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Summary file

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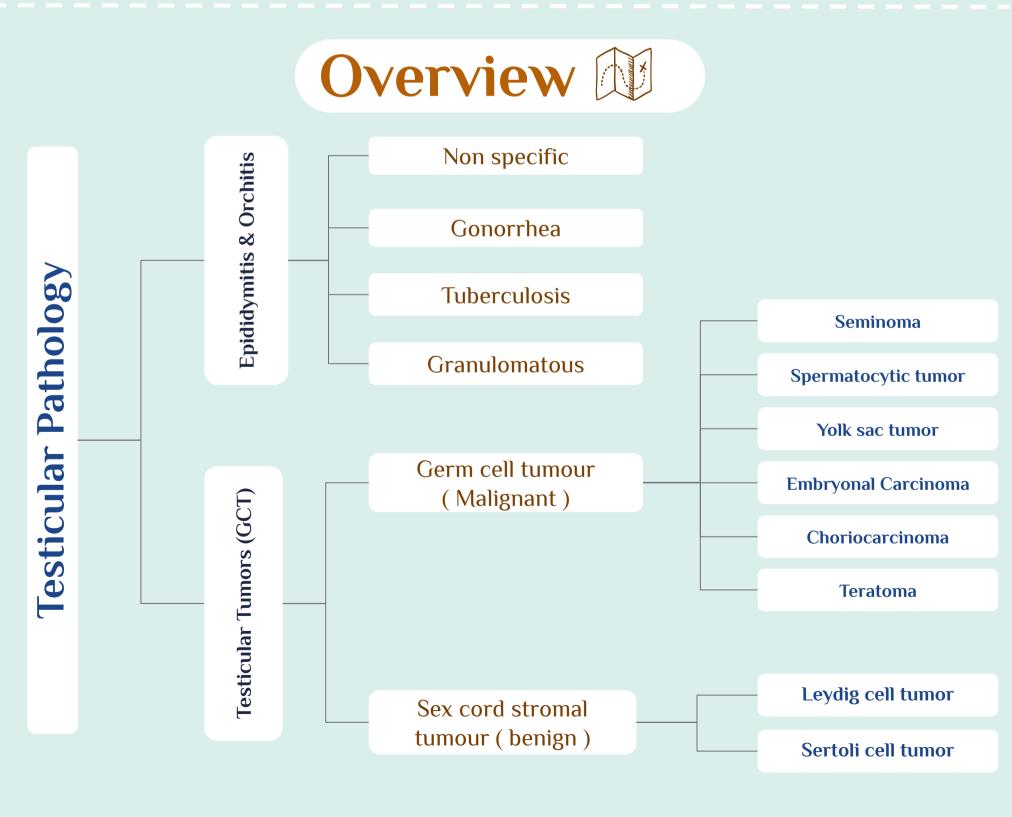
اللهم لا سهل الا ماجعلته سهلا و انت تجعل الحز ن إذا شئت سهلا





Know the predisposing factors and pathology of Epididymitis and orchitis:
-Non specific Epididymitis and orchitis
-Granulomatous/Autoimmune Orchitis
-Gonorrhea
-Tuberculosis

Be familiar with the basic classification and pathology of testicular tumors: Seminoma Yolk sac tumor Embryonal carcinoma Teratoma Choriocarcinoma



02

01

Epididymitis and orchitis

Introduction

- Epididymitis: Inflammation of epididymis
- Orch<u>itis</u>: Inflammation of testis
- Inflammatory conditions are more common in the epididymis than in testis.
- Some infections (e.g. Syphilis), may begin in testis with secondary involvement of epididymis.

Non specific¹ Epididymitis and Orchitis

Pathology

- Commonly related to **urinary tract infections** (cystitis, urethritis, and genitoprostatitis).
- Infections reach the epididymis/testis through:
 - > Vas deferens
 - > The **lymphatics** of spermatic cord

Causative organisms

- Children: Uncommon, but usually associated with Gram -ve rods and a congenital genitourinary abnormality.
- Men younger than 35: Chlamydia trachomatis and Neisseria gonorrhoeae.
- Men older than 35: E.Coli and pseudomonas.

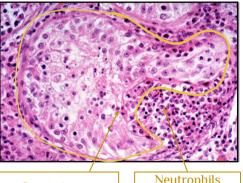
Microscopic findings "Uncommon to have samples, unless aggressive"

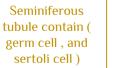
- Vascular congestion, edema, and infiltration by neutrophils, macrophages and lymphocytes.
- Initially involves interstitium but later involves the seminiferous tubules.
- May progress to frank abscess يتحول من الدلتهاب الى خراج, and heals by fibrous scarring. (in case resolution through antibiotics has not been achieved)
- ✤ fibrosis² (testis cannot regenerate)
- ✤ Leydig cells are not usually destroyed

Clinical features

- $\boldsymbol{\diamond}$ cardinal sign of inflammation (redness , heat , pain loss of function)
- Swelling of the organ , Painful nodule and tenderness

ORhitis





attack the seminiferous tubule + micrabses

1. Secondary involvement of the testis or epididymis.

2. Tissue will not be renewed and replaced.

Epididymitis and orchitis

Specific Epididymitis and Orchitis

Gonorrhea sexual transmitted disease

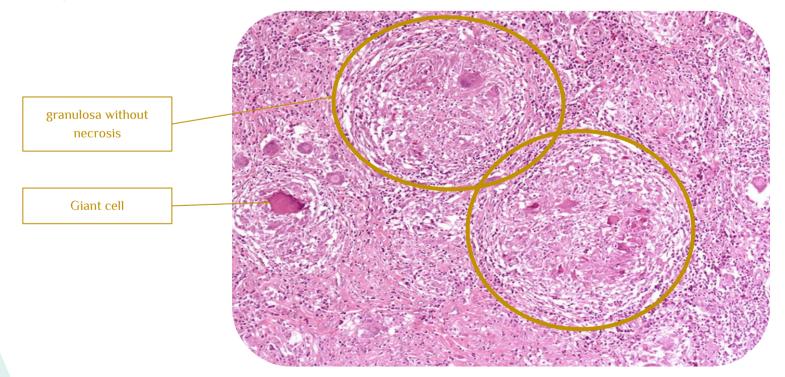
Gonococcal infection can spread from urethra to prostate, seminal vesicles then to epididymis/testis leading to suppurative orchitis (pus = قيح) and even abscess formation (collection of pus)

Tuberculosis

- It begins in epididymis and spreads to testis.
- There is associated tuberculous prostatitis and seminal vesiculitis.
- Microscopy: Caseating granulomas (granuloma with central necrosis), this feature characteristic for M.T in any organ
- **It mimics testicular tumors, and may accidentally undergo surgical excision if not diagnosed correctly.**

Granulomatous (autoimmune)

- ***** It affects **middle-aged men**.
- Present with unilateral testicular mass. mimics testicular tumor (may also accidentally undergo surgical excision).
- May be in response to **disintegrated sperm**¹, **post-infectious**, due to **trauma or sarcoidosis**².
- ♦ Microscopy: "No biopsy → risk of leakage"
 - Granulomatous³ (granuloma without cassation) inflammation with plasma cells and lymphocytes , gait cell .



1. Caused by injury to the seminiferous tubules \rightarrow sperms are out \rightarrow autoimmune reaction.

2. Autoimmune disease.

3. Culture to rule out TB, because they are similar microscopically.

Testicular tumors

Classification of testicular tumors

	Germ cell tumors 95% of testicular tumor in adults, <mark>malignant</mark>	Sex cord stromal tumors Uncommon and usually benign.
*	 Tumors with one histologic pattern (Pure form): Seminomatous germ cell tumors: Seminoma Most common type in all testicular tumors (50%) Spermatocytic seminoma Less common, older age >65 Nonseminomatous germ cell tumors (NSGCT) : (malignant) Embryonal carcinoma Yolk sac tumor Choriocarcinoma Teratoma: can be mature or immature (malignant transformation) Tumors with more than one histologic pattern (mixed form): mixed germ cell tumor 	 ✤ Leydig cell tumor ❖ Sertoli cell tumor

Clinical features :

-Inflammation of testis usually associated with pain and the cardinal inflammatory sign -Testicular Tumor usually painless , solid mass

Germ cell tumors

- ***** Testicular tumors are the most important cause of a **firm**, **painless enlargement** of testis.
- Peak incidence is between the age of **15 and 34 years**.
- Highly aggressive cancers, capable of extensive dissemination.
- With Current therapy most of them can be cured. (Responding well to treatment)
- Germ cell tumors may have:
 - ➤ Single component (Pure form).
 - Mixture (40% of cases) e.g. mixture of seminomatous and nonseminomatous components.
- Most GCTs originate from precursor lesion called intratubular germ cell neoplasia (it's like carcinoma-in-situ), except Spermatocytic tumors (not associated with precursor lesion) and we have to check pediatric patients for this lesion too.

Risk factors of all germ cell tumors

- Cryptorchidism: 3 to 5 fold increase in the risk of cancer in both undescended testis and contralateral descended testis.
 - > About 10% cases of testicular cancer have cryptorchidism.
- Testicular dysgenesis.
- Cenetic factors. (e.g. Klinefelter syndrome).
- Strong family predisposition: brothers, fathers and sons are at risk.
- Personal history : If Contralateral testis has cancer.
- **Race**: more common in whites than in blacks.

Germ cell tumors

Seminomatous germ cell tumors

1- Seminoma

- **Most common type** of testicular tumors.
- Most common type of germ cell tumors (50%).
- Identical tumor occurs in the ovary (called dysgerminoma).
- Secretes lactate dehydrogenase LDH.
- Peak incidence is between 40–50 years of age. Almost never occur in infants. (it is a bit higher than other GCT)
- Classic seminoma is **highly sensitive to radiation therapy**, overall 5-year survival is 90%-95%.

Morphology

Gross

- Enlarged Bulky masses, sometimes very large.
- Homogenous, gray-white, lobulated cut surface.
- Large tumors may contain foci of coagulative necrosis, usually without hemorrhage



Microscopic * Sheets of uniform cells (undifferentiated¹ germ cells) ✨ Divided into lobules by delicate fibrous septa containing lymphocytes (characteristics feature) Cells are large and round with large nucleus * and prominent nucleoli. Clear Cytoplasm of tumor cell because * containing of glycogen (Appears white and vacuolated). Tumor cells are positive for stains: PLAP, OCT4, and c-kit $(CD117)^2$ "these stains are special in Seminona, they help with diagnosis '

Lobules separated by fibrous septa that contain Lymphocytes

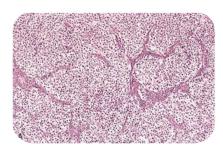
2- Spermatocytic tumor

- It was called previously spermatocytic seminoma.
- Uncommon: 1-2 % of testicular GCTs
- ✤ Men >65 years of age.

Prognosis

- Slowly growing tumor that does not metastasize.
- ✤ Not associated with intratubular germ cell neoplasia. (Precursor)
- Prognosis is excellent

Important feature to differentiate it from other germ cell tumors.
 These are stains used to diagnose certain types of tumors.



Fried egg Seminoma appearance = prominent nuclei with clear cytoplasm.

Non-seminomatous germ cell tumors

	Embryonal carcinoma (15 to 35% of testicular GCTs)				
Introduction	 Age group: 20 to 30 years Can be seen combined with other GCTs (in mixed GCTs). 				
Marker	Tumor cells are positive for cytokeratin (CK) and CD30 stain				
Morphology	Gross: smaller than seminomas and poorly demarcated.				
	 Microscopic: variegated surfaces with foci of <u>necrosis</u> and <u>hemorrhage</u>. 				
Prognosis	 More aggressive than seminomas Metastasizes early via both lymphatic and hematogenous routes New chemotherapeutic agents are very effective and greatly improve prognosis 				
Treatment	Treatment				
	Yolk sac tumors				
Introduction	 Yolk sac tumor is also called endodermal sinus tumor. Occur in two forms : Pure form: seen in young children, rare in adults. Combination: with other NSGCTs seen in adults, commonly mixed with embryonal carcinoma. 				
Prevalence	It is the most common tumor in infant and children up to 3 years of age with a very good prognosis				
Diagnosis	 Patients have elevated <u>serum alpha fetoprotein (AFP)</u> Used as a marker of disease progression and diagnosis. (helps in post treatment follow-up in case any recurrence happens) Tumor cell are positive for AFP and alpha-1-antitrypsin stain. 				
	 Gross: non encapsulated, homogenous, yellow white, mucinous. Not specific 				
Morphology	 ♦ Microscopic: > Tumor shows structures resembling endodermal sinuses called as Schiller-Duval bodies (characteristic features) > Hyaline-pink globules 				
Treatment	The biologic behavior of YST is similar to that of embryonal carcinoma. (But responds well to chemotherapy)				

Choriocarcinoma			
Introduction	 It is a highly malignant tumor, rare Pure form is extremely rare and it is the most aggressive non-seminomatous tumor. More common in female It is more common as a component of mixed GCT. Tumor cells are positive for human chorionic gonadotropin HCG stain "Helps in diagnosis" 		
Morphology	✤ Gross: small sized lesions.		
	 Microscopic: Malignant trophoblastic (placental) tissue: cytotrophoblastic and syncytiotrophoblastic cells. Prominent <u>hemorrhage</u> and <u>necrosis</u>. 		

Teratoma					
Introduction	 Composed of various different types of cells or organ components. They can occur at any age. Pure form: Second most common in infants and children after yolk sac tumor. VERY COMMON IN FEMALE Mixed form: usually in adults. 				
Morphology	Gross: Large (5 -10 cm). Solid and cystic areas. Heterogeneous: bizarrely distributed collection of different type of cells or organ structures (e.g: bone, cartilage and teeth). Microscopic: Any cell type can be present: neural, cartilage, bone, squamous epithelium, hair, glandular cells, smooth muscle, thyroid tissue, bronchial epithelium of lung, pancreatic tissue etc. 				
	A. Mature	If the cells/tissue is mature looking.			
	B. Immature ¹	If some of the cells/tissue component is immature.			
	Teratoma with malignant transformation	lf any of the cells/tissue undergoes non germ cell type of malignant transformation (e.g. squamous cells develop into squamous cell carcinoma).			
Prognosis	 ◆ Prepubertal males → benign. (mature only, immature is always malignant) ◆ Postpubertal males → malignant (regardless of maturity, unlike females) ◆ Mature and immature teratomas are both capable of metastasis. 				

Germ cell tumors

Mixed Germ Cell Tumors (GCTs)

- Common. Half of testicular tumors are composed of a mixture of GCTs.
- The common combinations or mixtures are:
 - Seminoma + embryonal carcinoma.
 - ➤ Teratoma + embryonal carcinoma +/- yolk sac tumor.

Clinical Features

- Painless solid enlarging mass in the testis
- They can secrete hormones and enzymes that can be detected in blood:
 - > HCG (in choriocarcinoma) , AFP (yolk sac tumor) , and lactate dehydrogenase LDH (seminoma)
- GCTs can metastasize by:
 - > **Direct** extension to the epididymis, spermatic cord, or scrotal sac.
 - > Lymphatic spread: Retroperitoneal and para-aortic nodes are first to be involved.
 - Hematogenous spread to Lung, liver, Brain, and bones
- A biopsy of a testicular tumor is not recommended because it is associated with a risk of tumor spillage.
 Using biopsy in the diagnosis is not recommended because it is painful and enhances the spread of tumour

Management

- The standard management: radical orchiectomy.
 - > Seminomatous tumors \rightarrow radiosensitive: respond well to radiotherapy.
 - 95% of patients can be cured.
 - > Non-seminomatous tumors \rightarrow chemosensitive: respond very well to chemotherapy.
 - 90% of patients achieve complete remission with aggressive chemotherapy.
 - > Pure choriocarcinoma has a poor prognosis (more common in females)

Seminomas	Nonseminomatous GCT
Radiosensitive	Non radiosensitive
Chemosensitive	Chemosensitive
Late metastasis	Early metastasis to retroperitoneal lymph nodes.
Excellent prognosis	More aggressive



Epididymitis and orchitis

Non specific	 Related to urinary tract infections. Causative organism: children (gram -ve rods), men younger than 35 (Chlamydia trachomatis and Neisseria), men older than 35 (E Coli and pseudomonas). Microscopy: Congestion and infiltration by neutrophils, macrophages & lymphocytes.
Gonorrhea	Spread from urethra to epididymis and testis leading to orchitis and abscess.
Tuberculosis	 Begins in epididymis and spread to testis. Microscopy: Caseating granuloma.
Granulomatous	 Mimic testicular tumor (unilateral mass). Microscopy: Granulomatous inflammation with plasma cells and lymphocytes.

Testicular tumors			
Seminomatous germ cell tumors			
Seminoma	 Most common type of testicular tumors and germ cell tumors (50%). Age group: peak incidence in 30s, almost never occur in infants. Secretes lactate dehydrogenase LDH 		
Spermatocytic tumor	 Uncommon: 1-2% of testicular GCTs It affects men over the age of 65 years. 		
Non-Seminomatous germ cell tumors			
Embryonal carcinoma	 They account for about 15% to 35% of testicular GCTs. The age group: 20 to 30 years. They are more aggressive than seminomas. Tumor cells are positive for cytokeratin (CK) and CD30 stain. 		
Choriocarcinoma	 It is a highly malignant tumor. Patients have elevated serum human chorionic gonadotropin (HCG). 		
Yolk sac tumor	 It is the most common tumor in infant and children up to 3 years of age and it has a very good prognosis in infants and children. In adults it occurs as mixed GCT. Elevated serum alpha fetoprotein (AFP) Microscopy: Schiller-Duval bodies. 		
Teratoma	 It is a tumor composed of various different types of cells or organ components. They can occur at any age Benign in prepubertal males and malignant in postpubertal males. Microscopy: it can be mature, immature, or with malignant transformation. 		
Mixed GCTS	• Common, half of testicular tumors are composed of a mixture of GCTs.		



MCQs

 $01\,|\text{A}\,32$ year old male present with testicular enlargement, microscopy findings shows granulomatous inflammation with plasma cells and lymphocytes, what's the most likely diagnosis?

A)Orchitis		B) Semino) Seminoma C) Embryonal carcinoma			ioma D	D) Urethritis		
appears solid orchiectomy	and homogen is performed, a	eous on ultras	ound examina al specimen sh	tion. No tumo ows Bulky m	d 2 weeks ago. T or markers are d asses with homo	etected on se	rologic testing.	An	
A) Choriocard	cinoma	B) Embry	onal carcinoma	a C) L	mphoma	D	D) Seminoma		
03 Which of	the following	is the most agg	gressive type o	f testicular t	umors?				
A) Seminoma		B) Sperma	atocytic tumor	C) C	horiocarcinoma	D	D) Teratoma		
04 Which of	the following	is the most co	nmon testicula	ar tumor in ir	fants and childr	ren?			
A) Yolk sac tu	Imor	B) Embry	onal cell Carci	noma C) S	eminoma	D	D) Teratoma		
05 A-26-yea is most likely		me to the hosp	ital with a righ	t testicular m	ass and history	of cryptorchi	dism. which of t	the following	
A) Sex cord tumors.		B) Germ o	B) Germ cell tumors.		C) Tuberculoma of the intrascrotal cord.		D) Vascular aneurysm.		
treatment wa		e. Later investig			patient was tre and CD30 posi		• •		
A) Yolk sac ce	ells	B) Semino	oma	C) E	nbryonal carcin	ioma D) Choriocarcino	oma	
examination, solid mass wi	the left testis i thin the body	is three times t	he size of the s. Laboratory	right testis ar studies inclue	in his scrotum f id is firm on pal le an elevated so s?	pation. An ulti	asound scan sh	nows a 6-cm	
A) Cytotrophoblasts B) Emb			onal carcinoma	a cells C) S	eminoma cells.	D	D) Yolk sac cell		
mass within t soft. A retrop composed of	he right testis eritoneal lymp cuboidal cells	. A right orchie oh node dissect intermingled w	ctomy is perfo tion is done. M vith large eosir	rmed, and gr icroscopic ex iophilic syncy	ne past 3 month oss examination amination show /tial cells contai ositive for which	shows the m s that areas o ning multiple	ass to be hemor f viable tumor a dark, pleomorp	rrhagic and are	
A) α-Feto	oprotein	B)CD20		· · · · · · · · · · · · · · · · · · ·	C)Carcinoembryonic antigen		D)Human chorionic gonadotropin.		
	01	00	07				07	0.0	
MCQs Answer	01	02	03	04	05 P	06	07 D	08	
key	A	D	C	A	B	C	D	D	

key

