ROSE in **EUS** guided FNA of **Pancreatic Lesions**

Guy's Hospital, London, 16 April 2018

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Imperial College NHS Trust Cytology Workload

- Cervical Cytology 57,500 (decreases 8-10%/year)
- Diagnostic Cytology 10,500 of which 30% FNA (increases 5%/year)
- FNA clinic managed by <u>cytopathologist</u> terminated
- Most FNA by U/S, EUS, EBUS, few by CT
- 600 EUS/EBUS in 2017
- The rest done by clinicians in the Rapid access clinics (Head&Neck, Thyroid, Breast, Lymphnodes)

No of FNA cases



Pancreatic Mass: Solid or Cystic?

- Solid Pancreatic masses
- Ductal Adenocarcinoma
 - typical
 - variant
- Chronic Pancreatitis
- Acinar Cell Carcinoma
- Pancreatic Endocrine Tumour (PNET)
- Pancreatoblastoma

- Cystic pancreatic masses
- Pseudocyst
- Serous Cystadenoma
- Solid pseudopapillary tumour
- Mucinous cyst
 - MCN
 - IPMN

Handling of ROSE samples: the **BMS**

- Direct air dried Diff Quick smears
- Assess whether there is material
- If yes, is it <u>representative</u> of the intended site?
- Is there contamination? (depends on <u>Pathway</u> of site)
- HOP(duodenal), TOP (gastric), Hilum (liver), adrenal, mesothelial
- Is it a Solid or cystic mass?

Role of the **BMS**

- Check Clinical Details
- Liaise <u>with</u> endoscopist *regarding* the query
- Check <u>whether</u> representative
- Suggest <u>further</u>...... studies (?lymphoma for Flow Cytometry)
- If atypical cells present, ask for **dedicated pass** in LBC

Adenocarcinoma



Difficult Differential Diagnosis: Reactive ductal atypia in chronic pancreatitis vs. better differentiated adenocarcinoma





BSCC Code of Practice--Fine Needle Aspiration Cytology. Kocjan G1, Chandra A, Cross P, Denton K, Giles T, Herbert A, Smith P, Remedios D, Wilson P. Cytopathology. 2009 Oct; 20(5):283-96.

- FNA cytology has been shown to be a cost-effective, reliable technique its accurate interpretation <u>depends</u> on obtaining <u>adequately</u> cellular samples <u>prepared</u> to a <u>high</u> standard.
- Its accuracy and cost-effectiveness can be seriously compromised by <u>inadequate</u> samples

Cont....

- Cytopathologists, Radiologists, Nurses or Clinicians may take FNAs, they must be adequately trained, experienced and subject to regular audit.
- The best results are obtained:
- when a pathologist or an experienced & trained Biomedical Scientist (cytotechnologist) provides immediate on-site assessment of <u>sample adequacy</u> &
- whether or not the FNA requires image-guidance.

EUS-guided FNA for diagnosis of **solid** pancreatic neoplasms: A meta-analysis GIE 2012

- 33 studies, 12 retrospective, **21 prospective**
- 4,984 patients
- Sensitivity for malignancy 85-91 %
- Specificity " 94-98%
- PPV 98-99%
- NPV 65-72%

EUS-guided FNA for diagnosis of solid pancreatic neoplasms

- False -Ve results up to 20-40 %
- False +ive very rare

Optimizing **Diagnostic yield** from EUS-FNA. Cytopathology June 2013

- ROSE increases diagnostic sensitivity & accuracy of FNA for solid pancreatic masses by up to 10-15 %
- Meta-analysis of 34 studies with 3644 patients : ROSE : p=0.001 for accuracy

Costs

- 1 EUS procedure = 1hour (45'+15')
- 1 session/week of a cytopathologist (3.5 hours=**£9700** gross/year)
- 1 session/week of a <u>BMS</u> gr7 = £2700

BMS Training Course in CT/US guided FNA Cytology Imperial College NHS Trust, Dept. of Cellular Pathology

• Aim of the course:

 Provide training to senior cytology BMSs in order to assist Radiologists and clinicians in the evaluation of cytological material obtained through CT/US guided FNAs including EUS and EBUS procedures

 Maximize the potential of <u>cytological material</u> for diagnostic ancillary techniques & research protocols The course will run in 3 hour sessions on Tuesday morning (half day) from 10.00 to 13.00 on a weekly basis including **lectures** by **BNISs**, **cytopathologists**, **radiologists** and **clinicians** March 11, 9 am- Cytology of respiratory tract

Dr Onn Kon - Indications and Clinical setting Dr C Wright - EBUS

March 18, 10 am - Cytology of respiratory tract Dr F Mauri – Lung Pathology Dr F Mauri - Cytology and ancillary techniques

March 26, 14.00 – 14.45 Lung and Thyroid Dr N Strickland - CT guided FNA Dr R Dina – Thyroid Cytology and ancillary techniques

April 1, 10 am - FNA of ThyroidMr F Palazzo - Clinical settingDr M Crofton - - US guided FNA of thyroid nodules

April 8, 10 am - FNA of pancreas and cytology of biliary tract Dr P Vlavianos - Clinical setting Dr R Dina - Cytology and ancillary techniques

April 15, 10 am – FNA of head and neck
Dr A Sandison - Clinical setting and Pathology
Dr D Blunt - US guided FNA of head and neck
Dr R Dina – Head and neck cytology
May – Assessment and Evaluation

Current setting

- All U/S-guided FNAs at HH if ROSE requested are attended by a senior BMS gr7
- All U/S-guided FNAs at SMH smeared by the Radiologists (trained)
- All EUS-guided FNAs attended by a BMS gr7
- EBUS-guided FNAs attended by a BMS **if** granulomas suspected (TB or sarcoid),
- But by a cytopathologist **if** cancer suspicion/staging

Diagn Cytopathol. 2018 Apr;46(4):293-298 (ROSE VS non ROSE)

230 specimens (218 patients) were obtained from:

- pancreas (114), lymph node (64), submucosal lesions of the GI tract (27), liver (8), and miscellaneous (17) sites.
- The results were classified as informative (77.8%) and non-informative (NI) (22.2%).

The NI rate was significantly high, when a <u>cytopathologist</u> was **absent** (P = .0008)

Diagn Cytopathol. 2018 Feb;46(2):154-159 (Cyto VS core biopsy)

A total of **48 patients** with solid pancreatic lesions were evaluated. The proportions of adequate samples were **48/48** (100%) for FNA and

45/48 (93.7%) for core biopsy (P = .24). The diagnostic yield was **42/48** (87.5%) and **33/48** (68.7%) for FNA and CNB respectively (P = .046). The incremental increase in diagnostic yield by combining both methods was **2/48** (4%).

The diagnostic yield for malignancy was 30/32 (93.7%) for FNA and 23/32 (71.8%) for CNB (P = .043).

The **sensitivity** for the <u>diagnosis</u> of malignancy for:

FNA 90.6% and <u>CNB</u> were **69%**, (P = .045).

TO ROSE OR NOT TO ROSE?

• J Gastroenterol Hepatol. 2014 Apr;29(4):697-705. (metanalysis)

The search produced **3822** original studies, of which 70 studies met our inclusion criteria. The overall <u>average adequacy rate</u> was **96.2%** (95% confidence interval: 95.5, 96.9).

ROSE was associated with a *statistically significant improvement* of up to **3.5%** in adequacy rates. There was heterogeneity in adequacy rates across all subgroups. No association between the assessor type and adequacy rates was found.

Studies with ROSE have **high per-case adequacy** and a **relatively high number of needle passes** in <u>contrast to non-ROSE</u> studies.

Causes of discordance between Cytology & Histology in pancreatic lesions: the experience at Imperial College NHS Trust. M. El Shiek, R.Dina

- All pancreatic FNA cytology specimens performed in our department from 2013 to 2016 with corresponding subsequent surgical specimens were identified.
- For each case the reported cytological category was recorded (C1 inadequate,C2 benign,C3 atypical; mucinous lesions, endocrine lesions, C4–suspicious for malignancy, C5–malignant).
- The final surgical diagnosis was recorded. <u>Discordant cases</u> (benign histo vs C4,C5 cytology or malignant histo vs C2,C3 cytology), were retrieved from filing archives and reviewed by a cytopathologist <u>blinded</u> to the previous results. The cytological categories on review were compared to those originally reported.

Causes of discordance between cytology and histology in pancreatic lesions: the experience at Imperial College NHS Trust. M. El Shiek, R.Dina

- A total of **75 cytology specimens** with <u>corresponding surgical</u> specimens were identified.
- A total of **17 cases** (22.6%) were discordant.
- Six out of 14 reviewed cases were confirmed to be correctly categorised (42.8%), the discordance <u>due to nonrepresentative</u> sampling.
- Remaining eight cases (67.2%), 2 were interpreted as inadequate (C1) while 6 were given a different cytological category on review which was at most <u>one tier above or</u> <u>below</u> the original cytological diagnosis.

