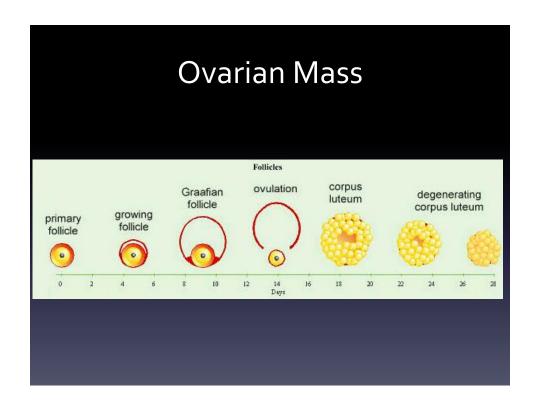
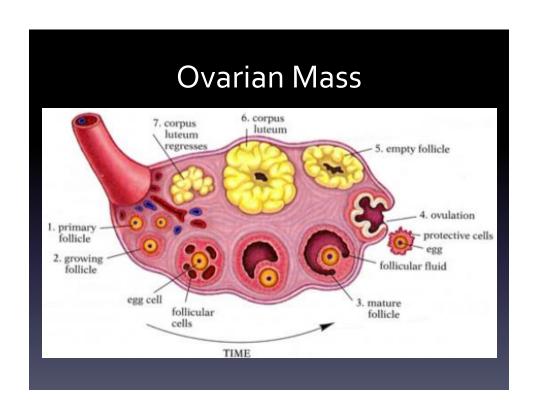
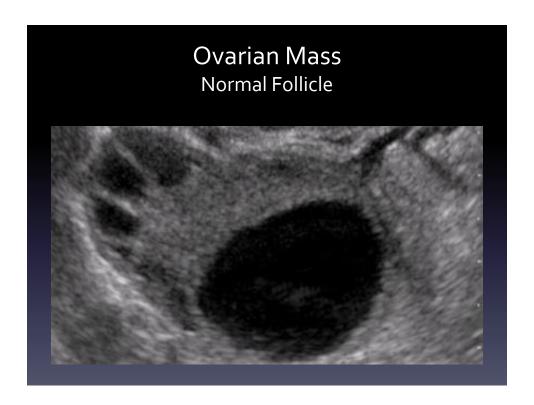
Ovarian Mass

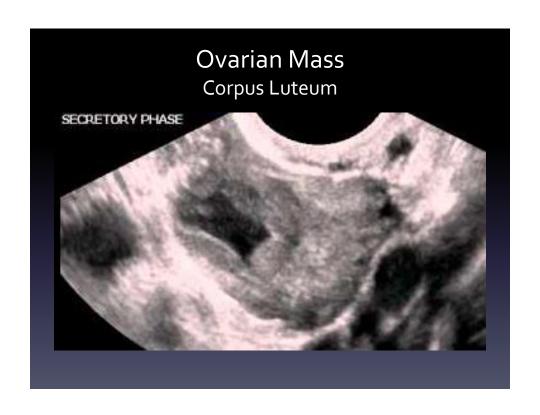
- Learning Targets
 - Normal ovarian function
 - Sources of ovarian masses
 - Evaluation of ovarian masses
 - Differential diagnosis
 - Benign or Malignant?
 - Appropriate Evaluation

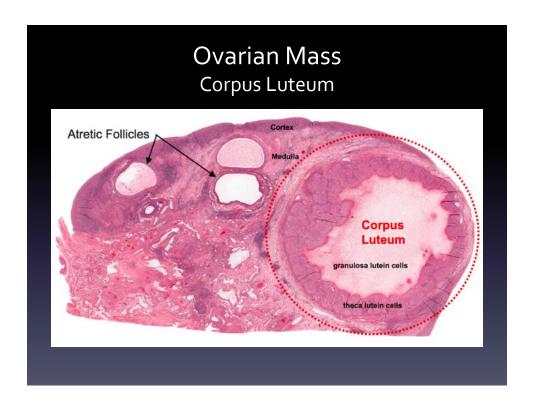
Normal Follicular Development OVARIAN MASSES

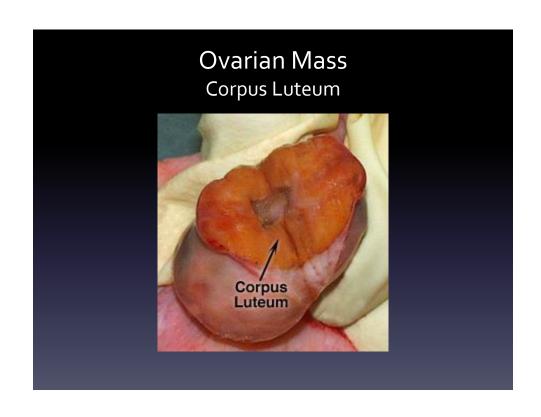


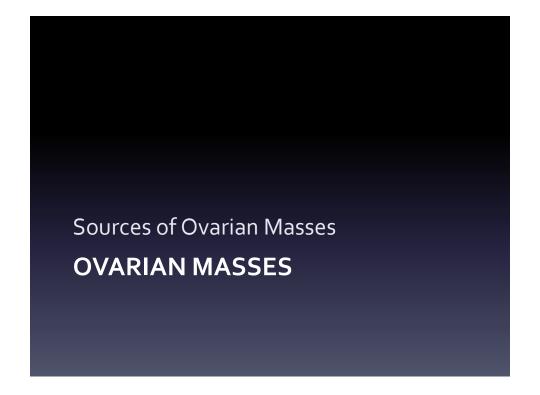




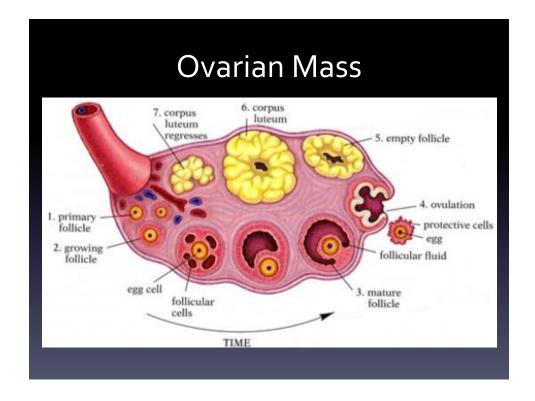








- Sources of Benign Ovarian Masses
 - Disorders of Normal Follicular Development
 - Disorders of Germ Cell Development
 - Disorders of Epithelial Cell Development



Evaluation of Ovarian Masses

OVARIAN MASSES

Ovarian Masses

- Evaluation of Ovarian Masses
 - Physical Exam
 - Ultrasonography
 - Color Doppler Ultrasonography
 - Other Imaging
 - Serum Marker Screening

- Physical Exam
 - "Limited ability to identify an adnexal mass, especially with increasing body mass index (BMI) > 30.
 - Features to be considered:
 - Regular versus Irregular
 - Solid versus soft
 - Fixed versus mobile
 - Smooth versus nodular
 - Unilateral or Bilateral
 - Ascites?

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Ovarian Masses

- Ultrasound Preferred imaging modality
 - Advantages
 - Widespread availability
 - Good patient tolerability
 - Cost-effective

- Ultrasound Preferred imaging modality
 - Limitations
 - Lack of specificity

$$-68 - 81\%$$

• Low positive predictive value for cancer

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Ovarian Masses

- Ultrasound Contents of Report
 - Size of the mass
 - Solid or cystic
 - Unilateral or bilateral
 - Presence or absence of septations
 - Mural nodules
 - Papillary excrescences
 - Free fluid in the pelvis

- Color Doppler Ultrasonography
 - Measurement of blood flow in and around the mass
 - Hypoxic tissue in tumors will recruit low resistance, high-flow blood vessels
 - Can measure pulsatility index, resistance index, maximum systolic velocity, etc.
 - Current role remains controversial

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Ovarian Masses

Modalities for the Evaluation of Adnexal Masses					
<u>Modality</u>	Sensitivity	Specificity			
Gray-scale transvaginal U/S	82-91%	68-81%			
Doppler ultrasonography	86%	91%			
Computed Tomography (CT)	90%	75%			
Magnetic resonance imaging	91%	88%			
Positron emission tomography	67%	79%			
CA 125 level measurement	78%	78%			

AHRQ. Management of adnexal mass.

- Serum Marker Screening
 - -CA 125-
 - Elevated in 80% of patients with epithelial ovarian cancer
 - Only elevated in 50% of patients with Stage 1 ovarian cancer
 - NOT useful as a screening test

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Ovarian Masses

- Serum Marker Screening
 - CA 125 False positive
 - Uterine leiomyoma
 - Endometriosis
 - Acute or chronic PID
 - Ascites, of any etiology
 - Inflammatory conditions SLE, IBD, etc.

- Serum Marker Screening
 - -CA 125
 - Useful in distinguishing between benign and malignant masses in postmenopausal women
 - Only extreme values helpful in premenopausal women
 - CA 125 values will increase over time when a cancer is present

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Ovarian Masses

- Serum Marker Screening
 - Germ Cell Tumor Markers

• Dysgerminoma Lactate dehydrogenase

Choriocarcinoma Beta-hCG level

• Granulosa cell Inhibin A and B

• Endodermal sinus Alpha-fetoprotein

Differential Diagnosis

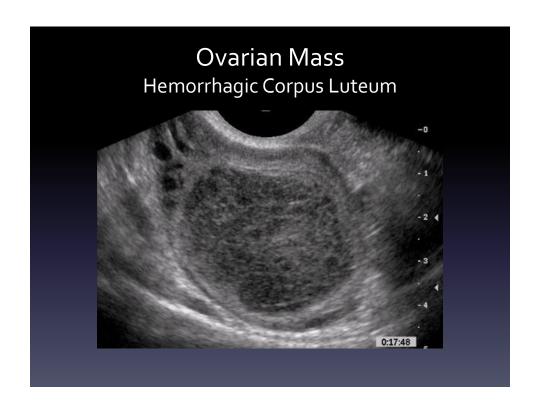
OVARIAN MASSES

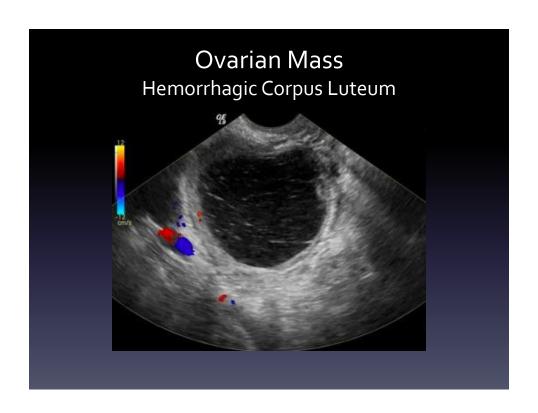
Ovarian Mass Differential Diagnosis

- Functional Cysts
 - Follicular cysts
 - Hemorrhagic corpus luteum
- Endometrioma
- Mature teratoma
- Germ Cell Tumors
- Benign neoplasm
- Malignant neoplasm

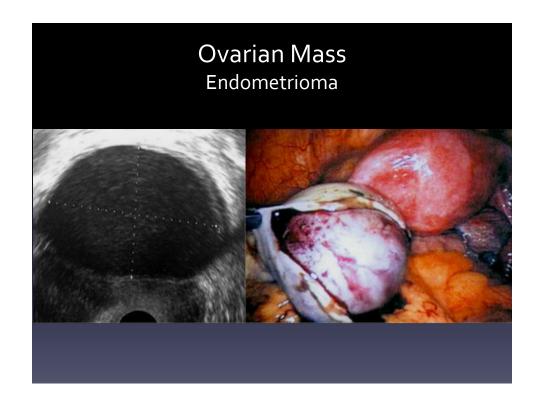


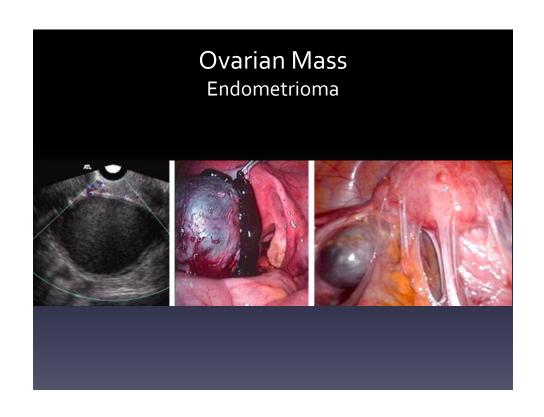


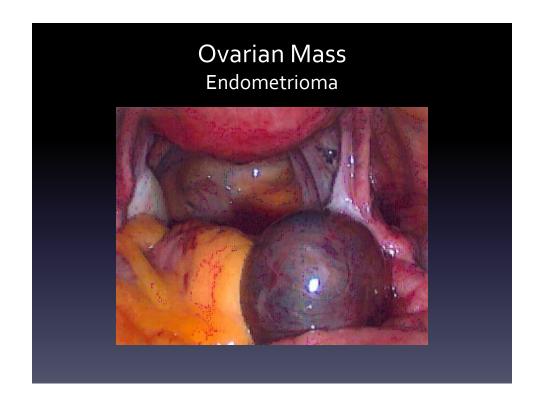


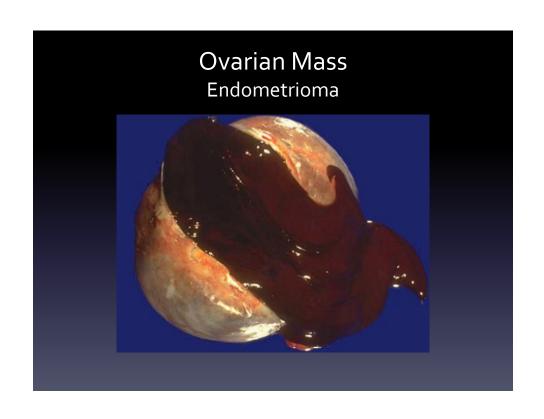


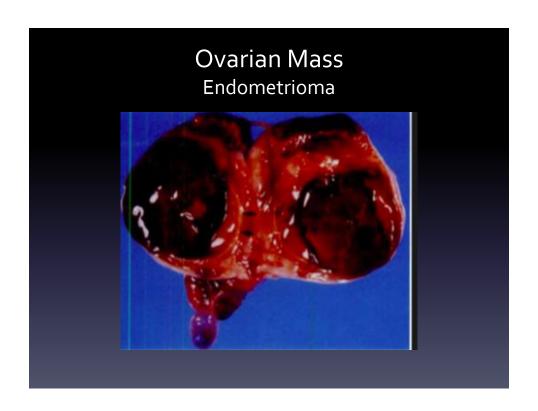


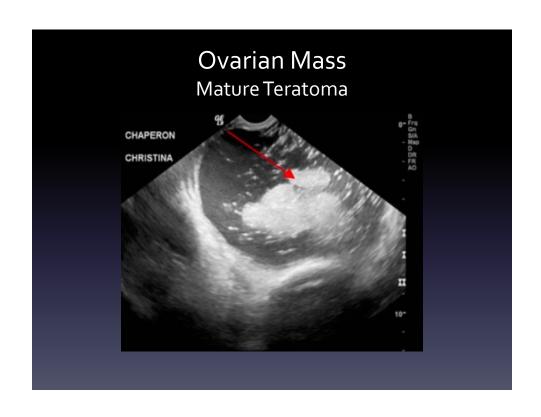


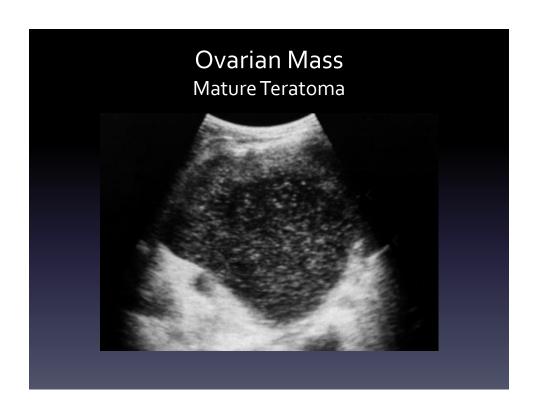








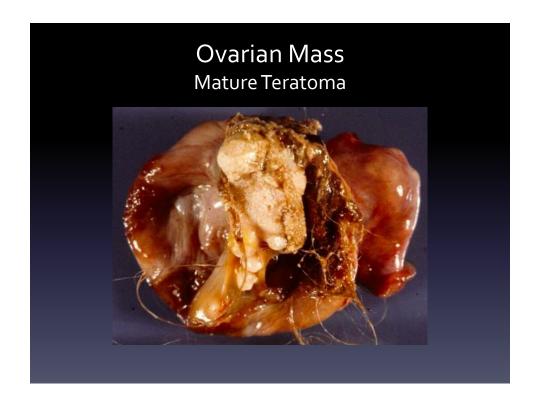


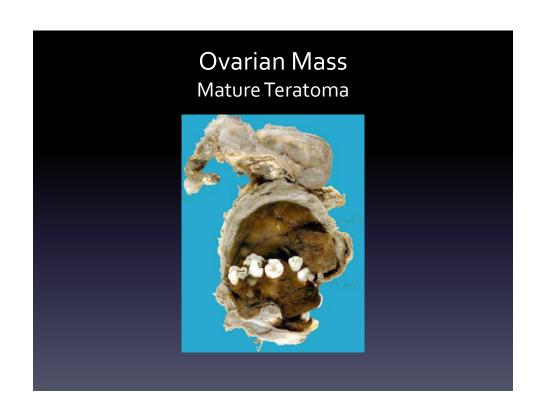


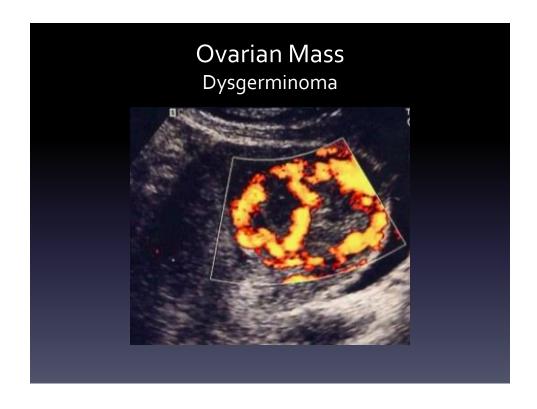


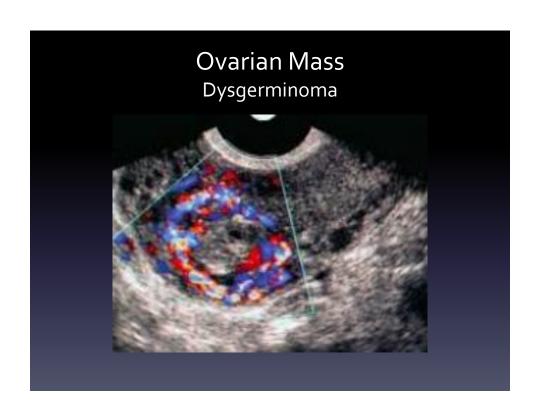


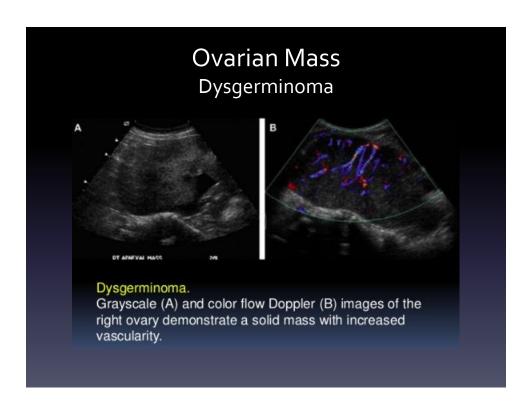


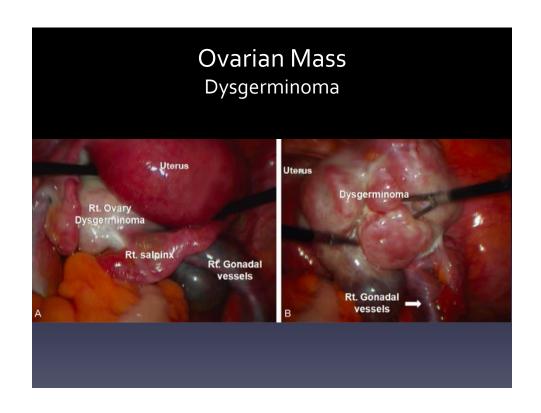


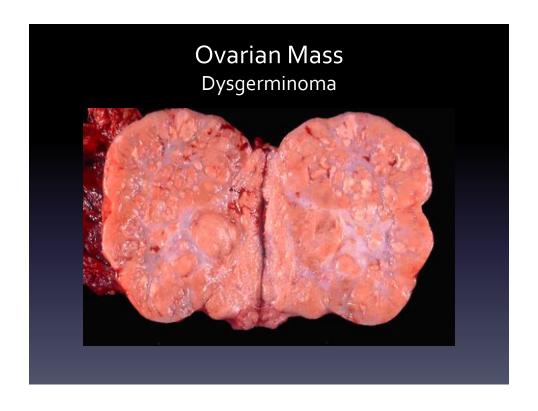


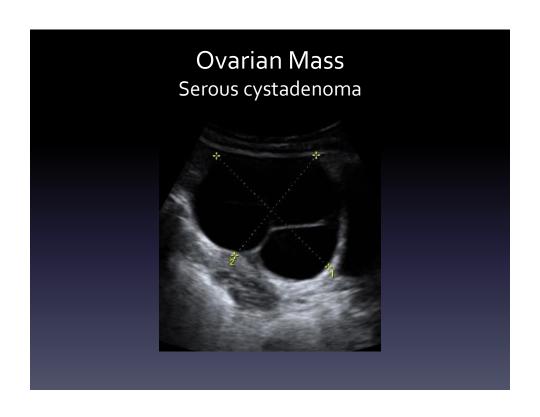


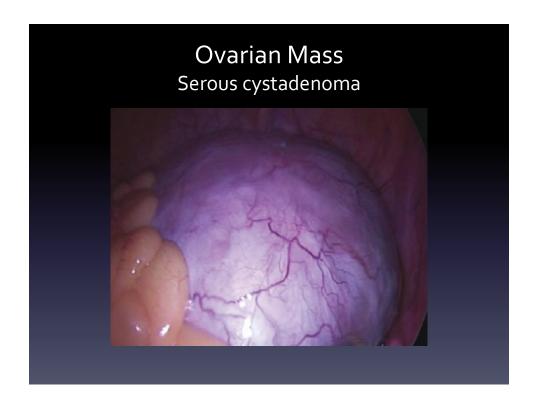


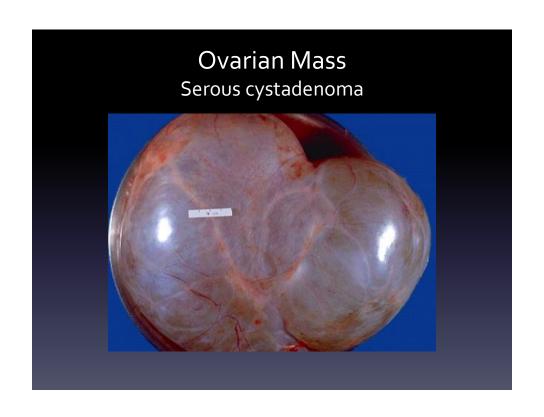


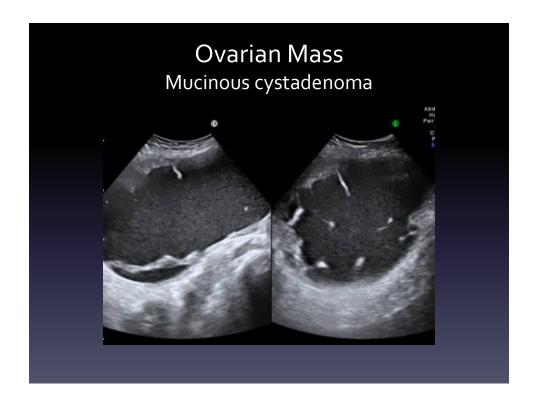


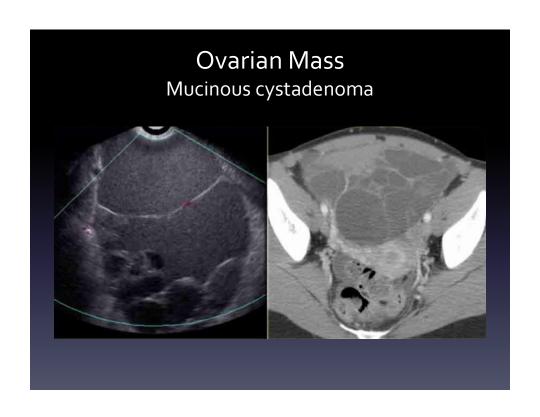


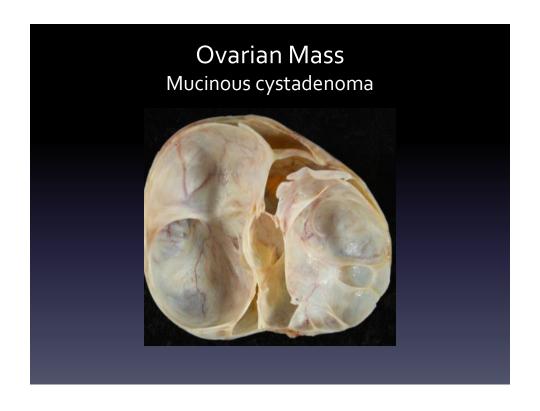














- Historical Risk Factors for Malignancy
 - Age
 - Family history of breast or ovarian cancer
 - Genetic susceptibility
 - Nulliparity
 - Primary infertility
 - Endometriosis

	Ovarian masses
Inte	ernational Ovarian Tumor Analysis
	10 SIMPLE RULES
RULES	FOR PREDICTING A MALIGNANT TUMOUR (M – RULES)
M1	IRREGULAR SOLID TUMOUR
M2	PRESENCE OF ASCITES
МЗ	AT LEAST 4 PAPILLARY STRUCTURES
M4	IRREGULAR MULTILOCULAR SOLID TUMOUR WITH LARGEST DIAMETER ≥ 100mm
M5	VERY STRONG BLOOD FLOW (COLOUR SCORE 4)
RULES	FOR PREDICTING A BENIGN TUMOUR (B – RULES)
B1	UNILOCULAR
B2	PRESENCE OF SOLID COMPONENT, LARGEST DIAMETER <7mm
B3	PRESENCE OF ACOUSTIC SHADOWS
B4	SMOOTH MULTILOCULAR TUMOUR WITH LARGEST DIAMETER < 100mm
B5	NO BLOOD FLOW (COLOUR SCORE 0)

International Ovarian To		_
iliterilational Ovarian i	umor Ana	lysi
TABLE 5 Model coefficients for 11 predictors obtained o data (n = 2445)	n development	
Predictor	Coefficient	SE
Intercept	-1.10	0.2
B1 (unilocular cyst)	-3.10	0.3
B2 (solid components present, but <7 mm)	-1.55	0.5
B3 (acoustic shadows)	-1.58	0.2
B4 (smooth multilocular tumor with largest diameter <100 mm)	-3.59	0.6
B5 (no blood flow; color score 1)	-1.96	0.24
M1 (irregular solid tumor)	2.38	0.39
M2 (ascites)	2.87	0.29
M3 (at least 4 papillary structures)	1.72	0.28
M4 (irregular multilocular-solid tumor with largest diameter ≥100 mm)	1.12	0.23
M5 (very strong flow; color score 4)	1.53	0.24
Oncology center	0.95	0.31

Ovarian masses International Ovarian Tumor Analysis

TABLE 1)				
Summar	classification	of Simple Rules	risk calculation	based on all d	ata (n = 4848)

Features	Observed malignancy rate	Estimated individual risk of malignancy	Classification
No M-features AND >2 B-features	1/175 (0.6%)	<0.01-0.29%	Very low risk
- No M-features AND 2 B-features - No M-features AND feature B1 present	20/1560 (1.3%)	0.19-2.7% 1.2-3.1%	Low risk
No M-features AND 1 B-feature present (except B1)	60/722 (8.3%)	2.4—15.2%	Intermediate risk
- No features - Equal no. of M- and B-features - >0 M-features, but more B- than M-features	451/1096 (41.1%)	27.5—48.7% 5.6—78.1% 1.3—28.4%	Elevated risk
More M- than B-features present	1133/1295 (87.5%)	42.0->99.9%	Very high risk

This simplified system only provides risk ranges for no. of B- and M-features present, but facilitates clinical triaging in absence of electronic devices. Personalized risk estimates can be obtained in second step.

B-feature, benign feature; M-feature, malignant feature

Timmerman et al. Simple ultrasound rules to predict risk of malignancy in adnexal masses. Am J Obstet Gynecol 2016.

Ovarian masses

International Ovarian Tumor Analysis (IOTA)

ACOG/SGO Referral Guidelines

- Premenopausal (< 50 years of age)
 - CA 125 levels > 200
 - Ascites
 - Evidence of metastasis
 - Family history of breast or ovarian malignancy (1st Degree)

International Ovarian Tumor Analysis (IOTA)

ACOG/SGO Referral Guidelines

- Postmenopausal (> 50 years of age)
 - Elevated CA 125
 - Ascites
 - Nodular or fixed pelvic mass
 - Evidence of metastasis
 - Family history of breast or ovarian malignancy (1st degree)

Appropriate Evaluation

OVARIAN MASSES

- Premenopausal
 - Medical history and physical examination
 - Quantitative B-hCG level
 - Complete Blood Count (CBC)
 - Transvaginal U/S
- Almost all pelvic masses in premenopausal women are benign

Ovarian Masses

- Postmenopausal
 - Medical history and physical examination
 - CA 125
 - Transvaginal U/S
- "With the exception of simple cysts on a transvaginal ultrasound, most pelvic masses in postmenopausal women will require surgical intervention."