Endoscopic ultrasound (EUS) has enabled detailed observation of lesions around the upper intestine, as well as aided the acquisition of pathological samples using fine-needle aspiration (FNA) under real-time EUS image guidance from the within the intestine. EUS is a highly sensitive imaging technique, and EUS-FNA is a reliable and safe biopsy procedure used in daily clinical practice. Moreover, both are commonly used for evaluating pancreatic lesions since the pancreas is located in the deep abdomen, making percutaneous imaging and biopsy difficult. On the other hand, abdominal ultrasound is a well-established modality for the evaluation of liver lesions, which is followed by percutaneous biopsy when necessary; however, the indications for EUS have also recently expanded to liver lesions.

A study in 1999 by Nguyen et al. provided an initial report of EUS-FNA for liver lesions. They reported that EUS revealed the presence of focal liver lesions in 14 of 574 (2.4%) patients with a history or with suspicion of malignancy, whereas previous computed tomography scans depicted the liver lesions in only 3 out of 14 (21%) patients. EUS-FNA using a 22-gauge needle was subsequently performed for these liver lesions (median size, 1.1 cm; range, 0.8 to 5.2 cm) with a mean number of passes of 2.0 (range, 1-5), yielding adequate samples for cytological evaluation without any adverse events. The authors then concluded that EUS and EUS-FNA established a definitive M stage. These study findings show that EUS can detect small focal liver lesions and confirm that a cytologic diagnosis of liver metastasis using FNA may impact future clinical management. Following this report, several groups have begun evaluating the efficacy and safety of EUS-FNA for focal liver lesions. A retrospective cohort study including 41 patients showed that EUS-FNA was feasible in 40 of them (97.5%). The sensitivity and specificity for malignancy were 94% and 100%, respectively, with an average needle pass of 1.4 passes, and minor adverse events (self-limited local bleeding) only occurred in two patients (4.8%). Since there have been no studies comparing EUS-FNA and other modalities, no conclusion has been made regarding the superiority of EUS-FNA for liver lesions; however, based on the results of these reports, EUS-FNA can be considered a feasible, reliable, and safe procedure for focal liver lesions.

Despite the excellent results of EUS-FNA for liver lesions, its indications for use remain unclear. One of the reasons for this might be the visualization capability of EUS for the liver. EUS is equipped with a high-frequency transducer that provides high-resolution images but has a limited visualization range which obscures deep liver observation. Furthermore, the liver can only be visualized from the stomach and duodenum during EUS, which results in a restricted visible area. These features of EUS only enables detailed observation of the liver’s left lateral, caudate, and partial right lobes, limiting the indica-
tion of EUS for liver lesion screening. On the other hand, EUS-FNA has advantages over percutaneous biopsy. First, since the EUS-FNA access route is from within the body, it is minimally affected by ascites surrounding the liver and is completely unaffected by subcutaneous fat. Second, the deep area in the percutaneous approach can be a shallow area in EUS-FNA. An example of this situation is the liver's caudate lobe, which is located in a deep area just behind the inferior vena cava using the percutaneous approach, but this becomes an area adjacent to the stomach using the EUS approach.

In this issue of Clinical Endoscopy, a study by Akay et al. evaluated the diagnostic capability of a single EUS-FNA puncture using a 22-gauge needle for focal liver lesions, which included 25 patients with liver lesions and a technical success rate of 88% (22/25). Of the 22 patients with successful EUS-FNA, the aspirate success and biopsy success rates were 94% (21/22) and 86% (19/22), respectively, even with a single pass of FNA. Moreover, the procedure itself had a sensitivity, specificity, and accuracy of 94%, 100%, and 86%, respectively, with no adverse post-procedure events occurring. The results of this study confirmed the efficacy and safety of EUS-FNA for treating liver lesions, and it also suggested that a single pass of FNA using a regular 22-gauge needle might be sufficient to obtain a diagnostic pathological specimen. Recently, fine-needle biopsy (FNB) needles with a Fransseen or Fork-tip shape have been developed, with reports showing a high diagnostic yield with fewer needle passes. Thus, a single pass of EUS-FNB may further improve the diagnostic capability of liver lesions.

Recent development of devices and deepening knowledge of EUS-related procedures have expanded EUS indications for liver disease diagnosis and treatment of liver. A meta-analysis of EUS-FNA for liver biopsy using a larger bore needle (19-gauge) showed a histologic diagnosis rate of 93.9% and an adverse event rate of 2.3%, concluding that EUS-guided liver biopsy is an effective and safe sampling method. EUS-guided portal pressure gradient (PPG) measurement using a 25-gauge FNA needle and a novel compact manometer showed a 100% technical success, no adverse events, and PPG with excellent correlation with clinical parameters. Other than these new indications, there have been reports of EUS-guided liver tumor treatments using thermal therapy, including radiofrequency ablation, laser ablation or cryoablation, brachytherapy, or photodynamic therapy, although most of them are still in research protocols. Considering these reports regarding new knowledge, techniques, and devices in EUS-guided interventions, the indication “light” of EUS-guided management for the liver will expand over the current negative side “shadow” of EUS.

Conflicts of Interest
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ORCID
Takaji Iwashita: https://orcid.org/0000-0003-4978-1787
Masahito Shimizu: https://orcid.org/0000-0002-1151-2058

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