

# Testicular Tumors

Tutorial handout for 4<sup>th</sup> year students in Alkindy Collage of Medicine / University of Baghdad

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## Epidemiology:

- ü They are the most common solid malignancy in men aged between 20 and 45 years.
- ü The peak incidence for NSGCT is 25 years and seminomas is 35 years.
- ü White : Black = 3:1
- ü Patient with Cryptorchidism are 3-14 times more likely to develop testicular cancer than normal individual
- ü Family history, maternal estrogen ingestion & HIV infection are risk factors for testicular carcinoma

## WHO histopathological classification of testicular tumors:

- ✓ Germ cell tumors (90%)
  - Ø Seminoma(48%)
    - ü Spermatocytic, classical, and anaplastic subtypes
  - Ø Non-seminomatous GCT (42%)
    - ü Teratoma
      - § Differentiated/mature
      - § Intermediate/immature
      - § Undifferentiated/malignant
    - ü Yolk sac tumor
    - ü Choriocarcinoma
    - ü Mixed NSGCT
  - Ø Mixed GCT(10%)
- ✓ Sex cord stromal tumors (3%) (10% malignant)
  - ü Leydig cell
  - ü Sertoli cell
  - ü Mixed or unclassified
- ✓ Mixed germ cell/sex cord tumors (rare)

✓ Other tumors (7%)

- ü Epidermoid cyst (benign)
- ü Adenomatoid tumor
- ü Adenocarcinoma of the rete testis
- ü Carcinoid
- ü Lymphoma (5%)
- ü Metastatic, from another site (1%)

Presentation:

- § Painless lump 86%
- § Pain 31%
- § Dragging sensation 29%
- § Secondary Hydrocele
- § Gynaecomastia from B-HCG production
- § O/E: The normal side is first examined, followed by the abnormal side. This may reveal asymmetry or slight scrotal skin discoloration with hard, non-tender, irregular, non-transilluminable mass in the testis or replacing the testis

Spread:

- ü Hematogenous: to the liver, lung, bone and brain (Teratoma).
- ü Lymphatic: to para- aortic nodes and produce back pain (Seminomas).
- ü Direct: through tunica albuginea and tunica vaginalis to the scrotal skin.

TNM staging of testicular tumors:

✓ T—Primary tumor

- ü TX: Cannot be assessed
- ü T0: No evidence of primary tumor
- ü Tis: Intratubular germ cell neoplasia (CIS)
- ü T1: Limited to testis and epididymis, no vascular invasion
- ü T2: Invades beyond tunica albuginea or has vascular invasion
- ü T3: Invades spermatic cord
- ü T4: Invades scrotum

✓ N—Regional lymph nodes

- ü NX: Cannot be assessed

- ü N0: No regional lymph node metastasis
- ü N1: Lymph node metastasis  $\leq 2$  cm, or multiple nodes, none more than 2 cm. and  $< 6$  nodes positive
- ü N2: nodal mass  $> 2$  cm and  $\leq 5$  cm. or  $\geq 6$  nodes positive
- ü N3: Nodal mass  $> 5$  cm.
- ✓ M—Distant metastasis
  - ü MX: Cannot be assessed
  - ü M0: No distant metastasis
  - ü M1: Distant metastasis present in nonregional lymph nodes or lungs
  - ü M2: Nonpulmonary visceral metastases
- ✓ S—Serum tumor markers
  - ü SX: Markers not available
  - ü S0: Marker levels within normal limits
  - ü S1: (LDH)  $< 1.5 \times$ normal and hCG  $< 5000$  mIU/mL and AFP  $< 1000$  ng/mL
  - ü S2: LDH  $1.5\text{--}10 \times$ normal or hCG  $5000\text{--}50,000$  mIU/mL or AFP  $1000\text{--}10,000$  ng/mL
  - ü S3: LDH  $> 10 \times$ normal or hCG  $> 50,000$  mIU/mL or AFP  $> 10,000$  ng/mL

### Differential Diagnosis of Scrotal mass:

#### *Painful mass:*

- ü *Epididymitis/orchitis; bacterial, STD, mumps, tuberculosis*
- ü *Incarcerated/strangulated hernia*
- ü *Testicular trauma: usually blunt; contusion, rupture; usually associated hematocele.*
- ü *Torsion (testicle, testicular or epididymal appendage)*
- ü *Tumor (pain infrequent unless traumatized or rapidly growing)*

#### *Painless mass:*

- ü *tumor of testis, epididymis, rete testis or tunica vaginalis*
- ü *Lipoma or hydrocele of the cord*
- ü *Other scrotal condition like Hydrocele, haematocele, Chylocele and Scrotal edema*
- ü *Sperm granuloma following vasectomy*
- ü *Spermatocele*

ü *Leukemia or lymphoma*

ü *Varicocele*

### Investigations:

Patients should be referred urgently and seen within 2 weeks if malignancy is suspected:

- Ø Ultrasound of Scrotum.
- Ø Tissue histology can follow an Inguinal Orchiectomy.
- Ø Disease can be staged by thoraco-abdominal CT scanning.
- Ø Tumour markers are useful in staging and assessing response to treatment.
  - ü Alpha-fetoprotein (AFP): is produced by yolk sac elements but not produced by seminomas.
  - ü Beta human chorionic gonadotrophin ( $\beta$ -HCG) is produced by trophoblastic elements and so may be elevated in both teratomas and seminomas.
  - ü Lactate dehydrogenase (LDH): less specific, correlate with tumor burden, and is most useful in monitoring treatment response in advanced seminoma.

### Treatment:

dependant on type of tumour and stage

### Seminomas:

Seminomas are radiosensitive:

Removal of primary tumour by Inguinal (Radical) orchiectomy plus:

- ü Stage I and II disease treated by inguinal orchiectomy plus radiotherapy to ipsilateral abdominal and pelvic nodes or surveillance
- ü Stage IIC and beyond are treated with chemotherapy (often cisplatin, etoposide and bleomycin - BEP)
- ü Surveillance
- ü Tumour markers are less reliable.

### NSGCT:

*NSGCT are not radiosensitive:*

Removal of primary tumour by Inguinal (Radical) orchiectomy plus:

- ü Chemotherapy for any who relapse or have metastasis at presentation (cisplatin, bleomycin and etoposide - BEP is standard regimen)

ü Surveillance

ü Tumour markers are very important.