**INTRODUCTION**

Endoscopic ultrasound–guided liver biopsy (EUS-LB) is gaining traction among hepatologists and endosonographers. It is safe and effective at delivering excellent liver biopsy cores and can be time saving if another endoscopic procedure is required at the same time. Although the first EUS-guided biopsy of a liver lesion was described in 1997, EUS-LB for routine assessment of elevated liver biochemistries or evidence of cirrhosis was later reported in 2008. There is no clear consensus on the optimal tools and techniques used in EUS-LB. Herein, we describe the various types of needles used in this procedure and suction techniques available.

**EUS-LB**

Before initiating the procedure, it is important to review the indications and contraindications. Indications are similar to those for percutaneous or transjugular liver biopsy and include identifying the etiology of complex liver disease, staging of liver disease (nonalcoholic steatohepatitis or cirrhosis), and tissue acquisition of focal hepatic lesions. Contraindications can be extrapolated from EUS-guided FNA in GI tract lesions and may include inability to tolerate sedation, gastric outlet obstruction, no clear biopsy path without collaterals, abnormal coagulation factors, or hemodynamic instability.

Both EUS-FNA and fine-needle biopsy (FNB) can be used in EUS-LB (Fig. 1). Three different EUS-FNB needle types are commercially available: ProCore, SharkCore, and Acquire. Each has a different needle type consisting of a reverse-bevel core, Fork tip (SharkCore; Medtronic, Minneapolis, Minn), and Franseen tip (Acquire; Boston Scientific, Natick, Mass), respectively (Fig. 1). A discrepancy in the literature exists regarding the superiority of FNA versus FNB needles based on the most recent meta-analyses available. However, data may favor FNB needles in terms of diagnostic yield. In a meta-analysis by Khan et al, diagnostic yield was similar between FNA and FNB needles only when FNA was accompanied by rapid onsite evaluation of the specimen. In addition, a 2020 meta-analysis by Baran et al found that FNB needles yielded a higher number of complete portal tracts.

There are 6 reported tissue acquisition techniques that can be used in EUS-LB (Fig. 2). The wet suction technique may improve diagnostic yield compared with dry suction when using FNA. Another useful FNA technique is wet heparin, which reportedly has less tissue fragmentation, more complete portal tracts, and increased aggregate specimen length compared with dry techniques. Heparin priming of the needle may decrease blood clots within the specimen, leading to less blood contamination, and enhance tissue processing and interpretation. Video 1 outlines the EUS-LB tools and techniques, a brief ex vivo cadaveric EUS-LB, and an in vivo real-world example of EUS-LB (Video 1, available online at www.giejournal.org; Fig. 2).

Adverse events may be related to sedation, may be well-recognized endoscopic adverse events, or may be biopsy-related, including bleeding, abdominal pain, infection, or needle tract seeding of malignant cells. Although FNB needles in 1 study conferred a higher likelihood of pain post-procedure compared with FNA, no studies to date have shown a higher bleeding risk in EUS-LB based on needle type or size.

**Figure 1.** Needle types. From left to right: ProCore FNB Needle, SharkCore FNB Needle, Acquire FNB Needle, and Expect FNA Needle.

**Figure 2.** Suction techniques. Ex vivo cadaveric liver.
SUMMARY

EUS-LB is now more routinely used for diagnostic purposes in liver pathology. Several suction techniques using different needle options have been described over the past decade. Outcomes are comparable, and there is currently no consensus regarding the optimal approach to EUS-LB.

DISCLOSURE

All authors disclosed no financial relationships.

Abbreviations: EUS-LB, EUS-guided liver biopsy; FNB, fine-needle biopsy.

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