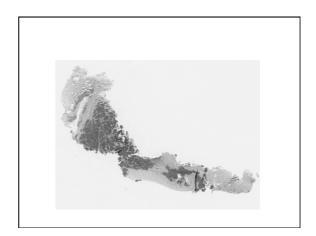
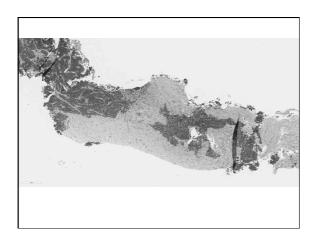
Slide seminar Case 12 Colin A Purdie BDIAP Symposium on Breast Pathology Saturday 25th November 2017

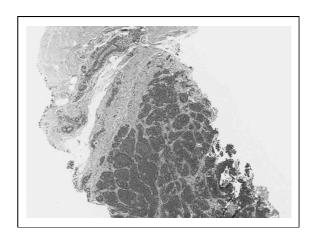
Case 12

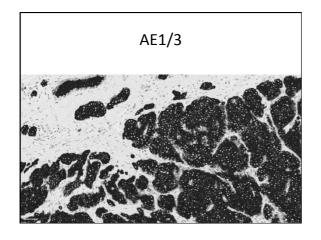
- Female, 72
- Asymptomatic
- Screen detected solid mass right breast
 - M3, 11mm
 - U3, 8mm

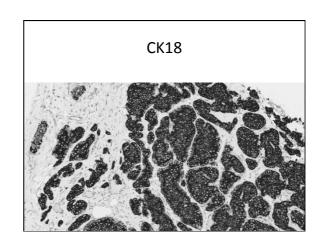


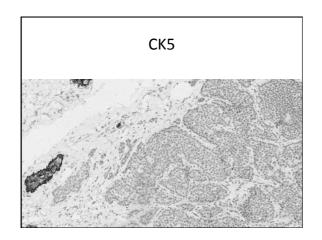


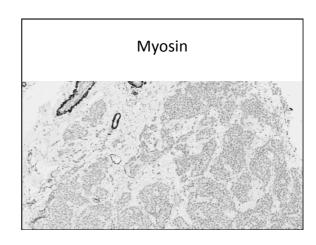


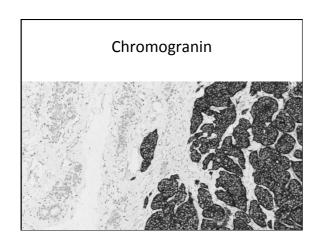


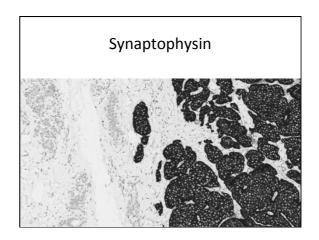


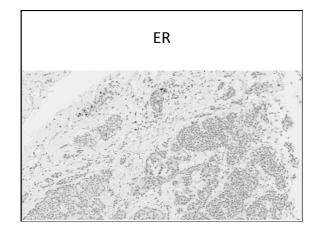


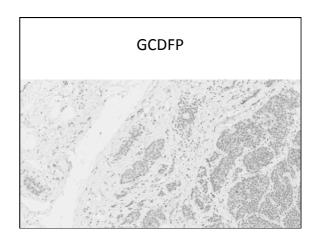


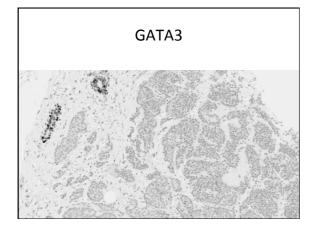












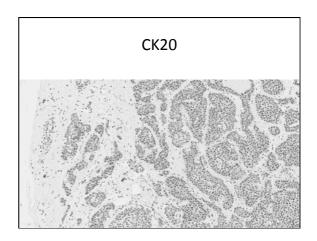
Neuroendocrine Carcinomas

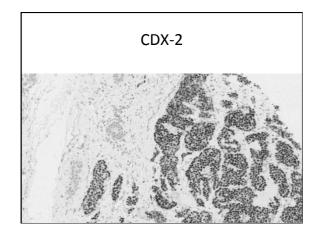
- Expression of NE markers
 - Chromogranin
 - Synaptophysin
 - NSE
- Dense core granules on EM
- Usually ER and PR positive

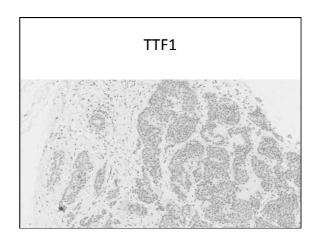
Carcinomas with neuroendocrine features

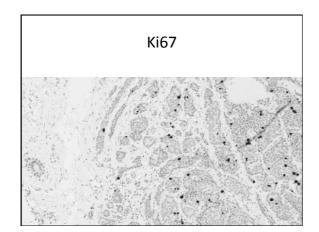
- Neuroendocrine tumour, well-differentiated Densely cellular, solid nests and trabeculae

 - G1 or G2
- Neuroendocrine carcinoma, poorly differentiated/small cell carcinoma
 - Identical to lung counterpart
- May be TTF1+
- Invasive carcinoma NST with neuroendocrine features
- Up to 30% of all breast carcinomas (esp mucinous Ca)
- Metastatic
 - Well-differentiated neuroendocrine tumour (carcinoid)
 - Poorly differentiated neuroendocrine carcinoma/small cell carcinoma









Management

- CT Thorax & abdo & pelvis with contrast

 Within the anterior mediastinum, there is a 4cm by 3.5cm by 5cm homogenous mass.

 There is diffuse nodal disease affecting the root of the mesentery particularly around the superior mesenteric artery and exploding down into the ileocolic region. The more confluent area measures 2.5cm.

 No clear evidence of focal bowel mass

 The pelvis appears unremarkable.

- ${\tt NM}$ Octreotide scan with SPECT :
 - History of right breast carcinoid tumour noted. No abnormal tracer uptake elsewhere to suggest metastatic disease or a primary tumour at another site.

Management

- Bloods
 - Chromogranin A 38.1 (nmol/l) (normal range 0-6)

Case 12

- Diagnosis
 - Metastatic well differentiated neuroendocrine carcinoma

References

• G. Bussolati and S. Badve, "Carcinomas with neuroendocrine features," in WHO Classification of Tumours of the Breast, S. R. Lakhani, I. O. Ellis, S. J. Schnitt, P. H. Tan, and M. J. van de Vijver, Eds., pp. 62–63, IARC Press, Lyon, France, 2012.