Ovarian Neoplasms

Ovarian Neoplasms- Introduction

- Common neoplasms.
- 80% are benign young (20-45)
- 20% are Malignant older (>40)
- 6% of all cancers in women.
- <u>50% deaths</u> due to <u>late detection</u>.

OVARIAN NEOPLASM

- NON-NEOPLASTIC functional cyst
- Primary
- Secondary

Non-neoplastic

- Corpus luteal:
- excessive bleeding into corpus luteum
- Cyst filled with blood
- Delayed period + pain
- Usually the following period is heavy

Non-neoplastic

- Granulosa-theca lutein cyst:
- in molar pregnancy or part of hyperstimulation syndrome
- due to excessive gonadotrophin
- Polycystic ovary
- Endometriotic cyst

Risk Factors

- Null parity
- Gonadal Dysgenesis
- Family History
- Ovarian cancer genes
 - BRCA1 (17q12) & BRCA2(13q12)
 - (Breast & ovary)

Classification

- Surface epithelial 65-70%
- stromal 15-20%
- Germ cell tumors 5-10%
- Metastatic tumors 5%

Surface Epithelial tumors:

- Coelomic mesothelium.
 - Serous(tubal), Mucinous (Cx) & endo
- Most common primary neoplasms
- 90% of malignant tumors of ovary
- Morphologically
 - Cystic Cystadenomas
 - Solid/cystic Cystadenofibromas
 - Solid adenofibromas

Surface Epithelial tumors

- Serous (tubal)
- **Mucinous** (endocx & intestinal)
- Endometrioid
- Transitional cell Brenners.
- Clear cell

Surface Epithelial tumors

All types can be benign, borderline , or malignant, depending upon;

- Benign ;
- gross: mostly cystic
- microscopic; fine papillae, single layer covering (no stratification), no nuclear atypia, no stromal invasion)

Surface Epithelial tumors

• Borderline

- gross; cystic / solid foci
- microscopic; papillary complexity, stratification, nuclear atypia, no stromal invasion
- Malignant ; -
- gross; mostly solid & hemorrhage / necrosis
 - microscopic; papillary complexity, stratification, nuclear atypia, stromal invasion

Serous Tumors:

- Frequently bilateral (30-66%).
- 75% benign/bord., 25% malignant.
- One unilocular cysts, papillary/less solid- **benign/borderline**
- Tall columnar ciliated epithelium.
- Papillary, solid, hemorrhage, necrosis or adhesions malignancy.
- Extension to peritoneum bad prognosis.

Serous Cystadenoma

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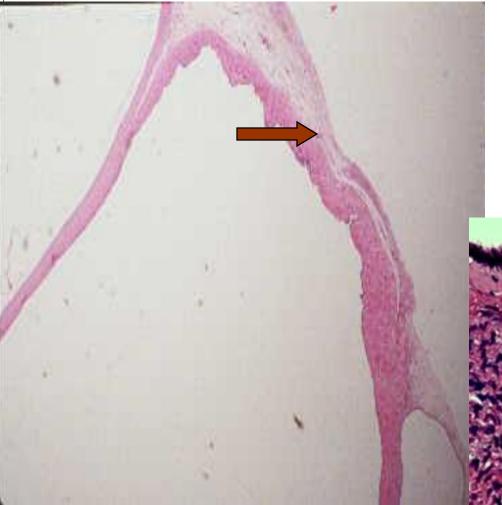
Bilateral cystadenoma



Serous Cystadenoma:

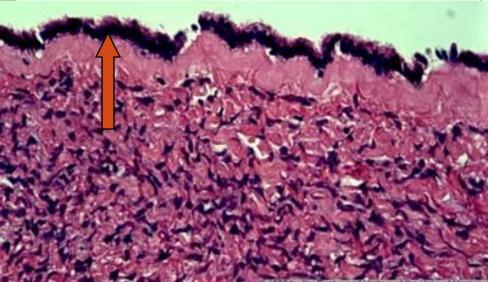


Serous Cystadenoma

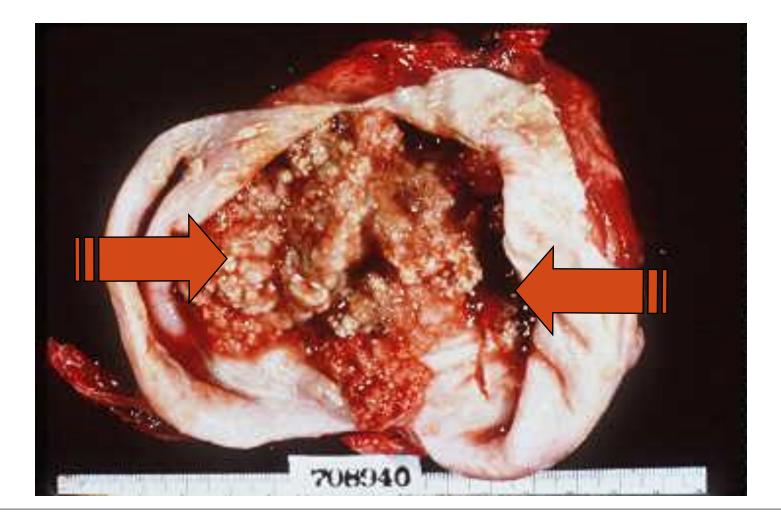


• single layer of columnar ciliated

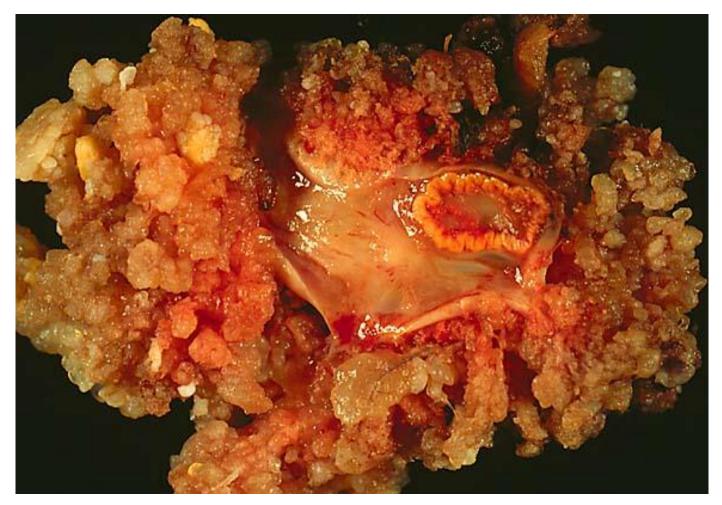
•Fine papillae



Papillary serous cystadenoma (solid/cystic)borderline

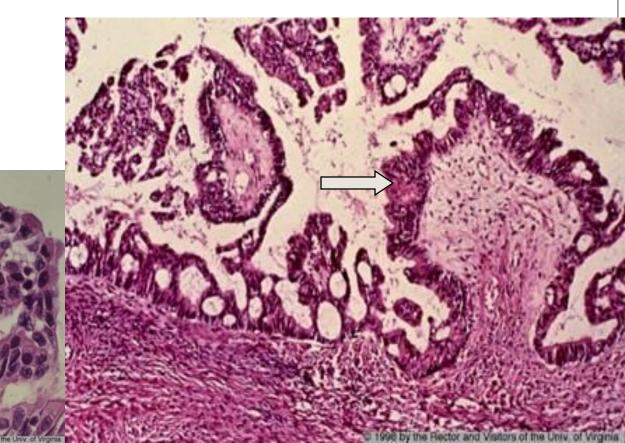


Papillary cystadenoma (bor)

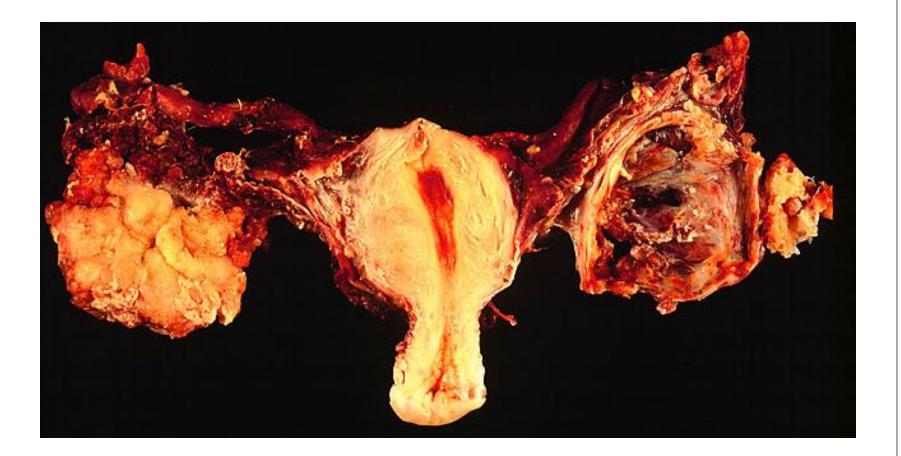


Papillary cystadenoma (bor)

- •Papillary complexity
- •Nuclear stratification& atypia
- •No stromal invasion



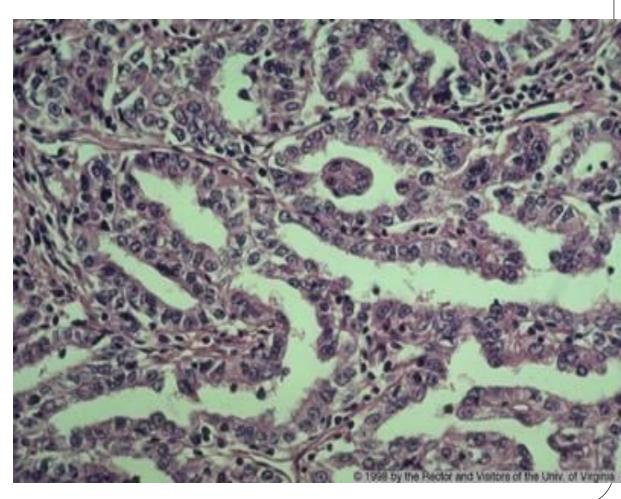
Serous cystadeno Ca – bilateral





Serous cystadenocarcinoma

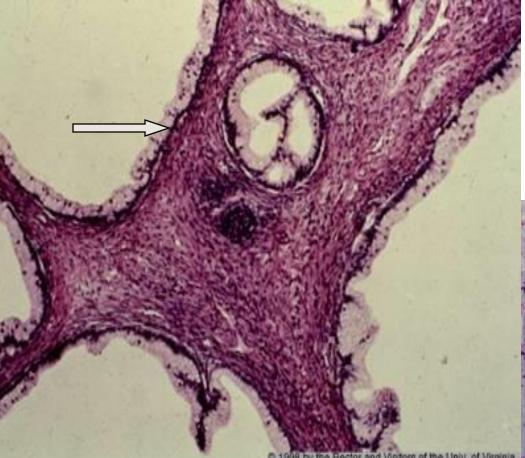
Papillary complexity
Nuclear
stratification& atypia
stromal invasion
Psammoma bodies



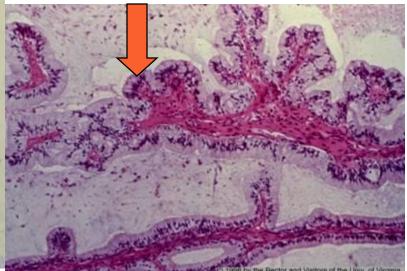
Mucinous Tumors:

- Less common 25%, very large.
- Rarely malignant 15%.
- Multiloculated, many small cysts.
- Rarely bilateral 5-20%.
- Tall columnar, apical mucin.
- <u>Pseudomyxoma peritonei</u>.

Mucinous cystadenoma



•Multilocular cyst lined by single layer of columnar cells with basally placed nuclei and apical mucin.



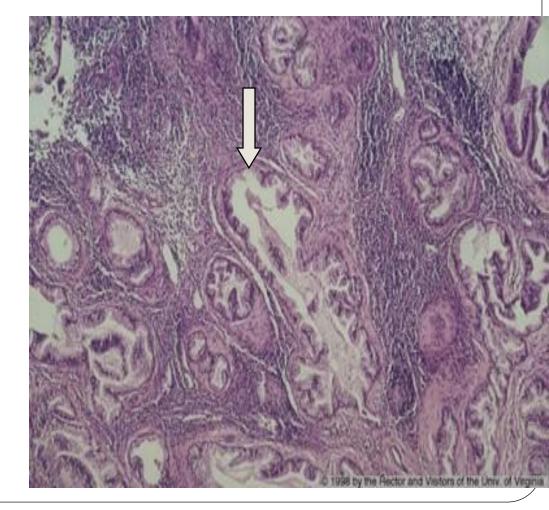
Mucinous cystadenoma-borderline

- Papillary complexityNuclear stratification& atypia
- No stromal invasion



Mucinous cystadenocarcinoma

Papillary complexity
Nuclear stratification& atypia
stromal invasion

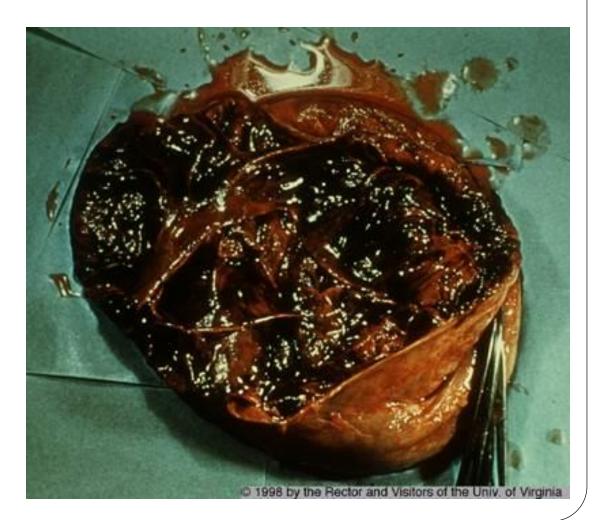


Endometrioid tumors

- most are unilateral (40% are bilateral)
- cells look like endometrium even though they are coming from the coelum of the ovary.
- almost all are malignant
- about 20% of all ovarian tumors
- many are associated with endometrial cancer (30%)
- patient may have concurrent endometriosis

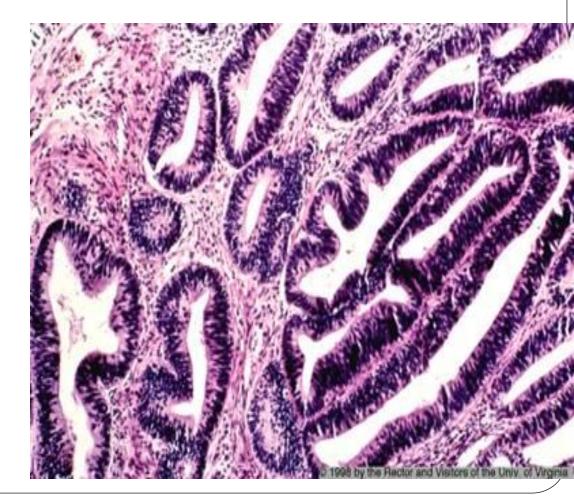
Endometrioid tumors

Solid / cyst filled by hemorrhage & necrosis



Endometrioid adenocarcinoma

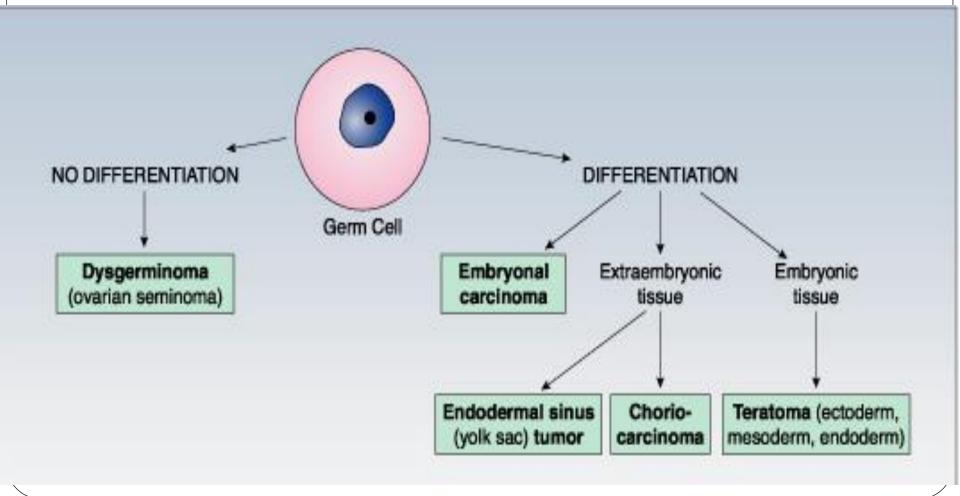
• stromal invasion by irregular malignant endometrial glands



Clinical course of coelomic surface epithelial tumors

- lower abdominal pain
- abdominal enlargement
- GI tract complaints
- urinary tract complaints
- malignant ones produce ascites
- serosal surfaces are seeded with cancer metastasis
- grow slowly and get very large

Germ cell tumor- classification



Germ cell Tumors

Teratoma

- Benign cystic (dermoid cysts)
- Solid immature
- Monodermal struma ovarii, carcinoid
- Dysgerminoma
- Yolk sac tumor (Endodermal sinus tumor)
- Choricarcinoma
- Mixed germ cell tumor

Dermoid (Benign cystic terratoma)

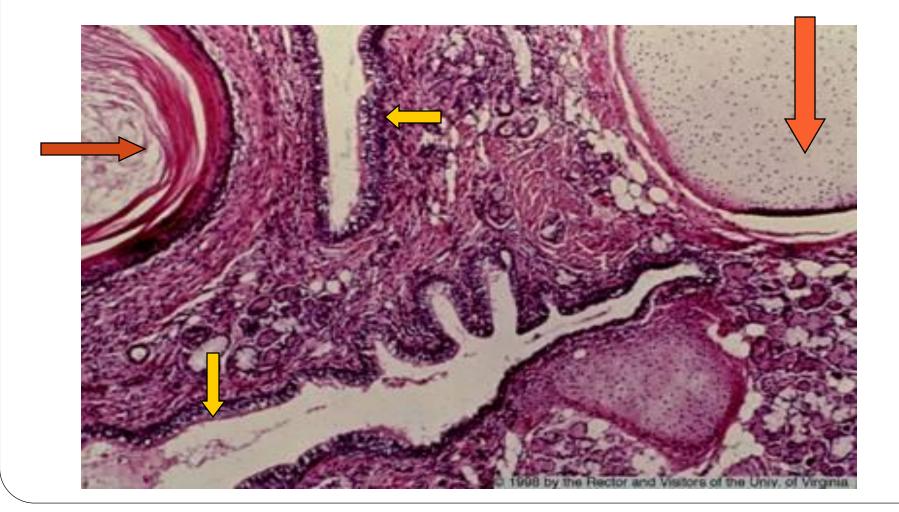
- Ectoderm + Mesoderm + Endoderm
- Supradiaphramatic structures
- Teeth permanent never deciduous
- Teeth canines,molars,incisors but never premolars
- No Gonadal tissue
- But assoc. with Granulosa

Theca Androblastoma

Cystic Teratoma (Dermoid Cyst)



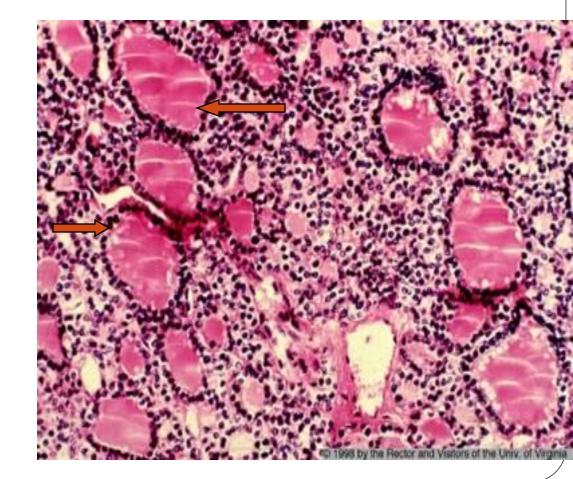
Dermoid Cyst



Monodermal teratomas

•Struma ovarii:

composed entirely of mature thyroid tissue



Dermoid (Benign cystic terratoma)

Complications

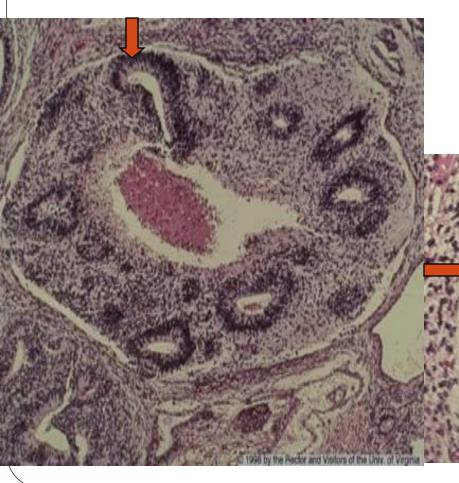
- Torsion 10%
- Rupture 1%
- Infection
- Hemolytic anemia-splenomegaly
- Malignancy

Immature Teratoma



•Solid/ necrosis &hemorrhage

Immature Teratoma



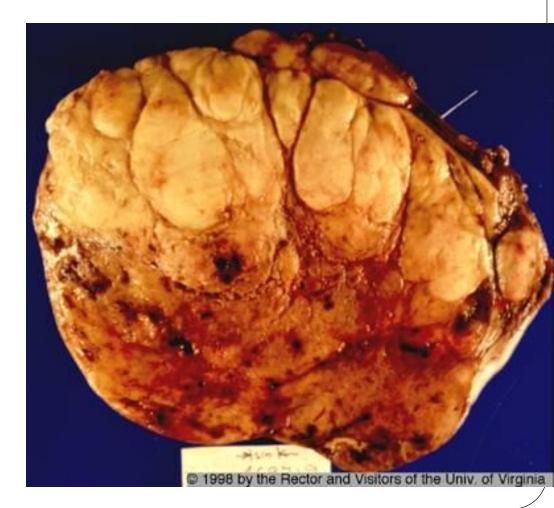
• primitive neuroepithelium with multiple neural tubes

Dysgerminoma

- The ovarian counterpart of the testicular seminoma
- 2% of all ovarian malignancies
- Most common malignant germ cell tumor
- Affects primarily younger females with the majority in the second and third decades.
- It is the <u>most frequently encountered ovarian</u> <u>malignancy in pregnancy</u>
- An excellent prognosis. Highly radiosensitive .

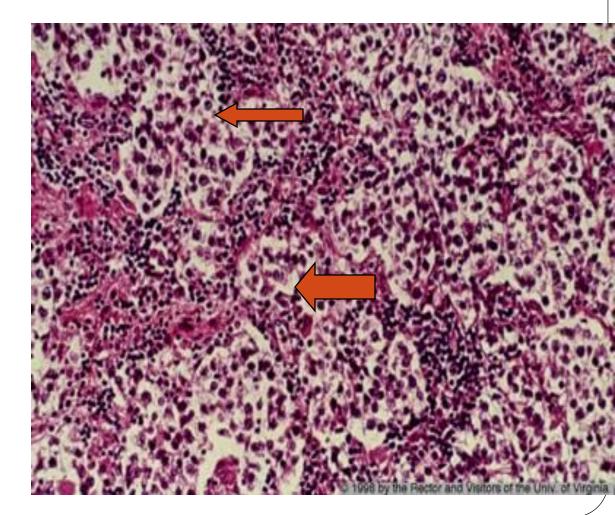
Dysgerminoma

•Solid/ lobulated mass with foci of hemorrhage



Dysgerminoma

•sheets of monotonous rounded cells with pale cytoplasm and central nuclei

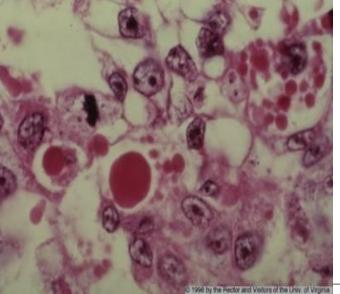


Endodermal sinus tumor (Yolk sac carcinoma)

- Tumor is a <u>highly malignant</u> and clinically aggressive neoplasm
- Most frequently in **children and young females**
- 20% of malignant germ cell tumors.
- Fatal within 2 years of diagnosis

carcinoma)

• Schiller-Duval body





Sex Cord - Stromal Tumors

- Granulosa-cell tumor
- Thecoma
- Fibroma
- Sertoli-Leydig cell tumors

Granulosa Cell Tumor

- Hormonally active tumor
- The most common <u>estrogenic ovarian neoplasm.</u>
- The <u>adult form</u> occurs mainly in postmenopausal women, associated with <u>endometrial</u> <u>hyperplasia and carcinoma</u>
- The juvenile type occurs in the first two decades, cause precocious sexual development.

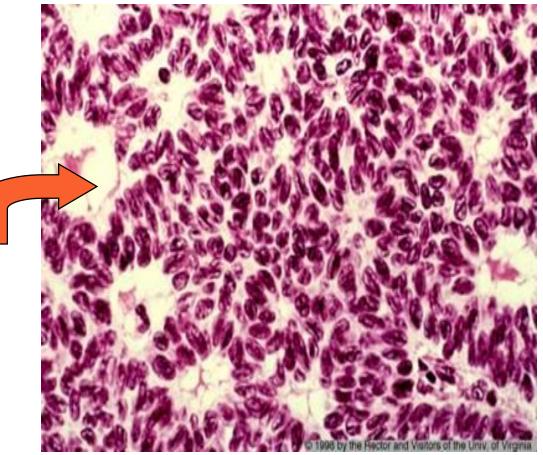
Granulosa Cell Tumor

•Solid with hemorrhage



Granulosa Cell Tumor

•Sheets of granulosa cells containing spaces lined by the cells to give a follicle-like appearance (Call-Exner bodies).



Thecoma

- Functional tumors producing estrogen
- It occur in postmenopausal women
- Endometrial hyperplasia or carcinoma may develop

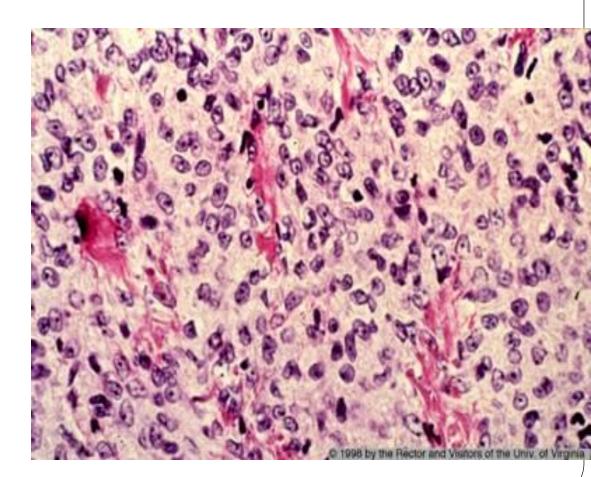
Thecoma

•Solid tumor with yellow - orange appearance.



Thecoma

• sheets of round to oval cells with pale cytoplasm containing lipid.

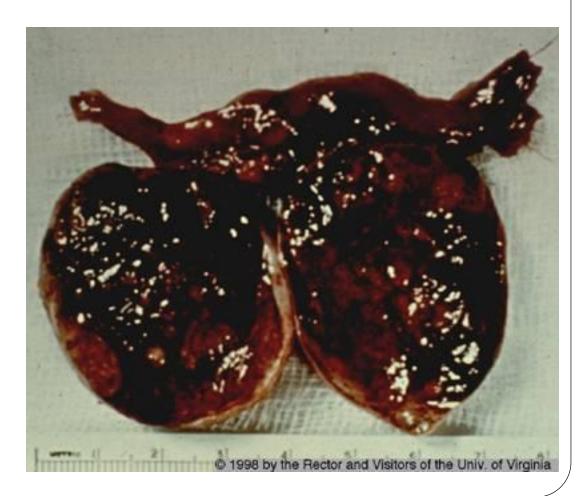


Sertoli-Leydig cell tumors

- 1% of ovarian neoplasms
- It occur predominantly in young women.
- Commonly androgenic, cause defeminization of women manifested as breast atrophy, amenorrhea, and loss of hair and hip fat, to virilization with hirsutism

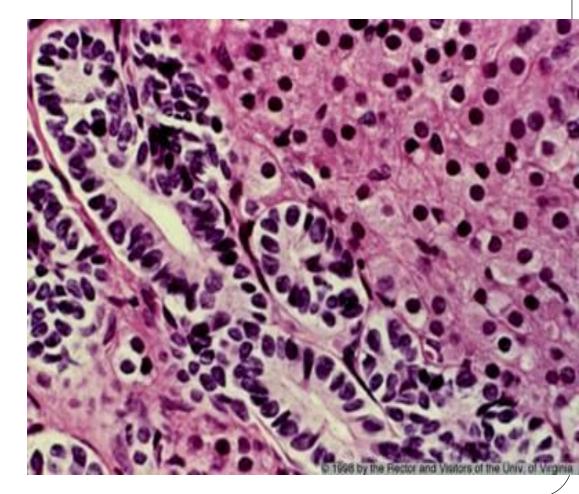
Sertoli-Leydig cell tumors

• solid & hemorrhage



Sertoli-Leydig cell tumors

•Tubules lined by Sertoli cells and sheet of Leydig cells



Metastases to ovary

- About 3% of malignant tumors in the ovary are metastatic
- The most common primary site is the breast followed by the large intestine, stomach, and other genital tract organs.

Krukenberg tumor

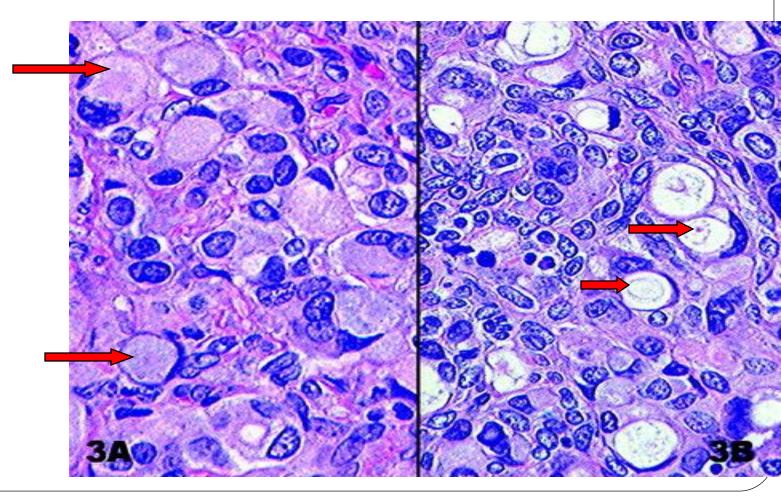
- It is applied to the uniform enlargement of the ovaries (usually bilaterally) due to diffuse infiltration of the ovarian stroma by **metastatic signet-ring cell carcinoma** .
- The commonest primary site is the stomach followed by the colon.

Krukenberg Tumor



Krukenberg Tumor

•Ovarian infiltration with signet ring cell



Physical signs

• <u>Benign</u>:

- usually mobile.unless large or complicated

• Malignant:

- Bilateral
- Ascites
- Hard deposit in pelvis
- Leg edema
- Signs of bowel obstruction of ureteric obstruction

Investigation

- Uss / CT scan
- Tumor markers(ca125,CEA, HCG,alpha FP
- Urea and electrolyte
- LFT
- Chest X ray
- Ascitic tap
- Calculate **RISK MALIGNANCY INDEX**

RISK MALIGNANCY INDEX

- CA 125 estimation
- Menopausal status

pre menopausal score = 1

post menopausal score= 3

• Ultrasound score

Multi locular, solid areas, bilateral, ascitis, intra abdominal mets.

if 0 or 1 score = 1

if 2-5 score= 3

RMI = CA125 X M X U

FIGO Staging

Stage 1	Growth limited to one or both ovaries
Stage 2	Growth limited to one or both ovaries with pelvic extension
Stage 3	Tumor involving one/both ovaries with peritoneal implants outside pelvis/positive retroperitoneal or inguinal nodes
Stage 4	Growth involving one or both ovaries with distant metastasis

MANAGMENT

- Surgery : primary interval debulking palliative second look surgery
- Chemotherapy

Primary surgery

- Primary cytoreduction
- TAH,BSO,OMETECTOMY,WASHINGS
- BOWEL SURGERY
- Optimal debulking: less than 2 cm residual tumors
- Staging once histology is available
- If confined to ovary and young age... conservative surgery

Interval debulking

- Alternative to primary surgery medically unfit large ascitis severe malnutrition
- 3 cycles of chemotherapy –surgery 3 more cycles of chemotherapy
- Aim : to improve patient condition less extensive surgery to achieve optimal debulking
- May improve survival

Chemotherapy

- Indication –
- stage 1c and above
- Platinium based
- Taxol
- 6 cycles at 3 weekly intervals
- Monitoring: examination CA125 FBC, U&E

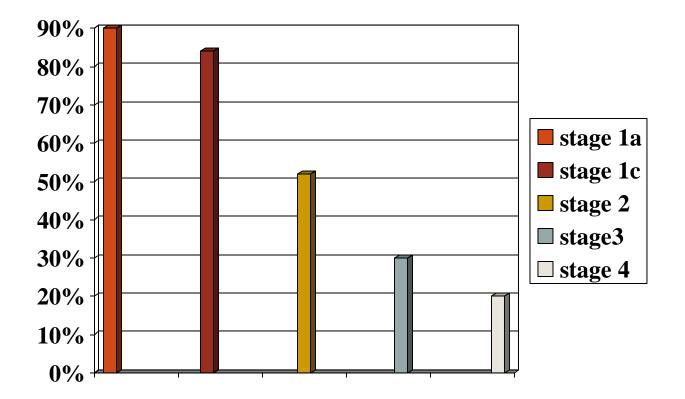
SECOND LOOK SURGERY

- Assess response to chemotherapy
- Plan future management
- Only in research context.

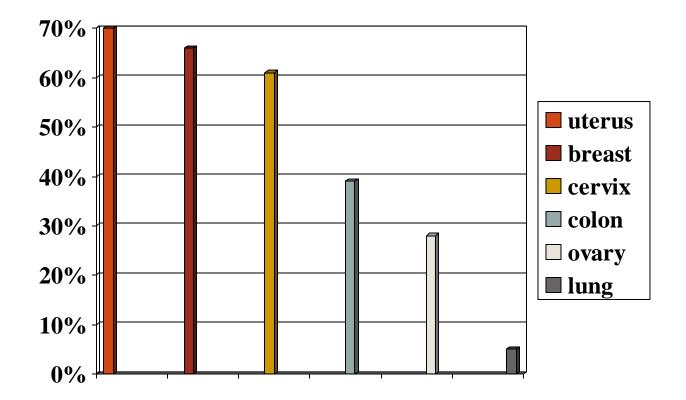
Palliative surgery

- Removal of intestinal obstruction
- Survival is very poor
- Quality of life considerations

Five year survival



Five year survival



Five year survival rates in England and Wales 1986-1990

Follow up

- How aggressive?.
- Three monthly for one year then six monthly then yearly
- History, examination and CA125
- Imaging if recurrence is suspected clinically or by CA125

Ovarian cancer screening

- Life time risk is 1%
- 5% of tumors are genetic
- History of breast cancer increases risk by factor of 2
- History of ca ovary increases the risk by factor of 3
- One first degree relative affected: risk 2.7%
- 2 first degree relatives affected : risk is 13%
- If BRCA1 mutation carrier : risk is 50%

screening

- Problems :
 - no pre-cancerous stage
 - unknown natural course
- TVS AND CA125 ON YEARLY BASIS
- ONGOING STUDY TO EVALUATE THIS.

