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 practical advantages and disadvantages associated with each of the techniques are described, and guidance is provided on optimization of scanning technique, image display, image interpretation, reporting of data and some of the known image artefacts. The basic principles and technology for elastography were developed by the academic research community before commercial translation, and it remains a heavily researched and rapidly developing field. EFSUMB recommends that users maintain an awareness of this field.
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 records 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.

Future perspectives

Future requirements in liver elastography include the need to compare the accuracy of different technologies (TE, pSWE and 2D-SWE) in a large cohort of patients and to analyse results in terms of the aetiologies of liver diseases. Robust cut-off values must be established for each of the different systems and diseases. Combination of elastographic methods, using ultrasound with biological tests, may improve the accuracy of the evaluation of patients.

References
