

Effectiveness of treatments for acute and subacute mechanical non-specific low back pain: a systematic review with network meta-analysis

Supplementary materials

Supplement A. PRISMA NMA Checklist	5
Supplement B. Difference between protocol and review	8
Supplement C. References of Included Studies	9
Supplement D. Interventions and Nodes	12
Box 1. Planned description interventions	12
Box 2. Nodes	13
Supplement E. Assessment of transitivity	15
Assessment of transitivity by trials.....	16
Table 1. Study and Patient characteristics (n=46)	16
Assessment of transitivity by interventions	21
Table 2. Stage of LBP	21
Table 3. Presence of leg pain or sciatica	22
Figure 1. Mean age.....	23
Figure 2. Percentage of male participants	23
Figure 3. Baseline severity (pain)	24
Figure 4. Length of treatment	24
Figure 5. Number of randomized	25
Table 4. Psychological assessment	26
Assessment of transitivity by head-to-head comparisons	27
Table 5. Stage of LBP	27
Table 6. Presence of leg pain or sciatica	28
Figure 6. Mean age.....	29
Figure 7. Percentage of male participants	29
Figure 8. Baseline severity (pain)	30
Supplement F. Risk of Bias	32
Figure 1. Aggregate Cochrane Risk-of-bias appraisal results	32
Table 1. Cochrane Risk-of-bias global judgement	33
Supplement G. Network Plots	35
Figure 1. Network Plot- Pain outcome	35
Figure 1a. Network for pain outcome at 1 month of FU	35
Figure 1b. Network for pain outcome at 3-6 months of FU	35

Figure 1c. Network for pain outcome at 12 months of FU.....	36
Figure 2. Network Plot- Disability outcome	37
Figure 2a. Network for disability outcome at 1 month of FU.....	37
Figure 2b. Network for disability outcome at 3-6 months of FU.....	37
Figure 2c. Network for disability outcome at 12 months of FU	38
Supplement H. Assessment of pairwise Meta-Analyses	39
Pairwise meta-analyses –Pain Outcome	39
Table 1. Pairwise meta-analyses at 1 week of FU for pain	39
Table 2. Pairwise meta-analyses at 1 month of FU for pain.....	39
Table 3. Pairwise meta-analyses at 3-6 months of FU for pain	40
Table 4. Pairwise meta-analyses at 12 months of FU for pain	40
Pairwise meta-analyses – Disability Outcome.....	41
Table 5. Pairwise meta-analyses at 1 week of FU for disability.....	41
Table 6. Pairwise meta-analyses at 1 month of FU for disability	41
Table 7. Pairwise meta-analyses at 3-6 months of FU for disability.....	42
Table 8. Pairwise meta-analyses at 12 months of FU for disability.....	42
Supplement I. Forest plot of network meta-analysis (network forest)	43
Figure 1. Network forest – pain outcome 1 week	43
Figure 2. Network forest – pain outcome 1 month.....	43
Figure 3. Network forest – pain outcome 12 months	44
Figure 4. Network forest – disability outcome 1 week.....	44
Figure 5. Network forest – disability outcome 1 month	45
Figure 6. Network forest – disability outcome 12 months.....	45
Supplement J. Incoherence estimation and evaluation	46
Table 1. Estimated Global Inconsistency in Networks.....	46
Table 2. Estimated Local Inconsistency for each pairwise comparison (side splitting) – pain outcome .	46
Table 2a. Nodesplit pain 1 week	46
Table 2b. Nodesplit pain 1 month	47
Table 2c. Nodesplit pain 12 months.....	48
Table 3. Estimated Local Inconsistency for each pairwise comparison (side splitting) – disability outcome	48
Table 3a. Nodesplit disability 1 week	48
Table 3b. Nodesplit disability 1 month.....	50
Table 3c. Nodesplit disability 12 months	50
Table 4. Strategy to explore global inconsistency – disability 1 week	51
Table 5. Strategy to explore global inconsistency – disability 1 month	52

Table 6a. Metaregression disability 1 week	53
Table 6b. Metaregression disability 1 month.....	53
Figure 1. Bubble plot disability 1 week	54
Figure 2. Bubble plot disability 1 month	55
Supplementary K. Subgroup analysis results	56
1. Subgroup meta-analysis (pharmacological and non-pharmacological)	56
Disability 1 week – non pharmacological treatments	56
Figure 1a. Network plot of non-pharmacological treatments.....	56
Figure 2a. Network forest of non-pharmacological treatments.....	56
Table 1a. Netleague of non-pharmacological treatments.....	57
Table 2a. SUCRA of non-pharmacological treatments	57
Disability 1 week – pharmacological treatments	58
Figure 1b. Network plot of pharmacological treatments	58
Figure 2b. Network forest of pharmacological treatments	58
Table 1b. Netleague of pharmacological treatments	59
Table 2b. SUCRA of pharmacological treatments.....	59
Disability 1 month – non pharmacological treatments.....	60
Figure 3a. Network plot of non-pharmacological treatments.....	60
Figure 4a. Network forest of non-pharmacological treatments.....	60
Disability 1 month – pharmacological treatments	62
Figure 3b. Network plot of pharmacological treatments	62
Figure 4b. Network forest of pharmacological treatments	62
Supplementary L. Network meta-analysis results- Interval plot	63
Figure 1. Interval Plot -Network Meta-Analyses – Pain outcome	63
Figure 1a. Interval plot all treatments against inert treatment for pain outcome at 1 month of FU	63
Figure 1b. Interval plot all treatments against inert treatment for pain outcome at 12 months of FU..	63
Figure 2. Interval Plot -Network Meta-Analyses – Disability Outcome	64
Figure 2a. Interval plot all treatments against inert treatment for disability outcome at 12 months of FU	64
Supplement M. All treatments against all treatments	65
Table 1. League table - pain	65
Table 1a. League table pain 1 month	65
Table 1b. League table pain 12 months	65
Table 2. Pain SUCRA	66
Figure 1. Cumulative ranking curve of pain 1 week	67
Figure 2. Cumulative ranking curve of pain 1 month	67

Figure 3. Cumulative ranking curve of pain 12 months.....	68
Table 3a. League table disability 12 months	69
Table 4. Disability SUCRA	70
Figure 4. Cumulative ranking curve of disability 12 months	71
Supplement N. Funnel Plot	72
Figure 1. Funnel plot-pain	72
Figure 1a. Pain Outcome 1 week.....	72
Figure 1b. Pain Outcome 1 month	73
Figure 2. Funnel plot- disability	74
Figure 2a. Disability Outcome 1 week.....	74
Figure 2b. Disability Outcome 1 month.....	75
Supplement O. Contribution matrix for the network on interventions	76
Figure 1. Contribution matrix for the network on interventions - Pain	76
Figure 1a. Contribution matrix for the network on interventions Pain Outcome 1 week	77
Figure 1b. Contribution matrix for the network on interventions Pain Outcome 1 month.....	78
Figure 1c. Contribution matrix for the network on interventions Pain Outcome 12 months.....	79
Figure 2. Contribution matrix for the network on interventions - Disability.....	80
Figure 2a. Contribution matrix for the network on interventions Disability Outcome 12 months.....	80
Supplement P. GRADE for Pain Outcome	81
1) Pain at 1 week.....	82
2) Pain at 1 month	87
3) Pain at 12 months.....	92
Supplement Q. GRADE for Disability Outcome	95
1) Disability at 12 months	96
Supplement R. Data check	99
Supplement S. References	100

Supplement A. PRISMA NMA Checklist

Section/Topic	Item #	Checklist Item	Reported on Page #
TITLE			
Title	1	Identify the report as a systematic review <i>incorporating a network meta-analysis (or related form of meta-analysis)</i> .	1
ABSTRACT			
Structured summary	2	Provide a structured summary including, as applicable: Background: main objectives Methods: data sources; study eligibility criteria, participants, and interventions; study appraisal; and <i>synthesis methods, such as network meta-analysis</i> . Results: number of studies and participants identified; summary estimates with corresponding confidence/credible intervals; <i>treatment rankings may also be discussed. Authors may choose to summarize pairwise comparisons against a chosen treatment included in their analyses for brevity.</i> Discussion/Conclusions: limitations; conclusions and implications of findings. Other: primary source of funding; systematic review registration number with registry name.	2
INTRODUCTION			
Rationale	3	Describe the rationale for the review in the context of what is already known, <i>including mention of why a network meta-analysis has been conducted</i> .	4
Objectives	4	Provide an explicit statement of questions being addressed, with reference to participants, interventions, comparisons, outcomes, and study design (PICOS).	4
METHODS			
Protocol and registration	5	Indicate whether a review protocol exists and if and where it can be accessed (e.g., Web address); and, if available, provide registration information, including registration number.	4
Eligibility criteria	6	Specify study characteristics (e.g., PICOS, length of follow-up) and report characteristics (e.g., years considered, language, publication status) used as criteria for eligibility, giving rationale. <i>Clearly describe eligible treatments included in the treatment network, and note whether any have been clustered or merged into the same node (with justification)</i> .	5
Information sources	7	Describe all information sources (e.g., databases with dates of coverage, contact with study authors to identify additional studies) in the search and date last searched.	5
Search	8	Present full electronic search strategy for at least one database, including any limits used, such that it could be repeated.	5
Study selection	9	State the process for selecting studies (i.e., screening, eligibility, included in systematic review, and, if applicable, included in the meta-analysis).	5
Data collection process	10	Describe method of data extraction from reports (e.g., piloted forms, independently, in duplicate) and any processes for obtaining and confirming data from investigators.	5-6

Data items	11	List and define all variables for which data were sought (e.g., PICOS, funding sources) and any assumptions and simplifications made.	5-6
Geometry of the network	S1	Describe methods used to explore the geometry of the treatment network under study and potential biases related to it. This should include how the evidence base has been graphically summarized for presentation, and what characteristics were compiled and used to describe the evidence base to readers.	7
Risk of bias within individual studies	12	Describe methods used for assessing risk of bias of individual studies (including specification of whether this was done at the study or outcome level), and how this information is to be used in any data synthesis.	6
Summary measures	13	State the principal summary measures (e.g., risk ratio, difference in means). <i>Also describe the use of additional summary measures assessed, such as treatment rankings and surface under the cumulative ranking curve (SUCRA) values, as well as modified approaches used to present summary findings from meta-analyses.</i>	7
Planned methods of analysis	14	Describe the methods of handling data and combining results of studies for each network meta-analysis. This should include, but not be limited to: <ul style="list-style-type: none"> • <i>Handling of multi-arm trials;</i> • <i>Selection of variance structure;</i> • <i>Selection of prior distributions in Bayesian analyses; and</i> • <i>Assessment of model fit.</i> 	7
Assessment of Inconsistency	S2	Describe the statistical methods used to evaluate the agreement of direct and indirect evidence in the treatment network(s) studied. Describe efforts taken to address its presence when found.	7
Risk of bias across studies	15	Specify any assessment of risk of bias that may affect the cumulative evidence (e.g., publication bias, selective reporting within studies).	7
Additional analyses	16	Describe methods of additional analyses if done, indicating which were pre-specified. This may include, but not be limited to, the following: <ul style="list-style-type: none"> • Sensitivity or subgroup analyses; • Meta-regression analyses; • <i>Alternative formulations of the treatment network; and</i> • <i>Use of alternative prior distributions for Bayesian analyses (if applicable).</i> 	7
RESULTS†			
Study selection	17	Give numbers of studies screened, assessed for eligibility, and included in the review, with reasons for exclusions at each stage, ideally with a flow diagram.	8
Presentation of network structure	S3	Provide a network graph of the included studies to enable visualization of the geometry of the treatment network.	10-11
Summary of network geometry	S4	Provide a brief overview of characteristics of the treatment network. This may include commentary on the abundance of trials and randomized patients for the different interventions and pairwise comparisons in the network, gaps of evidence in the treatment network, and potential biases reflected by the network structure.	10-11

Study characteristics	18	For each study, present characteristics for which data were extracted (e.g., study size, PICOS, follow-up period) and provide the citations.	8
Risk of bias within studies	19	Present data on risk of bias of each study and, if available, any outcome level assessment.	10
Results of individual studies	20	For all outcomes considered (benefits or harms), present, for each study: 1) simple summary data for each intervention group, and 2) effect estimates and confidence intervals. <i>Modified approaches may be needed to deal with information from larger networks.</i>	10-11
Synthesis of results	21	Present results of each meta-analysis done, including confidence/credible intervals. <i>In larger networks, authors may focus on comparisons versus a particular comparator (e.g. placebo or standard care), with full findings presented in an appendix. League tables and forest plots may be considered to summarize pairwise comparisons.</i> If additional summary measures were explored (such as treatment rankings), these should also be presented.	10-11-14
Exploration for inconsistency	55	Describe results from investigations of inconsistency. This may include such information as measures of model fit to compare consistency and inconsistency models, <i>P</i> values from statistical tests, or summary of inconsistency estimates from different parts of the treatment network.	10-11
Risk of bias across studies	22	Present results of any assessment of risk of bias across studies for the evidence base being studied.	10-11
Results of additional analyses	23	Give results of additional analyses, if done (e.g., sensitivity or subgroup analyses, meta-regression analyses, <i>alternative network geometries studied, alternative choice of prior distributions for Bayesian analyses, and so forth.</i>	10-11
DISCUSSION			
Summary of evidence	24	Summarize the main findings, including the strength of evidence for each main outcome; consider their relevance to key groups (e.g., healthcare providers, users, and policy-makers).	16-17
Limitations	25	Discuss limitations at study and outcome level (e.g., risk of bias), and at review level (e.g., incomplete retrieval of identified research, reporting bias). <i>Comment on the validity of the assumptions, such as transitivity and consistency. Comment on any concerns regarding network geometry (e.g., avoidance of certain comparisons).</i>	18
Conclusions	26	Provide a general interpretation of the results in the context of other evidence, and implications for future research.	18
FUNDING			
Funding	27	Describe sources of funding for the systematic review and other support (e.g., supply of data); role of funders for the systematic review. This should also include information regarding whether funding has been received from manufacturers of treatments in the network and/or whether some of the authors are content experts with professional conflicts of interest that could affect use of treatments in the network.	19

PICOS = population, intervention, comparators, outcomes, study design.

* Text in italics indicates wording specific to reporting of network meta-analyses that has been added to guidance from the PRISMA statement.

† Authors may wish to plan for use of appendices to present all relevant information in full detail for items in this section.

Supplement B. Difference between protocol and review

We extracted some important intervention details as suggested by the TIDieR checklist¹ in order to create consistent nodes, however, the poor reporting of included trials prevent the full reporting of their descriptions. We summarized some items in Table 1 of **Supplement E** (Assessment of transitivity) and full details are reported in the online repository OSF at the following link <https://osf.io/q24xh>.

We transparently edit the nodes according to the statement declaration in the published protocol². For instance, we build a new subgroup category “heat wrap” separated from “physical therapy” category. We also noted that “physical therapy” is represented only by TENS improving the homogeneity of treatment’s node. Then, we merged “Inert treatment” (e.g., placebo drug, sham therapy) and “No treatment” since only one study (Malmivaara 1995) reported no intervention in this control group described as: “the continuation of ordinary activities as tolerated.”

Supplement C. References of Included Studies

- 1 Amlie, E., Weber, H. & Holme, I. Treatment of acute low-back pain with piroxicam: results of a double-blind placebo-controlled trial. *Spine* 12, 473-476 (1987).
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- 17 Jellema, P., van der Windt, D.A., van der Horst, H.E., Twisk, J.W., Stalman, W.A. & Bouter, L.M. Should treatment of (sub)acute low back pain be aimed at psychosocial prognostic factors? Cluster randomised clinical trial in general practice. *BMJ (clinical research ed.)* 331, 84 (2005).

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- 46 Younes, M., Nowakowski, K., Didier-Laurent, B., Gombert, M. & Cottin, F. Effect of spinal manipulative treatment on cardiovascular autonomic control in patients with acute low back pain. *Chiropractic & manual therapies* 25, 33 (2017).

Supplement D. Interventions and Nodes

Box 1. Planned description interventions

Class	Example of individual treatments
Pharmacological	
Antidepressant drugs	Any kind of SSRI/SNRI or tricyclic drug
Muscle relaxant drugs	Any kind of skeletal muscle relaxant drug (e.g. flupirtin, orphenadrine, dantrolene, carisoprodol, tizanidine, incobotulinumtoxinA, cyclobenzaprine, metaxalone, baclofen, methocarbamol, chlorzoxazone)
Non-steroidal anti-inflammatory drugs (NSAIDs)	Any kind of NSAIDs drug, including COX-2 inhibitors (e.g. ibuprofen, naproxen, sulindac, ketoprofen, tolmetin, etodolac, fenoprofen, diclofenac, flurbiprofen, piroxicam, ketorolac, indomethacin, meloxicam, nabumetone, oxaprozin, mefenamic acid, diflunisal)
Opioid drugs	Any kind of strong or weak opioid analgesics (e.g. morphine, hydromorphone, oxycodone, fentanyl, methadone, buprenorphine, diamorphine, tapentadol, codeine, hydrocodone, tramadol, pentazocine, tilidine)
Paracetamol	
Steroids	Any kind of steroid drug (e.g. dexamethasone, methylprednisolone, prednisone)
Non-pharmacological treatments	
Acupuncture and dry needling	
Biopsychosocial rehabilitation	Any kind of cognitive behavioral treatment, multidisciplinary biopsychological rehabilitation and back school
Education	Any kind of advice to stay active, booklet, reassurance, ergonomics, workplace intervention, pain education (neurobiology and neurophysiology of pain)
Exercise	Any kind of exercise (aerobic or resistance training) single supervised or home exercise, including stretching and McKenzie therapy
Manual therapy	Any kind of mobilization or spinal manipulation (high velocity thrust techniques at or near to the end of the range of motion or low-grade velocity movements within the range of motion), myofascial therapy/trigger point, soft tissue massage
Physical Therapy	Any physical therapy (low-laser therapy, diathermy, transcutaneous electrical nerve stimulation, ultrasound therapy, heat wrap)
Taping	Kinesiotaping
Usual care	Any kind of treatment suggested by general medicine (minimal intervention: advice to stay active or to take drugs as needed)
Inert treatment	Any kind of sham or placebo therapy
No treatment	No treatment, waiting list control

Box 2. Nodes

Treatments	Nodes	Evidence and assumptions
Muscle relaxant drugs (Baclofen, Carisoprodol, Dantrolene, Tizanidine Thiocolchicoside)	Muscle relaxant	Separate assessment for muscle relaxants and for Benzodiazepines ³ . A metanalysis shown similar effects across muscle relaxant drugs versus placebo, $I^2=55\%$ ⁴ .
Non-Steroidal Anti-Inflammatory Drugs (NSAIDs), including COX-2 inhibitors (diclofenac, diflunisal, ibuprofen, indomethacin, loxoprofen, piroxicam, tenoxicam)	NSAIDs	Separate assessment for all NSAIDs ³ . No clear difference in short-term pain reduction when comparing selective COX-2 inhibitors to non-selective NSAIDs ⁵ .
Opioid analgesics (meptazinol)	Opioids	Separate assessment for opioids ³ . Inclusion criteria of SR: morphine, diamorphine, fentanyl, alfentanil, remifentanil, methadone, oxycodone, pethidine, tapentadol, tramadol, codeine, dihydrocodeine, meptazinol ⁶ . Inclusion criteria of SR: various opioid analgesics ⁷ .
Paracetamol	Paracetamol	Separate assessment for paracetamol ^{3,8} .
Steroids drugs (dexamethasone, methylprednisolone, prednisone)	Steroids	Separate assessment for steroids ³ . Systematic reviews found no evidence to suggest that a series of epidural injections was any more effective than a single injection (see Appendix 1 Table 3). Individual RCTs found no evidence of improvement in steroid benefits with increasing dose (see Appendix 1 Table 4) ⁹ . Individual RCTs found no consistent evidence of superior efficacy of one steroid over the others (see Appendix 1 Table 4) ⁹ . A meta-analysis included all type of steroids. ¹⁰
Acupuncture Cognitive behavioural treatment/multidisciplinary biopsychological rehabilitation (MBR) with or without exercise	Acupuncture Cognitive behavioral therapy	Inclusion criteria of Cochrane review, MBR program: the intervention included a physical component (e.g., pharmacological, physical therapy, exercise) in combination with either a psychological, social, or occupational component (or any combination of these) ¹¹ .
Back school Booklet, Information, ergonomics, any kind of advice, workplace intervention, pain education	Back school * Education	Findings suggest positive effects for education even if differ in terms of its contents such as health education, self-management, video education, and postural education ¹² .

		Many different types of patient education are widely used ¹³ .
McKenzie Any kind of exercise (aerobic or resistance training)	Exercise	No superior type of physical exercise for people with chronic non-specific neck pain ¹⁴ . Various exercise training approaches are effective ¹⁵ .
Stretching Spinal manipulation	Manual therapy	Inclusion criteria of SR: Studies investigating manual therapy using HVLA or non-HVLA techniques such as: joint mobilization, soft tissue focused techniques, myofascial release, longitudinal sliding, soft tissue mobilizations, deep-pressure massage, muscle energy, massage, hold relaxation technique, ischemic compression, and functional/fascial technique. therapy technique(s) ¹⁶ . Different forms of manual therapy did not lead to different outcomes in older persons with chronic LBP ¹⁷ .
Manual therapy (mobilization) Trigger point/myofascial therapy/massage		
Heat wrap	Heat wrap**	
TENS	Physical therapy	
Usual care or minimal treatment (general prescription such as drugs as needed, advice stay active)	Usual care	Usual care is a term used to describe the full spectrum of patient care practices in which clinicians have the opportunity (which is not necessarily seized) to individualize care ¹⁸ . Treatment reported: education and reassurance, exercise, bed rest, return to work ¹⁹ .
Sham therapy Placebo therapy No treatment	Inert treatment	

* This node was assessed only in the qualitative synthesis because of insufficient data (e.g., not reported outcome data)

**According to the protocol ² since we obtained a sufficient number of studies sharing the same description of the intervention, we created a new node (heat wrap) separated from the physical therapy node.

Supplement E. Assessment of transitivity

Before conducting the statistical analysis, we assessed whether the trials included in the NMA were on average similar in terms of characteristics that might modify the treatment effect (so that the transitivity assumption is plausible). Indirect comparisons, in contrast to direct comparisons, are not protected by randomisation and may be confounded by differences between the trials. In our analysis we deemed the following parameters as possible confounders²⁰ which were displayed as cumulative frequencies, boxplots or bar charts when appropriate: stage of NS-LBP, presence of leg pain or sciatica, mean age, percentage of male participants, baseline severity, length of treatment, number of randomized, psychological assessment. The plausibility of the transitivity assumption was evaluated by comparing the distribution of these potential effect modifiers across trials, interventions and head-to-head comparisons

Assessment of transitivity by trials

Table 1. Study and Patient characteristics (n=46)

ID	Author	Year	Setting	Stage of LBP	Presence of leg pain or sciatica	Length of treatment	Outcomes	Week of FU	Sample size	Treatments	Nodes	Age mean	Age variance (SD)	% of male
1	Amlie*	1987	Multi-center	Acute LBP (less than 6 weeks)	Not stated	1 week	Pain; disability	3 days; 7 days	282	1. Piroxicam 2. Placebo	NSAIDs Inert treatment	37,3 38,5	NA	58,6 59,2
2	Bergquist-ullman*	1977	Single center	Mixed LBP (less than 12 weeks)	Yes	2 weeks Max 10 trt	Pain; disability	10 days; 3 weeks; 6 weeks	145	1. Back school 2. Placebo	Back school Inert treatment	NA	NA	91,4 86,7
3	Berry	1988	Single center	Acute LBP (less than 6 weeks)	Yes	1 week	Pain	1 week	112	1. Tizanidine 2. Placebo	Muscle relaxant Inert treatment	44 38	13 13	51 50,9
4	Bertalanffy	2005	Single center	Acute LBP (less than 6 weeks)	No	1 day	Pain	30 minutes	63	1. TENS 2. Sham TENS	Physical therapy Inert treatment	47 49	7 14	53,3 51,5
5	Casale*	1988	Single center	Acute LBP (less than 6 weeks)	Not stated	4 days	Pain	Day 4	20	1. Dantrolene sodium 2. Placebo	Muscle relaxant Inert treatment	46,7 47,1	2,3 2,2	70 80
6	Cherkin*	1996	Single center	Mixed LBP (less than 12 weeks)	Yes	1 session	Pain; disability	1 week	299	1. Nurse education 2. Booklet 3. Usual care	Education Education Usual care	40,8 44,1 43,0	NA	57 49 51
7	Cherkin**	1998	Multi-center	Mixed LBP (less than 12 weeks)	No	1 month	Pain; disability	4 weeks; 12 weeks; 12 months	321	1. McKenzie 2. Manipulation 3. Booklet	Exercise Manual therapy Education	41,8 39,7 40,1	11,5 9,4 11,2	53 47 58
8	Dapas*	1985	Multi-center	Acute LBP (less than 6 weeks)	Not stated	14 days	Pain; disability	Day 4; Day 10	123	1. Baclofen 2. Placebo	Muscle relaxant Inert treatment	42,7 41,8	NA	52 44
9	Dreiser	2003	Multi-center	Acute LBP (less than 6 weeks)	No	1 week	Pain; disability	Day 3; day 8	372	1. Diclofenac-K 2. Ibuprofen 3. Placebo	NSAIDs NSAIDs Inert treatment	40,9 40,6 41	10,9 11,6	48,4 52,5 47,2

													11,3		
10	Eken*	2014	Silgle center	Acute LBP (less than 6 weeks)	No	1 day	Pain	30 minutes	137	1. Paracetamol 2. Dexketoprofen 3. Morphine	Paracetamol NSAIDs Opioid	31,5*	9,5*	60,6*	
11	Eskin*	2014	Single center	Acute LBP (less than 6 weeks)	Not stated	5 days	Pain	Day 5-7	79	1. Prednisone 2. Placebo	Steroids Inert treatment	39 41	8 9	67 73	
12	Faas*	1995	Multi-center	Acute LBP (less than 6 weeks)	Yes	5 weeks	Pain	1 week; 1 month; 12 month	363	1. Exercise 1. Usual care 2. Sham ultrasound	Exercise Usual care Inert treatment	35 34 37	NA	62 71 66	
13	Goldie*	1968	Single center	Acute LBP (less than 6 weeks)	Yes	14 days	Pain	1 week; 2 weeks	50	1. Indomethacin 2. Placebo	NSAIDs Placebo	NA	NA	52 52	
14	Haimovic*	1986	Single center	Acute LBP (less than 6 weeks)	Yes	7 days	Pain	1 week; 12 months	33	1. Dexamethasone 2. Placebo	Steroids Inert treatment	NA	NA	NA	
15	Hasegawa	2014	Single center	Acute LBP (less than 6 weeks)	No	1 week	Pain; disability	7 days; 28 days	80	1. Acupuncture 2. Sham acupuncture	Acupunctre Inert treatment	47 43,9	9,8 10,9	37,5 35	
16	Hindle*	1972	Single center	Acute LBP (less than 6 weeks)	Not stated	4 days	Pain; disability	2 days; 4 days	32	1. Carisoprodol 2. Placebo	Muscle relaxant Inert treatment	37 43,5	NA NA	56 62	
17	Jellema	2005	Multi-center	Mixed LBP (less than 12 weeks)	Not stated	5 days	Pain; disability	6, 26, 52 weeks	314	1. Behavioral therapy 2. Usual care	Cognitive behavioral therapy Usual care	43,4 42	11,1 12	52,4 52,6	
18	Ketenci	2005	Single center	Acute LBP (less than 6 weeks)	Not stated	1 week	Pain	Day 5-7	97	1. Thiocolchicoside 2. Tizanidine 3. Placebo	Muscle relaxant Muscle relaxant Inert treatment	37 37 40	NA NA NA	57,9 37,5 48,1	
19	Kettenmann*	2007	Single center	Mixed LBP (less than 12 weeks)	Not stated	4 days	Pain	Day 4	30	1. Heat wrap 2. Usual care	Heatwrap Usual care	56,2 57,9	14,9 11,7	46,7 25	
20	Lindstrom	1995	Single center	Subacute LBP (6-12 weeks)	Not stated	Until recovery	Pain; disability	12 months	103	1. Cognitive behavioral therapy 2. Usual care	Cognitive behavioral therapy Usual care	39,4 42,4	10,7 10,9	76,5 61,5	

21	Malmivaara	1995	Multi-center	Acute LBP (less than 6 weeks)	Yes	Not reported	Pain; disability	3 weeks; 12 weeks	119	1. Exercise 2. No treatment	Exercise Inert treatment	41,1 39,1	NA NA	29 30
22	Mayer	2005	Multi-center	Mixed (acute and subacute)	No	5 days	Pain; disability	1 week	76	1. Heat wrap 2. Exercise 3. Booklet	Heat wrap Exercise Education	29,3 32,6 31,3	9,9 10,3 10,9	32 40 7,7
23	Miki	2018	Single center	Acute LBP (less than 6 weeks)	No	4 weeks	Pain; disability	2 weeks, 1 month	127	1. Acetaminophen 2. Loxoprofen	Paracetamol NSAIDs	66,7 63,5	2,3 19,4	32,8 34,9
24	Nadler**	2002	Multi-center	Mixed (acute and subacute)	No	2 days or 1 day??	Pain; disability	4 days	371	1. Heat wrap 2. Acetaminophen 3. Ibuprofen 4. Unheated wrap 5. Oral placebo	Heat wrap Paracetamol NSAIDs Inert treatment Inert treatment	35,8 34,9 36,6 36,8 38,0	10,5 11,3 10,4 9,3 9,1	41,6 43,4 40,6 42,1 40
25	Nadler**	2003b	Multi-center	Mixed (acute and subacute)	No	3 days	Pain; disability	Days 2-4	76	1. Heat wrap 2. Oral placebo 3. Ibuprofen 4. Unheated wrap	Heat wrap Inert treatment NSAIDs	42,2 41,5 42,5 34,0	9,4 9,8 2,7 8,4	36,4 38,2 25 20
26	Nadler**	2003 a	Multi-center	Mixed (acute and subacute)	No	3 days	Pain; disability	Day 5	219	1. Heat wrap 2. Oral placebo 3. Ibuprofen 4. Unheated wrap	Heat wrap Inert treatment NSAIDs Inert treatment	35,6 36,7 36,3 34,9	11,6 10,8 11,6 11,3	45,7
27 a	Postacchini *	1988	Multi-center	Acute LBP (less than 6 weeks)	No	4 weeks 10-14 days 1 or 2 weeks	Pain; disability	3 weeks; 6 months	46	1. Manipulation 2. Diclofenac 3. Placebo gel	Manual therapy NSAIDs Inert treatment	36,3	NA	55
27 b	Postacchini *	1988	Multi-center	Acute LBP (less than 6 weeks)	No	4 weeks 10-14 days 1 week 1 or 2 weeks	Pain; disability	3 weeks; 6 months	66	1. Manipulation 2. Diclofenac 3. Back school 4. Placebo gel	Manual therapy NSAIDs Back school Inert treatment	40,3	NA	51,2
27 c	Postacchini *	1988	Multi-center	Acute LBP (less than 6 weeks)	Yes	4 weeks 10-14 days 1 or 2 weeks	Pain; disability	3 weeks; 6 months	53	1. Manipulation 2. Diclofenac 3. Placebo gel	Manual therapy NSAIDs Inert treatment	37,7	NA	45,8

28	Ralph*	2008	Multi-center	Acute LBP (less than 6 weeks)	No	7 days	Pain; disability	1 week	562	1. Carisoprodol 2. Placebo	Muscle relaxant Inert treatment	39,3 41,5	11,82 11,7	51,3 45
29	Sae-Jung	2016	Single center	Mixed (acute and subacute)	No	2 weeks	Pain; disability	1 month; 3 months	65	1. Diclofenac 2. Methylprednisolone	NSAIDs Steroids	49 44	8,7 9,3	55 53,1
30	Santilli	2006	Multi-center	Acute LBP (less than 6 weeks)	Yes	Until recovery (max 4 weeks)	Pain	15 days; 1, 3, 6 months	102	1. Active manipulation 2. Simulated manipulation	Manual therapy Inert treatment	NA NA	NA NA	69,8 55,1
31	Schrenk	2003	Single center	Mixed (acute and subacute)	Yes	Not reported	Pain; disability	3 visits	25	1. Exercise (McKenzie) 2. Mobilization	Exercise Manual therapy	40,1 44,8	17,1 12,7	46,7 80
32	Schneider	2015	Single center	Mixed (acute and subacute)	No	4 weeks	Pain; disability	4 weeks; 3 months; 6 months	112	1. Manual manipulation 2. Mechanical assisted manipulation 3. Usual care	Manual therapy Manual therapy Usual care	41,4 40,4 41,3	15,3 15,9 11,6	32,4 40 40
33	Seferlis	1998	Single center	Acute LBP (less than 6 weeks)	Yes	8 weeks	Pain; disability	1 months; 3 months; 12 months	180	1. Exercise 2. General practitioner program-usual care	Exercise Usual care	39	19-64 range	52,7
34	Serfer*	2009	Multi-center	Acute LBP (less than 6 weeks)	No	1 week	Pain; disability	1 week	828	1. Carisoprodol 250 mg 2. Carisoprodol 350 mg 3. Placebo	Muscle relaxant Muscle relaxant Inert treatment	40,9 40,5 40,7	11,7 12,4 13,1	47,7 44,3 39,4
35	Shin	2013	Multi-center	Acute LBP (less than 6 weeks)	Yes	1 day	Pain; disability	2 weeks; 4 weeks; 24 weeks	58	1. Acupuncture 2. Diclofenac	Acupuncture NSAIDs	37,9 38,7	7,4 8,6	66 52
36	Storheim	2003	Single center	Subacute LBP (6-12 weeks)	No	15 weeks 1 week	Pain; disability	18 weeks; 48 weeks	93	1. Exercise 2. Cognitive intervention 3. Usual care	Exercise Cognitive behavioral therapy Usual care	42,3 41,3 38,9	9,2 9,4 11,9	46,7 52,9 44,8
37	Suni*	2006	Multi-center	Mixed (acute and subacute)	Not stated	12 months	Pain; disability	6 months; 12 months	106	1. Exercise with cognitive goals 2. Control group	Cognitive behavioral therapy Usual care	47,6 46,9	5,8 5,3	100 100
38	Szpalski	1994	Single center	Acute LBP (less than 6 weeks)	Yes	1-2 weeks	Pain	8 days; 15 days	73	1. Tenoxicam 2. Placebo	NSAIDs Inert treatment	37,5 38,9	9,2 10,4	62,2 66,7

39	Takamoto	2015	Multi-center	Acute LBP (less than 6 weeks)	No	2 weeks	Pain; disability	1 week; 1 month	63	1. Compression at TP 2. Sham compression 3. Effleurage massage	Manual therapy Inert treatment Manual therapy	38 38,1 35,6	3 3,8 3	45,4 47,1 37,5
40	Traeger	2019	Multi-center	Acute LBP (less than 6 weeks)	Yes	2 sessions	Pain; disability	1 week, 6, 12 months	202	1. Education 2. Sham education	Education Inert treatment	46,5 43,8	14,7 14,1	47,5 50,5
41	Tuzun	2003	Multi-center	Acute LBP (less than 6 weeks)	Not stated	Until recovery, max 5 days	Pain	5 days	149	1.Thiocolchicoside 2. Placebo	Muscle relaxant Inert treatment	40,7 41	10,3 11	50 42
42	Veenema	2000	Single center	Acute LBP (less than 6 weeks)	Not stated	1 day	Pain	60 minutes	155	1. Meperidine 2. Ketorolac	Opioid NSAIDs	35,5 36,0	12,8 12,1	63,0 60,0
43	Videman*	1984	Single center	Acute LBP (less than 6 weeks)	No	Until recovery, max 3 weeks	Pain; disability	1 week; 3 weeks	70	1. Meptazinol 2. Diflunisal	NSAIDs Opioid	38,0 35,0	14,0 11,0	60,0 57,1
44	von Heymann**	2013	Multi-center	Acute LBP (less than 6 weeks)	Not stated	Not reported	Pain; disability	9 days	100	1. Manipulation 2. Diclofenac 3. Placebo-sham	Manual therapy NSAIDs Inert treatment	34,1* 37,5* 39,3*	9,5 10,9 10,2	63,9 10,9 10,2
45	Williams	2014	Multi-center	Acute LBP (less than 6 weeks)	Yes	Until recovery, max 4 weeks	Pain; disability	1week; 1 month; 3 months;	165 2	1. Paracetamol 2. Paracetamol as needed 3. Placebo	Paracetamol Paracetamol Inert treatment	44,1 45,5 45,4	14,8 16,7 15,9	52,0 53,0 55,0
46	Younes*	2017	Single center	Mixed (acute and subacute)	Not stated	1 week	Pain	1 week	22	1. Manipulation 2. Sham manipulation	Manual therapy Inert treatment	31,0 28,0	9,0 7,0	100,0 100,0 100,0

*studies were not included in quantitative analysis due to different reasons such as median and IQR, missing outcome data.

**not all treatment arms are reported in quantitative analysis (e.g., multi-arm trial reported 2 out 3 treatment arms with available outcome data).

Assessment of transitivity by interventions

Table 2. Stage of LBP

TREATMENT	FREQUENCIES (%)		
	Acute	Subacute	Mixed
A	76,5	0,0	23,5
B	100,0	0,0	0,0
C	50,0	0,0	50,0
D	0,0	50,0	50,0
E	20,0	0,0	80,0
F	42,9	14,3	42,9
G	0,0	0,0	100,0
H	58,3	0,0	41,7
I	100,0	0,0	0,0
J	77,8	0,0	22,2
K	100,0	0,0	0,0
L	80,0	0,0	20,0
M	100,0	0,0	0,0
N	66,7	0,0	33,3
O	22,2	22,2	55,6

Legend: A=Inert treatment; B=Acupuncture; C=Back school; D=Cognitive behavioral therapy; E=Education; F=Exercise; G=Heat wrap; H=Manual therapy; I=Muscle relaxant; J=NSAIDs; K=Opioids; L=Paracetamol; M=Physical therapy; N=Steroids; O=Usual care

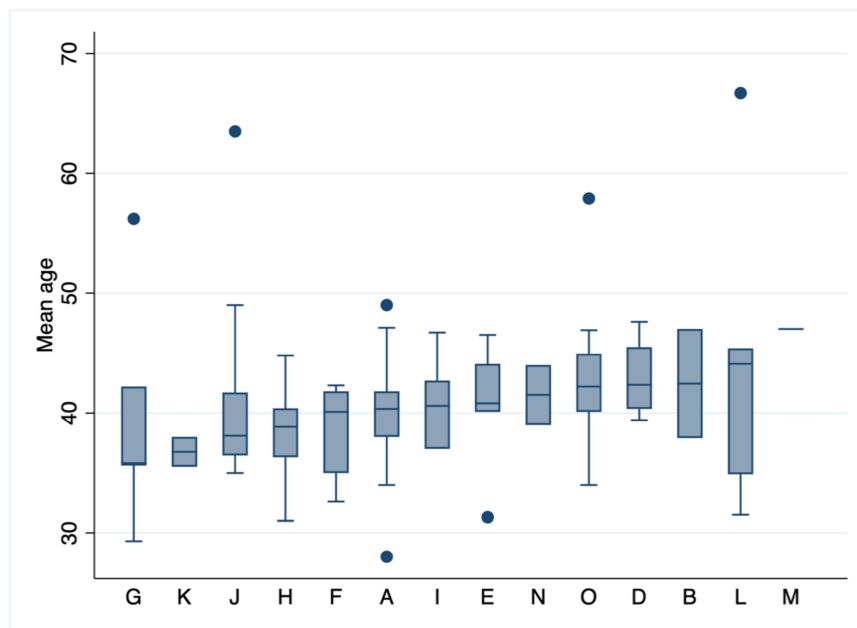
Table 3. Presence of leg pain or sciatica

TREATMENT	FREQUENCIES (%)		
	Yes	No	Not stated
A	32,4	41,2	26,5
B	50,0	50,0	0,0
C	50,0	50,0	0,0
D	0,0	25,0	75,0
E	60,0	40,0	0,0
F	57,1	42,9	0,0
G	0,0	80,0	20,0
H	25,0	58,3	16,7
I	10,0	30,0	60,0
J	22,2	61,1	16,7
K	0,0	66,7	33,3
L	40,0	60,0	0,0
M	0,0	100,0	0,0
N	33,3	33,3	33,3
O	33,3	22,2	44,4

Legend: A=Inert treatment; B=Acupuncture; C=Back school; D=Cognitive behavioral therapy; E=Education; F=Exercise; G=Heat wrap; H=Manual therapy; I=Muscle relaxant; J=NSAIDs; K=Opioids; L=Paracetamol; M=Physical therapy; N=Steroids; O=Usual care

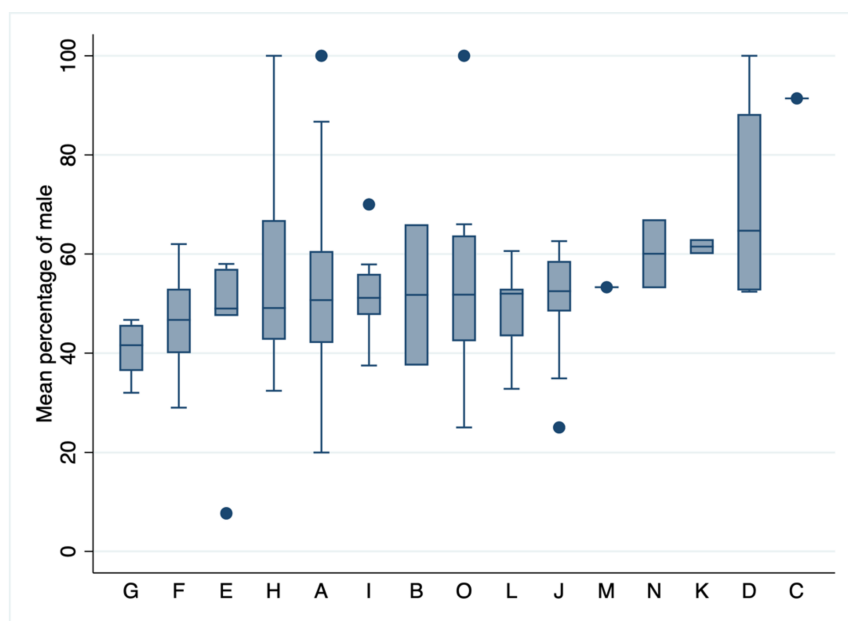
*Presence of leg pain or sciatica was reported in 15 studies out of 46 (31%) of which 6 were not included in quantitative analysis (qualitative analysis).

*Leg pain or sciatica is present in 32% (median, IQR 5-45%) of studies whereas 17% of studies did not report information (median, 0-33%).



Median age ranged from 35 to 48 years old with overlapping of 25-75 percentiles across interventions as already known by the Global Burden of Disease.²¹

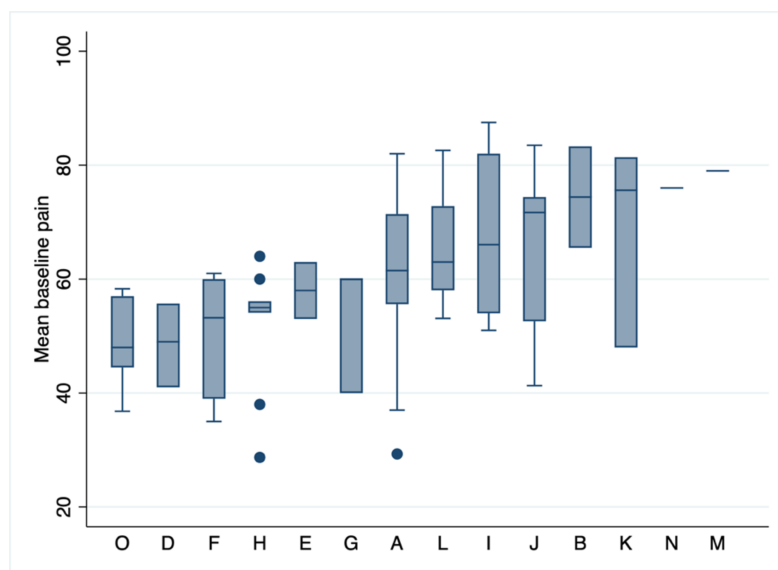
Figure 1. Mean age



Median percentage of male ranged from 42% to 90% with overlapping of 25-75 percentiles across interventions.*

Figure 2. Percentage of male participants

*Five studies did not report gender; outliers refers to 2 studies with a 100% male; however, these trials did not report outcome data and were not included in quantitative analysis (qualitative analysis). Excluding them, male and female can be equally distributed across interventions.



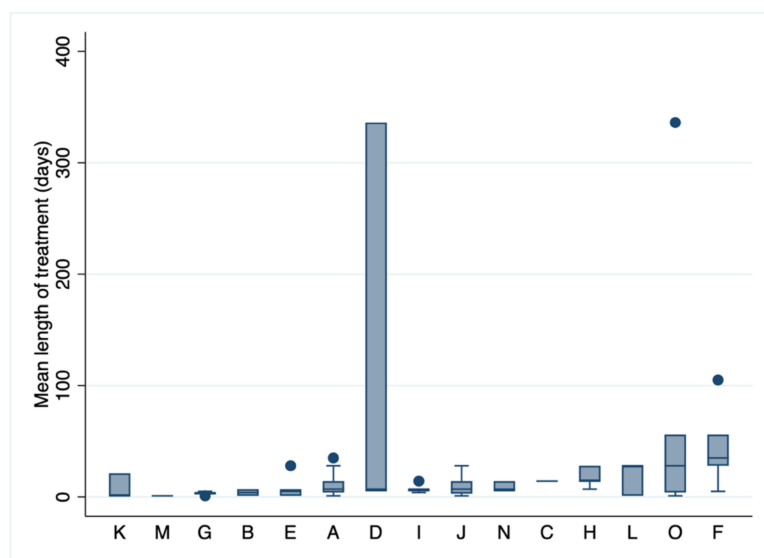
Median baseline pain ranged from 37 to 78 with overlapping of 25-75 percentiles across interventions.

Figure 3. Baseline severity (pain)

*A: 1 trial out of 34 had an outlier mean baseline value of 29.3, however this trial did not report outcome data and was not included in quantitative analysis (qualitative analysis);

**H: 1 trial out of 12 had an outlier mean baseline value of 28.7, however this trial did not report outcome data and was not included in quantitative analysis (qualitative analysis)

Severity of pain based on adapted scale 0-100



Median length of treatment ranged from 1 to 40 days with overlapping of 25-75 percentiles across interventions.

Figure 4. Length of treatment

*D: 1 trial out of 4 had an outlier mean length of treatment of 336 days, however this trial did not report outcome data and was not included in quantitative analysis (qualitative analysis); **O: 1 trial had out of 9 an outlier median length of treatment of 336 days, however this trial did not report outcome data and was not included in quantitative analysis (qualitative analysis)

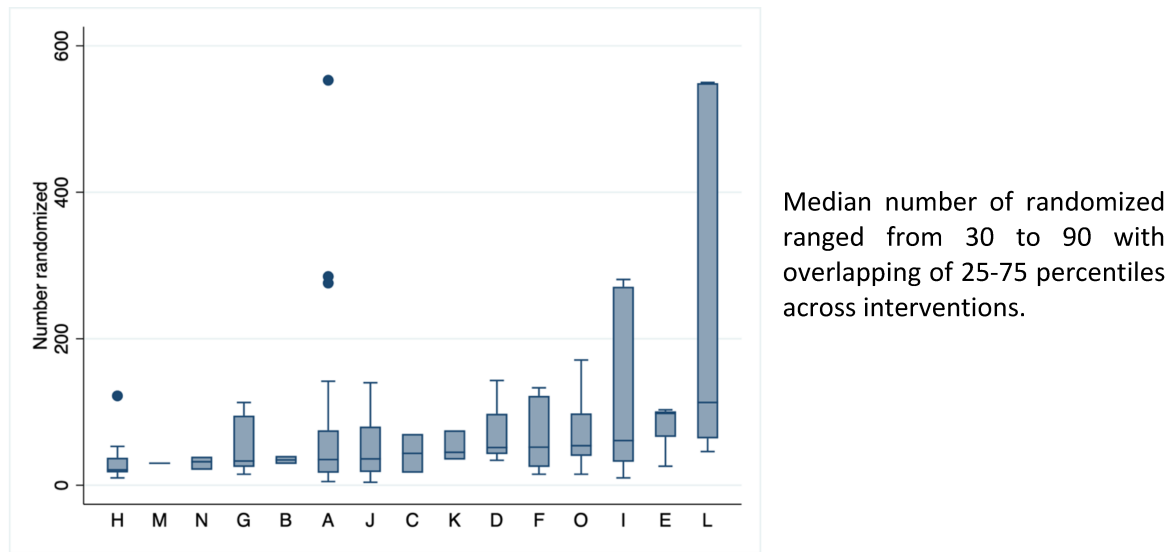


Figure 5. Number of randomized

*L: 1 trial out of 4 had an outlier number of randomized of 550, which represents less than 5% of the overall sample. However, we judged this reason insufficient to affect transitivity across interventions.

**A: 1 trial out of 34 had an outlier number of randomized of 545, which represents less than 5% of the overall sample. However, we judged this reason insufficient to affect transitivity across interventions.

Legend: A=Inert treatment; B=Acupuncture; C=Back school; D=Cognitive behavioral therapy; E=Education; F=Exercise; G=Heat wrap; H=Manual therapy; I=Muscle relaxant; J=NSAIDs; K=Opioids; L=Paracetamol; M=Physical therapy; N=Steroids; O=Usual care

Table 4. Psychological assessment

Overall, 10 RCTs (22%) reported a psychological assessment as baseline characteristics of samples. We found heterogeneity and poor reporting in outcome measurements with missing data; thus, we did not explore the heterogeneity across all included studies. We reported the psychological assessment in a table format.

PSYCHOLOGICAL ASSESSMENT				
ID	Author	Category of Intervention	Scores at baseline	Mean (SD)
4	Bertalanffy 2005	Physical therapy	Anxiety score ^a	82,0 (8,0)
4	Bertalanffy 2005	Inert treatment	Anxiety score ^a	85,0 (6,0)
6	Cherkin 1996	Education	Worry about pain ^b	6,0
6	Cherkin 1996	Education	Worry about pain ^b	6,0
6	Cherkin 1996	Usual care	Worry about pain ^b	5,7
12	Faas 1995	Usual care	NHP (emotion) ^c	7,4
12	Faas 1995	Inert treatment	NHP (emotion) ^c	7,2
12	Faas 1995	Exercise	NHP (emotion) ^c	7,7
16	Hindle 1972	Muscle relaxant	Anxiety and tension ^d	2,6
16	Hindle 1972	Inert treatment	Anxiety and tension ^d	2,2
17	Jellema 2005	Cognitive behavioral therapy	FABQpa ^e	14,3 (5,6)
			CSQ ^f	10,3 (6,6)
17	Jellema 2005	Usual care	FABQpa ^e	15,3 (5,2)
			CSQ ^f	11,2 (6,9)
23	Miki 2018	Paracetamol	PCS ^g	24,5 (1,5)
23	Miki 2018	NSAIDs	PCS ^g	30,7 (1,7)
32	Schneider 2015	Manual therapy	FABQ ^h	32,7 (15,3)
32	Schneider 2015	Manual therapy	FABQ ^h	33,0 (18,6)
32	Schneider 2015	Usual care	FABQ ^h	33,0 (17,8)
36	Storheim 2003	Exercise	FABQpa ^e	13,3 (5,2)
			FABQw ⁱ	25,9 (9,7)
36	Storheim 2003	Cognitive behavioral therapy	FABQpa ^e	14,1 (4,4)
			FABQw ⁱ	26,7 (9,1)
36	Storheim 2003	Usual care	FABQpa ^e	14,6 (3,8)
			FABQw ⁱ	29,1 (8,2)
40	Traeger 2019	Education	PCS ^g	18,3 (12)
			DASS ^j	4,1 (3,7)
40	Traeger 2019	Inert treatment	PCS ^g	19,9 (11,2)
			DASS ^j	5,1 (5)
45	Williams 2014	Paracetamol	Feelings of depression ^k	3,2 (2,9)
45	Williams 2014	Paracetamol	Feelings of depression ^k	3,1 (2,9)
45	Williams 2014	Inert treatment	Feelings of depression ^k	3,1 (2,9)

^a Visual analogue scale from 0 (no anxiety) to 100 (highest anxiety)

^b Numeric rating scale from 0 (no worry) to 10 (extremely worried)

^c NHP: Nottingham Health Profile – emotional reactions domains from 0 (good subjective health status) to 100 (poor subjective health status)

^d Four step severity rating scale from 1 (none) to 4 (severe)

^e FABQpa: Fear-avoidance belief questionnaire - four item physical activity subscale from 0 to 24, with higher score indicating more strongly held fear avoidance beliefs

^f CSQ: Coping strategies questionnaire - six item subscale from 0 to 36, with higher scores indicating greater use of coping strategies

^g PCS: Pain catastrophizing scale from 0 to 52, with higher scores indicating higher levels of catastrophizing

^h FABQ: Fear-avoidance belief questionnaire from 0 to 96, with higher score indicating more strongly held fear avoidance beliefs

ⁱ FABQw: Fear-avoidance belief questionnaire - seven item physical activity subscale from 0 to 42, with higher score indicating more strongly held fear avoidance beliefs

^j DASS: Depression severity scale of Depression, Anxiety and Stress Scale with range from 0 (no depressive symptoms) to 42 (high depressive symptoms)

^k Feelings of depression from 0 (not at all) to 10 (extremely).

Assessment of transitivity by head-to-head comparisons

Table 5. Stage of LBP

COMPARISONS	FREQUENCIES (%)		
	Acute	Subacute*	Mixed (acute and subacute)
AB	100,0	0,0	0,0
AC	50,0	0,0	50,0
AE	100,0	0,0	0,0
AF	100,0	0,0	0,0
AG	0,0	0,0	100,0
AH	85,7	0,0	14,3
AI	100,0	0,0	0,0
AJ	72,7	0,0	27,3
AL	50,0	0,0	50,0
AM	100,0	0,0	0,0
AN	100,0	0,0	0,0
AO	100,0	0,0	0,0
BJ	100,0	0,0	0,0
CH	100,0	0,0	0,0
CJ	100,0	0,0	0,0
DF	0,0	100,0	0,0
DO	0,0	50,0	50,0
EF	0,0	0,0	100,0
EG	0,0	0,0	100,0
EH	0,0	0,0	100,0
EO	0,0	0,0	100,0
FG	0,0	0,0	100,0
FH	0,0	0,0	100,0
FO	66,7	33,3	0,0
GJ	0,0	0,0	100,0
GL	0,0	0,0	100,0
GO	0,0	0,0	100,0
HJ	100,0	0,0	0,0
HO	0,0	0,0	100,0
JK	100,0	0,0	0,0
JL	66,7	0,0	33,3
JN	0,0	0,0	100,0
KK	100,0	0,0	0,0

Legend: A=Inert treatment; B=Acupuncture; C=Back school; D=Cognitive behavioral therapy; E=Education; F=Exercise; G=Heat wrap; H=Manual therapy; I=Muscle relaxant; J=NSAIDs; K=Opioids; L=Paracetamol; M=Physical therapy; N=Steroids; O=Usual care

*only 3 comparisons investigated subacute population:

DO: 50% was due to 2 studies (Lindstrom 1995 and Storheim 2003)

DF: 100% was due to 1 study (Storheim 2003)

FO: 33% was due to 1 study (Storheim 2003)

Generally, covariates were equally distributed across comparisons except for a very little percentage of comparisons (0.09%) represented by subacute population.

Moreover, these comparisons are present only in medium and long-terms of follow-ups:

- For both pain and disability at medium term no NMA was performed due to a disconnected network;
- For pain at long term, subacute population is present in 1 out of 4 head-to head comparisons;
- For disability at long term, subacute population is present in 3 out 5 head-to head comparisons.

Moreover, there is no consensus on the time-contingent traditional classification (acute, subacute, chronic) because this classification does not adequately reflect the prognostically highly important process of chronification²².

For all these reasons, stage of pain can not be considered a potential effect modifier.

Table 6. Presence of leg pain or sciatica

COMPARISONS	FREQUENCIES (%)		
	Yes*	No	Not stated
AB	0,0	100,0	0,0
AC	50,0	50,0	0,0
AE	100,0	0,0	0,0
AF	100,0	0,0	0,0
AG	0,0	100,0	0,0
AH	28,6	42,9	28,6
AI	12,5	25,0	62,5
AJ	27,3	54,6	18,2
AL	50,0	50,0	0,0
AM	0,0	100,0	0,0
AN	50,0	0,0	50,0
AO	100,0	0,0	0,0
BJ	100,0	0,0	0,0
CH	0,0	100,0	0,0
CJ	0,0	100,0	0,0
DF	0,0	100,0	0,0
DO	0,0	25,0	75,0
EF	0,0	100,0	0,0
EG	0,0	100,0	0,0
EH	0,0	100,0	0,0
EO	100,0	0,0	0,0
FG	0,0	100,0	0,0
FH	50,0	50,0	0,0
FO	66,7	33,3	0,0
GJ	0,0	100,0	0,0
GL	0,0	100,0	0,0
GO	0,0	0,0	100,0
HJ	25,0	50,0	25,0
HO	0,0	100,0	0,0
JK	0,0	66,7	33,3
JL	0,0	100,0	0,0
JN	0,0	100,0	0,0
KK	0,0	100,0	0,0

Legend: A=Inert treatment; B=Acupuncture; C=Back school; D=Cognitive behavioral therapy; E=Education; F=Exercise; G=Heat wrap; H=Manual therapy; I=Muscle relaxant; J=NSAIDs; K=Opioids; L=Paracetamol; M=Physical therapy; N=Steroids; O=Usual care

Presence of leg pain or sciatica was reported in 15 studies out of 46 (31%) of which 6 were not included in quantitative analysis.

*AE: 1 study

*AF: 2 studies, of which 1 was not included in quantitative analysis (qualitative analysis).

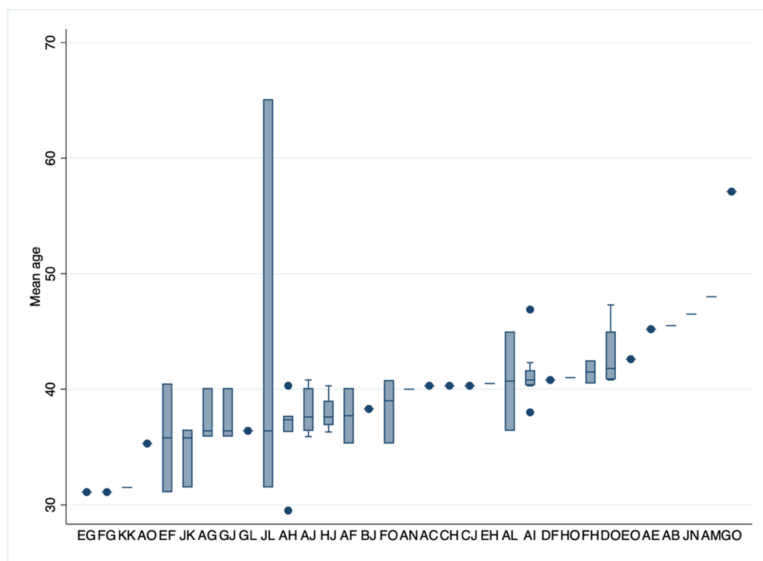
*EO: 1 study not included in quantitative analysis (qualitative analysis).

*BJ: 1 study

*AO: 1 study not included in quantitative analysis (qualitative analysis).

*FO: 2 studies of which 1 was not included in quantitative analysis (qualitative analysis).

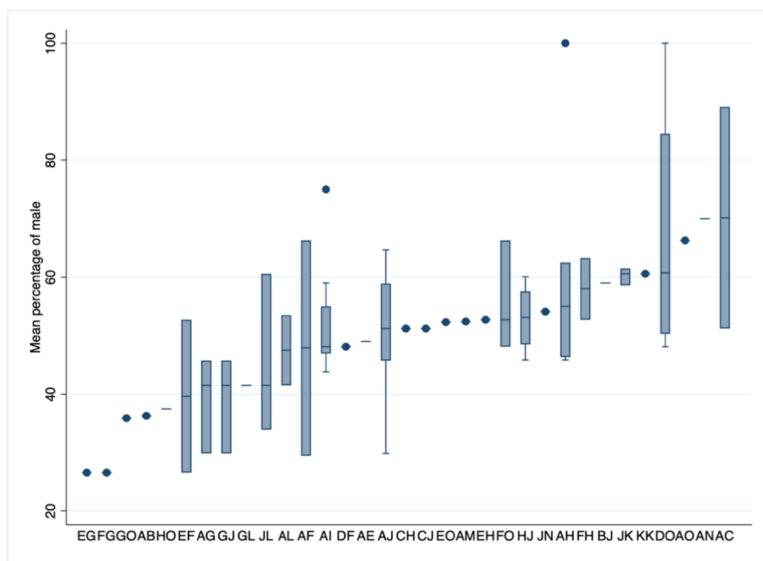
Overall, a very little percentage of leg pain or sciatica (0.09%) impact on global assessment.



Median of mean age ranged from 32 to 57 years old as already known by the Global Burden of Disease ²¹

Figure 6. Mean age

*JL: 1 out of 3 trials has a mean age of 65.1

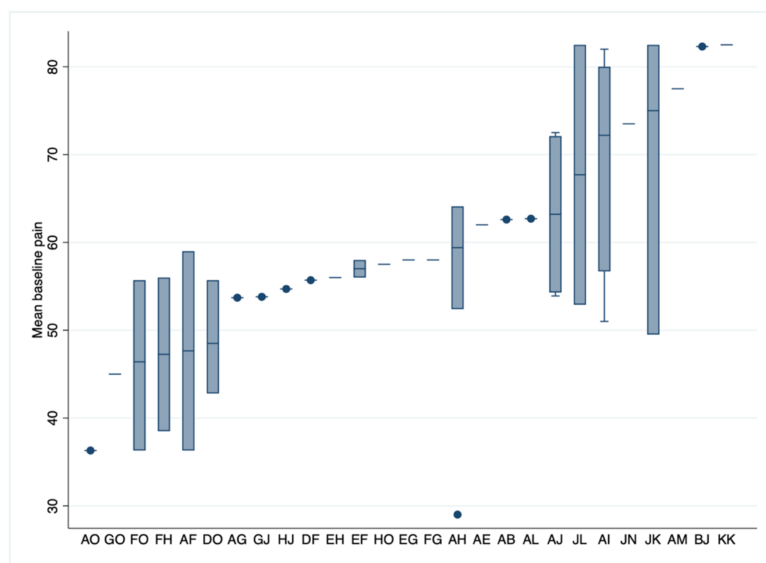


Median percentage of male ranged from 27 to 70 percent with overlapping of 25-75 percentiles across comparisons.

Figure 7. Percentage of male participants

*Five studies did not report gender

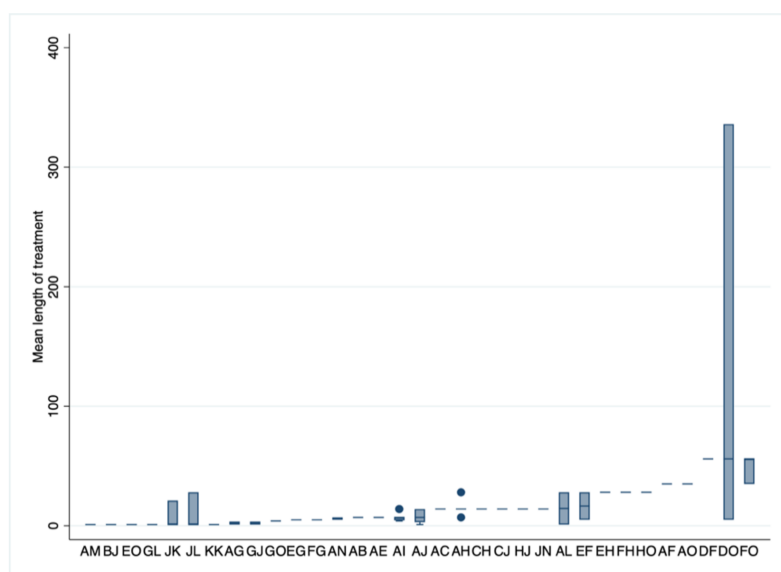
**AH and DO: outliers refer to 2 studies with a 100% male; however, these trials did not report outcome data and were not included in quantitative analysis (qualitative analysis). Excluding them, male and female can be equally distributed across interventions.



Median baseline pain ranged from 37 to 82 with overlapping of 25-75 percentiles across comparisons.

Figure 8. Baseline severity (pain)

*AH: 1 trial had an outlier mean baseline value of 29, however this trial did not report outcome data and was not included in quantitative analysis (qualitative analysis)



Median length of treatment ranged from 1 to 40 days with overlapping of 25-75 percentiles across comparisons.

Figure 9. Length of treatment

*DO: 1 trial had an outlier mean length of treatment of 336 days, however this trial did not report outcome data and was not included in quantitative analysis (qualitative analysis)

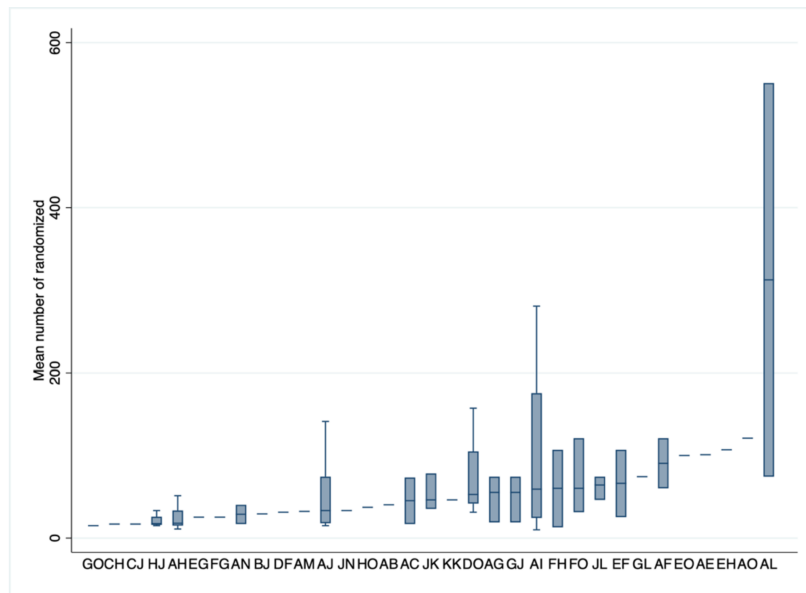


Figure 10. Number of randomized

*AL: 1 trial had an outlier number of randomized of 550, which represents less than 5% of the overall sample. However, we judged this reason insufficient to prejudice transitivity across interventions.

Legend: A=Inert treatment; B=Acupuncture; C=Back school; D=Cognitive behavioral therapy; E=Education; F=Exercise; G=Heat wrap; H=Manual therapy; I=Muscle relaxant; J=NSAIDs; K=Opioids; L=Paracetamol; M=Physical therapy; N=Steroids; O=Usual care

Supplement F. Risk of Bias

Figure 1. Aggregate Cochrane Risk-of-bias appraisal results

Risk of bias appraisal.²³

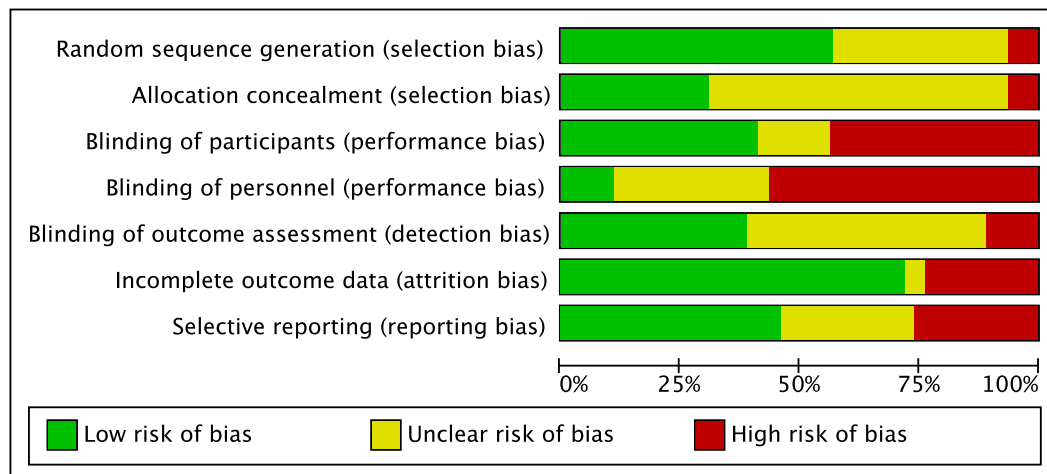


Table 1. Cochrane Risk-of-bias global judgement

Author, year	Random sequence generation	Allocation concealment	Blinding of participants	Blinding of personnel/ providers (performance bias)	Blinding of care	Blinding of outcome assessment	Incomplete outcome data	Selective Reporting	FINAL JUDGEMENT
Amlic 1987	unclear	unclear	low	unclear	unclear	low	low	low	unclear
Bergquist-Ullman 1977	low	unclear	high	high	unclear	high	low	high	high
Berry 1988	unclear	unclear	unclear	unclear	unclear	low	low	low	unclear
Bertalanffy 2005	low	low	low	high	low	low	low	low	low
Casale 1988	unclear	unclear	low	unclear	unclear	low	high	unclear	unclear
Cherkin 1996	high	unclear	high	high	low	low	unclear	unclear	unclear
Cherkin 1998	unclear	low	high	high	low	low	unclear	low	low
Dapas 1985	unclear	unclear	low	unclear	unclear	high	high	high	high
Dreiser 2003	low	low	low	unclear	unclear	low	low	low	unclear
Eken 2014	low	low	low	low	unclear	low	low	low	unclear
Eskin 2014	low	unclear	unclear	low	low	low	low	low	unclear
Faas 1995	high	unclear	high	high	high	low	low	low	high
Goldie 1968	unclear	unclear	low	low	unclear	low	low	low	unclear
Haimovic 1986	low	unclear	low	unclear	unclear	high	unclear	high	high
Hasagawa 2014	low	unclear	low	high	low	low	low	low	unclear
Hindle 1972	low	high	unclear	unclear	unclear	low	high	high	high
Jellema 2005	low	unclear	high	high	unclear	low	unclear	unclear	unclear
Ketenci 2005	unclear	unclear	low	unclear	unclear	low	low	low	unclear
Kettenmann 2007	high	high	high	unclear	high	high	unclear	high	high
Lindstrom 1995	unclear	unclear	high	unclear	unclear	low	high	unclear	unclear
Malmivaara 1995	low	low	high	high	low	low	low	low	low
Mayer 2005	low	unclear	high	high	unclear	low	high	unclear	unclear
Miki 2018	low	unclear	high	high	unclear	high	high	high	high
Nadler 2002	unclear	unclear	high	high	unclear	low	unclear	unclear	unclear
Nadler 2003b	unclear	unclear	high	high	unclear	low	unclear	unclear	unclear
Nadler 2003a	unclear	unclear	high	high	unclear	high	unclear	unclear	high
Postacchini 1988	unclear	unclear	unclear	unclear	unclear	unclear	high	unclear	unclear
Ralph 2008	unclear	unclear	unclear	unclear	unclear	low	high	unclear	unclear

Sae-Jung 2016	low	low	unclear	high	high	low	low	high
Santilli 2006	low	low	low	high	low	low	unclear	low
Schenk 2003	low	unclear	high	high	high	low	unclear	high
Schneider 2015	low	low	high	high	low	low	high	low
Seferlis 1998	unclear	unclear	high	high	unclear	high	low	high
Serfer 2010	low	unclear	low	low	high	low	low	high
Shin 2013	low	low	high	high	low	low	low	low
Storheim 2003	low	low	high	high	low	high	low	high
Suni 2006	low	unclear	high	high	low	unclear	unclear	unclear
Szpalski 1994	unclear	unclear	unclear	unclear	unclear	low	low	unclear
Takamoto 2015	low	unclear	high	high	low	high	high	high
Traeger 2019	low	low	low	high	low	low	low	low
Tuzun 2003	low	low	low	unclear	low	low	low	low
Veenema 2000	unclear	high	low	high	low	low	unclear	high
Videman 1984	unclear	unclear	low	unclear	unclear	low	unclear	unclear
Von Heymann 2013	low	low	low	high	low	high	high	high
Williams 2014	low	low	low	low	low	low	low	low
Younes 2017	low	unclear	low	high	low	high	high	high

Supplement G. Network Plots

Figure 1. Network Plot- Pain outcome

Note: The size of the nodes is proportional to the number of studies evaluating each intervention, and the thickness of the edges is proportional to the precision (the inverse of the variance) of each direct comparison.

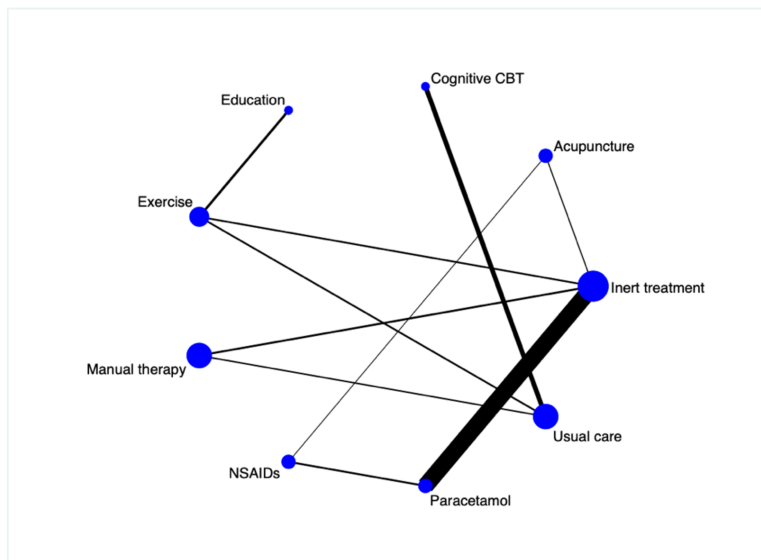


Figure 1a. Network for pain outcome at 1 month of FU

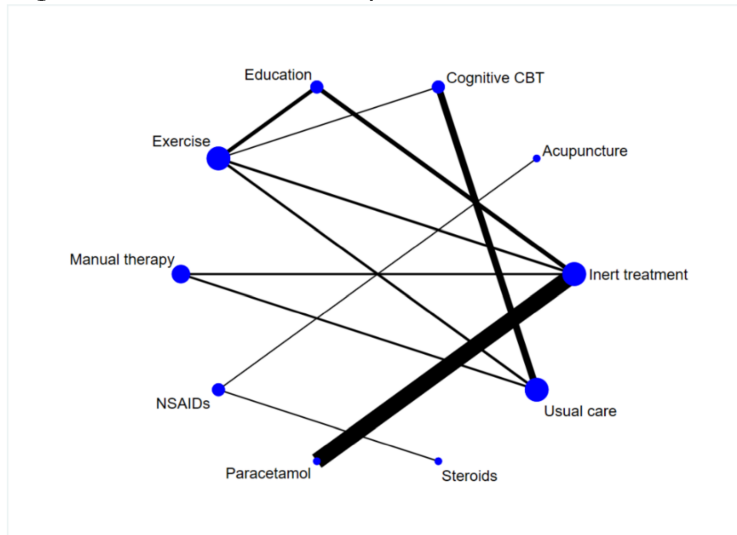


Figure 1b. Network for pain outcome at 3-6 months of FU

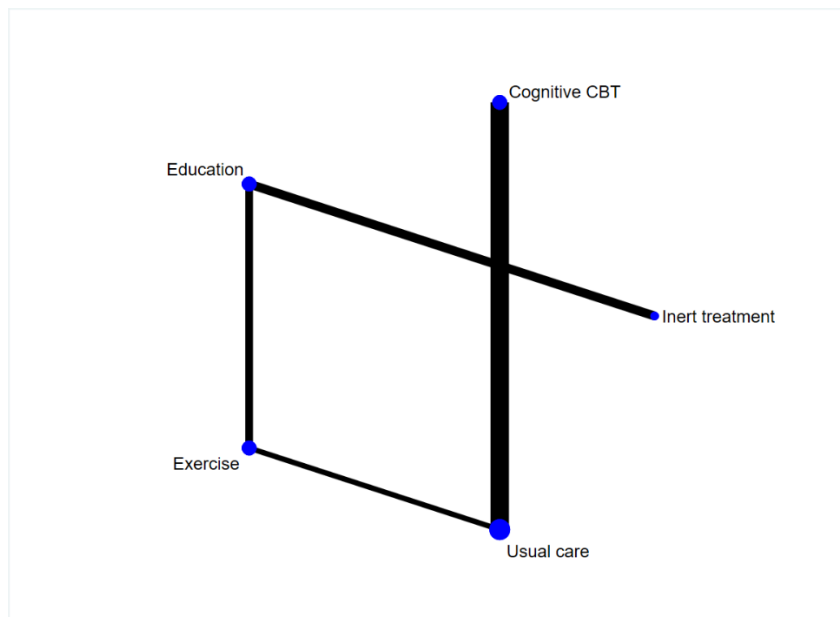


Figure 1c. Network for pain outcome at 12 months of FU

Figure 2. Network Plot- Disability outcome

Note: The size of the nodes is proportional to the number of studies evaluating each intervention, and the thickness of the edges is proportional to the precision (the inverse of the variance) of each direct comparison.

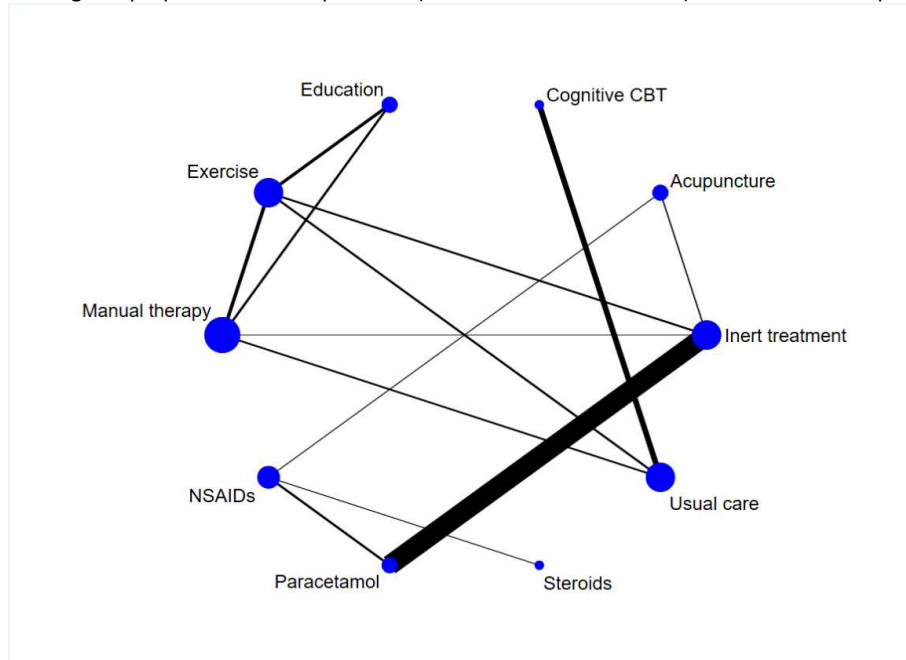


Figure 2a. Network for disability outcome at 1 month of FU

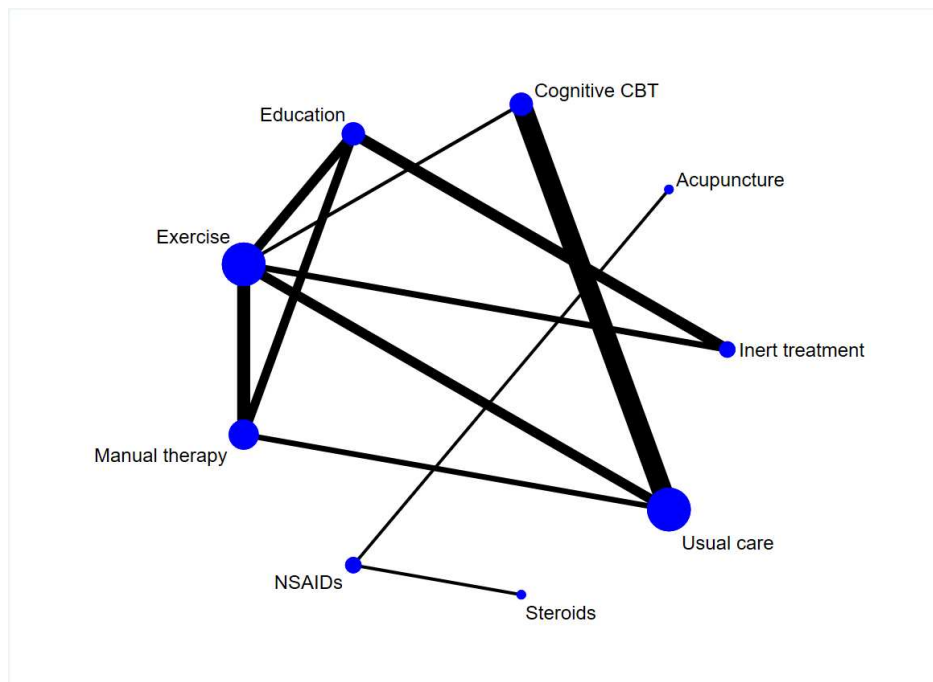


Figure 2b. Network for disability outcome at 3-6 months of FU

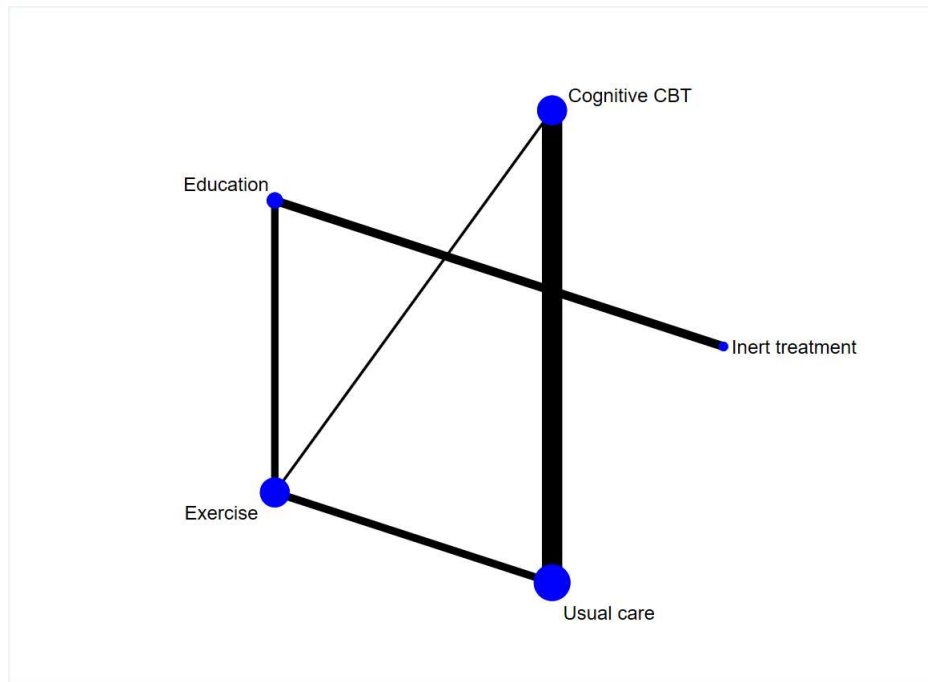


Figure 2c. Network for disability outcome at 12 months of FU

Supplement H. Assessment of pairwise Meta-Analyses

Pairwise meta-analyses –Pain Outcome

Table 1. Pairwise meta-analyses at 1 week of FU for pain

	Comparison	Number of studies	Effect size	Lower limit 95%	Upper limit 95%	Heterogeneity (I ²)	P value
1	Muscle relaxants vs Inert treatment	4	-1.06	-1.89	-0.24	91.1%	0.0000
2	Physical therapy vs Inert treatment	1	-2.85	-3.57	-2.14	Na	Na
3	NSAIDs vs Inert treatment	3	-0.84	-1.15	-0.53	54.2%	0.112
4	Opioid vs NSAIDs	2	-0.43	-0.71	-0.14	20.3%	0.263
5	Paracetamol vs NSAIDs	2	-0.21	-0.62	0.20	56.9%	0.128
6	Paracetamol vs Opioid	1	0.18	-0.24	0.59	Na	Na
7	Acupuncture vs Inert treatment	1	-0.30	-0.74	0.14	Na	Na
8	Exercise vs Education	1	-0.90	-1.47	-0.33	Na	Na
9	Heat wrap vs Education	1	-1.03	-1.60	-0.46	Na	Na
10	Heat wrap vs Exercise	1	-0.13	-0.68	0.43	Na	Na
11	Heat wrap vs Inert treatment	1	-4.77	-5.72	-3.81	Na	Na
12	Manual therapy vs Inert treatment	2	-1.20	-2.59	0.19	91.1%	0.000
13	Manual therapy vs Exercise	1	1.12	0.25	1.99	Na	Na
14	NSAIDs vs Acupuncture	1	-0.58	-1.11	-0.06	Na	Na
15	Education vs Inert treatment	1	0.04	-0.23	0.32	Na	Na
16	NSAIDs vs Manual therapy	1	0.67	0.20	1.13	Na	Na
17	Paracetamol vs Inert treatment	1	0.04	-0.08	0.16	Na	Na

Table 2. Pairwise meta-analyses at 1 month of FU for pain

	Comparison	Number of studies	Effect size	Lower limit 95%	Upper limit 95%	Heterogeneity (I ²)	P value
1	Exercise vs Education	1	-0.84	-1.14	-0.53	Na	Na
2	Acupuncture vs Inert treatment	1	-0.63	-1.08	-0.18	Na	Na
3	Usual care vs Cognitive CBT	1	0.04	-0.18	0.26	Na	Na

4	Exercise vs Inert treatment	1	0.00	-0.36	0.36	Na	Na
5	Paracetamol vs NSAIDs	1	-0.08	-0.43	0.27	Na	Na
6	Steroids vs NSAIDs	1	-1.51	-2.06	-0.95	Na	Na
7	Manual therapy vs Inert treatment	2	-0.86	-1.45	-0.27	59.7%	0.115
8	Usual care vs Manual therapy	2	0.61	-0.15	1.37	72.6%	0.056
9	Usual care vs Exercise	1	0.00	-0.36	0.36	Na	Na
10	NSAIDs vs Acupuncture	1	-0.55	-1.07	-0.02	Na	Na
11	Paracetamol vs Inert treatment	1	0.00	-0.12	0.12	Na	Na

Table 3. Pairwise meta-analyses at 3-6 months of FU for pain

	Comparison	Number of studies	Effect size	Lower limit 95%	Upper limit 95%	Heterogeneity (I ²)	P value
1	Exercise vs Education	1	-0.17	-0.47	0.13	Na	Na
2	Usual care vs Cognitive CBT	1	0.00	-0.22	0.22	Na	Na
3	Manual therapy vs Inert treatment	1	-0.80	-1.20	-0.40	Na	Na
4	Usual care vs Manual therapy	2	0.06	-0.62	0.73	66.6%	0.084
5	Usual care vs Exercise	1	0.00	-0.36	0.36	Na	Na
6	Exercise vs Cognitive CBT	1	-0.47	-0.97	0.03	Na	Na
7	Education vs Inert treatment	1	-0.08	-0.36	0.19	Na	Na
8	Paracetamol vs Inert treatment	1	-0.04	-0.16	0.07	Na	Na

Table 4. Pairwise meta-analyses at 12 months of FU for pain

	Comparison	Number of studies	Effect size	Lower limit 95%	Upper limit 95%	Heterogeneity (I ²)	P value
1	Exercise vs Education	1	-0.39	-0.68	-0.09	Na	Na
2	Usual care vs Cognitive CBT	2	0.09	-0.40	0.58	79.3%	0.028
3	Usual care vs Exercise	1	0.00	-0.36	0.36	Na	Na
4	Education vs Inert treatment	1	-0.30	-0.58	-0.03	Na	Na

Pairwise meta-analyses – Disability Outcome

Table 5. Pairwise meta-analyses at 1 week of FU for disability

	Comparison	Number of studies	Effect size	Lower limit 95%	Upper limit 95%	Heterogeneity (I ²)	P value
1	NSAIDs-Inert treatment	2*(3)	-0.432	-0.664	-0.199	22.3%	0.000
2	Acupuncture- Inert treatment	1	-0.385	-0.828	0.057	Na	0.088
3	Exercise-Education	1	-0.291	-0.842	0.260	Na	0.300
4	Heat Wrap-Education	1	-0.414	-0.967	0.140	Na	0.143
5	Heat Wrap-Exercise	1	-0.122	-0.677	0.432	Na	0.666
6	Paracetamol-NSAIDs	2	0.010	-0.201	0.221	0.0%	0.924
7	NSAIDs –Heat Wrap	1	-0.512	-0.780	-0.244	Na	0.000
8	Paracetamol–Heat Wrap	1	-0.466	-0.729	-0.202	Na	0.001
9	Heat Wrap- Inert treatment	1	-0.544	-0.792	-0.295	0.0%	0.000
10	Muscle Relaxant-Inert treatment	2*(3)	-0.235	-0.439	-0.031	70.6%	0.024
11	Manual therapy-Exercise	1	0.772	-0.063	1.606	Na	0.070
12	NSAIDs – Acupuncture	1	-0.732	-1.265	-0.199	Na	0.007
13	Manual therapy-Inert treatment	2	-0.660	-1.099	-0.221	19.6%	0.003
14	Education-Inert treatment	1	-0.271	-0.548	0.006	Na	0.055
15	NSAIDs –Manual Therapy	1	0.793	0.327	1.260	Na	0.001
16	Paracetamol-Inert treatment	1	-0.092	-0.210	0.026	Na	0.126

*3 comparisons from 2 studies

Table 6. Pairwise meta-analyses at 1 month of FU for disability

	Comparison	Number of studies	Effect size	Lower limit 95%	Upper limit 95%	Heterogeneity (I ²)	P value
1	Usual care – Manual therapy	1 *(2)	0.239	-0.333	0.810	53.5%	0.413
2	Acupuncture – Inert treatment	1	-0.709	-1.162	-0.257	Na	0.002
3	Usual care – Cognitive CBT	1	0.019	-0.203	0.241	Na	0.868
4	Exercise - Inert treatment	1	0.674	0.302	1.047	Na	0.000
5	Paracetamol - NSAIDs	1	-0.128	-0.476	0.220	Na	0.472
6	Steroids - NSAIDs	1	-1.215	-1.747	-0.682	Na	0.000

7	Usual care – Exercise	1	0.000	-0.358	0.358	Na	1.000
8	NSAIDs Acupuncture	1	-0.640	-1.169	-0.111	Na	0.018
9	Manual therapy - Inert treatment	1	-0.819	-1.438	-0.201	Na	0.009
10	Paracetamol - Inert treatment	1	-0.019	-0.137	0.099	Na	0.747
11	Exercise - Education	1	-0.426	-0.723	-0.129	Na	0.005
12	Manual therapy - Education	1	-2.158	-2.502	-1.815	Na	0.000
13	Manual therapy - Exercise	1	-1.732	-2.012	-1.452	Na	0.000

*2 comparisons from 1 study

Table 7. Pairwise meta-analyses at 3-6 months of FU for disability

	Comparison	Number of studies	Effect size	Lower limit 95%	Upper limit 95%	Heterogeneity (I ²)	P value
1	Usual care – Manual Therapy	1 *(2)	0.039	-0.348	0.426	0%	0.844
2	Usual care – Cognitive CBT	2	0.212	-0.333	0.757	75.4%	0.446
3	Exercise - Inert treatment	1	0.312	-0.052	0.677	Na	0.093
4	Steroids - NSAIDs	1	-0.794	-1.300	-0.287	Na	0.002
5	Usual care - Exercise	2	0.159	-0.229	0.547	38.0%	0.422
6	NSAIDs - Acupuncture	1	0.435	-0.087	0.956	Na	0.102
7	Exercise- Cognitive CBT	1	0.135	-0.356	0.627	Na	0.590
8	Education - Inert treatment	1	-0.096	-0.372	0.180	Na	0.496
9	Exercise- Education	1	-0.052	-0.347	0.243	Na	0.731
10	Manual therapy - Education	1	-0.896	-1.204	-0.588	Na	0.000
11	Manual therapy - Exercise	1	-0.844	-1.099	-0.590	Na	0.000

*2 comparisons from 1 study

Table 8. Pairwise meta-analyses at 12 months of FU for disability

	Comparison	Number of studies	Effect size	Lower limit 95%	Upper limit 95%	Heterogeneity (I ²)	P value
1	Exercise - Education	1	-0.437	-0.735	-0.138	Na	0.004
2	Usual care - Cognitive CBT	3	0.332	-0.142	0.806	80.4%	0.170
3	Usual care - Exercise	2	0.185	-0.249	0.619	49.5%	0.403
4	Exercise - Cognitive CBT	1	0.086	-0.405	0.577	Na	0.732
5	Education - Inert treatment	1	-0.163	-0.439	0.114	Na	0.249

Supplement I. Forest plot of network meta-analysis (network forest)

Figure 1. Network forest – pain outcome 1 week

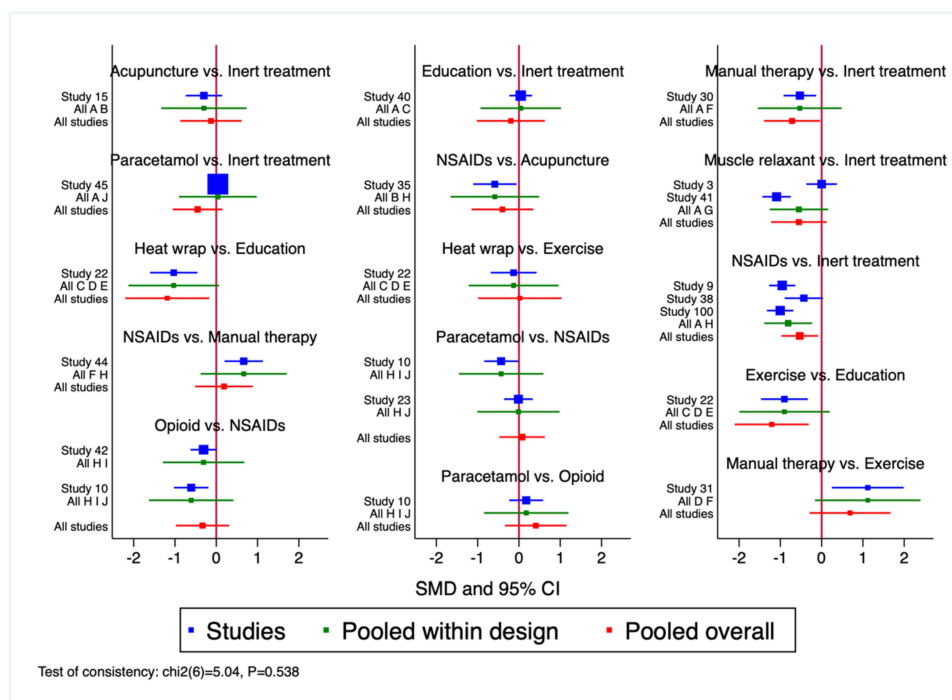


Figure 2. Network forest – pain outcome 1 month

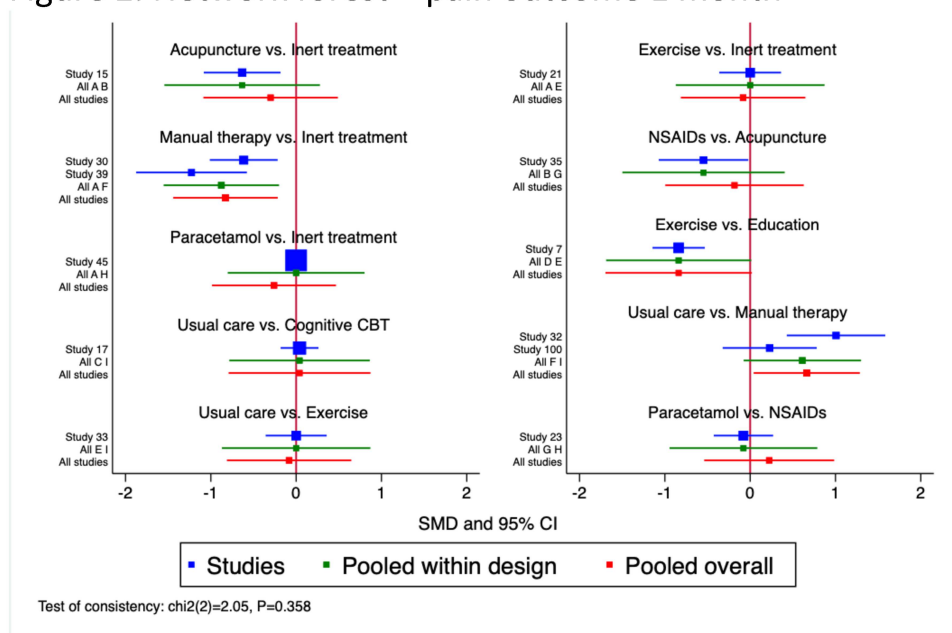


Figure 3. Network forest – pain outcome 12 months

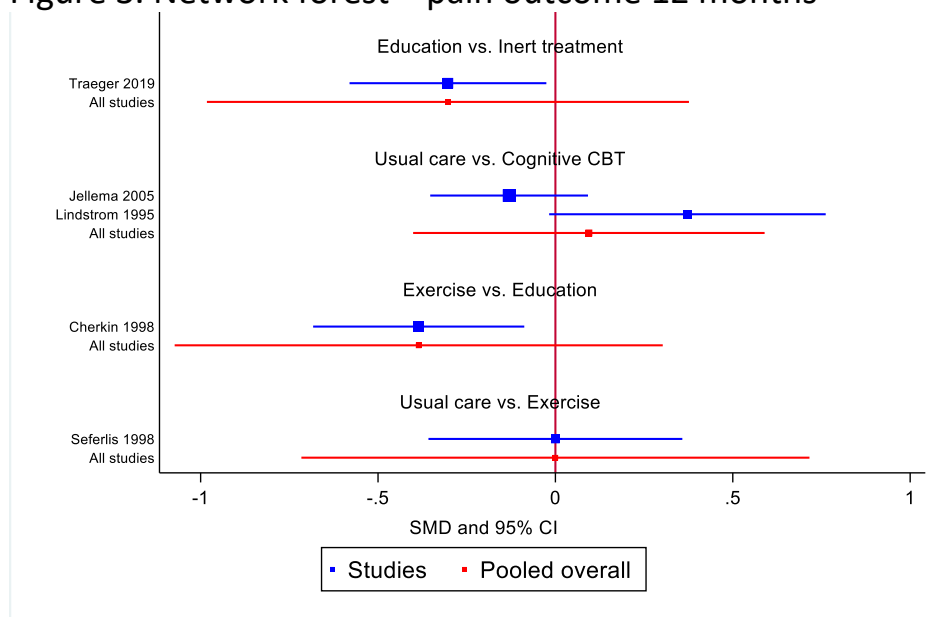


Figure 4. Network forest – disability outcome 1 week

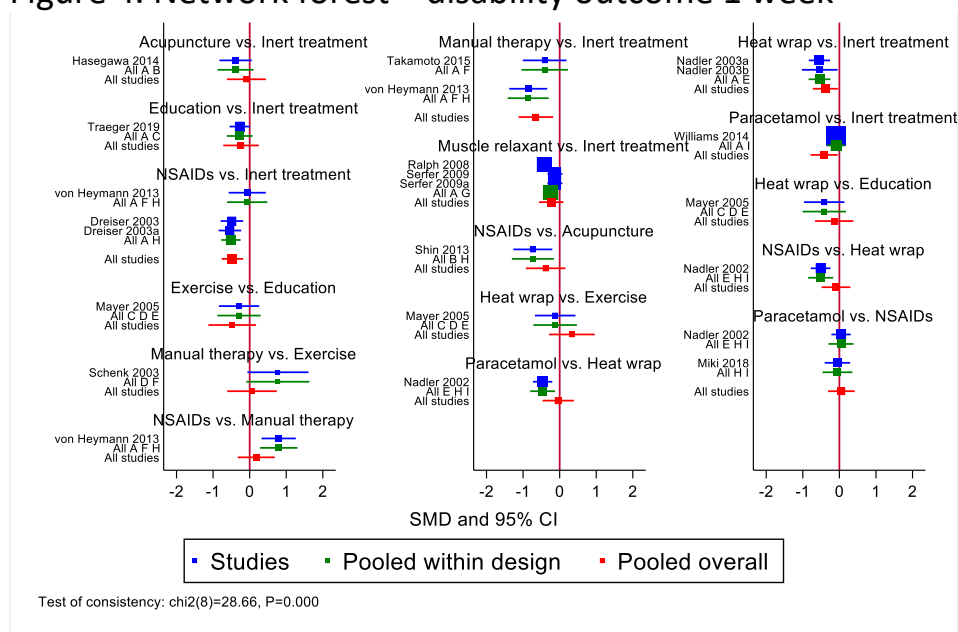


Figure 5. Network forest – disability outcome 1 month

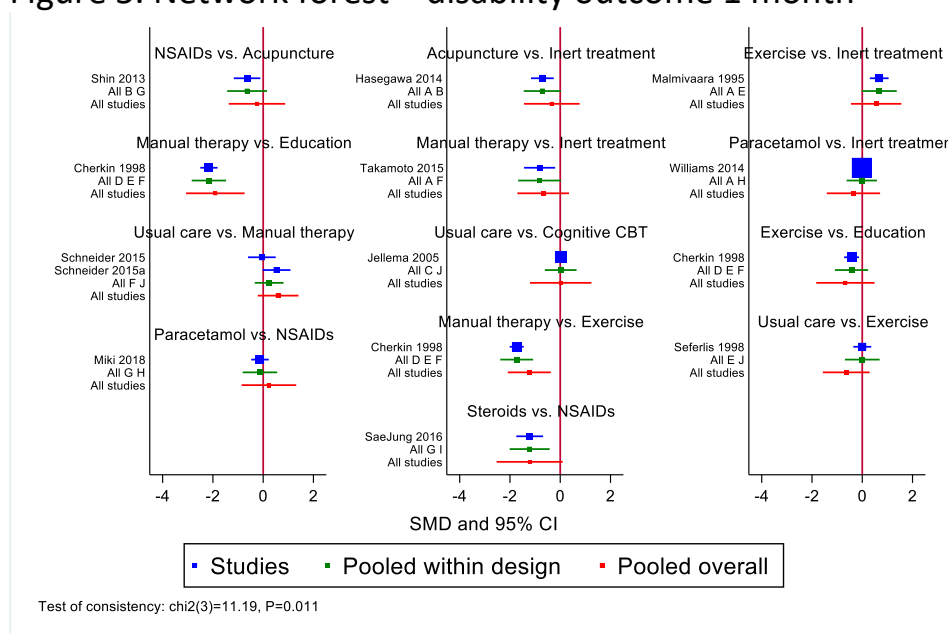
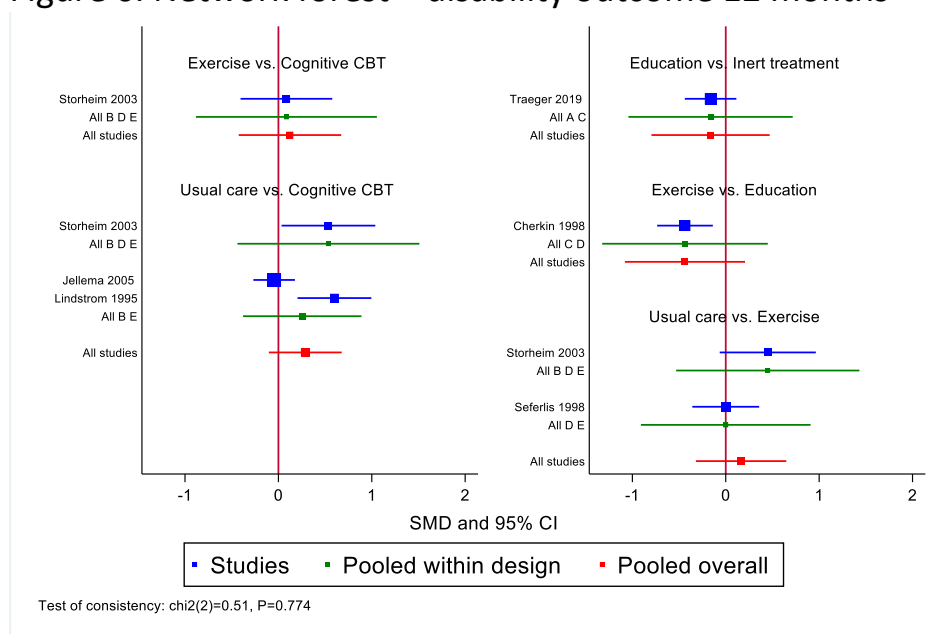


Figure 6. Network forest – disability outcome 12 months



Supplement J. Incoherence estimation and evaluation

Table 1. Estimated Global Inconsistency in Networks

OUTCOME	FOLLOW UP	Chi square	Prob > chi2	tau
PAIN	1 week	chi2 (7) = 9.48	Prob > chi2 = 0.5383	0.234
	1 month	chi2 (2) = 2.05	Prob > chi2 = 0.3583	0.169
	3-6 months	disconnected		-
	12 months	chi2 (1) = 0.00	Prob > chi2 = 1**	0.1
DISABILITY	1 week	chi2 (8) =28.66	Prob > chi2 = 0.0004*	-
	1 month	chi2 (3) =11.20	Prob > chi2 = 0.0107*	-
	3-6 months	disconnected		-
	12 months	chi2 (2) = 0.51	Prob > chi2 = 0.7737	0.097

* Global consistency is tested here using the 'design-by-interaction' test that infers consistency across an entire treatment network, using a chi square test. A p value <0.05 is taken to infer evidence of global inconsistency in the network.^{24 25}

**all the evidence about these contrasts comes from the trials which directly compare them

Table 2. Estimated Local Inconsistency for each pairwise comparison (side splitting) – pain outcome

Table 2a. Nodesplit pain 1 week

Side	Direct		Indirect		Difference			tau
	Coef.	Std. Err.	Coef.	Std. Err.	Coef.	Std. Err.	P>z	
Inert treatment - Acupuncture	-.2987834	.5246669	.0931138	.5981655	-.3918972	.7956616	0.622	.4740148
Inert treatment - Education	.0432741	.4689486	-1079062,00	.9044266	1122337,00	1018774,00	0.271	.4473322
Inert treatment - Manual therapy	-.5280427	.5132268	-.8939374	.5025075	.3658947	.7182726	0.610	.4719181
Inert treatment - Muscle relaxant

Inert treatment - NSAIDs	-.8159915	.2426794	-.0329156	.3199731	-.7830758	.4018672	0.051	.3754527
Inert treatment - Paracetamol	.0384353	.4065262	-.8652568	.3777104	.9036921	.5549132	0.103	.4020402
Acupuncture - NSAIDs	-.5837083	.5448436	-.1918109	.5798476	-.3918974	.7956619	0.622	.4740148
Education – Exercise *	-.9012443	.5332432	-2023588,00	.8680764	1122343,00	1018776,00	0.271	.4473321
Education - Heat wrap *	-1029994,00	.5348997	-3274667,00	1963983,00	2244673,00	2037546,00	0.271	.4473318
Exercise - Heat wrap *	-.1287492	.5293618	2115939,00	1968485,00	-2244688,00	2037552,00	0.271	.4473321
Exercise - Manual therapy	1117072,00	.6305311	-.005282	.8002101	1122354,00	1018777,00	0.271	.4473321
Manual therapy - NSAIDs	.6652757	.4944677	-.2694296	.4841419	.9347054	.69202	0.177	.4335961
NSAIDs - Opioid *	-.4512816	.3356582	.9098231	1082583,00	-1361105,00	1133386,00	0.230	.4358473

* All the evidence about these contrasts comes from the trials which directly compare them.

Table 2b. Nodesplit pain 1 month

Side	Direct		Indirect		Difference		P>z	tau
	Coef.	Std. Err.	Coef.	Std. Err.	Coef.	Std. Err.		
Inert treatment - Acupuncture	-.6327764	.3567964	.6254979	.5752867	-1.258.274	.6769479	0.063	.273273
Inert treatment - Exercise	-4.80e-12	.5233844	-.2740767	.7685576	.2740767	.9298451	0.768	.4896684
Inert treatment - Manual therapy	-.8871542	.3955099	-.613068	.8416375	-.2740862	.9298405	0.768	.4896674
Inert treatment - Paracetamol	-2.90e-12	.2798297	-1.258.269	.6164035	1.258.269	.6769475	0.063	.273273
Acupuncture - NSAIDs	-.5466608	.3826874	.7116145	.5583996	-1.258.275	.6769489	0.063	.2732733
Cognitive CBT - Usual care *	.0399034	.4245035	-.3263798	6.354.628	.3662832	6.354.629	1.000	.4090962
Education - Exercise *	-.8383118	.4379943	-.4467205	6.328.197	-.3915912	6.328.198	1.000	.4090963
Exercise - Usual care	-2.29e-08	.5225983	-.2740773	.7690965	.2740772	.9298486	0.768	.4896669
Manual therapy - Usual care	.6130723	.4016588	.8871557	.8387265	-.2740834	.9298459	0.768	.4896684
NSAIDs - Usual care	-.078838	.3258861	1.179.435	.5933446	-1.258.273	.6769487	0.063	.2732733

* All the evidence about these contrasts comes from the trials which directly compare them.

Table 2c. Nodesplit pain 12 months

Side	Direct		Indirect		Difference		P>z	tau
	Coef.	Std. Err.	Coef.	Std. Err.	Coef.	Std. Err.		
Inert treatment - Education*	-.3029187	.34666	.3777316	158.3944	-.6806503	158.3948	0.997	.3164487
Cognitive CBT - Usual care*	.0943039	.2527336	-1.379709	447.7409	1.474013	447.7409	0.997	.316448
Education - Exercise*	-.385339	.3509876	.3660218	174.4564	-.7513608	174.4568	0.997	.3164487
Exercise - Usual care*	-9.18e-11	.3653395	.8080591	209.9836	-.8080591	209.9839	0.997	.3164485

* All the evidence about these contrasts comes from the trials which directly compare them.

Table 3. Estimated Local Inconsistency for each pairwise comparison (side splitting) – disability outcome

Table 3a. Nodesplit disability 1 week

Side	Direct		Indirect		Difference		P>z	tau
	Coef.	Std. Err.	Coef.	Std. Err.	Coef.	Std. Err.		
Inert Treatment-Acupuncture	-0.3850695	0.3512901	0.318208	0.412454	-0.7032775	0.541778	0.194	0.269133
Inert Treatment- Education	-0.2712998	0.3261325	-0.18365	0.424351	-0.0876449	0.535197	0.87	0.293896
Inert Treatment-Heat wrap	-0.5423379	0.2294745	-0.17954	0.253958	-0.3627932	0.342356	0.289	0.259164
Inert Treatment-Manual therapy	-0.664142	0.2886231	-0.59046	0.501075	-0.0736865	0.581203	0.899	0.292533
Inert Treatment-Muscle relaxant
Inert Treatment-NSAIDs	-0.387447	0.2022145	-0.59797	0.251741	0.2105194	0.324018	0.516	0.293991
Inert Treatment-Paracetamol	-0.0922448	0.2390906	-0.67043	0.219723	0.5781899	0.324719	0.075	0.231374
Acupuncture- NSAIDs	-0.731988	0.38266	-0.02871	0.383529	-0.7032779	0.541778	0.194	0.269133
Education- Exercise	-0.2919225	0.4040913	-0.93469	0.632299	0.6427636	0.750304	0.392	0.290215
Education- Heat wrap	-0.4121889	0.3985883	0.083842	0.365582	-0.4960307	0.540926	0.359	0.281415
Exercise-Heat wrap	-0.1227089	0.3721725	1.177.067	0.505458	-1.299.776	0.627943	0.038	0.241674
Exercise- Manual therapy	0.7716	0.4925257	-0.52044	0.434413	1.292.041	0.656732	0.049	0.24743

Heat wrap- NSAIDs	-0.5127726	0.274752	0.1945	0.237414	-0.7072724	0.36315	0.051	0.238334
Heat wrap- Paracetamol	-0.4646165	0.2367674	0.3788	0.239479	-0.8434166	0.336712	0.012	0.195007
Manual therapy- NSAIDs	0.7923256	0.328629	-0.40012	0.328938	1.192.444	0.463877	0.01	0.226649
NSAIDs-Paracetamol	-0.0008166	0.2354043	0.15986	0.348297	-0.1606761	0.420353	0.702	0.293809

* All the evidence about these contrasts comes from the trials which directly compare them; inconsistency in bold contrast are >5% of the all comparisons

Table 3b. Nodesplit disability 1 month

Side	Direct		Indirect		Difference		P>z	tau
	Coef.	Std. Err.	Coef.	Std. Err.	Coef.	Std. Err.		
Inert Treatment -Acupuncture	-0.7093169	0.6236239	0.7481728	1.055.844	-145.749	122.626	0.235	0.579317
Inert Treatment-Exercise	0.6744899	0.7305522	0.3343372	0.9563461	0.3401527	1.203.455	0.777	0.705391
Inert Treatment-Manual Therapy	-0.819488	0.772666	-0.4793281	0.92265	-0.34016	1.203.452	0.777	0.705389
Inert Treatment- Paracetamol	-0.0194038	0.5824383	-1.476.859	1.079.109	1.457.455	1.226.259	0.235	0.579317
Acupuncture-NSAIDs	-0.6397983	0.6390752	0.8176958	1.046.569	-1.457.494	1.226.264	0.235	0.579317
Cognitive CBT-Usual care *	0.0188224	0.6228875	-0.1682687	6.329.995	0.1870911	6.329.998	1.000	0.612493
Education-Exercise *	-0.4262689	0.5999444	-2.366.002	1.562.167	1.939.733	1.667.265	0.245	0.580495
Education-Manual therapy *	-2.158.292	0.6063919	-0.2185552	155.468	-1.939.737	1.667.265	0.245	0.580495
Exercise- Manual therapy *	-1.732.024	0.5978718	-0.7621531	0.5809457	-0.9698712	0.8336358	0.245	0.580497
Exercise- Usual care	-1.82E-10	0.4822981	-1.423.537	0.5431255	1.423.537	0.7263586	0.05	0.446406
Manual Therapy-Usual care	0.2390929	0.3731235	1.662.631	0.6231943	-1.423.538	0.7263602	0.05	0.446407
NSAIDs- Paracetamol	-0.127779	0.6059484	1.329.688	1.066.091	-1.457.467	1.226.264	0.235	0.579317
NSAIDs- Steroids *	-1.214.723	0.6700337	1.142.942	630.608	-2.357.665	6.306.084	0.997	0.612493

* All the evidence about these contrasts comes from the trials which directly compare them.

Table 3c. Nodesplit disability 12 months

Side	Direct		Indirect		Difference		P>z	tau
	Coef.	Std. Err.	Coef.	Std. Err.	Coef.	Std. Err.		
Inert treatment-Education*	-0.162517	0.323069	0.382189	141.004	-0.54471	1.410.044	0.997	0.290697
Cognitive CBT-Exercise	0.088617	0.446814	0.174454	0.492926	-0.08584	0.6648704	0.897	0.369949
Cognitive CBT-Usual care*	0.3264051	0.226606	-0.35701	1.060.696	0.683413	1.086.459	0.529	0.336763
Education-Exercise*	-0.436679	0.328125	0.151605	1.535.627	-0.58828	153.563	0.997	0.290697
Exercise-Usual care *	0.2022777	0.296387	-0.12221	0.932483	0.32449	0.9785033	0.74	0.354265

* All the evidence about these contrasts comes from the trials which directly compare them.

Table 4. Strategy to explore global inconsistency – disability 1 week

	Study removed	Chi square	Prob > chi2	Resolving inconsistency
All studies		chi2 (8) = 28.66	Prob > chi2 = 0.0004*	
STRATEGY 1: nodesplitting				
All studies without inconsistent constast (Exercise-Heat wrap)	Mayer 2005	chi2 (6) = 21.33	Prob > chi2 = 0.0016*	Not resolved
All studies without inconsistent constast (Exercise- Manual therapy)	Shrenk 2003	chi2 (7) = 22.93	Prob > chi2 = 0.0018*	Not resolved
All studies without inconsistent constast (Heat wrap-Paracetamol)	Nadler 2002	chi2 (6) = 14.38	Prob > chi2 = 0.0257*	Not resolved
All studies without inconsistent constast (Manual therapy-NSAIDs)	von Heymann 2013	chi2 (6) = 19.47	Prob > chi2 = 0.0034*	Not resolved
All studies without the four previous inconsistent constasts	All studies above	chi2 (2) = 6.03	Prob > chi2 = 0.0491*	Not resolved
STRATEGY 2: inspection of covariates				
Metaregression	The effects of the investigated co-variates were not statistically significant. See Table 6a			Not resolved
STRATEGY 3: inspection of subgroups				
Subgroup analysis (splitting pharmacological from non-pharmacological intervention)	Dreiser 2003; Miki 2018; Nadler 2002; Ralph 2008; Serfer 2009; Shin 2013; von Heymann 2013 (arm NSAIDs); Williams 2014	chi2 (2) = 3.19	Prob > chi2 = 0.2030	Resolved
Subgroup analysis (splitting non-pharmacological from pharmacological intervention)	Hasegawa 2014; Mayer 2005; Nadler 2002 (arm heat wrap); Nadler 2003a; Nadler 2003b; Schenk 2003; Shin 2013; Takamoto 2015; Traeger 2019; von Heymann 2013 (arm manual therapy)	chi2 (1) = 2.14	Prob > chi2 = 0.1432	Resolved

* Global consistency is tested here using the 'design-by-interaction' test that infers consistency across an entire treatment network, using a chi square test. A p value <0.05 is taken to infer evidence of global inconsistency in the network. ^{24 25}

Table 5. Strategy to explore global inconsistency – disability 1 month

	Study removed	Chi square	Prob > chi2	Resolving inconsistency
All studies		chi2 (3) = 11.20	Prob > chi2 = 0.0107*	See network meta forest
STRATEGY 1: nodesplitting				
All studies without inconsistent constast	No contrast statistically significant			Not resolved
STRATEGY 2: inspection of covariates				
Metaregression	The effects of the investigated co-variables were not statistically significant. See Table 6b			Not resolved
STRATEGY 3: inspection of subgroups				
Subgroup analysis (splitting pharmacological from non-pharmacological intervention)	Miki 2008, Sea-Jung 2016; Shin 2013, Williams 2014	chi2 (2) = 7.15	Prob > chi2 = 0.0280*	Not resolved; See network meta forest
Subgroup analysis (splitting non-pharmacological from pharmacological intervention)	Cherkin 1998, Hasegawa 2014, Jellema 2005, Malmivaara 1995, Schneider 2015, Seferlis 1998, Shin 2013, Takamoto 2015	chi2 (1) = 19.69	Prob > chi2 = 0.0000*	Not resolved; See network meta forest

* Global consistency is tested here using the 'design-by-interaction' test that infers consistency across an entire treatment network, using a chi square test. A p value <0.05 is taken to infer evidence of global inconsistency in the network. ^{24 25}

Table 6a. Metaregression disability 1 week

Variable	Coeff.	St. error	P>[t]	Tau2	95% CI	
Age	0.003	0.008	0.699	0.067	-0.014	0.021
Gender	0.005	0.007	0.477	0.067	-0.010	0.021
Patients with subacute/acute pain	-0.022	0.077	0.782	0.067	-0.181	0.138
Baseline value of pain	-0.008	0.007	0.244	0.098	-0.023	0.006
Presence of leg pain or sciatica	-0.039	0.143	0.783	0.069	-0.337	0.257
Risk of bias	0.124	0.104	0.246	0.067	-0.092	0.342

Table 6b. Metaregression disability 1 month

Variable	Coeff.	St. error	P>[t]	Tau2	95% CI	
Age	0.014	0.034	0.677	0.664	-0.059	0.088
Gender	-0.043	0.022	0.071	0.504	-0.090	0.004
Patients with subacute/acute pain	-0.257	0.213	0.252	0.591	-0.721	0.207
Baseline value of pain	-0.017	0.026	0.533	0.651	-0.073	0.039
Presence of leg pain or sciatica	-0.113	0.235	0.638	0.660	-0.624	0.398
Risk of bias	0.008	0.259	0.976	0.674	-0.571	0.555

Figure 1. Bubble plot disability 1 week

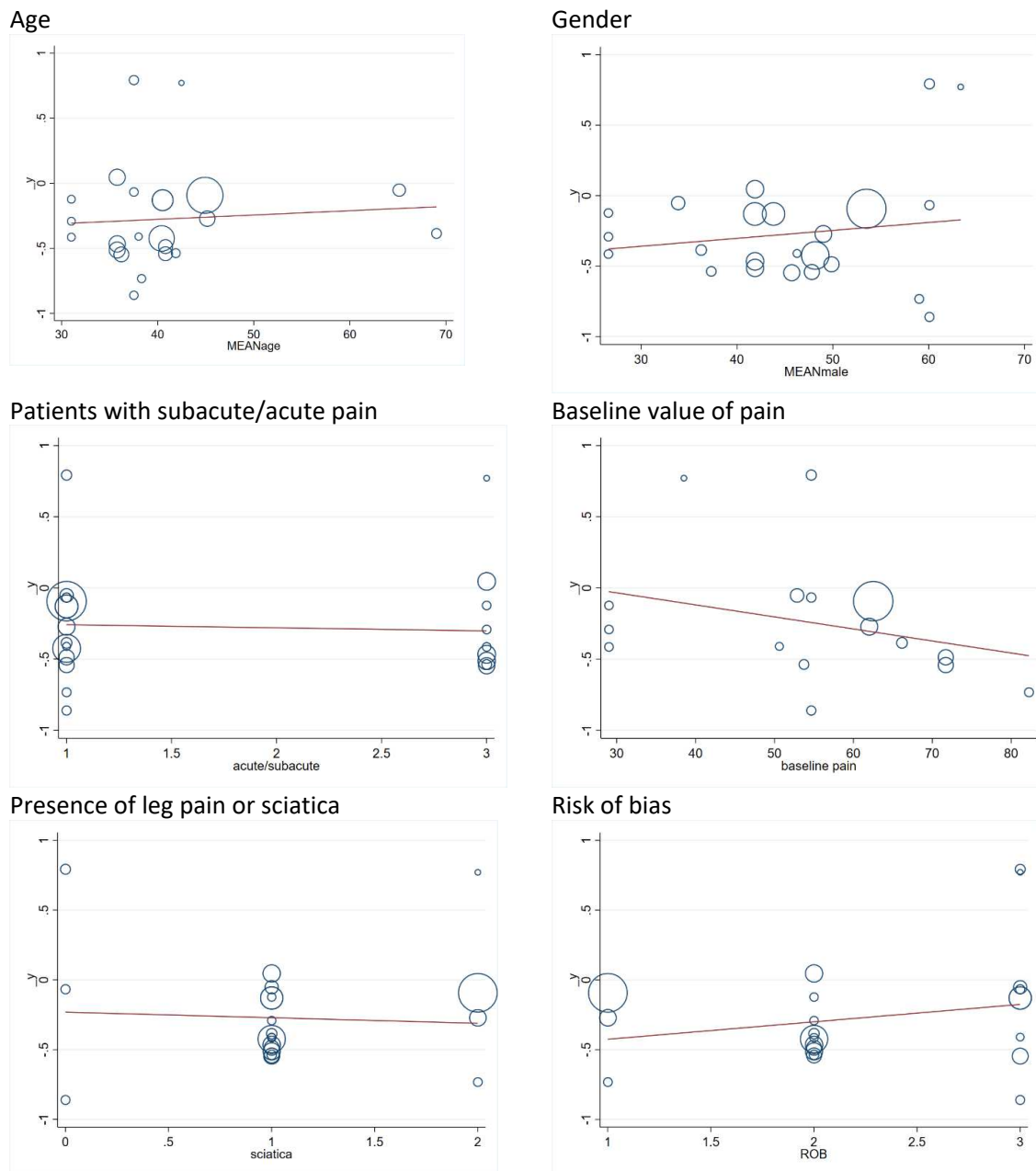
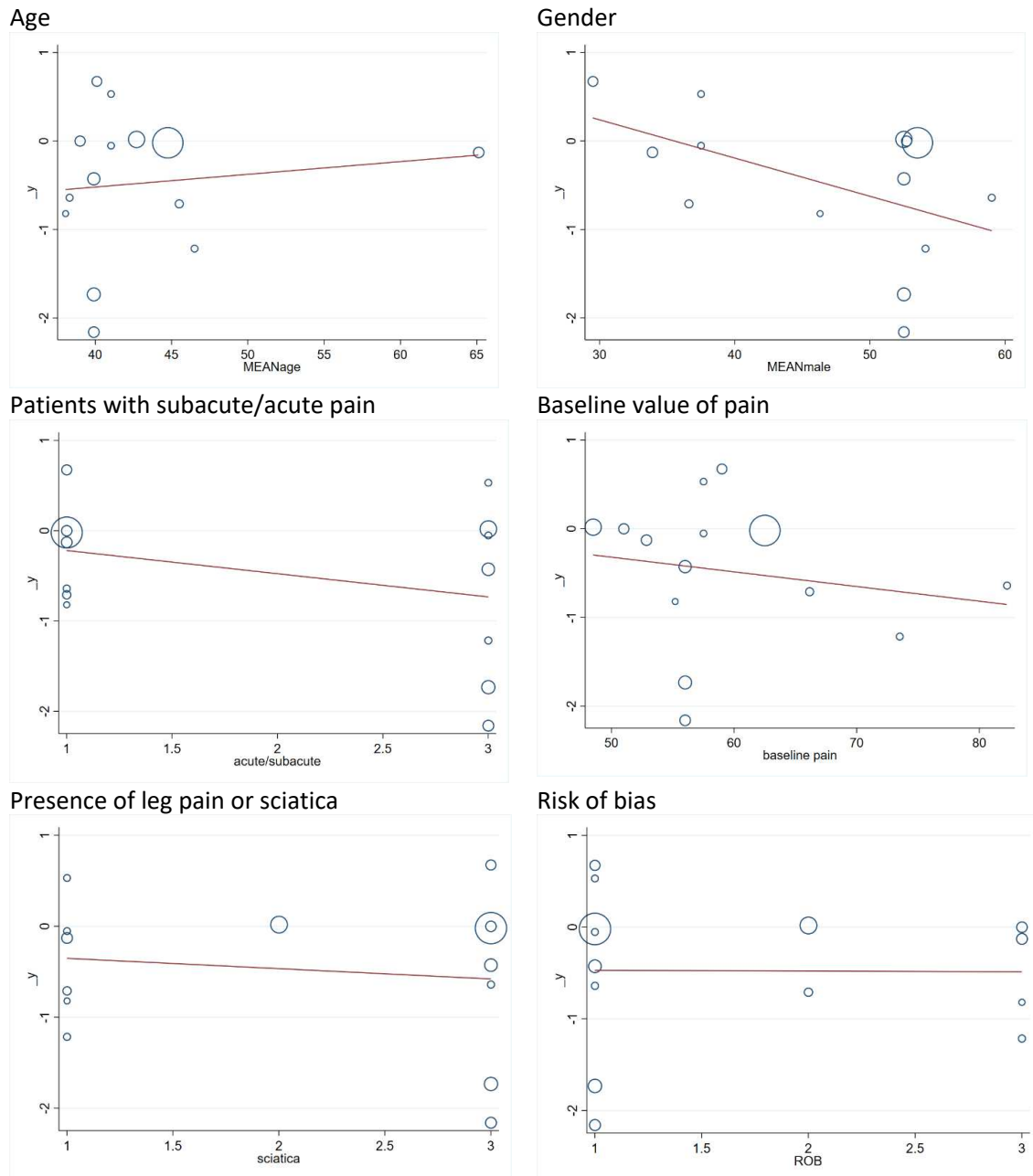


Figure 2. Bubble plot disability 1 month

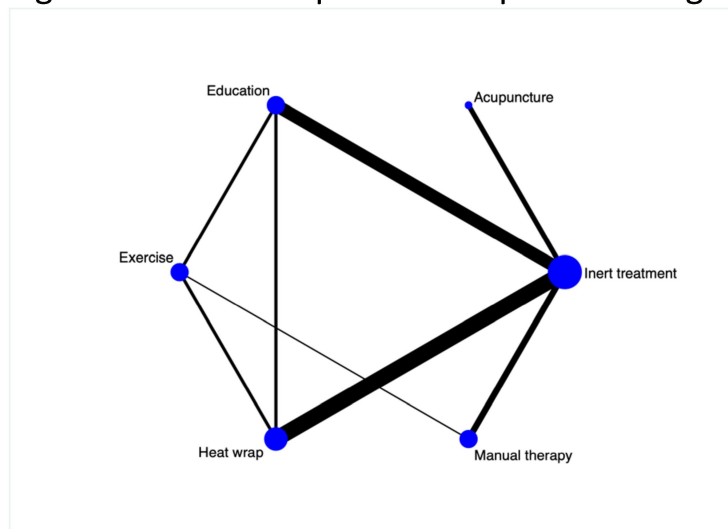


Supplementary K. Subgroup analysis results

1. Subgroup meta-analysis (pharmacological and non-pharmacological)

Disability 1 week – non pharmacological treatments

Figure 1a. Network plot of non-pharmacological treatments



Testing for inconsistency: $\chi^2(2) = 3.19$; Prob > $\chi^2 = 0.2030$

Figure 2a. Network forest of non-pharmacological treatments

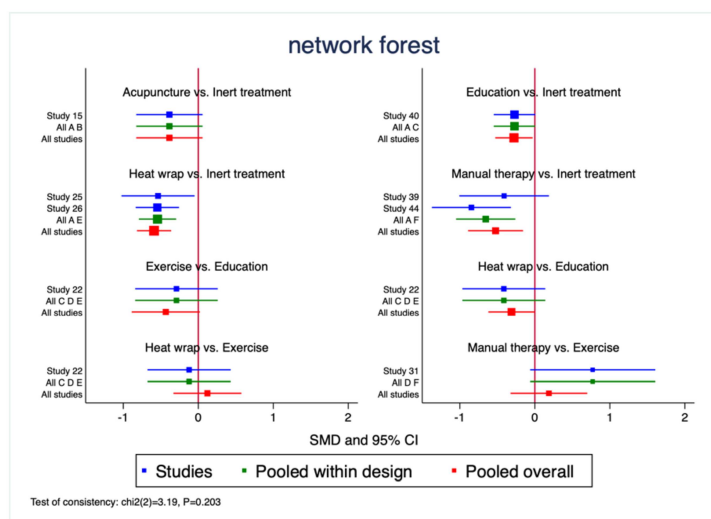


Table 1a. Netleague of non-pharmacological treatments

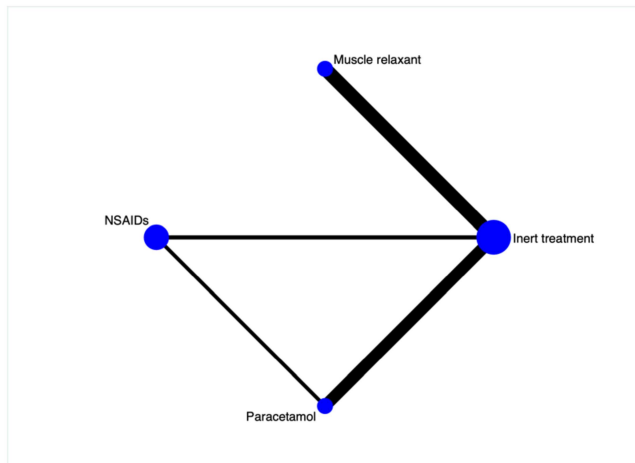
Inert treatment	-0.39 (-0.83,0.06)	-0.28 (-0.53,-0.03)	-0.71 (-1.16,-0.26)	-0.59 (-0.82,-0.36)	-0.52 (-0.89,-0.16)
0.39 (-0.06,0.83)	Acupuncture	0.11 (-0.40,0.61)	-0.33 (-0.96,0.30)	-0.20 (-0.70,0.29)	-0.14 (-0.71,0.44)
0.28 (0.03,0.53)	-0.11 (-0.61,0.40)	Education	-0.43 (-0.89,0.02)	-0.31 (-0.62,-0.00)	-0.25 (-0.68,0.19)
0.71 (0.26,1.16)	0.33 (-0.30,0.96)	0.43 (-0.02,0.89)	Exercise	0.12 (-0.33,0.57)	0.19 (-0.32,0.70)
0.59 (0.36,0.82)	0.20 (-0.29,0.70)	0.31 (0.00,0.62)	-0.12 (-0.57,0.33)	Heatwrap	0.07 (-0.36,0.49)
0.52 (0.16,0.89)	0.14 (-0.44,0.71)	0.25 (-0.19,0.68)	-0.19 (-0.70,0.32)	-0.07 (-0.49,0.36)	Manual therapy

Table 2a. SUCRA of non-pharmacological treatments

Treatment	SUCRA	PrBest	MeanRank
Manual therapy	80,3	43,6	2
Exercise	69,4	35,4	2,5
Heatwrap	67,9	12,6	2,6
Acupuncture	48,4	8,4	3,6
Education	31,2	0	4,4
Inert treatment	2,9	0	5,9

Disability 1 week – pharmacological treatments

Figure 1b. Network plot of pharmacological treatments



Testing for inconsistency: $\chi^2(1) = 2.14$; Prob > $\chi^2 = 0.1432$

Figure 2b. Network forest of pharmacological treatments

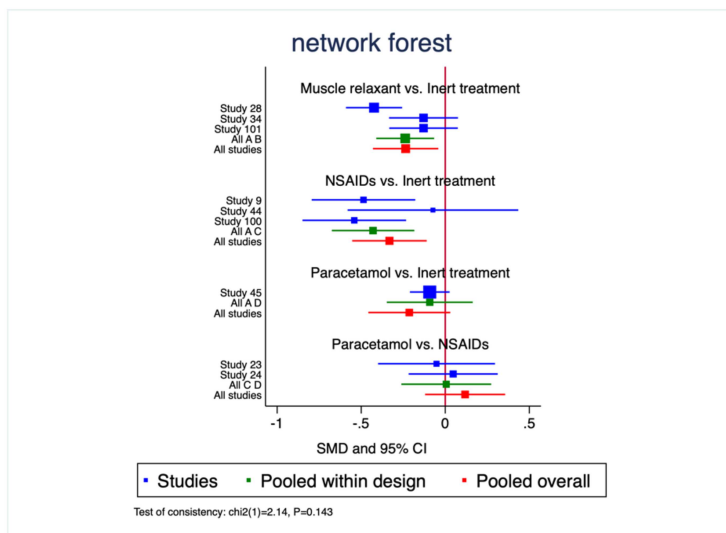


Table 1b. Netleague of pharmacological treatments

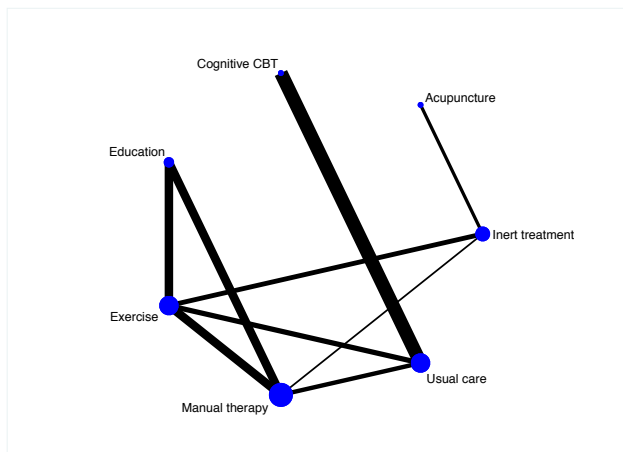
Inert treatment	-0.24 (-0.43,-0.04)	-0.33 (-0.55,-0.11)	-0.21 (-0.46,0.03)
0.24 (0.04,0.43)	Muscle relaxant	-0.10 (-0.39,0.20)	0.02 (-0.29,0.34)
0.33 (0.11,0.55)	0.10 (-0.20,0.39)	NSAIDs	0.12 (-0.12,0.36)
0.21 (-0.03,0.46)	-0.02 (-0.34,0.29)	-0.12 (-0.36,0.12)	Paracetamol

Table 2b. SUCRA of pharmacological treatments

Treatment	SUCRA	PrBest	MeanRank
NSAIDs	94,6	86	1,2
Muscle relaxant	64,1	11	2,1
Paracetamol	33,3	3	3
Inert treatment	7,9	0	3,8

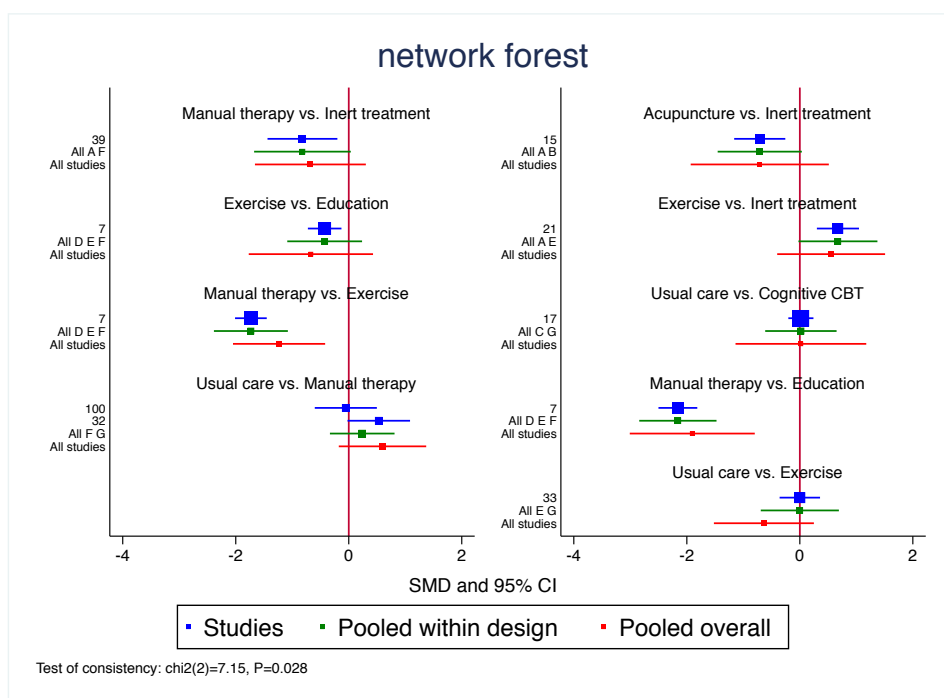
Disability 1 month – non pharmacological treatments

Figure 3a. Network plot of non-pharmacological treatments



Since we found sources of inconsistency (Prob > $\chi^2 = 0.0280$) in non-pharmacological network, we presented only pairwise meta-analyses and NMA

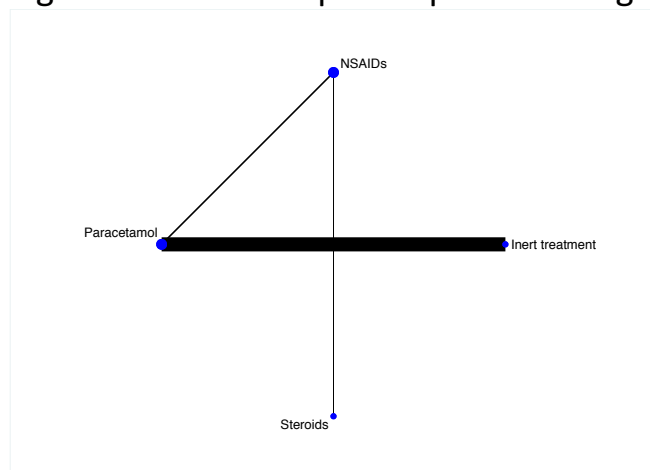
Figure 4a. Network forest of non-pharmacological treatments



Comparison	ES	[95% Conf.	Interval]	z	p value	I ²	Tau-squared
<i>Usual care-Manual Therapy</i>							
2 studies	-0.052	-0.601	0.497				
	0.531	-0.022	1.085				
overall	0.239	-0.333	0.81	z= 0.82	p = 0.413	53.5%	0.0910
<i>Acupuncture-Inert treatment</i>							
1 study	-0.709	-1.162	-0.257	z= 3.07	p = 0.002		
<i>Usual care-Cognitive CBT</i>							
1 study	0.019	-0.203	0.241	z= 0.17	p = 0.868		
<i>Exercise-Inert treatment</i>							
1 study	0.674	0.302	1.047	z= 3.55	p = 0.000		
<i>Usual care-Exercise</i>							
1 study	0	-0.358	0.358	z= 3.55	p = 0.000		
<i>Manual Therapy-Inert treatment</i>							
1 study	-0.819	-1.438	-0.201	z= 2.60	p = 0.009		
<i>Exercise-Education</i>							
1 study	-0.426	-0.723	-0.129	z= 2.81	p = 0.005		
<i>Manual Therapy - Education</i>							
1 study	-2.158	-2.502	-1.815	z= 12.31	p = 0.000		
<i>Manual Therapy-Exercise</i>							
1 study	-1.732	-2.012	-1.452	z= 12.10	p = 0.000		

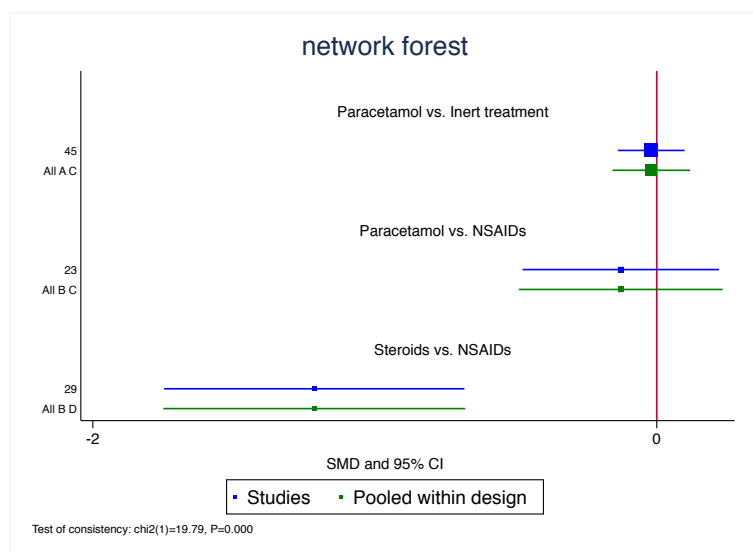
Disability 1 month – pharmacological treatments

Figure 3b. Network plot of pharmacological treatments



Since we found sources of inconsistency (Prob > chi2 = 0.000) in non-pharmacological network, we presented only pairwise meta-analyses and NMA

Figure 4b. Network forest of pharmacological treatments



Comparisons	ES	[95% Conf. Interval]	z	p-value
<i>Paracetamol-NSAIDs</i>				
1 study	-0.128	-0.476 0.22	z= 0.72	p = 0.472
<i>Steroids-NSAIDs</i>				
1 study	-1.215	-1.747 -0.682	z= 4.47	p = 0.000
<i>Paracetamol-Inert treatment</i>				
1 study	-0.019	-0.137 0.099	z= 0.32	p = 0.747

Supplementary L. Network meta-analysis results- Interval plot

Figure 1. Interval Plot -Network Meta-Analyses – Pain outcome

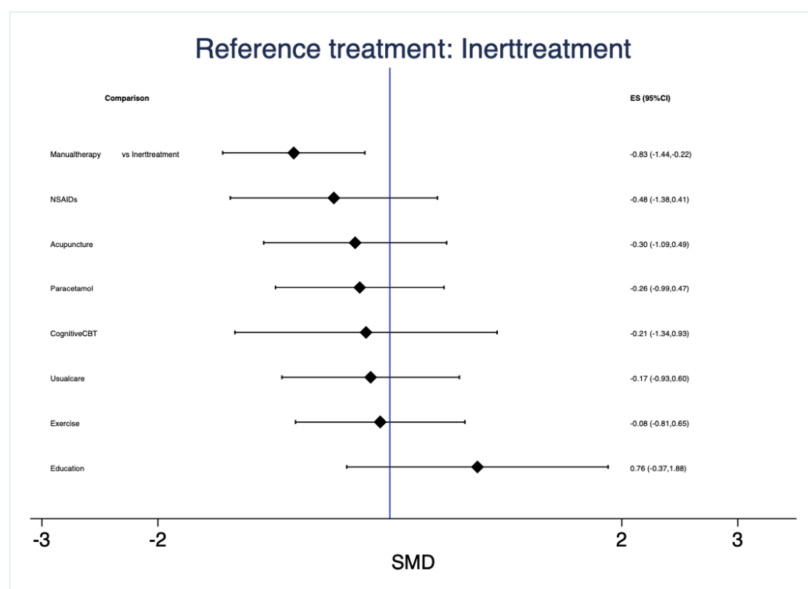


Figure 1a. Interval plot all treatments against inert treatment for pain outcome at 1 month of FU

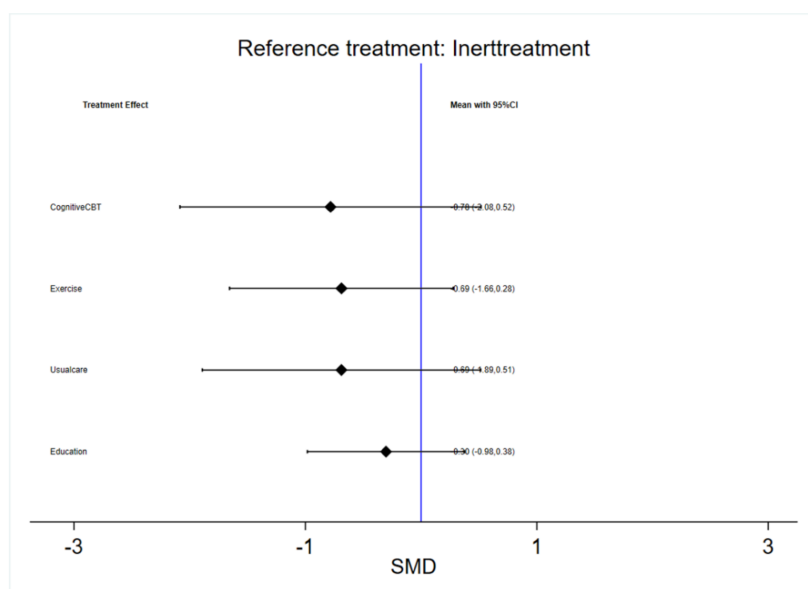


Figure 1b. Interval plot all treatments against inert treatment for pain outcome at 12 months of FU

Figure 2. Interval Plot -Network Meta-Analyses – Disability Outcome

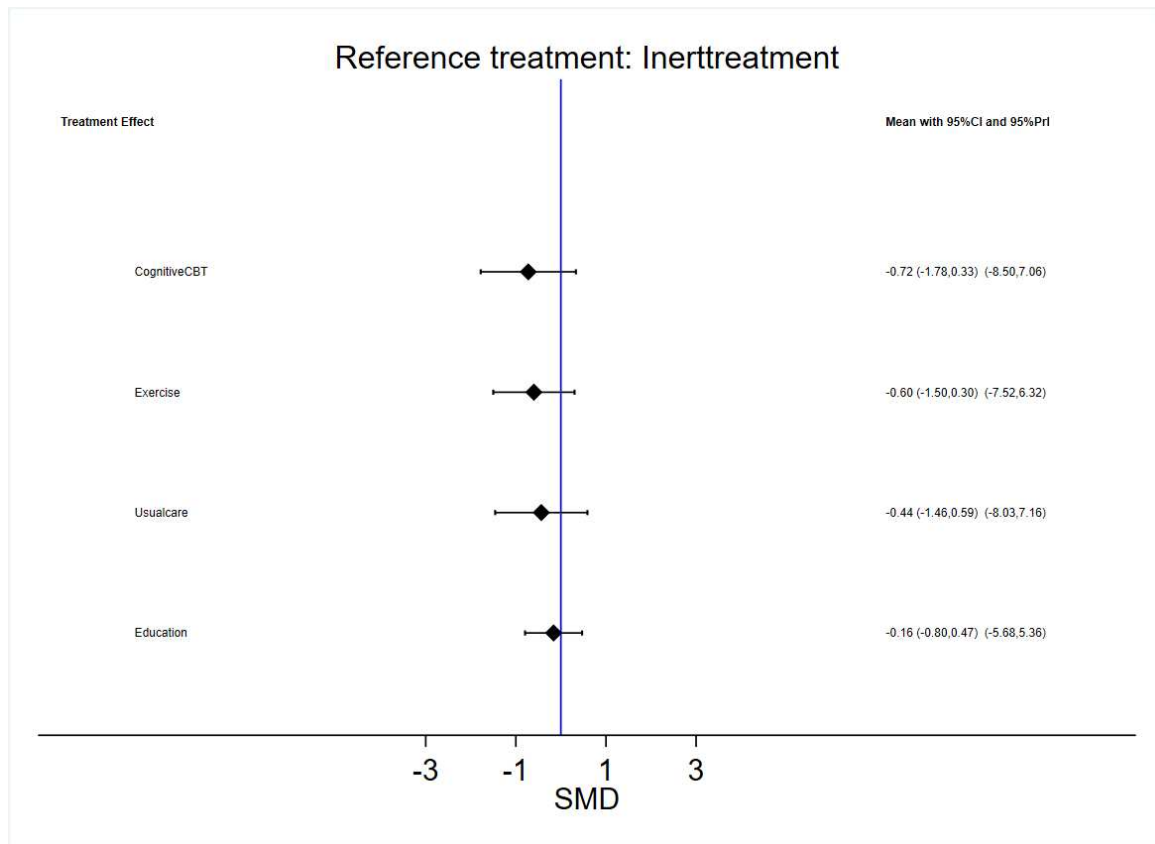


Figure 2a. Interval plot all treatments against inert treatment for disability outcome at 12 months of FU

Supplement M. All treatments against all treatments

Table 1. League table - pain

Table 1a. League table pain 1 month

Inert treatment	-0.30 (-1.09,0.49)	-0.21 (-1.34,0.93)	0.76 (-0.37,1.88)	-0.08 (-0.81,0.65)	-0.83 (-1.44,-0.22)	-0.48 (-1.38,0.41)	-0.26 (-0.99,0.47)	-0.17 (-0.93,0.60)
0.30 (-0.49,1.09)	Acupuncture	0.09 (-1.28,1.47)	1.05 (-0.32,2.43)	0.22 (-0.86,1.29)	-0.53 (-1.53,0.47)	-0.18 (-1.00,0.63)	0.04 (-0.85,0.93)	0.13 (-0.96,1.23)
0.21 (-0.93,1.34)	-0.09 (-1.47,1.28)	Cognitive CBT	0.96 (-0.44,2.36)	0.12 (-0.98,1.23)	-0.62 (-1.66,0.42)	-0.28 (-1.71,1.16)	-0.05 (-1.40,1.29)	0.04 (-0.79,0.87)
-0.76 (-1.88,0.37)	-1.05 (-2.43,0.32)	-0.96 (-2.36,0.44)	Education	-0.84 (-1.70,0.02)	-1.58 (-2.75,-0.42)	-1.24 (-2.67,0.20)	-1.02 (-2.35,0.32)	-0.92 (-2.05,0.21)
0.08 (-0.65,0.81)	-0.22 (-1.29,0.86)	-0.12 (-1.23,0.98)	0.84 (-0.02,1.70)	Exercise	-0.75 (-1.53,0.04)	-0.40 (-1.55,0.75)	-0.18 (-1.21,0.85)	-0.08 (-0.81,0.65)
0.83 (0.22,1.44)	0.53 (-0.47,1.53)	0.62 (-0.42,1.66)	1.58 (0.42,2.75)	0.75 (-0.04,1.53)	Manual therapy	0.35 (-0.73,1.42)	0.57 (-0.38,1.51)	0.66 (0.04,1.29)
0.48 (-0.41,1.38)	0.18 (-0.63,1.00)	0.28 (-1.16,1.71)	1.24 (-0.20,2.67)	0.40 (-0.75,1.55)	-0.35 (-1.42,0.73)	NSAIDs	0.22 (-0.54,0.99)	0.32 (-0.85,1.49)
0.26 (-0.47,0.99)	-0.04 (-0.93,0.85)	0.05 (-1.29,1.40)	1.02 (-0.32,2.35)	0.18 (-0.85,1.21)	-0.57 (-1.51,0.38)	-0.22 (-0.99,0.54)	Paracetamol	0.09 (-0.96,1.15)
0.17 (-0.60,0.93)	-0.13 (-1.23,0.96)	-0.04 (-0.87,0.79)	0.92 (-0.21,2.05)	0.08 (-0.65,0.81)	-0.66 (-1.29,-0.04)	-0.32 (-1.49,0.85)	-0.09 (-1.15,0.96)	Usual care

Table 1b. League table pain 12 months

Inert treatment	-0.69 (-1.89,0.51)	-0.69 (-1.66,0.28)	-0.30 (-0.98,0.38)	-0.78 (-2.08,0.52)
0.69 (-0.51,1.89)	Usual care	-0.00 (-0.72,0.72)	0.39 (-0.61,1.38)	-0.09 (-0.59,0.40)
0.69 (-0.28,1.66)	0.00 (-0.72,0.72)	Exercise	0.39 (-0.30,1.07)	-0.09 (-0.96,0.78)
0.30 (-0.38,0.98)	-0.39 (-1.38,0.61)	-0.39 (-1.07,0.30)	Education	-0.48 (-1.59,0.63)
0.78 (-0.52,2.08)	0.09 (-0.40,0.59)	0.09 (-0.78,0.96)	0.48 (-0.63,1.59)	Cognitive CBT

Table 2. Pain SUCRA

1 week of FU (immediate-term)			
Treatment	SUCRA	PrBest	MeanRank
Exercise	89,2	40,8	2
Heat wrap	85,8	45,2	2,3
Opioid	68,6	9,6	3,8
Manual therapy	60	1,4	4,6
Muscle relaxant	50,2	2	5,5
NSAIDs	47,9	0,2	5,7
Paracetamol	40,7	0,6	6,3
Education	25,1	0	7,7
Acupuncture	21,8	0,2	8
Inert treatment	10,7	0	9
1 month of FU (short-term)			
Treatment	SUCRA	PrBest	MeanRank
Manual therapy	91,1	57,2	1,7
NSAIDs	71,4	20,8	3,3
Acupuncture	55,7	7,4	4,5
Paracetamol	55,3	5	4,6
Cognitive CBT	50,8	8,6	4,9
Usual care	46,3	0,2	5,3
Exercise	40,3	0,6	5,8
Inert treatment	34,2	0	6,3
Education	4,9	0,2	8,6
12 months (long term)			
Treatment	SUCRA	PrBest	MeanRank
Cognitive CBT	73.7	45.0	2.1
Exercise	66.0	26.0	2.4
Usual care	61.4	16.8	2.5
Education	33.6	8.4	3.7
Inert treatment	15.3	3.8	4.4

Figure 1. Cumulative ranking curve of pain 1 week

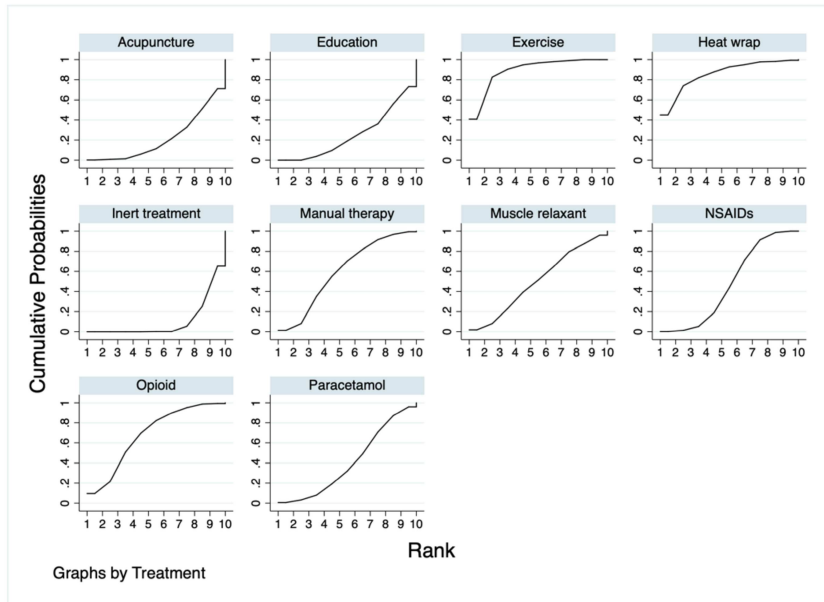


Figure 2. Cumulative ranking curve of pain 1 month

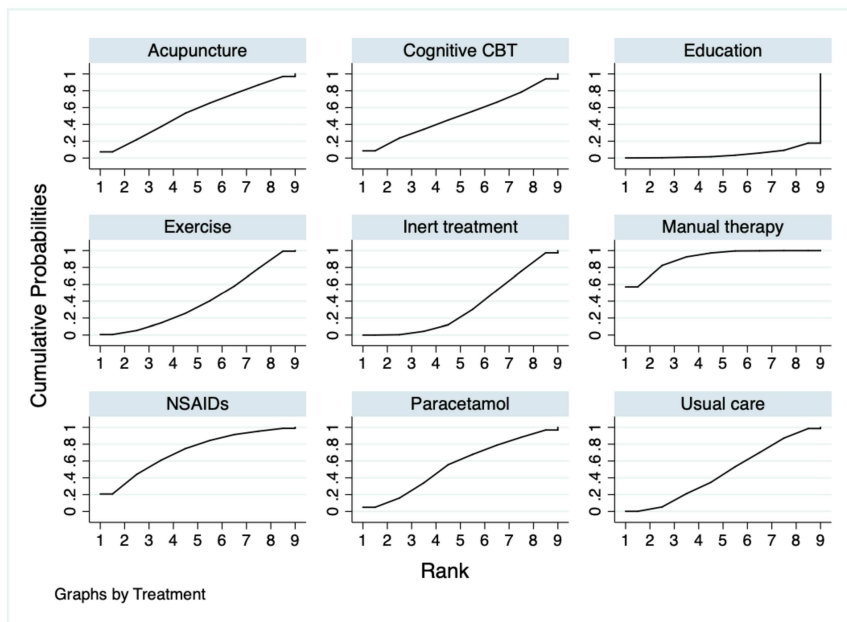


Figure 3. Cumulative ranking curve of pain 12 months

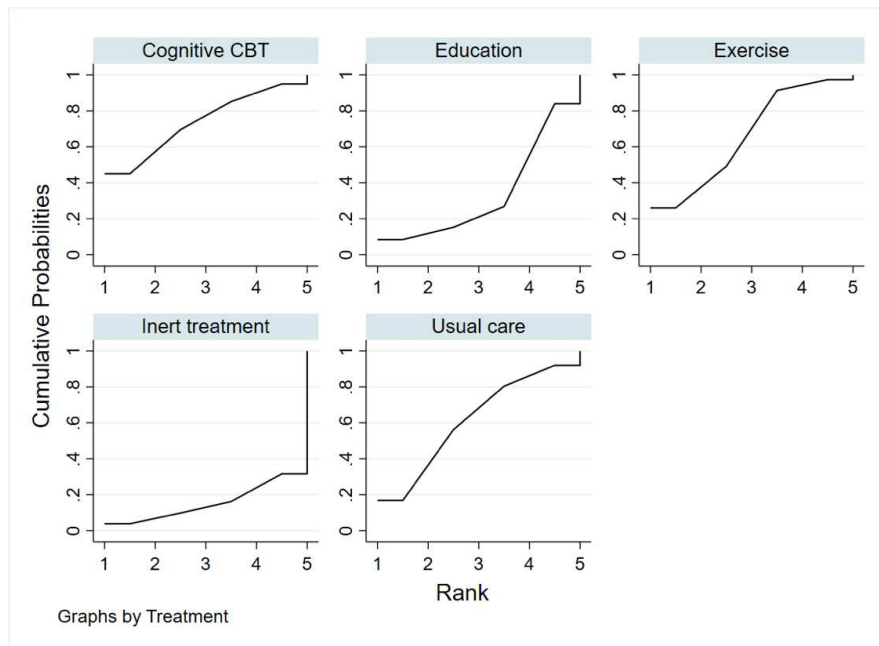


Table 3. League table - disability

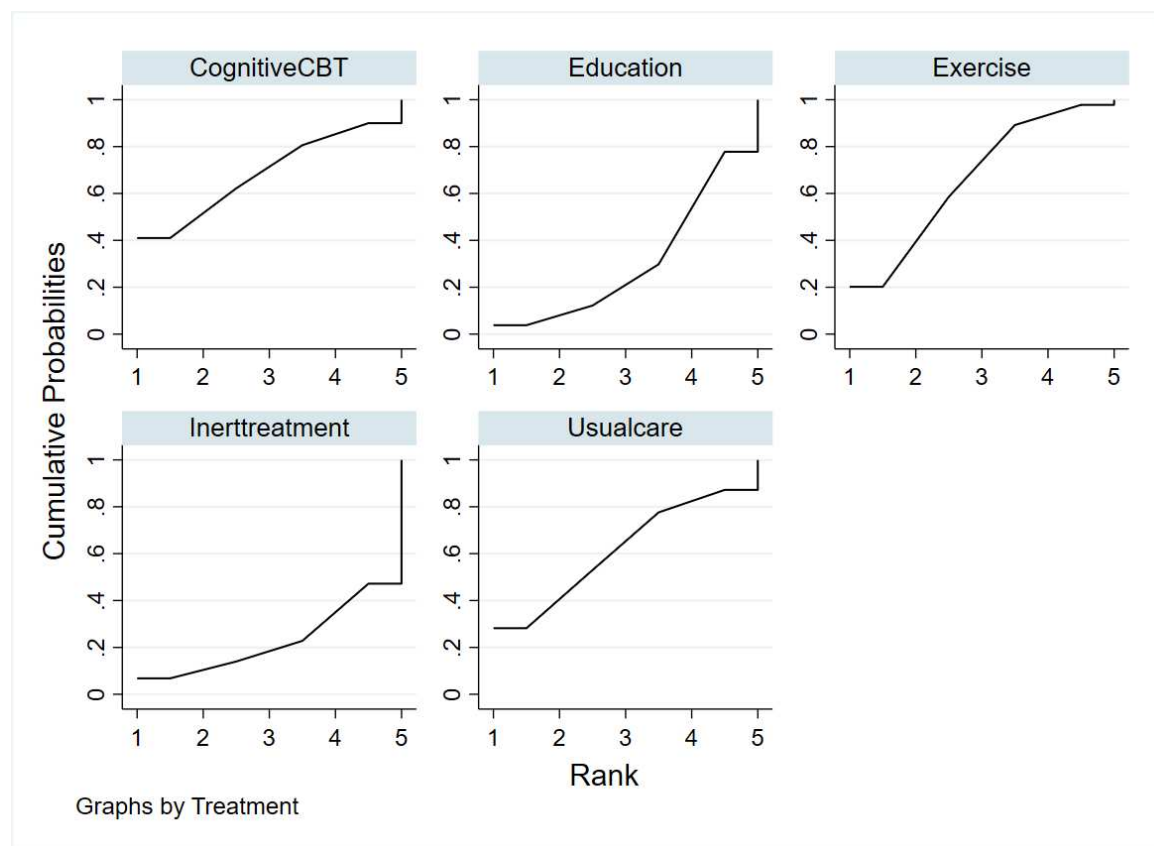
Table 3a. League table disability 12 months

Inert treatment	-0.44 (-1.46,0.59)	-0.60 (-1.50,0.30)	-0.16 (-0.80,0.47)	-0.72 (-1.78,0.33)
0.44 (-0.59,1.46)	Usual care	-0.16 (-0.65,0.32)	0.27 (-0.53,1.08)	-0.29 (-0.68,0.10)
0.60 (-0.30,1.50)	0.16 (-0.32,0.65)	Exercise	0.44 (-0.21,1.08)	-0.12 (-0.67,0.42)
0.16 (-0.47,0.80)	-0.27 (-1.08,0.53)	-0.44 (-1.08,0.21)	Education	-0.56 (-1.41,0.28)
0.72 (-0.33,1.78)	0.29 (-0.10,0.68)	0.12 (-0.42,0.67)	0.56 (-0.28,1.41)	Cognitive CBT

Table 4. Disability SUCRA

12 month of FU (long term)			
Treatments	SUCRA	PrBest	MeanRank
Cognitive CBT	68.5	41	2.3
Exercise	66.5	20.2	2.3
Usual care	61.5	28.2	2.5
Education	30.9	3.8	3.8
Inert treatment	22.7	6.8	4.1

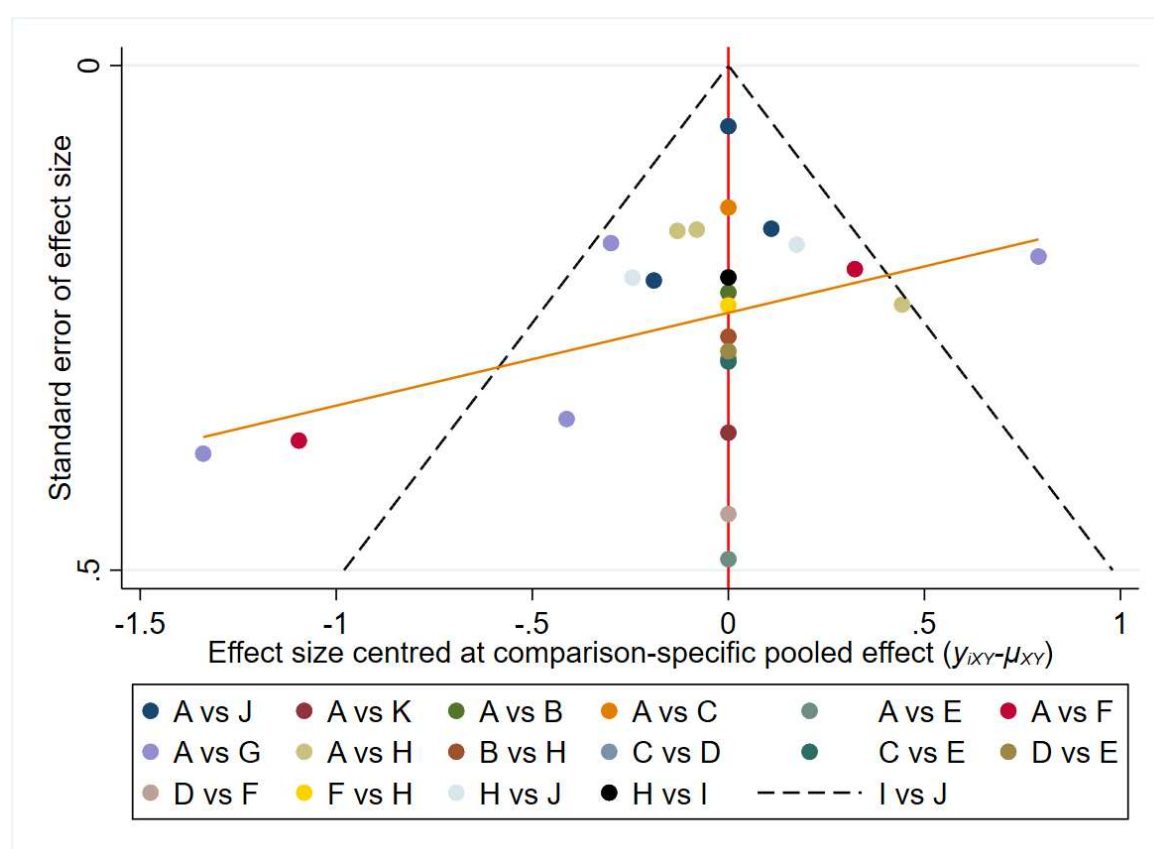
Figure 4. Cumulative ranking curve of disability 12 months



Supplement N. Funnel Plot

Funnel plot asymmetry was used to assess publication bias containing 10 or more trials reporting the outcome of interest. Thus, this was possible only for pain and disability outcomes at 1 week and 1 month of follow-up.

Figure 1. Funnel plot-pain



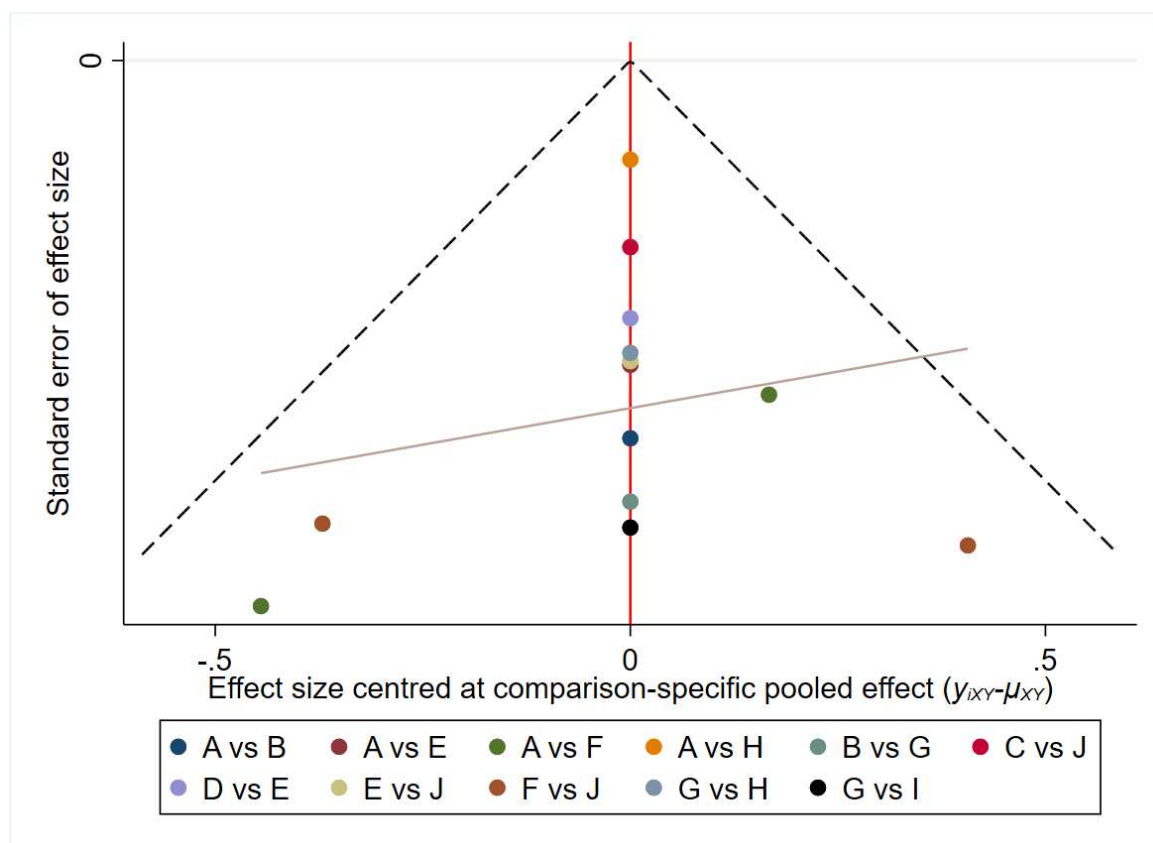
The red line represents the null hypothesis that the study-specific effect sizes do not differ from the respective comparison-specific pooled effect estimates. The orange line is the regression line.

Figure 1a. Pain Outcome 1 week

legend: Treatments used

- A (reference): Inert treatment
- B: Acupuncture
- C: Education
- D: Exercise
- E: Heat wrap
- F: Manual therapy
- G: Muscle relaxant

H: NSAIDs
 I: Opioid
 J: Paracetamol
 K: Physical therapy



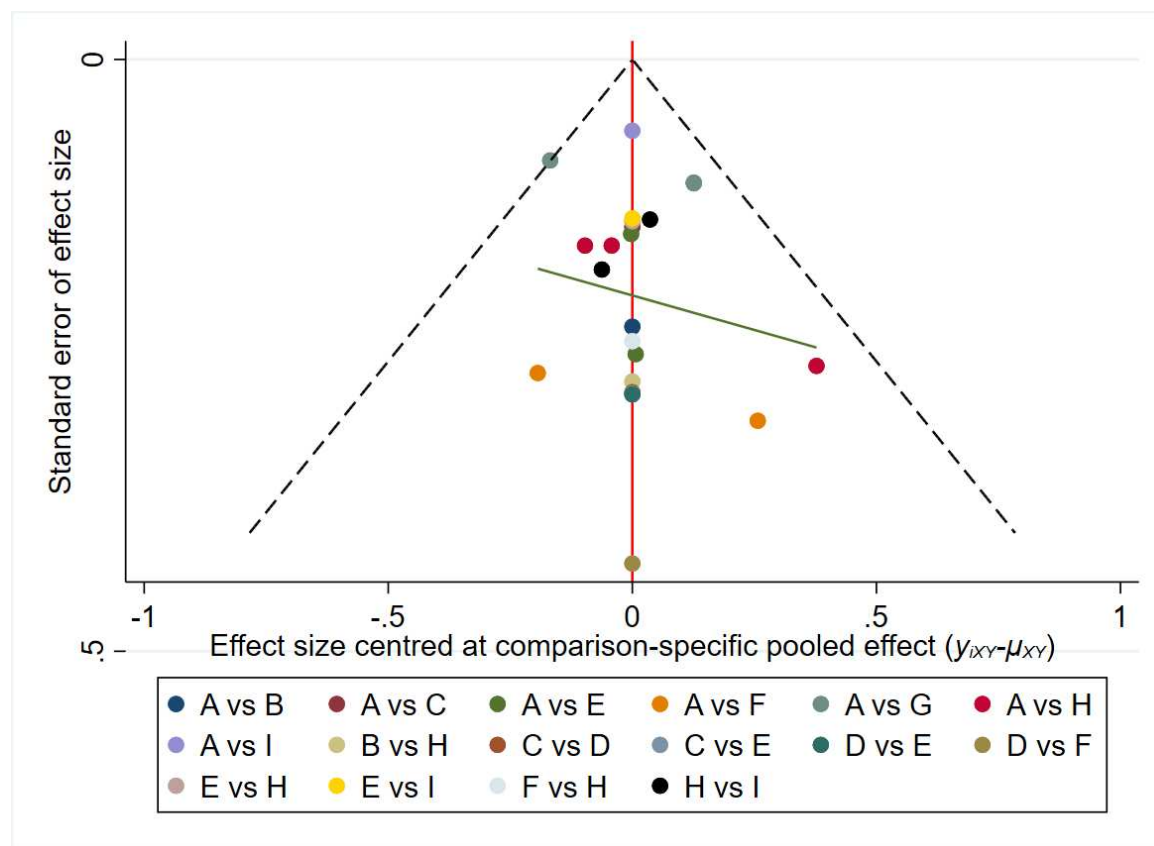
The red line represents the null hypothesis that the study-specific effect sizes do not differ from the respective comparison-specific pooled effect estimates. The gray line is the regression line.

Figure 1b. Pain Outcome 1 month

Legend: Treatments used

A (reference): Inert treatment
 B: Acupuncture
 C: Cognitive CBT
 D: Education
 E: Exercise
 F: Manual therapy
 G: NSAIDs
 H: Paracetamol
 I: Steroids
 J: Usual care

Figure 2. Funnel plot- disability



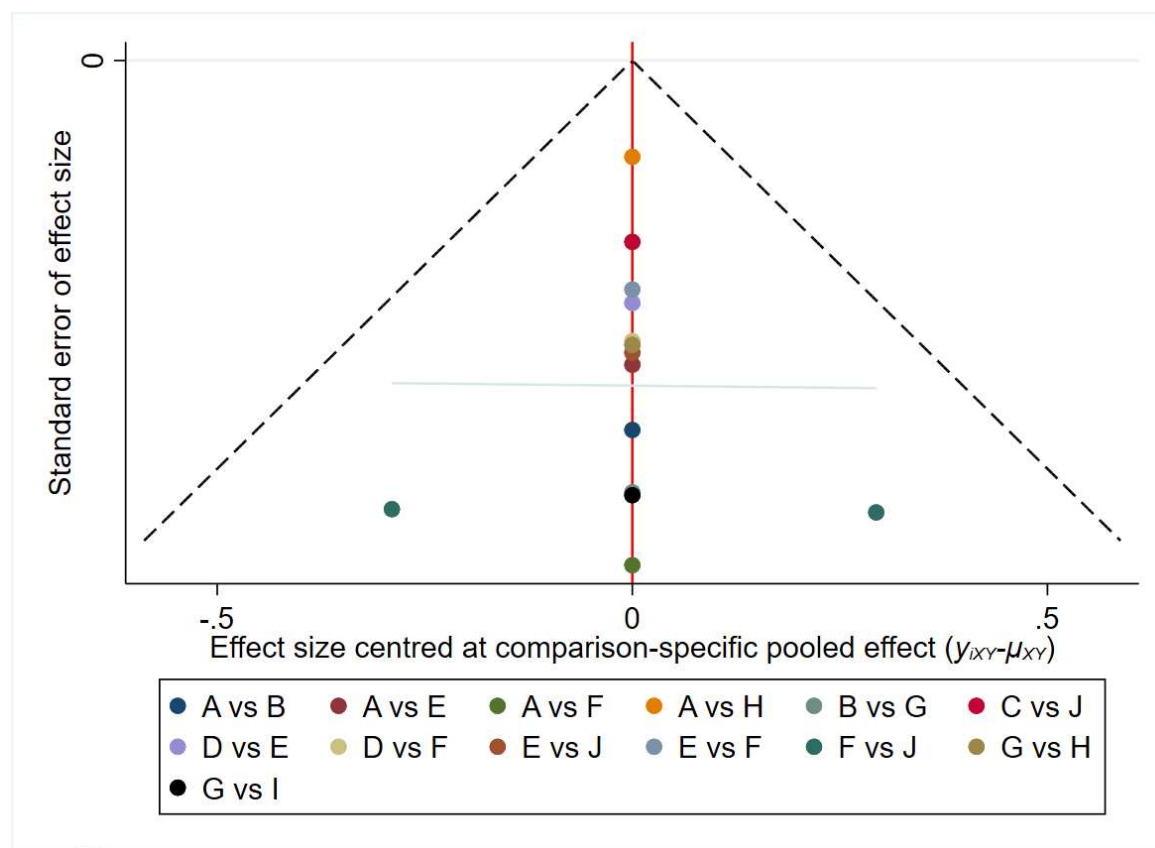
The red line represents the null hypothesis that the study-specific effect sizes do not differ from the respective comparison-specific pooled effect estimates. The green line is the regression line.

Figure 2a. Disability Outcome 1 week

Legend:

Treatments used

A (reference):	Inert treatment
B:	Acupuncture
C:	Education
D:	Exercise
E:	Heat wrap
F:	Manual therapy
G:	Muscle relaxant
H:	NSAIDs
I:	Paracetamol



The red line represents the null hypothesis that the study-specific effect sizes do not differ from the respective comparison-specific pooled effect estimates. The gray line is the regression line.

Figure 2b. Disability Outcome 1 month

Legend: Treatments used

- A (reference): Inert treatment
- B: Acupuncture
- C: Cognitive CBT
- D: Education
- E: Exercise
- F: Manual therapy
- G: NSAIDs
- H: Paracetamol
- I: Steroids
- J: Usual care

Supplement O. Contribution matrix for the network on interventions

Figure 1. Contribution matrix for the network on interventions - Pain

		Direct comparisons in the network																	
		AvsB	AvsC	AvsE	AvsF	AvsG	AvsH	AvsJ	AvsK	BvsH	CvsD	CvsE	DvsE	DvsF	FvsH	HvsI	HvsJ	IvsJ	
Mixed estimates	AvsB	38.0	0.7	0.1	0.4		11.7	9.7		22.6	0.5	0.2	0.3	0.8	1.3	3.9	5.8	3.9	
	AvsC	0.4	72.4	4.4	0.5		1.6	1.5		0.4	4.3	4.3	0.1	4.2	3.7	0.9	0.9	0.6	
	AvsE	0.4	23.5	10.0	3.8		2.2	1.3		0.4	8.5	21.0	13.6	6.0	3.4	0.7	1.1	0.7	
	AvsF	2.7	6.0	1.1	3.8		13.5	11.2		2.7	4.3	1.7	2.8	7.1	27.4	4.5	6.7	4.5	
	AvsG					100.0													
	AvsH	5.4	1.6	0.3	1.0		26.5	22.0		5.4	1.1	0.4	0.7	1.9	2.9	8.8	13.2	8.8	
	AvsJ	1.0	0.3	0.1	0.2		5.0	81.9		1.0	0.2	0.1	0.1	0.4	0.5	2.6	3.9	2.6	
	AvsK								100.0										
	BvsH	27.7	0.8	0.2	0.5		14.3	11.9		24.3	0.6	0.2	0.4	1.0	1.5	4.7	7.1	4.7	
	CvsD	0.7	11.7	3.5	1.0		3.6	3.0		0.7	27.9	12.4	16.0	8.2	7.2	2.2	2.2	2.2	
	CvsE	0.3	13.7	10.0	0.4		1.6	1.3		0.3	14.4	31.0	18.1	3.7	3.2	0.5	0.8	0.5	
	DvsE	0.6	0.4	6.3	0.7		2.5	2.1		0.6	17.9	17.5	37.6	5.9	5.2	0.8	1.3	0.8	
	DvsF	1.3	15.8	4.4	4.7		6.6	5.5		1.3	11.4	4.4	7.6	12.2	16.4	2.7	4.0	2.7	
	FvsH	1.8	0.5	0.1	0.3		8.9	5.5		1.8	5.4	2.1	3.4	8.8	44.3	2.2	3.3	2.2	
	HvsI	1.8	0.5	0.1	0.3		8.9	5.5		1.8	5.4	2.1	3.4	8.8	44.3	2.2	3.3	2.2	
	HvsJ	4.3	1.6	0.2	0.8		21.4	29.1		4.3	0.9	0.4	0.6	1.5	2.2	48.8	6.0	17.7	
	IvsJ	2.7	0.6	0.1	0.5		13.5	17.7		2.7	0.6	0.2	0.4	1.0	1.5	2.2	14.5	9.7	
Network meta-analysis estimates	AvsD	0.7	26.8	4.9	1.0		3.6	3.0		0.7	19.4	7.5	12.4	8.4	7.4	1.2	1.8	1.2	
	AvsI	2.9	0.9	0.2	0.5		14.5	24.5		2.9	0.6	0.2	0.4	1.0	1.6	25.2	6.2	19.3	
	BvsC	23.1	32.7	20.0	0.5		6.3	5.2		14.0	2.3	2.1	0.1	2.4	2.4	2.1	3.1	2.1	
	BvsD	17.4	18.8	6.4			2.8	2.4		11.2	5.2	5.2	8.7	3.6	3.4	1.0	1.0	1.0	
	BvsE	17.4	18.8	6.4			2.8	2.4		11.2	5.2	5.2	8.7	3.6	3.4	1.0	1.0	1.0	
	BvsF	19.4	5.1	0.6	3.2		3.6	4.6		15.5	3.7	1.4	2.3	6.0	28.8	1.9	2.8	1.9	
	BvsG	23.7	0.4	0.1	0.3		7.3	6.0		14.1	0.3	0.1	0.2	0.5	0.8	2.4	3.6	2.4	
	BvsI	21.2	0.3	0.1	0.2		5.3	15.3		16.7	0.2	0.1	0.1	0.4	0.6	24.3	1.7	13.5	
	BvsJ	21.0	0.4	0.1	0.2		6.0	33.6		16.9	0.3	0.1	0.2	0.4	0.6	4.1	6.2	4.1	
	BvsK	23.7	0.4	0.1	0.3		7.3	6.0		14.1	0.3	0.1	0.2	0.5	0.8	2.4	3.6	2.4	
	CvsF	2.1	22.9	0.6	2.3		10.3	8.5		2.1	35.1	3.0	2.1	2.9	3.4	0.4	0.5	0.4	
	CvsG	0.2	48.3	0.3	0.4		1.0	0.8		0.2	35.1	0.4	0.1	0.1	0.1	0.3	0.3	0.3	
	CvsH	3.2	31.3	0.4	0.4		16.1	13.4		3.2	2.6	2.2	2.2	3.0	3.0	5.4	8.0	5.4	
	CvsI	1.9	26.2	1.5	0.2		9.3	16.4		1.9	20.0	1.8	0.2	2.2	2.4	17.6	3.9	12.4	
	CvsJ	0.3	36.6	0.3	0.3		1.0	49.5		0.3	2.4	2.4	0.1	2.5	2.3	1.7	0.5	1.7	
	CvsK	0.2	40.0	0.3	0.3		1.0	49.5		0.3	2.4	2.4	0.1	2.5	2.3	1.7	0.5	1.7	
	DvsG	0.5	19.1	0.7	0.7		2.6	2.2		0.5	13.8	5.3	8.8	6.0	5.3	0.9	1.3	0.9	
	DvsH	2.2	18.3	0.4	0.2		10.9	9.1		2.2	13.4	5.2	7.0	6.8	3.6	3.6	5.4	3.6	
	DvsI	1.2	15.8	0.3	0.3		5.3	11.9		1.4	4.4	3.2	6.3	9.4	14.8	2.4	4.9	2.8	
	DvsJ	0.2	20.2	0.7	0.7		5.0	25.8		0.2	14.6	5.6	9.3	6.5	5.8	1.8	1.8	1.8	
	DvsK	0.5	19.1	0.7	0.7		2.6	2.2		0.5	13.8	5.3	8.8	6.0	5.3	0.9	1.3	0.9	
	EvsF	1.7	14.0	0.3	0.3		8.4	7.0		1.7	11.8	10.4	17.1	11.1	17.1	2.8	4.2	2.8	
	EvsG	0.3	20.4	0.4	0.4		11.9	1.3		0.3	5.9	14.6	9.4	3.5	3.5	0.8	0.8	0.8	
	EvsH	2.4	18.0	0.1	0.1		11.9	28.7		2.4	4.9	13.1	9.0	4.1	4.3	3.9	5.9	3.9	
	EvsI	1.4	19.3	0.3	0.3		6.9	12.7		1.4	4.5	11.3	7.6	3.3	3.3	14.4	2.8	9.8	
	EvsJ	0.3	20.4	0.4	0.4		11.9	1.3		0.3	5.9	14.6	9.4	3.5	3.5	0.8	0.8	0.8	
EvsK	0.3	20.4	0.4	0.4		11.9	1.3		0.3	5.9	14.6	9.4	3.5	3.5	0.8	0.8	0.8		
FvsG	2.0	4.3	0.8	2.7		9.8	8.1		2.0	3.1	1.2	2.0	5.1	1.9	3.2	4.9	3.2		
FvsI	0.1	5.1	0.9	3.2		0.7	10.1		0.1	3.7	1.4	2.3	6.0	27.9	25.8	1.4	11.3		
FvsJ	0.1	5.1	0.9	3.2		0.7	10.1		0.1	3.7	1.4	2.3	6.0	27.9	25.8	1.4	11.3		
FvsK	2.0	4.3	0.8	2.7		9.8	8.1		2.0	3.1	1.2	2.0	5.1	1.9	3.2	4.9	3.2		
GvsH	3.4	10.0	0.6	0.6		36.2	14.1		3.4	0.7	0.3	0.3	0.7	1.1	5.6	8.4	5.6		
GvsI	2.0	0.6	0.1	0.4		39.3	10.1		2.0	0.4	0.2	0.3	0.7	1.1	17.5	4.3	12.8		
GvsJ	0.5	0.2	0.1	0.1		46.9	2.7		0.5	0.1	0.1	0.1	0.2	0.3	1.4	2.1	1.4		
GvsK	3.4	1.0	0.2	0.6		16.9	14.1		3.4	0.7	0.3	0.5	1.2	1.8	5.6	8.4	5.6		
HvsK	3.0	0.6	0.1	0.4		20.7	36.2		3.0	0.4	0.2	0.3	0.7	1.1	17.5	4.3	12.8		
IvsK	0.5	0.2	0.1	0.1		2.7	43.4		0.5	0.1	0.1	0.1	0.2	0.3	1.4	2.1	1.4		
Entire network		5.4	13.1	2.5	0.9	7.0	7.1	12.1	7.0	4.0	5.3	4.9	4.7	3.9	7.2	6.3	3.8	4.7	
Included studies		1	1	1	2	4	3	1	1	1	1	1	1	1	1	2	2	1	

Figure 1a. Contribution matrix for the network on interventions Pain Outcome 1 week

Label: direct comparisons in the network are presented in the columns, and their contributions to the combined treatment effect are presented in the rows. The entries of the matrix are the percentage weights attributed to each direct comparison. The intervention labels are: A (reference): Inert treatment; B:Acupuncture; C: Education; D: Exercise; E: Heat wrap; F: Manual therapy; G: Muscle relaxant; H: NSAIDs; I: Opioid; J: Paracetamol; K: Physical therapy

		Direct comparisons in the network											
		AvsB	AvsE	AvsF	AvsH	BvsG	CvsJ	DvsE	EvsJ	FvsJ	GvsH	GvsI	
Mixed estimates	AvsB	40.4											
	AvsE		72.8										
	AvsF			9.1									
	AvsH				44.6								
	BvsG	2.2				99.5	2.2						
	CvsJ	23.7				23.7	29.0						
	DvsE							100.0					
	EvsJ								100.0				
	FvsJ		8.9	8.9						73.4	8.9		
	GvsH		24.7	24.7						24.7	25.8		
	GvsI	14.2			14.2	14.2						57.5	
												100.0	
	Network meta-analysis estimates	AvsC		26.1	7.3				33.3		26.1	7.3	
		AvsD		40.0	5.0				45.0		5.0	5.0	
AvsG		11.0			39.0	11.0						39.0	
AvsI		7.3			26.0	7.3						26.0	
BvsC		14.4	39.1	10.9					39.1	10.9			
BvsD		17.3	16.8	4.7	7.1	7.1	21.5		16.8	4.7	7.1		
BvsE		23.3	22.9	2.8	8.5	8.5		25.8	2.8	2.8	8.5		
BvsF		20.7	30.9	3.8	11.4	11.5			3.8	3.8	11.4		
BvsH		32.4	9.0	21.8	10.2	10.2			9.0	9.0	10.2		
BvsI		15.5			32.4	17.6					17.6		
BvsJ		18.3			15.5	19.0					15.5		
CvsD			21.4	6.0		9.0			21.4	6.0		9.0	
CvsE			3.4	3.4			31.1	31.1	27.7	3.4			
CvsF			4.9	4.9			49.1		40.3	4.9			
CvsG			16.4	16.4			33.6		16.4	17.2			
CvsH		4.4	15.6	4.4	15.6	4.4	20.0		15.6	4.4	15.6		
CvsI		0.6	19.3	5.4	24.2	0.6	24.7		19.3	5.4	0.6		
CvsJ		3.7	13.0	3.6	13.0	3.7	16.7		13.0	3.6	13.0	16.7	
DvsF			19.9	19.9				33.3	13.4	13.4			
DvsG		5.2	21.1	2.6	18.5	5.2		23.7	2.6	2.6	18.5		
DvsH		0.7	27.2	3.4	29.9	0.7		30.6	3.4	3.4	0.7		
DvsI		4.2	17.0	2.1	14.9	4.2		19.2	2.1	2.1	14.9	19.2	
DvsJ			4.9	4.9				49.1	40.3	4.9			
EvsF			29.8	29.8					29.2	29.2			
EvsG		6.8	27.6	3.4	24.2	6.8			3.4	3.4	24.2		
EvsH		1.0	39.2	4.9	43.1	1.0			4.9	4.9	1.0		
EvsI		5.2	21.1	2.6	18.5	5.2			2.6	2.6	18.5	23.7	
FvsG		6.1	8.2	19.7	21.7	6.1			8.2	8.2	21.7		
FvsH		0.9	11.1	28.9	37.1	0.9			11.1	11.1	0.9		
FvsI		4.8	6.4	15.4	17.0	4.8			6.4	6.4	17.0	21.8	
GvsJ	5.5	19.5	5.5	19.5	5.5			19.5	5.5	19.5			
HvsI	8.2			8.3	8.2					33.5	41.7		
HvsJ	0.7	25.7	7.2	32.1	0.7			25.7	7.2	0.7			
IvsJ	4.4	15.6	4.4	15.6	4.4			15.6	4.4	15.6	20.0		
Entire network		6.8	16.1	7.3	14.9	5.1	7.0	7.0	12.2	5.4	11.1	7.0	
Included studies		1	1	2	1	1	1	1	1	2	1	1	

Figure 1b. Contribution matrix for the network on interventions Pain Outcome 1 month

Label: direct comparisons in the network are presented in the columns, and their contributions to the combined treatment effect are presented in the rows. The entries of the matrix are the percentage weights attributed to each direct comparison. The intervention labels are: A (reference): Inert treatment; B: Acupuncture; C: Cognitive CBT; D: Education; E: Exercise; F: Manual therapy; G: NSAIDs; H: Paracetamol; I: Steroids; J: Usual care

			Direct comparisons in the network			
			AvsC	BvsE	CvsD	DvsE
Network meta-analysis estimates	Mixed estimates					
		AvsC	100.0	-	-	-
		BvsE	-	100.0	-	-
		CvsD	-	-	100.0	-
		DvsE	-	-	-	100.0

	Indirect estimates					
		AvsB	25.0	25.0	25.0	25.0
		AvsD	50.0	-	50.0	-
		AvsE	33.3	-	33.3	33.3
		BvsC	-	33.3	33.3	33.3
		BvsD	-	50.0	-	50.0
	CvsE	-	-	50.0	50.0	
Entire network			20.0	20.0	30.0	30.0
Included studies			1	2	1	1

Figure 1c. Contribution matrix for the network on interventions Pain Outcome 12 months

Label: direct comparisons in the network are presented in the columns, and their contributions to the combined treatment effect are presented in the rows. The entries of the matrix are the percentage weights attributed to each direct comparison. The intervention labels are: A (reference): Inert treatment; B: Cognitive CBT; C: Education; D: Exercise; E: Usual care

		Direct comparisons in the network				
		AvsC	BvsE	CvsD	DvsE	
Network meta-analysis estimates	Mixed estimates	AvsC	100.0			
		BvsE		100.0		
		CvsD			100.0	
		DvsE				100.0
	Indirect estimates	AvsB	25.0	25.0	25.0	25.0
		AvsD	50.0		50.0	
		AvsE	33.3		33.3	33.3
		BvsC		33.3	33.3	33.3
		BvsD		50.0		50.0
		CvsE			50.0	50.0
Entire network		20.0	20.0	30.0	30.0	
Included studies		1	2	1	1	

Figure 2. Contribution matrix for the network on interventions - Disability

Figure 2a. Contribution matrix for the network on interventions Disability Outcome 12 months

Label: direct comparisons in the network are presented in the columns, and their contributions to the combined treatment effect are presented in the rows. The entries of the matrix are the percentage weights attributed to each direct comparison. The intervention labels are: The intervention labels are: A (reference); B: Cognitive CBT; C: Education; D: Exercise; E: Usual care

		Direct comparisons in the network				
		AvsC	BvsD	BvsE	CvsD	DvsE
Network meta-analysis estimates	Mixed estimates					
	AvsC	100.0	-	-	-	-
	BvsD	-	46.2	26.9	-	26.9
	BvsE	-	25.6	48.8	-	25.6
	CvsD	-	-	-	100.0	-
	DvsE	-	22.3	22.3	-	55.3
	Indirect estimates					
	AvsB	29.7	18.7	10.9	29.7	10.9
	AvsD	50.0	-	-	50.0	-
	AvsE	30.4	8.8	8.8	30.4	21.7
	BvsC	-	26.7	15.6	42.2	15.6
CvsE	-	12.6	12.6	43.7	31.1	
Entire network		20.7	16.1	13.6	31.1	18.6
Included studies		1	1	3	1	2

Supplement P. GRADE for Pain Outcome

Introduction

CINeMA²⁶ considers 6 domains: (i) within-study bias, (ii) reporting bias, (iii) indirectness, (iv) imprecision, (v) heterogeneity, and (vi) incoherence. Features include the percentage contribution matrix, relative treatment effects for each comparison, estimation of the heterogeneity variance, prediction intervals, and tests for the evaluation of the assumption of coherence. In evaluating imprecision, heterogeneity, and incoherence, we consider the impact of these components of variability in forming clinical decisions.

Table of reasons for downgrading

We use the CINeMA software for GRADE assessment.^{26 27} We downgrade network estimate according to the following criteria.

(1) Study limitations: We downgraded by one level when the contributions from low RoB comparisons were less than 25% and contributions from moderate or high RoB comparisons were 75% or greater.

(2) Imprecision: We considered a clinically meaningful threshold for SMD to be 0.5²⁸ and downgraded the estimate if the SMD point estimate is 0 or more and the lower limit of its CrI is below 0.5; or if the SMD point estimate is less than 0 and the upper limit of its CrI is above 0.5.

(3) Inconsistency: We rated two concepts, heterogeneity and incoherence (inconsistency), in this domain.

For heterogeneity, we looked at the common tau and found that it is low compared to the expected value as reported in the literature,²⁹ so we did not downgrade any network estimate for heterogeneity. For inconsistency, we looked at the results of side splitting and we downgraded the comparisons with important inconsistency ($p < 0.10$), where we have not downgraded for imprecision (we did not downgrade the same network estimate for both imprecision and inconsistency).

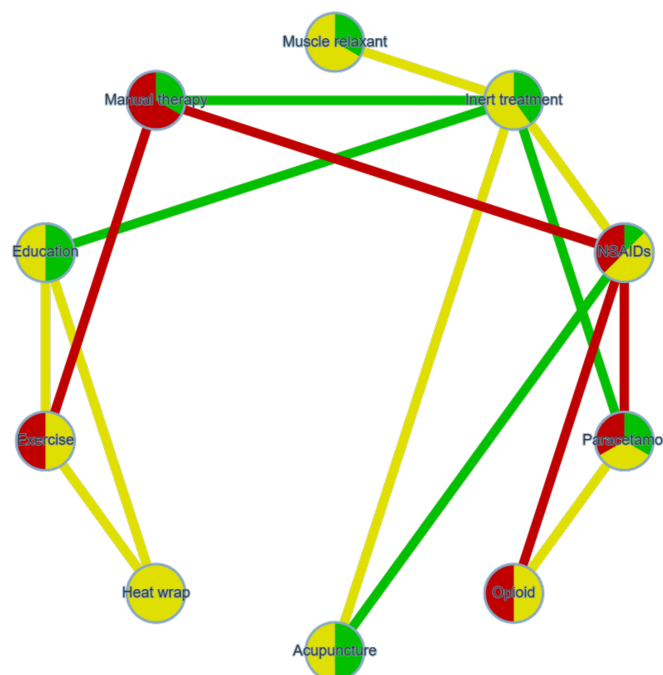
(4) Indirectness: We have assured transitivity in our network by limiting the included studies to acute and subacute population and to non-mixed treatments for NS-LBP. Thus, we did not downgrade for indirectness.

(5) Reporting bias: We cannot completely rule out the possibility that some studies are still missing. However, we assumed that publication bias was undetected.

1) Pain at 1 week

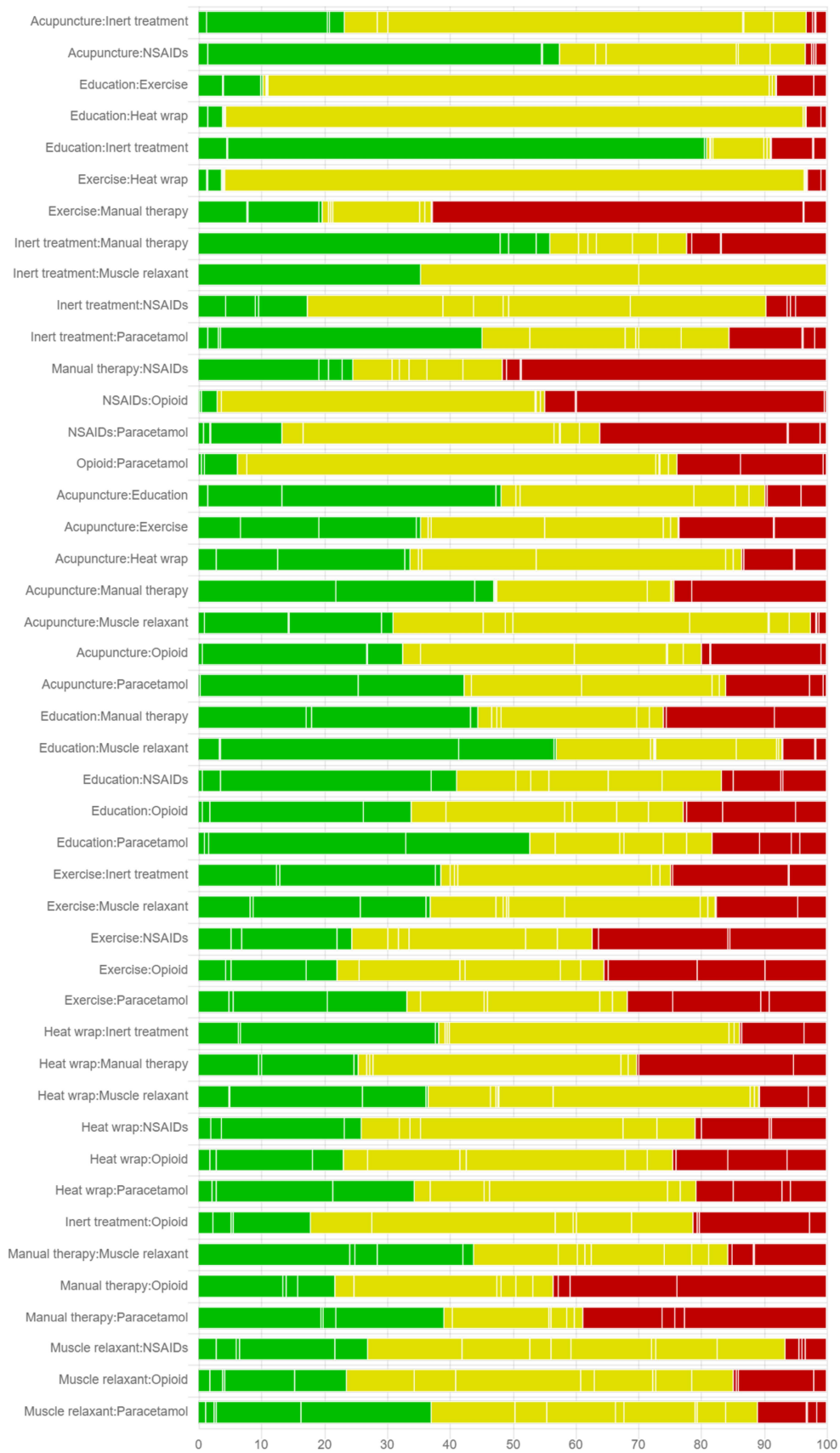
1) Summary of study limitations of the included studies

The colours in the circles indicate the percentage of low RoB studies [green], moderate RoB studies [yellow] and high RoB studies [red] involving each intervention. The colours of the line then indicate the average RoB assessment of each comparison based on the above information – low RoB comparison [green], moderate RoB comparison [yellow] and high RoB comparison [red].



2) Contribution of low or moderate RoB comparisons to each network estimate

Based on the above assessment of RoB for each comparison and the contribution matrix detailing contribution of each direct comparison to all network estimates, the following bar graphs show the percentage of low or moderate RoB contributions for each network estimate. The judgements about study limitations in each direct comparison is shown at the beginning of the graph. Each bar corresponds to a NMA relative treatment effect and shows how much information comes from comparisons at moderate risk of bias [yellow].



3) Summary grading of Evidence

Comparison	Number of studies	Within-study bias	Reporting bias	Indirectness	Imprecision	Heterogeneity	Incoherence	Confidence rating
Mixed evidence								
Acupuncture:Inert treatment	1	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Acupuncture:NSAIDs	1	No concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Moderate
Education:Exercise	1	Some concerns	Undetected	No concerns	No concerns	Some concerns	No concerns	Moderate
Education:Heat wrap	1	Some concerns	Undetected	No concerns	No concerns	Some concerns	No concerns	Moderate
Education:Inert treatment	1	No concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Low
Exercise:Heat wrap	1	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Exercise:Manual therapy	1	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Inert treatment:Manual therapy	1	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Inert treatment:Muscle relaxant	3	Some concerns	Undetected	No concerns	No concerns	Some concerns	No concerns	Moderate
Inert treatment:NSAIDs	3	Some concerns	Undetected	No concerns	No concerns	Some concerns	No concerns	Moderate
Inert treatment:Paracetamol	1	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Manual therapy:NSAIDs	1	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
NSAIDs:Opioid	2	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
NSAIDs:Paracetamol	2	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low

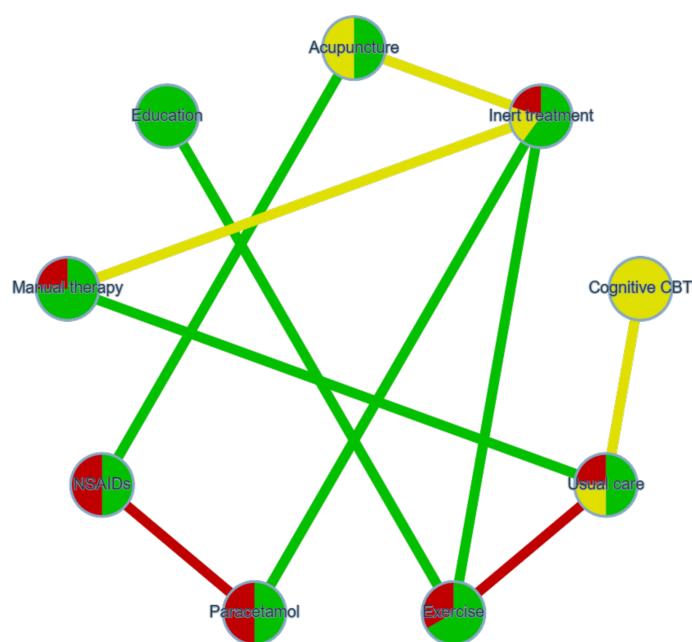
Opioid:Paracetamol	1	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Indirect evidence								
Acupuncture:Education	0	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Acupuncture:Exercise	0	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Acupuncture:Heat wrap	0	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Acupuncture:Manual therapy	0	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Acupuncture:Muscle relaxant	0	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Acupuncture:Opioid	0	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Acupuncture:Paracetamol	0	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Education:Manual therapy	0	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Education:Muscle relaxant	0	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Education:NSAIDs	0	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Education:Opioid	0	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Education:Paracetamol	0	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Exercise:Inert treatment	0	Some concerns	Undetected	No concerns	No concerns	Some concerns	No concerns	Moderate
Exercise:Muscle relaxant	0	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Exercise:NSAIDs	0	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Exercise:Opioid	0	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low

Exercise:Paracetamol	0	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Heat wrap:Inert treatment	0	Some concerns	Undetected	No concerns	No concerns	Some concerns	No concerns	Moderate
Heat wrap:Manual therapy	0	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Heat wrap:Muscle relaxant	0	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Heat wrap:NSAIDs	0	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Heat wrap:Opioid	0	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Heat wrap:Paracetamol	0	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Inert treatment:Opioid	0	Some concerns	Undetected	No concerns	No concerns	Some concerns	No concerns	Moderate
Manual therapy:Muscle relaxant	0	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Manual therapy:Opioid	0	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Manual therapy:Paracetamol	0	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Muscle relaxant:NSAIDs	0	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Muscle relaxant:Opioid	0	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Muscle relaxant:Paracetamol	0	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low

2) Pain at 1 month

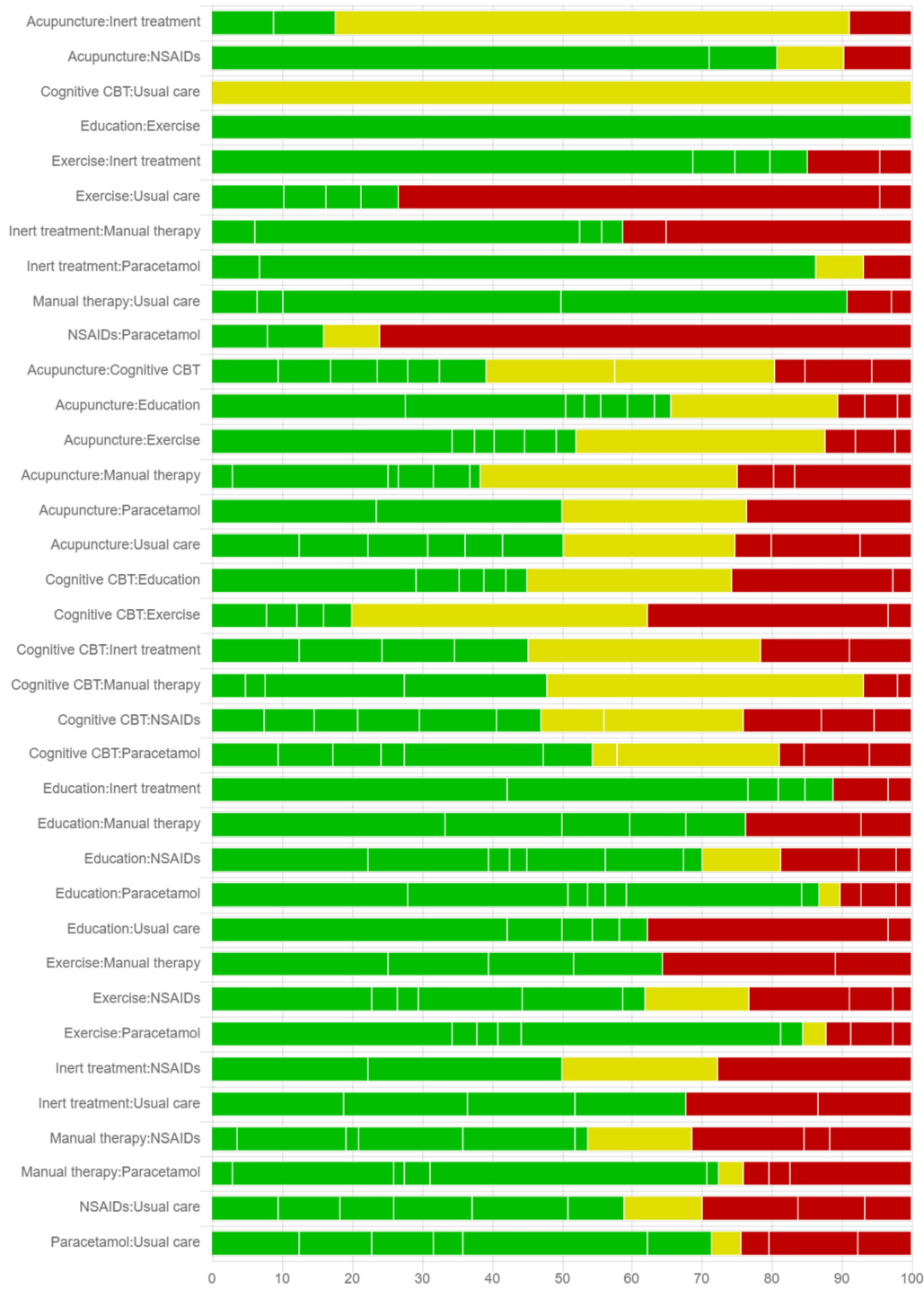
1) Summary of study limitations of the included studies

The colours in the circles indicate the percentage of low RoB studies [green], moderate RoB studies [yellow] and high RoB studies [red] involving each intervention. The colours of the line then indicate the average RoB assessment of each comparison based on the above information – low RoB comparison [green], moderate RoB comparison [yellow] and high RoB comparison [red].



2) Contribution of low or moderate RoB comparisons to each network estimate

Based on the above assessment of RoB for each comparison and the contribution matrix detailing contribution of each direct comparison to all network estimates, the following bar graphs show the percentage of low or moderate RoB contributions for each network estimate. The judgements about study limitations in each direct comparison is shown at the beginning of the graph. Each bar corresponds to a NMA relative treatment effect and shows how much information comes from comparisons at moderate risk of bias [yellow].



3) Summary grading of Evidence

Comparison	Number of studies	Within-study bias	Reporting bias	Indirectness	Imprecision	Heterogeneity	Incoherence	Confidence rating
Mixed evidence								
Acupuncture:Inert treatment	1	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Acupuncture:NSAIDs	1	No concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Low
Cognitive CBT:Usual care	1	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Education:Exercise	1	No concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Moderate
Exercise:Inert treatment	1	No concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Low
Exercise:Usual care	1	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Inert treatment:Manual therapy	2	Some concerns	Undetected	No concerns	No concerns	Some concerns	No concerns	Moderate
Inert treatment:Paracetamol	1	No concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Moderate
Manual therapy:Usual care	2	No concerns	Undetected	No concerns	No concerns	Some concerns	No concerns	High
NSAIDs:Paracetamol	1	Major concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Indirect evidence								
Acupuncture:Cognitive CBT		Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Acupuncture:Education		No concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Moderate
Acupuncture:Exercise		Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low

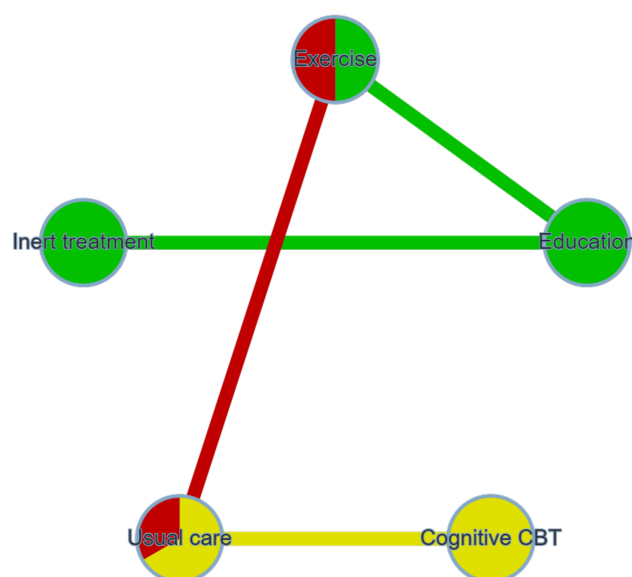
Acupuncture:Manual therapy	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Acupuncture:Paracetamol	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Acupuncture:Usual care	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Cognitive CBT:Education	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Cognitive CBT:Exercise	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Cognitive CBT:Inert treatment	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Cognitive CBT:Manual therapy	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Cognitive CBT:NSAIDs	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Cognitive CBT:Paracetamol	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Education:Inert treatment	No concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Moderate
Education:Manual therapy	No concerns	Undetected	No concerns	No concerns	Some concerns	No concerns	High
Education:NSAIDs	No concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Moderate
Education:Paracetamol	No concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Moderate
Education:Usual care	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Exercise:Manual therapy	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Exercise:NSAIDs	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Exercise:Paracetamol	No concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Low

Inert treatment:NSAIDs	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Inert treatment:Usual care	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
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Manual therapy:Paracetamol	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
NSAIDs:Usual care	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Paracetamol:Usual care	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low

3) Pain at 12 months

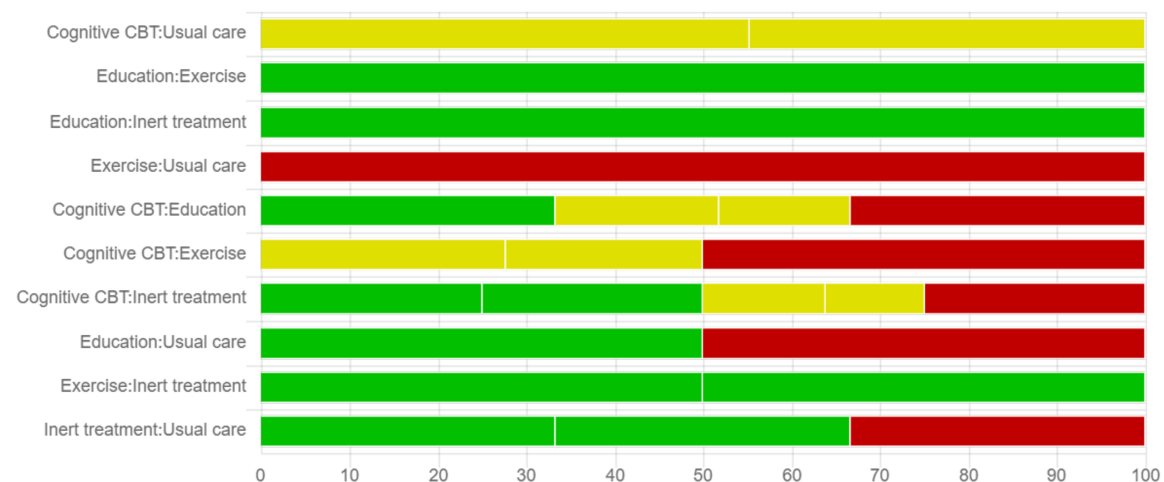
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3) Summary grading of Evidence

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Education:Exercise	1	No concerns	Undetected	No concerns	Some concerns	No concerns	No concerns	Moderate
Education:Inert treatment	1	No concerns	Undetected	No concerns	Some concerns	No concerns	No concerns	Moderate
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Indirect evidence								
Cognitive CBT:Education	-	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low

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Cognitive CBT:Inert treatment	-	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Education:Usual care	-	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Exercise:Inert treatment	-	No concerns	Undetected	No concerns	Some concerns	No concerns	No concerns	Moderate
Inert treatment:Usual care	-	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low

Supplement Q. GRADE for Disability Outcome

Introduction

CINeMA considers 6 domains: (i) within-study bias, (ii) reporting bias, (iii) indirectness, (iv) imprecision, (v) heterogeneity, and (vi) incoherence. Features include the percentage contribution matrix, relative treatment effects for each comparison, estimation of the heterogeneity variance, prediction intervals, and tests for the evaluation of the assumption of coherence. In evaluating imprecision, heterogeneity, and incoherence, we consider the impact of these components of variability in forming clinical decisions.

Table of reasons for downgrading

We use the CINeMA software for GRADE assessment.^{26 27} We downgrade network estimate according to the following criteria.

(1) Study limitations: We downgraded by one level when the contributions from low RoB comparisons were less than 25% and contributions from moderate or high RoB comparisons were 75% or greater.

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(3) Inconsistency: We rated two concepts, heterogeneity and incoherence (inconsistency), in this domain.

For heterogeneity, we looked at the common tau and found that it is low compared to the expected value as reported in the literature,²⁹ so we did not downgrade any network estimate for heterogeneity. For inconsistency, we looked at the results of side splitting and we downgraded the comparisons with important inconsistency ($p < 0.10$), where we have not downgraded for imprecision (we did not downgrade the same network estimate for both imprecision and inconsistency).

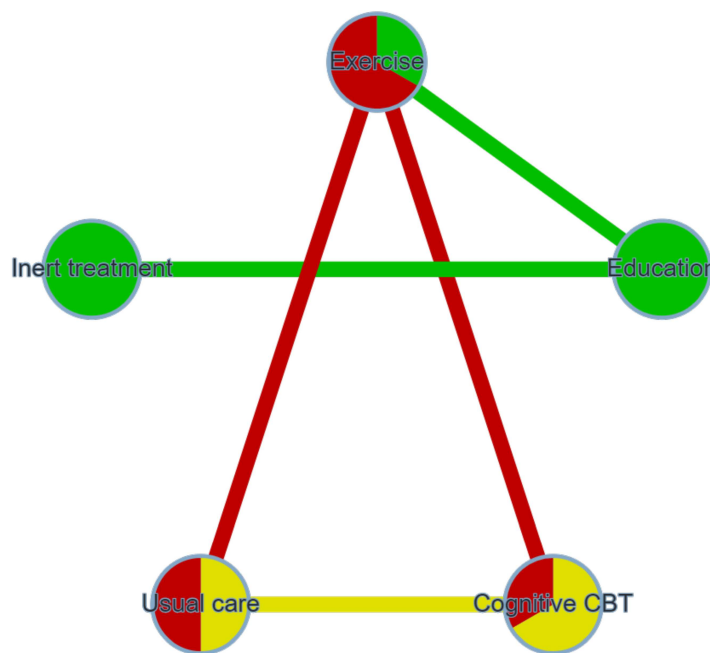
(4) Indirectness: We have assured transitivity in our network by limiting the included studies to acute and subacute population and to non-mixed treatments for LBP. Thus, we did not downgrade for indirectness.

(5) Reporting bias: We cannot completely rule out the possibility that some studies are still missing. However, we assumed that publication bias was undetected.

1) Disability at 12 months

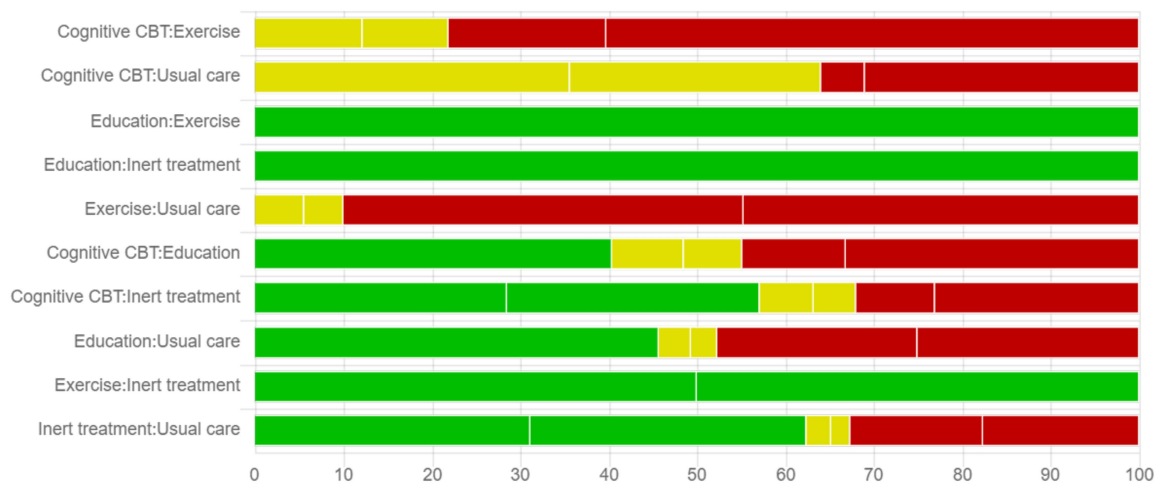
1- Summary of study limitations of the included studies

The colours in the circles indicate the percentage of low RoB studies [green], moderate RoB studies [yellow] and high RoB studies [red] involving each intervention. The colours of the line then indicate the average RoB assessment of each comparison based on the above information – low RoB comparison [green], moderate RoB comparison [yellow] and high RoB comparison [red].



2-Contribution of low or moderate RoB comparisons to each network estimate

Based on the above assessment of RoB for each comparison and the contribution matrix detailing contribution of each direct comparison to all network estimates, the following bar graphs show the percentage of low or moderate RoB contributions for each network estimate. The judgements about study limitations in each direct comparison is shown at the beginning of the graph. Each bar corresponds to a NMA relative treatment effect and shows how much information comes from comparisons at low [green] and high risk of bias [high].



3-Summary grading of Evidence

Comparison	Number of studies	Within-study bias	Reporting bias	Indirectness	Imprecision	Heterogeneity	Incoherence	Confidence rating
Mixed evidence								
Cognitive CBT:Exercise	1	Major concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Very low
Cognitive CBT:Usual care	3	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Education:Exercise	1	No concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Moderate
Education:Inert treatment	1	No concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Low
Exercise:Usual care	2	Major concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Very low
Indirect evidence								
Cognitive CBT:Education	-	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Cognitive CBT:Inert treatment	-	Some concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Education:Usual care	-	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low
Exercise:Inert treatment	-	No concerns	Undetected	No concerns	Some concerns	Some concerns	No concerns	Low
Inert treatment:Usual care	-	Some concerns	Undetected	No concerns	Major concerns	No concerns	No concerns	Very low

Supplement R. Data check

We checked the dataset for data extraction errors or “outlier effect sizes” having an influence on overall effects. We defined an “outlier effect sizes” of a study, visually inspecting forest plots of pairwise meta-analyses³⁰, when SMDs are greater than 1.5^{31 32} assuming 2 points of between population standard deviations across comparisons (resulting from the mean estimate of all final SD values in the control groups^{33 34}, see row dataset in OSF repository <https://osf.io/sjr4y> for 0-10 NRS scale). This calculation is coherent with literature where the MID between group difference is commonly set at 1 point (2 SD) on a NRS scale of 0-10³⁵. Coherently, in the Nice Guideline for Low Back Pain and Sciatica³⁶ the panel considered clinical important an improvement of 10% as a measure of clinical benefit e.g. 1 point decrease on a 0-10 scale for pain intensity³⁵.

Supplement S. References

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