796	0.8	0.805	0.807	0.809	0.812	0.815	0.82	0.824	0.827	0.829	0.83	0.829	0.832	0.837	0.843	0.848	0.852	0.856	0.859	0.861	0.865	0.867	0.868
Canada	0.802	0.805	0.808	0.811	0.814	0.818	0.823	0.828	0.832	0.836	0.841	0.846	0.85	0.853	0.857	0.859	0.861	0.862	0.864	0.867	0.871	0.874	0.877	0.878	0.879	0.88	0.881	0.882
Greenland	0.671	0.67	0.67	0.671	0.672	0.675	0.679	0.683	0.687	0.691	0.695	0.699	0.703	0.708	0.713	0.717	0.722	0.726	0.731	0.737	0.743	0.747	0.751	0.754	0.756	0.757	0.759	0.76
Alabama	0.745	0.749	0.787	0.79	0.793	0.798	0.803	0.804	0.806	0.809	0.812	0.783	0.821	0.823	0.826	0.826	0.826	0.829	0.854	0.84	0.846	0.85	0.853	0.857	0.839	0.803	0.800	0.867
Alaska	0.755	0.757	0.753	0.767	0.772	0.781	0.788	0.709	0.793	0.795	0.799	0.804	0.808	0.811	0.814	0.813	0.811	0.813	0.818	0.809	0.833	0.838	0.843	0.827	0.851	0.857	0.857	0.857
Arizona	0.751	0.753	0.755	0.758	0.76	0.764	0.769	0.77	0.771	0.772	0.775	0.781	0.784	0.787	0.79	0.791	0.791	0.797	0.806	0.815	0.823	0.828	0.832	0.836	0.839	0.842	0.845	0.845
Arkansas	0.723	0.727	0.733	0.738	0.741	0.746	0.751	0.751	0.753	0.754	0.757	0.763	0.767	0.77	0.773	0.772	0.77	0.774	0.78	0.789	0.797	0.802	0.807	0.811	0.816	0.821	0.825	0.826
California	0.771	0.771	0.773	0.776	0.78	0.787	0.794	0.798	0.803	0.807	0.812	0.819	0.823	0.826	0.829	0.83	0.831	0.835	0.84	0.846	0.852	0.856	0.859	0.863	0.865	0.869	0.871	0.872
Colorado	0.798	0.801	0.804	0.807	0.81	0.815	0.818	0.819	0.82	0.821	0.824	0.83	0.833	0.836	0.839	0.84	0.84	0.844	0.849	0.855	0.861	0.865	0.869	0.872	0.875	0.879	0.881	0.882
Connecticut	0.841	0.844	0.847	0.85	0.853	0.856	0.859	0.861	0.863	0.866	0.869	0.874	0.877	0.879	0.881	0.881	0.881	0.883	0.886	0.89	0.894	0.896	0.899	0.901	0.903	0.905	0.906	0.906
Delaware	0.801	0.804	0.808	0.811	0.813	0.816	0.819	0.819	0.82	0.822	0.825	0.83	0.833	0.836	0.838	0.839	0.838	0.841	0.846	0.852	0.858	0.861	0.864	0.866	0.868	0.871	0.873	0.874
Washington, DC	0.797	0.801	0.806	0.812	0.818	0.826	0.834	0.839	0.845	0.85	0.855	0.861	0.865	0.868	0.869	0.868	0.866	0.866	0.868	0.871	0.874	0.876	0.877	0.88	0.883	0.887	0.89	0.89
Florida	0.774	0.778	0.782	0.786	0.789	0.794	0.799	0.782	0.802	0.804	0.808	0.813	0.817	0.82	0.823	0.824	0.825	0.828	0.834	0.84	0.846	0.85	0.853	0.855	0.858	0.861	0.863	0.864
Hawaii	0.733	0.739	0.764	0.709	0.772	0.778	0.782	0.813	0.815	0.785	0.788	0.795	0.790	0.83	0.801	0.833	0.832	0.834	0.839	0.845	0.851	0.854	0.858	0.857	0.864	0.868	0.840	0.872
Idaho	0.756	0.76	0.764	0.769	0.773	0.778	0.783	0.783	0.784	0.785	0.787	0.791	0.794	0.796	0.799	0.798	0.797	0.801	0.807	0.814	0.821	0.824	0.827	0.83	0.832	0.836	0.84	0.841
Illinois	0.787	0.789	0.792	0.796	0.799	0.805	0.81	0.812	0.816	0.818	0.823	0.828	0.833	0.837	0.84	0.841	0.841	0.845	0.85	0.855	0.861	0.864	0.867	0.87	0.873	0.876	0.879	0.879
Indiana	0.772	0.774	0.778	0.781	0.783	0.787	0.791	0.791	0.793	0.794	0.797	0.802	0.806	0.808	0.81	0.809	0.807	0.809	0.813	0.819	0.825	0.829	0.832	0.836	0.839	0.844	0.847	0.848
Iowa	0.794	0.797	0.8	0.803	0.806	0.809	0.813	0.814	0.816	0.818	0.821	0.825	0.828	0.829	0.832	0.831	0.83	0.832	0.837	0.843	0.848	0.852	0.856	0.86	0.863	0.867	0.87	0.87
Kansas	0.782	0.785	0.788	0.791	0.794	0.798	0.802	0.803	0.804	0.806	0.809	0.814	0.817	0.819	0.821	0.819	0.817	0.819	0.823	0.83	0.837	0.842	0.847	0.852	0.856	0.861	0.864	0.864
Kentucky	0.743	0.747	0.751	0.755	0.759	0.763	0.768	0.769	0.77	0.772	0.776	0.782	0.786	0.789	0.791	0.79	0.787	0.789	0.793	0.799	0.806	0.81	0.814	0.817	0.821	0.827	0.83	0.831
Louisiana	0.735	0.739	0.743	0.748	0.752	0.758	0.762	0.764	0.765	0.767	0.771	0.777	0.782	0.786	0.789	0.788	0.787	0.79	0.796	0.803	0.811	0.815	0.819	0.823	0.826	0.831	0.835	0.835
Mane	0.813	0.795	0.821	0.805	0.809	0.814	0.817	0.838	0.821	0.823	0.826	0.85	0.854	0.857	0.84	0.84	0.84	0.842	0.846	0.851	0.856	0.859	0.886	0.864	0.800	0.869	0.871	0.872
Massachusetts	0.843	0.847	0.85	0.853	0.856	0.86	0.864	0.866	0.869	0.871	0.875	0.88	0.883	0.886	0.888	0.889	0.89	0.893	0.896	0.899	0.902	0.904	0.907	0.909	0.91	0.912	0.913	0.913
Michigan	0.788	0.791	0.795	0.799	0.803	0.808	0.813	0.815	0.817	0.82	0.823	0.829	0.832	0.835	0.838	0.838	0.837	0.839	0.841	0.845	0.849	0.852	0.854	0.857	0.86	0.864	0.867	0.868
Minnesota	0.818	0.821	0.825	0.828	0.83	0.834	0.837	0.839	0.841	0.843	0.845	0.849	0.852	0.855	0.857	0.858	0.858	0.86	0.865	0.87	0.874	0.877	0.881	0.884	0.886	0.89	0.892	0.893
Mississippi	0.719	0.722	0.727	0.732	0.736	0.741	0.746	0.747	0.748	0.75	0.753	0.758	0.762	0.764	0.765	0.764	0.76	0.763	0.77	0.779	0.789	0.794	0.8	0.804	0.809	0.814	0.818	0.819
Missouri	0.77	0.773	0.777	0.781	0.784	0.789	0.793	0.795	0.797	0.799	0.802	0.806	0.81	0.812	0.813	0.812	0.811	0.813	0.818	0.825	0.831	0.835	0.839	0.842	0.846	0.85	0.853	0.853
Montana	0.775	0.779	0.783	0.789	0.793	0.799	0.804	0.805	0.807	0.808	0.81	0.814	0.816	0.819	0.822	0.822	0.821	0.823	0.827	0.833	0.839	0.843	0.847	0.851	0.855	0.859	0.862	0.863
Nebraska	0.795	0.798	0.802	0.805	0.808	0.813	0.817	0.817	0.819	0.819	0.821	0.824	0.827	0.829	0.831	0.831	0.83	0.832	0.836	0.842	0.848	0.853	0.857	0.861	0.865	0.869	0.872	0.873
Nevada New Hampchira	0.76	0.761	0.762	0.765	0.767	0.773	0.779	0.781	0.784	0.787	0.791	0.798	0.802	0.805	0.808	0.809	0.81	0.814	0.82	0.827	0.833	0.836	0.838	0.84	0.842	0.845	0.84/	0.847
New Jersey	0.828	0.831	0.834	0.838	0.841	0.845	0.849	0.85	0.852	0.854	0.857	0.861	0.864	0.867	0.87	0.875	0.871	0.873	0.876	0.881	0.885	0.888	0.89	0.893	0.895	0.902	0.899	0.904
New Mexico	0.733	0.735	0.739	0.743	0.748	0.755	0.76	0.762	0.764	0.765	0.767	0.773	0.777	0.779	0.782	0.781	0.779	0.783	0.79	0.799	0.808	0.814	0.819	0.823	0.828	0.832	0.835	0.835
New York	0.816	0.819	0.822	0.825	0.828	0.832	0.836	0.838	0.841	0.843	0.847	0.852	0.855	0.858	0.86	0.861	0.861	0.864	0.868	0.872	0.877	0.88	0.883	0.886	0.889	0.892	0.893	0.893
North Carolina	0.763	0.766	0.77	0.774	0.777	0.781	0.785	0.785	0.786	0.788	0.791	0.796	0.8	0.802	0.805	0.805	0.805	0.809	0.815	0.822	0.829	0.833	0.836	0.84	0.843	0.846	0.849	0.85
North Dakota	0.79	0.792	0.796	0.799	0.802	0.807	0.813	0.816	0.82	0.823	0.827	0.83	0.833	0.834	0.835	0.835	0.833	0.835	0.839	0.844	0.85	0.854	0.859	0.864	0.87	0.875	0.879	0.88
Ohio	0.78	0.783	0.787	0.79	0.793	0.797	0.801	0.802	0.804	0.806	0.809	0.814	0.818	0.82	0.822	0.822	0.821	0.822	0.826	0.832	0.837	0.841	0.844	0.847	0.85	0.854	0.857	0.858
Oklahoma	0.749	0.751	0.755	0.758	0.76	0.764	0.768	0.768	0.768	0.769	0.772	0.777	0.781	0.784	0.786	0.785	0.784	0.786	0.792	0.8	0.808	0.813	0.818	0.824	0.829	0.835	0.838	0.838
Oregon	0.785	0.788	0.791	0.794	0.797	0.802	0.806	0.808	0.811	0.814	0.818	0.824	0.827	0.83	0.833	0.833	0.833	0.836	0.841	0.847	0.852	0.855	0.858	0.861	0.864	0.867	0.87	0.871
Pennsyivania Rhoda Island	0.815	0.804	0.808	0.812	0.816	0.821	0.824	0.826	0.828	0.83	0.833	0.837	0.84	0.842	0.845	0.845	0.845	0.847	0.851	0.856	0.86	0.863	0.866	0.869	0.871	0.875	0.878	0.879
South Carolina	0.752	0.757	0.322	0.323	0.772	0.777	0.781	0.782	0.783	0.785	0.787	0.793	0.796	0.799	0.801	0.858	0.859	0.802	0.808	0.815	0.875	0.876	0.83	0.834	0.838	0.842	0.845	0.846
South Dakota	0.769	0.772	0.777	0.783	0.788	0.794	0.799	0.801	0.804	0.805	0.808	0.811	0.813	0.814	0.816	0.814	0.812	0.814	0.819	0.826	0.833	0.838	0.842	0.847	0.851	0.856	0.859	0.86
Tennessee	0.749	0.752	0.757	0.761	0.765	0.77	0.774	0.775	0.777	0.779	0.781	0.786	0.788	0.789	0.79	0.789	0.786	0.789	0.795	0.803	0.81	0.815	0.819	0.823	0.827	0.832	0.836	0.837
Texas	0.743	0.745	0.747	0.75	0.752	0.757	0.761	0.763	0.764	0.766	0.769	0.775	0.779	0.782	0.784	0.783	0.782	0.785	0.792	0.801	0.809	0.815	0.82	0.824	0.829	0.834	0.837	0.838

Appendix Table 10: Socio-Demographic	Index val	ues for all	estimated	GBD 201	7 locations	, 1990-201	7								-													
Location	1990	1991	1992	1993	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015	2016	2017
Utah	0.761	0.765	0.769	0.773	0.776	0.781	0.786	0.787	0.79	0.792	0.795	0.8	0.803	0.806	0.808	0.808	0.807	0.811	0.817	0.825	0.832	0.837	0.841	0.845	0.848	0.852	0.855	0.856
Vermont	0.815	0.819	0.823	0.827	0.83	0.833	0.837	0.839	0.841	0.844	0.848	0.853	0.857	0.86	0.864	0.866	0.866	0.869	0.8/2	0.876	0.88	0.882	0.885	0.887	0.89	0.893	0.895	0.896
Washington	0.8	0.803	0.807	0.81	0.814	0.818	0.822	0.823	0.824	0.826	0.829	0.84	0.838	0.841	0.845	0.846	0.847	0.851	0.855	0.862	0.865	0.8/1	0.874	0.877	0.88	0.883	0.885	0.885
West Virginia	0.749	0.752	0.756	0.761	0.764	0.769	0.773	0.774	0.775	0.776	0.778	0.783	0.786	0.787	0.789	0.787	0.784	0.784	0.787	0.793	0.799	0.802	0.806	0.81	0.814	0.82	0.824	0.825
Wisconsin	0.801	0.804	0.808	0.812	0.815	0.819	0.823	0.825	0.826	0.828	0.831	0.835	0.839	0.841	0.843	0.843	0.843	0.845	0.849	0.853	0.858	0.862	0.865	0.868	0.871	0.874	0.877	0.878
Wyoming	0.766	0.771	0.777	0.782	0.786	0.792	0.796	0.797	0.799	0.801	0.804	0.809	0.813	0.816	0.818	0.819	0.819	0.823	0.831	0.838	0.846	0.851	0.855	0.858	0.862	0.866	0.869	0.869
Southern Latin America	0.594	0.6	0.607	0.613	0.619	0.626	0.632	0.638	0.643	0.648	0.652	0.655	0.658	0.662	0.667	0.673	0.677	0.679	0.682	0.685	0.69	0.695	0.7	0.704	0.707	0.713	0.717	0.72
Argentina	0.59	0.595	0.604	0.61	0.617	0.624	0.63	0.635	0.64	0.644	0.647	0.649	0.65	0.653	0.658	0.665	0.669	0.672	0.675	0.677	0.681	0.686	0.691	0.693	0.696	0.702	0.707	0.71
Chile	0.6	0.608	0.615	0.62	0.626	0.633	0.64	0.647	0.654	0.661	0.667	0.674	0.681	0.687	0.692	0.696	0.698	0.701	0.704	0.708	0.714	0.721	0.727	0.732	0.738	0.742	0.746	0.748
Uruguay	0.592	0.597	0.6	0.602	0.606	0.609	0.613	0.618	0.625	0.632	0.637	0.64	0.643	0.647	0.652	0.656	0.659	0.661	0.663	0.666	0.671	0.675	0.68	0.685	0.691	0.697	0.702	0.707
Western Europe	0.764	0.77	0.776	0.782	0.787	0.791	0.795	0.798	0.801	0.805	0.809	0.813	0.817	0.82	0.822	0.825	0.828	0.83	0.833	0.836	0.838	0.842	0.845	0.848	0.851	0.853	0.855	0.857
Andorra	0.85	0.854	0.856	0.857	0.858	0.859	0.86	0.863	0.866	0.868	0.871	0.873	0.875	0.878	0.881	0.883	0.885	0.886	0.888	0.89	0.891	0.894	0.896	0.897	0.899	0.9	0.901	0.902
Austria	0.776	0.778	0.78	0.785	0.79	0.795	0.8	0.805	0.809	0.815	0.818	0.822	0.825	0.828	0.851	0.834	0.858	0.841	0.845	0.847	0.85	0.854	0.857	0.859	0.862	0.863	0.865	0.800
Cyanus	0.724	0.73	0.74	0.75	0.758	0.765	0.323	0.778	0.335	0.789	0.795	0.803	0.81	0.817	0.824	0.83	0.837	0.842	0.847	0.851	0.854	0.857	0.859	0.861	0.862	0.863	0.864	0.865
Denmark	0.846	0.849	0.852	0.855	0.858	0.862	0.866	0.87	0.874	0.877	0.881	0.884	0.888	0.891	0.893	0.895	0.897	0.898	0.9	0.902	0.904	0.907	0.91	0.912	0.914	0.915	0.916	0.918
Finland	0.813	0.813	0.814	0.817	0.821	0.825	0.828	0.831	0.835	0.84	0.844	0.847	0.851	0.854	0.857	0.859	0.862	0.865	0.869	0.871	0.875	0.878	0.881	0.884	0.887	0.889	0.891	0.893
France	0.769	0.776	0.783	0.79	0.793	0.795	0.802	0.806	0.808	0.813	0.816	0.819	0.824	0.827	0.83	0.833	0.836	0.838	0.84	0.842	0.845	0.848	0.851	0.854	0.857	0.86	0.863	0.865
Germany	0.787	0.796	0.801	0.805	0.809	0.811	0.812	0.813	0.813	0.814	0.818	0.823	0.827	0.829	0.832	0.835	0.838	0.842	0.846	0.848	0.851	0.855	0.858	0.861	0.864	0.866	0.868	0.87
Greece	0.717	0.723	0.731	0.738	0.744	0.75	0.755	0.761	0.767	0.773	0.778	0.782	0.787	0.792	0.796	0.8	0.803	0.806	0.809	0.812	0.815	0.818	0.819	0.82	0.819	0.818	0.817	0.817
Iceland	0.814	0.818	0.821	0.825	0.828	0.83	0.833	0.835	0.839	0.843	0.848	0.854	0.859	0.862	0.865	0.869	0.872	0.876	0.88	0.883	0.886	0.889	0.892	0.895	0.899	0.902	0.905	0.907
Ireland	0.756	0.762	0.768	0.774	0.779	0.785	0.79	0.795	0.802	0.808	0.814	0.821	0.827	0.834	0.84	0.844	0.846	0.849	0.851	0.855	0.858	0.862	0.865	0.867	0.87	0.874	0.878	0.882
Israel	0.734	0.738	0.743	0.748	0.752	0.757	0.76	0.764	0.768	0.772	0.776	0.78	0.783	0.786	0.789	0.793	0.796	0.798	0.798	0.799	0.801	0.803	0.805	0.808	0.81	0.812	0.814	0.816
Italy	0.767	0.772	0.778	0.783	0.788	0.793	0.797	0.869	0.804	0.807	0.81	0.814	0.817	0.819	0.821	0.823	0.825	0.827	0.829	0.83	0.832	0.834	0.836	0.858	0.839	0.841	0.842	0.845
Malta	0.729	0.733	0.737	0.743	0.748	0.752	0.756	0.761	0.766	0.773	0.779	0.784	0.788	0.335	0.796	0.799	0.802	0.805	0.898	0.811	0.814	0.817	0.900	0.823	0.826	0.829	0.833	0.836
Netherlands	0.827	0.832	0.837	0.841	0.845	0.849	0.852	0.855	0.858	0.862	0.866	0.87	0.873	0.876	0.879	0.882	0.885	0.887	0.89	0.892	0.895	0.898	0.901	0.904	0.906	0.908	0.91	0.912
Norway	0.811	0.816	0.821	0.827	0.831	0.835	0.84	0.846	0.85	0.855	0.86	0.866	0.87	0.873	0.876	0.878	0.88	0.882	0.885	0.888	0.892	0.896	0.9	0.903	0.906	0.909	0.91	0.911
Portugal	0.642	0.65	0.659	0.667	0.675	0.682	0.688	0.694	0.699	0.705	0.71	0.716	0.722	0.727	0.732	0.736	0.741	0.744	0.748	0.751	0.755	0.76	0.764	0.768	0.771	0.773	0.775	0.778
Spain	0.715	0.723	0.731	0.738	0.745	0.752	0.758	0.763	0.768	0.773	0.778	0.782	0.786	0.79	0.794	0.797	0.799	0.802	0.805	0.809	0.812	0.815	0.818	0.819	0.82	0.822	0.823	0.825
Sweden	0.784	0.789	0.795	0.802	0.808	0.815	0.82	0.825	0.831	0.835	0.838	0.841	0.844	0.847	0.85	0.853	0.855	0.857	0.86	0.862	0.865	0.868	0.871	0.874	0.876	0.879	0.881	0.883
Stockholm	0.825	0.83	0.835	0.84	0.845	0.85	0.854	0.859	0.864	0.867	0.871	0.873	0.876	0.879	0.882	0.885	0.888	0.891	0.893	0.896	0.899	0.902	0.904	0.907	0.909	0.911	0.913	0.914
Sweden except Stockholm	0.773	0.778	0.785	0.792	0.798	0.805	0.811	0.816	0.821	0.825	0.829	0.832	0.834	0.838	0.841	0.843	0.845	0.848	0.85	0.852	0.854	0.858	0.86	0.863	0.866	0.868	0.87	0.873
Switzerland	0.841	0.842	0.844	0.846	0.847	0.848	0.758	0.85	0.851	0.852	0.854	0.857	0.859	0.86	0.862	0.865	0.865	0.808	0.8/1	0.873	0.875	0.877	0.88	0.882	0.884	0.880	0.841	0.889
England	0.725	0.729	0.745	0.743	0.758	0.763	0.756	0.702	0.776	0.782	0.78	0.794	0.791	0.801	0.804	0.807	0.81	0.813	0.816	0.819	0.823	0.827	0.832	0.838	0.842	0.845	0.847	0.849
East Midlands	0.71	0.716	0.723	0.73	0.737	0.741	0.745	0.75	0.755	0.762	0.768	0.773	0.778	0.781	0.785	0.788	0.791	0.794	0.797	0.8	0.804	0.808	0.813	0.819	0.823	0.826	0.828	0.83
Derby	0.715	0.722	0.73	0.738	0.745	0.75	0.754	0.758	0.764	0.771	0.778	0.785	0.79	0.793	0.797	0.801	0.803	0.806	0.809	0.811	0.815	0.82	0.826	0.832	0.837	0.841	0.844	0.846
Derbyshire	0.698	0.703	0.71	0.716	0.723	0.727	0.731	0.735	0.74	0.747	0.753	0.758	0.762	0.765	0.768	0.771	0.774	0.777	0.779	0.782	0.785	0.79	0.796	0.803	0.808	0.812	0.815	0.817
Leicester	0.708	0.715	0.723	0.73	0.737	0.743	0.747	0.751	0.756	0.763	0.769	0.775	0.78	0.785	0.79	0.794	0.798	0.802	0.805	0.809	0.813	0.817	0.822	0.828	0.832	0.835	0.837	0.839
Leicestershire	0.727	0.733	0.74	0.748	0.754	0.759	0.763	0.767	0.773	0.778	0.784	0.789	0.794	0.798	0.802	0.806	0.81	0.814	0.817	0.82	0.823	0.827	0.831	0.836	0.839	0.842	0.844	0.846
Lincolnshire	0.699	0.705	0.711	0.718	0.724	0.729	0.732	0.737	0.742	0.748	0.754	0.76	0.764	0.768	0.77	0.772	0.774	0.776	0.779	0.781	0.784	0.788	0.793	0.8	0.804	0.808	0.81	0.812
Northamptonshire	0.714	0.72	0.728	0.735	0.741	0.746	0.749	0.753	0.757	0.764	0.77	0.775	0.779	0.781	0.784	0.787	0.79	0.792	0.796	0.798	0.802	0.806	0.812	0.818	0.822	0.825	0.828	0.829
Nottinghamshire	0.735	0.742	0.749	0.737	0.703	0.709	0.774	0.78	0.780	0.795	0.8	0.800	0.812	0.817	0.822	0.820	0.829	0.832	0.855	0.859	0.842	0.792	0.832	0.803	0.839	0.801	0.802	0.803
Rutland	0.73	0.735	0.741	0.746	0.752	0.756	0.759	0.762	0.767	0.772	0.777	0.781	0.784	0.787	0.79	0.793	0.797	0.8	0.804	0.806	0.81	0.814	0.818	0.823	0.826	0.829	0.831	0.833
East of England	0.724	0.73	0.737	0.744	0.75	0.755	0.759	0.763	0.768	0.774	0.78	0.786	0.79	0.793	0.796	0.799	0.802	0.805	0.808	0.811	0.814	0.818	0.824	0.83	0.834	0.837	0.839	0.84
Bedford	0.73	0.736	0.743	0.749	0.756	0.76	0.764	0.768	0.774	0.78	0.785	0.79	0.794	0.797	0.8	0.803	0.805	0.807	0.81	0.812	0.816	0.82	0.825	0.83	0.833	0.836	0.837	0.838
Cambridgeshire	0.75	0.757	0.764	0.771	0.777	0.782	0.786	0.791	0.797	0.803	0.809	0.815	0.82	0.823	0.827	0.83	0.833	0.836	0.84	0.842	0.846	0.85	0.855	0.86	0.864	0.867	0.869	0.871
Central Bedfordshire	0.723	0.729	0.736	0.743	0.749	0.753	0.757	0.76	0.765	0.771	0.776	0.781	0.785	0.787	0.79	0.793	0.795	0.797	0.8	0.803	0.806	0.81	0.816	0.822	0.827	0.83	0.833	0.834
Essex	0.713	0.719	0.726	0.734	0.74	0.745	0.748	0.753	0.758	0.764	0.77	0.775	0.78	0.783	0.787	0.79	0.793	0.796	0.8	0.802	0.806	0.81	0.816	0.821	0.825	0.828	0.83	0.832
Hertfordshire	0.749	0.755	0.762	0.769	0.776	0.781	0.784	0.789	0.795	0.802	0.81	0.816	0.821	0.825	0.829	0.832	0.835	0.837	0.84	0.843	0.846	0.85	0.855	0.86	0.863	0.866	0.868	0.87
Luton	0.71	0.716	0.723	0.731	0.737	0.742	0.745	0.749	0.754	0.761	0.765	0.771	0.775	0.779	0.781	0.785	0.79	0.795	0.792	0.805	0.809	0.803	0.819	0.825	0.828	0.83	0.832	0.833
Peterborough	0.712	0.718	0.724	0.731	0.737	0.741	0.743	0.746	0.751	0.757	0.761	0.765	0.768	0.77	0.772	0.774	0.776	0.778	0.779	0.781	0.790	0.788	0.795	0.803	0.808	0.813	0.816	0.818
Southend-on-Sea	0.696	0.701	0.707	0.714	0.72	0.724	0.727	0.731	0.736	0.743	0.75	0.755	0.761	0.765	0.768	0.77	0.773	0.775	0.778	0.781	0.785	0.79	0.795	0.801	0.805	0.808	0.81	0.811
Suffolk	0.713	0.719	0.726	0.732	0.738	0.742	0.745	0.748	0.753	0.758	0.763	0.767	0.771	0.773	0.775	0.777	0.78	0.782	0.785	0.788	0.792	0.797	0.803	0.81	0.814	0.817	0.819	0.821
Thurrock	0.71	0.716	0.723	0.73	0.737	0.741	0.745	0.749	0.754	0.759	0.763	0.767	0.769	0.77	0.772	0.774	0.775	0.776	0.778	0.781	0.784	0.789	0.793	0.799	0.802	0.804	0.806	0.807
Greater London	0.78	0.786	0.793	0.8	0.806	0.81	0.814	0.818	0.823	0.83	0.836	0.842	0.846	0.849	0.852	0.855	0.858	0.861	0.863	0.866	0.87	0.875	0.88	0.885	0.889	0.891	0.893	0.894
Barking and Dagenham	0.704	0.709	0.716	0.723	0.73	0.734	0.737	0.741	0.746	0.752	0.756	0.76	0.762	0.762	0.763	0.763	0.763	0.763	0.764	0.765	0.768	0.773	0.78	0.787	0.793	0.797	0.8	0.802
Barnet	0.759	0.765	0.771	0.777	0.783	0.788	0.792	0.796	0.802	0.809	0.814	0.82	0.824	0.827	0.829	0.831	0.834	0.835	0.838	0.84	0.843	0.847	0.852	0.856	0.859	0.862	0.864	0.865
Bexley	0.721	0.727	0.734	0.74	0.746	0.75	0.753	0.757	0.761	0.767	0.772	0.777	0.781	0.784	0.787	0.789	0.791	0.793	0.796	0.798	0.801	0.804	0.809	0.815	0.819	0.822	0.824	0.826
Brent	0.753	0.758	0.764	0.7/1	0.777	0.781	0.785	0.789	0.795	0.801	0.806	0.811	0.814	0.816	0.818	0.819	0.821	0.821	0.822	0.823	0.825	0.828	0.833	0.839	0.842	0.845	0.847	0.849
Canden	0.838	0.750	0.762	0.854	0.858	0.862	0.762	0.760	0.873	0.798	0.885	0.809	0.814	0.817	0.898	0.822	0.825	0.827	0.829	0.85	0.913	0.855	0.857	0.924	0.926	0.843	0.979	0.040
Croydon	0.741	0.747	0.754	0.761	0.767	0.771	0.774	0.777	0.782	0.788	0.794	0.799	0.803	0.805	0.808	0.81	0.812	0.813	0.815	0.816	0.818	0.82	0.824	0.827	0.83	0.831	0.832	0.833
Ealing	0.764	0.77	0.776	0.783	0.789	0.794	0.797	0.801	0.807	0.814	0.82	0.825	0.828	0.831	0.833	0.834	0.835	0.836	0.837	0.837	0.839	0.843	0.848	0.853	0.857	0.86	0.863	0.865
Enfield	0.737	0.742	0.749	0.755	0.761	0.765	0.768	0.772	0.777	0.783	0.788	0.793	0.797	0.799	0.801	0.802	0.804	0.806	0.809	0.811	0.814	0.818	0.824	0.829	0.833	0.836	0.838	0.839
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Appendix Table 10: Socio-Demographic	Index val	ues for al	estimated	GBD 201	7 locations	, 1990-201	7																					
Location	1990	1991	1992	1993	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015	2016	2017
Greenwich	0.725	0.732	0.74	0.747	0.754	0.758	0.762	0.766	0.771	0.777	0.783	0.788	0.791	0.793	0.795	0.797	0.799	0.801	0.803	0.805	0.809	0.813	0.819	0.824	0.828	0.83	0.832	0.833
Hackney	0.767	0.774	0.781	0.789	0.795	0.798	0.8	0.801	0.805	0.811	0.817	0.822	0.825	0.828	0.831	0.834	0.838	0.843	0.848	0.853	0.858	0.864	0.871	0.877	0.882	0.884	0.886	0.887
Hammersmith and Fulham	0.825	0.833	0.839	0.845	0.851	0.856	0.86	0.864	0.869	0.875	0.879	0.883	0.887	0.889	0.892	0.895	0.898	0.901	0.904	0.907	0.91	0.914	0.917	0.921	0.923	0.924	0.925	0.927
Haringey	0.755	0.761	0.766	0.773	0.778	0.782	0.785	0.788	0.792	0.798	0.804	0.809	0.813	0.815	0.818	0.821	0.823	0.825	0.828	0.829	0.832	0.835	0.84	0.845	0.848	0.851	0.852	0.854
Harrow	0.747	0.754	0.76	0.767	0.773	0.777	0.78	0.784	0.79	0.796	0.802	0.808	0.812	0.815	0.819	0.821	0.824	0.825	0.827	0.829	0.83	0.832	0.835	0.839	0.842	0.844	0.846	0.848
Havering	0.719	0.725	0.731	0.737	0.743	0.747	0.75	0.754	0.759	0.765	0.77	0.775	0.78	0.783	0.786	0.789	0.792	0 794	0.797	0.8	0.802	0.805	0.809	0.814	0.817	0.82	0.823	0.824
Hillingdon	0.781	0.787	0.794	0.801	0.807	0.911	0.815	0.919	0.822	0.929	0.922	0.936	0.84	0.842	0.945	0.846	0.949	0.85	0.852	0.854	0.857	0.862	0.867	0.872	0.876	0.879	0.923	0.992
Hannalan	0.760	0.775	0.794	0.799	0.307	0.307	0.815	0.818	0.811	0.020	0.832	0.850	0.824	0.877	0.045	0.842	0.844	0.845	0.832	0.840	0.857	0.862	0.867	0.072	0.870	0.875	0.831	0.832
I-lineten	0.812	0.010	0.935	0.922	0.795	0.797	0.801	0.805	0.011	0.010	0.847	0.872	0.876	0.870	0.892	0.042	0.044	0.801	0.040	0.049	0.002	0.006	0.002	0.000	0.012	0.010	0.071	0.077
Istingion	0.812	0.015	0.823	0.852	0.858	0.845	0.047	0.851	0.850	0.862	0.807	0.872	0.870	0.879	0.002	0.885	0.0007	0.091	0.895	0.898	0.902	0.900	0.91	0.914	0.917	0.919	0.921	0.922
Kensington and Cheisea	0.839	0.845	0.851	0.857	0.861	0.865	0.869	0.875	0.8//	0.884	0.889	0.894	0.897	0.899	0.902	0.905	0.907	0.909	0.912	0.915	0.918	0.921	0.924	0.927	0.929	0.95	0.931	0.932
Kingston upon Thames	0.787	0.794	0.8	0.807	0.812	0.817	0.821	0.826	0.831	0.838	0.844	0.849	0.854	0.858	0.861	0.864	0.867	0.869	0.872	0.8/4	0.876	0.879	0.882	0.885	0.887	0.888	0.889	0.89
Lambeth	0.775	0.783	0.791	0.797	0.804	0.808	0.812	0.816	0.821	0.828	0.835	0.84	0.845	0.848	0.852	0.856	0.86	0.864	0.869	0.873	0.877	0.882	0.887	0.892	0.895	0.897	0.899	0.9
Lewisham	0.733	0.74	0.746	0.753	0.759	0.763	0.766	0.77	0.775	0.781	0.787	0.793	0.797	0.799	0.802	0.804	0.806	0.808	0.811	0.813	0.816	0.821	0.826	0.832	0.836	0.839	0.841	0.843
Merton	0.758	0.765	0.771	0.779	0.785	0.79	0.795	0.799	0.805	0.812	0.819	0.824	0.828	0.83	0.833	0.835	0.837	0.839	0.841	0.843	0.845	0.849	0.855	0.86	0.865	0.868	0.8/1	0.8/3
Newham	0.716	0.722	0.728	0.734	0.74	0.743	0.746	0.748	0.753	0.76	0.765	0.771	0.775	0.777	0.78	0.781	0.783	0.785	0.788	0.792	0.799	0.806	0.815	0.823	0.829	0.833	0.836	0.838
Redbridge	0.728	0.733	0.739	0.746	0.751	0.756	0.759	0.763	0.768	0.774	0.78	0.785	0.789	0.791	0.794	0.796	0.798	0.801	0.803	0.805	0.807	0.811	0.816	0.821	0.825	0.827	0.83	0.831
Richmond upon Thames	0.794	0.801	0.807	0.813	0.819	0.824	0.827	0.832	0.837	0.844	0.85	0.856	0.86	0.864	0.867	0.87	0.872	0.874	0.876	0.878	0.88	0.883	0.887	0.891	0.895	0.897	0.9	0.902
Southwark	0.793	0.8	0.807	0.815	0.821	0.827	0.83	0.835	0.84	0.847	0.852	0.857	0.86	0.863	0.866	0.869	0.872	0.875	0.879	0.883	0.887	0.893	0.898	0.904	0.907	0.909	0.911	0.912
Sutton	0.735	0.74	0.747	0.754	0.761	0.765	0.769	0.774	0.78	0.787	0.793	0.798	0.802	0.805	0.808	0.809	0.811	0.813	0.815	0.817	0.819	0.823	0.828	0.833	0.836	0.839	0.841	0.843
Tower Hamlets	0.766	0.773	0.78	0.788	0.795	0.801	0.805	0.81	0.817	0.825	0.832	0.839	0.844	0.848	0.852	0.857	0.861	0.866	0.871	0.876	0.881	0.886	0.891	0.896	0.9	0.902	0.904	0.905
Waltham Forest	0.712	0.718	0.724	0.731	0.738	0.742	0.745	0.749	0.754	0.76	0.765	0.77	0.773	0.775	0.777	0.778	0.78	0.781	0.783	0.784	0.787	0.792	0.798	0.805	0.81	0.814	0.817	0.819
Wandsworth	0.793	0.8	0.807	0.814	0.82	0.825	0.83	0.835	0.84	0.848	0.855	0.862	0.867	0.87	0.873	0.876	0.879	0.881	0.885	0.887	0.889	0.893	0.897	0.901	0.905	0.907	0.909	0.911
Westminster	0.831	0.837	0.843	0.849	0.854	0.858	0.862	0.865	0.87	0.875	0.881	0.885	0.889	0.891	0.895	0.898	0.901	0.903	0.905	0.907	0.91	0.913	0.916	0.92	0.922	0.924	0.925	0.927
North East England	0.697	0.703	0.711	0.718	0.725	0.73	0.733	0.738	0.744	0.75	0.757	0.763	0.767	0.771	0.774	0.778	0.781	0.784	0.787	0.79	0.794	0.798	0.804	0.81	0.814	0.817	0.819	0.821
County Durham	0.69	0.696	0.703	0.71	0.717	0.721	0.725	0.73	0.735	0.742	0.748	0.754	0.758	0.762	0.765	0.767	0.77	0.772	0.776	0.778	0.782	0.786	0.792	0.798	0.802	0.806	0.808	0.81
Darlington	0.703	0.709	0.716	0.723	0.73	0.734	0.737	0.74	0.745	0.752	0.758	0.764	0.769	0.773	0.777	0.78	0.783	0.785	0.788	0.791	0.795	0.8	0.806	0.812	0.817	0.821	0.823	0.825
Gateshead	0.7	0.706	0.713	0.72	0.727	0.732	0.735	0.739	0.745	0.751	0.758	0.764	0.769	0.773	0.777	0.781	0.784	0.787	0.791	0.794	0.798	0.803	0.808	0.814	0.818	0.821	0.824	0.826
Hartlepool	0.677	0.683	0.69	0.697	0.703	0.707	0.71	0.714	0.719	0.726	0.732	0.737	0.74	0.742	0.744	0.747	0.749	0.752	0.755	0.758	0.762	0.767	0.773	0.78	0.785	0.788	0.791	0.793
Middlesbrough	0.688	0.695	0.702	0.709	0.716	0.72	0.723	0.727	0.732	0.74	0.746	0.751	0.755	0.758	0.762	0.766	0.769	0.772	0.775	0.778	0.781	0.786	0.792	0.797	0.801	0.804	0.806	0.808
Newcastle upon Tyne	0.736	0.743	0.751	0.759	0.765	0.771	0.776	0.781	0.787	0.794	0.801	0.808	0.814	0.82	0.825	0.831	0.835	0.84	0.843	0.846	0.85	0.854	0.858	0.863	0.866	0.869	0.87	0.872
North Tyneside	0.696	0.703	0.71	0.718	0.724	0.729	0.733	0.737	0.743	0.749	0.756	0.762	0.768	0.772	0.776	0.78	0.784	0.787	0.79	0.793	0.797	0.802	0.807	0.813	0.817	0.821	0.823	0.825
Northumberland	0.697	0.703	0.709	0.716	0.722	0.727	0.73	0.735	0.74	0.746	0.752	0.757	0.761	0.764	0.767	0.77	0.773	0.776	0.779	0.781	0.784	0.787	0.793	0.798	0.802	0.805	0.807	0.808
Redcar and Cleveland	0.67	0.675	0.682	0.689	0.696	0.701	0.704	0.708	0.714	0.721	0.727	0.732	0.736	0.738	0.741	0.743	0.745	0.747	0.75	0.752	0.756	0.76	0.767	0.775	0.78	0.784	0.787	0.79
South Tyneside	0.664	0.671	0.678	0.685	0.692	0.697	0.7	0.705	0.711	0.718	0.725	0.731	0.737	0.74	0.744	0.748	0.751	0.754	0.757	0.759	0.762	0.767	0.773	0.78	0.785	0.789	0.792	0.794
Stockton-on-Tees	0.705	0.712	0.72	0.728	0.735	0.74	0.744	0.748	0.754	0.761	0.767	0.772	0.775	0.777	0.779	0.781	0.784	0.785	0.788	0.791	0.794	0.798	0.804	0.811	0.815	0.818	0.821	0.823
Sunderland	0.689	0.696	0.704	0.713	0.72	0.725	0.729	0.733	0.739	0.746	0.752	0.758	0.763	0.766	0.77	0.773	0.776	0.779	0.782	0.784	0.787	0.791	0.797	0.803	0.808	0.811	0.813	0.815
North West England	0.716	0.723	0.729	0.736	0.742	0.747	0.751	0.755	0.76	0.766	0.772	0.778	0.782	0.786	0.789	0.792	0.795	0.798	0.801	0.804	0.807	0.812	0.817	0.823	0.827	0.83	0.832	0.834
Blackburn with Darwen	0.691	0.695	0.701	0.707	0.713	0.717	0.719	0.723	0.728	0.734	0.74	0.745	0.748	0.75	0.753	0.756	0.758	0.761	0.764	0.766	0.769	0.774	0.78	0.788	0.793	0.797	0.8	0.802
Blackmool	0.68	0.685	0.691	0.698	0.703	0.707	0.71	0.713	0.718	0.724	0.729	0.733	0.737	0.738	0.741	0.743	0.745	0.747	0.748	0.75	0.752	0.756	0.762	0.768	0.773	0.776	0.779	0.781
Bolton	0.701	0.707	0.714	0.72	0.726	0.73	0.733	0.736	0.741	0.746	0.751	0.756	0.759	0.761	0.763	0.765	0.768	0.769	0.772	0.774	0.777	0.781	0.787	0 794	0.798	0.801	0.803	0.805
Burg	0.703	0.700	0.716	0.723	0.72	0.724	0.739	0.742	0.747	0.752	0.759	0.762	0.766	0.769	0.771	0.774	0.777	0.779	0.781	0.792	0.786	0.701	0.707	0.903	0.807	0.81	0.813	0.815
Chachina East	0.749	0.754	0.76	0.767	0.772	0.779	0.792	0.797	0.792	0.792	0.902	0.900	0.913	0.917	0.97	0.973	0.826	0.979	0.922	0.924	0.936	0.731	0.945	0.803	0.856	0.850	0.861	0.864
Chachira Wart and Chartar	0.74	0.747	0.754	0.761	0.769	0.772	0.776	0.78	0.795	0.792	0.707	0.803	0.807	0.811	0.815	0.919	0.822	0.825	0.828	0.921	0.834	0.939	0.842	0.847	0.050	0.852	0.854	0.855
Combrin	0.74	0.724	0.734	0.727	0.703	0.772	0.751	0.755	0.765	0.7/2	0.77	0.305	0.777	0.770	0.792	0.794	0.797	0.720	0.323	0.304	0.709	0.858	0.842	0.047	0.810	0.832	0.824	0.855
United	0.719	0.724	0.731	0.737	0.745	0.746	0.731	0.733	0.76	0.765	0.77	0.7/4	0.777	0.779	0.782	0.704	0.787	0.78	0.792	0.794	0.798	0.802	0.808	0.813	0.819	0.825	0.820	0.824
Hallon Kananalan	0.702	0.708	0.713	0.725	0.729	0.734	0.737	0.74	0.745	0.75	0.730	0.762	0.764	0.767	0.772	0.775	0.778	0.78	0.784	0.787	0.791	0.797	0.803	0.81	0.815	0.819	0.812	0.814
Knowsicy	0.091	0.090	0.702	0.708	0.747	0.753	0.722	0.720	0.751	0.737	0.745	0.749	0.754	0.737	0.76	0.704	0.707	0.77	0.7/4	0.777	0.782	0.787	0.795	0.82	0.800	0.827	0.815	0.810
Lancashire	0.72	0.720	0.734	0.741	0.747	0.752	0.750	0.76	0.769	0.77	0.773	0.780	0.705	0.780	0.789	0.792	0.794	0.797	0.799	0.802	0.803	0.809	0.813	0.82	0.847	0.827	0.829	0.851
Manahastan	0.721	0.752	0.755	0.742	0.748	0.755	0.795	0.702	0.706	0.775	0.785	0.789	0.935	0.801	0.800	0.811	0.813	0.840	0.852	0.020	0.051	0.035	0.840	0.874	0.047	0.849	0.892	0.002
Manchester	0.740	0.735	0.76	0.703	0.775	0.78	0.785	0.79	0.790	0.804	0.812	0.819	0.825	0.85	0.855	0.04	0.044	0.849	0.855	0.850	0.80	0.804	0.809	0.874	0.878	0.88	0.885	0.885
Didham	0.685	0.69	0.697	0.705	0.708	0.711	0.713	0.715	0.719	0.724	0.729	0.735	0.737	0.739	0.741	0.744	0.746	0.749	0.752	0.754	0.758	0.765	0.77	0.7/8	0.782	0.786	0.788	0.79
Rochuare	0.080	0.092	0.098	0.703	0.711	0.715	0.717	0.721	0.720	0.732	0.758	0.742	0.740	0.746	0.75	0.755	0.755	0.757	0.70	0.762	0.765	0.77	0.777	0.785	0.787	0.791	0.795	0.795
Saltord	0.708	0.714	0.721	0.728	0.735	0.74	0.743	0.747	0.753	0.761	0.768	0.775	0.781	0.785	0.79	0.794	0.797	0.8	0.803	0.806	0.809	0.814	0.819	0.826	0.83	0.834	0.836	0.838
Setton	0.706	0.713	0.72	0.727	0.733	0.738	0.741	0.746	0.751	0.757	0.763	0.768	0.773	0.776	0.78	0.783	0.786	0.788	0.79	0.792	0.794	0.796	0.8	0.803	0.806	0.809	0.81	0.812
St Helens	0.684	0.69	0.696	0.703	0.709	0.713	0.716	0.72	0.725	0.731	0.738	0.744	0.749	0.753	0.756	0.759	0.762	0.765	0.768	0.771	0.774	0.779	0.785	0.791	0.795	0.798	0.801	0.803
Stockport	0.727	0.734	0.741	0.748	0.755	0.759	0.763	0.767	0.772	0.779	0.785	0.791	0.796	0.799	0.803	0.806	0.809	0.812	0.814	0.817	0.82	0.823	0.828	0.833	0.837	0.839	0.841	0.843
Tameside	0.691	0.697	0.703	0.71	0.716	0.721	0.724	0.728	0.734	0.74	0.746	0.751	0.755	0.758	0.76	0.762	0.763	0.764	0.766	0.768	0.77	0.774	0.779	0.785	0.79	0.793	0.795	0.797
Trafford	0.751	0.757	0.764	0.771	0.778	0.782	0.786	0.79	0.795	0.802	0.809	0.815	0.82	0.824	0.829	0.832	0.836	0.839	0.842	0.844	0.848	0.852	0.856	0.862	0.865	0.868	0.871	0.873
Warrington	0.739	0.745	0.752	0.759	0.765	0.769	0.773	0.776	0.782	0.788	0.795	0.801	0.807	0.811	0.814	0.818	0.822	0.825	0.827	0.83	0.833	0.838	0.843	0.849	0.853	0.856	0.858	0.86
Wigan	0.691	0.697	0.703	0.71	0.716	0.72	0.723	0.727	0.731	0.737	0.742	0.747	0.75	0.753	0.755	0.757	0.76	0.762	0.764	0.766	0.769	0.774	0.78	0.786	0.79	0.793	0.796	0.798
Wirral	0.695	0.701	0.708	0.714	0.721	0.725	0.728	0.732	0.737	0.743	0.749	0.753	0.757	0.761	0.763	0.766	0.768	0.77	0.773	0.775	0.777	0.781	0.786	0.792	0.796	0.799	0.801	0.803
South East England	0.749	0.755	0.761	0.767	0.773	0.777	0.781	0.784	0.789	0.795	0.801	0.806	0.809	0.812	0.815	0.818	0.821	0.823	0.826	0.828	0.831	0.836	0.841	0.846	0.85	0.852	0.855	0.856
Bracknell Forest	0.759	0.764	0.77	0.776	0.781	0.785	0.788	0.791	0.797	0.804	0.811	0.817	0.822	0.825	0.829	0.832	0.835	0.838	0.841	0.843	0.845	0.849	0.854	0.859	0.862	0.865	0.867	0.869
Brighton and Hove	0.766	0.772	0.779	0.785	0.791	0.796	0.801	0.806	0.811	0.816	0.822	0.827	0.833	0.838	0.843	0.847	0.851	0.855	0.859	0.863	0.865	0.868	0.872	0.876	0.879	0.881	0.883	0.885
Buckinghamshire	0.764	0.769	0.775	0.782	0.788	0.792	0.795	0.799	0.804	0.81	0.815	0.82	0.824	0.826	0.829	0.832	0.834	0.836	0.838	0.84	0.842	0.846	0.851	0.855	0.859	0.861	0.863	0.865
East Sussex	0.712	0.718	0.724	0.73	0.736	0.74	0.742	0.745	0.749	0.754	0.759	0.763	0.766	0.769	0.771	0.773	0.776	0.778	0.781	0.783	0.787	0.791	0.797	0.803	0.807	0.81	0.812	0.814
Hampshire	0.744	0.75	0.756	0.762	0.768	0.772	0.775	0.778	0.782	0.788	0.793	0.798	0.802	0.804	0.807	0.81	0.812	0.815	0.818	0.82	0.824	0.828	0.834	0.839	0.843	0.846	0.848	0.85
Isle of Wight	0.704	0.709	0.715	0.722	0.727	0.732	0.735	0.739	0.744	0.749	0.756	0.761	0.765	0.767	0.768	0.771	0.773	0.776	0.779	0.781	0.784	0.788	0.794	0.801	0.806	0.809	0.812	0.814
Kent	0.723	0.728	0.734	0.74	0.746	0.75	0.752	0.756	0.76	0.765	0.77	0.774	0.777	0.779	0.782	0.785	0.787	0.79	0.793	0.796	0.8	0.805	0.811	0.817	0.822	0.824	0.826	0.828
Medway	0.703	0.709	0.715	0.722	0.728	0.731	0.734	0.737	0.742	0.747	0.752	0.756	0.76	0.762	0.765	0.768	0.771	0.773	0.776	0.778	0.781	0.785	0.791	0.797	0.802	0.805	0.807	0.809
•	•	•	•	•	•	-		-	•	-	-	•	•	-	•	•	•	•	•	•	-	•	-					

Appendix Table 10: Socio-Demographic	Index val	ues for al	l estimated	GBD 201	7 locations	, 1990-201	7																					
Location	1990	1991	1992	1993	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015	2016	2017
Milton Keynes	0.754	0.76	0.767	0.774	0.78	0.784	0.786	0.789	0.793	0.798	0.802	0.806	0.81	0.812	0.815	0.817	0.819	0.821	0.823	0.825	0.829	0.834	0.84	0.847	0.852	0.856	0.859	0.86
Oxfordshire	0.769	0.775	0.781	0.788	0.794	0.798	0.801	0.805	0.81	0.816	0.822	0.827	0.831	0.835	0.838	0.841	0.844	0.847	0.849	0.852	0.855	0.859	0.864	0.869	0.872	0.875	0.878	0.879
Portsmouth	0.75	0.756	0.763	0.77	0.776	0.781	0.785	0.79	0.795	0.8	0.805	0.81	0.815	0.818	0.822	0.824	0.827	0.829	0.832	0.835	0.838	0.842	0.846	0.851	0.854	0.857	0.858	0.86
Reading	0.785	0.791	0.797	0.803	0.809	0.813	0.817	0.821	0.827	0.834	0.84	0.847	0.852	0.856	0.86	0.864	0.866	0.868	0.87	0.872	0.874	0.877	0.882	0.887	0.89	0.892	0.894	0.895
Slough	0.764	0.77	0.777	0.784	0.79	0.793	0.796	0.799	0.802	0.806	0.81	0.813	0.815	0.816	0.817	0.818	0.819	0.82	0.822	0.825	0.829	0.834	0.841	0.848	0.852	0.855	0.858	0.859
Southampton	0.752	0.758	0.765	0.772	0.779	0.784	0.789	0.794	0.8	0.805	0.81	0.815	0.819	0.823	0.826	0.829	0.831	0.834	0.836	0.837	0.839	0.842	0.845	0.849	0.852	0.855	0.856	0.858
Surrey	0.773	0.779	0.785	0.791	0.797	0.801	0.805	0.809	0.814	0.821	0.827	0.833	0.838	0.841	0.845	0.847	0.85	0.853	0.856	0.858	0.861	0.864	0.868	0.873	0.876	0.879	0.881	0.883
West Berkshire	0.774	0.78	0.786	0.793	0.799	0.803	0.805	0.808	0.813	0.819	0.824	0.829	0.832	0.835	0.836	0.836	0.837	0.838	0.84	0.842	0.846	0.851	0.857	0.863	0.867	0.869	0.871	0.872
West Sussex	0.74	0.745	0.751	0.757	0.763	0.767	0.77	0.773	0.777	0.783	0.788	0.793	0.796	0.799	0.802	0.804	0.807	0.809	0.812	0.814	0.818	0.822	0.827	0.833	0.837	0.84	0.842	0.843
Windsor and Maidenhead	0.778	0.783	0.789	0.795	0.8	0.805	0.808	0.811	0.816	0.823	0.829	0.835	0.839	0.843	0.847	0.851	0.854	0.857	0.86	0.863	0.866	0.87	0.874	0.88	0.883	0.885	0.887	0.889
Wokingham	0.778	0.784	0.79	0.797	0.802	0.806	0.81	0.814	0.82	0.826	0.832	0.837	0.842	0.845	0.849	0.853	0.856	0.858	0.861	0.863	0.865	0.868	0.8/1	0.876	0.879	0.882	0.883	0.885
South west England	0.729	0.759	0.741	0.748	0.754	0.758	0.762	0.700	0.7/1	0.777	0.785	0.788	0.792	0.796	0.799	0.802	0.805	0.807	0.81	0.813	0.816	0.82	0.825	0.851	0.855	0.858	0.84	0.841
Baun and North East Somerset	0.732	0.738	0.75	0.77	0.763	0.762	0.780	0.79	0.790	0.803	0.809	0.803	0.822	0.812	0.835	0.838	0.824	0.878	0.83	0.833	0.836	0.839	0.803	0.807	0.851	0.872	0.856	0.873
Bristol City of	0.763	0.745	0.777	0.784	0.703	0.705	0.775	0.805	0.81	0.817	0.823	0.828	0.833	0.836	0.839	0.843	0.846	0.849	0.853	0.856	0.859	0.863	0.868	0.873	0.877	0.855	0.882	0.853
Cornwall	0.7	0.706	0.713	0.721	0.727	0.731	0.734	0.738	0.743	0.749	0.755	0.76	0.764	0.768	0.771	0.774	0.777	0.78	0.783	0.786	0.789	0.793	0.799	0.806	0.81	0.813	0.815	0.817
Devon	0.72	0.726	0.733	0.74	0.746	0.75	0.753	0.757	0.762	0.769	0.775	0.78	0.785	0.789	0.793	0.796	0.8	0.803	0.806	0.808	0.811	0.816	0.821	0.826	0.83	0.833	0.835	0.837
Dorset	0.716	0.721	0.727	0.734	0.74	0.744	0.747	0.751	0.756	0.762	0.769	0.773	0.777	0.779	0.781	0.783	0.786	0.788	0.791	0.793	0.797	0.802	0.808	0.814	0.818	0.821	0.823	0.825
Gloucestershire	0.735	0.741	0.747	0.754	0.76	0.765	0.768	0.772	0.777	0.783	0.79	0.795	0.8	0.804	0.808	0.811	0.813	0.816	0.818	0.82	0.824	0.828	0.833	0.839	0.843	0.846	0.848	0.85
North Somerset	0.714	0.72	0.727	0.733	0.739	0.743	0.746	0.75	0.755	0.76	0.766	0.771	0.776	0.78	0.783	0.786	0.789	0.792	0.795	0.798	0.801	0.806	0.813	0.819	0.824	0.827	0.83	0.832
Plymouth	0.724	0.73	0.737	0.744	0.75	0.754	0.758	0.762	0.767	0.772	0.778	0.783	0.787	0.79	0.793	0.796	0.799	0.802	0.805	0.807	0.81	0.814	0.819	0.825	0.829	0.832	0.834	0.836
Poole	0.727	0.733	0.74	0.746	0.753	0.758	0.761	0.765	0.771	0.777	0.783	0.789	0.793	0.796	0.798	0.801	0.804	0.806	0.809	0.811	0.814	0.818	0.824	0.83	0.835	0.838	0.84	0.842
Somerset	0.713	0.718	0.724	0.731	0.737	0.741	0.744	0.748	0.752	0.757	0.763	0.767	0.77	0.772	0.775	0.777	0.78	0.782	0.785	0.787	0.789	0.794	0.799	0.805	0.809	0.812	0.814	0.816
South Gloucestershire	0.747	0.752	0.758	0.765	0.771	0.775	0.779	0.783	0.789	0.796	0.802	0.808	0.813	0.817	0.821	0.824	0.827	0.831	0.834	0.837	0.84	0.844	0.849	0.855	0.859	0.862	0.865	0.867
Swindon	0.747	0.753	0.76	0.767	0.773	0.776	0.778	0.781	0.786	0.792	0.797	0.801	0.805	0.806	0.807	0.809	0.811	0.813	0.815	0.818	0.82	0.825	0.831	0.837	0.841	0.844	0.846	0.847
Torbay	0.699	0.705	0.711	0.717	0.723	0.727	0.73	0.733	0.736	0.741	0.745	0.749	0.75	0.751	0.753	0.755	0.757	0.759	0.761	0.762	0.764	0.767	0.773	0.779	0.783	0.786	0.789	0.79
Wiltshire	0.726	0.731	0.737	0.743	0.749	0.753	0.756	0.759	0.763	0.769	0.775	0.78	0.784	0.786	0.789	0.791	0.793	0.795	0.797	0.799	0.803	0.808	0.813	0.819	0.823	0.826	0.828	0.829
West Midlands	0.707	0.713	0.72	0.727	0.734	0.739	0.743	0.747	0.752	0.759	0.765	0.771	0.776	0.779	0.783	0.786	0.789	0.792	0.795	0.798	0.802	0.806	0.812	0.818	0.822	0.825	0.828	0.829
Birmingham	0.707	0.714	0.721	0.728	0.735	0.739	0.743	0.747	0.753	0.76	0.766	0.773	0.778	0.782	0.787	0.791	0.794	0.798	0.803	0.807	0.811	0.817	0.823	0.829	0.834	0.837	0.839	0.84
Coventry	0.722	0.729	0.737	0.745	0.751	0.757	0.762	0.766	0.772	0.779	0.785	0.791	0.796	0.8	0.803	0.807	0.81	0.812	0.815	0.818	0.821	0.825	0.831	0.837	0.841	0.844	0.847	0.848
Dudley	0.692	0.698	0.704	0.71	0.716	0.721	0.724	0.728	0.733	0.739	0.746	0.751	0.756	0.759	0.762	0.765	0.767	0.77	0.772	0.774	0.776	0.779	0.784	0.789	0.793	0.795	0.797	0.799
Herefordshire, County of	0.706	0.712	0.719	0.726	0.733	0.738	0.742	0.747	0.752	0.759	0.766	0.771	0.776	0.779	0.783	0.786	0.789	0.792	0.795	0.798	0.801	0.806	0.812	0.818	0.822	0.825	0.827	0.828
Sandwell	0.682	0.688	0.695	0.702	0.708	0.712	0.715	0.719	0.724	0.729	0.735	0.74	0.744	0.747	0.75	0.752	0.754	0.756	0.759	0.761	0.765	0.769	0.776	0.783	0.788	0.792	0.795	0.797
Shropshire	0.708	0.714	0.721	0.728	0.734	0.739	0.743	0.747	0.752	0.759	0.766	0.771	0.776	0.781	0.785	0.789	0.793	0.795	0.799	0.802	0.806	0.811	0.816	0.821	0.825	0.828	0.83	0.832
Solihull	0.733	0.74	0.747	0.755	0.762	0.767	0.771	0.776	0.783	0.789	0.795	0.801	0.807	0.811	0.815	0.818	0.822	0.825	0.828	0.83	0.832	0.835	0.84	0.845	0.848	0.851	0.853	0.855
Staffordshire	0.708	0.714	0.722	0.729	0.736	0.74	0.744	0.749	0.755	0.761	0.767	0.772	0.777	0.78	0.783	0.787	0.79	0.793	0.796	0.799	0.802	0.806	0.811	0.816	0.819	0.822	0.824	0.826
Stoke-on-Trent	0.689	0.696	0.703	0.71	0.716	0.721	0.724	0.728	0.733	0.738	0.743	0.747	0.75	0.751	0.753	0.754	0.756	0.757	0.76	0.763	0.767	0.772	0.78	0.788	0.794	0.798	0.801	0.804
Telford and Wrekin	0.713	0.719	0.726	0.733	0.74	0.745	0.749	0.754	0.759	0.764	0.769	0.773	0.776	0.777	0.779	0.781	0.783	0.784	0.787	0.789	0.791	0.796	0.802	0.809	0.814	0.817	0.82	0.822
Waisaii	0.681	0.080	0.692	0.752	0.76	0.707	0.709	0.712	0.717	0.725	0.729	0.734	0.738	0.74	0.743	0.746	0.749	0.752	0.755	0.757	0.761	0.700	0.772	0.779	0.783	0.787	0.789	0.791
Walvathampton	0.691	0.697	0.740	0.712	0.70	0.703	0.707	0.721	0.727	0.744	0.751	0.757	0.761	0.764	0.769	0.771	0.774	0.776	0.770	0.781	0.784	0.334	0.794	0.045	0.804	0.807	0.800	0.811
Worcestershire	0.707	0.713	0.72	0.727	0.713	0.725	0.742	0.747	0.754	0.744	0.771	0.757	0.782	0.786	0.791	0.794	0.798	0.8	0.803	0.805	0.808	0.811	0.816	0.822	0.826	0.879	0.831	0.833
Yorkshire and the Humber	0.711	0.717	0.724	0.731	0.737	0.742	0.745	0.75	0.755	0.762	0.768	0.773	0.778	0.781	0.785	0.788	0.791	0.794	0.797	0.8	0.803	0.808	0.813	0.819	0.823	0.826	0.828	0.83
Barnsley	0.675	0.68	0.686	0.692	0.698	0.702	0.705	0.709	0.715	0.721	0.726	0.731	0.735	0.738	0.741	0.743	0.746	0.748	0.75	0.752	0.755	0.759	0.765	0.772	0.778	0.781	0.785	0.787
Bradford	0.693	0.699	0.705	0.712	0.718	0.721	0.723	0.725	0.729	0.736	0.741	0.747	0.751	0.753	0.756	0.759	0.762	0.765	0.768	0.771	0.775	0.78	0.787	0.794	0,799	0.803	0.805	0.807
Calderdale	0.708	0.714	0.721	0.729	0.735	0.739	0.743	0.746	0.751	0.758	0.763	0.768	0.771	0.774	0.777	0.78	0.783	0.785	0.788	0.791	0.795	0.8	0.807	0.814	0.819	0.823	0.825	0.827
Doncaster	0.674	0.679	0.685	0.692	0.697	0.701	0.704	0.707	0.712	0.718	0.724	0.729	0.733	0.736	0.74	0.743	0.746	0.749	0.752	0.755	0.759	0.764	0.771	0.778	0.783	0.786	0.789	0.791
East Riding of Yorkshire	0.714	0.72	0.727	0.734	0.741	0.745	0.749	0.753	0.758	0.764	0.769	0.774	0.778	0.781	0.784	0.787	0.79	0.792	0.794	0.797	0.799	0.802	0.807	0.812	0.815	0.818	0.82	0.822
Kingston upon Hull, City of	0.689	0.695	0.702	0.71	0.717	0.722	0.726	0.73	0.736	0.742	0.749	0.755	0.76	0.763	0.767	0.77	0.773	0.776	0.779	0.782	0.785	0.789	0.795	0.801	0.806	0.809	0.811	0.813
Kirklees	0.703	0.709	0.716	0.723	0.73	0.734	0.737	0.741	0.746	0.752	0.758	0.763	0.767	0.77	0.773	0.776	0.779	0.781	0.783	0.785	0.788	0.793	0.798	0.805	0.809	0.812	0.814	0.816
Leeds	0.739	0.746	0.753	0.761	0.768	0.773	0.778	0.783	0.789	0.797	0.803	0.81	0.816	0.82	0.825	0.83	0.834	0.837	0.841	0.843	0.846	0.849	0.854	0.858	0.862	0.864	0.867	0.868
North East Lincolnshire	0.698	0.705	0.712	0.72	0.727	0.731	0.733	0.737	0.741	0.745	0.749	0.753	0.755	0.757	0.758	0.761	0.763	0.767	0.77	0.775	0.779	0.784	0.79	0.796	0.799	0.802	0.803	0.804
North Lincolnshire	0.702	0.707	0.714	0.72	0.727	0.731	0.734	0.737	0.742	0.748	0.752	0.757	0.76	0.762	0.764	0.766	0.768	0.77	0.773	0.776	0.78	0.785	0.792	0.799	0.803	0.807	0.809	0.811
North Yorkshire	0.724	0.73	0.737	0.743	0.75	0.754	0.758	0.762	0.767	0.774	0.78	0.785	0.79	0.794	0.798	0.801	0.803	0.806	0.808	0.81	0.813	0.817	0.822	0.827	0.831	0.834	0.836	0.839
Rotherham	0.681	0.687	0.693	0.7	0.707	0.711	0.714	0.717	0.722	0.728	0.734	0.739	0.744	0.747	0.75	0.753	0.756	0.759	0.761	0.763	0.766	0.77	0.776	0.782	0.787	0.791	0.794	0.796
Sheffield	0.727	0.733	0.74	0.747	0.753	0.758	0.762	0.767	0.773	0.78	0.786	0.792	0.798	0.803	0.807	0.811	0.815	0.819	0.823	0.826	0.83	0.834	0.839	0.843	0.847	0.849	0.852	0.853
Wakefield	0.692	0.698	0.705	0.712	0.718	0.722	0.726	0.729	0.734	0.741	0.747	0.752	0.755	0.758	0.76	0.763	0.765	0.766	0.768	0.769	0.772	0.777	0.784	0.791	0.797	0.801	0.804	0.806
York	0.762	0.769	0.776	0.783	0.789	0.795	0.799	0.804	0.81	0.816	0.822	0.829	0.834	0.839	0.844	0.848	0.851	0.854	0.857	0.859	0.862	0.865	0.868	0.872	0.874	0.876	0.878	0.879
Northern Ireland	0.71	0.718	0.726	0.735	0.742	0.747	0.751	0.755	0.761	0.767	0.774	0.78	0.786	0.79	0.795	0.798	0.801	0.805	0.808	0.812	0.815	0.818	0.822	0.825	0.828	0.831	0.833	0.835
Scotland	0.672	0.678	0.686	0.693	0.701	0.706	0.711	0.716	0.722	0.73	0.737	0.743	0.748	0.752	0.757	0.761	0.764	0.768	0.772	0.777	0.781	0.785	0.789	0.793	0.797	0.8	0.803	0.805
Wates	0.642	0.65	0.659	0.668	0.676	0.683	0.688	0.694	0.702	0.711	0.719	0.727	0.734	0.739	0.745	0.75	0.755	0.76	0.765	0.77	0.774	0.78	0.786	0.792	0.797	0.622	0.803	0.806
Andean Latin America	0.497	0.502	0.508	0.495	0.521	0.527	0.534	0.519	0.546	0.552	0.557	0.563	0.567	0.571	0.576	0.581	0.587	0.592	0.598	0.603	0.608	0.505	0.601	0.624	0.628	0.633	0.637	0.64
Bolivia	0.403	0.408	0.49	0.495	0.431	0.438	0.447	0.518	0.320	0.355	0.559	0.340	0.552	0.538	0.505	0.507	0.572	0.536	0.539	0.585	0.59	0.595	0.555	0.559	0.565	0.573	0.581	0.528
Ecuador	0.405	0.408	0.515	0.52	0.431	0.436	0.522	0.430	0.400	0.532	0.483	0.494	0.504	0.515	0.52	0.520	0.552	0.585	0.559	0.545	0.547	0.603	0.555	0.618	0.623	0.575	0.632	0.587
Peru	0.493	0.495	0.497	0.501	0.509	0.517	0.526	0.534	0.542	0.549	0.555	0.540	0.564	0.568	0.571	0.575	0.578	0.584	0.59	0.594	0.6	0.604	0.608	0.614	0.62	0.626	0.632	0.636
Caribbean	0.531	0.536	0.539	0.543	0.546	0.549	0.552	0.555	0.559	0.563	0.568	0.573	0.578	0.583	0.589	0.596	0.602	0,607	0.61	0.613	0.617	0.621	0.624	0.626	0.628	0.631	0.635	0.638
Antigua and Barbuda	0.61	0.616	0.622	0.627	0.631	0.633	0.636	0.64	0.644	0.649	0.653	0.657	0.661	0.666	0.67	0.675	0.68	0.685	0.69	0.694	0.696	0.698	0.7	0.703	0.706	0.709	0.712	0.715
The Bahamas	0.674	0.669	0.665	0.667	0.671	0.676	0.681	0.687	0.695	0.703	0.71	0.715	0.718	0.721	0.723	0.727	0.732	0.737	0.742	0.746	0.751	0.753	0.755	0.755	0.755	0.755	0.755	0.756

Appendix Table 10: Socio-Demographic	c Index val	ues for all	estimated	GBD 201	7 locations	, 1990-201	7				_	_						_		_	_	_			_			
Location	1990	1991	1992	1993	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015	2016	2017
Barbados	0.65	0.655	0.661	0.667	0.672	0.675	0.678	0.68	0.682	0.683	0.686	0.689	0.693	0.698	0.703	0.707	0.71	0.713	0.717	0.72	0.723	0.726	0.729	0.732	0.734	0.736	0.738	0.739
Belize	0.402	0.414	0.429	0.445	0.458	0.471	0.482	0.49	0.496	0.501	0.506	0.511	0.518	0.525	0.533	0.541	0.549	0.557	0.563	0.569	0.576	0.581	0.586	0.59	0.594	0.598	0.6	0.602
Bermuda	0.73	0.732	0.734	0.737	0.738	0.74	0.741	0.743	0.746	0.749	0.752	0.754	0.757	0.761	0.765	0.769	0.773	0.777	0.782	0.786	0.79	0.794	0.797	0.8	0.803	0.804	0.805	0.805
Dominica	0.534	0.535	0.539	0.605	0.555	0.563	0.001	0.577	0.584	0.592	0.599	0.606	0.614	0.623	0.629	0.636	0.641	0.645	0.65	0.656	0.652	0.658	0.673	0.608	0.68	0.670	0.684	0.687
Dominican Republic	0.442	0.443	0.445	0.449	0.454	0.459	0.464	0.469	0.475	0.48	0.484	0.49	0.495	0.499	0.511	0.525	0.536	0.547	0.555	0.562	0.568	0.572	0.575	0.576	0.578	0.583	0.589	0.593
Grenada	0.46	0.472	0.485	0.497	0.508	0.519	0.529	0.54	0.55	0.558	0.566	0.573	0.578	0.584	0.589	0.594	0.599	0.604	0.608	0.612	0.615	0.618	0.62	0.624	0.628	0.632	0.637	0.64
Guyana	0.442	0.446	0.45	0.454	0.458	0.465	0.472	0.481	0.489	0.497	0.503	0.51	0.516	0.521	0.526	0.53	0.533	0.537	0.542	0.547	0.552	0.557	0.563	0.568	0.573	0.576	0.58	0.584
Haiti	0.328	0.332	0.336	0.339	0.342	0.345	0.35	0.355	0.361	0.367	0.373	0.379	0.384	0.389	0.393	0.397	0.401	0.405	0.409	0.413	0.416	0.42	0.423	0.427	0.431	0.435	0.439	0.442
Jamaica	0.547	0.553	0.56	0.565	0.571	0.578	0.585	0.591	0.596	0.6	0.607	0.615	0.621	0.626	0.63	0.635	0.639	0.644	0.649	0.653	0.658	0.663	0.667	0.671	0.673	0.676	0.677	0.679
Puerto Rico	0.684	0.689	0.693	0.698	0.703	0.708	0.714	0.719	0.724	0.729	0.736	0.745	0.752	0.756	0.759	0.762	0.766	0.77	0.774	0.778	0.782	0.786	0.791	0.797	0.805	0.811	0.813	0.813
Saint Lucia	0.509	0.516	0.524	0.532	0.54	0.549	0.558	0.566	0.573	0.58	0.585	0.589	0.593	0.598	0.603	0.606	0.609	0.613	0.618	0.623	0.628	0.633	0.636	0.64	0.643	0.646	0.65	0.653
Saint Vincent and the Grenadines	0.465	0.472	0.479	0.486	0.492	0.499	0.506	0.512	0.518	0.524	0.53	0.537	0.543	0.549	0.553	0.557	0.561	0.566	0.571	0.575	0.579	0.584	0.588	0.593	0.597	0.601	0.605	0.608
Suriname	0.529	0.534	0.538	0.541	0.543	0.543	0.544	0.547	0.551	0.555	0.56	0.568	0.576	0.585	0.592	0.597	0.601	0.604	0.608	0.612	0.616	0.62	0.625	0.628	0.632	0.636	0.639	0.641
Trinidad and Tobago	0.601	0.604	0.608	0.611	0.615	0.619	0.624	0.629	0.634	0.64	0.646	0.652	0.657	0.663	0.669	0.675	0.68	0.685	0.689	0.691	0.693	0.694	0.695	0.696	0.697	0.698	0.698	0.698
Virgin Islands	0.673	0.683	0.691	0.699	0.706	0.713	0.719	0.725	0.73	0.734	0.737	0.74	0.748	0.755	0.762	0.768	0.774	0.78	0.786	0.79	0.795	0.799	0.802	0.804	0.805	0.806	0.806	0.807
Celembia	0.492	0.496	0.501	0.507	0.515	0.521	0.528	0.534	0.54	0.545	0.551	0.557	0.561	0.564	0.567	0.571	0.577	0.583	0.588	0.593	0.597	0.601	0.606	0.61	0.614	0.617	0.621	0.623
Costa Pica	0.524	0.465	0.485	0.49	0.490	0.505	0.511	0.518	0.566	0.528	0.555	0.550	0.594	0.598	0.548	0.555	0.558	0.303	0.373	0.581	0.588	0.590	0.64	0.645	0.619	0.654	0.658	0.654
FI Salvador	0.406	0.409	0.413	0.418	0.474	0.431	0.44	0.449	0.459	0.47	0.481	0.491	0.5	0.598	0.515	0.523	0.531	0.539	0.545	0.55	0.556	0.561	0.567	0.573	0.578	0.584	0.589	0.593
Guatemala	0.313	0.318	0.326	0.338	0.353	0.363	0.371	0.381	0.397	0.411	0.42	0.427	0.433	0.44	0.449	0.458	0.465	0.47	0.476	0.482	0.489	0.496	0.501	0.507	0.513	0.517	0.521	0.524
Honduras	0.34	0.344	0.347	0.352	0.358	0.365	0.372	0.381	0.39	0.4	0.41	0.419	0.428	0.437	0.445	0.452	0.459	0.465	0.47	0.475	0.481	0.486	0.49	0.495	0.499	0.504	0.508	0.512
Mexico	0.513	0.517	0.521	0.529	0.537	0.543	0.549	0.555	0.561	0.566	0.572	0.577	0.581	0.586	0.59	0.595	0.598	0.602	0.606	0.608	0.611	0.613	0.616	0.619	0.622	0.624	0.626	0.628
Aguascalientes	0.558	0.561	0.565	0.571	0.577	0.583	0.591	0.597	0.603	0.606	0.609	0.612	0.615	0.621	0.626	0.632	0.636	0.64	0.643	0.644	0.645	0.647	0.649	0.651	0.653	0.655	0.657	0.659
Baja California	0.57	0.572	0.575	0.579	0.582	0.585	0.589	0.594	0.599	0.604	0.608	0.611	0.615	0.618	0.622	0.626	0.63	0.634	0.639	0.642	0.644	0.646	0.649	0.651	0.653	0.654	0.656	0.657
Baja California Sur	0.569	0.573	0.577	0.582	0.586	0.589	0.593	0.598	0.602	0.605	0.609	0.612	0.614	0.617	0.621	0.624	0.628	0.633	0.637	0.641	0.644	0.647	0.649	0.652	0.654	0.656	0.657	0.659
Campeche	0.485	0.49	0.496	0.504	0.511	0.517	0.525	0.534	0.543	0.551	0.558	0.564	0.569	0.573	0.577	0.581	0.585	0.589	0.593	0.595	0.597	0.599	0.602	0.605	0.608	0.611	0.614	0.616
Chiapas	0.403	0.407	0.413	0.422	0.43	0.436	0.444	0.452	0.459	0.465	0.47	0.475	0.48	0.484	0.489	0.494	0.498	0.503	0.508	0.512	0.514	0.517	0.52	0.522	0.525	0.528	0.531	0.533
Chihuahua	0.546	0.549	0.552	0.555	0.558	0.559	0.562	0.566	0.57	0.574	0.577	0.581	0.585	0.589	0.594	0.599	0.604	0.61	0.615	0.619	0.622	0.625	0.628	0.63	0.633	0.635	0.637	0.639
Coahuila	0.549	0.555	0.561	0.567	0.573	0.576	0.581	0.587	0.592	0.597	0.601	0.604	0.607	0.611	0.616	0.62	0.624	0.628	0.631	0.633	0.634	0.635	0.637	0.638	0.64	0.642	0.644	0.645
Colima	0.538	0.543	0.549	0.556	0.561	0.565	0.571	0.577	0.583	0.588	0.594	0.598	0.603	0.607	0.611	0.616	0.621	0.626	0.631	0.634	0.637	0.64	0.643	0.645	0.648	0.65	0.652	0.654
Durango	0.602	0.606	0.61	0.617	0.629	0.64	0.65	0.656	0.542	0.666	0.671	0.676	0.681	0.685	0.689	0.693	0.696	0.698	0.7	0.701	0.702	0.704	0.706	0.708	0.71	0.712	0.714	0.716
Guanajuato	0.494	0.499	0.504	0.511	0.519	0.525	0.528	0.555	0.546	0.551	0.558	0.565	0.571	0.575	0.579	0.583	0.587	0.591	0.595	0.598	0.601	0.604	0.607	0.611	0.614	0.616	0.619	0.624
Guerrero	0.426	0.429	0.433	0.44	0.445	0.45	0.456	0.465	0.473	0.48	0.487	0.492	0.498	0.505	0.511	0.517	0.523	0.529	0.535	0.539	0.542	0.545	0.548	0.552	0.555	0.558	0.56	0.562
Hidalgo	0.417	0.424	0.433	0.444	0.454	0.462	0.471	0.481	0.491	0.501	0.511	0.52	0.527	0.531	0.535	0.538	0.543	0.548	0.554	0.558	0.562	0.567	0.572	0.576	0.58	0.583	0.585	0.587
Jalisco	0.532	0.538	0.545	0.552	0.558	0.563	0.569	0.576	0.582	0.588	0.593	0.598	0.602	0.606	0.611	0.615	0.619	0.624	0.628	0.63	0.632	0.635	0.637	0.64	0.642	0.645	0.647	0.649
México	0.529	0.524	0.525	0.539	0.558	0.568	0.574	0.577	0.58	0.585	0.59	0.596	0.6	0.603	0.607	0.611	0.613	0.614	0.614	0.614	0.615	0.617	0.621	0.625	0.628	0.631	0.633	0.635
Michoacán de Ocampo	0.464	0.47	0.476	0.483	0.489	0.494	0.501	0.508	0.515	0.522	0.527	0.532	0.537	0.542	0.546	0.551	0.555	0.559	0.563	0.566	0.568	0.571	0.573	0.576	0.579	0.581	0.584	0.586
Morelos	0.525	0.532	0.539	0.547	0.553	0.558	0.564	0.571	0.577	0.582	0.587	0.591	0.595	0.598	0.601	0.605	0.608	0.612	0.615	0.617	0.619	0.622	0.624	0.627	0.629	0.631	0.634	0.635
Nayarit	0.495	0.5	0.506	0.512	0.518	0.523	0.529	0.537	0.545	0.553	0.56	0.566	0.571	0.576	0.581	0.585	0.59	0.594	0.598	0.6	0.602	0.604	0.607	0.61	0.613	0.615	0.618	0.62
Nuevo León	0.598	0.602	0.606	0.611	0.615	0.617	0.62	0.625	0.63	0.635	0.64	0.644	0.647	0.65	0.653	0.656	0.659	0.662	0.665	0.666	0.667	0.669	0.67	0.672	0.673	0.675	0.676	0.677
Oaxaca	0.431	0.435	0.441	0.447	0.452	0.455	0.459	0.466	0.473	0.479	0.485	0.49	0.496	0.501	0.507	0.512	0.517	0.523	0.529	0.533	0.537	0.541	0.545	0.549	0.552	0.555	0.558	0.561
Puebla	0.456	0.463	0.47	0.478	0.487	0.493	0.496	0.497	0.498	0.499	0.504	0.509	0.516	0.524	0.532	0.538	0.544	0.549	0.554	0.558	0.563	0.567	0.571	0.575	0.577	0.58	0.582	0.584
Queretaro Quintana Roo	0.497	0.502	0.508	0.518	0.53	0.543	0.556	0.500	0.575	0.583	0.591	0.597	0.602	0.606	0.609	0.612	0.605	0.619	0.623	0.624	0.626	0.629	0.631	0.633	0.635	0.636	0.638	0.639
San Luis Potosí	0.32	0.486	0.492	0.541	0.548	0.555	0.502	0.53	0.538	0.545	0.551	0.556	0.561	0.556	0.571	0.575	0.58	0.586	0.591	0.595	0.599	0.602	0.606	0.61	0.613	0.616	0.619	0.621
Sinaloa	0.523	0.528	0.533	0.539	0.544	0.549	0.555	0.562	0.57	0.577	0.583	0.589	0.594	0.599	0.604	0.609	0.614	0.619	0.623	0.627	0.63	0.633	0.636	0.639	0.642	0.644	0.646	0.649
Sonora	0.553	0.557	0.562	0.566	0.57	0.573	0.578	0.583	0.588	0.593	0.597	0.601	0.605	0.608	0.612	0.616	0.621	0.625	0.629	0.632	0.635	0.637	0.64	0.643	0.645	0.647	0.649	0.65
Tabasco	0.474	0.479	0.486	0.493	0.5	0.507	0.515	0.524	0.533	0.541	0.548	0.553	0.558	0.563	0.568	0.573	0.578	0.583	0.588	0.591	0.594	0.596	0.599	0.602	0.604	0.607	0.609	0.611
Tamaulipas	0.548	0.553	0.558	0.564	0.568	0.571	0.574	0.579	0.586	0.592	0.598	0.602	0.606	0.609	0.613	0.616	0.62	0.624	0.628	0.63	0.633	0.635	0.637	0.64	0.642	0.643	0.645	0.647
Tlaxcala	0.478	0.482	0.487	0.495	0.506	0.515	0.524	0.531	0.536	0.541	0.545	0.55	0.556	0.561	0.567	0.573	0.578	0.583	0.587	0.59	0.591	0.594	0.596	0.598	0.6	0.601	0.603	0.604
Veracruz de Ignacio de la Llave	0.461	0.463	0.467	0.472	0.477	0.48	0.485	0.492	0.5	0.509	0.517	0.523	0.529	0.534	0.54	0.546	0.551	0.557	0.563	0.567	0.571	0.574	0.578	0.581	0.584	0.587	0.59	0.592
Yucatán	0.497	0.502	0.508	0.516	0.524	0.531	0.54	0.548	0.556	0.562	0.569	0.575	0.581	0.587	0.592	0.596	0.599	0.602	0.605	0.608	0.61	0.613	0.617	0.62	0.623	0.626	0.628	0.63
Zacatecas	0.483	0.489	0.495	0.502	0.509	0.514	0.52	0.527	0.534	0.54	0.546	0.551	0.555	0.56	0.564	0.568	0.572	0.577	0.581	0.584	0.586	0.589	0.592	0.595	0.598	0.602	0.605	0.608
Nicaragua	0.357	0.363	0.368	0.374	0.381	0.389	0.397	0.406	0.415	0.424	0.432	0.439	0.446	0.453	0.46	0.466	0.47	0.475	0.481	0.486	0.492	0.499	0.504	0.509	0.514	0.52	0.525	0.53
Panama	0.542	0.546	0.55	0.555	0.56	0.565	0.569	0.573	0.578	0.583	0.589	0.595	0.6	0.604	0.608	0.611	0.614	0.618	0.622	0.626	0.63	0.635	0.641	0.648	0.656	0.664	0.671	0.677
Venezuela	0.528	0.536	0.553	0.559	0.566	0.578	0.587	0.592	0.591	0.587	0.591	0.602	0.602	0.588	0.576	0.579	0.594	0.61	0.621	0.625	0.629	0.633	0.64	0.646	0.648	0.651	0.654	0.655
rropicai Latin America Brazil	0.494	0.5	0.507	0.514	0.521	0.529	0.537	0.545	0.551	0.556	0.562	0.567	0.571	0.577	0.582	0.588	0.594	0.602	0.608	0.615	0.622	0.628	0.635	0.642	0.649	0.655	0.659	0.662
Acre	0.494	0.301	0.308	0.315	0.322	0.33	0.337	0.345	0.351	0.356	0.362	0.367	0.372	0.377	0.583	0.589	0.595	0.517	0.508	0.536	0.622	0.556	0.030	0.643	0.583	0.055	0.00	0.602
Alagoas	0.355	0.363	0.371	0.379	0.387	0.395	0.404	0.412	0.419	0.425	0.431	0.436	0.442	0.448	0.455	0.462	0.47	0.478	0.487	0.496	0.505	0.514	0.523	0.531	0.539	0.546	0.552	0.556
Amapá	0.467	0.475	0.483	0.491	0.5	0.508	0.517	0.526	0.534	0.54	0.546	0.552	0.558	0.564	0.57	0.576	0.583	0.591	0.598	0.605	0.613	0.621	0.629	0.636	0.643	0.65	0.655	0.659
Amazonas	0.438	0.447	0.457	0.466	0.475	0.483	0.492	0.499	0.505	0.51	0.514	0.519	0.523	0.528	0.533	0.539	0.546	0.553	0.561	0.568	0.577	0.585	0.594	0.602	0.611	0.618	0.625	0.629
Bahia	0.402	0.41	0.419	0.427	0.435	0.443	0.451	0.459	0.465	0.47	0.475	0.48	0.485	0.491	0.496	0.503	0.51	0.518	0.526	0.534	0.542	0.551	0.559	0.567	0.575	0.582	0.587	0.591
Ceará	0.411	0.419	0.426	0.433	0.44	0.448	0.455	0.463	0.469	0.475	0.48	0.486	0.492	0.498	0.505	0.512	0.52	0.528	0.536	0.544	0.553	0.561	0.569	0.577	0.584	0.591	0.596	0.6
Distrito Federal	0.63	0.636	0.642	0.649	0.656	0.663	0.671	0.679	0.685	0.691	0.696	0.702	0.707	0.713	0.719	0.725	0.731	0.738	0.744	0.75	0.756	0.763	0.769	0.775	0.78	0.785	0.789	0.792
Espírito Santo	0.499	0.507	0.515	0.524	0.532	0.54	0.549	0.557	0.564	0.57	0.576	0.582	0.588	0.593	0.599	0.606	0.612	0.618	0.625	0.631	0.638	0.644	0.651	0.657	0.663	0.669	0.673	0.677
Goiás	0.46	0.468	0.476	0.484	0.493	0.501	0.51	0.518	0.526	0.532	0.538	0.545	0.551	0.558	0.564	0.571	0.579	0.586	0.594	0.601	0.608	0.616	0.623	0.63	0.636	0.642	0.647	0.65

Appendix Table 10: Socio-Demographic	Index val	ues for all	estimated	GBD 201	7 locations	, 1990-201	7										_											
Location	1990	1991	1992	1993	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015	2016	2017
Maranhão	0.313	0.322	0.33	0.339	0.347	0.355	0.364	0.371	0.377	0.38	0.383	0.386	0.389	0.392	0.396	0.402	0.409	0.418	0.427	0.436	0.446	0.456	0.467	0.477	0.486	0.495	0.502	0.507
Mato Grosso	0.475	0.484	0.492	0.501	0.509	0.518	0.527	0.535	0.543	0.548	0.554	0.559	0.564	0.57	0.576	0.582	0.589	0.596	0.604	0.611	0.618	0.626	0.633	0.641	0.648	0.654	0.659	0.662
Mato Grosso do Sul	0.465	0.473	0.481	0.489	0.497	0.506	0.515	0.523	0.531	0.537	0.543	0.549	0.555	0.56	0.566	0.573	0.58	0.587	0.594	0.6	0.607	0.614	0.622	0.629	0.636	0.642	0.647	0.65
Minas Gerais Pará	0.491	0.498	0.506	0.513	0.521	0.53	0.538	0.545	0.551	0.557	0.562	0.567	0.573	0.579	0.585	0.591	0.598	0.604	0.611	0.618	0.624	0.631	0.637	0.643	0.649	0.654	0.658	0.661
Paraiba	0.399	0.406	0.413	0.432	0.427	0.434	0.441	0.401	0.453	0.47	0.473	0.465	0.469	0.474	0.48	0.435	0.493	0.501	0.509	0.521	0.529	0.535	0.543	0.551	0.559	0.565	0.571	0.579
Paraná	0.513	0.519	0.525	0.532	0.539	0.548	0.556	0.564	0.572	0.578	0.585	0.591	0.597	0.603	0.609	0.615	0.622	0.628	0.634	0.64	0.646	0.652	0.658	0.664	0.67	0.675	0.679	0.682
Pernambuco	0.416	0.423	0.43	0.437	0.444	0.451	0.458	0.466	0.472	0.477	0.481	0.486	0.492	0.497	0.503	0.51	0.517	0.525	0.533	0.54	0.548	0.556	0.564	0.572	0.579	0.585	0.59	0.594
Piauí	0.365	0.372	0.379	0.386	0.393	0.4	0.408	0.415	0.42	0.425	0.429	0.434	0.439	0.444	0.45	0.457	0.465	0.473	0.482	0.491	0.5	0.51	0.518	0.527	0.535	0.542	0.548	0.552
Rio de Janeiro	0.576	0.581	0.585	0.59	0.595	0.601	0.608	0.614	0.619	0.624	0.628	0.632	0.637	0.641	0.645	0.65	0.655	0.66	0.665	0.67	0.675	0.681	0.686	0.692	0.697	0.702	0.706	0.709
Rio Grande do Norte	0.415	0.422	0.429	0.436	0.444	0.451	0.46	0.467	0.474	0.48	0.485	0.491	0.497	0.503	0.509	0.516	0.524	0.532	0.541	0.549	0.558	0.567	0.575	0.583	0.59	0.597	0.602	0.605
Rio Grande do Sul	0.543	0.549	0.555	0.561	0.567	0.574	0.581	0.587	0.593	0.598	0.603	0.608	0.614	0.619	0.624	0.63	0.635	0.641	0.647	0.653	0.659	0.665	0.67	0.676	0.681	0.686	0.69	0.693
Rondônia	0.423	0.433	0.441	0.45	0.458	0.467	0.475	0.484	0.491	0.497	0.502	0.508	0.515	0.521	0.528	0.535	0.543	0.551	0.559	0.567	0.575	0.584	0.592	0.599	0.606	0.613	0.618	0.622
Roraima	0.428	0.438	0.447	0.456	0.465	0.474	0.483	0.492	0.499	0.504	0.509	0.514	0.521	0.527	0.534	0.543	0.552	0.562	0.572	0.581	0.591	0.601	0.611	0.62	0.628	0.636	0.642	0.646
Santa Catarina	0.541	0.548	0.554	0.56	0.567	0.574	0.582	0.589	0.595	0.601	0.606	0.612	0.618	0.623	0.629	0.635	0.641	0.647	0.653	0.659	0.665	0.672	0.678	0.684	0.69	0.695	0.699	0.702
Sergine	0.338	0.433	0.372	0.379	0.387	0.393	0.473	0.481	0.488	0.494	0.65	0.506	0.512	0.518	0.632	0.531	0.538	0.546	0.554	0.68	0.685	0.578	0.586	0.594	0.601	0.607	0.612	0.72
Tocantins	0.396	0.404	0.412	0.443	0.430	0.436	0.445	0.453	0.46	0.466	0.471	0.477	0.312	0.491	0.324	0.507	0.538	0.540	0.537	0.502	0.558	0.578	0.530	0.586	0.594	0.601	0.607	0.611
Paraguay	0.467	0.471	0.475	0.48	0.485	0.491	0.497	0.504	0.512	0.519	0.525	0.532	0.538	0.544	0.548	0.553	0.558	0.562	0.566	0.569	0.573	0.577	0.582	0.589	0.598	0.606	0.613	0.619
North Africa and Middle East	0.456	0.464	0.472	0.478	0.485	0.492	0.499	0.506	0.513	0.521	0.531	0.539	0.547	0.553	0.561	0.568	0.574	0.582	0.589	0.596	0.602	0.607	0.612	0.617	0.622	0.628	0.635	0.639
North Africa and Middle East	0.456	0.464	0.472	0.478	0.485	0.492	0.499	0.506	0.513	0.521	0.531	0.539	0.547	0.553	0.561	0.568	0.574	0.582	0.589	0.596	0.602	0.607	0.612	0.617	0.622	0.628	0.635	0.639
Afghanistan	0.149	0.15	0.151	0.152	0.151	0.151	0.152	0.152	0.154	0.155	0.157	0.159	0.164	0.171	0.177	0.184	0.191	0.199	0.208	0.218	0.228	0.238	0.248	0.258	0.267	0.276	0.284	0.29
Algeria	0.495	0.504	0.513	0.522	0.531	0.54	0.549	0.558	0.567	0.574	0.583	0.591	0.599	0.608	0.617	0.625	0.633	0.641	0.648	0.653	0.659	0.665	0.67	0.675	0.68	0.685	0.691	0.696
Bahrain	0.612	0.618	0.624	0.63	0.635	0.639	0.644	0.648	0.653	0.657	0.662	0.668	0.673	0.677	0.68	0.684	0.687	0.69	0.693	0.695	0.696	0.697	0.698	0.7	0.702	0.706	0.709	0.712
Egypt	0.441	0.451	0.461	0.47	0.477	0.485	0.493	0.5	0.507	0.514	0.52	0.526	0.532	0.537	0.542	0.547	0.55	0.553	0.556	0.559	0.563	0.565	0.564	0.566	0.572	0.584	0.596	0.604
Iran	0.503	0.513	0.522	0.532	0.541	0.548	0.553	0.556	0.56	0.572	0.589	0.604	0.616	0.626	0.634	0.641	0.649	0.656	0.662	0.667	0.672	0.677	0.68	0.683	0.687	0.691	0.696	0.7
Iraq	0.433	0.433	0.437	0.441	0.444	0.446	0.45	0.455	0.463	0.473	0.481	0.488	0.494	0.496	0.502	0.507	0.512	0.517	0.522	0.528	0.535	0.542	0.551	0.559	0.566	0.572	0.58	0.585
Jordan	0.552	0.556	0.561	0.566	0.57	0.574	0.578	0.584	0.589	0.594	0.597	0.6	0.603	0.607	0.612	0.616	0.621	0.626	0.633	0.045	0.655	0.062	0.671	0.679	0.685	0.09	0.094	0.097
Lehanon	0.519	0.528	0.536	0.544	0.633	0.632	0.569	0.578	0.588	0.597	0.604	0.611	0.618	0.626	0.635	0.643	0.65	0.658	0.75	0.737	0.743	0.748	0.734	0.762	0.717	0.770	0.781	0.780
Libva	0.645	0.657	0.669	0.679	0.688	0.696	0.705	0.713	0.72	0.725	0.729	0.732	0.736	0.742	0.747	0.753	0.759	0.765	0.771	0.777	0.783	0.781	0.784	0.783	0.779	0.773	0.766	0.761
Могоссо	0.389	0.397	0.404	0.411	0.419	0.426	0.434	0.442	0.449	0.456	0.463	0.47	0.477	0.484	0.49	0.496	0.503	0.51	0.517	0.525	0.532	0.54	0.547	0.554	0.561	0.567	0.574	0.579
Palestine	0.362	0.366	0.371	0.377	0.384	0.39	0.397	0.405	0.413	0.421	0.427	0.431	0.433	0.435	0.439	0.444	0.448	0.453	0.46	0.468	0.479	0.491	0.503	0.513	0.521	0.528	0.536	0.541
Oman	0.473	0.485	0.497	0.509	0.521	0.533	0.546	0.558	0.57	0.582	0.594	0.606	0.618	0.629	0.639	0.649	0.659	0.668	0.678	0.686	0.695	0.703	0.711	0.72	0.727	0.734	0.739	0.744
Qatar	0.603	0.615	0.626	0.637	0.647	0.656	0.665	0.674	0.681	0.688	0.695	0.701	0.706	0.712	0.717	0.721	0.725	0.728	0.731	0.735	0.738	0.743	0.747	0.751	0.755	0.759	0.762	0.766
Saudi Arabia	0.436	0.457	0.477	0.496	0.514	0.53	0.545	0.56	0.573	0.586	0.598	0.61	0.621	0.633	0.645	0.658	0.67	0.682	0.695	0.706	0.716	0.728	0.738	0.748	0.757	0.765	0.773	0.779
Sudan	0.221	0.228	0.236	0.244	0.252	0.26	0.267	0.275	0.283	0.291	0.3	0.308	0.317	0.326	0.336	0.346	0.357	0.367	0.378	0.389	0.401	0.414	0.425	0.437	0.448	0.46	0.47	0.478
Syria	0.389	0.394	0.401	0.409	0.419	0.429	0.439	0.448	0.458	0.466	0.474	0.481	0.489	0.496	0.506	0.518	0.527	0.536	0.545	0.556	0.566	0.576	0.584	0.589	0.595	0.601	0.606	0.611
Turkay	0.453	0.465	0.473	0.482	0.491	0.548	0.511	0.521	0.532	0.545	0.555	0.504	0.572	0.582	0.592	0.625	0.609	0.617	0.625	0.652	0.659	0.645	0.688	0.656	0.001	0.000	0.671	0.675
United Arab Emirates	0.507	0.510	0.524	0.555	0.54	0.548	0.556	0.304	0.372	0.379	0.387	0.394	0.749	0.755	0.010	0.625	0.033	0.043	0.032	0.84	0.788	0.078	0.088	0.793	0.700	0.713	0.723	0.729
Yemen	0.203	0.208	0.214	0.22	0.228	0.238	0.247	0.256	0.266	0.275	0.284	0.294	0.304	0.313	0.324	0.336	0.349	0.362	0.373	0.382	0.393	0.401	0.408	0.415	0.422	0.426	0.429	0.43
South Asia	0.312	0.317	0.324	0.331	0.338	0.345	0.352	0.358	0.363	0.371	0.38	0.388	0.395	0.403	0.412	0.421	0.43	0.438	0.446	0.455	0.462	0.468	0.476	0.489	0.504	0.517	0.526	0.534
South Asia	0.312	0.317	0.324	0.331	0.338	0.345	0.352	0.358	0.363	0.371	0.38	0.388	0.395	0.403	0.412	0.421	0.43	0.438	0.446	0.455	0.462	0.468	0.476	0.489	0.504	0.517	0.526	0.534
Bangladesh	0.256	0.264	0.272	0.281	0.289	0.298	0.306	0.314	0.322	0.329	0.335	0.34	0.345	0.35	0.356	0.362	0.369	0.377	0.385	0.393	0.401	0.41	0.418	0.427	0.435	0.443	0.451	0.458
Bhutan	0.324	0.333	0.343	0.352	0.362	0.372	0.382	0.392	0.402	0.412	0.422	0.432	0.442	0.452	0.461	0.47	0.479	0.489	0.498	0.507	0.516	0.525	0.534	0.542	0.549	0.557	0.564	0.57
India	0.32	0.324	0.331	0.339	0.346	0.353	0.36	0.365	0.37	0.378	0.387	0.395	0.403	0.412	0.422	0.431	0.44	0.448	0.457	0.465	0.472	0.477	0.485	0.5	0.517	0.532	0.542	0.55
Andhra Pradesh	0.28	0.286	0.294	0.303	0.312	0.322	0.331	0.337	0.344	0.352	0.363	0.374	0.384	0.395	0.406	0.415	0.424	0.434	0.442	0.451	0.459	0.466	0.473	0.487	0.503	0.517	0.528	0.536
Arunachal Pradesh	0.311	0.317	0.323	0.33	0.336	0.345	0.354	0.36	0.366	0.374	0.381	0.388	0.395	0.405	0.416	0.424	0.432	0.44	0.447	0.458	0.467	0.473	0.479	0.498	0.52	0.536	0.548	0.556
Bihar	0.327	0.332	0.339	0.343	0.298	0.338	0.303	0.307	0.371	0.291	0.369	0.397	0.400	0.299	0.303	0.308	0.459	0.324	0.435	0.439	0.404	0.467	0.365	0.485	0.5	0.515	0.325	0.55
Chhattisgarh	0.292	0.295	0.297	0.303	0.298	0.311	0.316	0.319	0.322	0.328	0.336	0.344	0.352	0.362	0.303	0.382	0.392	0.403	0.413	0.421	0.427	0.333	0.305	0.455	0.474	0.49	0.503	0.455
Delhi	0.486	0.493	0.504	0.516	0.522	0.526	0.531	0.535	0.537	0.545	0.555	0.565	0.575	0.584	0.593	0.602	0.612	0.62	0.628	0.637	0.644	0.649	0.656	0.67	0.686	0.7	0.709	0.715
Goa	0.499	0.505	0.512	0.519	0.526	0.534	0.542	0.55	0.56	0.571	0.58	0.588	0.597	0.606	0.616	0.626	0.634	0.643	0.655	0.665	0.674	0.683	0.691	0.702	0.714	0.724	0.733	0.74
Gujarat	0.35	0.353	0.361	0.37	0.379	0.388	0.396	0.402	0.409	0.418	0.426	0.433	0.438	0.447	0.457	0.469	0.478	0.487	0.496	0.506	0.515	0.521	0.528	0.541	0.556	0.569	0.577	0.584
Haryana	0.342	0.348	0.356	0.364	0.372	0.38	0.388	0.394	0.398	0.405	0.414	0.423	0.431	0.44	0.451	0.462	0.472	0.481	0.491	0.5	0.508	0.512	0.521	0.54	0.564	0.582	0.593	0.6
Himachal Pradesh	0.34	0.347	0.356	0.366	0.376	0.386	0.397	0.408	0.418	0.431	0.445	0.457	0.468	0.479	0.491	0.502	0.513	0.524	0.534	0.544	0.553	0.562	0.571	0.585	0.601	0.614	0.625	0.633
Jammu and Kashmir	0.334	0.337	0.341	0.349	0.356	0.363	0.37	0.38	0.389	0.402	0.417	0.43	0.442	0.454	0.466	0.476	0.485	0.493	0.503	0.511	0.519	0.527	0.535	0.547	0.562	0.574	0.582	0.59
Jharkhand	0.281	0.283	0.286	0.29	0.293	0.297	0.3	0.304	0.307	0.312	0.319	0.325	0.333	0.342	0.353	0.362	0.371	0.38	0.388	0.397	0.405	0.411	0.419	0.434	0.451	0.466	0.478	0.487
Kamataka	0.323	0.331	0.339	0.349	0.357	0.366	0.375	0.38	0.386	0.396	0.406	0.415	0.424	0.434	0.444	0.454	0.463	0.472	0.481	0.49	0.498	0.504	0.512	0.526	0.542	0.556	0.566	0.574
Kerala Madhan Pradach	0.404	0.408	0.418	0.429	0.438	0.447	0.456	0.463	0.472	0.483	0.496	0.506	0.515	0.525	0.536	0.548	0.557	0.565	0.574	0.581	0.587	0.592	0.599	0.612	0.629	0.642	0.652	0.659
Maharashtra	0.284	0.288	0.293	0.299	0.303	0.309	0.315	0.31/	0.319	0.320	0.554	0.342	0.548	0.355	0.363	0.509	0.573	0.578	0.584	0.538	0.594	0.598	0.559	0.425	0.588	0.403	0.611	0.487
Manipur	0.381	0.387	0.393	0.4	0.405	0.411	0.417	0.421	0.427	0.436	0.445	0.454	0.461	0.469	0.479	0.49	0.498	0.505	0.512	0.52	0.526	0.532	0.539	0.549	0.561	0.572	0.582	0.59
Meghalaya	0.334	0.339	0.345	0.352	0.357	0.364	0.37	0.376	0.382	0.39	0.401	0.411	0.42	0.43	0.441	0.45	0.46	0.469	0.478	0.487	0.495	0.501	0.507	0.52	0.534	0.547	0.557	0.565
Mizoram	0.392	0.399	0.408	0.417	0.424	0.433	0.443	0.449	0.456	0.465	0.475	0.483	0.492	0.502	0.511	0.52	0.526	0.531	0.537	0.545	0.55	0.552	0.557	0.57	0.587	0.6	0.609	0.616
Nagaland	0.395	0.399	0.406	0.414	0.421	0.429	0.435	0.441	0.445	0.452	0.462	0.472	0.482	0.492	0.503	0.513	0.522	0.532	0.543	0.552	0.56	0.567	0.574	0.587	0.602	0.614	0.625	0.633
Odisha	0.283	0.287	0.292	0.299	0.305	0.314	0.32	0.325	0.331	0.34	0.349	0.357	0.364	0.373	0.384	0.393	0.404	0.415	0.425	0.434	0.441	0.447	0.456	0.472	0.49	0.504	0.515	0.524
Punjab	0.383	0.388	0.396	0.404	0.413	0.422	0.429	0.435	0.441	0.449	0.459	0.468	0.475	0.483	0.492	0.501	0.511	0.52	0.53	0.539	0.547	0.554	0.563	0.576	0.592	0.605	0.614	0.622

Appendix Table 10: Socio-Demographic	Index val	ues for all	estimated	GBD 201	7 locations	, 1990-201	7																					
Location	1990	1991	1992	1993	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015	2016	2017
Rajasthan	0.263	0.268	0.274	0.28	0.287	0.295	0.303	0.309	0.314	0.322	0.332	0.342	0.349	0.359	0.369	0.377	0.385	0.391	0.398	0.404	0.41	0.415	0.421	0.438	0.458	0.473	0.484	0.492
Sikkim	0.343	0.349	0.355	0.362	0.368	0.375	0.382	0.388	0.395	0.403	0.413	0.421	0.431	0.442	0.453	0.463	0.473	0.482	0.492	0.512	0.528	0.541	0.555	0.574	0.593	0.608	0.619	0.628
Tamil Nadu	0.35	0.354	0.363	0.373	0.383	0.391	0.399	0.406	0.414	0.424	0.436	0.446	0.455	0.465	0.475	0.486	0.497	0.506	0.514	0.522	0.529	0.535	0.543	0.559	0.579	0.595	0.607	0.615
Telangana	0.288	0.294	0.3	0.307	0.315	0.323	0.331	0.337	0.344	0.352	0.363	0.373	0.383	0.395	0.407	0.418	0.429	0.441	0.454	0.465	0.478	0.488	0.5	0.517	0.536	0.552	0.565	0.575
Tripura	0.333	0.336	0.34	0.346	0.35	0.356	0.363	0.368	0.373	0.385	0.398	0.409	0.419	0.43	0.44	0.449	0.457	0.462	0.469	0.476	0.48	0.483	0.488	0.5	0.515	0.527	0.536	0.543
Uttar Pradesh	0.272	0.276	0.281	0.286	0.291	0.297	0.303	0.307	0.31	0.317	0.325	0.333	0.34	0.348	0.358	0.366	0.374	0.382	0.39	0.398	0.404	0.41	0.419	0.436	0.454	0.469	0.48	0.488
Uttarakhand	0.304	0.306	0.31	0.315	0.321	0.329	0.336	0.341	0.347	0.355	0.367	0.378	0.391	0.405	0.42	0.434	0.449	0.463	0.478	0.494	0.509	0.522	0.535	0.552	0.57	0.586	0.598	0.607
West Bengal	0.321	0.322	0.329	0.338	0.345	0.353	0.36	0.366	0.371	0.38	0.389	0.398	0.406	0.416	0.425	0.434	0.441	0.448	0.455	0.462	0.467	0.471	0.476	0.49	0.506	0.519	0.53	0.538
Union Territories other than Delhi	0.406	0.408	0.412	0.417	0.423	0.43	0.439	0.446	0.453	0.463	0.474	0.486	0.498	0.511	0.524	0.537	0.548	0.559	0.569	0.579	0.586	0.591	0.597	0.61	0.624	0.636	0.646	0.653
Nepal	0.216	0.222	0.228	0.235	0.242	0.249	0.257	0.265	0.273	0.281	0.29	0.298	0.307	0.315	0.323	0.331	0.339	0.347	0.355	0.364	0.373	0.382	0.39	0.399	0.407	0.415	0.422	0.429
Pakistan	0.305	0.31	0.314	0.319	0.324	0.33	0.337	0.343	0.35	0.359	0.367	0.375	0.382	0.389	0.396	0.404	0.412	0.42	0.428	0.436	0.443	0.449	0.458	0.466	0.474	0.48	0.486	0.492
Southeast Asia, East Asia, and Oceania	0.468	0.479	0.489	0.498	0.506	0.517	0.528	0.538	0.546	0.553	0.561	0.568	0.574	0.58	0.588	0.596	0.606	0.615	0.624	0.632	0.641	0.649	0.656	0.662	0.67	0.673	0.675	0.685
East Asia	0.465	0.476	0.480	0.495	0.504	0.516	0.528	0.54	0.55	0.559	0.568	0.577	0.584	0.591	0.599	0.61	0.621	0.632	0.642	0.652	0.661	0.671	0.679	0.685	0.694	0.696	0.696	0.709
North Korne	0.430	0.409	0.479	0.469	0.524	0.51	0.525	0.555	0.545	0.534	0.503	0.575	0.38	0.387	0.393	0.600	0.502	0.05	0.699	0.65	0.601	0.6/1	0.078	0.065	0.695	0.695	0.625	0.529
Taiwan (Province of China)	0.512	0.317	0.32	0.324	0.722	0.322	0.739	0.313	0.759	0.300	0.303	0.3	0.498	0.498	0.499	0.501	0.505	0.300	0.308	0.933	0.939	0.842	0.52	0.324	0.854	0.551	0.355	0.338
Oceania	0.406	0.408	0.41	0.412	0.415	0.418	0.421	0.423	0.426	0.428	0.43	0.421	0.432	0.733	0.435	0.436	0.439	0.44	0.442	0.033	0.446	0.449	0.452	0.455	0.459	0.463	0.467	0.471
American Samoa	0.609	0.612	0.615	0.62	0.624	0.679	0.633	0.636	0.639	0.643	0.45	0.651	0.655	0.455	0.662	0.666	0.668	0.671	0.673	0.675	0.678	0.682	0.687	0.691	0.694	0.697	0.407	0.702
Federated States of Micronesia	0.462	0.469	0.476	0.483	0.489	0.496	0.501	0.505	0.509	0.514	0.518	0.523	0.527	0.532	0.536	0.54	0.543	0.547	0.549	0.552	0.555	0.559	0.562	0.565	0.567	0.57	0.573	0.575
Fiii	0.533	0.537	0.542	0.547	0.552	0.558	0.563	0.568	0.574	0.58	0.585	0.589	0.594	0.598	0.602	0.604	0.607	0.61	0.611	0.613	0.615	0.617	0.619	0.623	0.627	0.632	0.637	0.641
Guam	0.698	0.695	0.693	0.694	0.698	0.703	0.709	0.717	0.727	0.739	0.75	0.759	0.765	0.768	0.77	0.77	0.769	0.77	0.771	0.774	0.776	0.78	0.784	0.788	0.792	0.794	0.794	0.794
Kiribati	0.355	0.357	0.359	0.361	0.363	0.365	0.368	0.37	0.373	0.376	0.379	0.382	0.385	0.388	0.39	0.393	0.395	0.397	0.398	0.399	0.401	0.403	0.406	0.41	0.414	0.418	0.423	0.427
Marshall Islands	0.413	0.421	0.429	0.436	0.444	0.451	0.457	0.462	0.465	0.469	0.473	0.477	0.482	0.485	0.489	0.493	0.498	0.503	0.507	0.512	0.518	0.523	0.528	0.533	0.538	0.542	0.547	0.55
Northern Mariana Islands	0.738	0.744	0.748	0.75	0.752	0.754	0.756	0.757	0.758	0.759	0.763	0.766	0.767	0.767	0.767	0.766	0.765	0.764	0.763	0.761	0.759	0.758	0.757	0.756	0.756	0.756	0.757	0.758
Papua New Guinea	0.318	0.321	0.324	0.33	0.335	0.34	0.344	0.348	0.351	0.354	0.356	0.358	0.36	0.361	0.363	0.365	0.368	0.371	0.375	0.379	0.383	0.388	0.393	0.398	0.404	0.409	0.415	0.419
Samoa	0.538	0.539	0.538	0.536	0.535	0.536	0.536	0.537	0.538	0.54	0.542	0.544	0.546	0.548	0.55	0.552	0.554	0.556	0.558	0.559	0.561	0.563	0.56	0.558	0.562	0.567	0.572	0.576
Solomon Islands	0.316	0.32	0.326	0.331	0.337	0.344	0.35	0.355	0.36	0.364	0.365	0.366	0.366	0.366	0.367	0.369	0.371	0.375	0.38	0.384	0.388	0.394	0.4	0.406	0.411	0.416	0.421	0.425
Tonga	0.522	0.528	0.533	0.538	0.544	0.549	0.553	0.557	0.56	0.564	0.566	0.569	0.573	0.577	0.581	0.584	0.587	0.589	0.592	0.595	0.599	0.603	0.607	0.61	0.614	0.617	0.621	0.625
Vanuatu	0.38	0.384	0.388	0.391	0.395	0.398	0.402	0.406	0.409	0.413	0.416	0.419	0.421	0.423	0.425	0.428	0.431	0.435	0.44	0.445	0.449	0.453	0.458	0.462	0.465	0.469	0.472	0.475
Southeast Asia	0.467	0.477	0.486	0.495	0.504	0.513	0.521	0.529	0.534	0.538	0.543	0.548	0.552	0.557	0.562	0.567	0.572	0.578	0.584	0.589	0.595	0.602	0.609	0.616	0.622	0.629	0.635	0.641
Cambodia	0.259	0.266	0.273	0.279	0.286	0.294	0.302	0.31	0.317	0.326	0.335	0.344	0.353	0.362	0.372	0.382	0.392	0.401	0.411	0.42	0.427	0.435	0.443	0.451	0.459	0.467	0.475	0.482
Indonesia	0.465	0.476	0.487	0.499	0.51	0.52	0.53	0.537	0.542	0.546	0.55	0.554	0.558	0.562	0.566	0.571	0.576	0.581	0.587	0.594	0.6	0.608	0.615	0.622	0.629	0.636	0.642	0.648
Laos	0.307	0.313	0.32	0.326	0.332	0.339	0.345	0.352	0.358	0.364	0.371	0.378	0.385	0.392	0.4	0.407	0.416	0.425	0.434	0.444	0.453	0.463	0.473	0.483	0.493	0.502	0.511	0.519
Malaysia	0.57	0.575	0.581	0.589	0.598	0.606	0.616	0.626	0.638	0.654	0.661	0.664	0.671	0.679	0.684	0.689	0.697	0.704	0.709	0.714	0.721	0.727	0.733	0.739	0.744	0.749	0.754	0.759
Maldives	0.386	0.399	0.412	0.424	0.435	0.446	0.46	0.477	0.493	0.507	0.518	0.528	0.537	0.545	0.554	0.56	0.567	0.575	0.584	0.593	0.602	0.611	0.62	0.629	0.636	0.643	0.65	0.655
Mauritius	0.554	0.561	0.568	0.576	0.585	0.595	0.603	0.609	0.613	0.616	0.621	0.627	0.633	0.639	0.646	0.651	0.657	0.663	0.669	0.675	0.68	0.687	0.694	0.701	0.708	0.713	0.717	0.72
Myanmar	0.33	0.333	0.337	0.341	0.347	0.353	0.36	0.367	0.375	0.383	0.392	0.402	0.412	0.423	0.434	0.446	0.458	0.47	0.481	0.492	0.501	0.51	0.518	0.527	0.535	0.542	0.549	0.556
Philippines	0.511	0.516	0.521	0.525	0.53	0.534	0.539	0.542	0.545	0.547	0.55	0.553	0.555	0.557	0.559	0.561	0.563	0.566	0.568	0.569	0.572	0.579	0.586	0.593	0.599	0.605	0.612	0.617
Sri Lanka	0.49	0.495	0.501	0.508	0.516	0.524	0.532	0.54	0.547	0.553	0.559	0.565	0.571	0.578	0.584	0.59	0.597	0.604	0.611	0.618	0.626	0.634	0.642	0.65	0.658	0.666	0.673	0.68
Seyeneiles	0.549	0.557	0.565	0.573	0.582	0.589	0.597	0.605	0.613	0.62	0.626	0.631	0.636	0.64	0.645	0.640	0.65	0.653	0.656	0.658	0.641	0.647	0.667	0.671	0.675	0.68	0.680	0.692
Timer Lett	0.302	0.314	0.525	0.554	0.342	0.352	0.361	0.367	0.309	0.372	0.379	0.587	0.394	0.0	0.605	0.61	0.616	0.623	0.629	0.635	0.641	0.647	0.654	0.66	0.067	0.673	0.679	0.684
Vietnam	0.276	0.285	0.29	0.290	0.302	0.307	0.452	0.321	0.323	0.321	0.32	0.323	0.332	0.543	0.502	0.519	0.525	0.419	0.437	0.547	0.40	0.4/1	0.481	0.49	0.495	0.592	0.504	0.505
Sub-Saharan Africa	0.304	0.307	0.311	0.314	0.317	0.32	0.324	0.328	0.332	0.335	0.339	0.343	0.348	0.353	0.359	0.365	0.371	0.379	0.386	0.393	0.554	0.407	0.414	0.421	0.428	0.435	0.441	0.446
Central sub-Saharan Africa	0.298	0.303	0.307	0.309	0.311	0.313	0.316	0.318	0.32	0.323	0.325	0.328	0.332	0.336	0.341	0.348	0.355	0.364	0.373	0.382	0.391	0.402	0.413	0.423	0.433	0.443	0.452	0.457
Angola	0.235	0.24	0.245	0.249	0.253	0.258	0.263	0.269	0.276	0.282	0.288	0.293	0.299	0.305	0.312	0.32	0.329	0.34	0.351	0.363	0.375	0.389	0.401	0.414	0.428	0.441	0.453	0.461
Central African Republic	0.22	0.225	0.228	0.232	0.236	0.24	0.242	0.245	0.249	0.254	0.257	0.261	0.265	0.268	0.271	0.275	0.28	0.285	0.29	0.296	0.304	0.313	0.323	0.325	0.328	0.33	0.333	0.334
Congo (Brazzaville)	0.382	0.39	0.398	0.405	0.41	0.416	0.421	0.426	0.43	0.434	0.439	0.444	0.449	0.455	0.46	0.467	0.475	0.482	0.49	0.499	0.509	0.52	0.531	0.542	0.552	0.561	0.569	0.574
DR Congo	0.293	0.296	0.298	0.298	0.296	0.294	0.291	0.288	0.283	0.279	0.274	0.269	0.265	0.263	0.262	0.264	0.264	0.265	0.267	0.27	0.278	0.288	0.3	0.315	0.33	0.344	0.356	0.364
Equatorial Guinea	0.2	0.204	0.212	0.22	0.229	0.241	0.26	0.292	0.316	0.339	0.363	0.388	0.41	0.429	0.449	0.467	0.483	0.499	0.516	0.53	0.544	0.559	0.573	0.587	0.599	0.61	0.62	0.625
Gabon	0.433	0.443	0.453	0.462	0.472	0.481	0.49	0.498	0.506	0.514	0.522	0.529	0.535	0.542	0.549	0.556	0.562	0.569	0.576	0.582	0.589	0.598	0.607	0.616	0.625	0.634	0.644	0.651
Eastern sub-Saharan Africa	0.23	0.233	0.236	0.239	0.241	0.245	0.249	0.254	0.259	0.262	0.266	0.271	0.276	0.282	0.288	0.294	0.301	0.308	0.316	0.324	0.332	0.34	0.348	0.356	0.365	0.373	0.381	0.387
Burundi	0.247	0.252	0.257	0.258	0.263	0.265	0.265	0.266	0.268	0.268	0.268	0.267	0.268	0.269	0.271	0.272	0.274	0.276	0.278	0.282	0.286	0.29	0.295	0.299	0.303	0.306	0.308	0.31
Comoros	0.272	0.279	0.286	0.293	0.298	0.303	0.306	0.31	0.314	0.319	0.325	0.331	0.338	0.344	0.351	0.358	0.365	0.372	0.378	0.384	0.39	0.396	0.403	0.41	0.417	0.423	0.429	0.434
Djibouti	0.313	0.317	0.32	0.322	0.325	0.329	0.333	0.337	0.339	0.341	0.342	0.347	0.359	0.374	0.388	0.4	0.407	0.412	0.419	0.425	0.432	0.439	0.446	0.454	0.462	0.47	0.478	0.485
Eritrea	0.202	0.214	0.223	0.234	0.247	0.26	0.272	0.285	0.296	0.306	0.315	0.323	0.331	0.337	0.343	0.348	0.353	0.357	0.36	0.364	0.368	0.372	0.378	0.383	0.39	0.396	0.403	0.409
Ethiopia	0.138	0.141	0.143	0.146	0.148	0.15	0.155	0.161	0.166	0.169	0.172	0.177	0.183	0.189	0.195	0.202	0.21	0.221	0.233	0.245	0.257	0.268	0.28	0.292	0.303	0.314	0.325	0.334
Kenya	0.341	0.349	0.357	0.364	0.372	0.377	0.382	0.387	0.392	0.398	0.401	0.403	0.406	0.411	0.416	0.42	0.425	0.432	0.438	0.445	0.452	0.459	0.465	0.473	0.481	0.488	0.494	0.499
Baringo	0.254	0.266	0.278	0.289	0.299	0.307	0.313	0.319	0.326	0.333	0.338	0.341	0.345	0.352	0.358	0.362	0.368	0.376	0.384	0.393	0.401	0.408	0.414	0.421	0.428	0.434	0.439	0.444
Bomet	0.306	0.315	0.325	0.333	0.341	0.347	0.351	0.355	0.361	0.367	0.371	0.373	0.378	0.385	0.392	0.398	0.406	0.414	0.423	0.433	0.442	0.449	0.456	0.465	0.475	0.483	0.49	0.496
Bungoma	0.316	0.325	0.333	0.341	0.348	0.353	0.357	0.36	0.365	0.37	0.373	0.373	0.376	0.38	0.384	0.387	0.391	0.397	0.403	0.41	0.417	0.423	0.429	0.436	0.445	0.451	0.458	0.463
Busia	0.297	0.304	0.312	0.32	0.327	0.332	0.336	0.339	0.344	0.349	0.352	0.353	0.356	0.361	0.367	0.37	0.375	0.381	0.386	0.393	0.4	0.404	0.409	0.415	0.423	0.428	0.434	0.438
Eigeyo Marakwet	0.292	0.302	0.312	0.321	0.329	0.336	0.342	0.348	0.355	0.362	0.368	0.372	0.378	0.386	0.394	0.4	0.408	0.417	0.425	0.435	0.443	0.451	0.458	0.467	0.475	0.483	0.49	0.496
Emou	0.375	0.384	0.393	0.4	0.407	0.413	0.417	0.422	0.427	0.431	0.434	0.437	0.44	0.444	0.449	0.452	0.458	0.464	0.47	0.478	0.486	0.493	0.499	0.507	0.514	0.521	0.527	0.533
Garissa Homa Pau	0.153	0.16	0.168	0.242	0.184	0.19	0.195	0.201	0.207	0.213	0.202	0.219	0.223	0.228	0.233	0.237	0.242	0.249	0.255	0.263	0.272	0.28	0.288	0.298	0.309	0.318	0.326	0.334
Iciala	0.214	0.222	0.232	0.243	0.253	0.20	0.205	0.2/1	0.279	0.288	0.292	0.293	0.297	0.305	0.313	0.319	0.328	0.338	0.346	0.330	0.300	0.374	0.382	0.392	0.403	0.411	0.419	0.425
Kajjado	0.394	0.202	0.270	0.407	0.414	0.42	0.425	0.420	0.424	0.429	0.442	0.308	0.449	0.452	0.516	0.321	0.320	0.331	0.357	0.491	0.35	0.492	0.30	0.505	0.514	0.521	0.539	0.524
Kakamega	0.295	0.392	0.311	0.319	0.326	0.332	0.337	0.342	0.348	0.356	0.36	0.361	0.365	0.37	0.375	0.378	0.383	0.389	0.394	0.4	0.407	0.412	0.417	0.425	0.433	0.439	0.445	0.354
	0.275	0.505	0.011	0.017	0.520	0.002	0.001	0.042	0.510	0.000	0.50	0.501	0.505	0.57	0.575	0.570	0.505	0.505	0.074	0.4	0.107	0.112	0.117	0.125	0.455	0.107	0.110	0.45

Appendix Table 10: Socio-Demographic	Index val	ues for all	estimated	GBD 201	7 locations	, 1990-201	7												_			-						
Location	1990	1991	1992	1993	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015	2016	2017
Kericho	0.266	0.277	0.288	0.299	0.309	0.317	0.324	0.331	0.339	0.348	0.353	0.356	0.362	0.37	0.378	0.385	0.394	0.404	0.414	0.425	0.436	0.445	0.454	0.464	0.475	0.485	0.493	0.5
Kiambu	0.435	0.443	0.45	0.457	0.464	0.469	0.473	0.476	0.48	0.484	0.487	0.489	0.492	0.496	0.5	0.504	0.509	0.516	0.521	0.528	0.535	0.541	0.548	0.555	0.562	0.569	0.575	0.58
Kilth	0.292	0.3	0.307	0.314	0.321	0.327	0.331	0.336	0.34	0.346	0.348	0.349	0.352	0.357	0.361	0.365	0.371	0.378	0.385	0.392	0.4	0.408	0.415	0.424	0.434	0.442	0.45	0.456
Kirinyaga	0.389	0.396	0.402	0.407	0.411	0.415	0.418	0.422	0.425	0.429	0.432	0.434	0.437	0.442	0.447	0.451	0.457	0.464	0.471	0.479	0.486	0.493	0.5	0.507	0.514	0.521	0.527	0.533
Kisumu	0.315	0.325	0.334	0.342	0.349	0.355	0.36	0.364	0.37	0.376	0.381	0.384	0.388	0.395	0.402	0.407	0.415	0.424	0.432	0.442	0.451	0.459	0.466	0.475	0.484	0.491	0.497	0.503
Kitui	0.28	0.288	0.297	0.304	0.311	0.317	0.322	0.327	0.332	0.338	0.343	0.346	0.351	0.357	0.363	0.369	0.376	0.383	0.391	0.399	0.408	0.416	0.423	0.432	0.44	0.448	0.455	0.461
Kwale	0.294	0.301	0.308	0.314	0.321	0.326	0.33	0.334	0.338	0.342	0.344	0.346	0.348	0.352	0.357	0.36	0.366	0.374	0.381	0.39	0.399	0.407	0.414	0.424	0.433	0.442	0.45	0.457
Laikipia	0.346	0.354	0.361	0.368	0.375	0.38	0.385	0.389	0.395	0.401	0.405	0.409	0.415	0.423	0.433	0.442	0.451	0.462	0.472	0.483	0.494	0.502	0.511	0.521	0.531	0.54	0.549	0.556
Lamu	0.295	0.303	0.312	0.321	0.328	0.334	0.338	0.343	0.347	0.352	0.355	0.356	0.359	0.364	0.369	0.373	0.378	0.384	0.39	0.397	0.405	0.411	0.417	0.425	0.434	0.441	0.448	0.453
Machakos	0.352	0.362	0.372	0.38	0.389	0.396	0.403	0.411	0.418	0.426	0.431	0.435	0.44	0.444	0.449	0.452	0.456	0.461	0.465	0.471	0.476	0.482	0.487	0.493	0.5	0.507	0.512	0.518
Makueni	0.3	0.307	0.314	0.322	0.328	0.333	0.336	0.339	0.343	0.348	0.35	0.351	0.354	0.36	0.367	0.373	0.381	0.391	0.401	0.413	0.423	0.432	0.439	0.447	0.454	0.46	0.465	0.469
Mandera	0.102	0.109	0.118	0.126	0.135	0.142	0.149	0.156	0.164	0.172	0.178	0.181	0.187	0.194	0.201	0.206	0.212	0.218	0.224	0.231	0.238	0.244	0.251	0.261	0.271	0.28	0.288	0.295
Marsabit	0.22	0.223	0.227	0.231	0.236	0.239	0.241	0.244	0.247	0.252	0.255	0.256	0.259	0.264	0.268	0.271	0.276	0.281	0.286	0.293	0.299	0.304	0.309	0.315	0.323	0.329	0.335	0.34
Meru	0.338	0.346	0.354	0.362	0.369	0.376	0.381	0.387	0.393	0.399	0.403	0.407	0.411	0.417	0.423	0.428	0.434	0.441	0.447	0.454	0.462	0.469	0.475	0.482	0.49	0.496	0.503	0.508
Migori	0.229	0.237	0.246	0.255	0.262	0.267	0.27	0.2/4	0.2/9	0.285	0.287	0.286	0.289	0.296	0.303	0.307	0.315	0.325	0.333	0.344	0.355	0.364	0.372	0.383	0.395	0.404	0.412	0.419
Murang'a	0.390	0.405	0.414	0.422	0.408	0.437	0.445	0.449	0.433	0.401	0.403	0.408	0.471	0.473	0.478	0.465	0.480	0.491	0.490	0.302	0.31	0.317	0.525	0.534	0.545	0.532	0.501	0.508
Nairobi	0.499	0.506	0.513	0.519	0.525	0.53	0.534	0.539	0.544	0.549	0.553	0.556	0.56	0.566	0.572	0.577	0.585	0.593	0.602	0.611	0.619	0.627	0.634	0.643	0.652	0.52	0.667	0.528
Nakuru	0.337	0.347	0.357	0.367	0.375	0.383	0.389	0.394	0.399	0.405	0.409	0.413	0.417	0.423	0.43	0.436	0.445	0.454	0.464	0.474	0.484	0.493	0.502	0.511	0.521	0.53	0.538	0.545
Nandi	0.323	0.333	0.344	0.354	0.362	0.369	0.375	0.381	0.388	0.394	0.399	0.402	0.406	0.412	0.418	0.422	0.428	0.435	0.441	0.449	0.456	0.462	0.468	0.475	0.483	0.49	0.496	0.501
Narok	0.217	0.225	0.234	0.242	0.25	0.255	0.259	0.264	0.272	0.28	0.284	0.285	0.289	0.297	0.305	0.31	0.318	0.326	0.333	0.342	0.35	0.356	0.362	0.371	0.381	0.389	0.396	0.402
Nyamira	0.361	0.37	0.379	0.387	0.395	0.401	0.406	0.41	0.414	0.419	0.421	0.423	0.426	0.431	0.437	0.443	0.451	0.46	0.468	0.478	0.488	0.496	0.504	0.513	0.523	0.531	0.538	0.544
Nyandarua	0.353	0.361	0.369	0.376	0.382	0.388	0.392	0.397	0.402	0.409	0.413	0.416	0.421	0.428	0.436	0.442	0.449	0.458	0.466	0.475	0.484	0.491	0.498	0.506	0.514	0.521	0.528	0.534
Nyeri	0.407	0.414	0.421	0.427	0.434	0.439	0.444	0.449	0.454	0.459	0.463	0.466	0.47	0.475	0.479	0.484	0.489	0.496	0.502	0.509	0.516	0.522	0.527	0.533	0.539	0.544	0.549	0.554
Samburu	0.205	0.21	0.215	0.221	0.225	0.228	0.23	0.232	0.235	0.239	0.24	0.239	0.24	0.243	0.247	0.249	0.253	0.258	0.262	0.267	0.273	0.277	0.28	0.287	0.294	0.299	0.304	0.308
Siaya	0.227	0.236	0.246	0.255	0.265	0.272	0.279	0.287	0.294	0.303	0.309	0.313	0.318	0.326	0.334	0.34	0.349	0.359	0.368	0.379	0.39	0.4	0.409	0.421	0.432	0.443	0.452	0.46
Taita Taveta	0.352	0.362	0.371	0.38	0.389	0.396	0.403	0.409	0.416	0.422	0.427	0.43	0.434	0.438	0.443	0.447	0.452	0.458	0.464	0.472	0.48	0.487	0.494	0.502	0.509	0.516	0.523	0.529
Tana River	0.231	0.237	0.244	0.25	0.255	0.26	0.263	0.267	0.27	0.274	0.276	0.276	0.278	0.281	0.285	0.288	0.292	0.298	0.304	0.311	0.319	0.326	0.333	0.342	0.353	0.362	0.371	0.379
I naraka Nitni Trans Nizoia	0.342	0.35	0.357	0.364	0.371	0.377	0.382	0.387	0.392	0.397	0.402	0.405	0.392	0.417	0.423	0.429	0.437	0.445	0.453	0.465	0.472	0.48	0.488	0.497	0.506	0.514	0.522	0.528
Turkana	0.211	0.216	0.221	0.226	0.231	0.333	0.237	0.239	0.242	0.36	0.334	0.333	0.392	0.25	0.252	0.252	0.253	0.255	0.257	0.76	0.448	0.266	0.40	0.403	0.281	0.484	0.291	0.490
Uasin Gishu	0.363	0.373	0.382	0.391	0.399	0.405	0.41	0.416	0.421	0.428	0.433	0.436	0.441	0.447	0.453	0.458	0.465	0.472	0.48	0.488	0.496	0.503	0.51	0.517	0.526	0.533	0.539	0.545
Vihiga	0.328	0.337	0.346	0.355	0.362	0.367	0.371	0.374	0.376	0.378	0.378	0.378	0.379	0.382	0.387	0.392	0.399	0.408	0.417	0.427	0.437	0.444	0.45	0.457	0.464	0.469	0.474	0.477
Wajir	0.104	0.111	0.12	0.128	0.136	0.142	0.146	0.151	0.156	0.162	0.165	0.166	0.168	0.172	0.176	0.178	0.181	0.185	0.189	0.195	0.201	0.206	0.21	0.218	0.226	0.233	0.238	0.243
West Pokot	0.213	0.219	0.226	0.233	0.24	0.245	0.249	0.253	0.258	0.265	0.268	0.27	0.274	0.28	0.286	0.291	0.298	0.305	0.312	0.319	0.327	0.333	0.34	0.349	0.359	0.368	0.376	0.382
Madagascar	0.262	0.264	0.265	0.265	0.265	0.265	0.266	0.268	0.27	0.272	0.274	0.278	0.282	0.286	0.291	0.294	0.295	0.295	0.298	0.3	0.302	0.303	0.306	0.31	0.315	0.321	0.326	0.331
Malawi	0.199	0.202	0.204	0.206	0.208	0.212	0.216	0.221	0.225	0.23	0.234	0.238	0.244	0.249	0.256	0.262	0.27	0.279	0.288	0.296	0.305	0.313	0.32	0.327	0.333	0.339	0.345	0.349
Mozambique	0.133	0.135	0.136	0.138	0.14	0.142	0.147	0.154	0.162	0.169	0.175	0.182	0.189	0.197	0.206	0.215	0.224	0.235	0.246	0.257	0.268	0.279	0.29	0.301	0.311	0.322	0.332	0.34
Rwanda	0.267	0.269	0.272	0.274	0.268	0.267	0.269	0.272	0.276	0.28	0.284	0.29	0.297	0.304	0.311	0.319	0.327	0.334	0.342	0.349	0.356	0.363	0.371	0.378	0.386	0.393	0.401	0.407
Somalia	0.153	0.156	0.157	0.159	0.16	0.161	0.162	0.163	0.163	0.164	0.165	0.167	0.169	0.172	0.176	0.179	0.183	0.188	0.193	0.198	0.203	0.208	0.212	0.217	0.222	0.227	0.231	0.235
South Sudan	0.179	0.18	0.182	0.183	0.184	0.186	0.187	0.189	0.191	0.194	0.197	0.2	0.204	0.207	0.211	0.215	0.22	0.225	0.229	0.234	0.239	0.244	0.249	0.254	0.259	0.265	0.27	0.275
Landa	0.273	0.28	0.285	0.289	0.173	0.178	0.185	0.193	0.304	0.212	0.222	0.313	0.32	0.323	0.33	0.334	0.339	0.344	0.33	0.335	0.30	0.339	0.373	0.357	0.366	0.397	0.403	0.412
Zambia	0.312	0.314	0.316	0.319	0.321	0.324	0.326	0.329	0.332	0.335	0.338	0.343	0.348	0.353	0.358	0.365	0.373	0.382	0.392	0.402	0.412	0.421	0.431	0.44	0.449	0.458	0.466	0.472
Southern sub-Saharan Africa	0.521	0.525	0.53	0.535	0.539	0.543	0.546	0.55	0.555	0.56	0.565	0.569	0.574	0.578	0.582	0.585	0.59	0.596	0.603	0.607	0.612	0.616	0.619	0.622	0.627	0.633	0.637	0.64
Botswana	0.463	0.473	0.483	0.492	0.501	0.509	0.517	0.524	0.532	0.539	0.546	0.553	0.56	0.567	0.574	0.581	0.588	0.597	0.605	0.612	0.619	0.626	0.633	0.639	0.646	0.652	0.658	0.663
Lesotho	0.333	0.34	0.347	0.354	0.362	0.369	0.376	0.383	0.389	0.396	0.402	0.407	0.413	0.417	0.422	0.426	0.431	0.436	0.441	0.446	0.452	0.458	0.464	0.47	0.476	0.482	0.488	0.493
Namibia	0.454	0.459	0.465	0.471	0.477	0.483	0.489	0.495	0.501	0.507	0.512	0.516	0.521	0.525	0.53	0.534	0.54	0.546	0.552	0.558	0.565	0.572	0.579	0.587	0.595	0.603	0.61	0.616
South Africa	0.557	0.56	0.565	0.57	0.574	0.576	0.579	0.583	0.587	0.593	0.598	0.602	0.607	0.611	0.615	0.619	0.624	0.631	0.639	0.644	0.649	0.653	0.655	0.658	0.664	0.669	0.674	0.677
Swaziland	0.426	0.435	0.444	0.452	0.459	0.466	0.473	0.478	0.483	0.489	0.494	0.499	0.503	0.508	0.513	0.518	0.523	0.528	0.533	0.538	0.543	0.547	0.552	0.557	0.563	0.569	0.574	0.578
Zimbabwe	0.408	0.417	0.424	0.43	0.436	0.44	0.446	0.45	0.453	0.456	0.457	0.459	0.458	0.456	0.453	0.449	0.443	0.438	0.43	0.426	0.424	0.425	0.43	0.436	0.444	0.451	0.458	0.463
Western sub-Saharan Africa	0.293	0.296	0.299	0.303	0.306	0.309	0.313	0.316	0.32	0.324	0.328	0.333	0.337	0.343	0.35	0.357	0.365	0.373	0.381	0.389	0.396	0.403	0.411	0.417	0.424	0.43	0.437	0.441
Burkina Faso	0.218	0.223	0.228	0.233	0.238	0.158	0.163	0.253	0.258	0.263	0.268	0.275	0.278	0.284	0.29	0.296	0.301	0.307	0.313	0.32	0.326	0.332	0.338	0.345	0.353	0.36	0.367	0.373
Cameroon	0.332	0.338	0.343	0.347	0.35	0.352	0.355	0.357	0.359	0.36	0.361	0.361	0.361	0.362	0.366	0.373	0.38	0.388	0.396	0.405	0.414	0.423	0.433	0.443	0.453	0.464	0.474	0.482
Cape Verde	0.307	0.315	0.322	0.329	0.337	0.347	0.357	0.367	0.377	0.387	0.399	0.41	0.42	0.431	0.441	0.451	0.461	0.472	0.482	0.491	0.5	0.509	0.517	0.524	0.531	0.538	0.544	0.549
Chad	0.119	0.123	0.128	0.131	0.135	0.138	0.141	0.144	0.146	0.148	0.15	0.152	0.154	0.156	0.161	0.166	0.171	0.178	0.185	0.192	0.2	0.207	0.216	0.224	0.232	0.24	0.248	0.253
Cote d'Ivoire	0.273	0.28	0.287	0.292	0.297	0.301	0.306	0.312	0.317	0.322	0.326	0.33	0.334	0.338	0.342	0.346	0.349	0.353	0.357	0.361	0.367	0.371	0.377	0.384	0.391	0.398	0.406	0.412
The Gambia	0.245	0.251	0.257	0.263	0.269	0.275	0.282	0.288	0.294	0.3	0.307	0.314	0.319	0.325	0.332	0.338	0.343	0.349	0.355	0.361	0.367	0.372	0.378	0.384	0.389	0.395	0.401	0.405
Ghana	0.38	0.386	0.392	0.396	0.4	0.403	0.408	0.413	0.416	0.418	0.422	0.425	0.43	0.435	0.439	0.443	0.45	0.459	0.467	0.474	0.482	0.49	0.498	0.505	0.513	0.521	0.53	0.537
Guinea	0.164	0.168	0.171	0.175	0.178	0.183	0.188	0.194	0.2	0.206	0.211	0.217	0.222	0.228	0.234	0.24	0.246	0.252	0.26	0.266	0.273	0.281	0.288	0.295	0.303	0.31	0.318	0.325
Guinea-Bissau	0.183	0.189	0.194	0.199	0.205	0.211	0.217	0.224	0.229	0.235	0.241	0.246	0.252	0.257	0.263	0.268	0.274	0.28	0.286	0.293	0.3	0.308	0.315	0.322	0.329	0.336	0.343	0.349
Liberia	0.207	0.209	0.21	0.206	0.199	0.19	0.178	0.174	0.175	0.183	0.198	0.214	0.228	0.234	0.239	0.245	0.251	0.257	0.264	0.27	0.276	0.284	0.292	0.301	0.31	0.318	0.324	0.328
Mali	0.13	0.134	0.137	0.141	0.145	0.148	0.152	0.157	0.161	0.166	0.171	0.176	0.181	0.187	0.192	0.198	0.203	0.209	0.215	0.22	0.226	0.232	0.238	0.243	0.249	0.255	0.262	0.267
Mauritania	0.299	0.307	0.314	0.321	0.328	0.335	0.341	0.346	0.351	0.356	0.359	0.363	0.367	0.371	0.377	0.382	0.39	0.398	0.406	0.412	0.42	0.427	0.435	0.442	0.45	0.458	0.465	0.471
Niger	0.092	0.097	0.101	0.105	0.108	0.111	0.113	0.115	0.118	0.121	0.123	0.125	0.127	0.13	0.134	0.137	0.14	0.144	0.147	0.151	0.156	0.16	0.165	0.17	0.176	0.181	0.186	0.191
Nigeria	0.344	0.346	0.347	0.349	0.352	0.355	0.358	0.362	0.365	0.369	0.374	0.379	0.385	0.391	0.4	0.41	0.419	0.429	0.437	0.445	0.452	0.459	0.466	0.473	0.478	0.484	0.49	0.493
Sao 1 ome and Principe	0.287	0.292	0.296	0.3	0.305	0.309	0.314	0.32	0.326	0.332	0.338	0.345	0.353	0.362	0.371	0.381	0.39	0.399	0.408	0.416	0.426	0.435	0.444	0.453	0.463	0.472	0.481	0.488

Appendix Table 10: Socio-Demographic	Index val	ues for al	estimated	GBD 201	7 locations	pendix Table 10: Socio-Demographic Index values for all estimated GBD 2017 locations, 1990-2017																						
Location	1990	1991	1992	1993	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003	2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015	2016	2017
Senegal	0.244	0.25	0.257	0.262	0.267	0.272	0.277	0.281	0.286	0.291	0.295	0.299	0.303	0.307	0.311	0.316	0.319	0.324	0.328	0.333	0.338	0.342	0.347	0.352	0.357	0.362	0.368	0.373
Sierra Leone	0.202	0.205	0.207	0.211	0.216	0.219	0.222	0.225	0.226	0.228	0.23	0.233	0.238	0.244	0.249	0.256	0.262	0.27	0.278	0.286	0.294	0.302	0.312	0.325	0.336	0.344	0.351	0.357
Togo	0.263	0.27	0.276	0.281	0.286	0.292	0.297	0.303	0.307	0.311	0.315	0.318	0.321	0.325	0.329	0.333	0.337	0.343	0.348	0.353	0.36	0.366	0.373	0.381	0.39	0.398	0.406	0.413