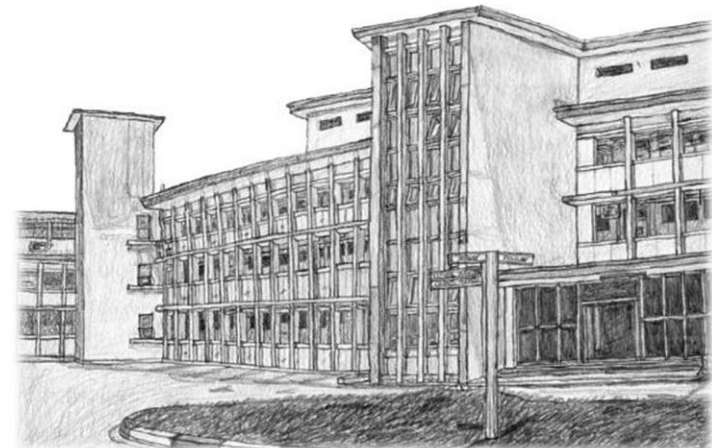
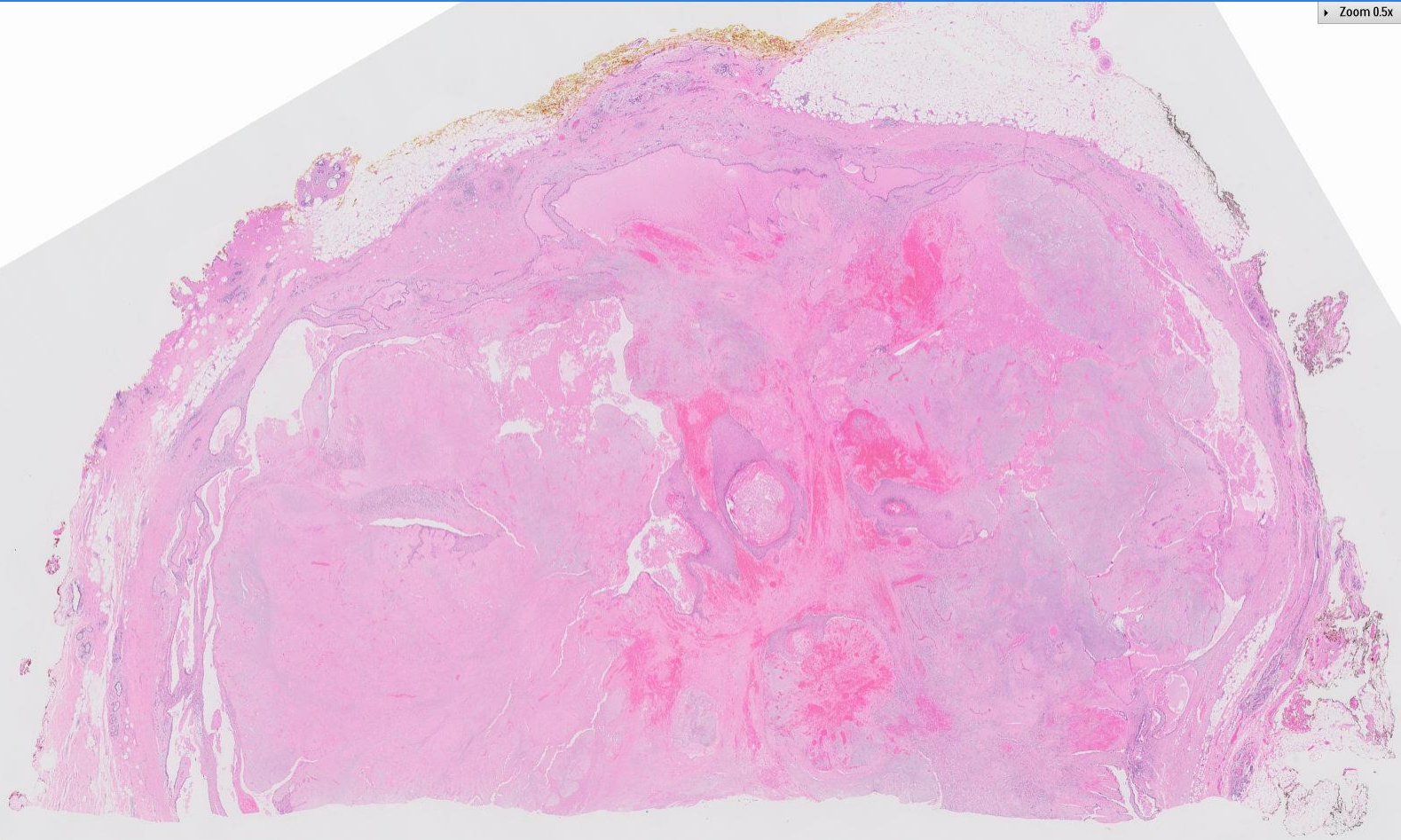


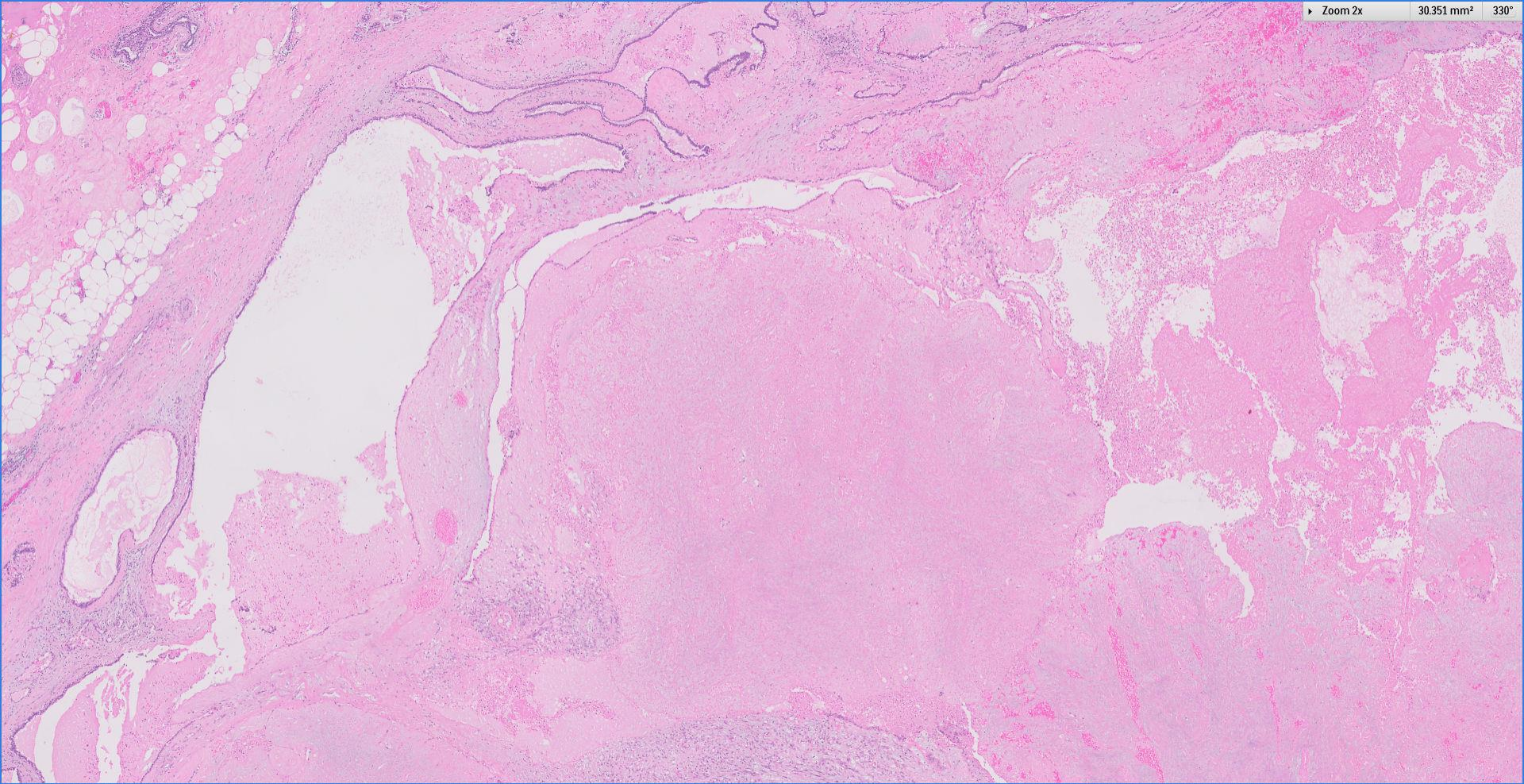
Case 30

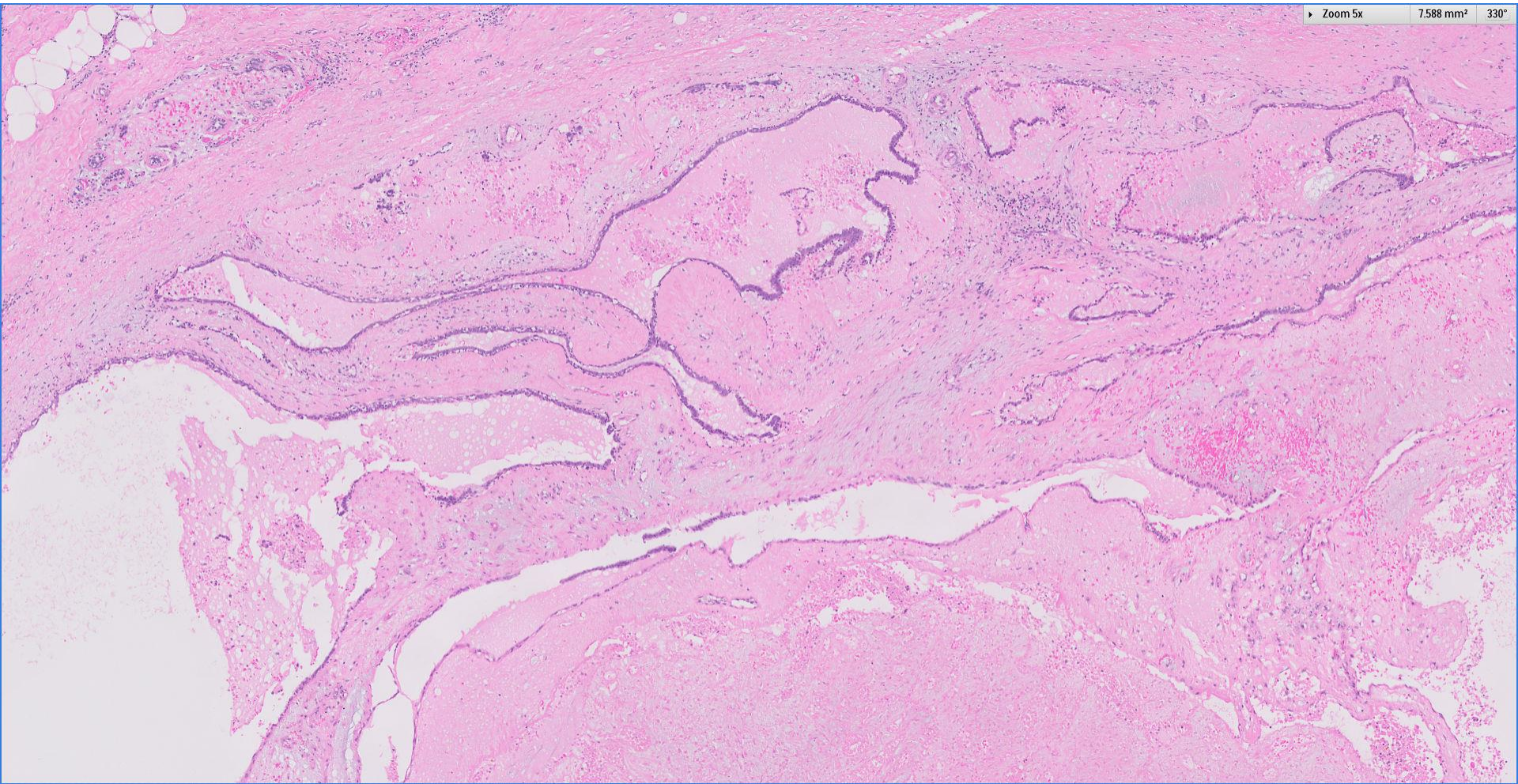
40 year old female.

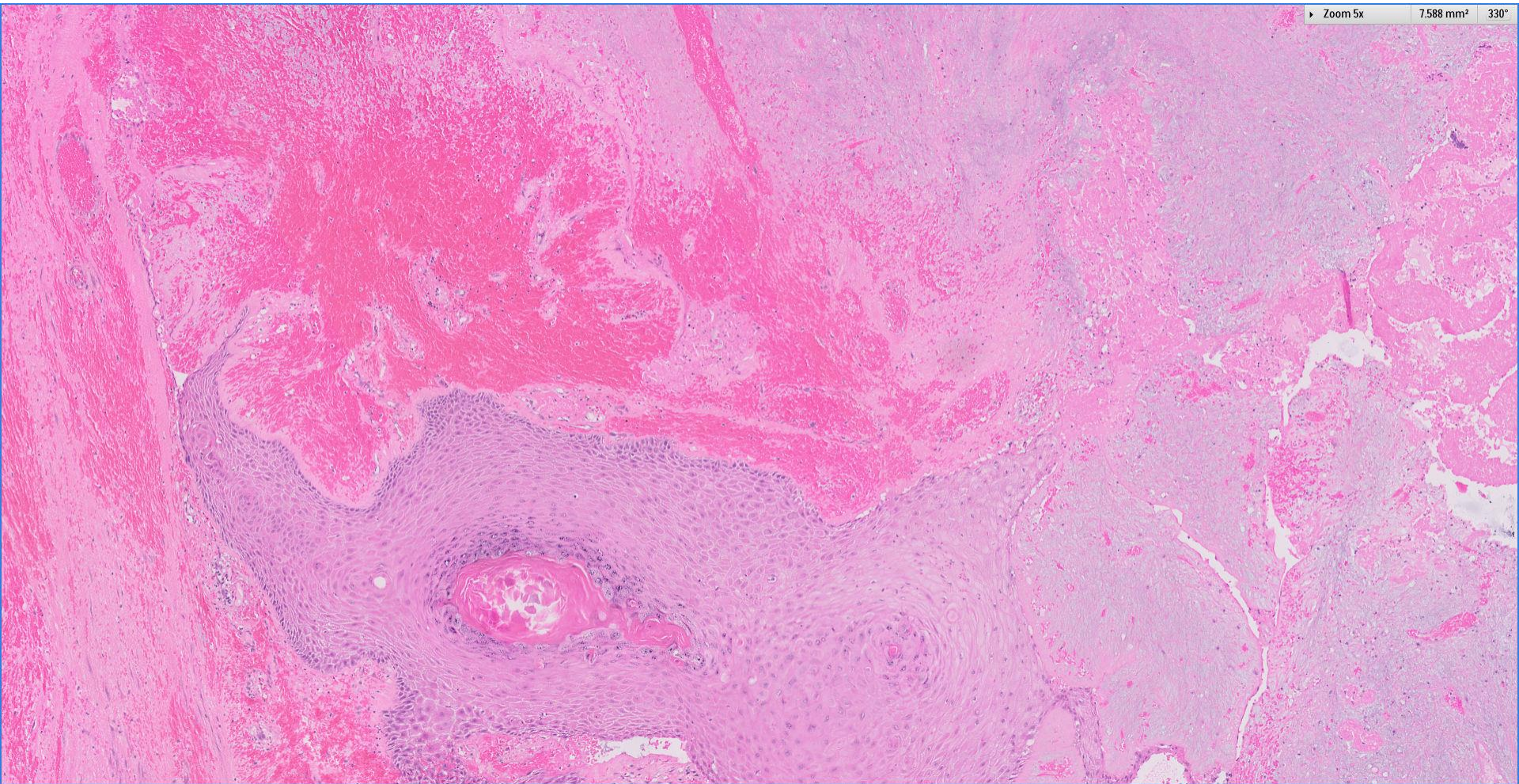
Right breast 10 o'clock mass, excised.

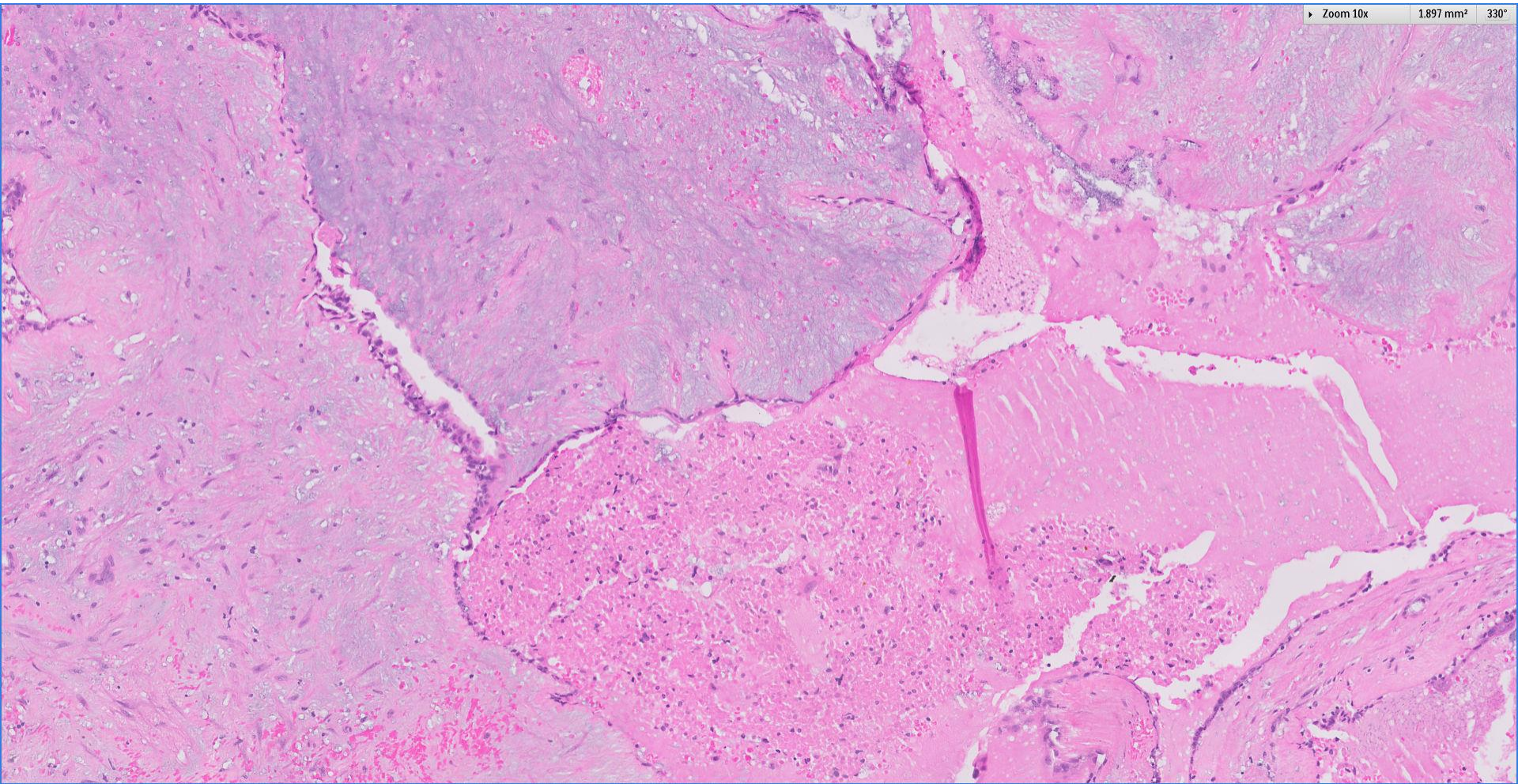




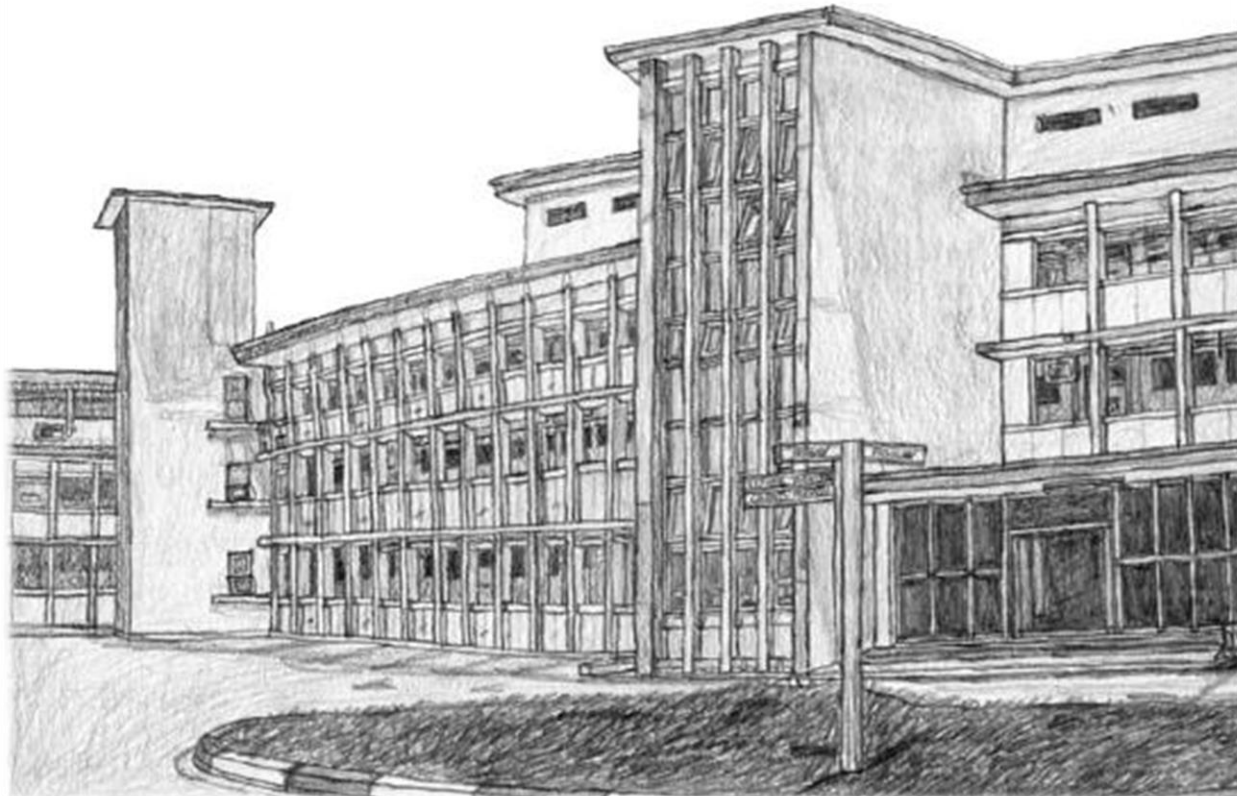


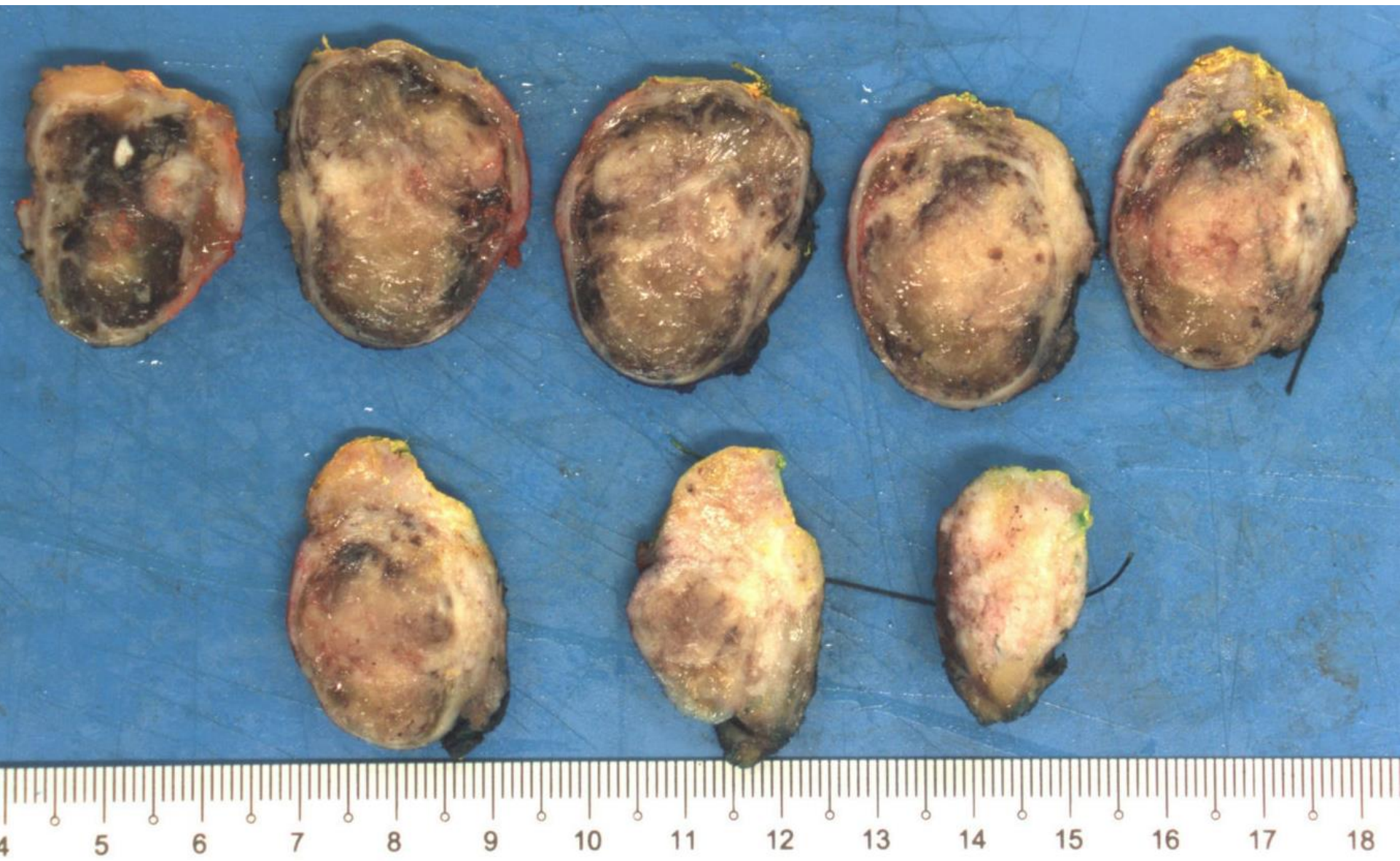






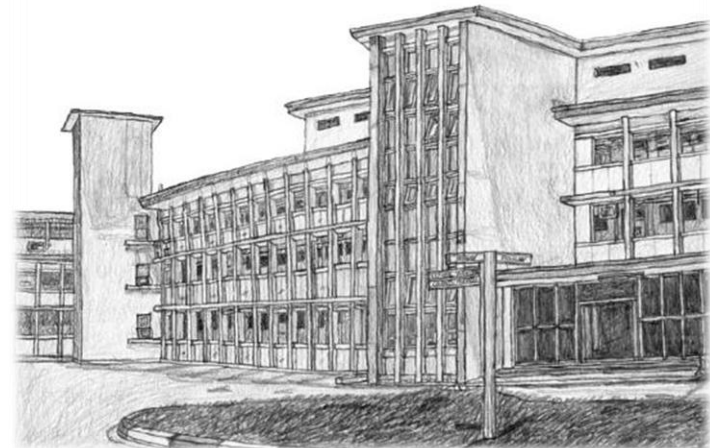
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Diagnosis

Right breast 10 o'clock mass, excision ~
**Benign phyllodes tumour with haemorrhagic
infarction and squamous metaplasia**



Phyllodes tumour & squamous metaplasia

- Squamous metaplasia is sometimes encountered in the epithelial component of breast phyllodes tumours.
- Squamous cysts may be observed.
- Squamous metaplasia may also be seen around areas of infarction.

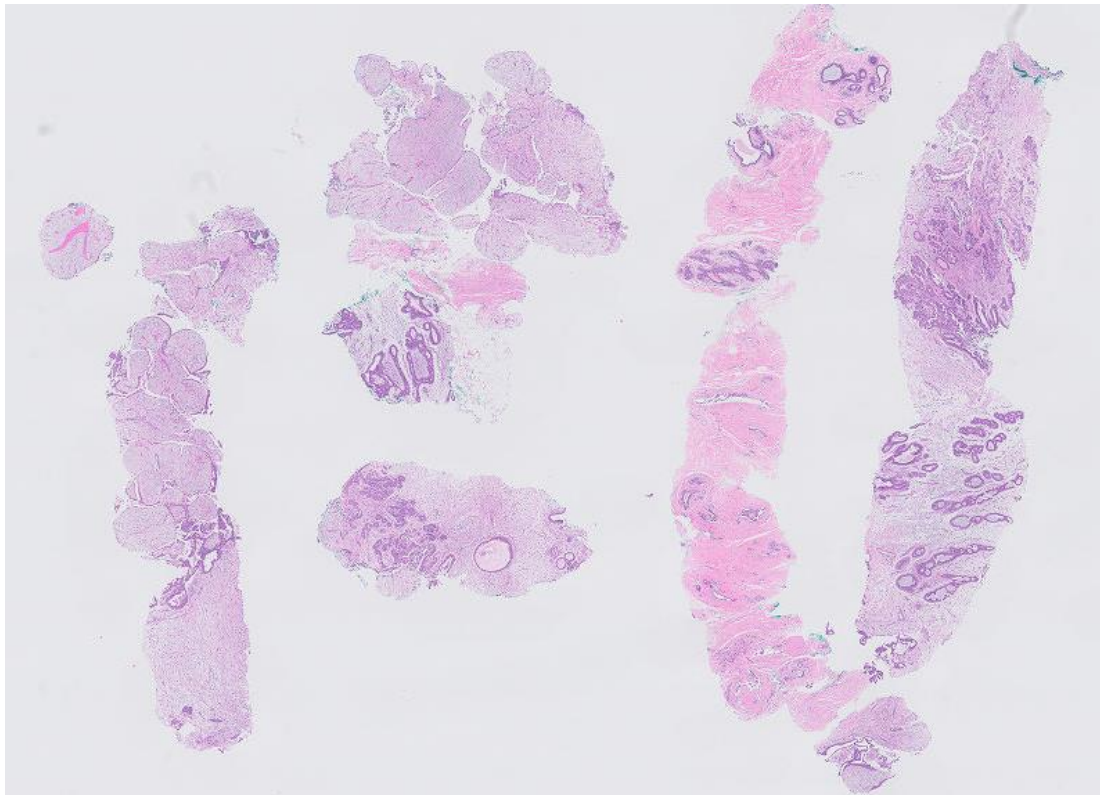


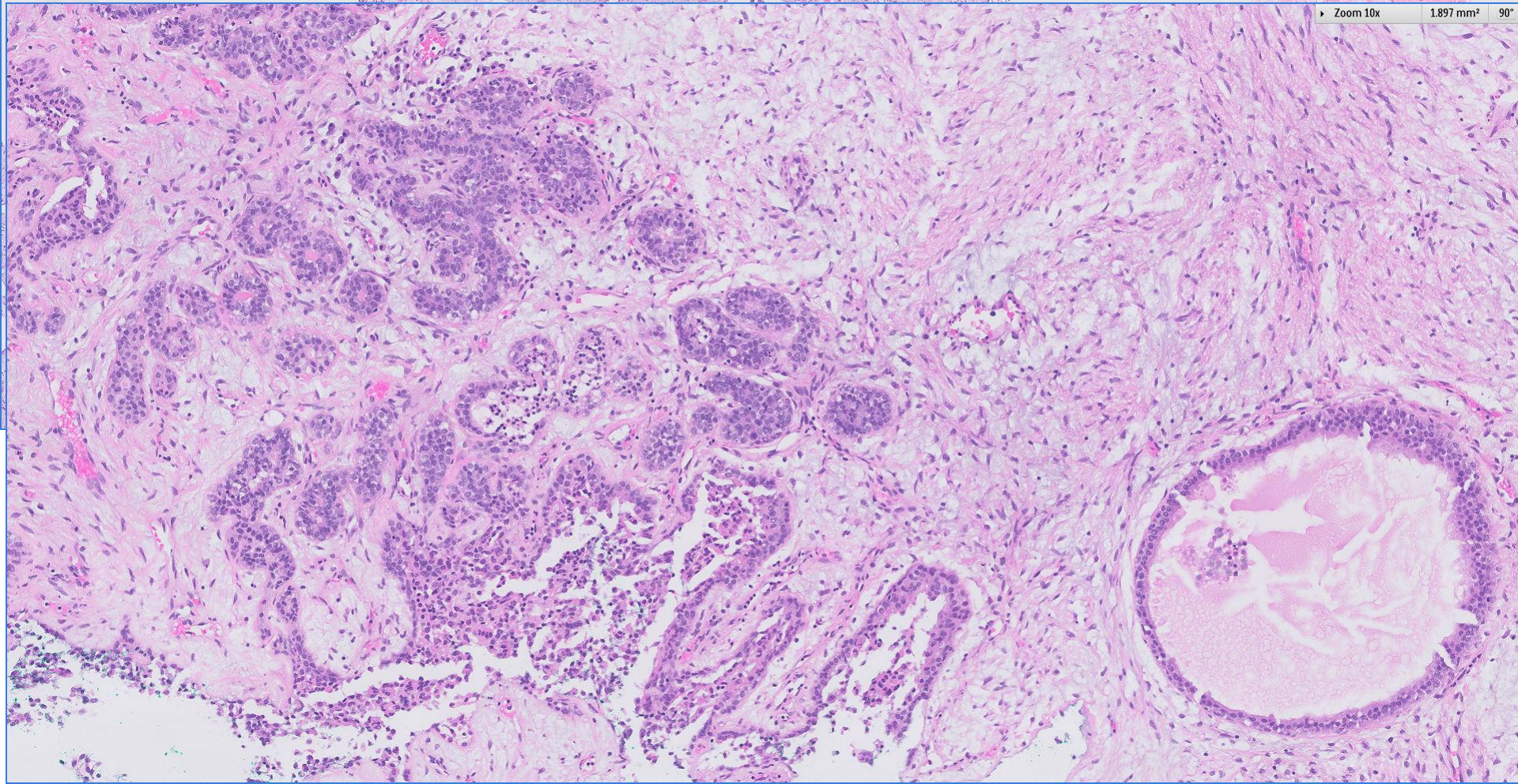
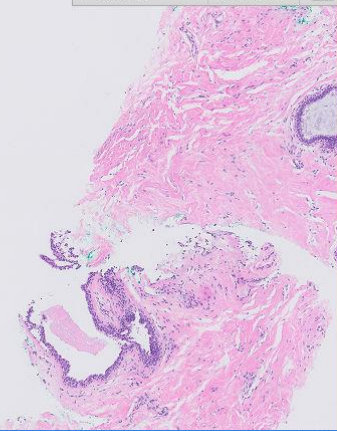
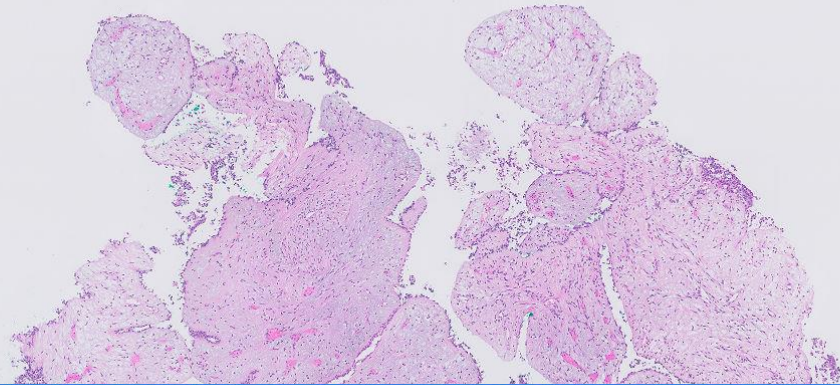
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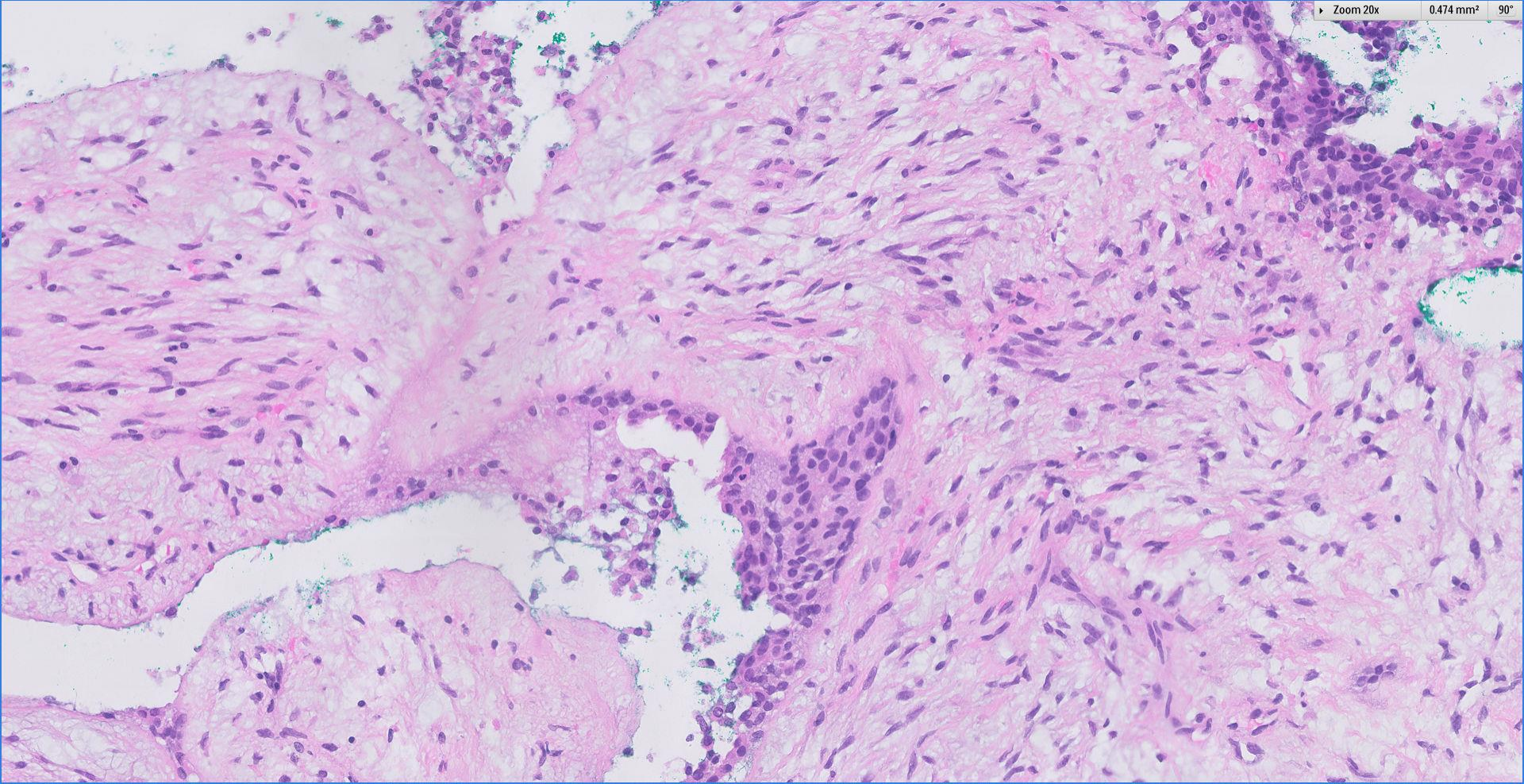
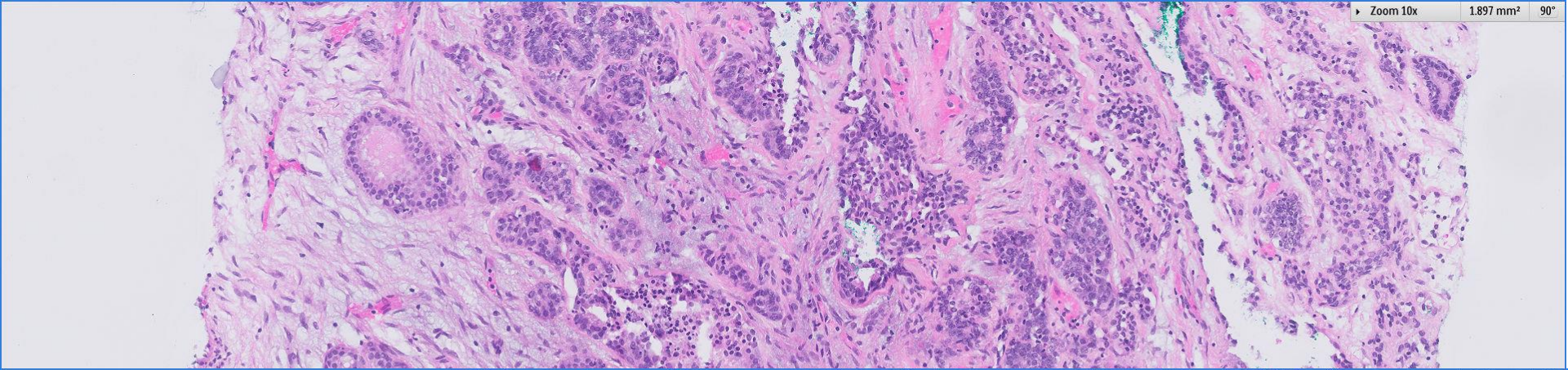


Previous core biopsy

- 7 weeks previously.
- Diagnosed as 'mildly cellular fibroepithelial lesion'





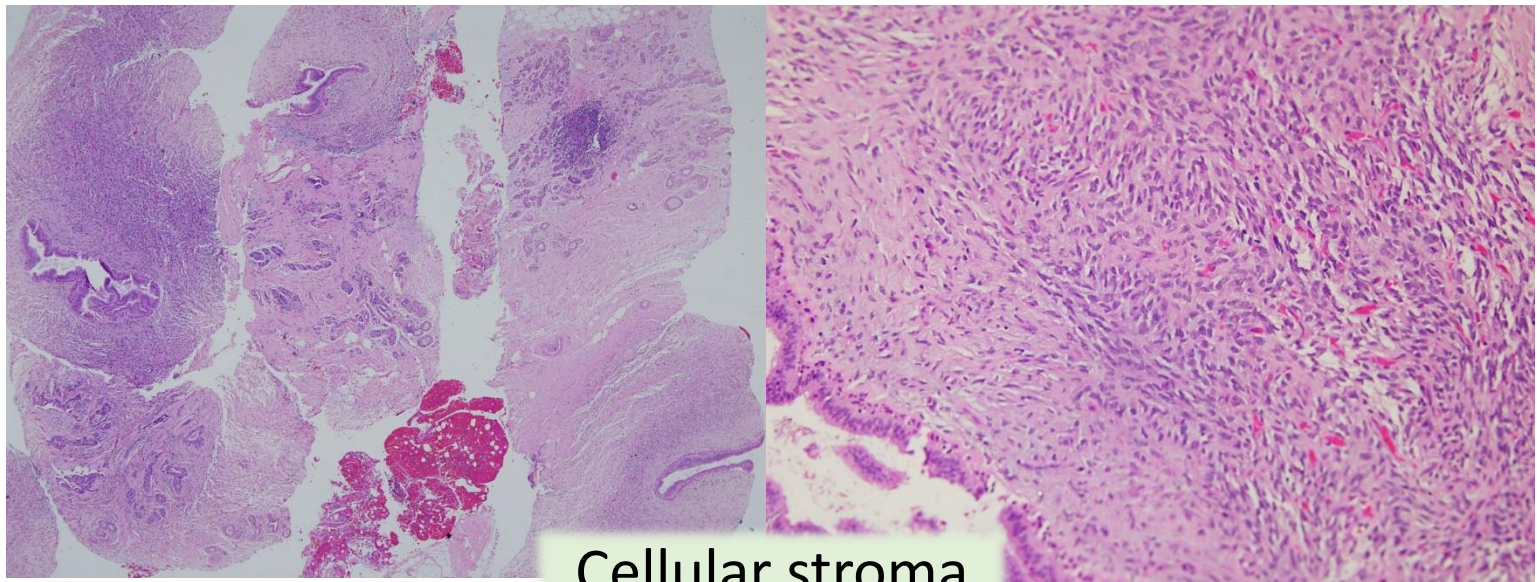


Additional breast lumps

- Right breast 1230 ~ core biopsy diagnosed fibroadenoma.
- Left breast 1000 ~ core biopsy diagnosed fibroadenoma.



Core biopsy diagnosis of ***cellular***
fibroepithelial neoplasms:
fibroadenoma or phyllodes?



Core biopsy diagnosis of **cellular fibroepithelial lesions** – prediction of phyllodes tumour

Author	Reference	Key findings predicting phyllodes tumour
Jacobs et al	<i>Am J Clin Pathol</i> 2005; 124: 342-354	Marked stromal cellularity, mitoses in moderate stromal cellularity Ki67 & topoisomerase II α indices
Lee et al	<i>Histopathology</i> 2007; 51: 336-344	Stromal cellularity \geq 50% stroma, stromal overgrowth, fragmentation, adipose within stroma
Resetkova et al	<i>Breast J</i> 2010; 16:573-80.	No predictive value of clinical, radiologic or pathologic data Suggested follow-up alone for a patient subset
Jara-Lazaro et al	<i>Histopathology</i> 2010; 57: 220-232	Marked stromal cellularity/atypia, stromal overgrowth, mitoses \geq 2 per 10 hpf, ill-defined lesional borders, Ki67 & topoisomerase II α indices \geq 5%, reduced CD34 staining
Yasir et al	<i>Am J Clin Pathol</i> 2014; 142: 362-369	Mitoses, stromal overgrowth, fragmentation, adipose infiltration, heterogeneity, subepithelial condensation, nuclear pleomorphism

RESEARCH ARTICLE

Open Access



A five-gene reverse transcription-PCR assay for pre-operative classification of breast fibroepithelial lesions

Wai Jin Tan¹, Igor Cima¹, Yukti Choudhury¹, Xiaona Wei¹, Jeffrey Chun Tatt Lim², Aye Aye Thike²,
Min-Han Tan¹ and Puay Hoon Tan^{2,3*}

Methods: We profiled the transcriptome of a training set of 48 formalin-fixed, paraffin-embedded fibroadenomas and phyllodes tumors and further designed 43 quantitative polymerase chain reaction (qPCR) assays to verify differentially expressed genes. Using machine learning to build predictive regression models, we selected a five-gene transcript set (*ABCA8*, *APOD*, *CCL19*, *FN1*, and *PRAME*) to discriminate between fibroadenomas and phyllodes tumors. We validated our assay in an independent cohort of 230 core biopsies obtained pre-operatively.

Results: Overall, the assay accurately classified 92.6 % of the samples (AUC = 0.948, 95 % CI 0.913–0.983, $p = 2.51E-19$), with a sensitivity of 82.9 % and specificity of 94.7 %.

FibroPhyllo™ Tissue Test

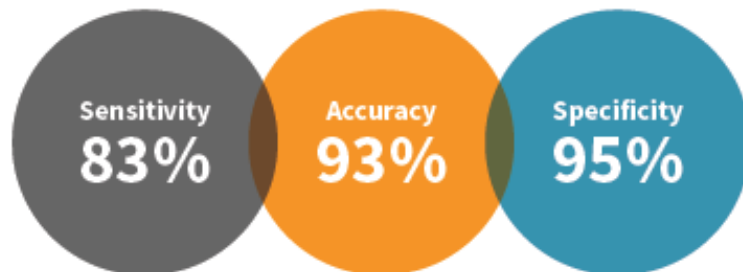
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The performance of the FibroPhyllo™ Tissue Test in pre-operative classification of breast fibroepithelial lesions was validated in a cohort study of 230 core biopsies with at least 2 years of follow-up⁵.

LUCENCE
DIAGNOSTICS

TEST PERFORMANCE



TEST REQUIREMENTS



- 1) FFPE Tissues - Core / Excisional Biopsies
 - Minimum 50 microns equivalent (e.g. 10 slides of 5 micron-thickness or 5 slides of 10 micron-thickness)
 - Slides uncharged and uncoated
- 2) Matched H&E slide with tumor region marked out
- 3) Matched histology report of tissue biopsy
- 4) Tissue curls are not **be** accepted

TURNAROUND TIME



7 days

Call our sales hotline: +65 6592 5102 or email: enquiry@lucencedx.com



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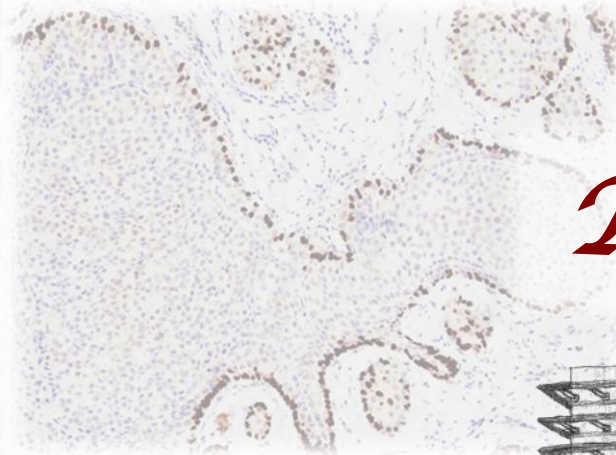
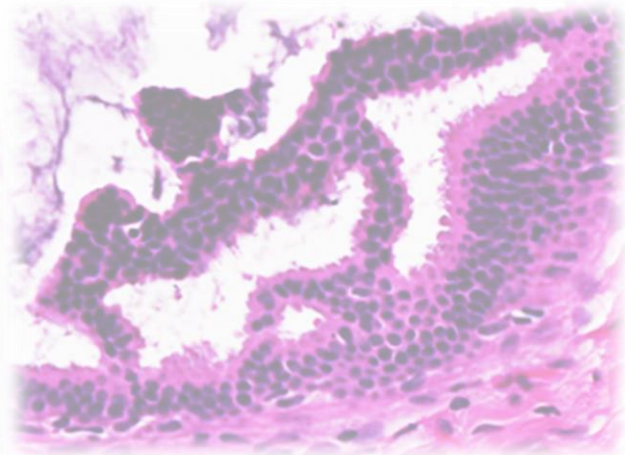
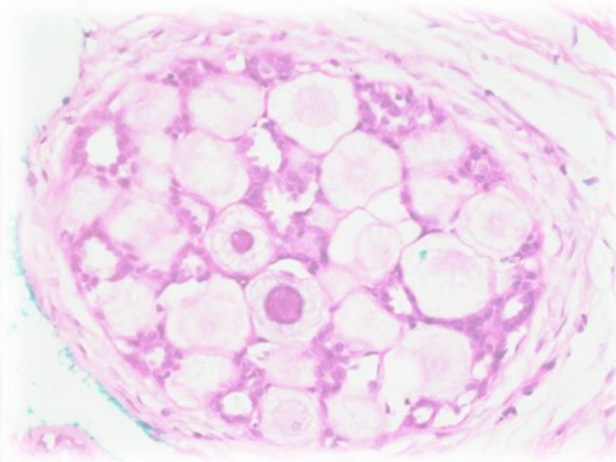
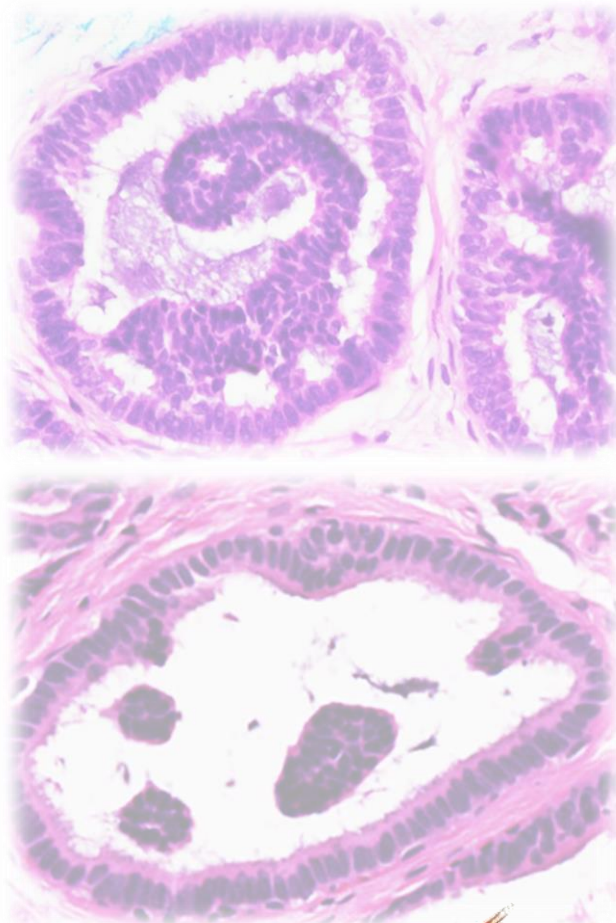


* The breast pathology team does not have any commercial or financial association with this test

Core biopsies of fibroepithelial neoplasms ~ an approach

- Benign, consistent with FA ~ may leave alone.
- Benign, favouring or unable to rule out phyllodes tumour ~ excise.
- Benign, 2 cm or larger ~ consider excision.
- Benign, but patient experiences rapid tumour growth ~ excise.
- Malignant ~ resection.

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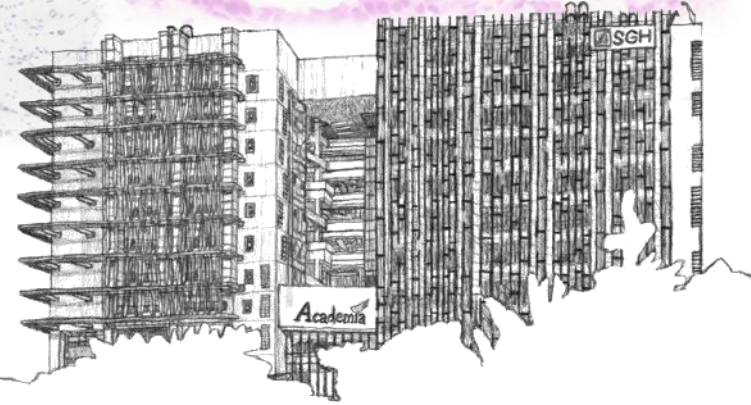


Thank you!

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General Hospital
SingHealth
Division of Pathology

 SingHealth DukeNUS
ACADEMIC MEDICAL CENTRE
PATHOLOGY

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International Academy of Pathology
Singapore Division



THE ACADEMIA,
SINGAPORE GENERAL HOSPITAL 09.05.2014

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