EFSUMB Guidelines and Recommendations on the Clinical Use of Liver Ultrasound Elastography, update 2017 (long and short version)

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Introduction
Over the last decade the European Federation of Societies for Ultrasound in Medicine and Biology (EFSUMB) has produced a series of guidelines and recommendations regarding different ultrasound applications including:
• contrast enhanced ultrasound (CEUS) [1–5],
• dynamic contrast enhanced ultrasound (DCE-US) for quantification of tumour perfusion [6],
• elastography [7, 8],
• interventional ultrasound [9–19],
• student education [20, 21],
• pediatric use of CEUS [22, 23] and
gastrointestinal ultrasound [24, 25].

EFSUMB is working to promote high quality in ultrasound education and sustain excellent professional standards in training and practice [26]. Each of these guidelines can be considered as a chapter in a growing book. These guidelines are available on the EFSUMB website and are provided to guide both novice and expert users in performing examinations with ultrasound technology [27, 28].

Content
The first elastography guidelines worldwide were introduced and published by EFSUMB in 2013 [7, 8] followed by the WFUMB guidelines [29–33]. Most recently the first update on the 2013 published EFSUMB Guidelines and Recommendations on the clinical use of elastography, focused on the assessment of diffuse liver disease, was released. The first part (long version) of these Guidelines and Recommendations deals with the basic principles of elastography and provides an update of where the technology has changed [34]. The basic principles of elastography remain unchanged since they were outlined in the first part of the original EFSUMB and WFUMB guidelines on this subject [7, 29]. This paper therefore aims to provide an update of where the technology has changed, as of 2017, in this rapidly moving field. Sufficient recapitulations are provided to allow the present paper to be understood without reference to the earlier work, although the purpose is not to reproduce the material of the 2013 paper in detail.

Elastography guidelines
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The second part provides clinical information about the practical use of elastography techniques and interpretation of results in the assessment of diffuse liver disease, and analyzes the main findings based on published studies, stressing the evidence from meta-analyses. The role of elastography in different etiologies of liver disease and in several clinical scenarios is also discussed. This updated document is intended to form a reference and to provide a practical guide for both beginners and advanced clinical users [34, 35]. The liver is an important target organ for the use of elastography; stiffness correlates with the degree of fibrosis and indirectly with complications including portal hypertension. Transient elastography (TE), point shear wave elastography (pSWE), two-dimensional shear wave elastography (2D-SWE), strain elastography (SE) and strain-rate imaging (SRI) are discussed according to the following chapter criteria. The following chapters were prepared to include summaries of the examination procedure, number of cases for which there is evidence, measurement technique (including fasting and resting), normal values, reproducibility, quality parameters in patients and healthy subjects, the evaluation of factors other than liver fibrosis which influence liver stiffness (confounders) and comparison of results between systems. Elastography is part of a clinical decision making process. Therefore, “clinical decision making before elastography” was a chapter of its own. The evaluation of liver diseases included the initial evaluation of chronic liver disease (as prerequisites for SWE) and the evaluation of liver diseases including the fibrosis staging of chronic hepatitis C (CHC), chronic hepatitis B (CHB), non-alcoholic fatty liver disease (NAFLD), alcoholic liver disease (ALD), cholestatic liver disease, autoimmune he-
patitis (AIH) and portal hypertension. The prognostic relevance, monitoring (evaluation of) response to treatment and prediction of hepatic complications were analysed as well, according to levels of evidence (LoE), grade of recommendation (GoR) and the consensus criteria. Investigators can easily use the level of evidence to identify areas in need of additional studies. Reimbursement issues and the value of SWE in social health care systems, are also discussed.

Methodology

A steering committee was appointed, whose role was to define the general content of the guidelines, with subsequent invitation of experts from member organizations of EFSUMB, based on their publications and expertise in the different fields, to participate in the guideline development. Section leaders were selected from the steering committee; the section leaders defined the subchapters and key topics of the new guidelines sections. Literature search was performed systematically in PubMed using predefined key words and MeSH terms and, in addition, by complimentary “hand search” using reference lists of articles retrieved by systematic search. Search in principle was defined for guidelines, meta-analyses and systematic reviews, original research articles (randomized controlled trials, prospective studies, retrospective studies, case series). Evidence tables were generated for each key topic according to EFSUMB requirements. All the recommendations were judged with regard to their evidence-based strength according to the Oxford Centre for Evidence-Based Medicine Levels of Evidence and grade of recommendations (GoR) [http://www.cebm.net/oxford-centre-evidence-based-medicine-levels-evidence-march-2009].

Drafts of overview, recommendations and comments were provided by the authors for each section, revised by the members of the steering committee.

The revised drafts were submitted to the whole expert group before the expert meeting, held in London, UK on 1 July 2016. Representatives of manufacturers were invited to participate in the meeting and were allowed to comment but had no influence on the writing of the guidelines. At the expert meeting the review and recommendations were presented to the entire organ group. All evidence-based recommendations were discussed, improved, and the draft document was improved according to the level of evidence. Grade of recommendation and the level of consensus was also documented. Consensus was graded using the proposed and published system: strong consensus (> 95% of experts votes), broad agreement (> 75% – 95%) and majority consensus (> 50% – 75%). The steering committee first, and entire group of authors thereafter, reviewed and revised the draft document in a step-wise fashion for consistency and accuracy. Following the consensus meeting, comments were adapted to the final recommendations and shortened. The LoE and GoR were checked by the authors and steering committee.