Macroscopy course Ovary and Fallopian tube

B. 202'i



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- Some anatomy and terminology
- Concept of peritoneal cavity and why it is important to understand it
- Most common clinical context
- Salpinx
- Ovary

Adnexa = appendage of the uterus adnectere: what is attached to



GRAY H. GRAY'S ANATOMY. LEA & FEBIGER, NEW YORK, 1918.

35 & BSI

Adnexa = appendage of the uterus adnectere: what is attached to



GRAY H. GRAY'S ANATOMY. LEA & FEBIGER, NEW YORK, 1918.

Adnexa = appendage of the uterus A world about orientation

Posterior view



GRAY H. GRAY'S ANATOMY. LEA & FEBIGER, NEW YORK, 1918.



Looking from above

- Salpingectomy = removal of the Fallopian tube
- Oophorectomy = ovariectomy
- Adnexectomy = salpingo-oophorectomy
- It can be unilateral or bilateral



Because you might read this..: BSO

Bilateral salpingo-oophorectomy



What's the name of all the fluffy things around? Image: Itigament? Adipose and fibrous tissue, holding things together and carrying blood supply and

lymphatics



Posterior

Peritoneal cavity

- Virtual cavity that the surgeon makes real by inflating it, or where liquid can accumulate in some pathological conditions
- From the liver to the diaphragm to the guts to the uterus, peritoneum is a thin membranous glistening layer, like a sac
- Important to understand the concept because it means anything that happens to the ovary can very quickly spread through this cavity





Peritoneal cavity

• What does the surgeon see ?





When do we remove the adnexa?

- Salpingectomy:
 - Tubal pregnancy
 - Abcess, hydrosalpinx, hematosalpinx
 - Often context of young patient
- Adnexectomy:
 - Cyst, mass
 - Is it benign? Is it malignant? Radiologist and clinicians have a guess, we have the last word
 - Prophylactic: remove it before cancer develops
- What about ovariectomy alone?

Somehow a rare situation > if you remove the ovary you have no reason to keep the salpinx

Opposite is not true ! Especially in patient before menopause, you want to preserve the ovary as much as possible Always check the age and clinical informations !

Salpingectomy



Salpingectomy







Ovarian's surface and fimbriae are intimately connected



Fimbriae are complex and numerous infoldings of tubal mucosa

Salpingectomy-incidental findings

What you might see often:

- little (<1cm) cyst at the infundibulum
- sterilisation procedures (metal clips (1 or 2) or threads)

What you might see sometimes:

- tubal phymosis







Tubal pregnancy

- Pregnancy can be found in any segment of the salpinx
- Specify number of fragments
- Integrity of tubal surface
- Report not 2 but 3 dimensions (smallest and largest diameters)
- Describe content if identified (blood, placental villi, embryonal remnants...)

What's at stake?

- Identify the tubal implantation of the pregnancy
- Exclude a placental pathology (hydatiform mole)





Prophylactic bilateral salpingectomy:

- Hereditary context linked to a high risk of ovarian cancer
- Mutation of BRCA gene
- Ovarian cancer comes from ... the fimbriae of the salpinx!

Remember...?







Very early cancer will not translate macroscopically: occult malignancy

You're dealing with a complex tridimensionnal structure : the fimbriae

You are going to analyze on slide in a bidimensional way

Remember our concept of peritoneal cavity, anything happening in the adnexa can very quickly spread !





How to cut your specimen in a way to maximize your chances to find Nemo?













What is a normal ovary? ... depends of the age of the patient

Wide range of morphologically different but normal gross aspects depending on the age

Expected size of an adult ovary:



What is a normal ovary? ... depends of the age of the patient

Wide range of morphologically different but normal gross aspects depending on the age

Inspection: pink to whiter exterior surface that can be smooth or convoluted







Three ill defined zones are discernible



What is a normal ovary? ... depends of the age of the patient

Wide range of morphologically different but normal gross aspects depending on the age

Cut section



Pre-ovulatory follicle reaches up to 25 mm physiologically

Wide range of morphologically different but normal gross aspects depending on the age

Cut section



It means you are going to see « cyst » ! And it's NORMAL Clues in favour of physiological: cortical, unilocular,gelatinous content, around one centimeter If in doubt: check your patient AGE + take PHOTOS



« Usual suspect »: The Corpus luteum

Scalopped/festooned contours Yellowish ribbon Hemorragic content

Wide range of morphologically different **but** normal gross aspects depending on the age

Cut section



Nevertheless.....

The « normal » ovary of hormonally active woman is NOT what you 're gonna see the most, because we try not to remove them!

You'll see a lot of: -perimenopausal

-post menopausal ovary > atrophic

Whitish, smaller

Medulla and hilum expand to the detriment of cortex



How to sample a « normal » ovary:





- Examine cut surface
- If normal: one or two sections where cortex/medulla and hilum are visible





Cyst.. I mean a mass in general Remember, often, no prior histological diagnosis. Remain vigilant! It could be anything !

ALWAYS look for clinical and radiological information

INTEGRITY of the specimen: INTACT or COLLAPSED or FRAGMENTED

SURFACE

SIZE and WEIGHT

When in doubt : take a PHOTO !



INTEGRITY of the specimen

INTACT



COLLAPSED



FRAGMENTED or overtly OPENED/BREECHED



INTEGRITTY of the specimenINTATCOLLAPSEDINTATCOLLAPSEDINTAT

You don't know what you're looking at yet!



Keep in mind the concept of peritoneal cavity and quick spreading of adnexal tumor

Why is it so important?





A 2014 meta-analysis assessed the impact of intraoperative rupture on prognosis, after analysing nine eligible studies which included 2,382 patients.¹³ Patients with preoperative capsular rupture showed poorer progression-free survival (PFS) than those with no rupture or intraoperative rupture. In sub-analyses, preoperative rupture was associated with a worse prognosis, and intraoperative rupture had a poorer PFS than no rupture.

Surface of the specimen

Why is it so important?

Pretty much the same reasons as for the integrity...



- IC Tumour limited to 1 or both ovaries or fallopian tubes, with any of the following:
 - IC1 Surgical spill
 - IC2 Capsule ruptured before surgery or tumour on ovarian or fallopian tube surface
 - IC3 Malignant cells in the ascites or peritoneal washings

Surface aspect:

Smooth Roughened Adhesions Excrescences Cyst





Surface of the specimen

Why is it so important?



- Should we ink the surface?
- IC Tumour limited to 1 or both ovaries or fallopian tubes, with any of the following:
 - O IC1 Surgical spill

Pretty much the same reasons as for the integrity...

IC2 Capsule ruptured before surgery or tumour on ovarian or fallopian tube surface

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Surface aspect:

Smooth Roughened Adhesions Excrescences Cyst



Opening of the specimen !!!

Broadly speaking two main appearances/aspects





Opening of the specimen !!!

Cystic

Unilocular or multilocular?



Septations separating locules

Thickness of the cyst wall: thin and delicate, thick and irregular

Content? Clear/watery/serous Blood-filled Old blood « chocolate » Gelatinous/mucoid Colloïd





Opening of the specimen !!!

Cystic

Unilocular or multilocular?



Content?

Keratinous debris Hair





Opening of the specimen !!!

Examine the inner surface









Opening of the specimen !!!

Examine the inner surface



BUT WHY ????

Cystic







Opening of the specimen !!!

Examine the inner surface





Cystic



We all know that this is rather benign

Opening of the specimen !!!

Examine the inner surface, why? Somehow you already know

Cystic

Excrescences you see grossly will translate by cellular proliferation that will define the diagnosis of malignancy !

Opening of the specimen !!!





Cystic

How to sample?





Sample both thin walled part and excrescences

(to illustrate continuum of the disease)

- Focus on breaches of the capsule, adhesion
- If small: embed entire tumour
- If <10cm: at least one block per cm
- If > 10cm: 2 blocks per cm

Opening of the specimen !!!



Cystic

- Trick to sample thin walled cyst
- Swiss roll technique, a bit like with placental membrane
 - Carefull embedding on the edge to make sure to analyze the full

thickness of the cyst wall





Opening of the specimen !!!





Solid

Debulking

Definition:

When cancer is at an advanced stage, there is spreading in the peritoneal cavity and on the surface of viscera

Aim of the surgeon:

If cancer is obvious: to remove as much cancer as possible

If cancer is not obvious: to make sure there's no microscopic spreading and evaluate staging

Long and tedious surgery and long and tedious gross examination

Type of specimen in a debulking (beside hyterectomy with BSO):

- Omentum
- Patches of peritoneal membrane sampled by the surgeon at different level of the peritoneal cavity
- Peritoneum covering diaphragm, liver, guts, pelvic floor....



Peritoneal

Debulking - Omentum

Definition:

Apron like structure of adipose and fibrous peritonealized tissue that is covering the viscera

Weight Measure

Describe if present metastatis deposit and their SIZE

20 mm limit

IIIA2 Microscopic extrapelvic (above the pelvic brim) peritoneal involvement with or without positive retroperitoneal lymph nodes

IIIB Macroscopic peritoneal metastasis beyond the pelvis up to 2 cm in greatest dimension, with or without metastasis to the retroperitoneal lymph nodes

IIIC Macroscopic peritoneal metastasis beyond the pelvis more than 2 cm in greatest dimension, with or without metastasis to the retroperitoneal lymph nodes (includes extension of tumor to capsule of liver and spleen without parenchymal involvement of either organ)



Debulking - Omentum

Definition:

Apron like structure of adipose and fibrous peritonealized tissue that is covering the viscera

Weight Measure

IF omentum is macroscopically normal after carefull inspection and palpation of cut section:

5 to 6 blocks are necessary



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Debulking – Peritoneal biopsy /patches

Simple specimen

No orientation, no anatomical structures to consider

Measure

DESCRIBE, if nodule suspicious for tumor deposit : MEASURE (20mm cut off)

How to sample ?

General principal:

If you see something obvious macrocopically > Your gross examination is the most important. Sample the largest nodule

If you don't see anything > Microscopy is going to be the most important. Entire embedding is legit if number of blocks is reasonnable.



References (free access):



- <u>https://www.rcpa.edu.au/Manuals/Macroscopic-Cut-Up-</u>
 <u>Manual/Gynaecology-and-perinatal/Ovary-and-fallopian-tube-benign-setting</u>
- https://www.iccr-cancer.org/datasets/published-datasets/femalereproductive/ovary-ft-pp/

Data element	Response							
Fresh tissue received	No	Yes	If yes, descri	be any addit	ional tests/	frozen secti	ons/biob	anking performed
Specimen labelled as	Text	As stated by the clinician						
Specimen type (select all that apply)	Right	Right ovary Left ovary Right ovarian cystectomy Left ovarian cystectom						ian cystectomy
	Bight fallopian tube Left fallor			pian tube Literus Cervix				
	Omentum		Paritonnal hippsier		Peritoneal washings/ascitic fluid			uid .
	lump	a nodor u	posify number and site/s		Other specify (a a howal bladder appendix)			
The second se	Cymph nodes, specify number and s				, other, specify (e.g. bower, biduder, uppendix)			
specimen integrity	Ovaria	fragmen	tad spaciman	Overse oth	other specify			
	Eallon	ian tuhe	serosa intact	Fallonian	Fallonian tube tumour on serosal surface			
	Fallopian tube fragmented Fallopian tube other, specify							
	Fallopian tube fimbriae Not identified Present Fallopian tube description							
Specimen weight	g This will primarily be the ovary/adnexal mass weight State what is included							
Specimen dimensions	Length x width x mm If abnormal length x width x thickness x x mm							mm
	Uterus: superior to inferior x distance between cornu x anterior to posteriorx_x_mm							
Number of tumours	1							
For each turnour: (if >1 designate accordingly)								
Tumour dimensions	Length x width x thickness _x_x_mm							
Tumour description(s)	Text							
Macroscopic description of omentum								
Omentum dimensions	Length x width x thickness _ x _ x _ mm							
Omental Involvement	Involved Not involved							
Maximum dimension of largest deposit	mm	1						
Number of metastatic deposit(s)								
Size of metastatic deposit(s)	_mm							
Macroscopic tumour site	Left o	vary F	Right ovary	Left fallop	ian tube	Fimbrial		Non fimbrial
	Right	fallopian	tube Fimbri	al No	n fimbrial	Perito	neum	Other, specify
Lymph nodes	Not re	ceived	Received	Describe	Rec	ord number	per cass	ette
Other macroscopic information	Text	Text						
Block identification key	Text							



Thank you for your attention



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