(EUS-FNA) samples produced by the 19g procore needle, standard 19g needle and



To compare the tissue diagnostic yield of solid lesion biopsies based on the histopathological Analysis of endoscopic ultrasound guided fine Needle aspiration

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22g procore needle: a single centre, observational study.



## Abstract

Endoscopic ultrasound (EUS) is a sensitive method for detecting and extra-intestinal mass lesions including lymphadenopathy.FNA allows evaluating cellular findings suggestive of malignancy but inflammation causes cellular changes undistinguishable from neoplasia solely based on cytological evaluation, because tissue architecture and cell morphology essential for accurate pathological assessment. Various EUS-guided techniques have been explored to retrieve tissue specimens with variable success and complication rates. Currently, the data are conflicting and more randomized trials comparing these needles to standard needles are required. The study was conducted at Medanta - The Medicity, Gurgaon as Single Centre, prospective, observational study. All the Patients, above 18 years of age, having intestinal and extraintestinal solid mass lesions including lymphadenopathy, were subjected to EUS guided FNA. The study was conducted from June 2016 to May 2017. Patients with cystic lesions, refused to sign the informed consent and with coagulopathy (INR>1.5, Platelets <50000) were excluded from study. Total 215 patients were evaluated, out of which EUS-FNA was technically feasible in 210 (97.67%) cases. Three needle passes were made in every case. There was no significant difference between these three groups with regard to the age (p-value-0.676), gender (p-value-0.856), location (pvalue-0.998), echogenicity (p-value-0.123), border (p-value-0.216), size (p-value-0.735 & 0.374) of the lesions and presence of calcification (pvalue-0.093)or necrosis (pvalue0.729). Sample suitable for pathological evaluation were

Obtained in 90.5% cases with a tissue core in 45.7% cases. 28.1% lesions were malignant,62.4% were benign and 9.5% remained undiagnosed. The histopathological diagnoses were possible in 87.1%, 90.0% and 94.3% cases respectively with 22G Procore, 19G Procore and 19G Standard needles(p-valu-.350). Samples for the presence of blood clot in order of 19G procore(70.00%) > 22G procore (50.00%) > 19 G Standard (42. 8%), (P-value0.003). There was no post procedure complications noted in any groups.

## **Bibliography**

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