25-gauge histology needle versus 22-gauge cytology needle in endoscopic ultrasonography-guided sampling of pancreatic lesions and lymphadenopathy

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Background and study aims: A new 25-gauge (G) endoscopic ultrasound-guided fine-needle aspiration (EUS-FNA) device (EchoTip ProCore; Cook Medical, Bloomington, Indiana, USA) has recently been developed, which features a hollowed-out reverse bevel to trap core samples. However, data on the differences between the diagnostic yield of the 25G EchoTip ProCore and that of a 22G standard needle are limited.

Patients and methods: This pilot study included 27 patients referred during an 11-month period for EUS-FNA of pancreatic masses and enlarged lymph nodes adjacent to the upper gastrointestinal tract. Each lesion was punctured once by both a 25G EchoTip ProCore needle and a 22G standard needle (EchoTip; Cook Medical) with capillary sampling. Blinded histocytologic analyses were conducted. The final diagnosis was based on FNA histocytologic results.

Results: A total of 28 EUS-FNA procedures targeting masses of the pancreas (n = 19) and lymph nodes (n = 9) were performed. No complications were encountered. Single-pass sensitivity rates for pancreatic and lymph node malignancy were equal for the needle types: 89.5% (95% CI 66.82 – 98.39) and 66% (95% CI 24.1 – 94), respectively. There were no significant differences between the needles in terms of EUS visualization (P = 0.125), amount of blood contamination (P = 0.705), macroscopic quantity of the material (P = 0.858), quality of the cytology (P = 0.438), and adequacy and accuracy of the cell block material (P = 0.220).

Conclusions: Both needles were safe and successful in terms of a high diagnostic yield, with similar histocytologic results.

The results of this study were presented at Digestive Disease Week (DDW) 2014, Chicago, Illinois. This trial was registered at ClinicalTrials.gov (B027201316271).

Introduction

Endoscopic ultrasound-guided fine-needle aspiration (EUS-FNA) has become a mainstay diagnostic technique for the evaluation of lesions arising from the pancreas and upper gastrointestinal tract as well as adjacent structures, including lymph nodes and the liver. EUS-FNA is safe and has a diagnostic accuracy of 60% to 90% [1–3]. The diagnostic yield of EUS-FNA is influenced by several factors: the nature of the target lesion, experience of the endoscopist, presence of an onsite cytopathologist, and various technical aspects, such as needle size and number of passes. In order to limit the influencing parameters, a novel needle design was introduced in 2010–2011 that features a hollowed-out reverse bevel to trap core samples (EchoTip ProCore; Cook Medical, Bloomington, Indiana, USA). A potential advantage of this type of needle is the acquisition of larger amounts of tissue with preserved architecture. Preliminary results on the performance of the 19-gauge (G), 22G, and 25G ProCore needles were promising, with reported high diagnostic accuracy rates of 89.6% [4], 88.5% [5], and 86% [6], respectively. However, comparative studies of the 22G ProCore needle versus the 22G standard cytology needle showed similar results for the cytology parameters, amount of blood contamination, and diagnostic adequacy and accuracy of the cell block material [7–9]. Fewer data are available on the diagnostic yield of the 25G ProCore needle. Preliminary data demonstrated an 83% first-pass sensitivity for pancreatic malignancy, with a cumulative sensitivity of 96% after three passes [6]. Histologic diagnosis was possible in 63% after a single pass and in 80% after multiple passes [6]. Taking into account the results of a recent meta-analysis [10] that shows a potential advantage of the 25G needle over the 22G needle...
for the diagnosis of pancreatic malignancy, the idea of a 25G needle that can obtain histologic specimens sounds attractive. Therefore, we designed a prospective study comparing the most commonly used 22G standard cytology needle versus the 25G EchoTip ProCore needle in the same group of lesions in terms of differences in accuracy, technical performance, and quality and quantity of the cytology and cell block specimen obtained.

Materials and methods

This pilot study included patients with pancreatic mass lesions or lymphadenopathy who were referred for EUS-FNA sampling at our hospital (Grand Hôpital de Charleroi, Charleroi, Belgium) between December 2012 and October 2013. Exclusion criteria included the following: cystic lesion, coagulation disorder (international normalized ratio > 1.5, platelet count < 50 000/mm³), pregnancy, age < 18 years, and refusal or inability to provide informed consent. Patients were monitored closely for possible complications after the procedure. The study was approved by the Grand Hôpital de Charleroi review board, and written informed consent was obtained from all patients for participation in the study. This trial was registered at ClinicalTrials.gov (027201316271).

A linear array echoendoscope (UCT160; Olympus Medical Systems, Tokyo, Japan) connected to a processor featuring color Doppler function (ProSound Alpha 7; Hitachi Medical Systems Europe [Aloka], Zug, Switzerland) was used for all procedures, which were carried out by four experienced endosonographers (A.S., P.W., H.H., and C.G.) and one gastroenterology fellow (G.M.) which were carried out by four experienced endosonographers and one gastroenterology fellow. A monolayer technique. It is also easily reproducible because one needs only to record the magnification at which diagnostic cells are seen filling the microscopic field.

The quality of the obtained cytologic and histologic specimens was reported by the cytopathologist according to the following grading system: insufficient material or contaminated material (score of 0), rare diagnostic cells (score of 1), diagnostic cells at every field with 20 times magnification (score of 3). The primary outcome parameter was the percentage of cases in which the pathologist classified the quality of the sample as sufficient for histocytologic evaluation (score ≥ 1). This grading system, although not published as such, is very easy to apply to a cytologic slide, especially when the specimen has been obtained with a monolayer technique. It is also easily reproducible because one needs only to record the magnification at which diagnostic cells are seen filling the microscopic field.

Secondary outcomes included comparisons of several performance parameters. The visibility of each needle was qualitatively scored as no blood (score of 0), small quantity of blood (score of 2), or diagnostic cells at every field with 10 times magnification (score of 2), or diagnostic cells at every field with 20 times magnification (score of 3). The primary outcome parameter was the percentage of cases in which the pathologist classified the quality of the sample as sufficient for histocytologic evaluation (score ≥ 1). This grading system, although not published as such, is very easy to apply to a cytologic slide, especially when the specimen has been obtained with a monolayer technique. It is also easily reproducible because one needs only to record the magnification at which diagnostic cells are seen filling the microscopic field.

Fig. 1  The 25G EchoTip ProCore histology needle (left), which has a hollowed-out reverse bevel, and the 22G EchoTip standard cytology needle (right). Photos supplied by Cook Medical.
in the presence of fragments with no visible core, medium when
one or two cores were visible (score of 2), or excellent when at
least three cores were visible (score of 3).
A final diagnosis of malignancy or benignancy was made accord-
ing to one of the following reference methods: (1) definite benign
or malignant histologic diagnosis based on surgical resection
specimens from patients who had undergone surgery, (2) cytology
or histology findings with definite proof of malignancy in pa-
tients with unresectable tumors according to imaging findings
and compatible clinical follow-up, and (3) cytology or histology
findings without proof of malignancy and a minimum clinical
and radiologic follow-up of 7 months.
For comparison of continuous data, a paired t test was performed
if a normal distribution was shown, and the Wilcoxon rank sum
test was carried out if normality could not be demonstrated.
McNemar’s test was used for dichotomous categorical data. For
all tests, a P value of less than 0.05 was regarded as statistically
significant with SPSS 17.0 for Windows (SPSS Inc., Chicago, Illi-
nois, USA).

Results

A total of 28 EUS-FNA procedures targeting lesions of the pan-
creas (n = 19) and lymph nodes (n = 9) were performed in 27 pa-
tients (18 women, 9 men) with a median age of 69 years (range,
38–88). The final diagnoses were pancreatic adenocarcinoma
(n = 18), pancreatic neuroendocrine tumor (n = 1), malignant
lymphadenopathy (n = 6), and benign lymphadenopathy (n = 3).
No benign pancreatic lesions were encountered.

In the subgroup of patients with pancreatic lesions, the median
tumor size was 39 mm (range, 10–70). Of these pancreatic les-
ions, 10 were punctured through the duodenum and 9 were
punctured through the stomach.
In the subgroup of patients with lymphadenopathy, the median
lymph node size was 24 mm (range, 15–45). Of these lesions, 6
were punctured through the esophagus and 3 through the du-
odenum.

Final diagnoses were made on the basis of surgery in 3 cases, po-
sitive FNA for malignancy with a compatible clinical course in 23
cases, and negative FNA for malignancy with at least 7 months of
follow-up in 2 cases. No procedure-related complications were
seen.

In terms of EUS visualization, visualization was suboptimal in 16
% of punctures with the 25G needle versus 0% of punctures with
the 22G needle; however, this difference was not statistically sig-
nificant (P = 0.125) (Videos 1, 2). No relevant differences were
found regarding the ease of puncture (P = 0.688), amount of blood
contamination (P = 0.705), macroscopic quantity of the material
(P = 0.858), and quality of the cytologic (P = 0.438) and histologic
(P = 0.220) specimens (Table 1). Subgroup characteristics re-
garding the adequacy of histocytologic material for the two need-
le types and different types of lesions are shown in Table 2.

In the subgroup of patients with pancreatic cancer, each needle
missed two cases. The first case was a patient with adenocarcino-
ma of the head of the pancreas. Transduodenal puncture with the
25G needle showed rare benign cells (cytology score = 1, cell
block score = 1), while puncture with the 22G needle was non-
contributive (cytology score = 0, cell block score = 0). Because of
the strong suspicion of malignancy, a follow-up EUS-FNA a few
weeks later was performed with a standard 22G needle and con-
ﬁrmed the diagnosis of pancreatic adenocarcinoma. Each needle
missed one other case of pancreatic adenocarcinoma of the head
of the pancreas punctured through the duodenum because of in-
sufﬁcient histocytologic material (total histocytologic score = 0).
Therefore, the single-pass sensitivity for pancreatic neoplasia for
both needles was 89.5% (95% CI 66.82–98.39).
In the subgroup of patients with lymphadenopathy, a total of two
false-negative results for malignancy were obtained with both
needles that concerned the same patients. The ﬁrst patient had
gallbladder cholangiocarcinoma and perihepatic lymph nodes
suspicous for malignancy that were 2 cm in size. Transduodenal
puncture with both needles resulted into noncontributive histo-
cytologic material (total histocytologic score = 0). Follow-up im-
aging was compatible with metastatic lymph nodes. The second
patient had lung cancer and mediastinal lymph nodes suspicious
for malignancy (4 cm in size). EUS-FNA with both needles
showed rare benign cells (cytology score = 1, histology score = 1).
However, clinical and radiologic follow-up was compatible with
metastatic lymph nodes. Therefore, the single-pass sensitivity,
speciﬁcity, positive predictive value, negative predictive value,
and accuracy for malignancy were equal for the needle types:
66% (95% CI 24.1–94), 100% (95% CI 30.9–100), 44.4% (95% CI
39.5–100), 60% (95% CI 17–92.7), and 84.8% (95% CI 67.3–94.2),
respectively.

Discussion

Overall, EUS-FNA is highly effective for most pancreatic tumors
and solid malignancies adjacent to the upper gastrointestinal
tract, with reliable sensitivity, speciﬁcity, and overall diagnostic
accuracy of 60% to 90% [1–3]. However, FNA cytology specimens
may not be adequate in cases in which the diagnosis relies on tis-
sue architecture, such as autoimmune pancreatitis, lymphomas,
gastrointestinal stromal tumors, and well-differentiated adenocarci-
nomas [1,13].
In order to procure larger amounts of tissue with preserved ar-
chitecture that would enable histologic analysis, a novel needle
assembly with a reverse-bevel technology (EchoTip ProCore)
was introduced in 2010–2011.According to the designer, the
side notch (not Tru-Cut) should provide additional “cheese-grat-
ing” action, making it possible to obtain more specimens.
Initially, a 19G version was introduced to the market. It had the
ability to obtain full histology in 89.5% of cases, with an overall
diagnostic accuracy of 93% [4]. However, because technical diffi-
culties were encountered during transduodenal passes, the same
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The 25G ProCore needle has been introduced. A PubMed search revealed only three studies of ProCore needles comparing the diagnost ic yield of the most commonly used 22G FNA needle with that of the 22G ProCore needle. No advantage of the ProCore design was reported in terms of cytology and histology. However, in the study by Witt et al., fewer passes were needed to achieve diagnosis with the 22G ProCore needle (2.11 with the 22G ProCore needle vs. 2.94 with the 22G standard needle). Nevertheless, this result should be interpreted with caution because of the small number of different groups of lesions (n = 18). Interestingly, in the study by Strand et al., technical failure was encountered in 5 of 32 patients because of significant resistance to advancement of the needle, with deflection of the echoendoscope (GF-UCT140, Olympus) while in the transduodenal position. Most recently, the 25G ProCore needle has been introduced. Potential advantages of the smaller caliper are greater flexibility, less friction in the needle sheath, easier penetration of hard pancreatic tumors, and fewer bloody aspirates. This argument was suggested in a meta-analysis by Madhoun et al. but not confirmed in a recent large randomized controlled trial. Preliminary data on the performance of the 25G ProCore needle suggest an 83% first-pass sensitivity for pancreatic malignancy, with a cumulative sensitivity of 96% after three passes. Histologic diagnosis was possible in 63% after a single pass and in 80% after multiple passes. In our study, both the 22G standard needle and the 25G ProCore needle had a slightly better single-pass performance. Cytologic and histologic diagnoses were possible in 84.2% of the cases with the ProCore needle. The 25G needle was not visualized optimally in 16% of punctures when used in the intestinal tract. Our pilot study compared different needle characteristics and the adequacy of cytology and histology. Although the 25G needle was not visualized optimally in 16% of punctures, no effect on the quality of cytology or histology, amount of blood contamination, and technical difficulty was observed. We found no significant differences between the two needles in terms of diagnostic yield, with similar diagnostic rates and similar numbers of successful procedures.

Our prospective, single-group, paired design has the intrinsic strength that performance comparisons were made on the same 

| Author          | Type of study | Number of lesions | Target                        | Comparison of histology specimen | Overall diagnostic yield | Technical performance | Comment                              |
|-----------------|---------------|-------------------|-------------------------------|---------------------------------|--------------------------|-----------------------|--------------------------------------|
| Bang et al. 2012 [7] | Randomized    | 28 lesions per needle type | Pancreas                     | No significant difference        | Equivalent               | No significant difference | Needles of different manufacturers (Boston Scientific and Cook) |
| Witt et al. 2013 [8] | Retrospective | 18 lesions per needle type | Pancreas, lymph nodes, other masses | No significant difference        | Equivalent; fewer passes needed with ProCore needle | No reported difficulties | Retrospective study, small number per group |
| Strand et al. 2014 [9] | Randomized    | 32 lesions punctured by both needles | Pancreas                     | No significant difference        | Lower overall diagnostic yield for the ProCore needle | Technical failure in 16% cases with the ProCore needle | Only two passes permitted for the 22G ProCore group |

| Table 1 | Histocytologic and endoscopic assessment of the performance of the two needles. |
|-----------------------|-----------------------|-----------------------|-----------------------|-----------------------|-----------------------|-----------------------|-----------------------|
| Target lesion         | Adequate material for cytologic assessment | Adequate material for histologic assessment | Target comparison    | Overall diagnostic yield | Technical performance | Comment                              |
| Pancreatic mass       | 25G EchoTip ProCore | 22G EchoTip | P value | 25G EchoTip ProCore | 22G EchoTip | P value | Technical performance | Comment | Comment |
| Lymph node            | 8/9                   | 7/9                   | 1                   | 7/9                   | 7/9                   | 1                   | Equivalent; fewer passes needed with ProCore needle | No reported difficulties | Retrospective study, small number per group |
| All lesions           | 24/28                 | 22/28                 | 0.5                 | 22/28                 | 24/28                 | 0.5                 | Equivalent               | No significant difference | Needles of different manufacturers (Boston Scientific and Cook) |

1. Wilcoxon rank sum test.
2. Paired t test.
3. McNemar’s test.

| Table 2 | Adequacy of material for cytologic and histologic assessment. |
|-----------------------|-----------------------|-----------------------|-----------------------|-----------------------|-----------------------|-----------------------|-----------------------|
| Target lesion         | Adequate material for cytologic assessment | Adequate material for histologic assessment | Target comparison    | Overall diagnostic yield | Technical performance | Comment | Comment |
| Pancreatic mass       | 25G EchoTip ProCore | 22G EchoTip | P value | 25G EchoTip ProCore | 22G EchoTip | P value | Technical performance | Comment | Comment |
| Lymph node            | 8/9                   | 7/9                   | 1                   | 7/9                   | 7/9                   | 1                   | Equivalent; fewer passes needed with ProCore needle | No reported difficulties | Retrospective study, small number per group |
| All lesions           | 24/28                 | 22/28                 | 0.5                 | 22/28                 | 24/28                 | 0.5                 | Equivalent | No significant difference | Needles of different manufacturers (Boston Scientific and Cook) |
group of lesions, eliminating in this way the bias of differences in lesion type, size, and location. This is in contrast with most EUS-FNA studies, which compare different needles in different populations. The main limitation of our study is the small number of cases. A larger number of cases may be needed to detect subtle differences between the two needles. Furthermore, no benign pancreatic lesion was encountered during the study period, such as focal autoimmune pancreatitis, in which an EUS-FNA diagnosis is challenging.

In conclusion, this pilot study demonstrates that the diagnostic yield of the new 25G EchoTip ProCore needle is comparable with that of a 22G standard FNA assembly. Both needles performed equally in terms of the quality of cytologic and histologic specimens, blood contamination, and ease of puncture.

Competing interests: This study was sponsored by Cook Medical.

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