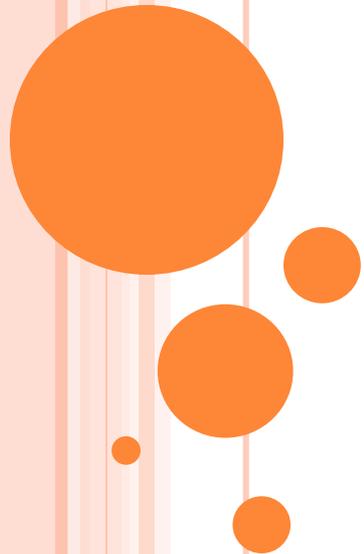


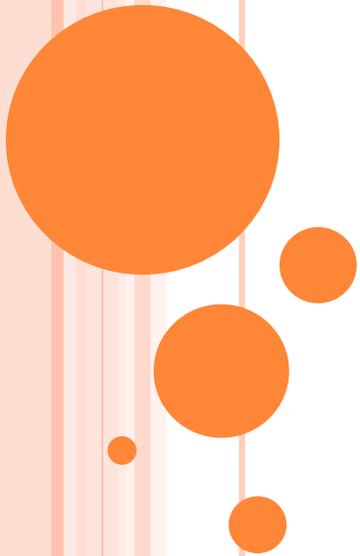


# FEMALE GENITAL SYSTEM, PATHOLOGY LAB

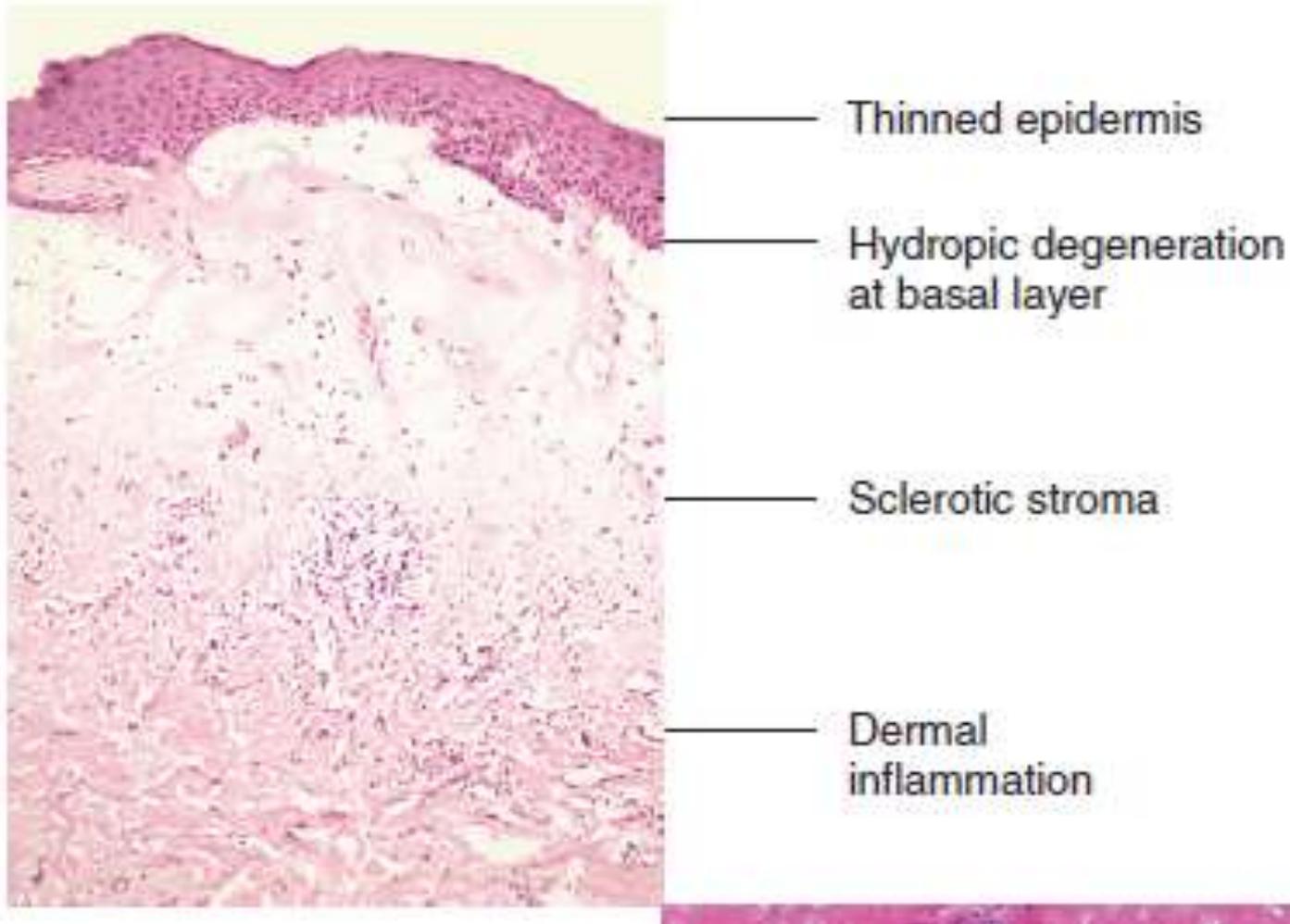


**Dr. Bushra AlTarawneh, MD**  
**Anatomical pathology**  
**Mutah University**  
**School of Medicine- Department of Microbiology &**  
**Pathology**  
**UGS lectures 2023**

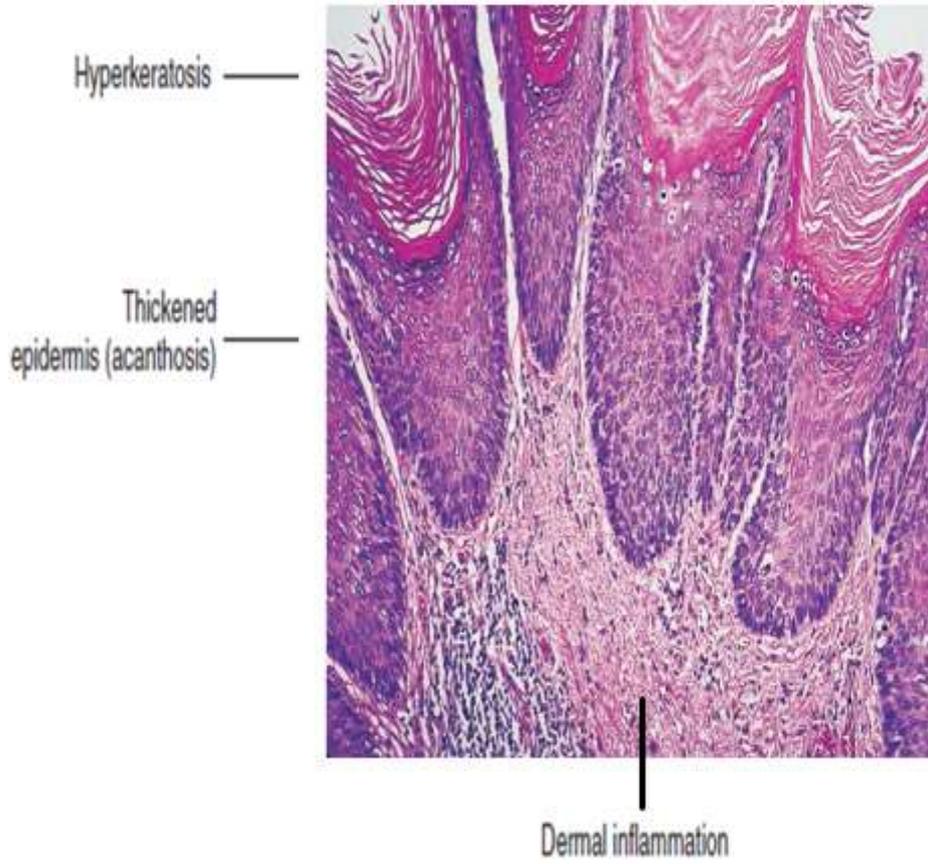
# VULVA



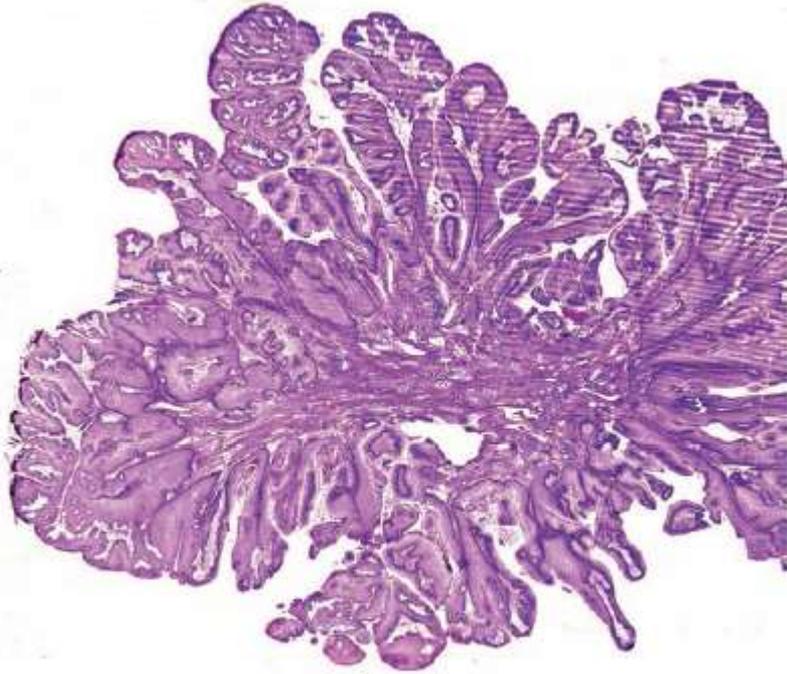
Lichen sclerosus is characterized by thinning of the epidermis, disappearance of rete pegs, hydropic degeneration of the basal cells, dermal fibrosis, and a scant perivascular mononuclear inflammatory cell infiltrate.



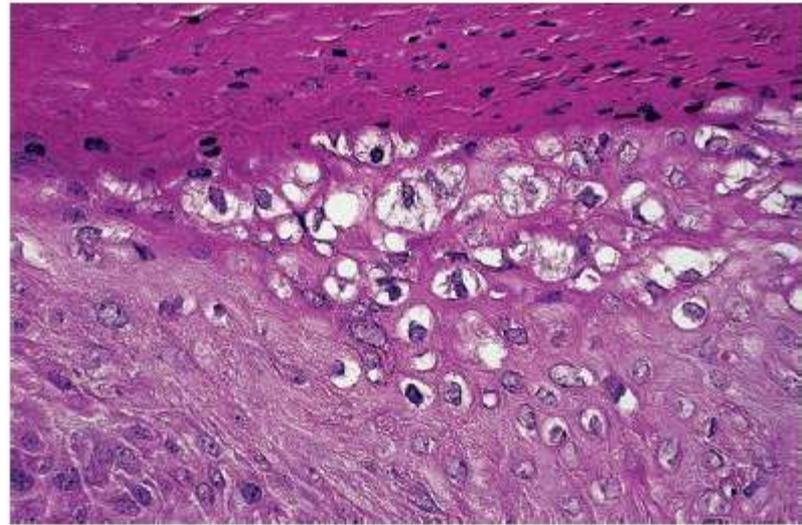
- Lichen simplex chronicus is marked by epithelial thickening (particularly of the stratum granulosum) and hyperkeratosis. Increased mitotic activity is seen in the basal and suprabasal layers; however, there is no epithelial atypia. Leukocytic infiltration of the dermis is sometimes pronounced.



On histologic examination, the characteristic cellular feature is **koilocytosis**, a cytopathic change characterized by perinuclear cytoplasmic vacuolization and wrinkled nuclear contours that is a hallmark of HPV.



Whole mount of condyloma acuminatum of vulva.



Prominent koilocytotic changes in vulvar epithelium.



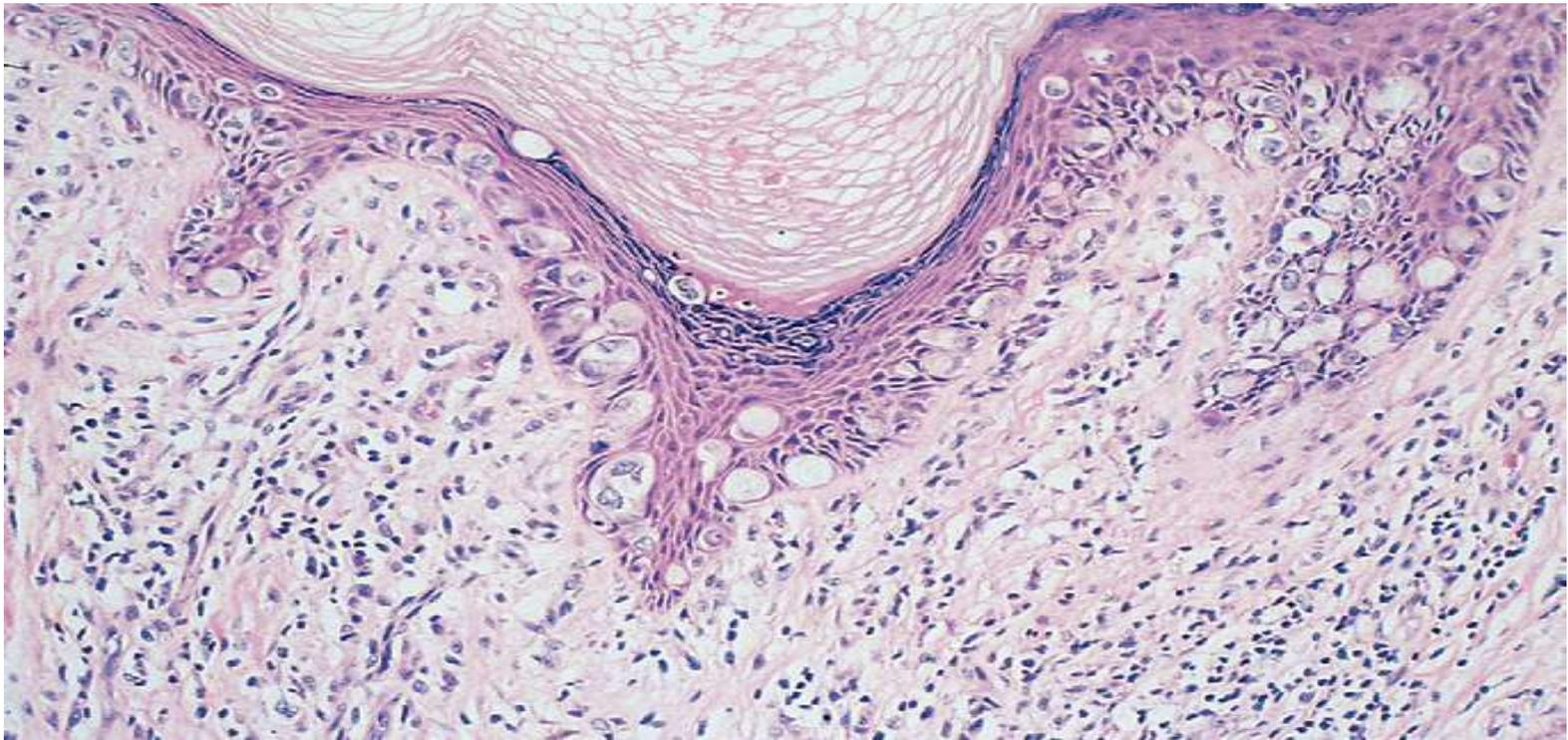


Large condyloma of vulva.

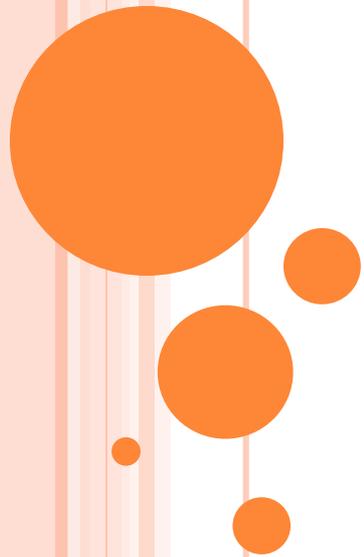


# EXTRAMAMMARY PAGET DISEASE

On histologic examination, large epithelioid cells with abundant pale, finely granular cytoplasm and occasional cytoplasmic vacuoles infiltrate the epidermis, singly and in groups. The presence of mucin, as detected by periodic acid–Schiff (PAS) staining, is useful in distinguishing Paget disease from vulvar melanoma, which lacks mucin.



# VAGINA



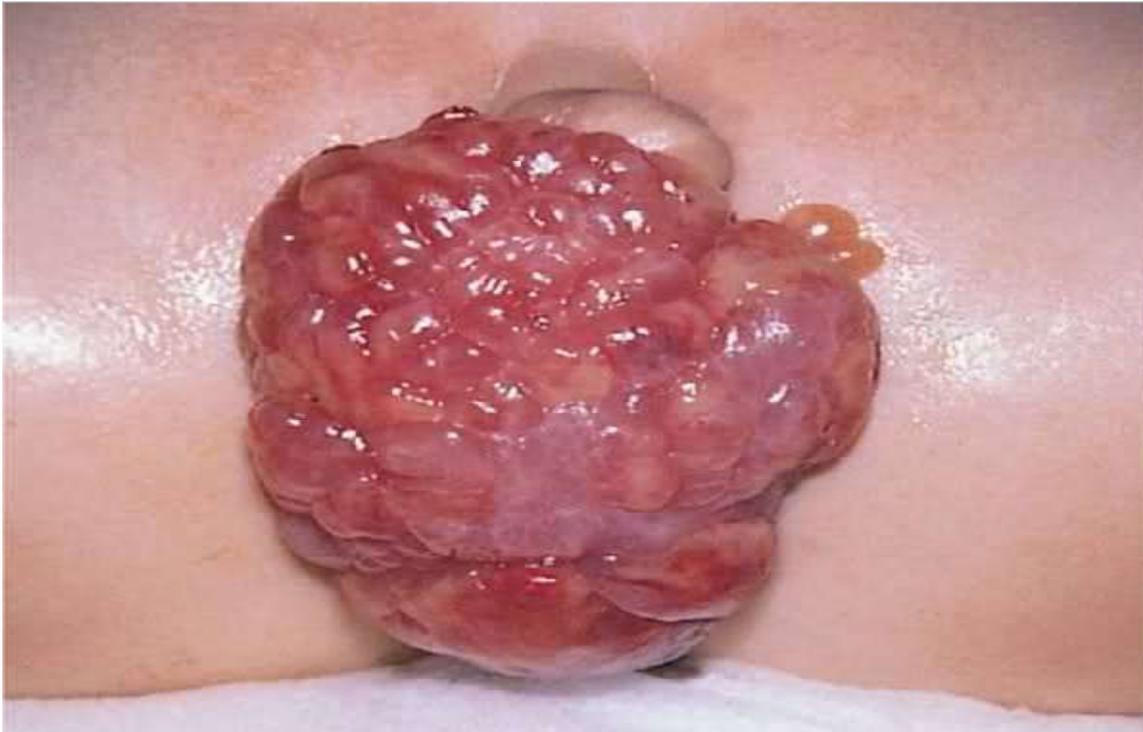
# SARCOMA BOTRYOIDES

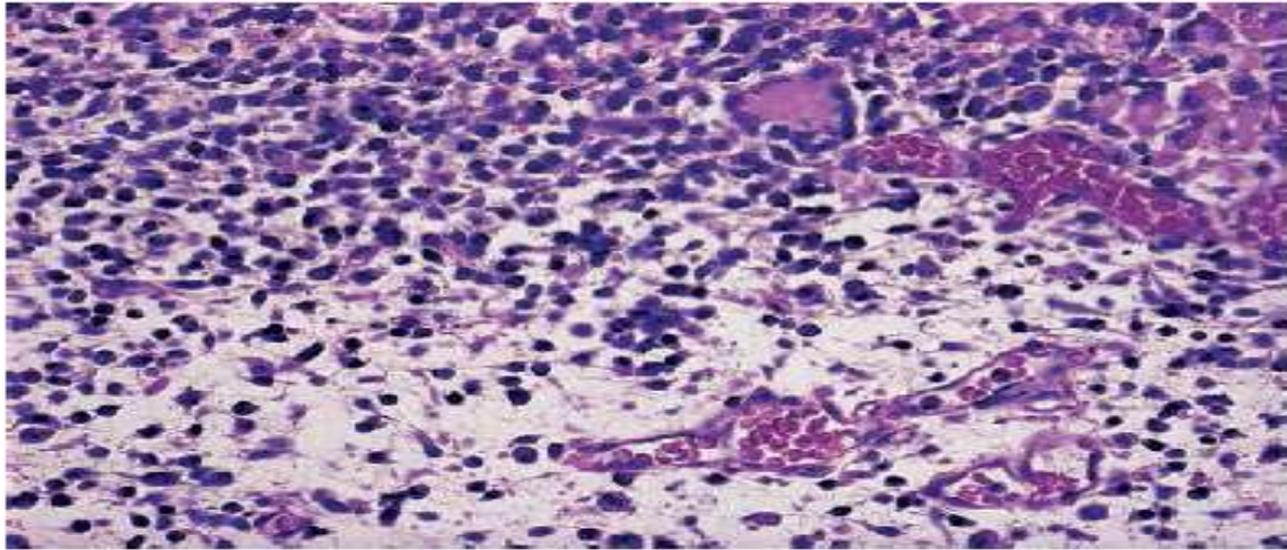
- is a rare form of primary vaginal cancer.
- It usually is encountered in infants and children younger than 5 years of age. It also may occur in other sites, such as the urinary bladder and bile ducts.
- Grossly, it presents as a conglomerate of soft polypoid masses resembling a bunch of grapes—hence its designation as “botyroid.”



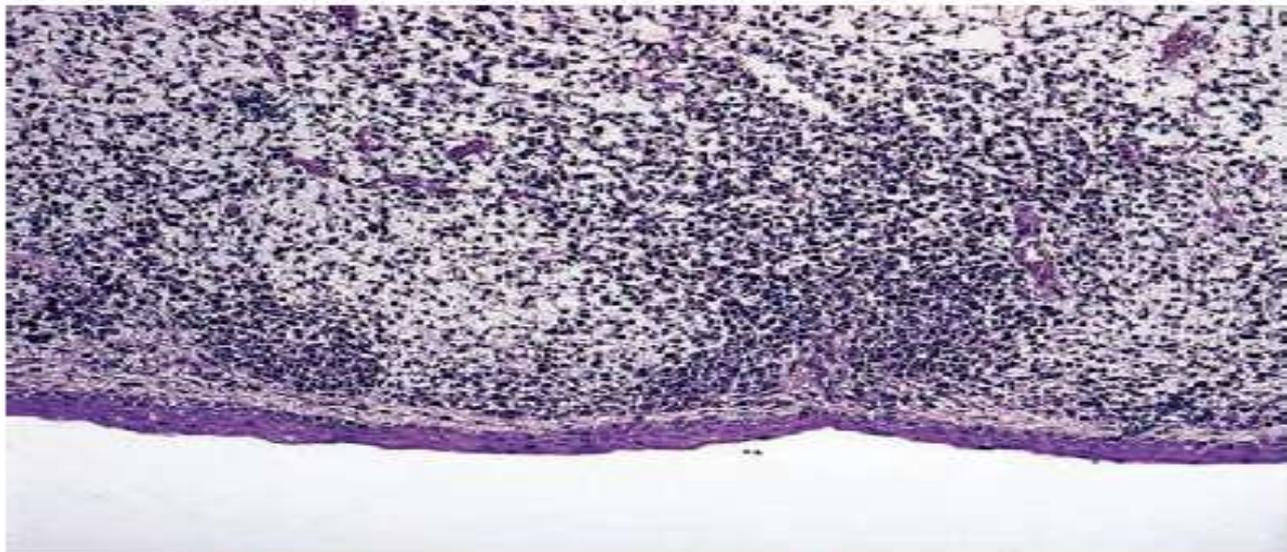
# SARCOMA BOTRYOIDES (EMBRYONAL RHABDOMYOSARCOMA)

THE GRAPE-LIKE CONFIGURATION OF BOTRYOID EMBRYONAL RHABDOMYOSARCOMA OF VAGINA. IS CHARACTERISTIC.





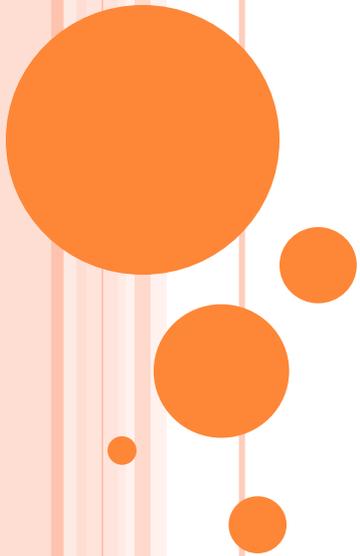
Microscopic Appearance of Embryonal Rhabdomyosarcoma. The differential diagnosis is that of small round cell tumors.



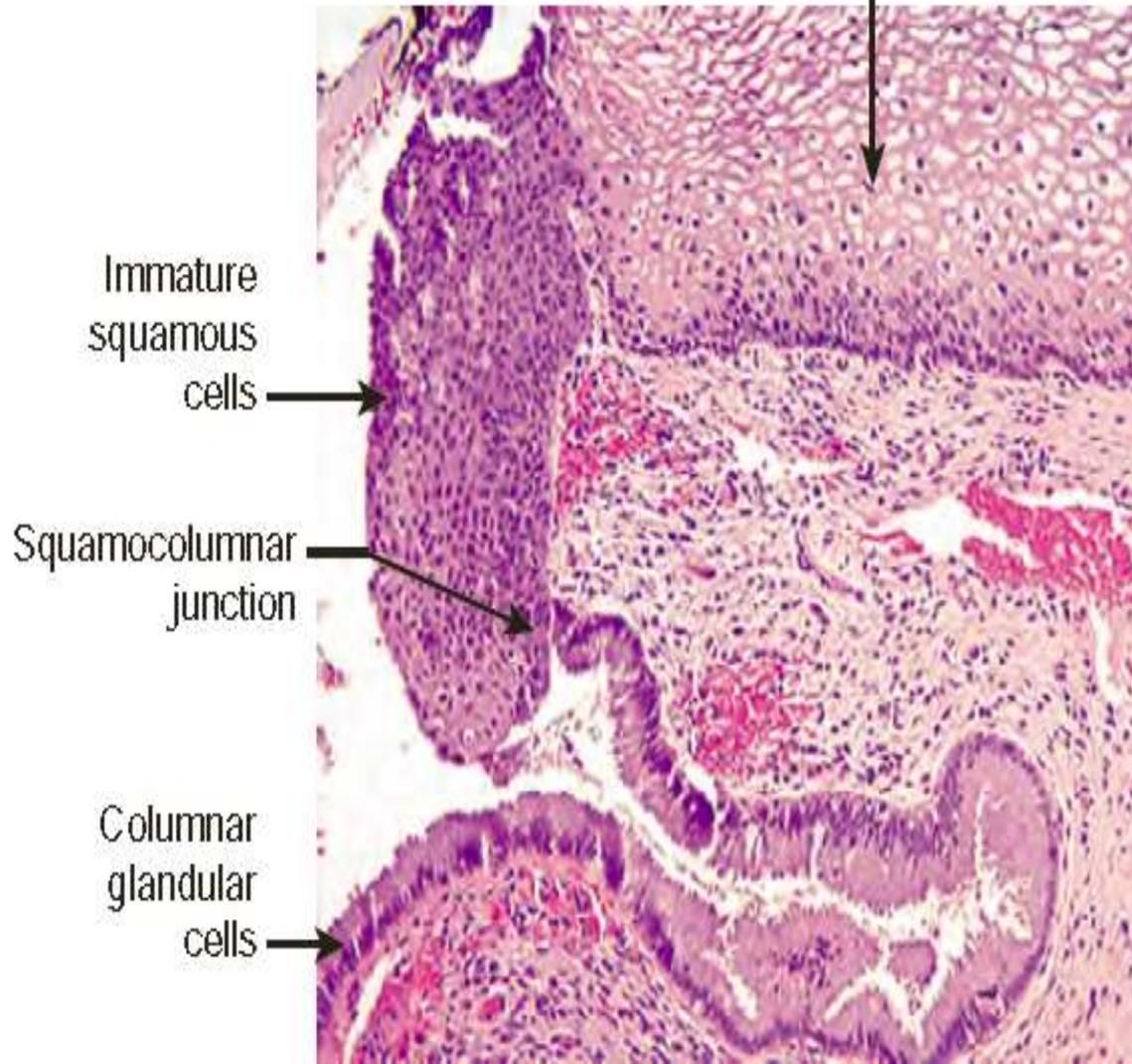
So-called cambium layer beneath non-neoplastic epithelium in embryonal rhabdomyosarcoma.



# CERVIX



Mature squamous cells



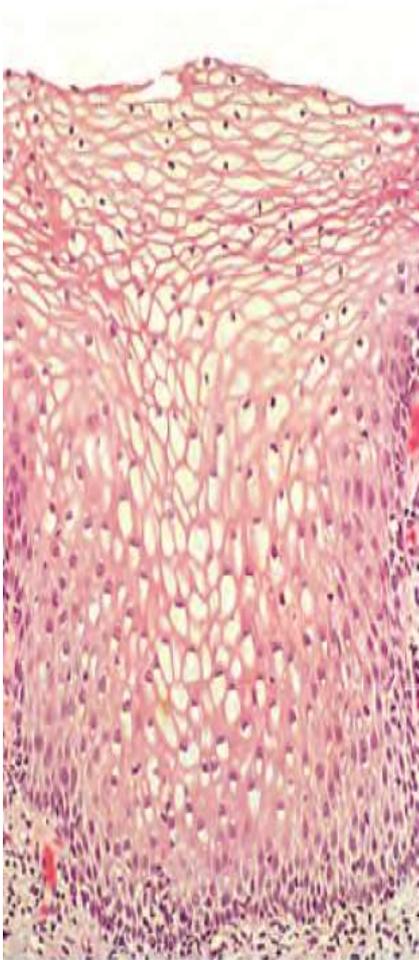
Immature squamous cells

Squamocolumnar junction

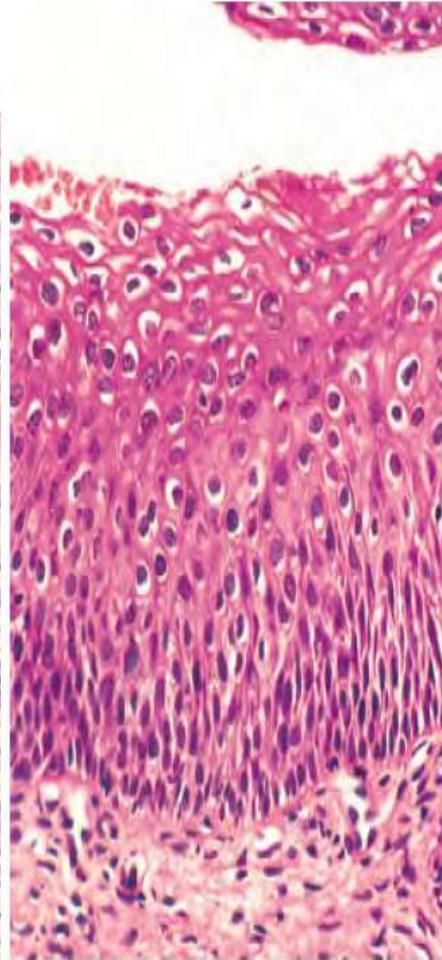
Columnar glandular cells



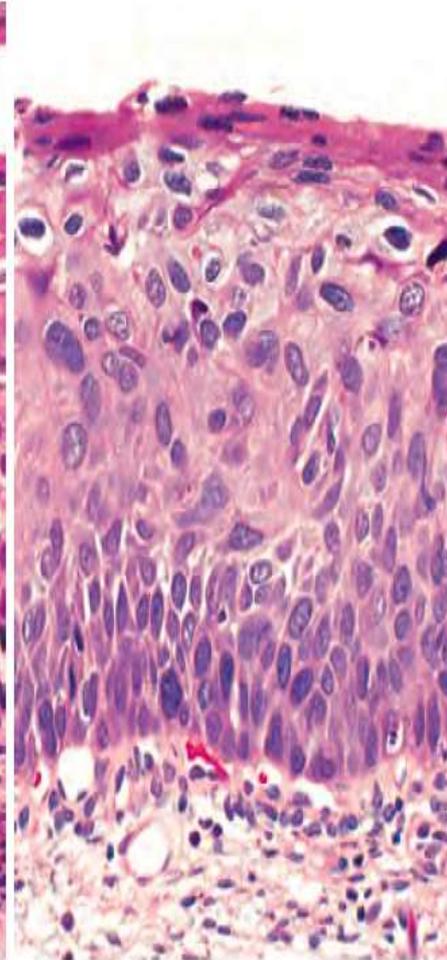
**CIN → DYSPLASIA: NUCLEAR ENLARGEMENT, HYPERCHROMASIA (DARKER), COARSE CHROMATIN, & VARIATION IN NUCLEAR SIZE & SHAPE**



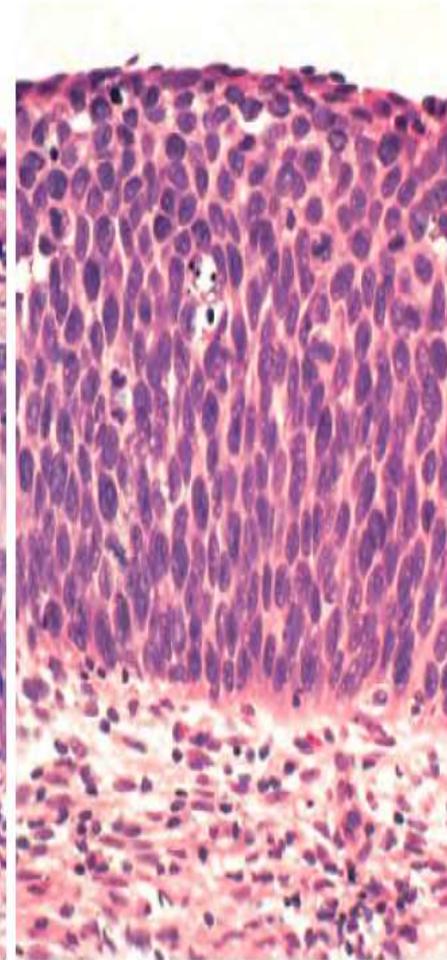
Normal



CIN I

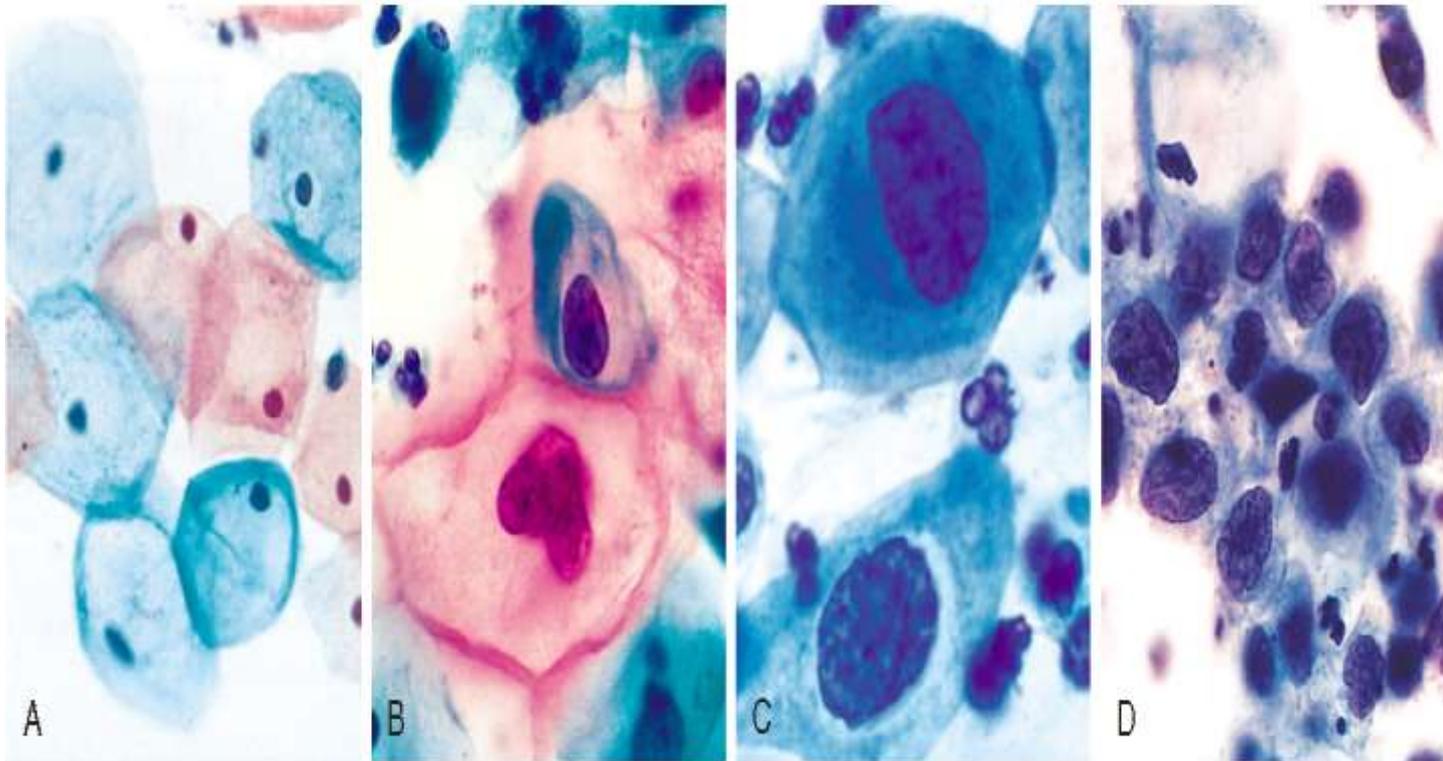


CIN II



CIN III

**CIN → DYSPLASIA:** NUCLEAR ENLARGEMENT, HYPERCHROMASIA (DARKER), COARSE CHROMATIN, & VARIATION IN NUCLEAR SIZE & SHAPE

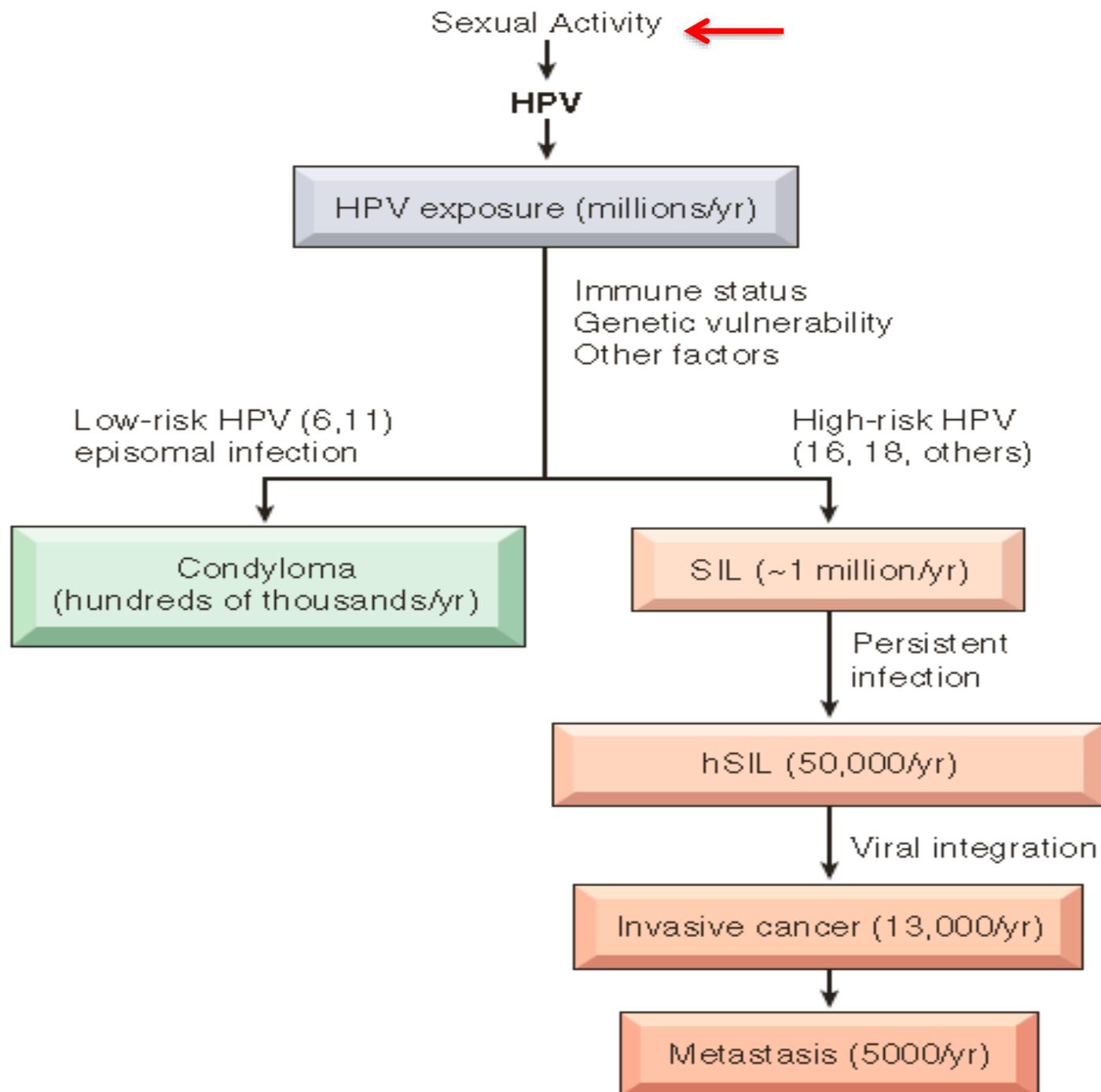


# CIN → SIL (SQUAMOUS INTRAEPITHELIAL LESION)

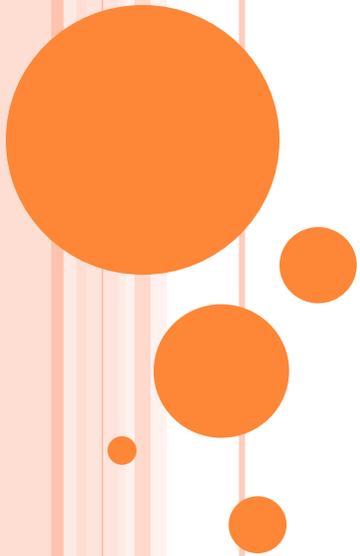
Dysplasia/Carcinoma in Situ	Cervical Intraepithelial Neoplasia (CIN)	Squamous Intraepithelial Lesion (SIL), Current Classification
Mild dysplasia	CIN I	Low-grade SIL (LSIL)
Moderate dysplasia	CIN II	High-grade SIL (HSIL)
Severe dysplasia	CIN III	High-grade SIL (HSIL)
Carcinoma in situ	CIN III	High-grade SIL (HSIL)

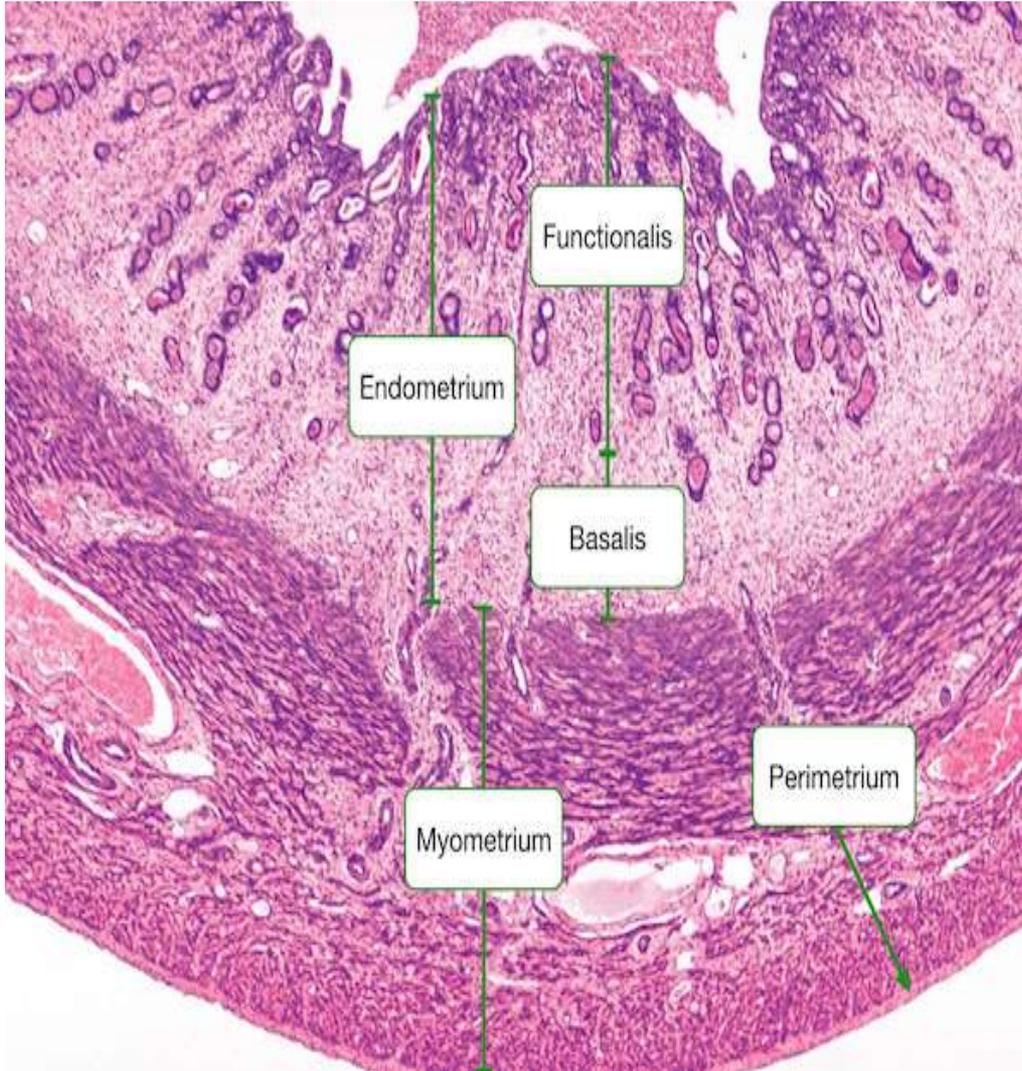
CIN, Cervical intraepithelial neoplasia; SIL, squamous intraepithelial lesion.





# BODY OF UTERUS



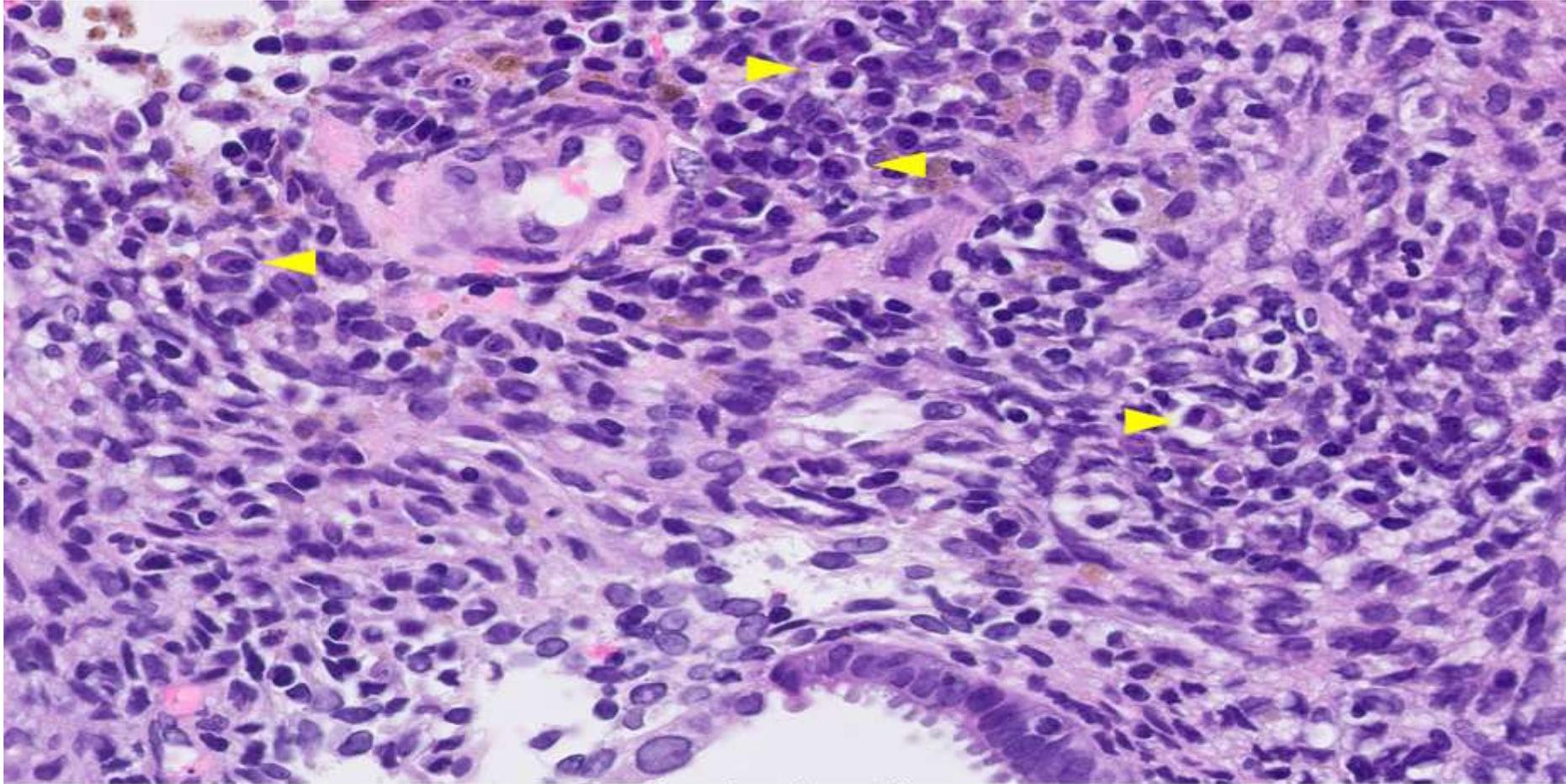


# UTERUS

Uterine body (corpus) is composed of:  
**endometrium**, consisting of glands & stroma. + **myometrium**, made up of smooth muscle.

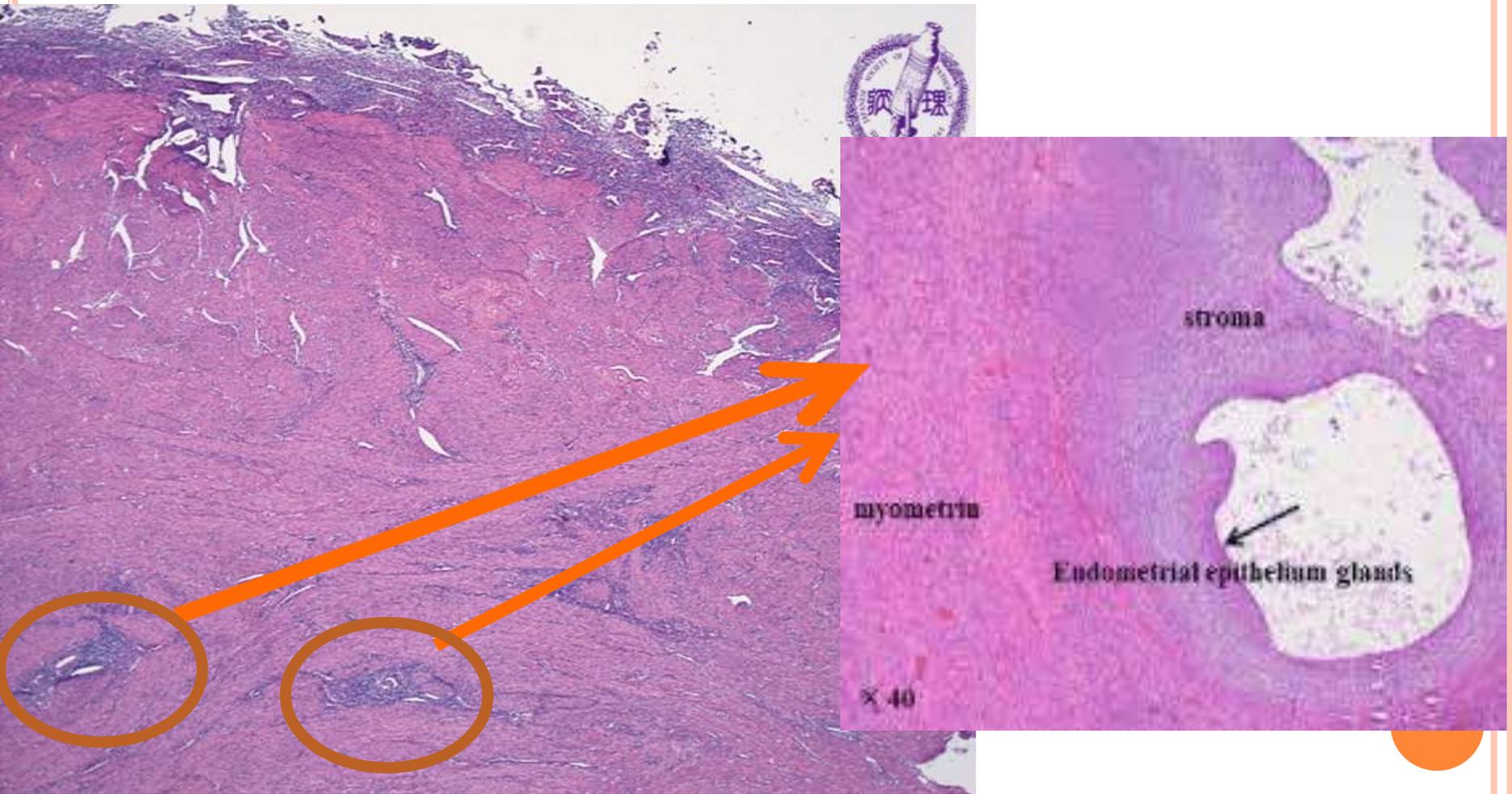


# ENDOMETRITIS



Chronic endometritis:  
presence of plasma cells (abundant here, *arrows*) is a major diagnostic feature (H&E,  $\times 20$ )

# UTERINE PATHOLOGY - ADENOMYOSIS



# ENDOMETRIOSIS - MICROSCOPICALLY

Diagnosis; **2 of 3 features:**

endometrial glands,  
endometrial stroma,  
or hemosiderin  
pigment.



Normal Colon



# ENDOMETRIOSIS - GROSS

Ovarian endometriosis: ovary + a large endometriotic cyst with degenerated blood (“chocolate cyst”).

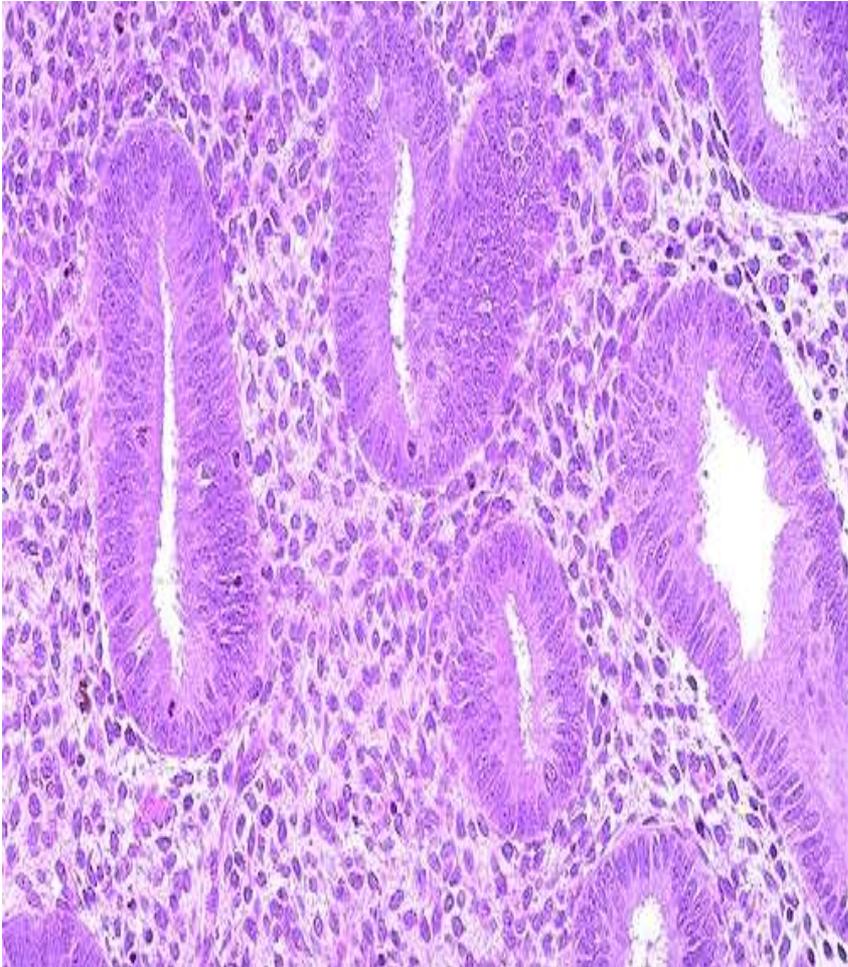


# UTERINE PATHOLOGY - ENDOMETRIAL HYPERPLASIA

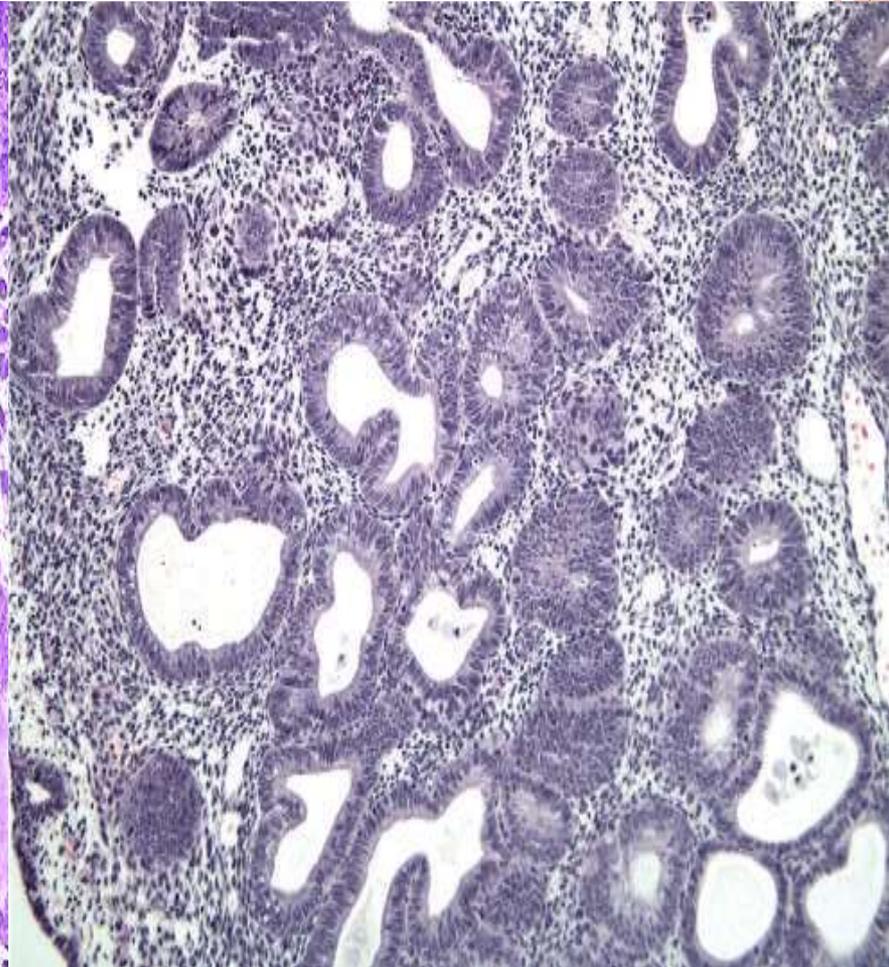
- **Pathogenesis:** prolonged or marked excess of estrogen relative to progestin → exaggerated proliferation.
- An important **precursor** of endometrial carcinoma.
- Two categories based on the presence of cytologic atypia:
  1. Hyperplasia without atypia; low risk for progression to endometrial Ca.
  2. Hyperplasia with atypia(**endometrial intraepithelial neoplasia (EIN)**) higher risk for progression to endometrial Ca. → 20%.



# UTERUS- HYPERPLASIA W/O ATYPIA

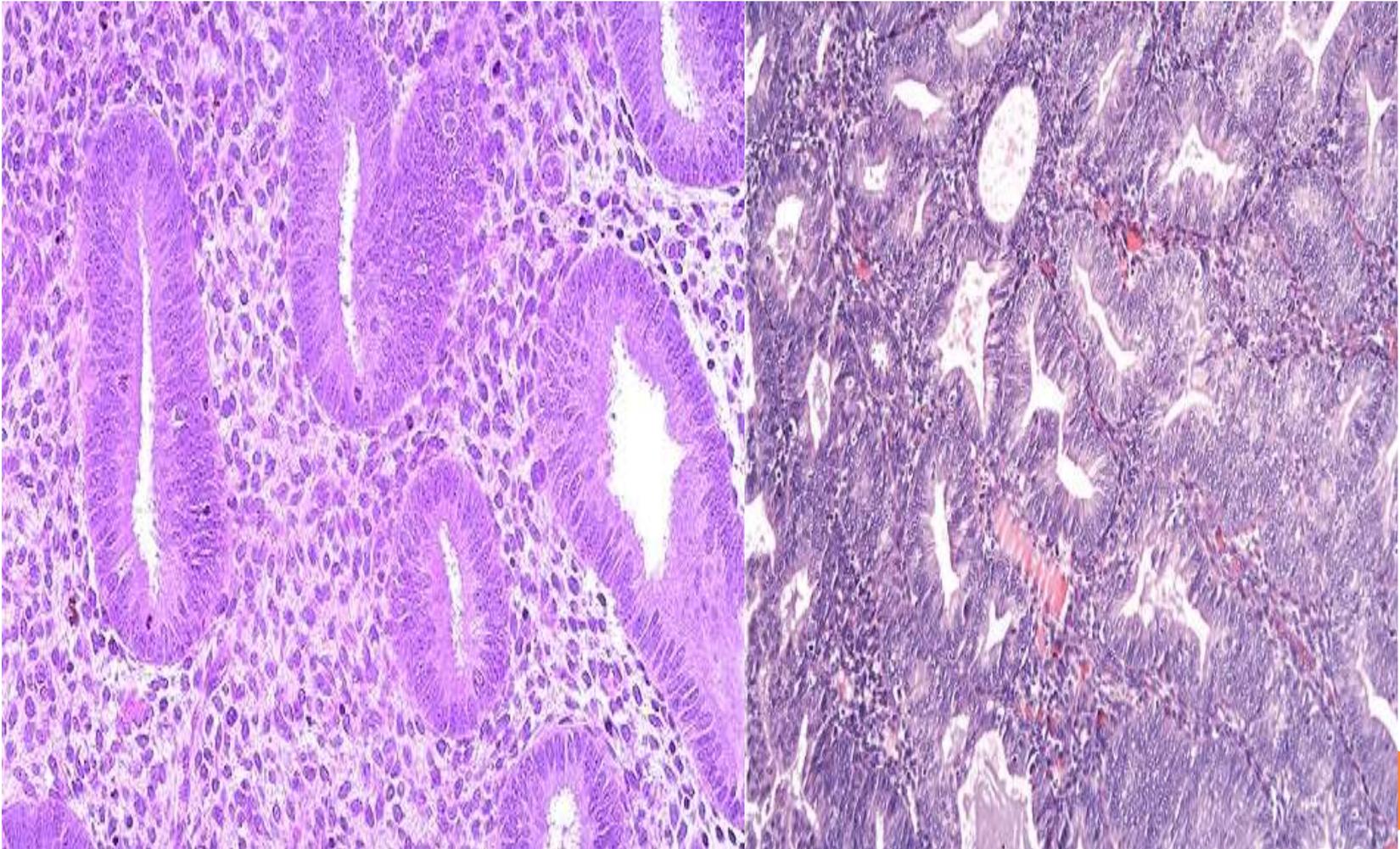


Normal



Hyperplasia without atypia

# UTERUS- HYPERPLASIA WITH ATYPIA



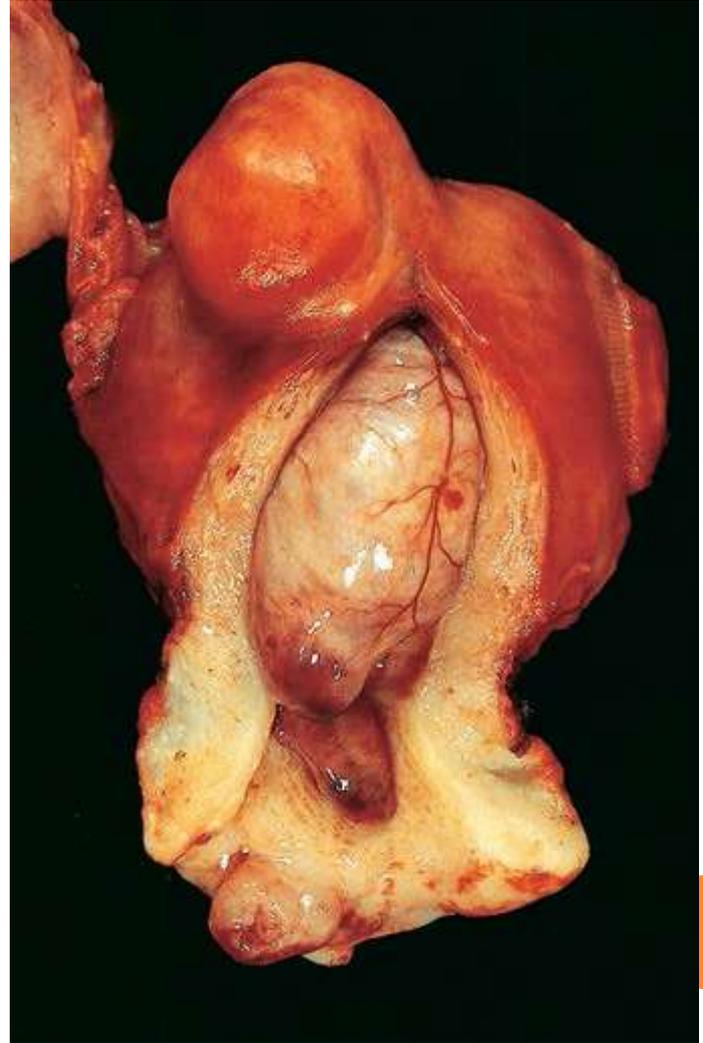
**Normal**

**Hyperplasia with atypia**

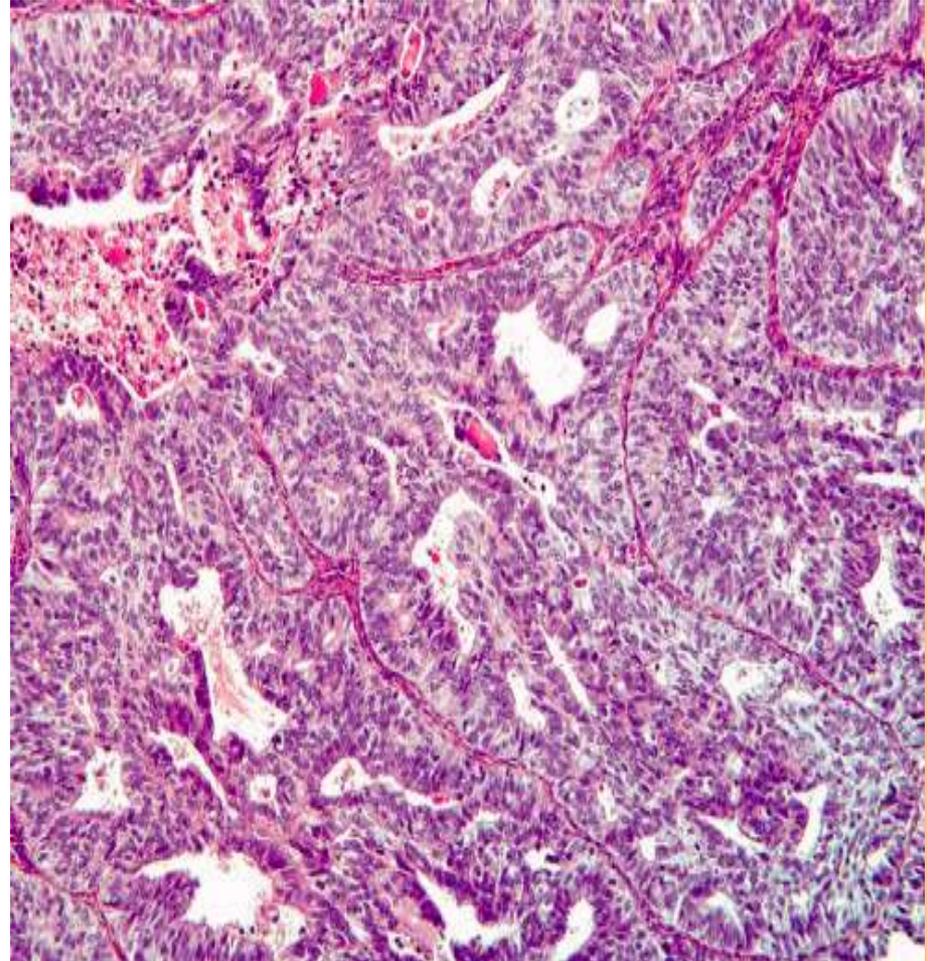
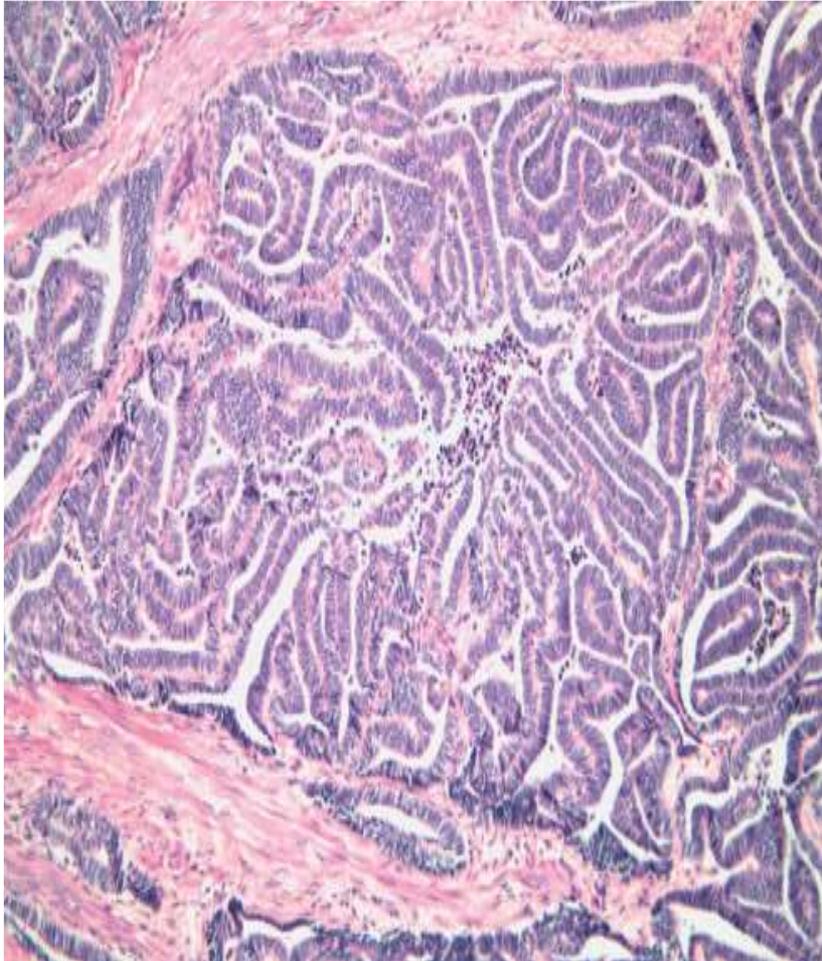
# TUMORS OF THE ENDOMETRIUM



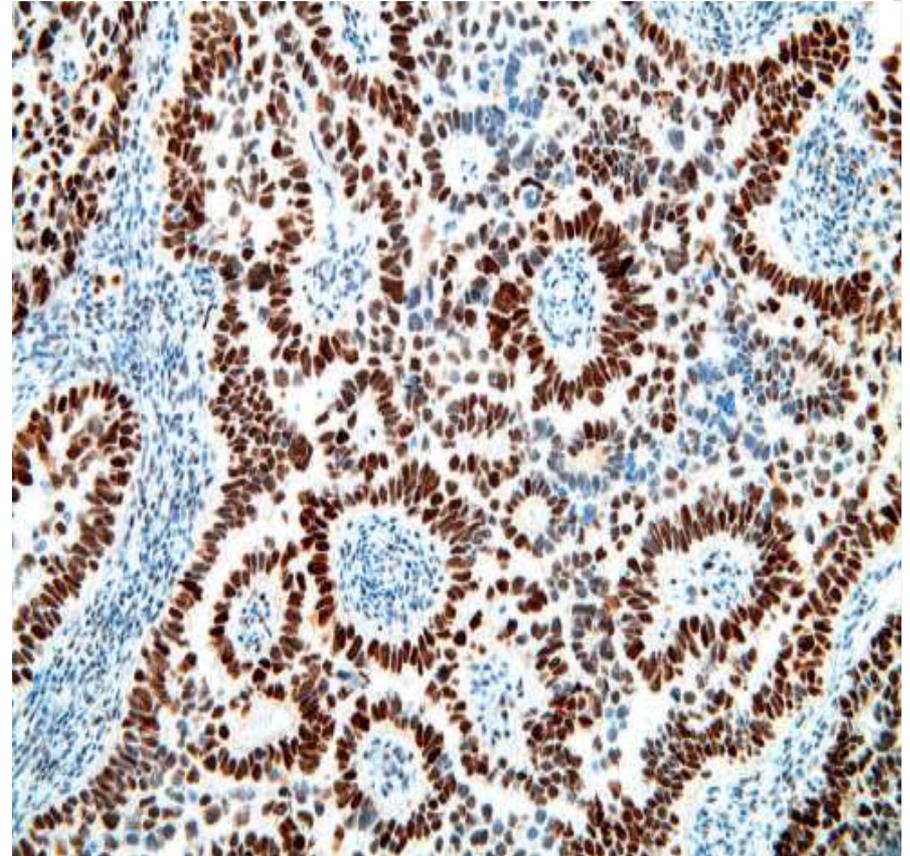
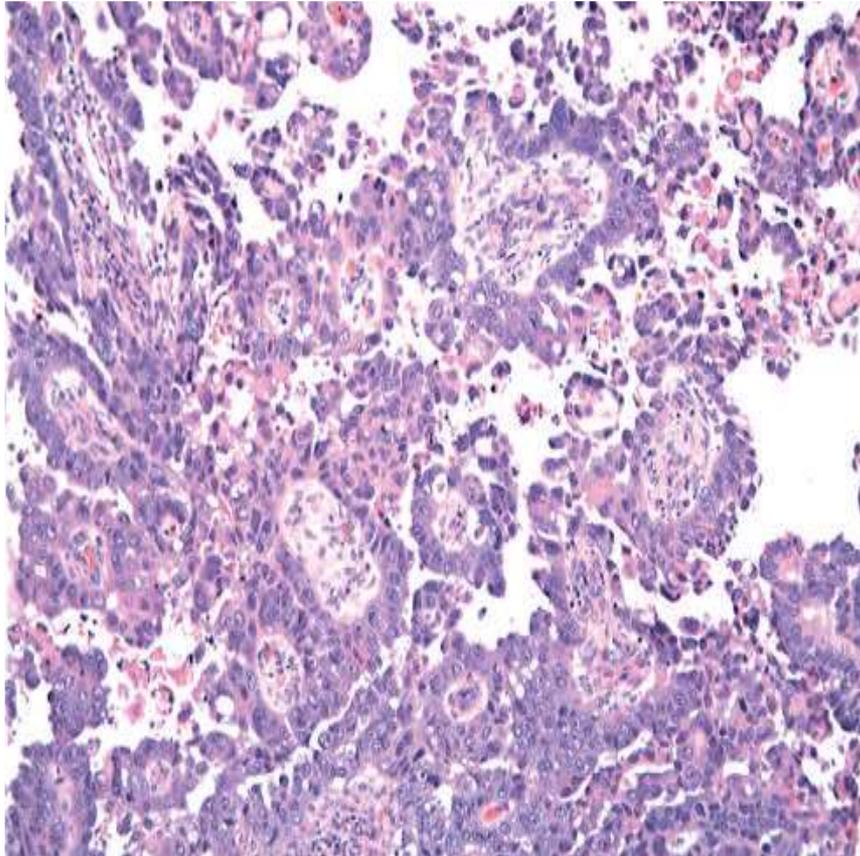
# ENDOMETRIAL POLYP



# TUMORS OF ENDOMETRIUM - ENDOMETRIOID CARCINOMA



# TUMORS OF ENDOMETRIUM - SEROUS CARCINOMAS



Microscopic: typically grow in small papillae with marked cytologic atypia. Immunohistochemical staining shows accumulation of p53, a finding associated with TP53 mutation.

# TUMORS OF THE MYOMETRIUM



# TUMORS OF MYOMETRIUM – LEIOMYOMAS (FIBROIDS)

**Location:** within the myometrium (intramural), beneath the endometrium (submucosal) or or the serosa (subserosal)



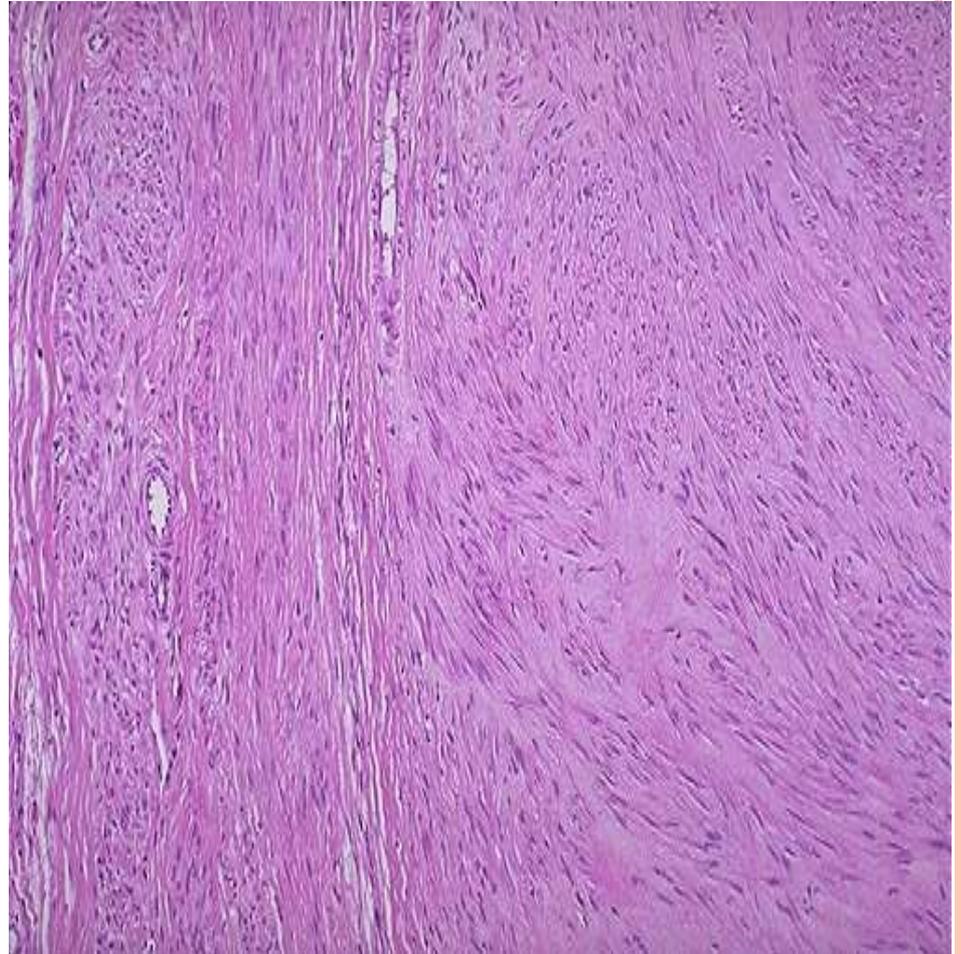
# TUMORS OF MYOMETRIUM – LEIOMYOMAS (FIBROIDS)

**Gross:** typically sharply circumscribed, firm gray white masses with a characteristic whorled cut surface, often occur as multiple tumors.



# TUMORS OF MYOMETRIUM – LEIOMYOMAS (FIBROIDS)

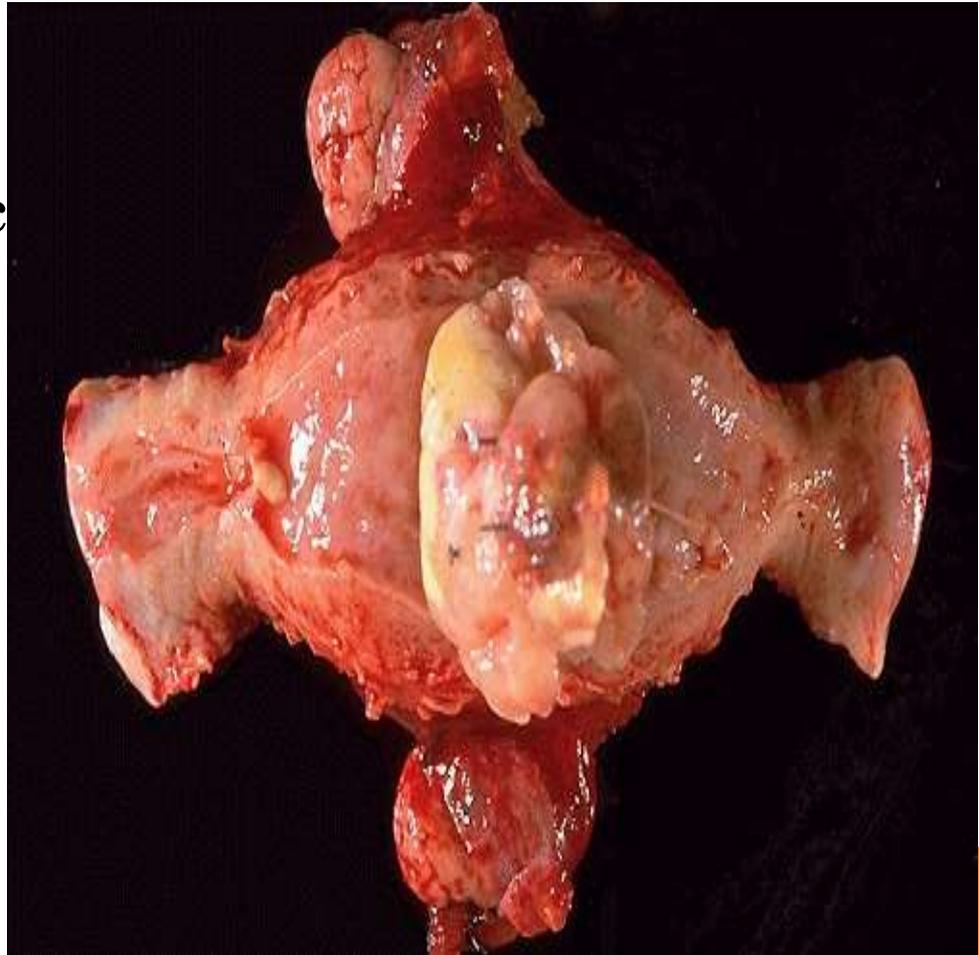
**Histologic examination,**  
**bundles of smooth**  
**muscle cells** mimicking  
the appearance of normal  
myometrium



# TUMORS OF MYOMETRIUM – LEIOMYOSARCOMA

**Gross:** soft,  
hemorrhagic, necrotic  
masses.

Irregular borders.



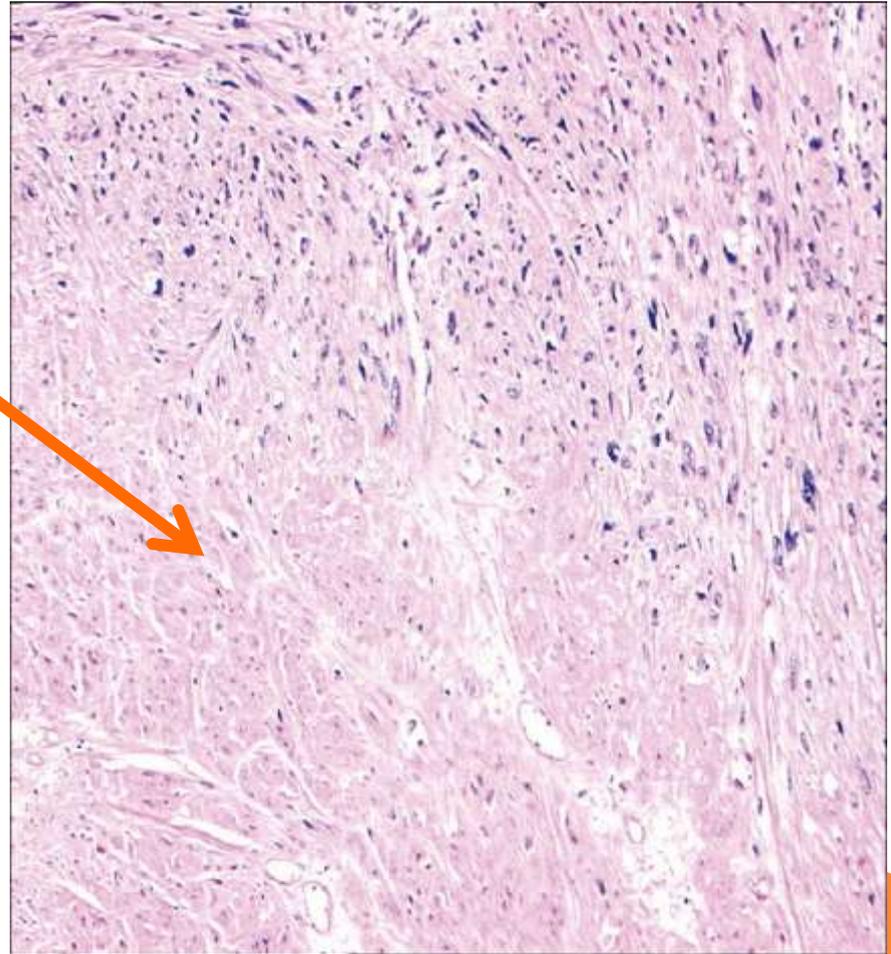
# TUMORS OF MYOMETRIUM – LEIOMYOSARCOMA

## Microscopic:

Diagnostic features of leiomyosarcoma;

(1) **tumor necrosis**,  
(2) cytologic atypia, and  
(3) mitotic activity.

Assessment of all three is necessary to make a diagnosis.



# TUMORS OF MYOMETRIUM – LEIOMYOSARCOMA

## Microscopic:

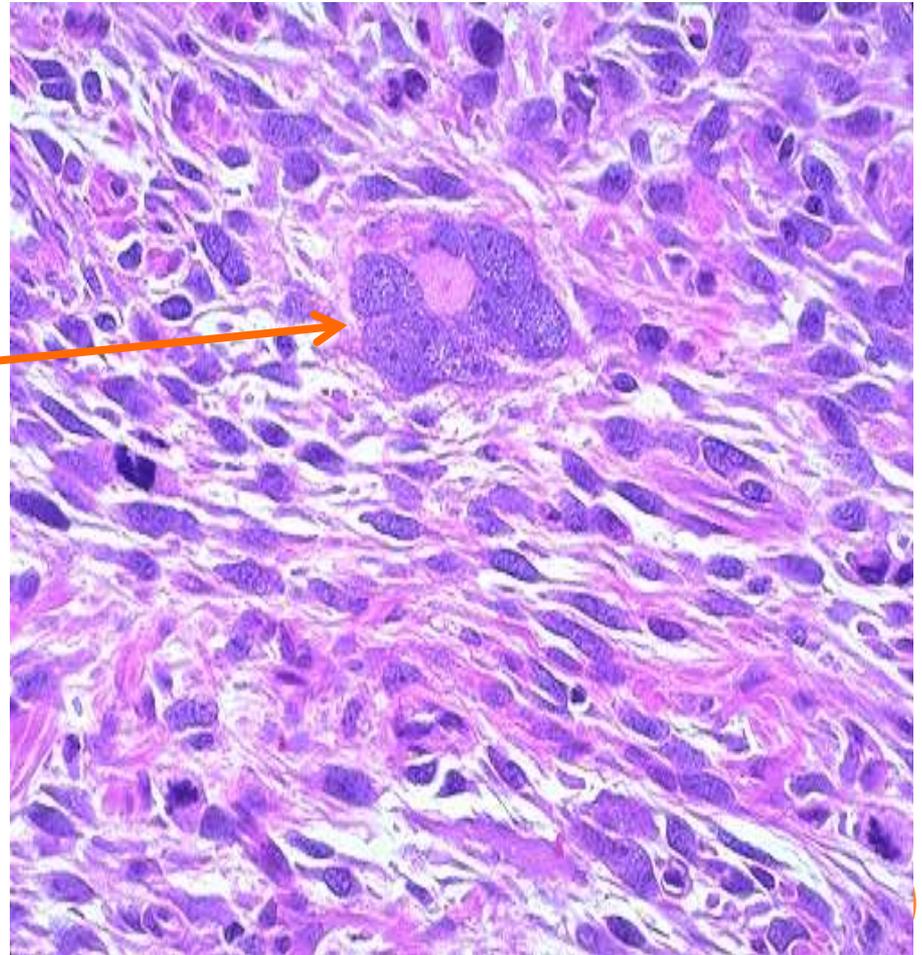
Diagnostic features of leiomyosarcoma;

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and (3) mitotic activity.

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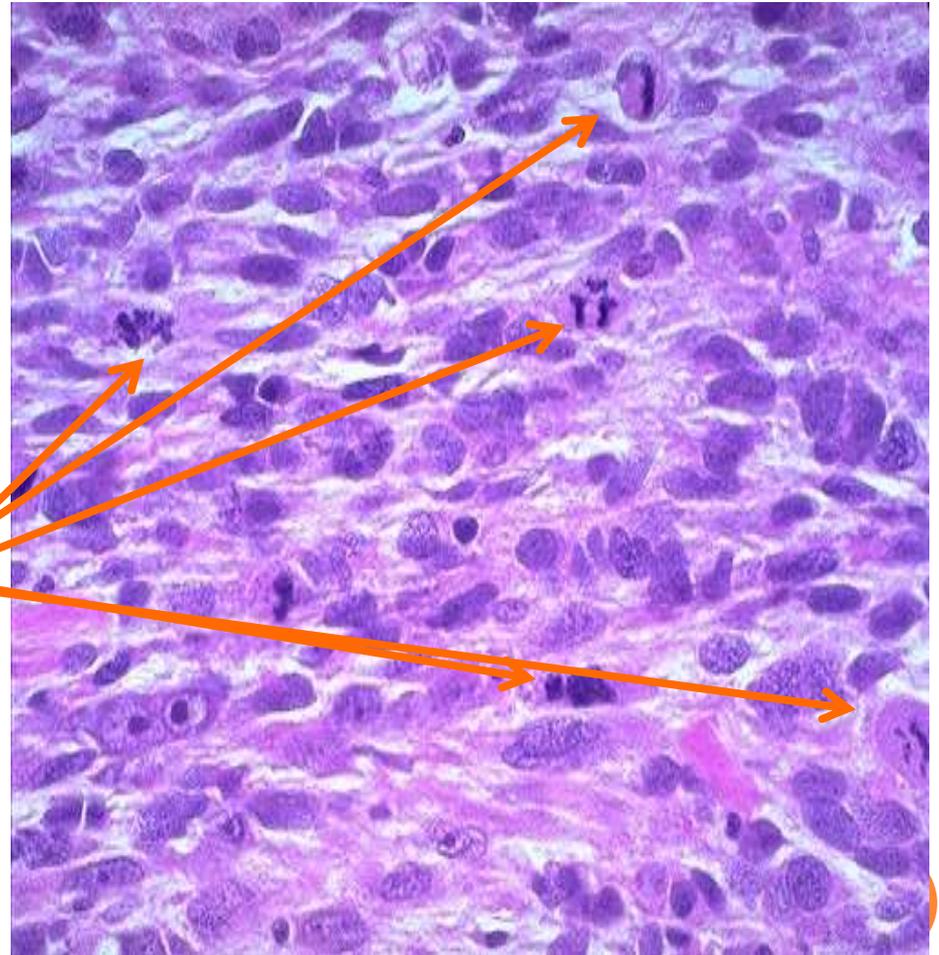
# TUMORS OF MYOMETRIUM – LEIOMYOSARCOMA

## Microscopic:

Diagnostic features of leiomyosarcoma;

- (1) tumor necrosis,
- (2) cytologic atypia, and
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Assessment of all three is necessary to make a diagnosis.



# OVARIES

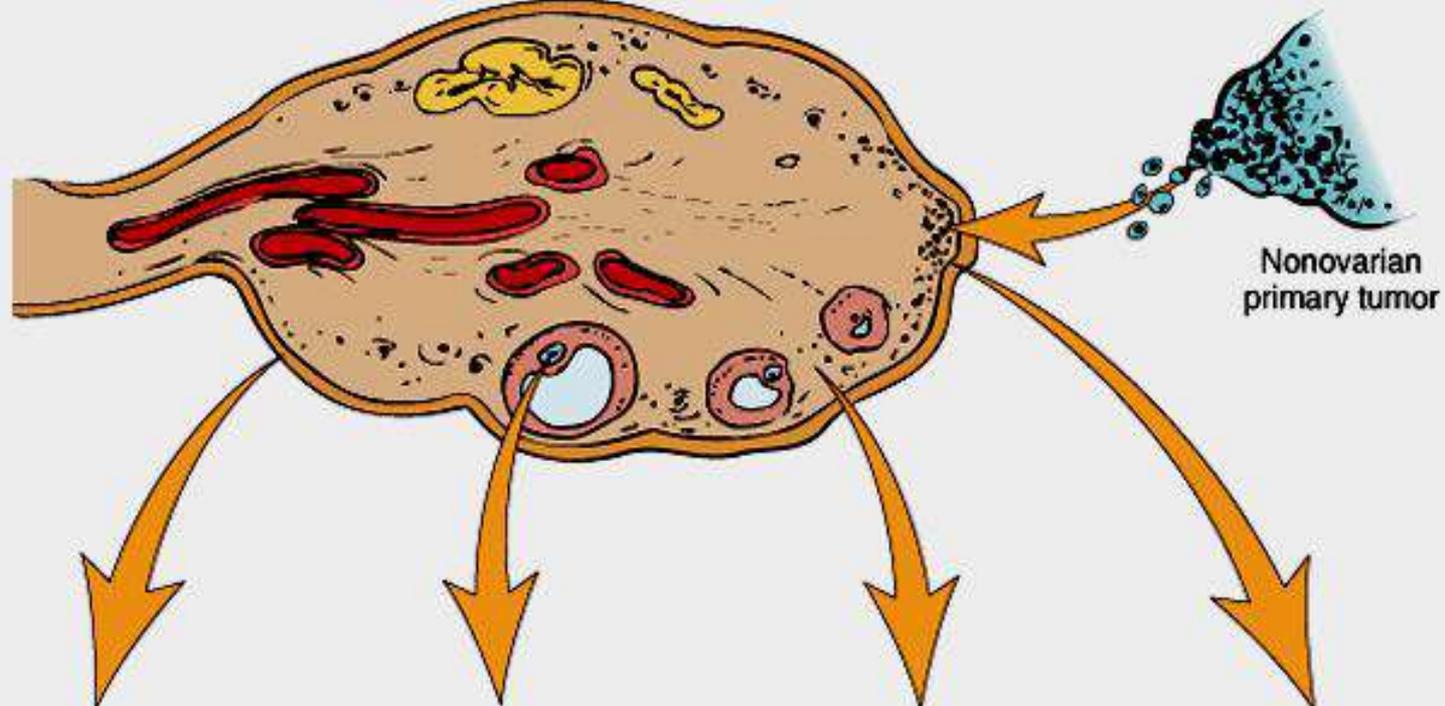


# OVARIES- POLYCYSTIC OVARIAN SYNDROME



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ORIGIN	SURFACE EPITHELIAL CELLS (Surface epithelial-stromal cell tumors)	GERM CELL	SEX CORD-STROMA	METASTASIS TO OVARIES
Overall frequency	65%–70%	15%–20%	5%–10%	5%
Proportion of malignant ovarian tumors	90%	3%–5%	2%–3%	5%
Age group affected	20+ years	0–25+ years	All ages	Variable
Types	<ul style="list-style-type: none"> <li>• Serous tumor</li> <li>• Mucinous tumor</li> <li>• Endometrioid tumor</li> <li>• Clear cell tumor</li> <li>• Brenner tumor</li> <li>• Cystadenofibroma</li> </ul>	<ul style="list-style-type: none"> <li>• Teratoma</li> <li>• Dysgerminoma</li> <li>• Endodermal sinus tumor</li> <li>• Choriocarcinoma</li> </ul>	<ul style="list-style-type: none"> <li>• Fibroma</li> <li>• Granulosa-theca cell tumor</li> <li>• Sertoli-Leydig cell tumor</li> </ul>	

# SEROUS TUMORS - BENIGN

## SEROUS TUMORS

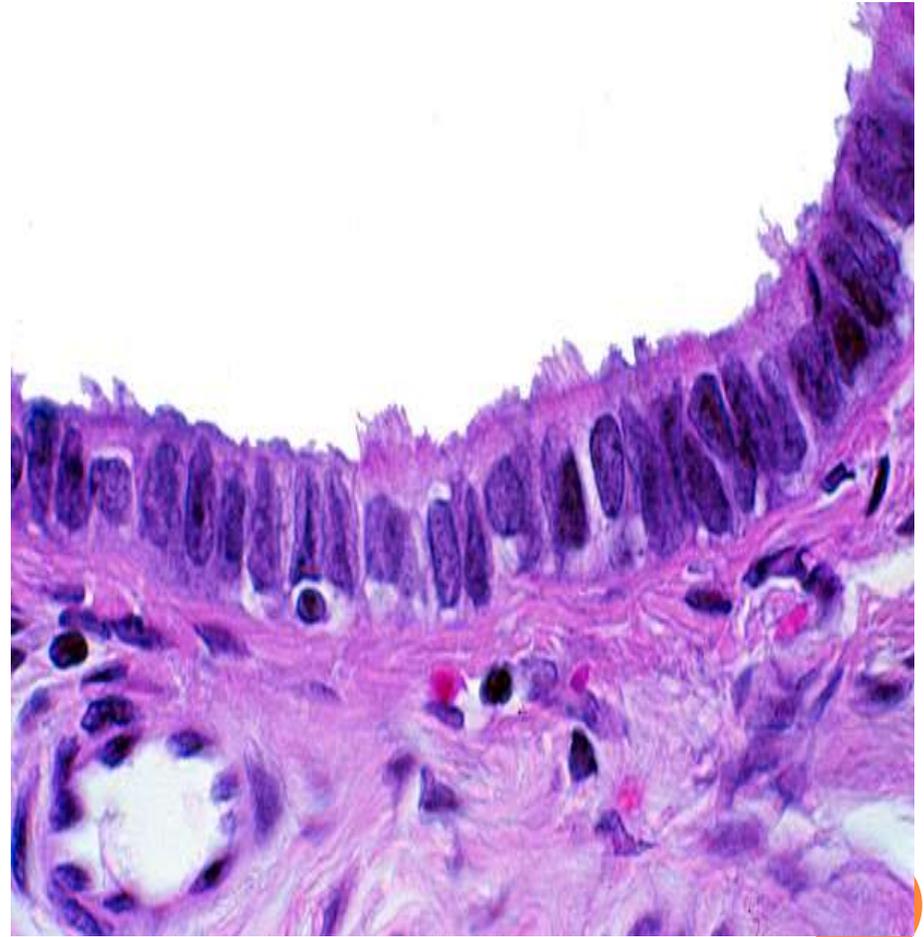
- Gross: Large & cystic ( up to 30 cm), filled with a clear serous fluid
- May be bilateral.
- Called serous cystadenoma



# SEROUS TUMORS - BENIGN

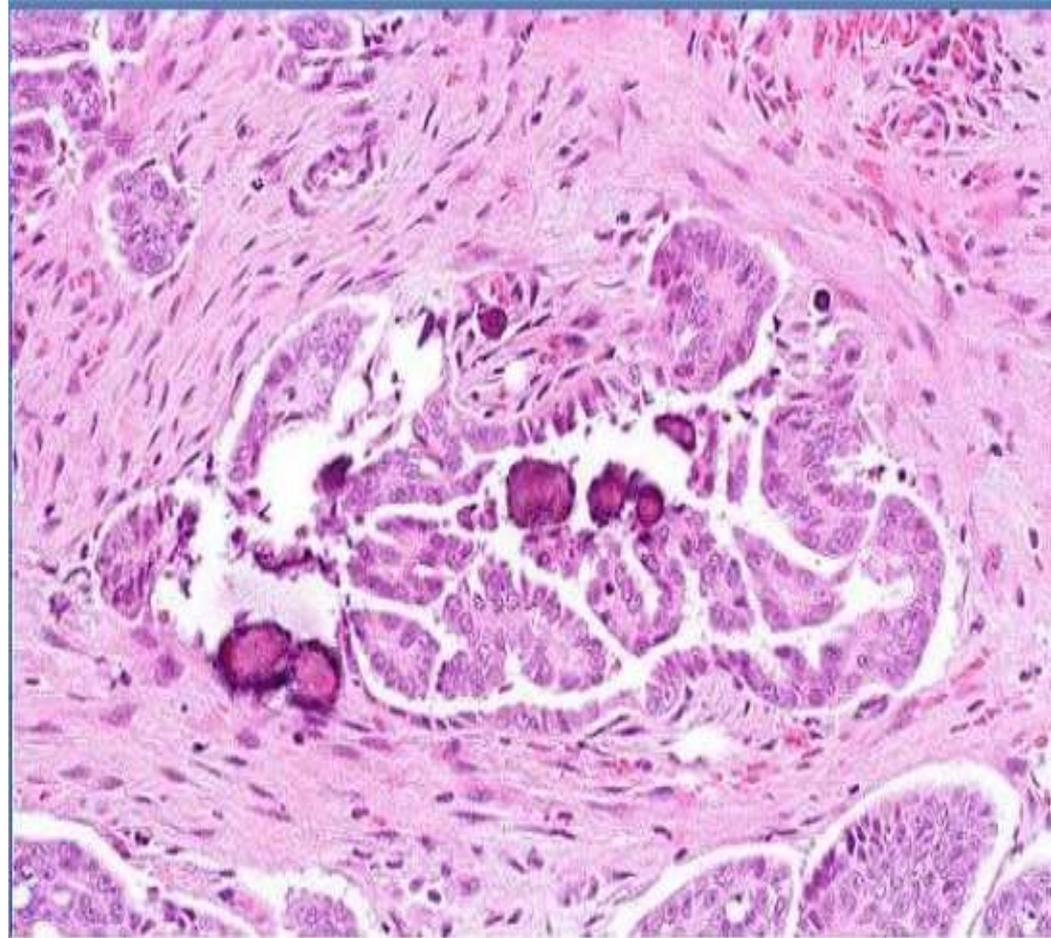
## SEROUS TUMORS

**Microscopy:** Single layer of columnar epithelium. Some cells are ciliated.



# SEROUS TUMORS - SEROUS TUMORS

Psammoma bodies (laminated calcified concretions) are common in tips of papillae of **all serous tumors**



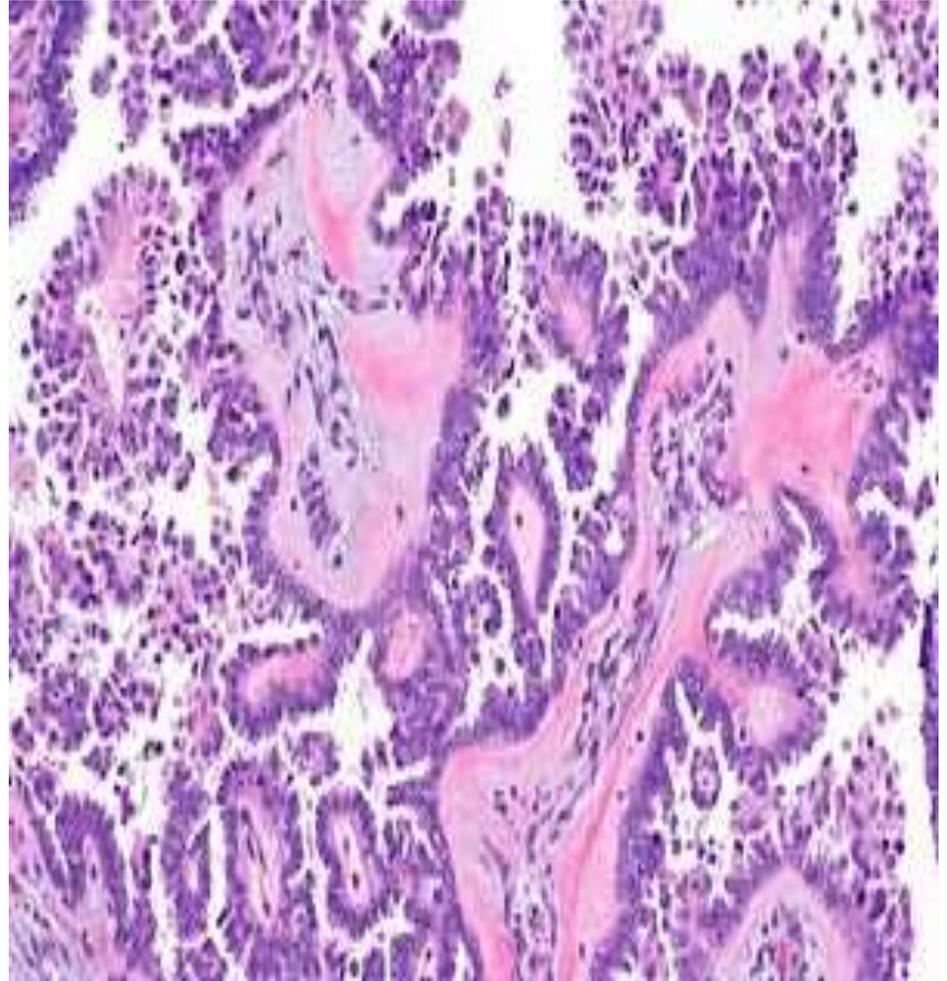
# SEROUS TUMORS - BORDERLINE SEROUS TUMORS

- complex architecture.  
(Protruding papillary projections)
- might be associated with  
peritoneal implants.



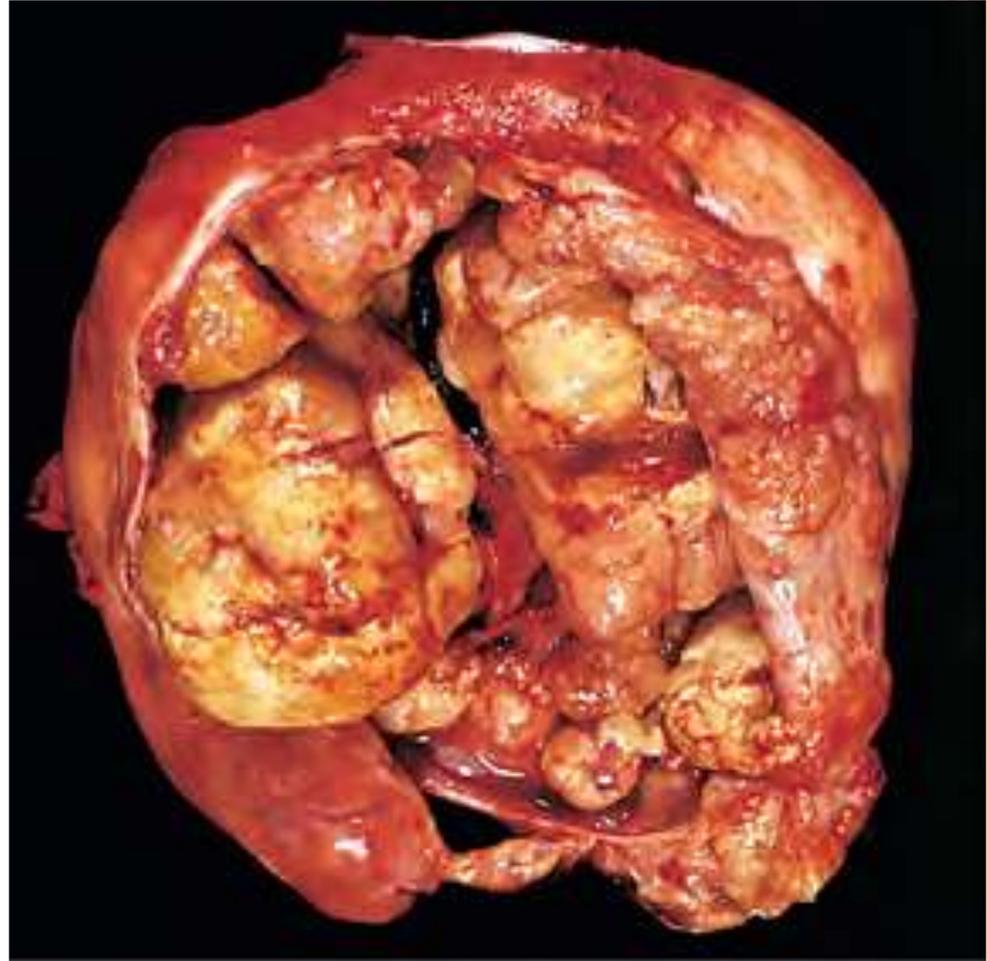
# SEROUS TUMORS - BORDERLINE SEROUS TUMORS

- complex architecture.
- mild cytologic atypia, but no stromal invasion.
- Prognosis *intermediate* between benign & malignant.



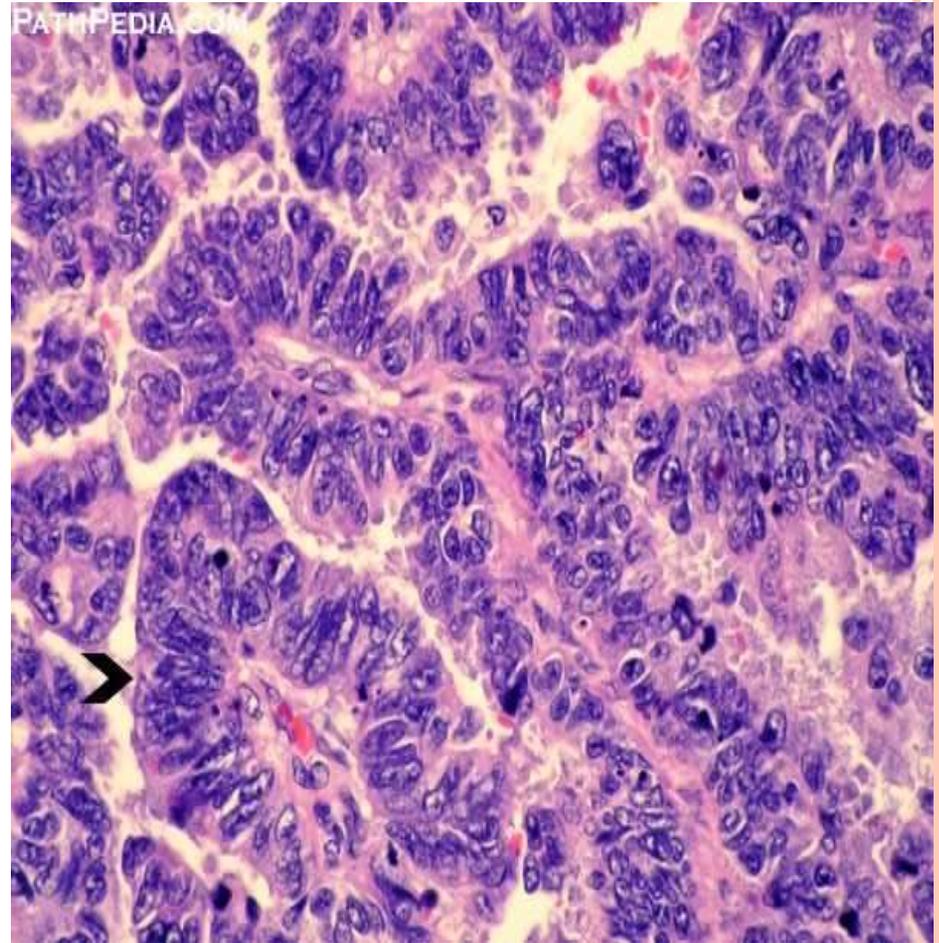
# SEROUS TUMORS - SEROUS CARCINOMA

- papillary formations are usually more complex
- tumor has **invaded the serosal surface**.
- prognosis **poor**, depends on stage at the time of diagnosis.

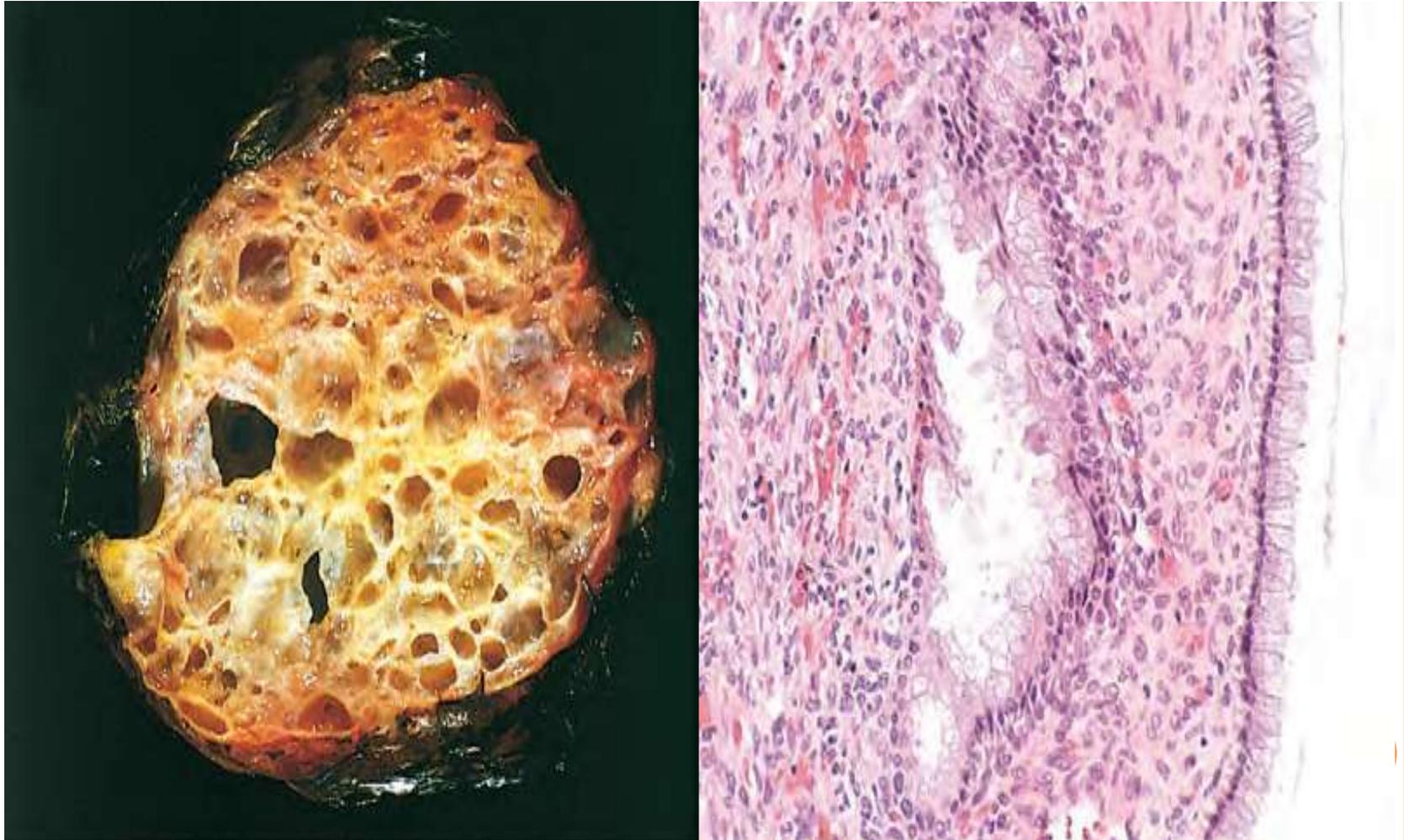


# SEROUS TUMORS - SEROUS CARCINOMA

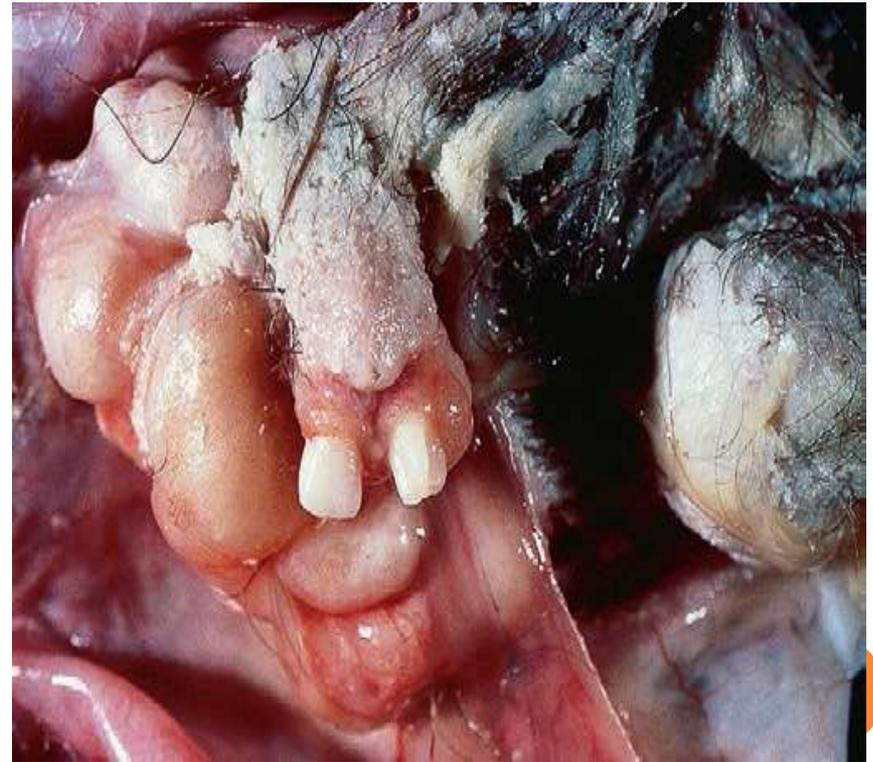
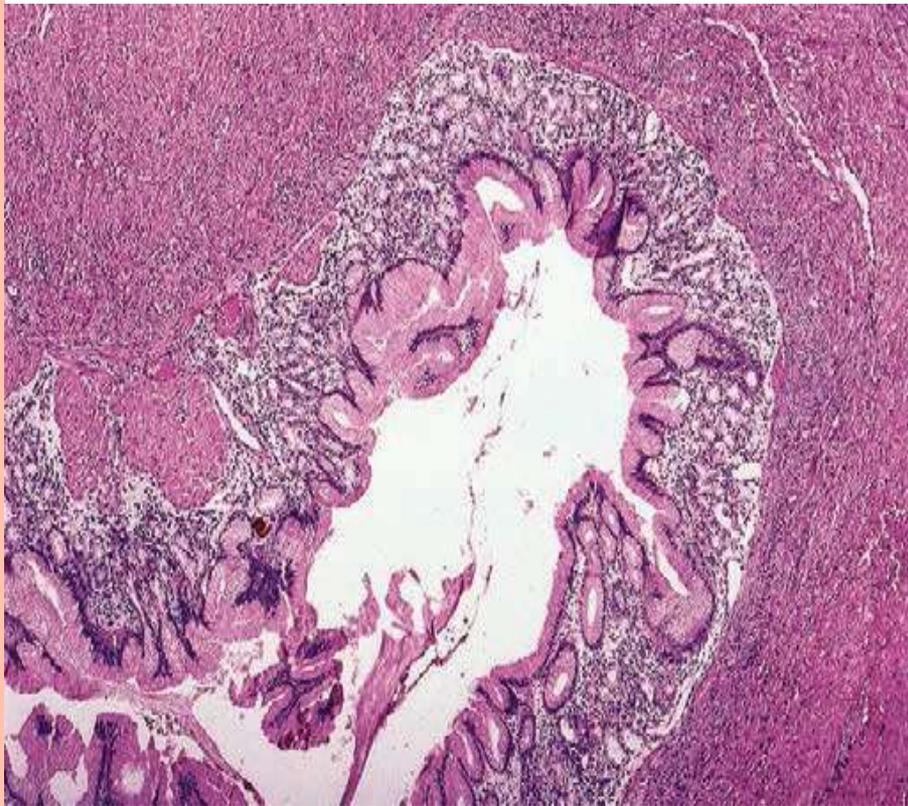
- complex papillary formations (multilayered)
- markedly cytological atypia
- By definition nests of malignant cells invade the stroma.



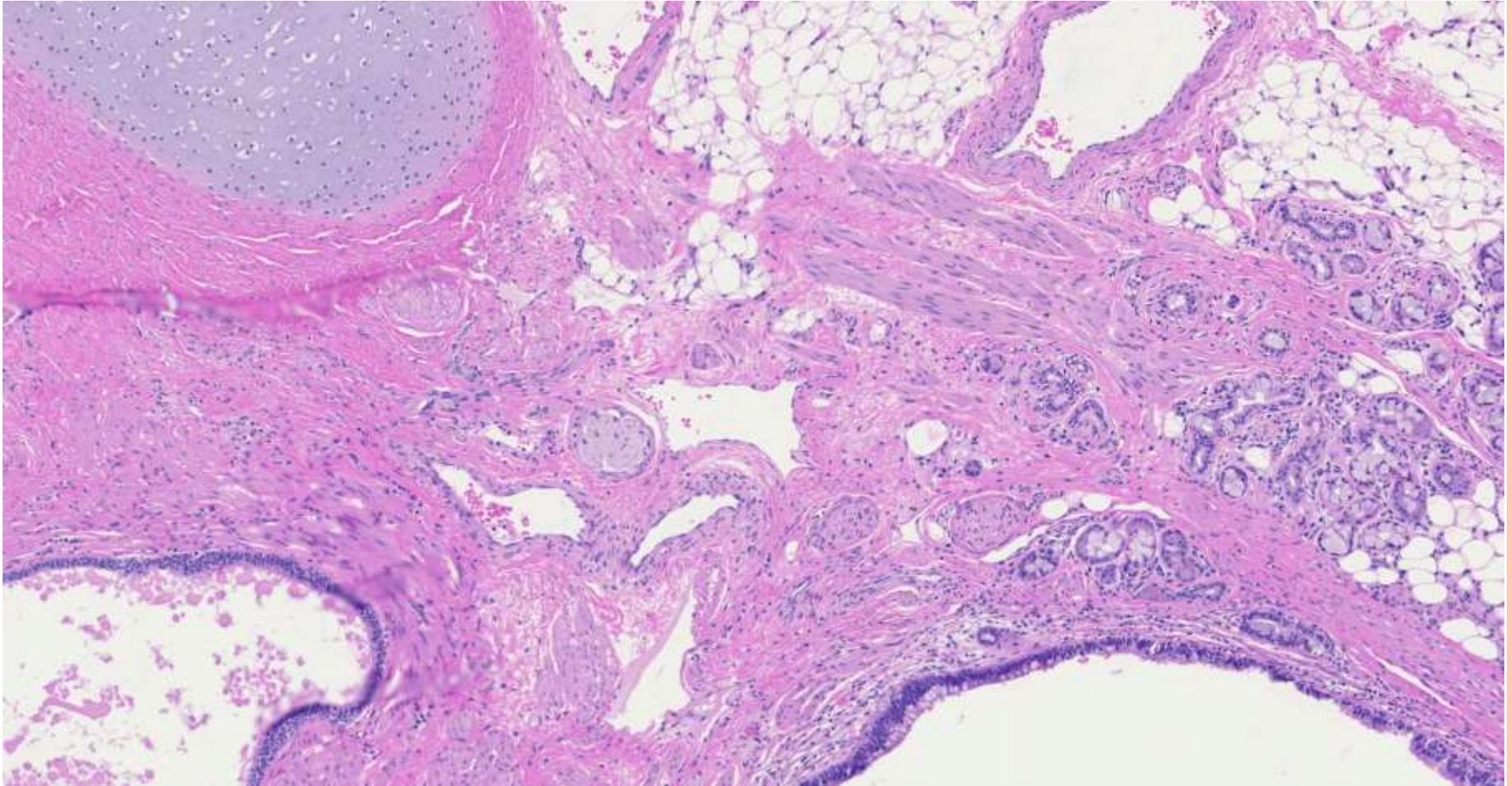
# OVARIES- MUCINOUS CYSTADENOMA



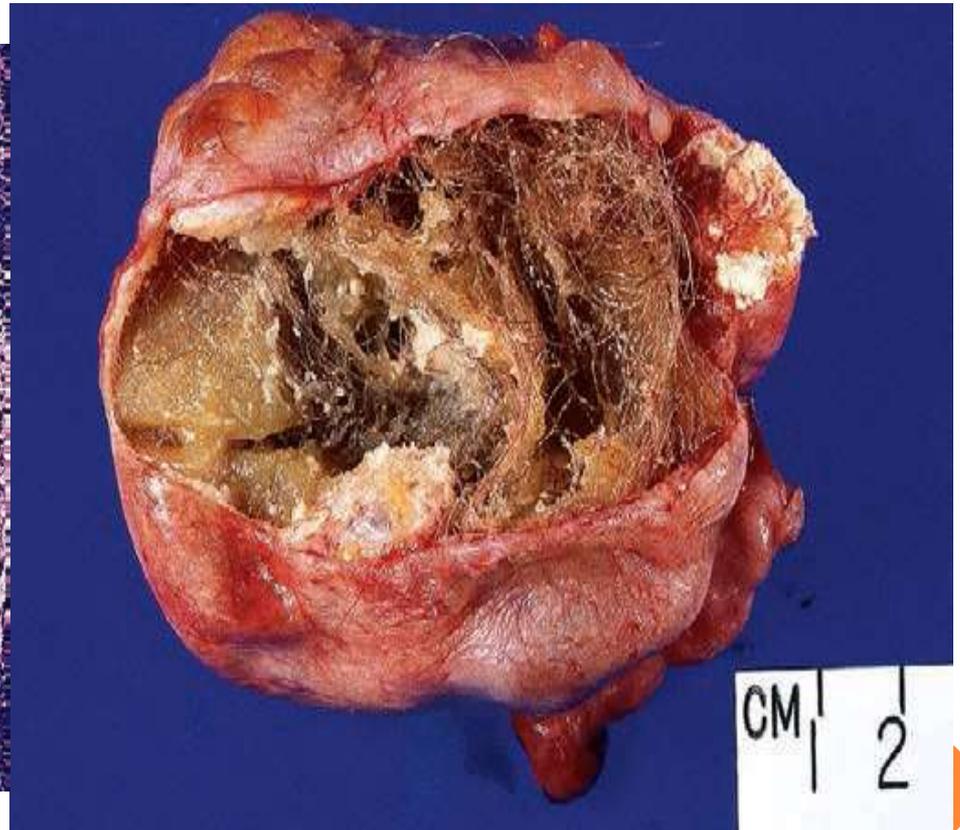
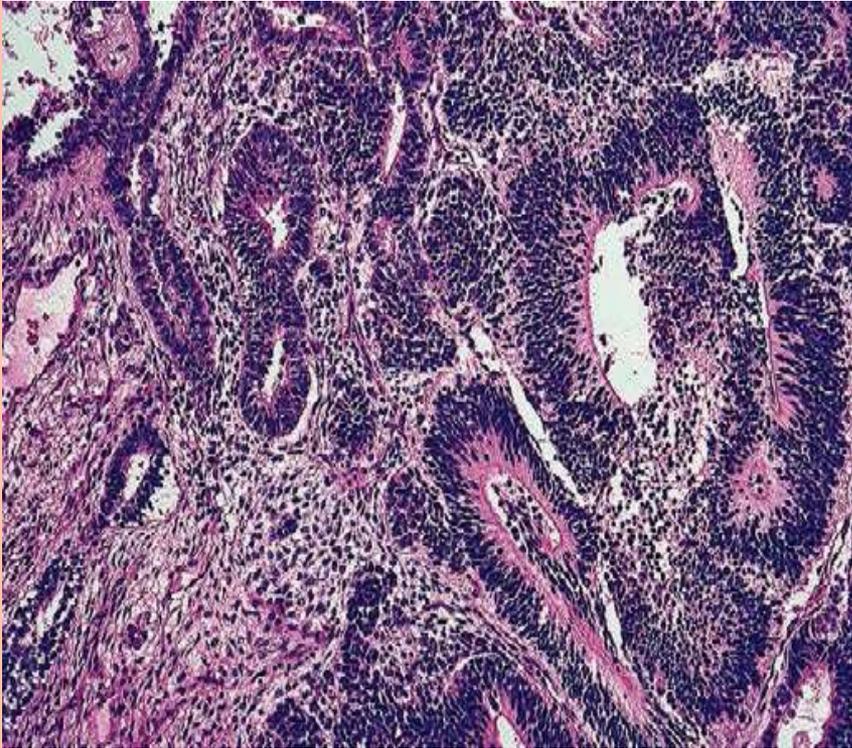
# BENIGN MATURE CYSTIC TERATOMAS



Mature cystic teratoma with multiple tissue types, including cartilage, endocervical epithelium, nerves, adipose and salivary gland



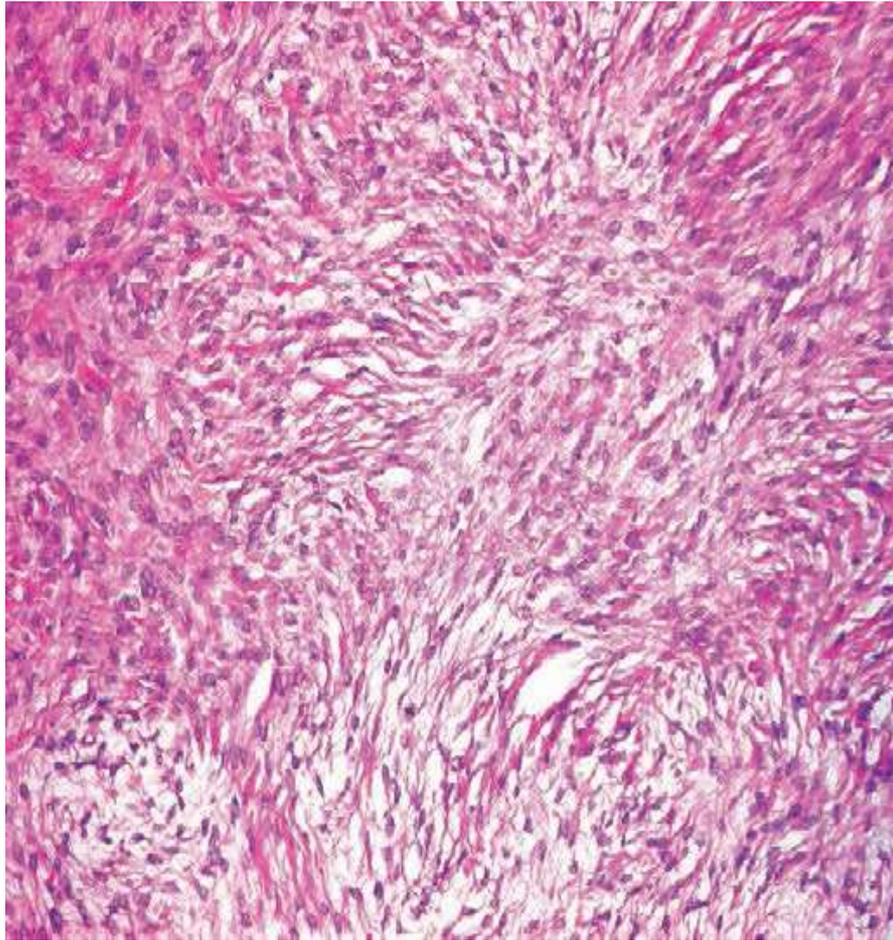
# IMMATURE MALIGNANT TERATOMA



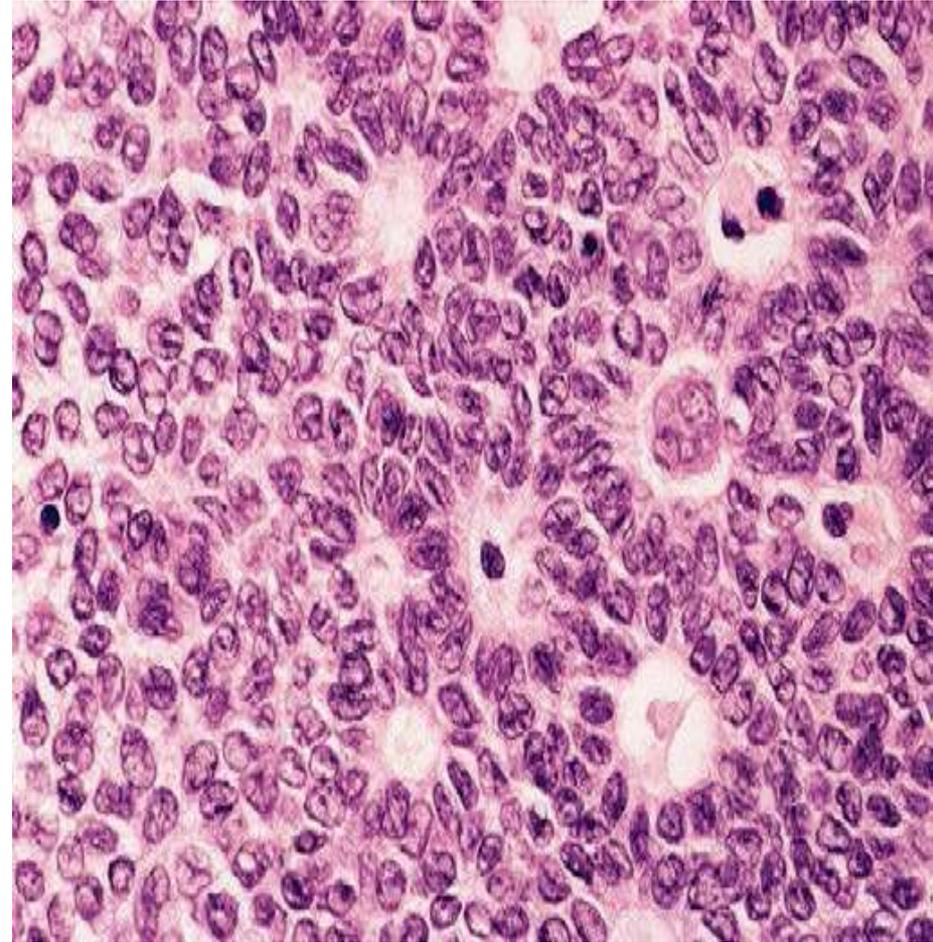
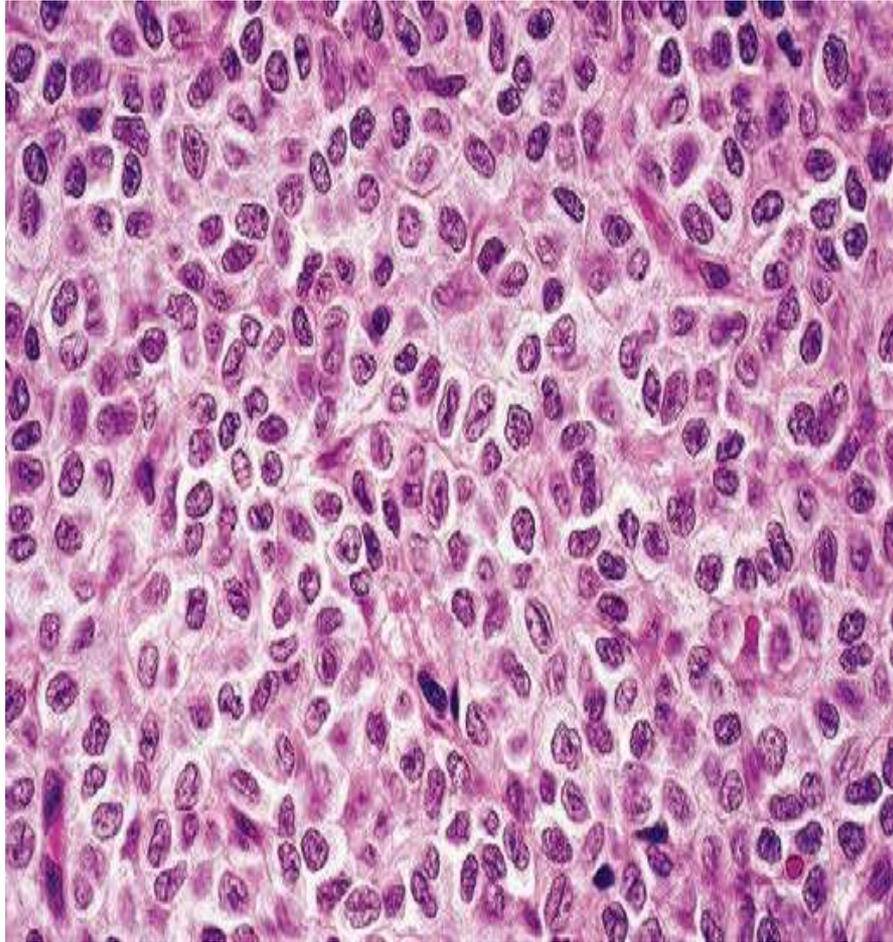
# OVARIES – SEX CORD TUMORS

Neoplasm	Peak Incidence	Usual Location	Morphologic Features	Behavior
<b>Sex Cord Tumors</b>				
Granulosa-theca cell	Most postmenopausal, but may occur at any age	Unilateral	<p>May be tiny or large, gray to yellow (with cystic spaces)</p> <p>Composed of mixture of cuboidal <u>granulosa cells</u> in cords, sheets, or strands and spindled or plump lipid-laden theca cells</p> <p>Granulosa elements may recapitulate ovarian follicle as <u>Call-Exner bodies</u></p>	<p>May elaborate large amounts of ★ estrogen (from thecal elements) and so may promote endometrial or breast carcinoma</p> <p>Granulosa element may be malignant (5% to 25%)</p>
Thecoma-fibroma	Any age	Unilateral	<p>Solid gray fibrous cells to yellow (lipid-laden) plump thecal cells</p>	<p>Most hormonally <u>inactive</u></p> <p>A few elaborate estrogens</p> <p>About 40%, for obscure reasons, produce ascites and hydrothorax</p> <p>★ (Meigs syndrome)</p> <p>Rarely malignant</p>
Sertoli-Leydig cell	All ages	Unilateral	<p>Usually small, gray to yellow-brown, and solid</p> <p>Recapitulates development of testis with tubules or cords and plump pink Sertoli cells</p>	<p>Many masculinizing or defeminizing</p> <p>Rarely malignant</p>

# OVARIAN FIBROMA

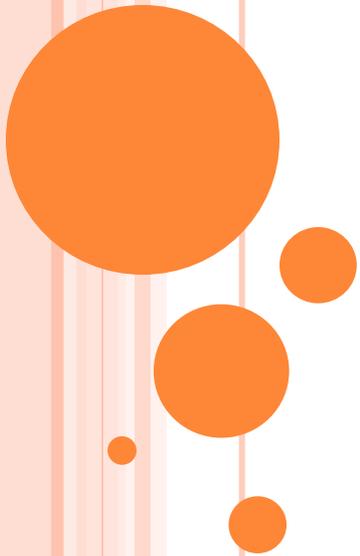


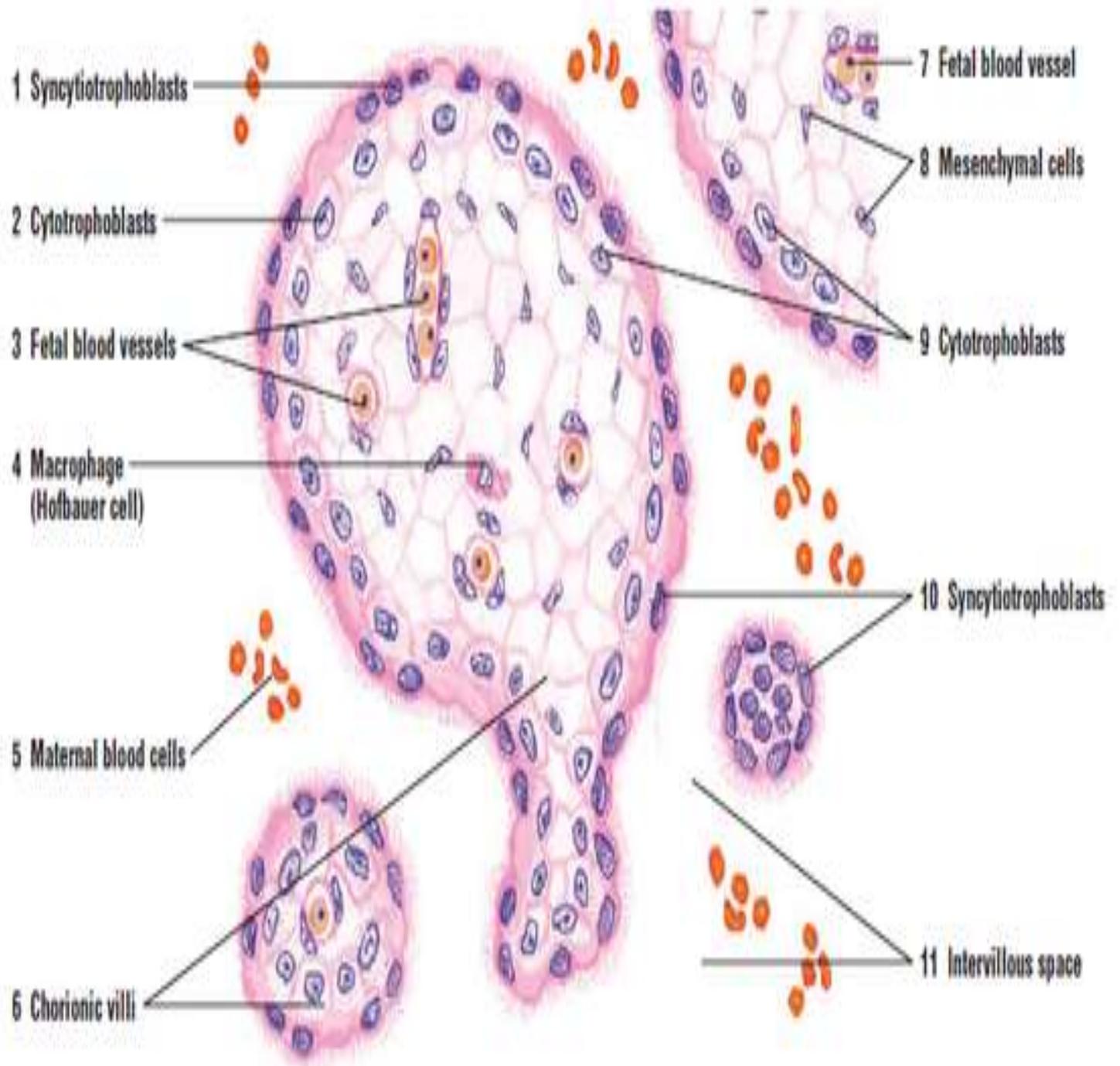
# OVARIES-GRANULOSA CELL TUMOR.

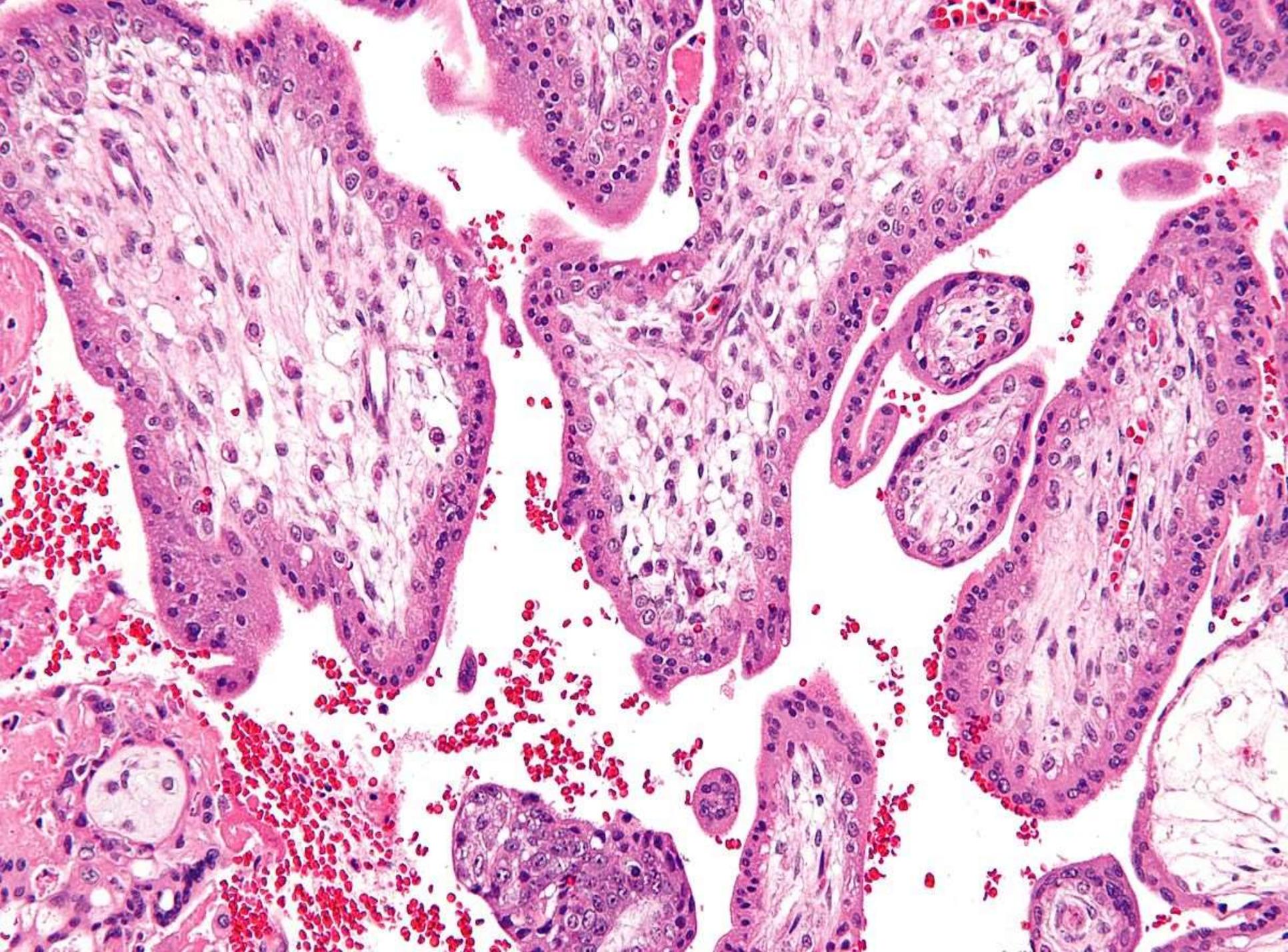


# **FEMALE GENITAL SYSTEM, LECTURE 5**

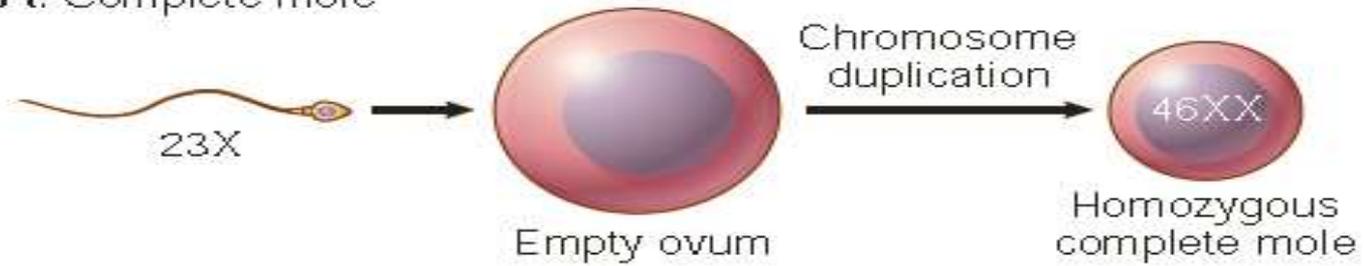
## **GESTATIONAL TROPHOBLASTIC DISEASE**



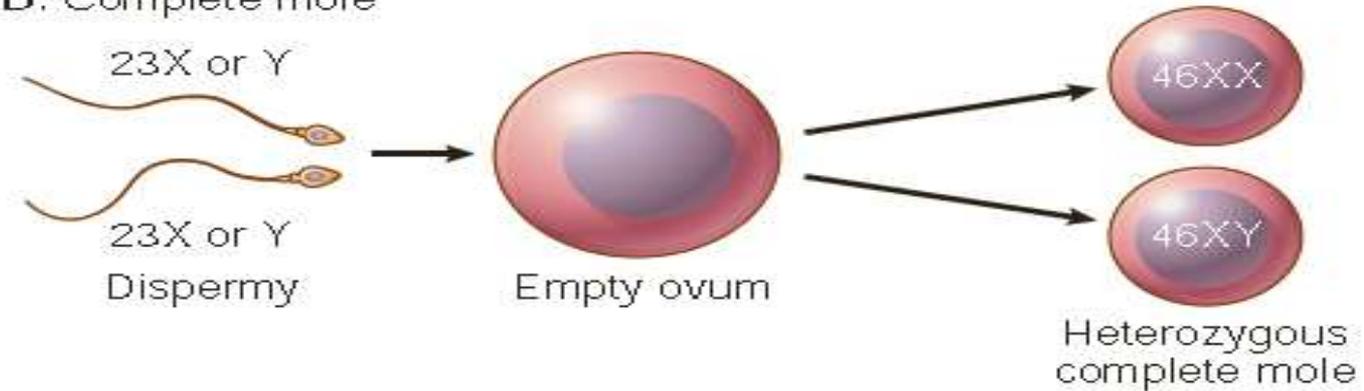
PLACENTAL DISC  
HISTOLOGY



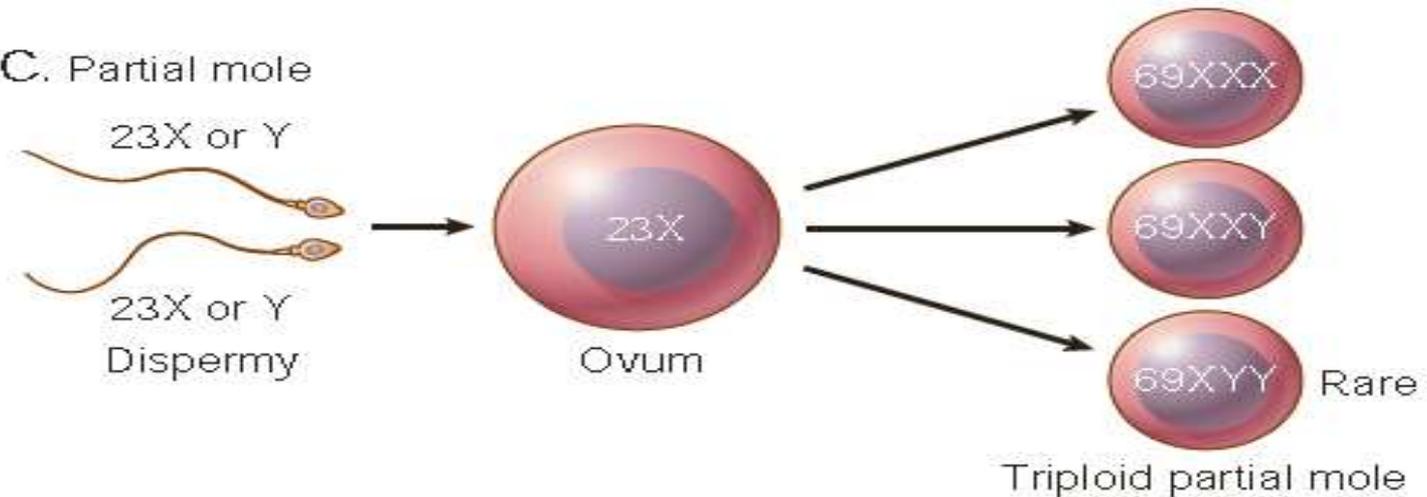
**A. Complete mole**



**B. Complete mole**

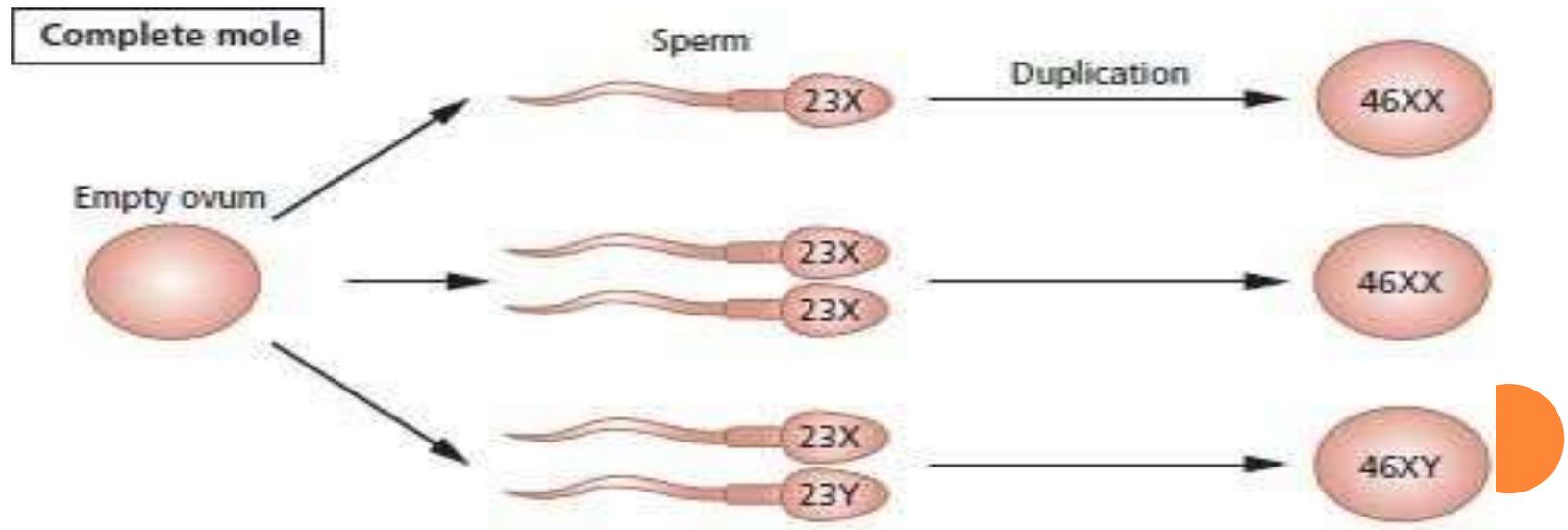


**C. Partial mole**



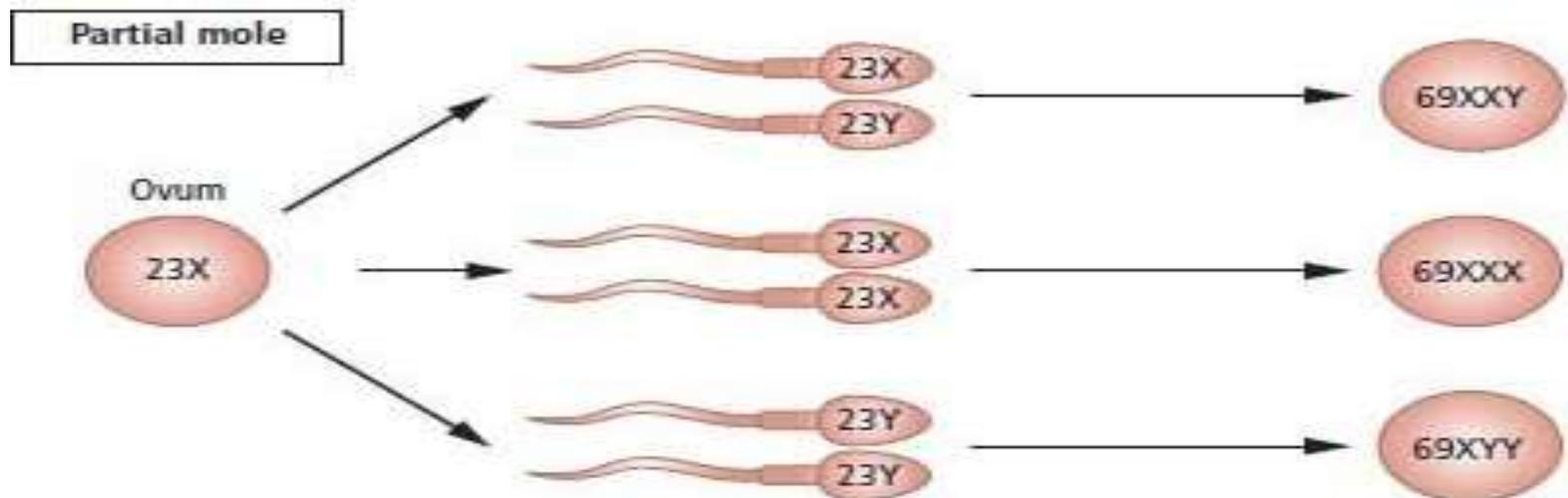
# HYDATIDIFORM MOLE - COMPLETE

- 🕒 Complete mole are not compatible with embryogenesis & **does not contain fetal parts.** The chorionic epithelial cells are diploid (46,XX or, uncommonly, 46,XY)



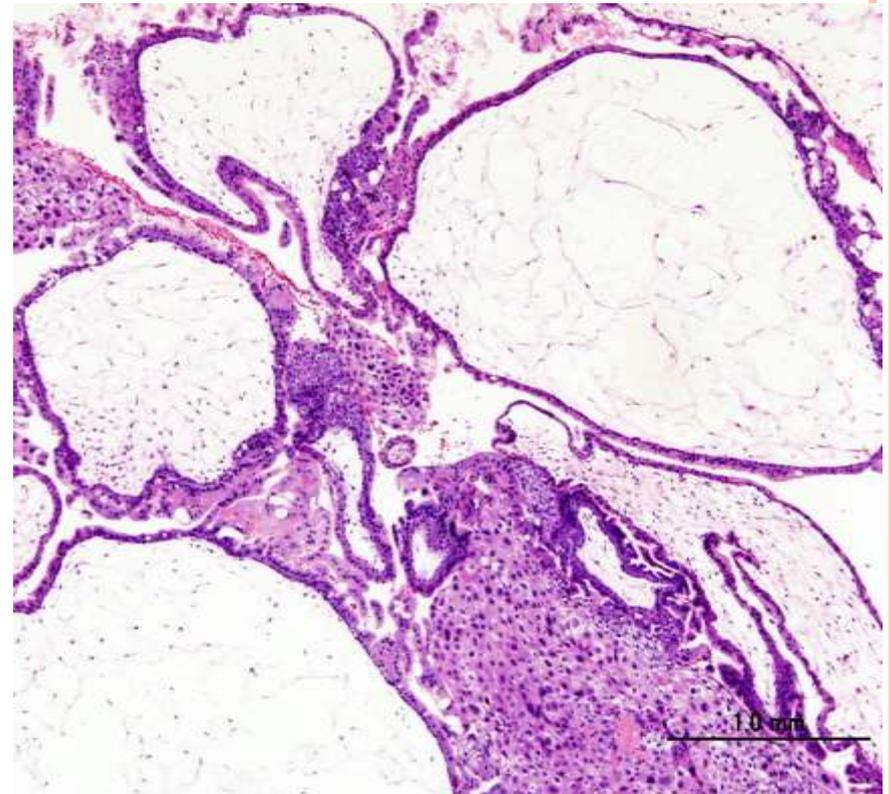
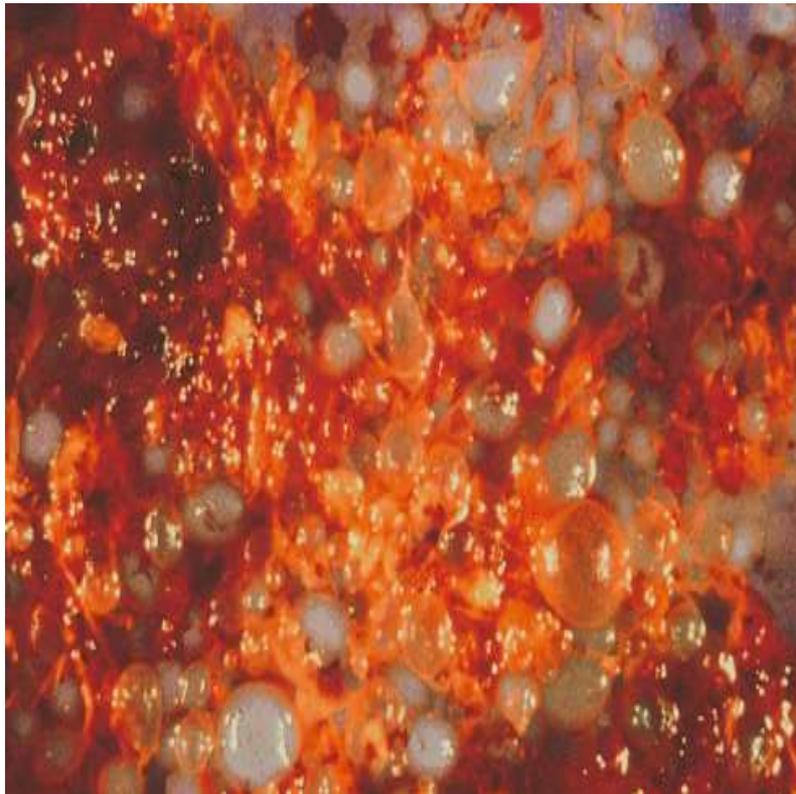
# HYDATIDIFORM MOLE - PARTIAL

- ⌚ Partial mole is compatible with early embryo formation → **may contain fetal parts & some normal chorionic villi**. Chorionic epithelial cells almost always triploid (e.g., 69,XXY)



# HYDATIDIFORM MOLE – MORPHOLOGY

Uterine cavity is expanded by friable mass (**Grape-like villi**) composed of thin-walled, cystically dilated chorionic villi covered by varying amount of atypical chronic epithelium.



# HYDATIDIFORM MOLE – ULTRASOUND SNOW STORM



<b>Feature</b>	<b>Complete Mole</b>	<b>Partial Mole</b>
<b>Karyotype</b>	<b>46,XX (46,XY)</b>	<b>Triploid (69,XXY)</b>
<b>Villous edema</b>	<b>All villi</b>	<b>Some villi</b>
<b>Trophoblast proliferation</b>	<b>Diffuse; circumferential</b>	<b>Focal; slight</b>
<b>Atypia</b>	<b>Often present</b>	<b>Absent</b>
<b>Serum hCG</b>	<b>Elevated</b>	<b><u>Less elevated</u></b>
<b>hCG in tissue</b>	<b>++++</b>	<b>+</b>
<b>Behavior</b>	<b>2% choriocarcinoma</b>	<b>Rare choriocarcinoma</b>

Syncytiotrophoblasts with multinucleation



Cytotrophoblasts



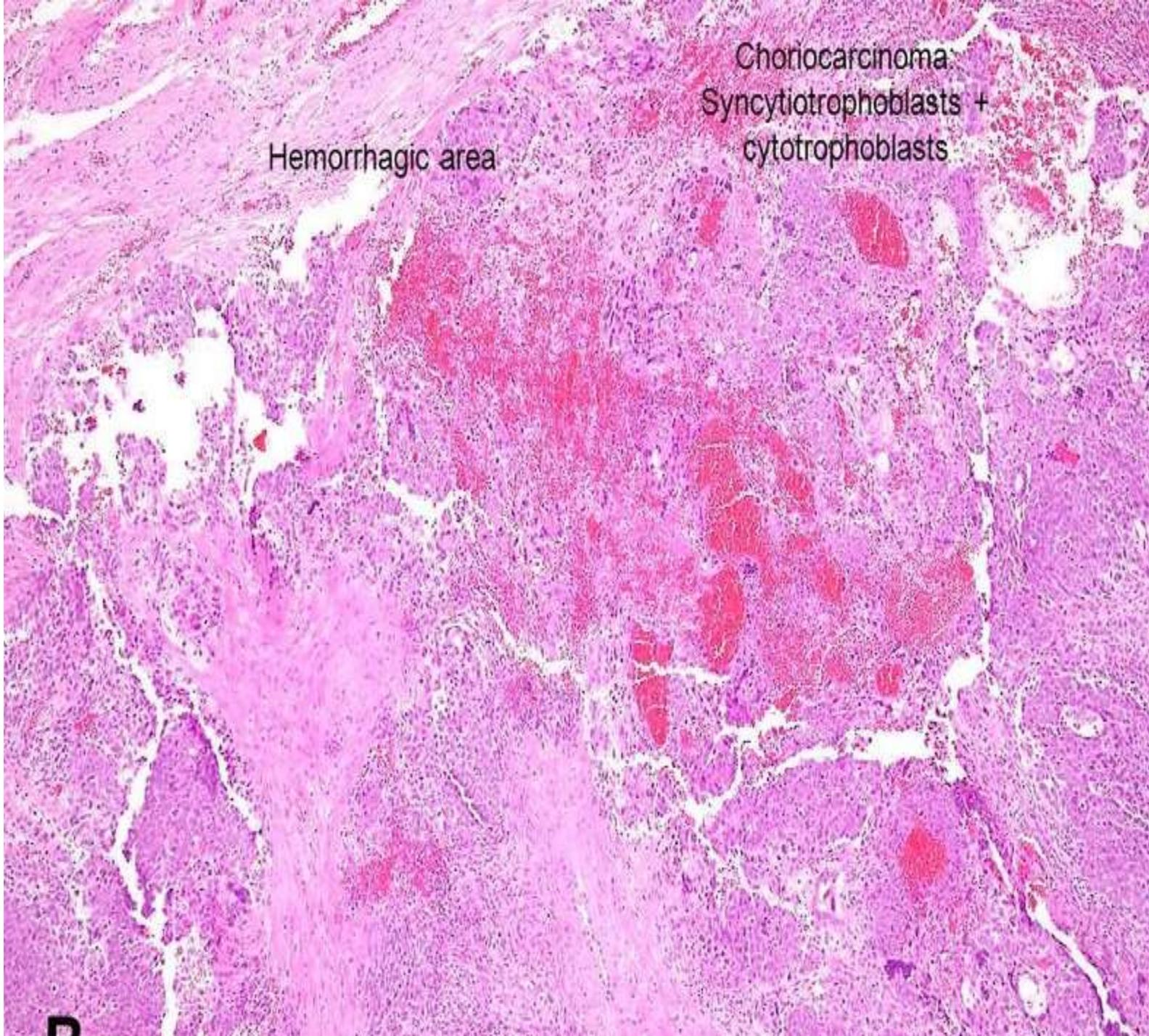
**C**



Hemorrhagic area

Choriocarcinoma:  
Syncytiotrophoblasts +  
cytotrophoblasts

D



GOOD LUCK IN YOUR EXAMS

