An experimental study to assess the best maneuver when using a reverse side-bevel histology needle for EUS-guided fine-needle biopsy

**Background and study aims:** Recently, ProCore™ was developed as an endoscopy ultrasound (EUS)-guided histology needle designed to address several current limitations of EUS-guided fine-needle aspiration (FNA). Nevertheless, tissue yield with the ProCore™ is not consistent. No standard technique has been established. This experimental study was conducted to ascertain the best maneuver when using the ProCore™.

**Patients and methods:** We performed fine-needle aspiration and biopsy (FNAB) with a 22-gauge (G) ProCore™ using chicken tenderloin and liver. Six methods were used, with two needle movement techniques (natural speed and whipping back) and three negative pressures (no suction (NS), slow pull (SP), and 10-mL suction). Regarding the liver, no significant difference was found with suction pressures (P=0.1126, P=0.0718), but a significant difference was found with SP (P=0.0028). Comparing “natural speed” with “whipping back” using the tenderloin, no significant difference in tissue yield was found with NS and 10 mL of pressure (P=0.1508; SP P=0.0873; 10 mL P=0.6667).

**Conclusions:** EUS-FNAB using ProCore™ can be performed with negative pressure with any needling technique. Although ProCore™ has a reverse side-bevel, results in using it with a whipping-back technique were inconclusive.

**Introduction**

Endoscopic ultrasound (EUS)-guided fine-needle aspiration (FNA) is efficacious for sampling solid mass lesions. The technique of EUS-FNA has spread rapidly, finding wide use as a less-invasive, reliable, and safe technique for obtaining tissue specimens from pancreatic tumors, lymphadenopathy, and gastrointestinal subepithelial and other lesions. Its diagnostic yield, however, could be improved as past studies have shown it to be around 90% [1–4].

To achieve adequate tissue sampling, various EUS-FNA needle devices have been developed. Recently, a needle (Echo Tip® HD ProCore™; Wilson-Cook Medical Inc., Winston-Salem, NC) that combines the attributes of both FNA and fine-needle biopsy (FNB) has been introduced and rapidly achieved wide usage [5]. The device has a reverse bevel side fenestration that allows tissue to be suctioned into the bevel, yielding a core of tissue upon withdrawal (Fig. 1). The theoretical advantages of ProCore™ include: 1) a “core trap” within the needle, which holds the core specimen; 2) a “reverse side-bevel,” which serves to cut the specimen and allows tissue to be scraped into the bevel so that core tissue can be obtained upon device withdrawal. Procore™ was developed as a biopsy needle that can absorb material, which clearly distinguishes it from a core cut needle.

A study of EUS-FNA and biopsy (FNB) using the ProCore™ needle has shown an increase in diagnostic accuracy of approximately 15% compared to EUS-FNA conducted with a standard needle [6]. However, a recent study demonstrated that the diagnostic sufficiency and technical performance of a standard FNA needle and the FNAB needle (ProCore™) are comparable. No significant difference was found between the two needle types in the yield or quality of the histologic core [7]. However, the evaluation of the FNAB needle is
not clear. The tissue yield with the ProCore™ is not regarded as consistent because no standard FNAB technique has been established.

Because of the unique characteristics of the ProCore™ needle, we theorized that moving it with a whipping back technique might yield more tissue than moving it at natural speed because the needle has a “reverse side-bevel” that cuts the specimen. It also was unclear whether application of suction pressure would be beneficial when using ProCore™. A slow-pull technique, whereby simultaneous minimal negative pressure is provided by pulling the stylet slowly and continuously, has been reported to be useful for EUS-FNAB [8, 9], but its value remains controversial. Therefore, we conducted an experiment to ascertain which maneuver was better for EUS-FNAB using the ProCore™ needle. This study was undertaken to ascertain the best movements for using the ProCore™ in EUS-FNAB.

Materials and methods

Two endoscopists (AY and AI) alternated performance of FNAB on fresh chicken tenderloin and liver using 22-gauge (G) ProCore™ needles without using an endoscope. To effectively use the “reverse side-bevel” that characterizes ProCore™, movement of the needle with a whipping back technique might be important. Therefore, the following two methods were evaluated: the “whipping back” technique and the conventional method using “natural speed.” To perform whipping back of the needle, we constructed a handmade device with a stopper attached to the top (Fig. 2). When the needle is pulled vigorously, it stops at a constant distance. Furthermore, the hole in the top of the device enables penetration of the extension tube. When the needle is pushed forward using the whipping back technique, the puncture speed was natural. The length of punctures in the chicken tenderloin and the liver were uniform (20 mm); 10 passes were made for each puncture.

As in previous reports, three types of suction pressures were used: “without suction” [10], “with slow pull technique” [9], and “with 10-mL suction force” [11]. We standardized our techniques for core biopsy as follows. Six techniques with 10 back-and-forth movements of the needle per pass were evaluated: 1) needling with “natural speed” without suction; 2) needling with “natural speed” with slow-pull technique; 3) needling with “natural speed” with 10-mL suction; 4) needling with “whipping back” without suction; 5) needling with “whipping back” with the slow-pull technique; and 6) needling with “whipping back” with 10-mL suction. After FNAB, the tissues obtained were expressed entirely onto a chart by reinsertion of the stylet. In most cases, tissue was obtained from the needlepoint and reverse side-bevel. The materials were evaluated based on whether a visible core for histology was obtained. An electronic balance was used to weight the specimens and differences in weight for materials obtained with the “whipping back” technique and the conventional method with “natural speed” were assessed. In addition, the effect of suction was assessed by examining the difference in the weight of tissue obtained “without suction,” “with slow pull technique,” and “with 10-mL suction” for each technique.

The differences in the weights of obtained materials between two groups were analyzed using the Mann-Whitney U test. Confidence interval (95% CI) was calculated for differences in proportions for categorical data, and the approximate 95% confidence interval (CI) for differences in medians for nonparametric data was calculated. All data analyses were performed using GraphPad Prism 6 (GraphPad Software Inc., La Jolla, CA, USA). All P values were two-sided, and statistical significance was set at P<0.05.

Results

For each tissue type and for all techniques, more specimens were obtained with higher negative pressure. Moreover, visible core samples (Fig. 3) were obtained consistently in the groups with negative pressure.
FNAB for chicken tenderloin

**Cases using “natural speed” technique**

The weights of obtained tissues averaged 0.35 mg (median) [range, 0.3 – 0.5 mg] in the no-suction group, 0.7 mg [range, 0.6 – 1.1 mg] in the slow-pull group, and 2.45 mg [range, 1.8 – 3.8 mg] in the 10-mL suction group (Fig. 4). Significant differences were found between different suction pressures (no-suction vs. slow-pull; 95 % CI, 0.3000 – 0.6000; P < 0.0001, slow-pull vs. 10-mL suction; 95 % CI, 1.200 – 2.400; P < 0.0001, no-suction vs. 10-mL suction; 95 % CI, 1.600 – 2.800; P < 0.0001).

**Cases using “whipping back” technique**

The weights of obtained tissues averaged 0.2 mg [range, 0 – 0.6 mg] in the no-suction group, 0.4 mg [range, 0.1 – 0.8 mg] in the slow-pull group, and 2.0 mg [range, 1.2 – 3.4 mg] in the 10-mL suction group (Fig. 4).

No significant difference was found between the no-suction and the slow-pull groups (95 % CI, 0.0 – 0.4000; P = 0.0596), but a significant difference was found between slow-pull and 10-mL suction (95 % CI, 0.9000 – 2.0000; P < 0.0001), and between no-suction and 10-mL suction (95 % CI, 1.100 – 2.2000; P < 0.0001).

**Comparison between needling methods with the same suction pressure**

Comparison of “Natural speed” with “Whipping back” techniques using the same suction pressure showed no significant difference at no-suction and 10-mL suction (no-suction; 95 % CI, – 0.1000 – 0.3000; P = 0.1126, 10-mL suction; 95 % CI, – 0.1000 – 1.4000; P = 0.0718), but a significant difference at slow pull (95 % CI, 0.1000 – 0.5000; P = 0.0028), as shown in Fig. 5.

**FNAB for chicken liver**

**Cases using “natural speed” technique**

The weights of obtained tissues averaged 0.8 mg [range, 0.3 – 2.7 mg] in the no-suction group, 4.6 mg [range, 3.1 – 8.8 mg] in the slow-pull group, and 18.7 mg [range, 13.6 – 25 mg] in the 10-mL suction group (Fig. 6).
Significant differences were found between different suction pressures (no-suction vs. slow-pull; 95% CI, 1.900 – 8.000; \( P = 0.0079 \), slow-pull vs. 10-mL suction; 95% CI, 8.700 – 20.40; \( P = 0.0079 \), no-suction vs. 10-mL suction; 95% CI, 12.80 – 24.20; \( P = 0.0079 \)).

**Cases using “whipping back” technique**

The weights of obtained tissues averaged 2.1 mg [range, 1.1 – 3.3 mg] in the no-suction group, 7.5 mg [range, 5.2 – 8.9 mg] in the slow-pull group, and 15.8 mg [range, 13.2 – 22 mg] in the 10-mL suction group (Fig. 6).

Significant differences were found between different suction pressures (no-suction vs. slow-pull; 95% CI, 3.100 – 6.600; \( P = 0.0079 \), slow-pull vs. 10-mL suction; 95% CI, 5.500 – 14.50; \( P = 0.0079 \), no-suction vs. 10-mL suction; 95% CI, 0.90 – 19.90; \( P = 0.0079 \)).

**Comparison of needling methods using the same suction pressure**

Comparison between “natural speed” and “whipping back” using the same suction pressure showed no significant differences at any suction pressure (no-suction; 95% CI, –2.5000 – 0.6000; \( P = 0.1508 \), slow pull; 95% CI, –4.400 – 1.300; \( P = 0.0873 \), 10mL suction; 95% CI, –6.0000 – 9.2000; \( P = 0.6667 \)), as shown in Fig. 7.

**Discussion**

EUS-FNA is a highly accurate method of obtaining tissue to diagnose various diseases. Cytology of samples obtained with FNA is generally adequate for diagnosis of malignancy, but histologic examination may be necessary for tumors such as lymphoma, gastrointestinal stromal tumor (GIST), and metastatic carcinoma. In addition, pathologists may specifically request core tissue to establish a definitive diagnosis in challenging cases [12,13]. An FNB specimen includes core tissue with better preservation of cellular architecture than an aspirate [14]. In general, use of an FNB specimen achieves greater diagnostic accuracy and the technique provides more tissue for ancillary testing than does an FNA sample [15]. EUS-guided trucut needle biopsy (EUS-TNB; Cook Endoscopy Inc., Winston-Salem, NC, USA) was used to overcome the limitations of EUS-FNA by acquisition of histology specimens to support evaluation of tissue architecture and the use of immunohistochemistry. The overall diagnostic accuracy of EUS-TNB for evaluating suspicious lesions at various sites in the body is reportedly 75% to 84% and 61% to 67.5% for pancreatic masses [16,17]. Although the EUS-TNB technique exhibits some advantages over FNA, criticisms of EUS-TNB include difficult maneuverability because of needle rigidity and its spring-loaded mechanism, increased cost, uncertain safety, and a paucity of clinical data to develop practice guidelines [18].

“EUS-guided ProCore™ biopsy” recently was introduced, combining the attributes of both FNA and FNB and a new design of core needles incorporating a hollowed-out reverse bevel to trap core specimens [5]. The device has a 2-mm reverse bevel side fenestration 3.9 mm from the needle tip, which enables suctioning of tissue into the bevel and acquisition of core tissue on withdrawal. ProCore™ has three needle sizes (19-G, 22-G, 25-G), and when tight angulation of the echoendoscope or the use of a forceps elevator is necessary, the 22-G or 25-G needles appear to be suitable for insertion. Iglesias-Garcia et al reported that the technique of EUS-FNAB with ProCore™ 19G is feasible and safe for histopathologic diagnosis. The sample quality, according to the pathologist, was adequate for full histologic assessment in 85.9% of lesions (98 of 114) [5]. To obtain better samples (especially for histology), various suction techniques using ProCore™ have been explored. Nakai et al reported the usefulness of the slow-pull technique, which provides less bloody specimens without reducing cellularity in EUS-FNA for pancreatic malignant lesions. Moreover, the sensitivity of the slow-pull technique is similar in terms of cytology but higher than that for histology [9]. In another report, Bang et al described the possibility of obtaining tissue for cytolc analysis in 89.3% and histologic analysis in 83.3% of cases using the 22-G ProCore™ with 10 mL suction [10]. In addition, Fabbri et al reported that the overall sensitivity, specificity, and accuracy of EUS-FNB using a 22-G ProCore™ in solid pancreatic lesions of ≤2 cm with 10 mL suction pressure were 80%, 100%, and 82%, respectively [19]. Witt et al. described that 73% of cases had corresponding cell blocks and contained representative diagnostic material using ProCore™ with or without suction [20]. Strand et al. demonstrated that EUS-FNB using suction with strength entrusted to the discretion of the acting endoscopist was successful in 84.4% of cases, and that core tissue for histologic examination could be obtained in 70.3% of cases [21]. As described above, the standard method of suction for ProCore™ remains controversial and data on it are inconclusive.

Past reports from relevant literature suggest that use of a ProCore™ needle with conventional EUS-FNA technique provides high tissue yields. Although the needle movement is done at natural speed or rapidly at the moment of needling with conventional EUS-FNA technique [22], controversy exists about which needling technique is best for obtaining samples for histology. Given the characteristics of the ProCore™ needle, moving it with a whipping back technique might produce more tissue than doing so at a natural speed because it has a “reverse side-bevel,” that serves to cut the specimen. In addition, the suction best suited for ProCore™ use remains undetermined. Therefore, we assessed how much material could be obtained using two needling techniques and three suction techniques. In our experimental study, the unique structure of ProCore™ did not necessitate a particular needling method. With regard to the suction technique, higher negative pressure yielded more specimen from both tissue targets (chicken tenderloin and liver) regardless of movement technique.

Several reports have described the usefulness of the slow-pull technique for obtaining good-quality samples when using a ProCore™ needle [9]. Katanuma et al. showed that the slow-pull technique offers fundamentally no suction in their bench-top testing, and questioned its usefulness [23]. In our experimental data, the quantities of tissues obtained were much greater in the slow-pull group than in the no-suction group. This result shows that the slow-pull technique has some absorption effect when obtaining tissues from a solid mass. In addition, the capillary pressure might affect the results described. The quality of samples is regarded as an advantage of the slow-pull technique, but for this study, we are unable to provide insight on that because we did not evaluate our specimens under a microscope. However, because obtaining more tissue for histopathologic diagnosis is extremely important, reliable absorption appears to be necessary, even with use of a ProCore™ needle.

For this study, we used fresh chicken liver and were conscious of the potential for blood contamination because fresh liver includes blood. Macroscopic observation of all specimens obtained
during our procedures showed little or no adhesion of blood. Although we did not perform microscopic evaluation of sample quality, the presence of blood did not obstruct sampling of an adequate or large amount of tissue with any technique. Nevertheless, an important limitation of our experimental study is that the fresh chicken tenderloin and liver that we used were not live organs. From a clinical perspective, results with the techniques might differ depending on the quantity of blood in the target and differences in tissues subject to sampling. A prospective, randomized, controlled trial must be conducted using live organs or tumors to ascertain the optimal technique for use of ProCore™ in EUS-guided biopsy.

In conclusion, results of our experimental study indicate that EUS-FNAB using ProCore™ might be performed best with negative pressure (at least 10 mL) in any needling technique or those using other FNA needles. Although ProCore™ has a reverse side-bevel, it does not appear that the “whipping back” technique would be useful. In addition, no blood vessels were seen in the target materials, a fact that makes it difficult to draw clear conclusions about the effects and usefulness of the slow-pull technique.

Competing interests: None

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