**ATH-04**

**COMPARING THE DIAGNOSTIC YIELD AND SAFETY OF ENDOSONOGRAPHY GUIDED USE OF SHARK-CORE AND PRO-CORE NEEDLES**

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**Background**

Endoscopic ultrasound (EUS) guided fine needle aspiration and biopsy (EUS-FNA/FNB) is the standard technique for diagnosis and evaluation of pancreatic lesions. Core biopsy specimens provide accurate diagnoses. ProCore needle (Wilson-Cook Medical Inc. NC, USA) and Shark-Core needle (Covidien, Ireland) were designed to obtain histological and cytological samples. Each claim superiority over other for better diagnostic acquisition and safety. There is limited head-to-head comparison data available in literature about different needles.

**Aim**

The aim of our study was to compare the diagnostic yield and safety of 22G FNB needles for sampling of pancreatic lesions.

**Methods**

We prospectively randomised patients with pancreatic lesions either to Shark-Core or to Pro-Core needle sampling. Data collection included demographics, needle type, number of passes.

**Results**

A total of 143 patients having 151 pancreatic lesions were analysed. 74 lesions were biopsied in Shark-Core group from 69 patients. Four patients had their procedure repeated. Mean age was 62.4 years and 51% were males. Mean number of needle passes were four. 33 samples had malignancy and 26 had other diagnoses and 10 had insufficient tissue. The diagnostic yield was 86.3%. 2 patients developed mild pancreatitis and one required admission with epigastric pain and vomiting.

Pro-Core group had 77 lesions biopsied from 74 patients. 3 had two procedures each and one patient had both Shark core and Procore sampling. Mean age was 63 with males 50.8%. Median passes were four. 37 had malignancy and 23 had other diagnoses. Inadequate tissue sample was obtained in 13. The overall diagnostic yield was 63/77 (81.81%). 3 patients had epigastric pain and vomiting and needed admission.

**Discussion**

Though Shark-Core needle demonstrated better diagnostic yield and had marginally more complications, both of which were not statistically different from Pro-Core needle. In our experience, the time required for tissue acquisition is in favor of Shark-Core needle as it allows inner needle to be withdrawn than the whole system, maintaining the scope position unlike Pro-Core system that requires the whole biopsy needle system to be withdrawn from the scope.

With many more core sampling needles now commercially available, further studies may be required to evaluate other types of needles for tissues yield, safety as well as time required to acquire samples.

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**ATH-05**

**CAN PROTOCOLISED MEASUREMENTS WITH BARIUM RADIOLOGY PREDICT SEVERITY AND TREATMENT OUTCOMES IN ZENKER’S DIVERTICULUM?**

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**Introduction**

Barium swallow is an established investigation for Zenker’s diverticulum (ZD), yet, no agreed measurement protocol exists for the evaluation of ZD on Barium radiology. We developed a standardised protocol to measure ZD dimensions on Barium radiology, and aimed to correlate ZD dimensions with symptoms, procedural difficulty and treatment outcomes.
Methods This prospective single-centre study included patients with symptomatic ZD undergoing flexible endoscopic septal division (FESD) as day case procedures between 2014–2018. Patients underwent Barium swallow imaging; ZD dimensions were measured and agreed by two expert radiologists using a predefined protocol. Symptom severity pre- and post-FESD was recorded using the Dysphagia, Regurgitation, Complications (DRC) scale. Procedure difficulty was rated on a three-point scale: easy, moderate, difficult. The primary outcome was therapeutic success, defined as remission following single episode FESD with a DRC score of 1 or less 6 months of follow-up. ZD dimensions were subject to Mann-Whitney tests and logistic regression analyses.

Results In total, 68 patients (mean age 74.2, SD 11.8) underwent Barium radiology. Male gender comprised 60.9% of the cohort and was associated with larger pouch height (P=0.008), width (P=0.004) and depth (P=0.045). A positive correlation was identified between baseline DRC score and pouch depth (rho 0.319, P=0.012), particularly the symptom of regurgitation (P=0.009). No significant associations were found between ZD dimensions and procedural difficulty. Overall, each patient underwent an average of 1.4 FESD procedures. The outcome of therapeutic success at 6-months was achieved in 69% and was associated with poorer pouch height (median 14.5 mm vs. 19 mm, P=0.030) and pouch width (median 20 mm vs. 28 mm, P=0.046), with smaller criopharyngeal length tending towards significance (median 20.8 mm vs. 26.3 mm, P=0.051). On multivariable analysis using a forward stepwise approach, pouch height was the sole dimension affecting the study outcome, with each additional mm in pouch height associated with a decrement in the probability of therapeutic success (OR 0.946, 95% CI: 0.897–0.997, P=0.031).

Conclusions ZD dimensions may be feasibly evaluated using Barium radiology. Specific parameters, especially those relating to the pouch, appear to correlate with baseline severity and post-FESD patient outcomes. These results may inform the planning of FESD in day case patients with ZD.

REFERENCE

ATH-06 FIRST CLINICAL EXPERIENCE WITH SPEEDBOAT RS2, A NEW MULTIMODALITY DEVICE FOR COLONIC SUBMUCOSAL DISSECTION

1. Zacharias P Tsiamoulas*, 2Ioannis Stasinos, 1Aristeidis Oikonomakis, 1Joseph Sebastian, 1Nipin Bagla, 3Christopher Hancock, 2Brian P Saunders, 1East Kent Hospitals University, Margate, UK; 2St Mark’s Hospital, London, UK; 3Bangor University, UK

1/20 case was abandoned during dissection. Endoscopically en-bloc resection was achieved in 11/19 (57.8%) cases and 10/19 cases (52.6%) resections were R0 histologically. In 8/19 patients snare excision was used to facilitate resection with a median of 2 resection pieces. Microwave energy was utilized to pre-coagulate and control active bleeding successfully in 17/19 (89.5%) cases and haemostatic monopolar forceps was required in 2 cases. Histology showed dysplasia in 15/19, NET in 1/19 and T1 cancer in 3/19. 2/3 patients with T1 cancers had endoscopic surveillance with no recurrence and 1 had surgery and no residual tissue. 3/19 patients experienced...