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Update of the simplified criteria for autoimmune hepatitis: evaluation of the methodology for immunoserological testing

Johanna Galaski[1]*, Christina Weiler-Normann[1,2,7]*, Miriam Schakat[1], Kalliopi Zachou[3], Paolo Muratori[4,7], Sibylle Lampalzer[1], Friedrich Haag[5], Christoph Schramm[1,2,7], Marco Lenzi[6,7], George N. Dalekos[3], Ansgar W. Lohse[1,7]

[1]I. Department of Medicine, University Medical Center Hamburg-Eppendorf, Hamburg, Germany; [2]Martin Zeitz Center for Rare Diseases, University Medical Center Hamburg-Eppendorf, Hamburg, Germany; [3]Institute of Internal Medicine and Hepatology, Department of Medicine and Research Laboratory of Internal Medicine, National Expertise Center of Greece in Autoimmune Liver Diseases, General University Hospital of Larissa, Larissa, Greece; [4]Department for Life Quality Studies, Alma Mater Studiorum, University of Bologna, Bologna, Italy; [5]Institute of Immunology, University Medical Center Hamburg-Eppendorf, Hamburg, Germany; [6]Center for the Study and Treatment of Autoimmune Diseases of the Liver and Biliary System, Department of Medical and Surgical Sciences (DIMEC), University of Bologna, Italy; [7]European Reference Network on Hepatological Diseases (ERN RARE-LIVER).

*Equally contributing authors

Corresponding author

Ansgar W. Lohse

I. Department of Medicine, University Medical Center Hamburg-Eppendorf, Martinistraße 52, 20246 Hamburg, Germany.

Tel. +49 (0)40 741053910; Fax. +49 (0)40 741058531

Email: a.lohse@uke.de

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Conflict of Interest

Weiler-Normann C reports speaker's fees from Euroimmun and Werfen (Inova) to her institution. All other authors declare no conflict of interest with respect to this study.

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Data availability statement

The dataset generated during this study is available from the corresponding author upon reasonable request.

Author contributions

Galaski J: substantial contribution to conception and design, data acquisition and analysis, interpretation of data, drafting of the article

Weiler-Normann C: substantial contribution to conception and design, data acquisition and interpretation of data, critical revision of the article for important intellectual content

Schakat M, Zachou K, Muratori P, Lampalzer S, Haag F, Lenzi M, Dalekos GN: substantial contribution to data acquisition, critical revision

Schramm, C: critical revision of the article for important intellectual content

Lohse AW: substantial contribution to conception and design, interpretation of data, critical revision of the article for important intellectual content

All authors approved submission.

Lay summary

Autoantibodies are a hallmark of autoimmune hepatitis and are traditionally tested for by immunofluorescence assays on rodent tissue sections. Herein, we demonstrate that both HEp-2 cells as substrate for ANA IFT and ELISA-based testing are potentially reliable alternatives for autoantibody assessment in autoimmune hepatitis. We propose the implementation of these testing methods into the simplified criteria for the diagnosis of autoimmune hepatitis.

Highlights

- IFT on HEp-2 cells is a valid alternative to the standard ANA assessment on rodent tissue sections in AIH when cutoffs titers are increased
- ANA ELISA and F-actin ELISA represent potential alternatives to IFT in the diagnosis of AIH
- ANA ELISA kits should include HEp-2 nuclear extracts to account for unrecognized autoantigens
- ELISA cutoffs need to be validated locally to be predictive in diagnosing AIH

Abstract

Background & Aims: The simplified criteria for the diagnosis of autoimmune hepatitis (AIH) include immunofluorescence testing (IFT) of antinuclear and smooth muscle autoantibodies (ANA and SMA) on rodent tissue sections. We aimed to establish scoring criteria for implementation of ANA IFT on HEp-2 cells and ELISA-based testing. **Methods:** ANA and SMA reactivity of 61 AIH sera and 72 non-alcoholic fatty liver disease (NAFLD) controls were separately assessed on tissue sections and human epithelioma (HEp-2) cells to compare the diagnostic value at increasing titers. A total of 113 AIH patients at diagnosis and 202 controls from three European centers were assessed by IFT as well as three different commercially available ANA ELISA and one anti-F-actin ELISA. **Results:** ANA assessment by IFT on liver sections had 83.6% sensitivity and 69.4% specificity for AIH at a titer of 1:40. On HEp-2 cells, sensitivity and specificity were 75.4% and 73.6%, respectively, at an adjusted titer of 1:160. Area under the curve (AUC) values of ANA ELISA ranged from 0.70 – 0.87, with ELISA coated with HEp-2 extracts in addition to selected antigens performing significantly better. SMA assessment by IFT had the highest specificity for the SMA-VG/T pattern and anti-MF reactivity on HEp-2 cells. ELISA-based anti-F-actin evaluation was a strong predictor of AIH (AUC 0.88) and performed better than SMA assessment by IFT (AUC 0.77 – 0.87). **Conclusion:** At adjusted cutoffs, both ANA IFT using HEp-2 cells and ELISA-based autoantibody evaluation for ANA and SMA are potential alternatives to tissue-based IFT for the diagnosis of AIH.

Introduction

105 Autoimmune hepatitis (AIH) is a chronic immune-mediated liver disease. Due to heterogeneity of the presentation, the diagnosis remains challenging. An early diagnosis is, however, critical for timely initiation of life-saving immunosuppressive therapy. To assist diagnostic evaluation, a simplified diagnostic score was established by the International Autoimmune Hepatitis Group (IAIHG) in 2008 for use in clinical practice [1]. Scoring criteria include characteristic findings on liver histology, the absence of viral hepatitis, an elevation of immunoglobulin G (IgG), and circulating autoantibodies.

Autoantibodies associated with AIH include antinuclear antibodies (ANA), smooth muscle antibodies (SMA), liver kidney microsomal type 1 (LKM1) antibodies, liver cytosol type 1 (LC1) antibodies, and soluble liver antigen/liver pancreas (SLA/LP) antibodies. Screening for liver disease-associated autoantibodies is traditionally performed by immunofluorescence testing (IFT) on rodent tissue sections. Accordingly, the simplified AIH score refers to autoantibody titers as measured by IFT using tissue sections at a cutoff titer of 1:40. However, in several laboratories, there has been a shift of autoantibody assessment towards human epithelioma (HEp-2) cells rather than tissue sections as substrate for IFT. Furthermore, enzyme-linked immunosorbent assays (ELISA), for which the score does not account for, are frequently used in some countries. In order to make the simplified AIH score usable across the world, adaptation of the score to different immunoserology methods is urgently needed.

HEp-2 cells are widely used as substrate for ANA evaluation. In addition to a higher sensitivity, characteristic staining patterns evaluated on HEp-2 cells are useful in guiding further confirmatory testing. However, a consensus statement by the IAIHG committee for autoimmune serology advises against the use of HEp-2 cells at a screening stage [2] because of a high positivity rate in healthy individuals at low

cutoff titers [3]. If HEp-2 cells are used, the IAIHG suggests titers should be halved for the simplified score to be applicable [1]. However, this possible correction factor suggestion has never been validated by comparative studies [4].

SMA constitute a heterogeneous group of autoantibodies that primarily target F-actin, [5]. On kidney tissue sections, Bottazzo and colleagues distinguished three immunofluorescence patterns: SMA-V (vessels), SMA-VG (vessels/glomeruli), and SMA-VGT (vessels/ glomeruli/ tubuli) [6]. In contrast to the SMA-V pattern, SMA-VG/T correlates with F-actin reactivity and is more specific for AIH [6-8]. Similarly, anti-F-actin antibodies stain microfilaments (MF) on HEp-2 cells [9]. Overall, sensitivity and specificity of SMA positivity strongly depend on fluorescence patterns, which is not taken into consideration by current AIH scoring systems.

Since IFT is time-consuming, requires experienced technicians and lacks standardization, ELISA have emerged as a widely used alternative for routine autoantibody testing in many laboratories, especially in the United States. These tests were originally developed for use in the evaluation of rheumatic diseases and their diagnostic value in liver disease is unknown. ELISA testing can minimize interobserver variability inherent to IFT. However, it is unclear whether ELISA can replace IFT for the detection of the heterogeneous autoantibodies ANA and SMA with their range of antigenic specificities. To complicate matters even further, up to 30% of ANA-positive AIH patients do not react with any known nuclear antigens [10] and might thus be missed by ELISA testing, which are based primarily on known nuclear antigens. In addition, commercially available ANA ELISA lack standardization – they differ in their antigenic profiles and assay-specific cutoff values.

Taken together, the AIH simplified score does not account for ANA and SMA as evaluated by IFT on HEp-2 cells or for ELISA, even though these tests are widely used. We therefore set out to study the diagnostic validity of IFT and ELISA-based

autoantibody testing for the diagnosis of AIH to make these applicable in diagnosing AIH.

Patients and methods

Study population

This multicenter study included a total of 113 patients with AIH at diagnosis and 202 controls (82 NAFLD patients, 99 primary sclerosing cholangitis (PSC) patients and 21 healthy controls) from three centers: Hamburg (Germany), Bologna (Italy), and Larissa (Greece). A flow-chart of patient cohorts is shown in Figure 1. The large majority of AIH patients (106/113, 93.8%) were treatment-naïve at the time of sampling. In addition, sera from 26 patients with primary biliary cholangitis (PBC) were tested and analyzed separately. Sera were collected between December 2006 and March 2020 and stored at -80°C until use. The study was approved by the local ethics committee (PV4081-0005, PV 4081-0008).

The diagnosis of AIH was based on clinical, serological, and histopathological criteria, consistent with the EASL clinical practice guidelines [11], and confirmed by long-term follow-up in all patients. Patients with AIH and features of PSC or PBC were excluded from the study. Diagnoses of disease controls were based on established diagnostic criteria [12-14]. Blood donors with liver enzymes within the normal range, negative for HBV/HCV, and negative for autoantibodies by IFT were included as healthy controls.

Autoantibody assessment by IFT

IFT was performed in the respective center in which sera were collected. At the University Medical Center Hamburg-Eppendorf sera were tested using a Biochip

183 Mosaic of primate liver, rat kidney, and rat stomach tissue sections as well as human 184 epithelioma (HEp-2) cells (Mosaic Basic Profile 3, Euroimmun, Germany). The assay 185 was performed manually according to the manufacturer's instructions at a dilution of 186 1:40. Further dilutions up to 1:1280 were processed by the Helios automated IFA 187 system (Aesku Diagnostics, Wendelsheim, Germany), using the same substrates 188 and conditions. Reactivity patterns were assessed under a fluorescence microscope 189 (Eurostar, Euroimmun, Germany). ANA and SMA reactivity were separately 190 evaluated on all four substrates. SMA reactivity on kidney sections was assessed 191 according to Bottazzo et al. [6]. The observers were blinded to clinical data.

192 Sera from the University Hospital of Bologna, Italy, were tested by IFT on both tissue 193 sections and HEp-2 cells (Euroimmun, Germany) and were automatically processed 194 at a starting dilution of 1:80 up to 1:640. ANA titers were mainly reported as 195 assessed on HEp-2 cells and thus these data were used for comparison with ANA 196 ELISA.

197 Sera from the University Hospital of Larissa, Greece, were tested by 198 immunofluorescence on in-house fresh cryostat liver, kidney and stomach rat 199 sections and HEp-2 cells (Inova Diagnostics). ANA titers were mainly reported as 200 assessed on tissue sections and thus these data were used for comparison with ANA 201 ELISA. Sera were processed manually at a starting dilution of 1:40 up to 1:640.

202

203 **Detection of antinuclear and F-actin antibodies by ELISA**

204 All ELISA testing was performed at the University Medical Center Hamburg-205 Eppendorf. Antinuclear antibodies were assessed using enzyme immunoassays from 206 three different manufacturers (Quanta Lite ANA ELISA, Inova Diagnostics, US; ANA 207 Screening Test, Bio-Rad, US; ANA Screen ELISA, Euroimmun, Germany). All assays 208 detect autoantibodies of IgG subtype and display antigenic specificities to dsDNA,

histones, Sm/RNP, SS-A, SS-B, Scl-70, centromere, and Jo-1. The Quanta Lite ANA ELISA is additionally coated with highly purified proliferating cell nuclear antigen (PCNA), mitochondrial M2 antigen, and ribosomal-P proteins. Besides individual antigens, immunoassays from both Inova Diagnostics and Bio-Rad include HEp-2 cell nuclei extracts.

Antibodies to F-actin were detected using a commercial ELISA (Quanta Lite Actin IgG, Inova Diagnostics, US). All enzyme immunoassays were performed in duplicates according to the manufacturer's recommendations. Investigators who carried out immunoassays were blinded to clinical data and the results of IFT.

Statistical analyses

Data was expressed as median (range), or n (%) as appropriate. Statistical significance between groups was assessed with Fisher's exact test for categorical variables and the Mann-Whitney *U* test for continuous variables. Correlations were evaluated using Spearman correlation coefficients. The diagnostic value of variables in discriminating AIH from controls was assessed by receiver operating characteristic (ROC) analysis. Statistical significance between area under the curve (AUC) values was assessed by the DeLong test. All reported *P* values are based on two-sided tests and a P value < 0.05 was considered statistically significant. Statistical analyses were performed using GraphPad Prism (version 6), IBM SPSS (version 23), and R software (version 3.5.1).

Results

Comparison of HEp-2 cells and tissue sections as substrates for ANA IFT

We first investigated the diagnostic value of HEp-2 cells in comparison to tissue sections as substrates for ANA IFT in the context of AIH. To this end, sera from 61 AIH patients and 72 patients with biopsy-proven NAFLD treated at the University Medical Center Hamburg-Eppendorf were evaluated for autoantibodies by IFT. Clinical characteristics of the patient groups at the time of sampling are summarized in supplemental Table 1.

Sensitivity and specificity of ANA IFT for HEp-2 cells and tissue sections are shown in Table 1. Among tissue sections, primate liver showed the highest diagnostic value for ANA evaluation. Sensitivity and specificity were 83.6% and 69.4% at a titer of 1:40, respectively, and 68.9% and 80.6% at a titer of 1:80, respectively. Specificity increased to 91.7% at a titer of 1:160 at the cost of a lower sensitivity of 47.5%. As expected, the use of HEp-2 cells led to higher titers. Specificity was inadequate at a 1:40 dilution. At a titer of 1:80, sensitivity was 91.8% at a low specificity of 36.1%. At higher titers, sensitivity and specificity were comparable to those observed on liver sections: 75.4% and 73.6%, respectively, at a titer of 1:160; 72.1% and 76.4%, respectively, at a titer of 1:320. The homogenous pattern was significantly more frequent in AIH patients (41.0%) than in NAFLD patients (6.9%, $P < 0.001$).

Sensitivity and specificity of SMA fluorescence patterns on tissue sections and HEp-2 cells

We next assessed the diagnostic value of several SMA fluorescence patterns at different titers (Table 2). As expected, at a 1:40 titer, the SMA-V pattern on kidney sections, staining of smooth muscle on stomach sections as well as consideration of any SMA positivity resulted in a low specificity of 33.3% – 45.8%. In contrast, the SMA-VG pattern was more specific for the diagnosis of AIH even at low titers. Sensitivity and specificity were 72.1% and 70.8%, respectively, at a titer of 1:40, and

65.6% and 88.9%, respectively, at a titer of 1:80. The highest specificity was seen for the SMA-VGT pattern and anti-MF reactivity on HEp-2 cells. At a 1:40 dilution, specificity was 93.1% – 94.4% at a sensitivity of 52.5% – 60.7%. Of note, with increasing titers, staining of the SMA-VGT pattern first faded for tubuli, then glomeruli, and finally vessels. In other terms, the SMA-VGT pattern changed to SMA-VG and finally to SMA-V with increasing dilutions. Taken together, SMA positivity was highly specific even at low titers for SMA-VG/T and anti-MF reactivity on HEp-2 cells, but only at higher titers for other SMA patterns.

ELISA-based autoantibody testing for the diagnosis of AIH

We next assessed the diagnostic value of ELISA-based autoantibody evaluation to discriminate between AIH and controls. Sera from three European centers were reassessed by three different ANA ELISA and one F-actin ELISA. Clinical characteristics of the patient groups at the time of sampling are summarized in supplemental Tables 1 – 3.

ANA testing by the Bio-Rad and Inova ANA ELISA had a similar diagnostic accuracy (AUC 0.85 and 0.87, respectively; P = 0.32) and performed significantly better compared to the ANA Euroimmun ELISA (AUC 0.70; $P < 0.001$) (Figure 2A).

Correlation analyses between the ANA ELISA results found the strongest correlation between the Bio-Rad and Inova ANA ELISA ($r_s = 0.72$; $P < 0.001$) (Supplemental Figure 1). Test characteristics of the ANA ELISA kits varied greatly at cutoffs recommended by the manufacturers. In fact, sensitivity and specificity were 65.5% and 88.6% for the Bio-Rad assay (recommended cutoff ≥ 1 RU), 79.6% and 78.2% for the ANA Inova assay (recommended cutoff ≥ 20 RU), and 22.1% and 95.0% for the ANA Euroimmun assay (recommended cutoff ≥ 1 RU), respectively (Table 3).

Like for ANA, we assessed the diagnostic value of a F-actin ELISA. ROC analysis revealed anti-F-actin as a strong predictor of AIH (AUC 0.89) (Figure 2B). At a cutoff of 20 RU, sensitivity and specificity were 81.4% and 82.2%, respectively; at a cutoff of 30 RU, sensitivity and specificity were 66.4% and 92.6%, respectively (Table 3). Importantly, anti-F-actin was still a predictor of AIH in the subgroup of patients with normal range IgG (≤ 16 g/l; n = 35/109) (AUC 0.79).

ELISA- compared to IFT-based evaluation of autoantibodies

We next compared ELISA- and IFT-based ANA evaluation. To account for the inter-laboratory variability inherent to IFT, ELISA assessment was compared to IFT results obtained by the respective centers according to local standards. Figure 3 and 4 show the diagnostic performance of ELISA *vs.* IFT for ANA and SMA/F-actin, respectively, for each center. ANA testing by ELISA and IFT performed similarly for all cohorts, except for the Euroimmun ELISA that showed a significantly lower AUC compared to IFT for the Hamburg cohort (Euroimmun ANA ELISA, AUC 0.65; ANA IFT, AUC 0.82 – 0.83; $P < 0.001$).

In addition to the patient groups shown in Figure 1, we tested sera from 26 PBC patients known to frequently present with ANA. Clinical characteristics of PBC patients are detailed in supplemental Table 4. While 17/26 (65.4%) of PBC patients tested positive for ANA by IFT on HEp-2 cells at a cut-off of 1:80, 23/26 (88.4%) and 25/26 (96.2%) tested positive by the Bio-Rad and Inova ANA ELISA, respectively. Importantly, median values of the Inova ANA ELISA were significantly higher in PBC patients compared to AIH patients (49.6 RU AIH *vs.* 161.7 RU PBC; $P < 0.001$) while there was no statistical significant difference for the Bio-Rad ELISA (1.6 RU AIH *vs.* 2.0 RU PBC; $P = 0.25$).

310 The F-actin ELISA yielded higher AUC values compared to IFT for each center, 311 reaching statistical significance for the Hamburg cohort when compared to anti-MF 312 reactivity on HEp-2 cells (F-actin ELISA, AUC 0.86; anti-MF AUC 0.79; $P = 0.003$) 313 and for the Bologna cohort when compared to any SMA reactivity (F-actin ELISA, 314 AUC 0.93; any SMA, AUC 0.77; $P = 0.002$).

315 We further assessed the performance of ELISA-based autoantibody testing in the 316 subgroup of patients with a histological diagnosis of liver cirrhosis. Overall, 24 AIH 317 patients and 15 controls (4 PSC patients, 11 NAFLD patients) with cirrhosis were 318 identified. ANA IFT assessed on tissue sections (available for n = 35; 20 AIH patients 319 *vs.* 15 controls) reached an AUC of 0.84 whereas ELISA-based ANA assessment 320 yielded higher AUC values of 0.88 – 0.93, without reaching statistical significance 321 (supplemental Figure 2A). In contrast, anti-F-actin (n = 39) was again a strong 322 predictor of AIH (AUC 0.91) and performed significantly better than SMA assessment 323 by IFT (SMA-VG/T; AUC 0.80; $P = 0.049$) (supplemental Figure 2B).

324

325 **Concordance between IFT- and ELISA-based ANA testing**

326 We next assessed concordance between IFT- and ELISA-based autoantibody testing 327 and were specifically interested in the proportion of AIH patients that tested positive 328 by IFT but were missed when tested by ELISA. Of 51 AIH patients from the Hamburg 329 cohort that tested positive for ANA by IFT on liver tissue sections, the ANA ELISA by 330 Inova, Bio-Rad and Euroimmun detected 40/51 (78.4%), 28/51 (54.9%), and 10/51 331 (19.6%) cases at recommended cut-offs, respectively. Conversely, of 10 AIH patients 332 that tested negative for ANA by IFT, 6 (60%) tested positive by the Inova ELISA and 333 4 (40%) by the Bio-Rad ELISA. Furthermore, the Inova and Bio-Rad assays detected 334 all but one of ANA-positive AIH cases from the Larissa and Bologna cohorts.

335 Together, the ROC analysis indicates that ELISA represent a potential alternative to 336 IFT-based autoantibody assessment. However, assays vary considerably in their 337 performance and cut-offs need to be validated for the diagnosis of AIH. If these 338 aspects are taken under consideration and local cut-offs established, ELISA-based 339 autoantibody testing as proposed in Table 4 can be used in the diagnostic work-up of 340 liver disease patients.

341

342 Discussion

343

344 This is the first study to comprehensively evaluate IFT- and ELISA-based 345 assessment of ANA and SMA/anti-F-actin in AIH. In analogy to the simplified IAIHG 346 diagnostic score that largely refers to autoantibody assessment as evaluated by IFT 347 on tissue sections, we propose the implementation of autoantibody testing as 348 measured by IFT on HEp-2 cells and ELISA.

349 We first aimed to validate the use of HEp-2 cells as substrate for ANA IFT in patients 350 with AIH. As expected, at low titers, ANA as evaluated on HEp-2 cells showed a high 351 sensitivity at the expense of a low specificity. It is precisely the low specificity at a 352 1:40 titer that led the IAIHG to advise against use of HEp-2 cells for ANA evaluation 353 at a screening stage [2]. However, to our knowledge, the diagnostic value of ANA IFT 354 on HEp-2 cells has not been assessed at higher titers in the context of liver disease. 355 A previous study investigating ANA IFT in liver disease reported an increased 356 sensitivity of ANA IFT using HEp-2 cells, but was restricted to a 1:40 dilution [15]. 357 Our results suggest that HEp-2 cells are a valid alternative to tissue sections, if 358 threshold titers are adapted. We here propose increasing cutoff titers to 1:160 and 359 1:320 for the simplified diagnostic score to be applicable. As outlined above, a cutoff 360 titer of 1:160 is also the recommended cutoff for ANA screening in rheumatic

1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25 26 27 28 29 30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46 47 48 49 50 51 52 53 54 55 56 57 58 59 60 61 62 63 64 65

diseases [16]. However, titers vary depending on reagents and equipment used and should be validated locally. In addition, the difference in immunofluorescence intensity between tissue sections and HEp-2 cells is not the same for all subtypes of ANA, but highly dependent on the respective ANA pattern. Nevertheless, overall, HEp-2 cells are a valid alternative to tissue sections for ANA evaluation in AIH.

We further compared the diagnostic value of different SMA patterns for the diagnosis of AIH. In line with a study by Muratori and colleagues [9], we found that specificity was highest for SMA-VGT and anti-MF reactivity at a titer of 1:40. Complementing this previous study, we additionally assessed SMA patterns at further dilutions. Interestingly, sensitivity and specificity of generic SMA at higher titers was comparable to the diagnostic value of SMA-VG/T and anti-MF reactivity at a 1:40 titer. Furthermore, as previously described [6], we observed a shift from SMA-VGT to SMA-G and then SMA-V with increasing dilutions for individual samples. It thus appears likely that the SMA-VGT pattern is a reflection of high-titer SMA with specificity for F-actin. In contrast, the SMA-V pattern can be seen for both low-titer SMA with anti-F-actin reactivity or SMA targeting other cytoskeletal components. Taken together, our results add to the literature [6, 7, 9] that highlights the importance of reporting SMA patterns, in both the scientific and clinical context.

Several studies have assessed ANA evaluation by ELISA in rheumatic diseases [17-21], but analogous studies in AIH are lacking. To fill this gap, we assessed the diagnostic value of three different ANA ELISA in AIH patients. We observed significant differences depending on the ELISA used, with the Bio-Rad and Inova assays performing best. In contrast, at the cut-off recommended by the manufacturer, the Euroimmun ANA ELISA had a low sensitivity of 22.1% at a 95% specificity. These results might be explained by differing ELISA formulations. Indeed, both the Inova and Bio-Rad ANA ELISA include HEp-2 nuclear extracts in addition to

387 recombinant and purified nuclear antigens to account for unrecognized autoantigens. 388 In contrast, the Euroimmun assay is only comprised of selected nuclear antigens. Its 389 antigenic specificities are therefore better defined, ensuring high specificity for the 390 diagnosis of rheumatic diseases. However, our data suggest that this comes at the 391 cost of a low diagnostic value in autoimmune hepatitis. With regard to ELISA 392 formulations, it is also worth mentioning that the Inova ANA ELISA is the only assay 393 in this study including purified ribosomal P and mitochondrial M2 antigen. In a study 394 by Calich and colleagues, autoantibodies against ribosomal P were found in 9/93 395 (9.7%) AIH patients and none of the healthy controls [22]. In contrast, the 396 incorporation of mitochondrial antigens is not expected for an ANA screening assay 397 and carries considerable potential for confusion. Indeed, if the Inova ANA ELISA 398 were to be used for the diagnostic workup of elevated liver enzymes, distinction 399 between ANA and antimitochondrial antibodies (AMA) would not be possible in a 400 reasonable fashion. Incorporation of mitochondrial antigens also likely explains the 401 significantly higher values of the Inova ANA ELISA in PBC patients compared to AIH 402 patients. Overall, while the careful choice of ELISA formulation and validation of cut-403 offs is critical, our data suggest that in principle ELISA testing represents a potentially 404 good alternative to ANA IFT. Importantly, if ELISA-based autoantibody assessment is 405 negative despite clinical suspicion of AIH, additional IFT should be performed.

406 In the present study, we further compared IFT-based SMA evaluation to an anti-F-407 actin ELISA. Consistent with previous results [23], we found that anti-F-actin had a 408 significantly higher diagnostic value for the diagnosis of AIH. Interestingly, while 409 hypergammaglobulinemia potentiated the predictive value of anti-F-actin for the 410 diagnosis of AIH, F-actin autoantibodies were still a strong predictor of AIH in the 411 subgroup of AIH patients with IgG within the normal range (AUC 0.79).

412 Several limitations to the present study warrant further discussion. First, IFT allows
413 for the detection of additional autoantibodies such as AMA and provides
414 characteristic staining patterns that point towards antigenic specificities of ANA. The
415 benefit of this relevant information was not assessed in the present study. While ANA
416 ELISA do not provide such additional information, some specific and reliable tests
417 exist to further assess antigen specificity of ANA-positive sera. Indeed, most of the
418 PBC sera we tested were highly positive both in the Inova ANA ELISA, which does
419 however also include M2 antigen, the key target of antimitochondrial antibodies
420 characteristic of PBC, as well as in the Bio-Rad ANA ELISA. Thus, for discrimination
421 between AIH and PBC sera, further systematic testing by a specific M2-AMA ELISA
422 and by sp100 and gp210 ELISA would be required. However, this would have been
423 beyond the scope of the present study.

424 Second, we included only one F-actin ELISA. However, compared to the
425 heterogeneous group of ANA, F-actin is a defined antigen and the F-actin ELISA
426 used in this study was investigated in two previous studies [7, 23].

427 Furthermore, while control cohorts were well characterized, relevant patient groups
428 such as drug-induced liver injury patients were not included in the present study.
429 Finally, the gender distribution between AIH and controls was somewhat unbalanced
430 reflecting the natural sex differences in these conditions. Although this potentially
431 influenced the frequency of autoantibodies in patient groups, it most probably did not
432 affect how the various autoantibody assays compared to one another.

433 In conclusion, our results suggest that both IFT evaluation on HEp-2 cells as well as
434 ELISA-based autoantibody assessment are potential alternatives to IFT on tissue
435 sections. Our data indicate that (1) HEp-2 cells can be used for ANA assessment in
436 AIH if scoring cutoff titers are increased, (2) The SMA-VG/T pattern and anti-MF
437 reactivity on HEP-2 cells are highly specific even at low titers while generic SMA is

specific only at higher titers, (3) ANA and F-actin ELISA show at least equivalent diagnostic performance compared to IFT, but ELISA kits for ANA assessment should include HEp-2 nuclear extracts to account for unknown nuclear antigens and cutoffs need to be validated for the use in AIH. In the future, cut-off values for autoantibody testing should be determined and validated by industry on standardized AIH sera and controls and re-validated by diagnostic laboratories, as technical details may influence the exact values. Nonetheless, the objective nature of these tests will make them more attractive in the future avoiding observation errors due to the subjective assessment of staining patterns as in SMA testing on tissue sections. Based on our results, under the prerequisite of careful choice of ELISA formulation and validation of cut-offs, we propose an adaptation of the simplified diagnostic score for AIH as summarized in Table 4 for everyday use in different laboratory settings.

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525 # Tables

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Table 1. Sensitivity and specificity of ANA IFT for different tissue sections

Substrate	Titer	Sensitivity (%)	Specificity (%)	PPV (%)	NPV (%)	Accuracy (%)
HEp-2 cells	1:40	95.1	8.3	46.8	66.7	48.1
	1:80	91.8	36.1	54.9	83.9	61.7
	1:160	75.4	73.6	70.8	77.9	74.4
	1:320	72.1	76.4	72.1	76.4	74.4
	1:640	60.7	87.5	80.4	72.4	75.2
Primate liver	1:40	83.6	69.4	69.9	83.3	75.9
	1:80	68.9	80.6	75.0	75.3	75.2
	1:160	47.5	91.7	82.9	67.4	71.4
	1:320	47.5	91.7	82.9	67.4	71.4
	1:640	29.5	94.4	81.8	61.3	64.7
Rat kidney	1:40	75.4	73.6	70.8	77.9	74.4
	1:80	65.6	81.9	75.5	73.8	74.4
	1:160	52.5	87.5	78.1	68.5	71.4
	1:320	47.5	91.7	82.9	67.4	71.4
	1:640	34.4	93.1	80.8	62.6	66.2
Rat stomach	1:40	78.7	70.8	69.6	79.7	74.4
	1:80	67.2	81.9	75.9	74.7	75.2
	1:160	52.5	88.9	80.0	68.8	72.2
	1:320	44.3	91.7	81.8	66.0	69.9
	1:640	36.1	93.1	81.5	63.2	66.9
Any tissue positivity (primate liver, rat kidney, rat stomach)	1:40	85.3	65.3	67.5	83.9	74.4
	1:80	73.8	77.8	73.8	77.8	75.9
	1:160	52.5	87.5	78.1	68.5	71.4
	1:320	50.8	91.7	83.8	68.8	72.9
	1:640	37.7	93.1	82.1	63.8	67.7

AIH n=61; NAFLD n=72; ANA, antinuclear antibodies; HEp-2 cells, human epithelioma-2 cells; IFT, immunofluorescence test; NPV, negative predictive value; PPV, positive predictive value.

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Table 2. Sensitivity and specificity of SMA IFT for different patterns

Substrate	Titer	Sensitivity (%)	Specificity (%)	PPV (%)	NPV (%)	Accuracy (%)
HEp-2 (microfilaments)	1:40	60.7	94.4	90.2	73.9	79.9
	1:80	59.0	98.6	97.3	74.0	80.5
	1:160	54.1	98.6	97.1	71.7	78.2
	1:320	52.5	98.6	97.0	71.0	77.4
	1:640	41.0	100	100	66.7	72.9
Kidney SMA-V (vessels)	1:40	78.7	45.8	55.2	71.7	60.9
	1:80	73.8	72.2	69.2	76.5	72.9
	1:160	68.9	80.6	75.0	75.3	75.2
	1:320	62.3	88.9	82.6	73.6	76.7
	1:640	49.2	98.6	96.8	69.6	75.9
Kidney SMA-VG (vessels, glomeruli)	1:40	72.1	70.8	67.7	75.0	71.4
	1:80	65.6	88.9	83.3	75.3	78.2
	1:160	63.9	94.4	90.7	75.6	80.5
	1:320	55.7	97.2	94.4	72.2	78.2
	1:640	36.1	100	100	64.9	70.7
Kidney SMA-VGT (vessels, glomeruli tubuli)	1:40	52.5	93.1	86.5	69.8	74.4
	1:80	49.2	93.1	85.7	68.4	72.9
	1:160	44.3	95.8	90.0	67.0	72.2
	1:320	31.2	97.2	90.5	62.5	66.9
	1:640	23.0	100	100	60.5	64.7

Kidney SMA-VG or HEp2 microfilaments	1:40	75.4	69.4	67.7	76.9	72.2
	1:80	68.9	88.9	84.0	77.1	79.7
	1:160	65.6	94.4	90.9	76.4	81.2
	1:320	62.3	97.2	95.0	75.3	81.2
	1:640	44.3	100	100	67.9	74.4
Liver (bile canaliculi)	1:40	59.0	83.3	75.0	70.6	72.2
	1:80	49.2	95.8	90.9	69.0	74.4
	1:160	42.6	98.6	96.3	67.0	72.9
	1:320	39.3	98.6	96.0	65.7	71.4
	1:640	26.2	100	100	61.5	66.2
Stomach (tunica muscularis, lamina muscularis mucosa, interglandular fibrils)	1:40	83.6	45.8	56.7	76.7	63.2
	1:80	75.4	72.2	69.7	77.6	73.7
	1:160	72.1	80.6	75.9	77.3	76.7
	1:320	68.9	90.3	85.7	77.4	80.5
	1:640	54.1	97.2	94.3	71.4	77.4
Any SMA positivity	1:40	86.9	37.5	54.1	77.1	60.2
	1:80	80.3	69.4	69.0	80.7	74.4
	1:160	72.1	79.2	74.6	77.0	75.9
	1:320	72.1	88.9	84.6	79.0	81.2
	1:640	60.7	97.2	94.9	74.5	80.5

AIH n=61; NAFLD n=72; HEp-2 cells, human epithelioma-2 cells; IFT, immunofluorescence test; NPV, negative predictive value; PPV, positive predictive value; SMA, smooth muscle antibodies; VGT, vessel, glomeruli, tubuli.

Table 3. Diagnostic value of ANA and F-Actin ELISA at cut-offs recommended by manufacturers

ELISA	Assay	Cutoff	Sensitivity (%)	Specificity (%)	PPV (%)	NPV (%)	Accuracy (%)
ANA ELISA	Bio-Rad	≥ 1.0	65.5	88.6	76.3	82.1	80.3
	Inova	≥ 20	79.6	78.2	67.2	87.3	78.7
		≥ 30	69.0	86.6	74.3	83.3	80.3
	Euroimmun	≥ 1.0	22.1	95.0	71.4	68.6	68.9
F-Actin ELISA	Inova	≥ 20	81.4	82.2	71.9	88.8	81.9
		≥ 30	66.4	92.6	83.3	83.1	83.2

AIH n=113; controls n=202; distribution of diagnoses as shown in Figure 1; ANA, antinuclear antibodies; NPV, negative predictive value; PPV, positive predictive value.

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Table 4. Simplified criteria for autoimmune hepatitis

– Update of serological criteria

Variable	Cutoff	Points[1]
ANA or SMA/F-Actin	Positive[2]	1
ANA or SMA/F-Actin	Strongly positive[3]	
or LKM	≥1:40	2
or SLA	Positive	
IgG	>Upper normal limit	1
	>1.1 times upper normal limit	2
Liver histology (with evidence of	Compatible with AIH	1
hepatitis)	Typical AIH	2
Absence of viral hepatitis	Yes	2
		≥6: probable AIH
		≥7: definite AIH

[1]Addition of points achieved (maximum 2 points for autoantibodies);
[2]IFT: ≥1:40 when assessed on tissue sections; ≥ 1:80 or 1:160 for ANA when assessed on HEp-2 cells, depending on local standards. ELISA with locally established cut-offs;
[3]IFT: ≥1:80 when assessed on tissue sections; ≥ 1:160 or 1:320 for ANA when assessed on HEp-2 cells. ELISA with cut-offs established locally;
Note: if ELISA-based autoantibody assessment is negative despite high clinical suspicion of autoimmune hepatitis, IFT should be performed in addition.

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Figure legends

Figure 1. Flow-chart of patient cohorts included in this study.

Figure 2. Receiver-operating-characteristic (ROC) curves showing the diagnostic value of ELISA for the diagnosis of AIH. Diagnostic performance of (A) three different ANA ELISA and (B) a F-actin ELISA to discriminate between AIH and controls (distribution of diagnoses as shown in Figure 1). Area under the curve (AUC) values are indicated.

Figure 3. Receiver-operating-characteristic (ROC) curves showing the diagnostic performance of three different ANA ELISA in comparison with ANA immunofluorescence for the diagnosis of AIH. Diagnostic performance is separately shown for cohorts from (A–B) Hamburg, (C–D) Larissa, and (E–F) Bologna. The distribution of diagnoses is shown in Figure 1. Area under the curve (AUC) values are indicated.

Figure 4. Receiver-operating-characteristic (ROC) curves showing the diagnostic performance of a F-actin ELISA in comparison with SMA immunofluorescence for the diagnosis of AIH. Diagnostic performance is separately shown for cohorts from (A–B) Hamburg, (C–D) Larissa, and (E–F) Bologna. The distribution of diagnoses is shown in Figure 1. Area under the curve (AUC) values are indicated.

590 **Figures**

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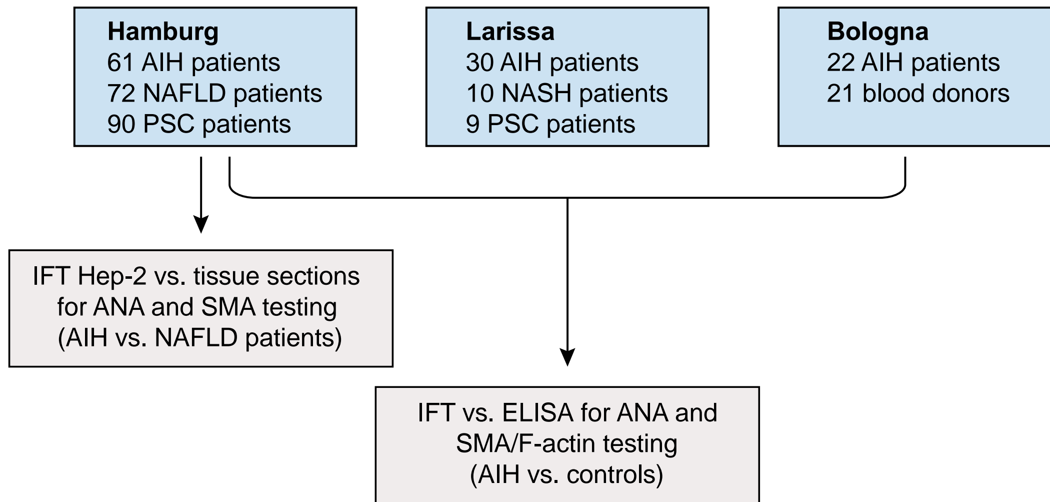

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24 594 **Figure 2:**
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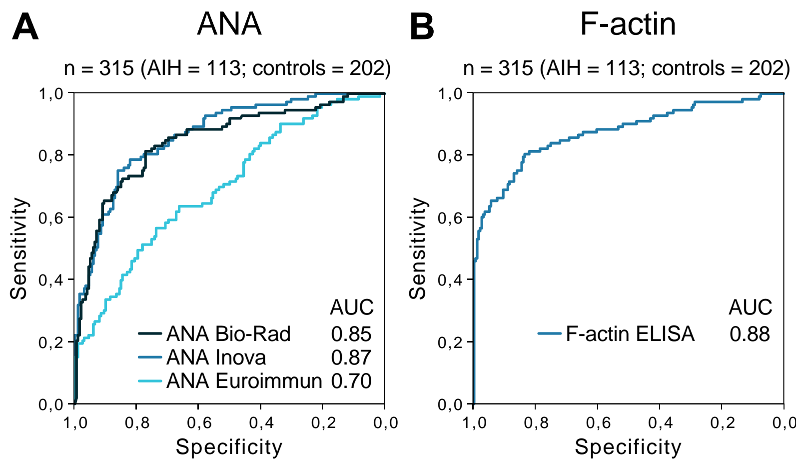

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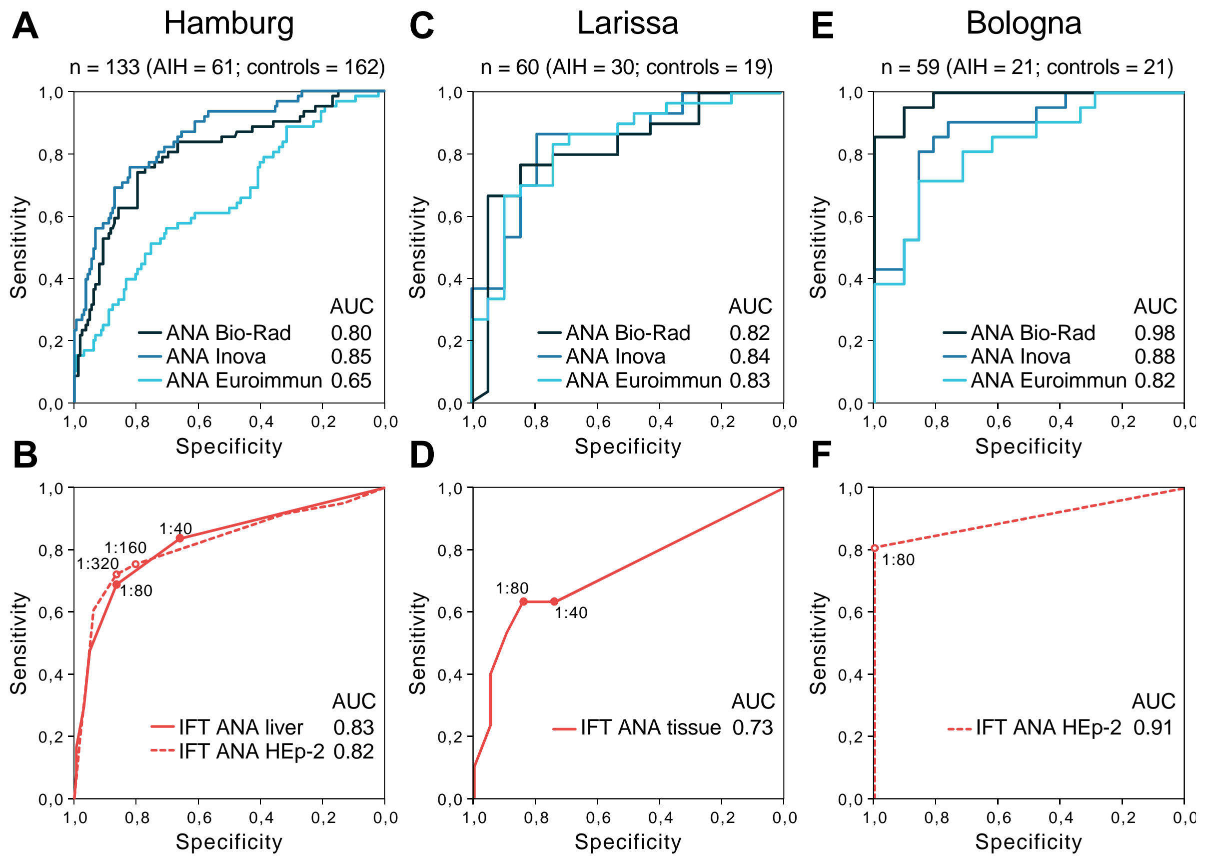
608 **Figure 3:**



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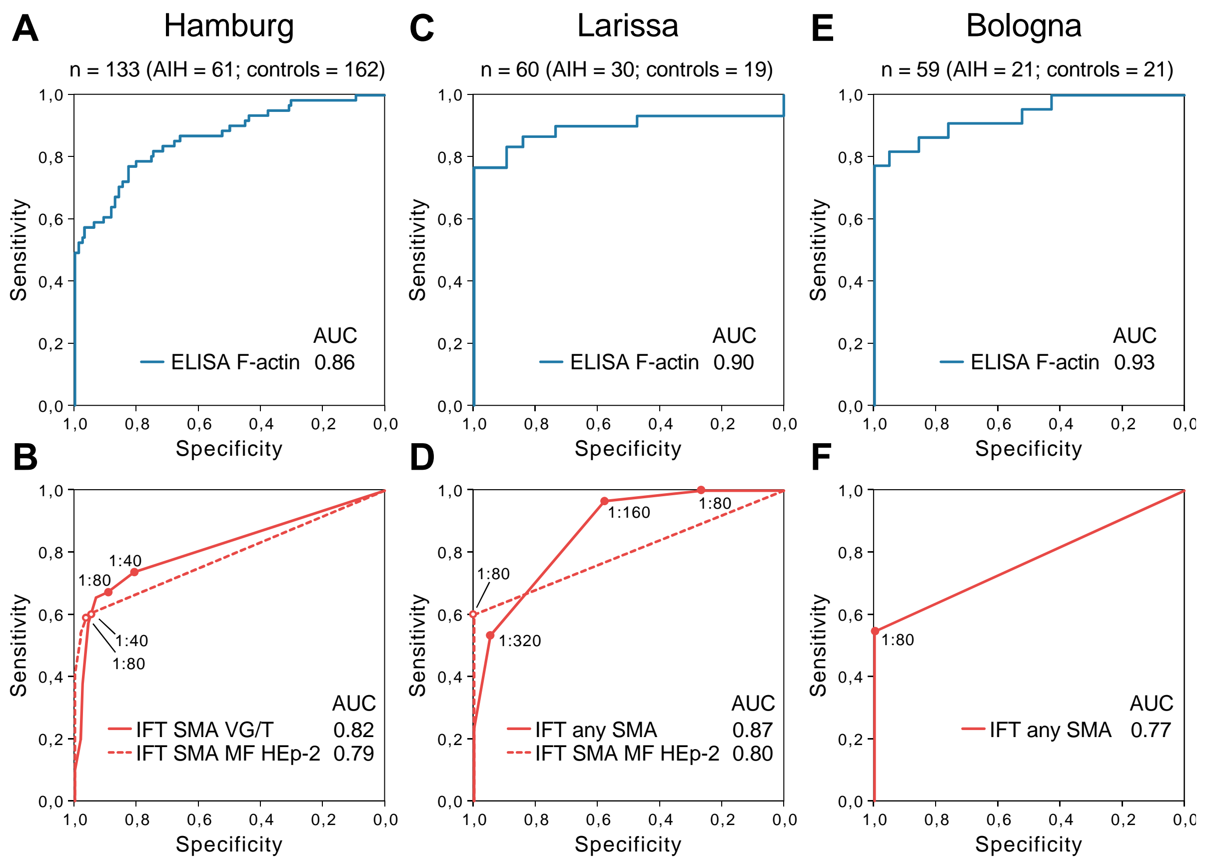

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Table 1

Table 1. Sensitivity and specificity of ANA IFT for different tissue sections

Substrate	Titer	Sensitivity (%)	Specificity (%)	PPV (%)	NPV (%)	Accuracy (%)
HEp-2 cells	1:40	95.1	8.3	46.8	66.7	48.1
	1:80	91.8	36.1	54.9	83.9	61.7
	1:160	75.4	73.6	70.8	77.9	74.4
	1:320	72.1	76.4	72.1	76.4	74.4
	1:640	60.7	87.5	80.4	72.4	75.2
Primate liver	1:40	83.6	69.4	69.9	83.3	75.9
	1:80	68.9	80.6	75.0	75.3	75.2
	1:160	47.5	91.7	82.9	67.4	71.4
	1:320	47.5	91.7	82.9	67.4	71.4
	1:640	29.5	94.4	81.8	61.3	64.7
Rat kidney	1:40	75.4	73.6	70.8	77.9	74.4
	1:80	65.6	81.9	75.5	73.8	74.4
	1:160	52.5	87.5	78.1	68.5	71.4
	1:320	47.5	91.7	82.9	67.4	71.4
	1:640	34.4	93.1	80.8	62.6	66.2
Rat stomach	1:40	78.7	70.8	69.6	79.7	74.4
	1:80	67.2	81.9	75.9	74.7	75.2
	1:160	52.5	88.9	80.0	68.8	72.2
	1:320	44.3	91.7	81.8	66.0	69.9
	1:640	36.1	93.1	81.5	63.2	66.9
Any tissue positivity (primate liver, rat kidney, rat stomach)	1:40	85.3	65.3	67.5	83.9	74.4
	1:80	73.8	77.8	73.8	77.8	75.9
	1:160	52.5	87.5	78.1	68.5	71.4
	1:320	50.8	91.7	83.8	68.8	72.9
	1:640	37.7	93.1	82.1	63.8	67.7

AIH n=61; NAFLD n=72; ANA, antinuclear antibodies; HEp-2 cells, human epithelioma-2 cells; IFT, immunofluorescence test; NPV, negative predictive value; PPV, positive predictive value.

Table 2

Table 2. Sensitivity and specificity of SMA IFT for different patterns

Substrate	Titer	Sensitivity (%)	Specificity (%)	PPV (%)	NPV (%)	Accuracy (%)
HEp-2 (microfilaments)	1:40	60.7	94.4	90.2	73.9	79.9
	1:80	59.0	98.6	97.3	74.0	80.5
	1:160	54.1	98.6	97.1	71.7	78.2
	1:320	52.5	98.6	97.0	71.0	77.4
	1:640	41.0	100	100	66.7	72.9
Kidney SMA-V (vessels)	1:40	78.7	45.8	55.2	71.7	60.9
	1:80	73.8	72.2	69.2	76.5	72.9
	1:160	68.9	80.6	75.0	75.3	75.2
	1:320	62.3	88.9	82.6	73.6	76.7
	1:640	49.2	98.6	96.8	69.6	75.9
Kidney SMA-VG (vessels, glomeruli)	1:40	72.1	70.8	67.7	75.0	71.4
	1:80	65.6	88.9	83.3	75.3	78.2
	1:160	63.9	94.4	90.7	75.6	80.5
	1:320	55.7	97.2	94.4	72.2	78.2
	1:640	36.1	100	100	64.9	70.7
Kidney SMA-VGT (vessels, glomeruli tubuli)	1:40	52.5	93.1	86.5	69.8	74.4
	1:80	49.2	93.1	85.7	68.4	72.9
	1:160	44.3	95.8	90.0	67.0	72.2
	1:320	31.2	97.2	90.5	62.5	66.9
	1:640	23.0	100	100	60.5	64.7
Kidney SMA-VG or HEp2 microfilaments	1:40	75.4	69.4	67.7	76.9	72.2
	1:80	68.9	88.9	84.0	77.1	79.7
	1:160	65.6	94.4	90.9	76.4	81.2
	1:320	62.3	97.2	95.0	75.3	81.2
	1:640	44.3	100	100	67.9	74.4
Liver (bile canaliculi)	1:40	59.0	83.3	75.0	70.6	72.2
	1:80	49.2	95.8	90.9	69.0	74.4
	1:160	42.6	98.6	96.3	67.0	72.9
	1:320	39.3	98.6	96.0	65.7	71.4
	1:640	26.2	100	100	61.5	66.2
Stomach (tunica muscularis, lamina muscularis mucosa, interglan-dular fibrils)	1:40	83.6	45.8	56.7	76.7	63.2
	1:80	75.4	72.2	69.7	77.6	73.7
	1:160	72.1	80.6	75.9	77.3	76.7
	1:320	68.9	90.3	85.7	77.4	80.5
	1:640	54.1	97.2	94.3	71.4	77.4
Any SMA positivity	1:40	86.9	37.5	54.1	77.1	60.2
	1:80	80.3	69.4	69.0	80.7	74.4
	1:160	72.1	79.2	74.6	77.0	75.9
	1:320	72.1	88.9	84.6	79.0	81.2
	1:640	60.7	97.2	94.9	74.5	80.5

AIH n=61; NAFLD n=72; HEp-2 cells, human epithelioma-2 cells; IFT, immunofluorescence test; NPV, negative predictive value; PPV, positive predictive value; SMA, smooth muscle antibodies; VGT, vessel, glomeruli, tubuli.

Table 3

Table 3. Diagnostic value of ANA and F-Actin ELISA at cut-offs recommended by manufacturers

ELISA	Assay	Cutoff	Sensitivity (%)	Specificity (%)	PPV (%)	NPV (%)	Accuracy (%)
ANA ELISA	Bio-Rad	≥ 1.0	65.5	88.6	76.3	82.1	80.3
	Inova	≥ 20	79.6	78.2	67.2	87.3	78.7
		≥ 30	69.0	86.6	74.3	83.3	80.3
	Euroimmun	≥ 1.0	22.1	95.0	71.4	68.6	68.9
F-Actin ELISA	Inova	≥ 20	81.4	82.2	71.9	88.8	81.9
		≥ 30	66.4	92.6	83.3	83.1	83.2

AIH n=113; controls n=202; distribution of diagnoses as shown in Figure 1; ANA, antinuclear antibodies; NPV, negative predictive value; PPV, positive predictive value.

Table 4

Table 4. Simplified criteria for autoimmune hepatitis

– Update of serological criteria

Variable	Cutoff	Points[1]
ANA or SMA/F-Actin	Positive[2]	1
ANA or SMA/F-Actin	Strongly positive[3]	
or LKM	≥1:40	2
or SLA	Positive	
IgG	>Upper normal limit	1
	>1.1 times upper normal limit	2
Liver histology (with evidence of	Compatible with AIH	1
hepatitis)	Typical AIH	2
Absence of viral hepatitis	Yes	2
		≥6: probable AIH
		≥7: definite AIH

[1]Addition of points achieved (maximum 2 points for autoantibodies);
[2]IFT: ≥1:40 when assessed on tissue sections; ≥ 1:80 or 1:160 for ANA when assessed on HEp-2 cells, depending on local standards. ELISA with locally established cut-offs;
[3]IFT: ≥1:80 when assessed on tissue sections; ≥ 1:160 or 1:320 for ANA when assessed on HEp-2 cells. ELISA with cut-offs established locally;
Note: if ELISA-based autoantibody assessment is negative despite high clinical suspicion of autoimmune hepatitis, IFT should be performed in addition.

Figure 1

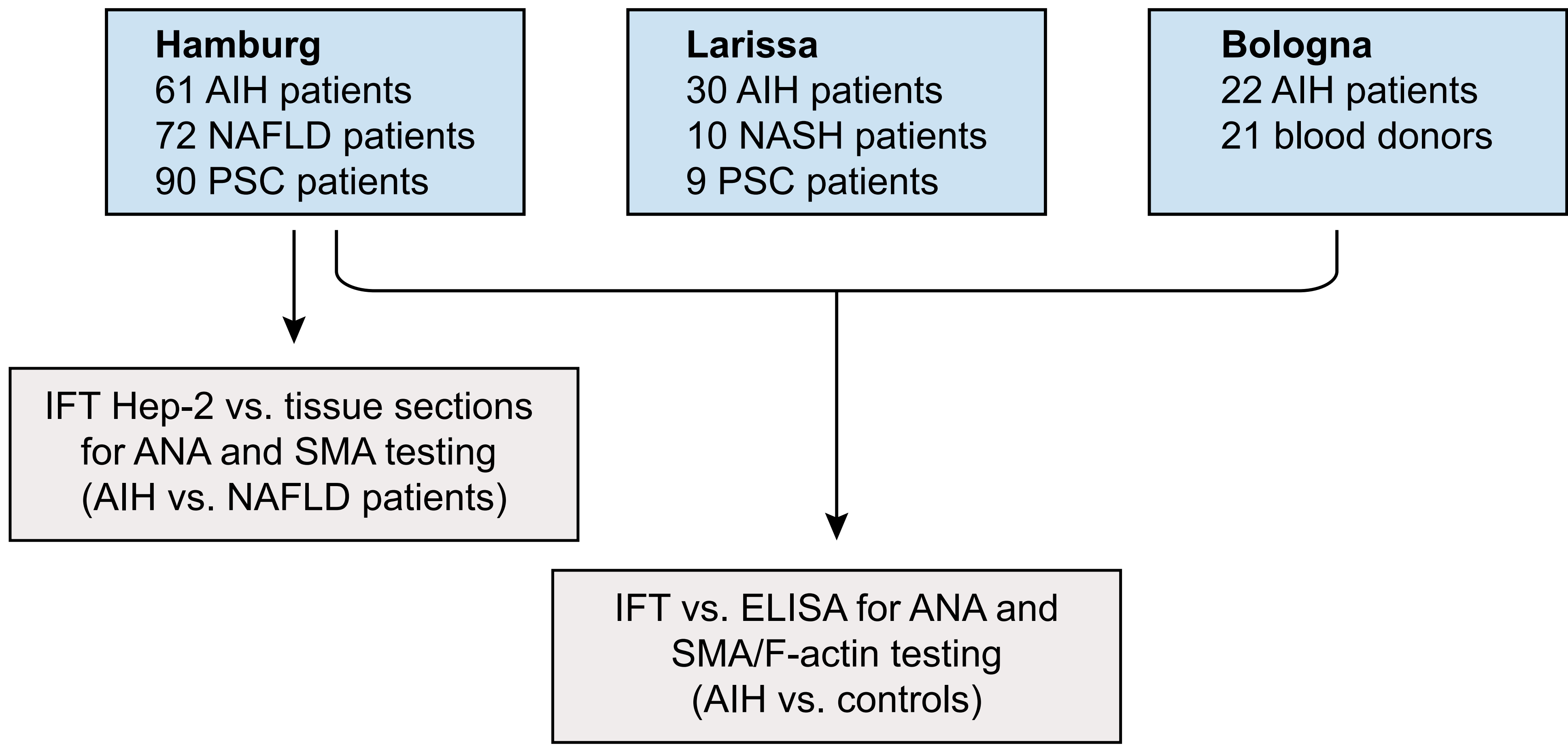


Figure 2

A

ANA

n = 315 (AIH = 113; controls = 202)

1,0
0,8
0,6
0,4
0,2
0,0

Sensitivity

AUC
ANA Bio-Rad 0.85
ANA Inova 0.87
ANA Euroimmun 0.70

1,0 0,8 0,6 0,4 0,2 0,0

Specificity

B

F-actin

n = 315 (AIH = 113; controls = 202)

1,0
0,8
0,6
0,4
0,2
0,0

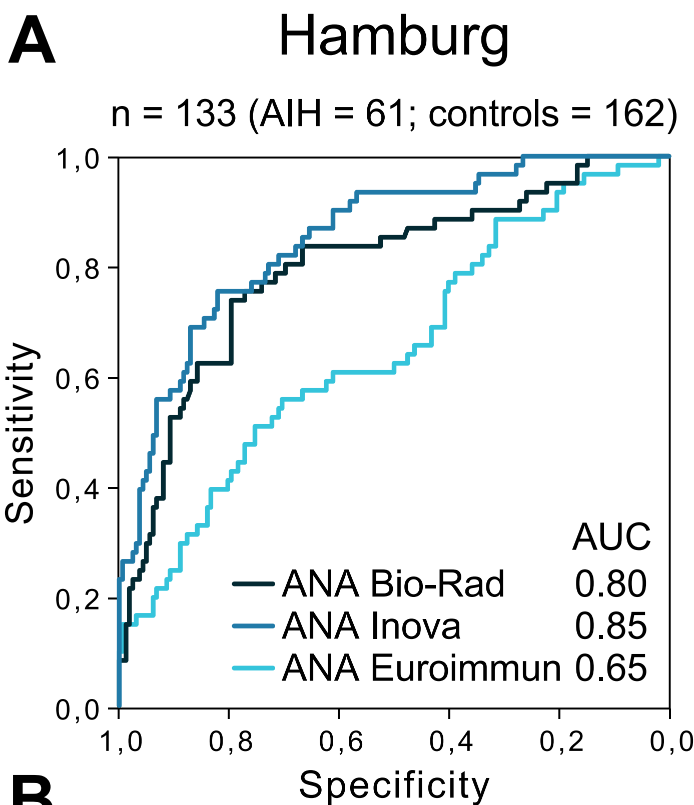
Sensitivity

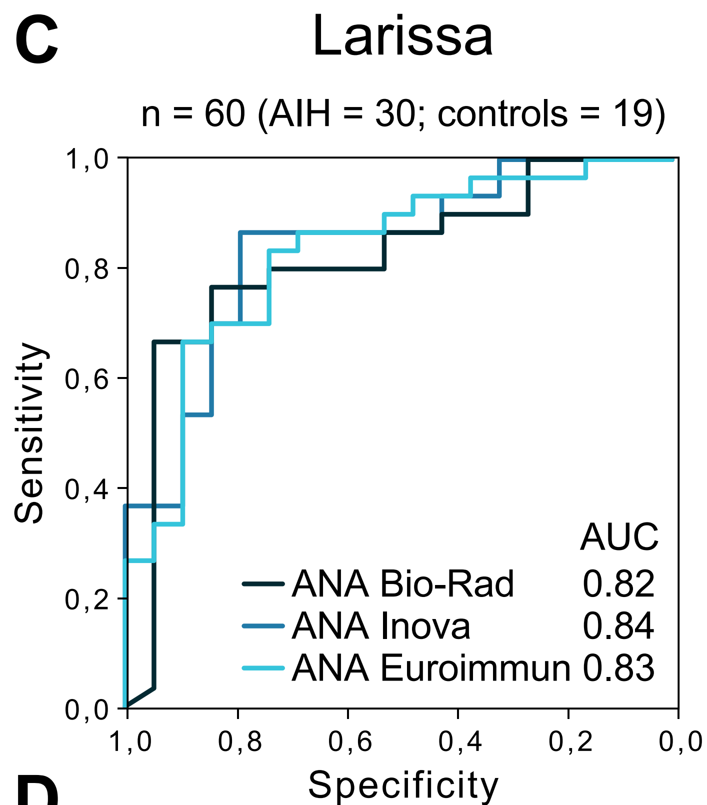
AUC
F-actin ELISA 0.88

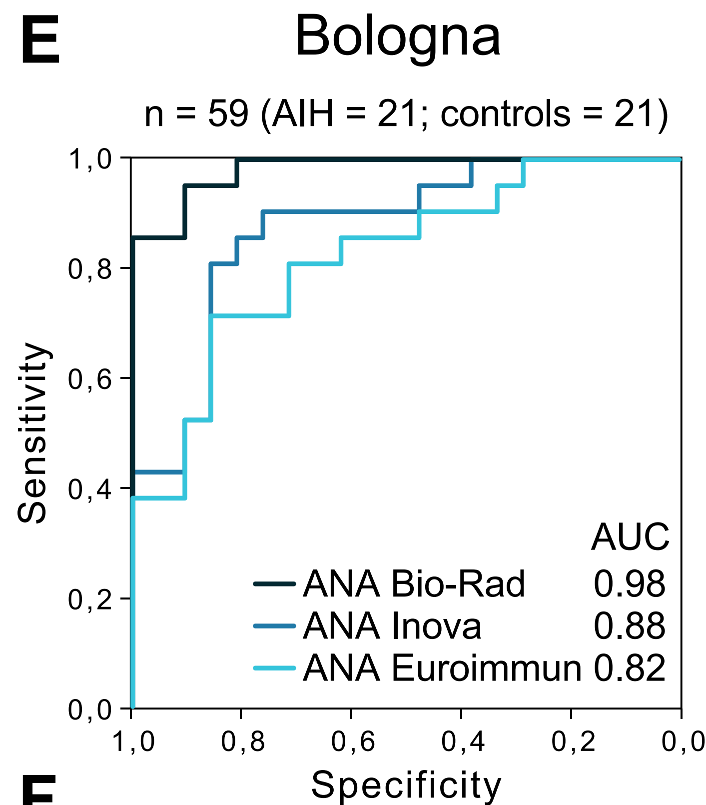
1,0 0,8 0,6 0,4 0,2 0,0

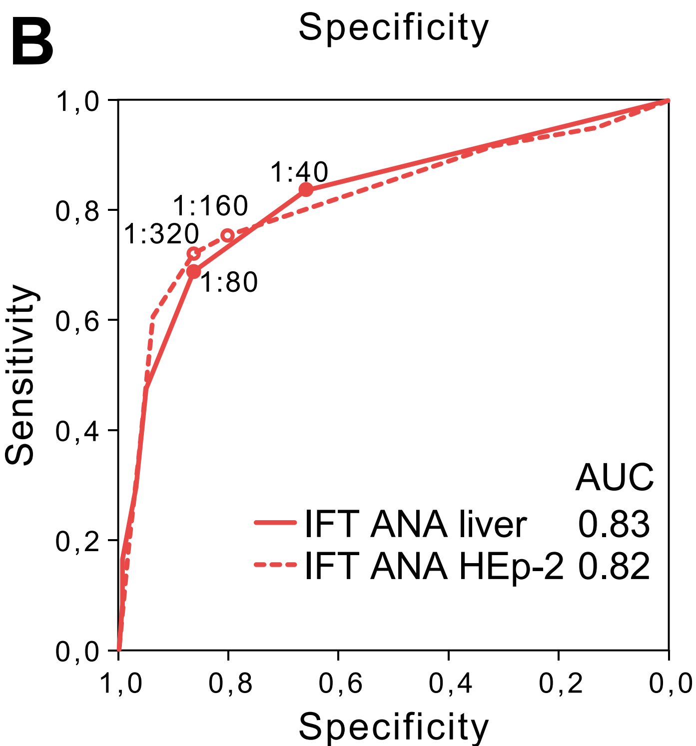
Specificity

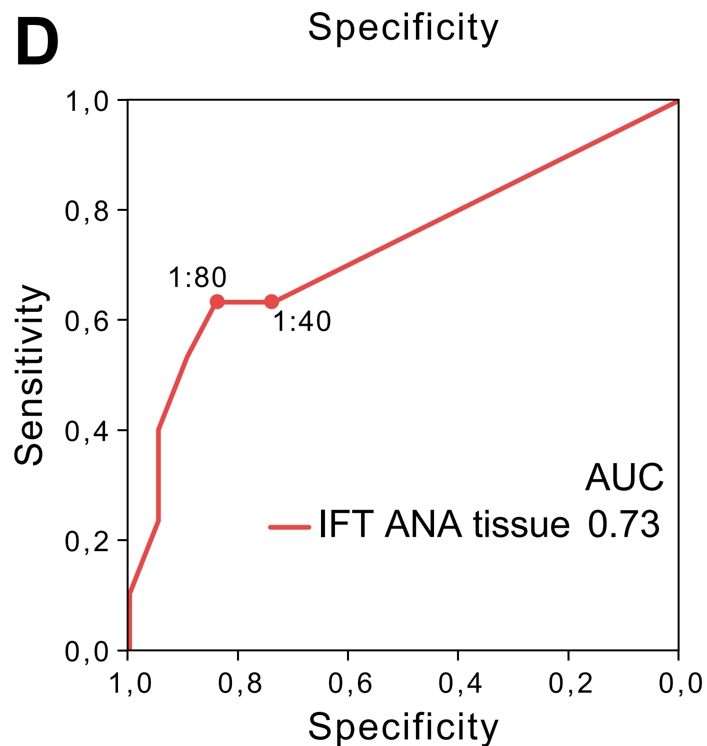
Figure 3











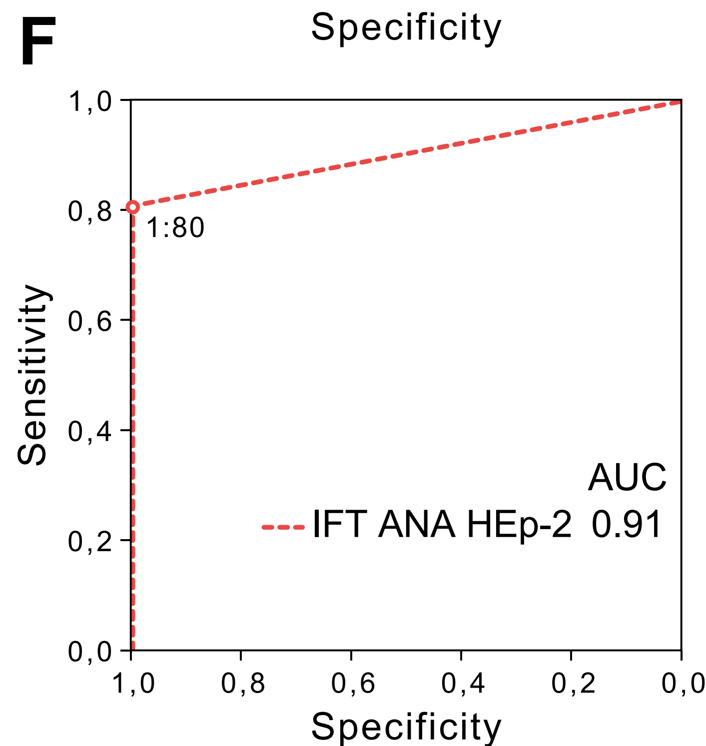
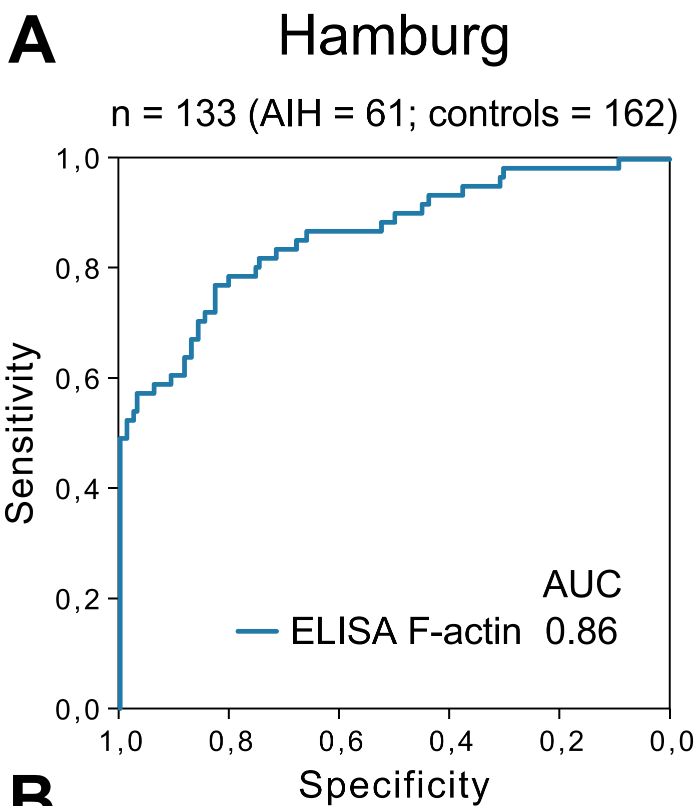
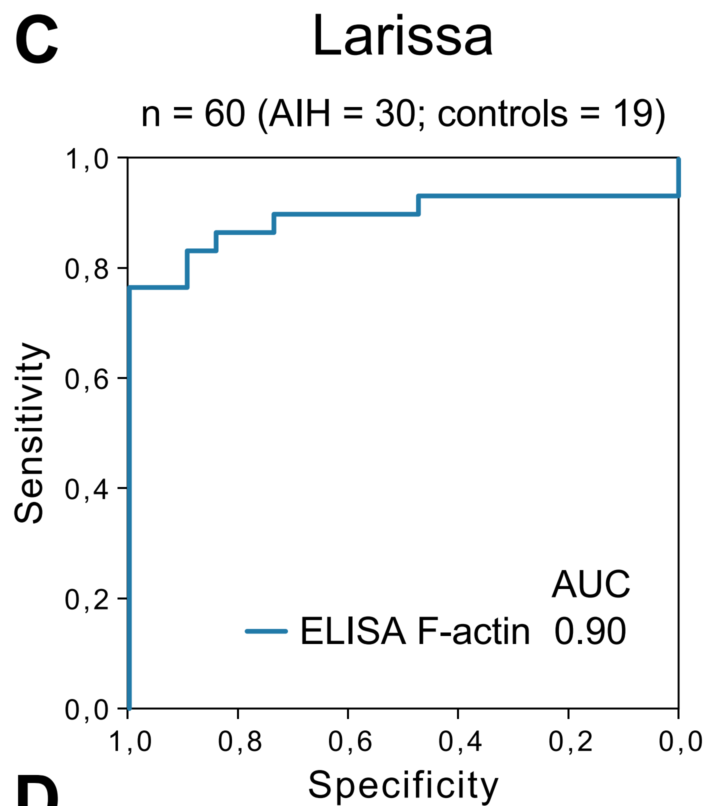
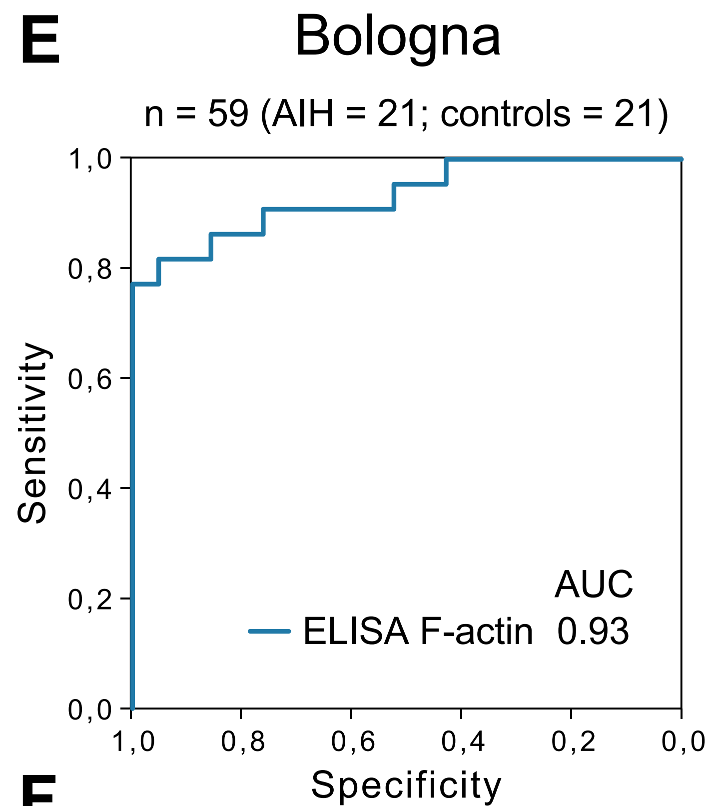
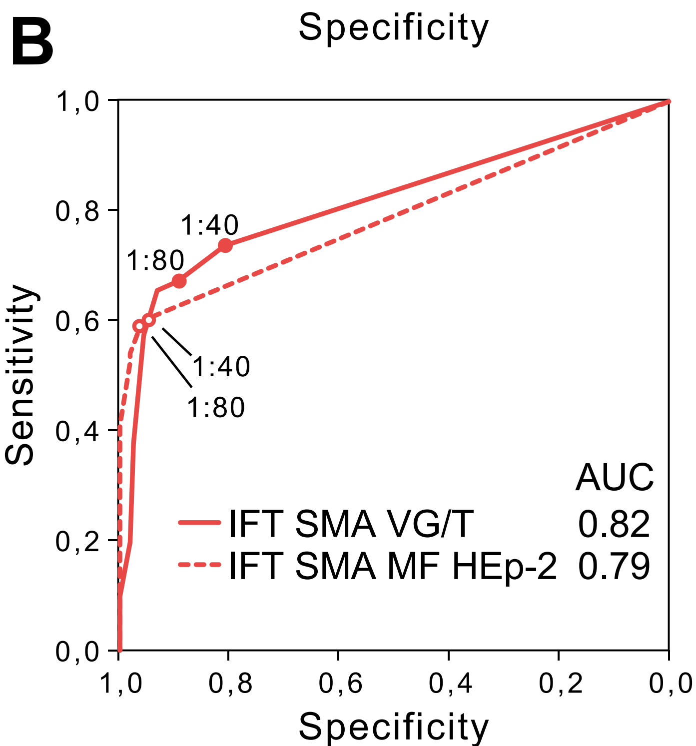


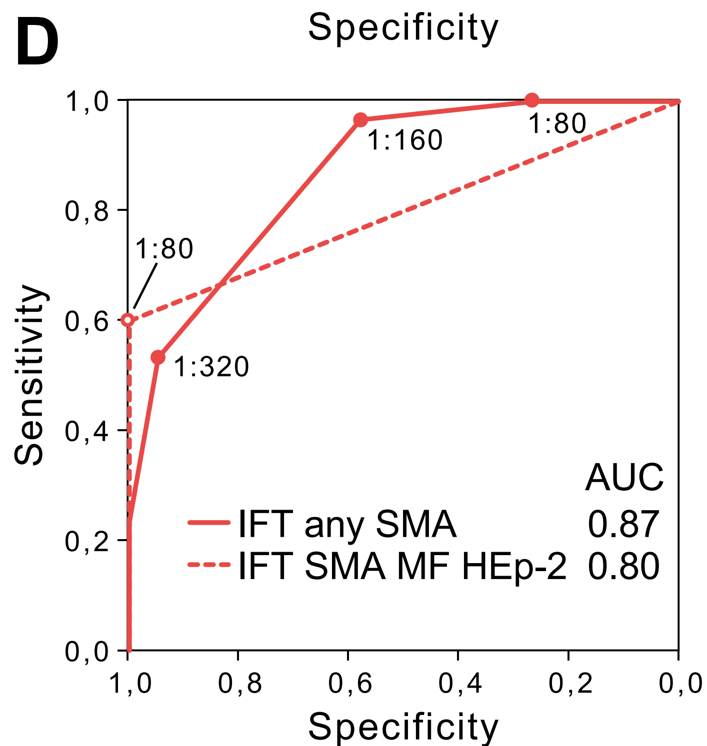
Figure 4

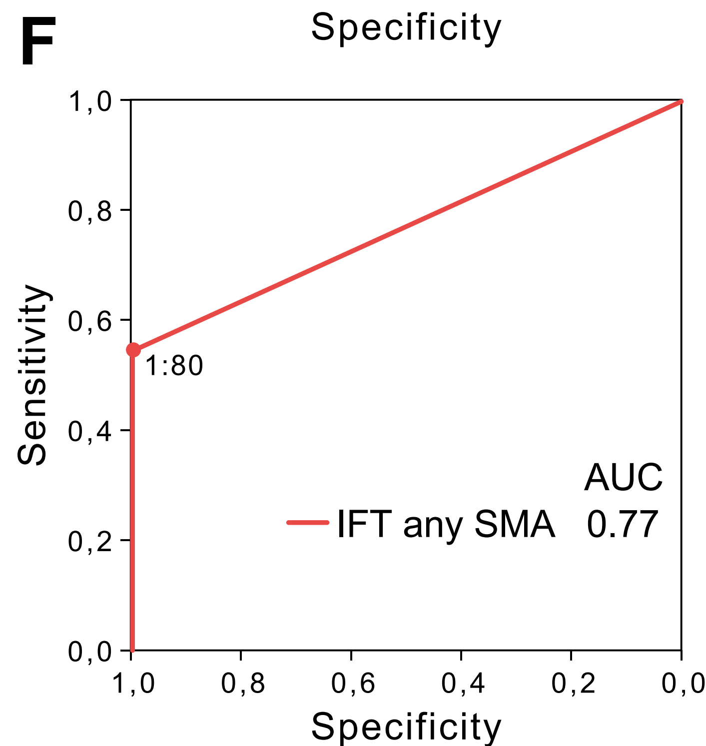




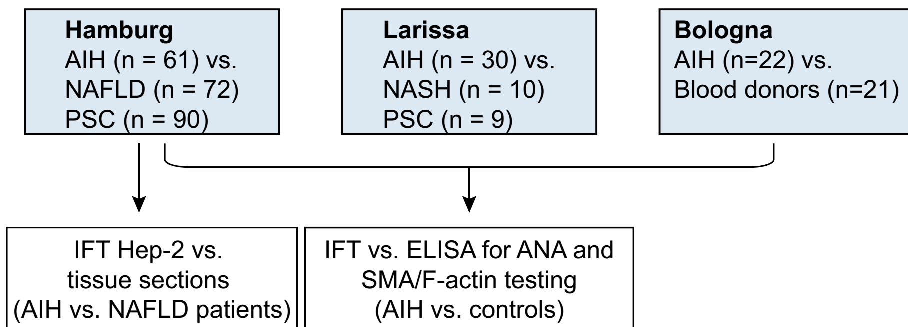




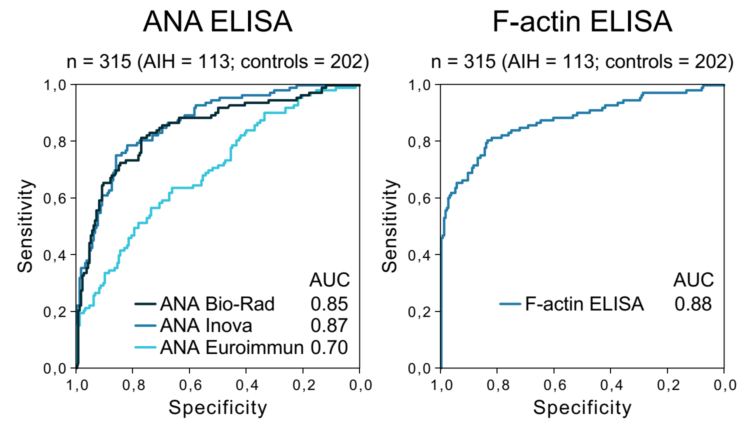




Graphical Abstract



Diagnostic performance of ANA and F-actin ELISA for the diagnosis of AIH



The simplified criteria for the diagnosis of AIH – update of serological criteria

Variable	Cutoff	Points
ANA or SMA/F-Actin	Positive[1]	1
ANA or SMA/F-Actin or LKM or SLA	Strongly positive[2] ≥1:40 Positive	2
IgG	>Upper normal limit >1.1 times upper normal limit	1 2
Liver histology (with evidence of hepatitis)	Compatible with AIH Typical AIH	1 2
Absence of viral hepatitis	Yes	2
		≥6: probable AIH ≥7: definite AIH

[1]IFT: ≥1:40 when assessed on tissue sections; ≥ 1:80 or 1:160 for ANA when assessed on HEp-2 cells, depending on local standards. ELISA with cut-offs validated locally;
[2]IFT: ≥1:80 when assessed on tissue sections; ≥ 1:160 or 1:320 for ANA when assessed on HEp-2 cells. ELISA with cut-offs validated locally;
Note: if ELISA-based autoantibody assessment is negative despite of a high clinical suspicion for autoimmune hepatitis, IFT should be performed.

Highlights

- IFT on HEp-2 cells is a valid alternative to the standard ANA assessment on rodent tissue sections in AIH when cutoffs titers are increased
- ANA ELISA and F-actin ELISA represent potential alternatives to IFT in the diagnosis of AIH
- ANA ELISA kits should include HEp-2 nuclear extracts to account for unrecognized autoantigens
- ELISA cutoffs need to be validated locally to be predictive in diagnosing AIH