Introduction

This review deals with the four sets of international contrast-enhanced ultrasonography (CEUS) guidelines that have been published from 2004 to 2013 [1–5] and discusses the clinical implications thereof, with an emphasis on daily clinical work. The intention is to focus on common clinical scenarios in which CEUS plays a decisive role and to illustrate these scenarios with typical cases. According to the World Health Organization (WHO), two thirds of the world’s population has no access to any kind of medical imaging [6]. Ultrasonography, including refinements such as elastography and CEUS, has tremendous potential to rectify this inequity [7]. We discuss this possibility in light of four sections in which we comment on each set of guidelines separately with regard to their most important aspects. In addition, pictorial explanations drawing on clinical cases are provided to emphasize selected points.

2004 Guidelines for the Use of Contrast Agents in Ultrasonography

The first generation of international CEUS guidelines was issued in 2004 by the European Federation of Societies for Ultrasound in Medicine and Biology (EFSUMB) [1]. This article was important for two major reasons. First, it presented a scientific endorsement of the clinical use of contrast agents during ultrasonography as a decisive imaging modality in the work-up of focal liver lesions (FLLs).

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and as an ideal tool for all phases of liver tumor ablation. Second, it established the acronym of CEUS to describe this technique. A search on Scopus performed on August 28, 2015 showed that the article had been cited 308 times in 135 different journals.

These guidelines provided a detailed explanation of how to detect and characterize liver tumors and how to differentiate among various benign entities based on the enhancement pattern after the administration of the contrast agents, and demonstrated that while some refinements may be necessary, a simple algorithm can be used for the most common findings (Table 1). The main criterion for malignancy on CEUS is contrast hypo-enhancement in the late phase, whereas benign lesions typically remain isoenhancing to hyperenhancing in comparison with the surrounding normal liver tissue. Benign entities can reliably be differentiated from each other by carefully scrutinizing the arterial contrast uptake phase; for instance, a globular filling-in pattern is typical for hemangioma and, along with continued late-phase enhancement, is pathognomonic for this diagnosis.

FLLs represent a very common incidental finding in modern medicine due to the frequent use of cross-sectional imaging in most high-income countries. This comes as no surprise to anyone familiar with abdominal ultrasonography. It may, however, be surprising to discover that despite the frequency of this finding, the literature contains no prospective data from clinically controlled studies describing the frequency and characterization of sonographically detected incidental FLLs. In a retrospective computed tomography (CT) study, Volk et al. [8] evaluated 1,892 examinations from a single institution that were performed for various reasons, but specifically without previously known or suspected FLL, liver cirrhosis, or underlying malignant disease. They identified a total of 100 cases with unexpected FLL, and a CT diagnosis of incidentally discovered benign liver lesions was made in 33% of those 100 cases [8]. In contrast, a somewhat surprising 17% risk of malignancy was found by Little et al. [9] in a study of 64 patients admitted over a 5-year period to a surgical referral department for the diagnostic work-up of solid liver lesions incidentally diagnosed at scanning of otherwise “relatively well” patients. The truth most likely lies somewhere between these scenarios; and in daily clinical practice FLLs may be encountered either incidentally or in follow-up scans of oncology patients. In any case, the basic questions that arise are identical: “Are the lesions benign or malignant?” and “If the lesion is malignant, then what kind of malignancy is it?”

Diagnostic ultrasonography has several important qualities that cannot be equally or easily achieved using CT or magnetic resonance imaging (MRI). Among these are features crucial to the two questions above, including spatial and, in particular, temporal resolution, as well as the practical aspects of real-time examinations and scanner availability. In three different European multicenter studies, CEUS was found to be comparable to contrast-enhanced CT or MRI [10–16]. The most extensive material from the German Ultrasound Society drew on 1,349 patients and reported a positive predictive value of 95.4% and a negative predictive value of 95.7% in differentiating between malignant and benign lesions [10]. In two 2011 meta-analyses of CEUS versus contrast-enhanced CT and MRI, the overall performance of the three modalities was statistically equal, with reported sensitivity and specificity ratios ranging from 81% to 89%. Interestingly, however, CEUS had the highest diagnostic odds ratio [17,18].

In the most recently published systematic review from the National Institute for Health Research of the United Kingdom [19], which identified a total of 854 references and included 21 studies, Westwood et al. [19] concluded that SonoVue (Bracco, Milan, Italy) CEUS can provide similar diagnostic performance to contrast-enhanced CT and MRI for the major clinical applications: (1) the characterization of FLLs detected in the sonographic surveillance of cirrhosis patients, (2) the detection of liver metastases in patients with colorectal cancer, and (3) the characterization of incidentally detected FLLs. They made a strong statement that deserves the

| Type                        | Arterial phase (10–20 sec) | Portal phase (30–45 sec) | Late phase (>120 sec) |
|-----------------------------|----------------------------|--------------------------|-----------------------|
| Benign                      | Hemangioma                 | Globular enhancement    | Centripetal filling   | Iso-/hyper-enhancement |
| Focal nodular hyperplasia   | Hyper-enhancement spoke wheel appearance centrifugal filling | Moderately hyper-/iso-enhancement | Is-o-enhancement scar (40%) |
| Hepatocellular adenoma      | Hyper-enhancement           | Transition               | Iso-enhancement       |
| Malignant                   | Hepatocellular carcinoma   | Hyper-enhancement        | Iso-/slightly hypo-enhancement | Hypo-enhancement |
| Metastasis, hypovascular    | Hyper-enhancement with/without central necrosis | Iso-/slightly hypo-enhancement | Strong hypo-enhancement |
| Metastasis, hypovascular    | No enhancement or peripheral rim | Iso-/slightly hypo-enhancement | Strong hypo-enhancement |

Time ranges in the parentheses are the delayed time for each imaging after contrast agent injection.

Table 1. Algorithm for using contrast-enhanced ultrasonography to characterize focal liver lesions based on the enhancement pattern after the administration of contrast agents.
full attention of medical professionals working in this field: “If the main use of liver imaging is considered to be the rapid rule-out of malignancy, equivalent diagnostic performance may be sufficient for SonoVue CEUS to be preferred over other imaging modalities. A potential advantage of using SonoVue CEUS would be the option of completing the assessment at the same time as the initial unenhanced ultrasonography.” These conclusions are broadly supported [20–22] and elegantly framed by Friedrich-Rust et al. [23] in the following statement: “Use CEUS as first method of choice for the diagnostic work up of focal liver lesions if B-mode and Doppler-ultrasonography are not conclusive.”

In addition, CEUS has an excellent safety profile [24] and patient compliance, as well as no issues regarding organ function or radiation exposure. Therefore, when available, CEUS should be the first method to answer the above question of “Are the lesions benign or malignant?” However, CEUS cannot provide a reliable answer to the question “If the lesion is malignant, then what kind of malignancy is it?” If a histological diagnosis is deemed necessary for treatment planning, a tissue sample must be obtained, in which case an ultrasound-guided biopsy is the obvious choice [25]. Fig. 1 (with Video clips 1–3) provides a pictorial illustration of this concept of the clinical work-up of newly diagnosed liver lesions if the scenario includes the suspicion of malignant disease. In the case of malignancy, a standard protocol for staging and treatment planning should always include full-body imaging modalities such as contrast-enhanced CT or MRI according to the specific malignancy.

Fig. 1. A 60-year-old woman referred for an ultrasonography to rule out gallstones incidentally diagnosed with two focal solid lesions in the right liver lobe located laterally in segment 6 and very closely separated.

A. The two lesions show different appearances on B-mode imaging; one is hyperechoic (arrows) with slightly posterior enhancement and peripheral lie and the other is echo-poor with a halo around it (arrowheads). From the B-mode imaging alone, one would suspect the echo-rich lesion to be a hemangioma and the echo-poor lesion to be a metastasis. A routine contrast-enhanced ultrasonography was performed to further characterize these lesions. B–D. The hyperechoic lesion shows globular filling-in contrast enhancement, whereas the hypoechoic lesion demonstrates arterial hyperenhancement (B) and contrast wash-out pattern on portal (C) and late phase (D) images, thus confirming the B-mode diagnosis of hemangioma and metastasis. Since the patient had no known primary cancer, the institutional tumor board decided that an ultrasound-guided biopsy was indicated, and a TruCut 1.2-mm core biopsy provided the diagnosis of metastatic hepatic adenocarcinoma of colorectal origin.
in question.

A more common problem, however, is incidentally diagnosed, but sonographically benign liver lesions and the further clinical work-up, if any, of such lesions. In other words, the question is how to handle the unexpected finding of an FLL that, both sonographically and with respect to the patient’s symptoms and medical record, is deemed a benign finding. Here, as in many other times in life, a good piece of advice is to “keep it simple,” and the simplest rule one can apply is to follow the guidelines. The current guidelines are identical to the 2004 guidelines with regard to this issue, and state that CEUS is indicated for “incidental findings on routine ultrasonography.” This recommendation makes no further differentiation, and no doubt is left as to whether it is indicated in all cases, including the typical benign B-mode appearance. In the scenario of an incidentally detected lesion, this means that, unless the lesion is a sonographically typical cyst, a CEUS exam should be

Fig. 2. A 53-year-old man with no history or clinical suspicion of cancer referred to rule out gallstones.
A. B-mode transverse sonogram shows two hyperechoic lesions (between crosshairs) in both lobes of the liver. B. Another hyperechoic lesion (arrowheads) is defined in lower section of right hepatic lobe. C–E. Contrast-enhanced ultrasonography (CEUS) sequences illustrate arterial contrast uptake with very early (before normal liver tissue) enhancement of the centrally located lesion with a centrifugal pattern, a feeding artery consistent with the focal nodular hyperplasia (FNH) arterial pattern, and the slower centripetal globular filling-in enhancement pattern of the other two lesions (one in each liver lobe), consistent with the arterial pattern of hemangioma.
performed. This is the simplest and safest rule to follow, and it is also exactly what the guidelines have recommended from the first set of guidelines issued in 2004 to the latest guidelines issued in 2012. If one follows this simple rule, it is not necessary to establish complicated algorithms to determine whether an incidental FLL should be further characterized on B-mode ultrasonography in order to reach a final decision or if a CEUS study should be next step.

However, many experienced ultrasonography diagnosticians, including myself, feel that there is an issue with hemangiomas and that this issue deserves further commentary. Hemangioma is the most common tumorous condition of the liver and was found in an autopsy study [26] to be present in no less than 20% of men, although with a mean size of 5.2 mm, meaning that many of them were very small and most likely would have remained undiscovered on routine ultrasonography. In the abovementioned study of patients referred for further assessment [9], hemangiomas constituted 52% of the lesions. Hemangioma is also the most common benign tumor and the most common echo-rich tumor of the liver [27]. Most hemangiomas have very typical B-mode features, including a well-demarcated, round, and uniformly echo-rich appearance with a usually peripheral lie, posterior enhancement, and a diameter less than 3 cm. Provided that the tumor is detected in a patient without any history of cancer or suspicion thereof, a diagnosis of benignity can be established using conventional ultrasonography with an estimated probability of >95%, and no further imaging is indicated [20,21,27]. The definitive characterization of every incidentally found hemangioma with CEUS would lead to a substantial workload and disproportionate cost. Moreover, all such efforts would be in vain because, according to Dietrich et al. [21], neither CEUS nor any other imaging modality would provide the correct diagnosis with a higher certainty than the ultrasonography techniques already available [9]. In approximately 30% of cases, however, hemangiomas demonstrate atypical sonographic features, and a CEUS study should be the next step for further characterization and to rule out malignancy. The same is true in the clinical scenario where a B-mode exam demonstrates an apparently typical hemangioma in an oncological patient or a patient suspected of malignancy. In such cases, CEUS is also the recommended next step. Fig. 2 (with Video clips 4–7) shows a case with a coincidental diagnosis of multiple simultaneous large hemangiomas in coexistence with focal nodular hyperplasia.

Guidelines and Good Clinical Practice Recommendations for CEUS: 2008 Update

The second generation of guidelines was published as an article in the February 2008 issue of European Journal of Ultrasound/Ultraschall in der Medizin [2]. The most important element of this update was the inclusion of extrahepatic applications for CEUS. A search on Scopus on August 28, 2015 showed that the 2008 CEUS guidelines article had been cited a total of 459 times in 139 journals, thus exceeding the citations of the 2004 guidelines by 50%. In addition to the initial indications, this article presented the best-documented newly accepted indications for the use of contrast agents, which were CEUS of the kidney and urinary tract, including the vesico-ureteric reflux, pancreas, and cerebral circulation, and CEUS in abdominal blunt trauma. Guidelines dealing with CEUS of
liver lesions and CEUS in ablative procedures were updated with the newest data available and the technical aspects were adjusted, but the algorithms for the interpretation of enhancement patterns remained fundamentally unchanged.

One application stands out in comparison with the others, and that is CEUS in blunt abdominal trauma. First, the trauma patient is one of the most commonly encountered clinical scenarios. Second, this new CEUS indication has a tremendous potential for reducing the population-level exposure to ionizing radiation. Third, CEUS for the evaluation of trauma patients has proven beneficial in clinical use on a broad scale. Prior to its inclusion in the 2008 guidelines, only few articles with small study populations were available [28–30]. In recent years, several larger studies have been published confirming that positive outcomes were associated with this technique [31–40], including one prospective multicenter trial by Catalano et al. [41] that enrolled 156 patients with contrast-enhanced CT used as the gold standard, and found that per-patient sensitivity increased from 79% for baseline ultrasonography to 94% for CEUS. In another study of 256 patients [40] admitted to an emergency department with low-energy blunt abdominal trauma, Sessa et al. [40] found that CEUS had a much higher sensitivity than baseline ultrasonography in the detection of abdominal organ lesions (59% vs. 96%) and in the diagnosis of hemoperitoneum (91% vs. 95%).

In a pediatric study of 76 children [42] admitted with isolated low-energy abdominal trauma, Menichini et al. [42] reported that the sensitivity and overall accuracy of CEUS in diagnosing solid organ lesions was 100%, as compared to the discouraging values of 39% and 44% for ultrasonography alone, with both sets of outcomes evaluated in comparison to contrast-enhanced CT. It was, however, noted that CT was more accurate than CEUS in identifying prognostic factors such as urinoma and active bleeding.

These conclusions concur perfectly with the recommendations found in the EFSUMB CEUS guidelines on non-hepatic applications published in 2012 [5], which place a special emphasis on being aware of the availability of CEUS as an alternative to CT for patients with low- to moderate-energy blunt abdominal trauma as a supplement to the FAST protocols, especially in the pediatric population and as follow-up imaging modality. However, emphasis was also put on the fact that for high-energy trauma, CT is the modality of choice and that in the case of CEUS-positive findings, contrast-enhanced CT should most likely be performed to exclude active bleeding and obtain a better overall view of the entire abdomen [40,42]. Fig. 3 (with Video clip 8) and Fig. 4 (with Video clips 9, 10) illustrate the CEUS evaluation procedure in trauma cases.

The EFSUMB Guidelines and Recommendations on the Clinical Practice of CEUS: 2011 Update on Non-Hepatic Applications

This non-hepatic component of the third generation of the CEUS guidelines was published in the February 2012 issue of European

![Fig. 3. A 26-year-old man referred for 2 days of pain in the left flank after a physical encounter in an ice hockey match. The patient's condition was stable, his blood pressure was normal, his hemoglobin levels were normal, and he was not in severe pain. The incident was categorized as moderate blunt trauma, and a contrast-enhanced ultrasonography (CEUS) using SonoVue was performed instead of contrast-enhanced computed tomography. A. B-mode ultrasonography in the left flank vaguely gives the impression of a minor intrasplenic hematoma (arrow) and demonstrates a tiny amount of perisplenic fluid collection. B. CEUS of the left flank clearly shows a minor hematoma in the lower pole of the spleen (crosshairs). The patient was treated conservatively and discharged after 2 days. Follow-up, including B-mode imaging and CEUS, showed healing. This figure illustrates the use of CEUS in minor to moderate blunt trauma.](image-url)
The key importance of this document was the fact that it placed further emphasis on the inclusion of extra-hepatic applications for CEUS. In addition, each recommendation was graded with respect to type and level of evidence. A search on Scopus on August 28, 2015 showed that the non-hepatic CEUS guidelines had been cited a total of 252 times in 104 different scientific journals.

The diversity of applications covered in this document is huge, and while many ultrasonography professionals may see a range of different clinical entities, no single expert is likely to master the evaluation of all entities. One point that deserves special mention with respect to the clinical use of contrast agents is the application of CEUS in pediatric patients, as it represents a unique medico-legal challenge. This application is controversial and is considered an ethical dilemma by many clinicians. No ultrasound contrast agent has been approved for pediatric use, but in many clinical situations, CEUS could substitute for an alternative imaging modality that would likely include exposure to ionizing radiation or the use of potentially harmful iodinated contrast agents. Most drugs are not specifically approved for pediatric use, but when indicated, most physicians do not hesitate to administer the relevant drug off-label after informed consent from the parents. Indeed, if a necessary medication is not administered out of fear of using an off-label drug, the physician may face serious medico-legal and, it should be added, ethical consequences. Additionally, it should be remembered that most indications mentioned in the non-hepatic guidelines represent off-label use of contrast agents and the CEUS examination is performed under the responsibility of the physician. The benefit of reducing ionizing radiation is obviously most important in children and fertile women, but should always be weighed against its possible disadvantages in the actual clinical context of the individual patient. This important question has been discussed in the literature [43,44], and while there is no easy way out of this medico-legal dilemma, it should always be borne in mind that the physician’s top priority must be the patient’s well-being. For reasons that are not always clear, children admitted acutely in the middle of the night with abdominal pain may sometimes undergo a CT scan to rule out serious conditions that would require surgery, which may result in the diagnosis of an incidental lesion that eventually requires further work-up in the morning. Incidentally, detected focal splenic lesions are one such clinical scenario that is frequently encountered and represents a particular dilemma. Tumors of the spleen are rare findings, and although most of these lesions are hemangiomas or hamartomas, their differentiation from malignant tumors must be confirmed before they can be disregarded. The spleen has a similar sonographic appearance to the liver and this fact sometimes brings about the misperception that focal spleen lesions have similar contrast uptake to FLLs and can therefore be interpreted according to the algorithm used for the liver (Table 1). Unfortunately, this is not the case. Although the spleen, like the liver, actually does have sinusoids, a very important difference from the liver is that the spleen has no equivalent to the portal vein and thus no dual blood flow.
supply. Further complicating matters, unfortunately, benign spleen lesions frequently show wash-out in the parenchymal phase, and the fact that this is a major indicator of malignancy in the work-up of FLLs may lead to the false-positive diagnosis of a malignant splenic tumor [45,46]. If presented with an otherwise healthy patient with a rare incidental finding that is most often benign but may lead to an erroneous diagnosis of malignancy on CEUS, it would seem prudent to perform a follow-up CEUS one and three months later. If malignancy still cannot be ruled out with certainty, positron emission tomography-CT has been suggested as a definitive step [47] before performing ultrasound-guided biopsy with its low but not negligible risk. Fig. 5 (with Video clips 11, 12) demonstrates the clinical problem of the false-positive CEUS diagnosis of a malignant spleen lesion based on parenchymal hypoenhancement. The final diagnosis, verified by an ultrasound-guided biopsy, was the benign finding of a hamartoma.

Fig. 5. A 57-year-old man with abdominal pain with coincidental focal lesion of the spleen diagnosed on contrast-enhanced computed tomography (CECT). The patient had no previous history of cancer. However, in a subsequent work-up, contrast-enhanced ultrasonography (CEUS) found the lesion suspicious and the local tumor board decided to refer the patient to our unit for a percutaneous biopsy to rule out malignancy. A. CECT from the referring tumor board shows a focal hypodense spleen lesion (arrow). B. B-mode ultrasonography performed in our unit demonstrates a slightly hypoechoic lesion (arrow) in the spleen. C. CEUS during the arterial phase demonstrates enhancement of the lesion, which is slightly delayed and hypoenhancing (arrows) compared to the surrounding normal spleen tissue. D. CEUS demonstrates strong hypoenhancement (arrows), also known as wash-out, in the late parenchymal phase. From CEUS-guided biopsy, the final diagnosis was splenoma, which is a benign hamartoma-like tumor of the spleen. In conclusion, CEUS wash-out is not indicative of a strong suspicion of malignancy, and a clinical examination and positron emission tomography-computed tomography should be considered before biopsy in patients with no other indications of malignant disease. This figure illustrates that wash-out in spleen lesion CEUS is not equivalent to the CEUS of focal liver lesions and cannot be taken as a strong indicator of malignancy.
Guidelines and Good Clinical Practice Recommendations for CEUS in the Liver: Update 2012: A WFUMB-EFSUMB Initiative in Cooperation with Representatives of AFSUMB, AIUM, ASUM, FLAUS and ICUS

This third generation of CEUS liver guidelines was published simultaneously in the February 2013 issues of Ultrasound in Medicine and Biology and European Journal of Ultrasound/Ultraschall in der Medizin [3,4]. The key importance of this document is the fact it presented a worldwide consensus from a wide and truly representative range of medical professional bodies on the usage of CEUS and the interpretation of enhancement patterns with clinically approved contrast agents for liver applications. The article describes the use of available different contrast agents licensed across the world for use in the work-up of clinical scenarios related to the liver.

These contrast agents are all fluorocarbons, sometimes referred to as perfluorocarbons, and also sometimes called organofluorine compounds. They are biologically inert gases at atmospheric pressure that are encapsulated in a lipid membranous shell. Thus, they do not interact with metabolism, can be administered regardless of renal and liver functionality, and have an extremely high safety profile, as specifically documented for SonoVue [24] (marketed as Lumason in the United States but not licensed for the liver), which is a sulfur hexafluoride introduced in 2001 by Bracco SpA (Milan, Italy) and licensed in Europe, China, India, Korea, Hong Kong, New Zealand, Singapore, and Brazil. The other two contrast agents are chemically very closely related and marketed under the names Definity/Luminity, A B

**Fig. 6.** A 77-year-old woman under surveillance for cirrhosis.

A. B-mode and color Doppler ultrasonography demonstrate an interim nodule (crosshairs) in the right hepatic lobe. B. Sonazoid contrast-enhanced ultrasonography (CEUS) shows arterial hyper-enhancement (arrows) with earlier onset than the surrounding liver tissue. C. Sonazoid CEUS-guided 1.2-mm TruCut biopsy of an isoenhancing lesion in the postvascular phase was performed at 13 minutes post-contrast. The biopsy revealed well-differentiated hepatocellular carcinoma. D. An image from a Sonazoid CEUS shows sustained isoenhancement in the postvascular phase at 31 minutes post-contrast. This figure illustrates the fact that with Sonazoid CEUS of well-differentiated hepatocellular carcinomas, in approximately 50% of cases, no enhancement defect is noted at 30 minutes post-contrast, potentially warranting a biopsy.
which is an octafluoropropane compound introduced in 2001 by Lantheus Medical (Billerica, MA, USA) and licensed in Canada and Australia, and Sonazoid, a perfluorobutane introduced in 2007 by Daiichi-Sankyo (GE, Tokyo, Japan) and licensed in Japan and South Korea, and recently also in Denmark and Norway.

A search on Scopus on August 28, 2015 showed that the 2012 update of the liver CEUS guidelines had been cited a total of 170 times in 97 scientific journals, with 80 citations in 52 journals of the version of the article printed in Ultrasound in Medicine and Biology and 90 citations in 45 journals of the version printed in European Journal of Ultrasound/Ultraschall in der Medizin. Compared to the two previous generations of CEUS guidelines, this most recent update provided recommendations on two new topics: information related to Sonazoid and a recommendation to use CEUS for biopsy guidance to increase the retrieval rate of percutaneous ultrasound-guided biopsy and to reduce the frequency of negative microscopy results [48–52].

Sonazoid, remarkably and completely unlike the other two contrast agents, has a so-called postvascular phase in which the contrast agent leaves the vascular bed and enters the parenchymal interstitium. This postvascular phase starts 10 minutes after injection and may last more than 30 minutes, and is caused by the Kupffer cell uptake of this particular contrast agent, which is very similar to that of the liver-specific MRI contrast agents often referred to as superparamagnetic iron oxide (SPIO) [53]. This mechanism ensures the uniform enhancement of normal liver tissue. Sonazoid, according to Kudo et al. [54–56] has been used with particular frequency in the work-up of hepatocellular carcinoma (HCC) in the cirrhotic liver because the typical HCC contains no Kupffer cells and thus stands out as an enhancement defect in the postvascular phase [57]. Unfortunately, however, between 50% and 70% of well differentiated HCCs—corresponding to approximately 10%–15% of HCCs in the cirrhotic population—may not demonstrate this typical enhancement defect [58,59], and in this situation a biopsy may be indicated to achieve the correct diagnosis [56,60,61]. Fig. 6 (with Video clips 13–15) demonstrates a clinical scenario such as this, in which a CEUS-guided biopsy was performed utilizing Sonazoid. The benefit of CEUS-guided biopsy was reported by Bang et al. [50] as early as 2000 and later was proven by Wu et al. [49] to increase the diagnostic accuracy rate by up to 10% by directing the biopsy towards contrast-enhanced areas. Schlottman et al. [48] suggested a different approach, in which CEUS was used to identify previously non-visualized lesions that were more easily accessible for biopsy, and this technique later received support from Yoon et al. [51,52].

In addition to the work-up of liver lesions, which has been described in detail above, the single other major indication for CEUS that has existed since the initial development of this technique is assistance in percutaneous ablative procedures. In many institutions around the world, ultrasonography is the preferred guidance modality for ablation. However, with regard to treatment evaluation, it is not sufficient to examine morphology alone, since tissue viability cannot be judged without knowledge of perfusion, which necessitates the use of contrast agents. Additionally, the ablation volume itself occasionally may present on B-mode ultrasonography very similarly to the surrounding normal tissue, again making the use of contrast agents necessary. Thus, before the availability of CEUS,
it was mandatory to include CT or MRI in the follow-up protocol of ablation therapy. With the advent of CEUS, ultrasonography can play a pivotal role in all aspects of percutaneous ablation from diagnosis, through guidance and procedural monitoring [62–64], to follow-up examinations after percutaneous ablation. Recent reports, including one multicenter study, have concluded that CEUS is comparable in performance to both contrast-enhanced CT and MRI [65,66] in the evaluation of ablation treatment for local recurrence, the assessment of ablation volume viability, and the identification of new metastases. Fig. 7 (with Video clip 16) and Fig. 8 (with Video clip 17) illustrate how CEUS is an indispensable tool in ultrasonography follow-up after percutaneous ablation, both with respect to imaging and as guidance for percutaneous biopsy in the case of relapse.

**Conclusion**

The introduction of contrast agents for ultrasonography has had an overwhelming impact on modern medicine. CEUS has dramatically expanded the field of medical ultrasound and opened up a completely new world in patient management. In liver imaging, CEUS has come to play a decisive role, due to its excellent temporal resolution, and the use of CEUS has changed the classic indications for biopsies and limited biopsy to equivocal cases. CEUS in many aspects is comparable to contrast-enhanced CT and MRI [1–4,10,15–19,22,23,65], and CEUS has changed the algorithms for medical imaging in high-income industrialized societies. According to the WHO, two thirds of the world’s population has no access to medical imaging [6]. Ultrasonography including CEUS has the potential to rectify this inequality, and if CEUS were incorporated as an integral part of ultrasonography, it would be realistic to bring state-of-the-art medical imaging to the entire world. Just imagine how the lack of availability of contrast agents would limit the value and reduce the impact of CT or MRI! This is the scenario ultrasonography must face in many countries where contrast agents have not been approved. The World Federation for Ultrasound in Medicine and Biology has as its overall purpose to bring the use of medical ultrasound to every corner of the world. Used together with all its technical refinements, including CEUS, ultrasonography is a second-to-none multipotent imaging tool capable of providing state-of-the-art imaging to the entire world in the new millennium.

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**Abbreviations:** AFSUMB, Asian Federation of Societies for Ultrasound in Medicine and Biology; AIUM, American Institute of Ultrasound in Medicine; ASUM, Australasian Society for Ultrasound in Medicine; CEUS, contrast-enhanced ultrasonography; CT, computed tomography; EFSUMB, European Federation of Societies for Ultrasound in Medicine and Biology; FLAUS, Latin American Federation of Societies for Ultrasound in Medicine and Biology; FLL, focal liver lesion; HCC, hepatocellular carcinoma; ICUS, International Contrast Ultrasound Society; MRI, magnetic resonance imaging;
WFUMB, World Federation for Ultrasound in Medicine and Biology; WHO, World Health Organization.

Conflict of Interest
No potential conflict of interest relevant to this article was reported.

Supplementary Material
Video clip 1. In the arterial phase, the hyperechoic lesion shows globular filling-in contrast enhancement, whereas the hypoechoic lesion demonstrated arterial hyperenhancement (http://dx.doi.org/10.14366/usg.15057.v001).

Video clip 2. During the portal phase, the hyperechoic lesion continues the same pattern of contrast filling-in enhancement, whereas the hypoechoic lesion shows a contrast wash-out pattern, thus confirming the B-mode diagnosis of hemangioma and metastasis (http://dx.doi.org/10.14366/usg.15057.v002).

Video clip 3. During the late phase, the same contrast enhancement pattern as in Video clip 2C continues (http://dx.doi.org/10.14366/usg.15057.v003).

Video clip 4. Contrast-enhanced ultrasonography sequence illustrates arterial contrast uptake with very early (before normal liver tissue) enhancement of the central lesion with a centrifugal pattern, a feeding artery consistent with the focal nodular hyperplasia arterial pattern, and the slower centripetal globular filling-in enhancement pattern of the larger lesion consistent with the arterial pattern of hemangioma (http://dx.doi.org/10.14366/usg.15057.v004).

Video clip 5. Contrast-enhanced ultrasonography sequence captured immediately after Video clip 4, illustrates arterial contrast uptake with slow centripetal globular filling-in enhancement pattern of another more peripherally located lesion consistent with the arterial pattern of hemangioma (http://dx.doi.org/10.14366/usg.15057.v005).

Video clip 6. Contrast-enhanced ultrasonography from the portal phase shows the typical slow filling-in of a large hemangioma in which the central areas do not display enhancement, as well as the sustained hyperenhancement of focal nodular hyperplasia, with both features extending long into the parenchymal phase (http://dx.doi.org/10.14366/usg.15057.v006).

Video clip 8. In a patient of minor spleen trauma, contrast-enhanced ultrasonography of the left flank clearly shows a small hematoma in the lower pole of the spleen (http://dx.doi.org/10.14366/usg.15057.v008).

Video clip 9. Ultrasonography-computed tomography (CT) fusion at follow-up two days after the accident shows the perisplenic fluid collection with a similar size as found in the CT imaging two days previously. In addition, a minor laceration is present in the spleen parenchyma near the medial-cranial pole, which is not as clearly visualized in the sonogram as in the corresponding fused CT (http://dx.doi.org/10.14366/usg.15057.v009).

Video clip 10. A fused contrast-enhanced ultrasonography and computed tomography image clearly demonstrates the small laceration, and also confirms that no extravasation of the ultrasound contrast agent was present, thus demonstrating that bleeding from the spleen had ceased (http://dx.doi.org/10.14366/usg.15057.v010).

Video clip 11. A contrast-enhanced sonogram in the arterial phase performed in our unit demonstrates arterial enhancement, which is slightly delayed and slightly hypoenhancing compared to the surrounding normal spleen tissue (http://dx.doi.org/10.14366/usg.15057.v011).

Video clip 12. By a contrast-enhanced ultrasonography-guided biopsy, the final diagnosis was splenoma, a benign hamartoma-like tumor of the spleen (http://dx.doi.org/10.14366/usg.15057.v012).

Video clip 13. A Sonazoid contrast-enhanced ultrasonography shows arterial hyperenhancement with earlier onset than the surrounding liver tissue (please note that there is in addition a simple cyst in the patients right upper corner of the Video clip which appears typically totally non-enhancing and echopoor) (http://dx.doi.org/10.14366/usg.15057.v013).

Video clip 14. A Sonazoid contrast-enhanced ultrasonography-guided 1.2-mm TruCut biopsy of an isoenhancing lesion in the postvascular phase at 13 minutes post-contrast revealed well-differentiated hepatocellular carcinoma (http://dx.doi.org/10.14366/usg.15057.v014).
**Video clip 15.** A Sonazoid contrast-enhancing ultrasonography shows sustained isoenhancement in the postvascular phase at 31 minutes post-contrast (http://dx.doi.org/10.14366/usg.15057.v015).

**Video clip 16.** SonoVue contrast-enhanced ultrasonography shows a well-demarcated ablation cavity without any enhancement, indicating successful ablation with no suspicion of skipped areas and no viable tissue (http://dx.doi.org/10.14366/usg.15057.v016).

**Video clip 17.** After the reinjection of SonoVue, a contrast-enhanced ultrasonography-guided cytology biopsy was directed towards the suspected recurrence area, and microscopy confirmed the diagnosis of local recurrence (http://dx.doi.org/10.14366/usg.15057.v017).

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