

Testicular cancer

- The testis developed in the retroperitoneum , around the T10... Reach the scrotum in the ninth month
- Supplied by collateral from **cremasteric artery** and **artery to ductus deferens**.
- Venous drainage ; from the right testis to **IVC**, from the left testis to **left renal vein**.
- Lymphatic drainage to **para-aortic LN**, epididymis, spermatic cord, and scrotum to **Inguinal LN** .

→ RFs :- **15- 35 yrs** , **Family history**, white, **Cryptorchidism**, CA in one testis, Intersex syndromes, Trauma, HIV, recurrent infections, Maternal estrogen ingestion.

→ Presentation : painless, painful if with necrosis or hemorrhage, Unilateral , more in Rt one

→ Physical examination :- enlarged testis, nodularity, firm to hard consistency, Secondary hydrocele, flat and difficult to feel epididymis , Examination for mets, gynecomastia?

Non-Germinal cell ca

Leydig cell ca

- **MC**
- Not with cryptorchidism.
- Produce **androgens**
- Precocious puberty,
- Gynecomastia, low libido
- **Painless**
- **bilateral orchiectomy + RPLND**

Sertoli cell ca

- Any age,
- Not with cryptorchidism
- Excess **estrogen**
- large cell calcifying / sclerosing
- **10% malignant**
- **bilateral orchiectomy + RPLND**

Gonadoblastoma ca

- Mixed germ cell, sex cord, stromal ca.
- within dysgenetic gonads.
- **IS malignant >> GCT**
- **Bilateral orchidectomy**
- ~ radical to the affected one

Germinal cell ca 90%.. more Malignant

Seminoma ~ 48%

- MC GCT
- 25-35 y/o
- Unilateral , Right > left
- Rare in prepubertal
- progressive painless enlargement of the testis, histologically identical to ovarian dysgerminomas —> **Pale** and **homogenous** , **Soft, well-demarcated**, without hemorrhage / unlike NSGCTs
- Secrete **B-hcg** , never secrete AFP
- types; classic, anaplastic, spermatocytic
- Treatment: **radical inguinal orchiectomy + RTX** / sensitive

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|--|
| o Stage 1 : bilateral inguinal orchiectomy+RPLND +/-RTX |
| o Stage 2a/2b : bilateral inguinal orchiectomy +RPLND+/-CTX |
| o Stage 2c/3 : bilateral inguinal orchiectomy +high doseCTX |

Non Seminoma 42%

| Embryonal | Yolk sac | Teratoma / 3 G.cells | Choriocarcinoma |
|---|--|--|--|
| 25-35 Secretes AFP and β-hCG | Infants and children Secretes AFP and β-hCG | 25-35 Does not secrete AFP nor β-hCG | 20-30 Always secretes β-hCG and never secretes AFP |
| | Hematogeneous spread <i>AFP > 1000 ng/ml ↑ kumer ←</i> | <i>• Benign in children • Malignant in Adult</i> | Hematogeneous spread, especially to <u>lungs</u> and <u>liver</u> . No LN involvement (the only one) |
| Poor response to CTX and RTX | Most common tumor in infants and children | | WORST PROGNOSIS |

- Higher metastasis
- ill-defined, invasive masses. containing foci of H & N
- AFP level correlates with disease extent.
- **Schiller-Duval bodies** "glomeruloid"
- poor response to Cx / Rx
- Rare & the **worst**
- small, no testicular enlargement.

→ Mets

- o abdominal and lumbar pain.(GI) chest pain, hemoptysis, dyspnea .(lungs)
- o jaundice (liver) hydronephrosis.(kidneys)
- o troiser's sign.(palpable left supraclavicular LN)

→ Ddx : Hydrocele, Spermatocele , Hematoma, Hernia, Torsion, Epididymitis, TB

→ Investigations:-

- Labs: CBC, LFT, KFT, LDH
- Serum assay: **AFP, B-HCG** & for follow up after surgery
- Radiology: chest x-ray, CT for abdomen and pelvis, ultrasound for contralateral testis.
- Histopathology always should be done.

| α-fetoprotein | B-hCG | LDH |
|---|---|--|
| Normally expressed by early embryo, liver and GIT Expressed by <u>embryonal</u> and <u>yolk sac</u> tumors | Normally produced by placenta Expressed by 100% of <u>choriocarcinomas</u> , 40% of <u>teratomas</u> , and 10% of <u>seminomas</u> <i>All except Teratoma</i> | Normally present in smooth muscle cells, cardia, SKM, liver and bones Expressed in <u>seminomas</u> |
| Elevated AFP suggests non-seminomatous element | | Mainly used to determine tumor <u>burden (size)</u> . High tumor burden is usually associated with high levels of LDH. |
| Other causes of increased AFP: liver dysfunction, viral hepatitis, alcohol | Other causes of increased B-hCG: hypogonadism and marijuana | Elevated in serum due to various reasons, so not specific and carries high risk of false positive |

- Any **solid testicular mass** should be managed as malignancy until proven otherwise and

histopathology should always be done. → *After Resection to determine T-stage*

Radical orchiectomy : testis, epididymis and spermatic cord, with Sperm cryopreservation

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|------|--|
| pTX | Primary tumor cannot be assessed. |
| pT0 | No evidence of primary tumor (e.g., histologic scar in testis). |
| pTis | Intratubular germ cell neoplasia (carcinoma <i>in situ</i>). |
| pT1 | Tumor limited to the testis and epididymis without vascular/lymphatic invasion ; tumor may invade into the tunica albuginea but not the tunica vaginalis. |
| pT2 | Tumor limited to the testis and epididymis with vascular/lymphatic invasion , or tumor extending through the tunica albuginea with involvement of the tunica vaginalis. |
| pT3 | Tumor invades the spermatic cord with or without vascular/lymphatic invasion. |
| pT4 | Tumor invades the scrotum with or without vascular/lymphatic invasion. |
| NX | Regional lymph nodes cannot be assessed. |
| N0 | No regional lymph node metastasis. |
| N1 | Metastasis with a lymph node mass ≤2 cm ; or multiple lymph nodes, none >2 cm in greatest dimension. |
| N2 | Metastasis with a lymph node mass >2 cm but not >5 cm ; or multiple lymph nodes, any one mass >2 cm but not >5 cm in greatest dimension. |
| N3 | Metastasis with a lymph node mass >5 cm in greatest dimension. |
| M0 | No distant metastasis. |
| M1 | Distant metastasis. |
| M1a | Nonregional nodal or pulmonary metastasis. |
| M1b | Distant metastasis other than to nonregional lymph nodes and lung. |

Respond for All ... **LN & Follicle**
 Surgical or chemo ... **No Radio**

| Stage | Histology | Treatment | 5 Year Survival Rate |
|-----------------------|-------------|--|----------------------|
| I | Seminoma | Orchiectomy, radiation therapy | 97% |
| I | Nonseminoma | Orchiectomy, RPLND, surveillance 1 year | 95% |
| II as I + Chx | Seminoma | Nonbulky Tumor: Orchiectomy and radiation therapy Bulky Tumor: Orchiectomy and combination chemotherapy (cisplatin-based regimen) or by radiation therapy | 90% 70% |
| | Nonseminoma | Orchiectomy and RPLND, followed by combination chemotherapy (cisplatin, bleomycin, etoposide) | 95% |
| III orchiect & M. Chx | Seminoma | Orchiectomy and multidrug chemotherapy (cisplatin, bleomycin, etoposide) | 95% |
| | Nonseminoma | Orchiectomy and multidrug chemotherapy (cisplatin, bleomycin, etoposide) | 70% |