A Private Investigation: Radiologic-Pathologic Correlation of Testicular Tumors
Disclosure

None of the authors have conflicts of interest to disclose
Learning Objectives

1. Review sonographic findings of seminoma and nonseminomatous tumors of the testis, as well as less common tumors including lymphoma, epidermoid cyst and gonadal stromal tumor.

2. Direct comparison of sonographic findings with gross and histologic pathology findings.

3. Discuss pearls and pitfalls in accurately diagnosing testicular tumors.
Testicular Tumors

• **Demographics**
  - 1% of all solid tumors in males.
  - Most common male solid tumor malignancy between 15-35 years.
  - Most common are germ cell tumors (95%) followed by sex cord-stromal tumors.

• **Risk factors**
  - Cryptorchidism
  - History of prior testicular malignancy
  - Age (20-34) and ethnicity (Whites)
  - Infertility
  - Intersex syndrome
  - HIV infection
  - Family history
Classification

- **Germ-cell tumors**
  - Seminoma
  - Nonseminomatous germ cell tumor (GCT)
    - Pure or mixed malignant GCT (polyembryonal)
    - Embryonal cell
    - Teratoma
    - Yolk sac (endodermal sinus tumor)
    - Choriocarcinoma

- **Non Germ-cell tumors**
  - Leydig (interstitial cell)
  - Sertoli (andoblastoma)

- Metastasis
- Lymphoma
- Epidermoid cyst
- Paratesticular tumors
- Mimicks/pitfalls
Germ Cell Tumors
Seminoma

Demographics

- Most common single cell-type tumor and most common tumor in undescended testis
- Age 40-50
- 1-3% bilateral
- Increased hCG
- 25% metastasis at presentation
- Good prognosis
- Spermatocytic subtype: older age group, no symptoms, no tumor marker, no metastasis

Imaging, pathology and treatment

- Well-defined, hypoechoic, solid mass
- Small tumors (<1.5 cm) avascular; larger tumors hypervascular
- May have cystic component
- Calcifications may be present
- Treatment: radiotherapy ± chemotherapy
  - Unless spermatocytic subtype, treatment is orchiectomy
Seminoma

Imaging: Enlarged left testicle with numerous heterogeneous and hypoechoic nodules and masses with hyperemic intervening parenchyma between the nodules and masses

Pathology: seminoma
Embryonal Cell Carcinoma

**Demographics**

- **Pure:**
  - Rare, represents 2-3% all testicular tumors
- **Mixed:**
  - Common, present in 87% mixed germ cell tumors
- 3rd and 4th decades
- Often small at presentation
- Aggressive

**Imaging and pathology**

- Heterogeneous, mostly solid mass
- Poorly defined margins
- May demonstrate necrosis
- +/- coarse calcifications
- Can invade tunica albuginea and cause abnormal testicular contours
- Anaplastic epithelial cells
Pure GCT, Embryonal Cell Carcinoma Predominant

Imaging: Ill-defined hypoechoic intratesticular mass with coarse and fine calcifications (white arrow) resulting in abnormal contour of the testicle (yellow arrow)

Pathology: Embryonal cell carcinoma, pure (green arrow)
Teratoma

Demographics

- 4-9% all testicular tumors
- Pure:
  - Very young children (<2 years)
- Mixed:
  - Young adults (3rd and 4th decade)
- Present as painless testicular mass

Imaging, pathology and treatment

- Well-defined anechoic/complex heterogeneous cystic intratesticular mass
- Cystic areas, calcification, and/or fibrosis can suggest teratoma
- May contain mucinous or sebaceous material, hair follicles
- Treatment:
  - Varies depending on stage
  - Surgical → chemotherapy
Imaging: 2 year old patient with asymmetrically enlarged testicle with painless, firm, heterogeneously hypoechoic testicular mass demonstrating intermittent vascular flow

Pathology (image not available):
Malignant GCT, nonseminoma (60% immature teratoma, 40% yolk sac tumor)
Yolk Sac Tumor (Endodermal Sinus Tumor)

Demographics
- Common
- 80% childhood testicular tumors
- <2 years
- Pure:
  - Rare in adults
- Mixed:
  - Present in 44% adult cases
- AFP elevated >90%

Imaging, pathology and treatment
- Nonspecific imaging features
- May only have testicular enlargement without discrete mass
- Totipotential germ cells
- Treatment:
  - Varies depending on stage
  - Often confined to testis at time of orchiectomy
    - If serum AFP is not elevated, orchiectomy may be curative
Mixed GCT, Yolk Sac Tumor Predominant

Imaging: Asymmetrically enlarged testicle with complex solid and cystic intratesticular mass with vascularity to the solid components in background of microlithiasis

Pathology: Malignant mixed GCT, nonseminomatous (40% yolk sac tumor, 30% embryonal cell carcinoma, 30% immature teratoma with rare syncytiotrophonlasts)
Choriocarcinoma

Demographics

- Rare
- Pure:
  - Represents <1% testicular tumors
- Mixed:
  - Present in 8% mixed germ cell tumors
- Often present with widespread, early metastases
  - Lung, liver, GI tract, brain
- HCG elevated in 10%

Imaging and treatment

- Heterogeneous solid intratesticular mass
- Commonly with hemorrhage and focal necrosis
- Calcification and cystic necrosis also common
- Metastases also hemorrhagic
- Treatment:
  - Worst prognosis
  - Death usually within 1 year of diagnosis (pure)
  - 5 year survival rate of 48% (mixed)
Mixed GCT, containing Choriocarcinoma

Imaging: Heterogenously hypoechoic mass containing coarse and punctate calcifications (white arrow) with increased vascularity

Pathology: Malignant mixed GCT nonseminomatous (40% yolk sac tumor, 30% embryonal carcinoma, 20% immature teratoma, and 10% choriocarcinoma)
Non-Germ Cell Tumors
Sertoli Cell Tumor

**Demographics**
- <1% of testicular tumors
- Mean size: 3.5 cm, majority benign; malignant > 5 cm
- Mean age 45 years; up to 20% occur in childhood
- May produce estrogen/Müllerian inhibiting factor
- Association with Peutz-Jegher or Carney syndromes in younger ages.
- Some bilateral
- Presentation: slowly enlarging testicular mass

**Imaging and treatment**
- Solid hypoechoic mass with cystic component +/- punctate calcifications.
- Large calcifications associated with syndromes
- Internal or perinodular flow
- Treatment: orchiectomy
Sertoli Cell Tumor

Imaging: Small, heterogeneous, hypoechoic, solid lesion involving the lateral aspect of the right testicle with increased color Doppler flow

Pathology: Sertoli cell tumor
Lymphoma

Demographics

- 5% of testicular tumors
- Most common testicular malignancy in >60 years
- Median age: 66 - 68 years
- Most common bilateral testicular neoplasm
- Presents as firm painless mass
- Constitutional symptoms uncommon. If present, strongly suggests systemic disease

Imaging, pathology and treatment

- Hypoechoic mass with increased vascularity
- Hydrocele in ~40% of cases
- Involves epididymis and spermatic cord in 1/2 of cases
- Majority are diffuse large B-cell lymphoma
- Treatment: orchiectomy + chemotherapy
Lymphoma

Imaging: Hypoechoic focal intratesticular masses with high vascularity and associated hydrocele

Pathology: lymphoma
Epidermoid Cyst

Demographics

- 1% of all testicular tumors
- 0.5-10.5 cm in diameter
- Most common in 2nd-4th decade
- No malignant transformation

Imaging, pathology and treatment

- Well-circumscribed encapsulated round mass
- Alternating hypo and hyperechoic rings (onion skin appearance) or echogenic center (bull’s eye or target appearance)
- No blood flow
- Keratinizing squamous epithelium within a fibrous wall
- Treatment: local excision
Imaging: Well-circumscribed predominantly hypoechoic lesion with an echogenic rim and lamellated periphery with heterogeneous internal echotexture in the medial aspect of the left testicle abutting the mediastinum

Pathology: epidermoid cyst
Paratesticular Masses

- 3-5\textsuperscript{th} decade
- Usually slow-growing
- Most are benign
  - Adenomatoid, most common (30%)
  - Papillary cystadenomas
  - Leiomyomas
- Malignant masses, extremely rare in adults
  - Adenocarcinomas
  - Sarcomas
    - Rhabdomyosarcomas
    - Leiomyosarcoma
    - Liposarcoma
Adenomatoid Tumor

Demographics

- Benign solid tumor of epididymis
- Most common solid mass of epididymal tail
- > 3rd decade
- 98% asymptomatic
- Can slowly enlarge over time

Imaging and treatment

- Solid round or oval mass
- Most often in epididymal tail (4x more common)
- Mostly iso- or hypoechoic
- Rarely cystic
- Typically hypovascular
- Treatment: benign, although most are surgically excised to confirm diagnosis
Scrotal Liposarcoma

Demographics

- Solid, bulky lipomatous malignant tumor
- 2nd most common soft tissue tumor in adults, 10-16% incidence
- Lipoma of spermatic cord
  - ~7% paratesticular sarcomas
- Middle aged and elderly
- Up to 1/4 recur, 1/10 metastasize
- Round cell type: poorly differentiated and highly metastatic

Imaging and treatment

- Nonspecific imaging appearance on US. If can identify fat, helpful
- Often contain calcification
- CT and MR more specific for recognition of fatty tissue
- Treatment: excision including inguinal lymph nodes
  - Additional treatment depends on stage and histologic profile
Scrotal Liposarcoma

CT: Fat density mass in the left inguinal canal extending into the left hemiscrotum

Ultrasound: Nonspecific minimally vascular heterogeneous echogenic tissue in the inguinal canal and left hemiscrotum

Pathology: well-differentiated liposarcoma abutting but not involving the testes and epididymis
Mimics/Pitfalls

**Testicular**
- Infarct
- Rete testis cyst
- Hematoma
- Abscess

**Paratesticular**
- Paratesticular cystic lesions can rarely mimic solid tumors
  - Spermatocele
  - Complicated epididymal cyst
  - Tubular ectasia of rete testis
  - Tunica albuginea cyst
  - Hematocele
  - Pyocele
  - Complicated hydrocele
Heterogeneously hypoechoic solid and cystic lesion of the testis without definite blood flow to the solid component

Pathology: Small circumscribed infarct without evidence of malignancy
Imaging: Several small cystic lesions in the periphery of the testis, consistent with cystic dilation of the rete testis
Testicular Tumor Mimic: Testicular Hematoma

Imaging: Avascular, heterogeneous parenchymal echogenicity of testis in a patient with history of trauma
Paratesticular Tumor Mimic: Complicated Epididymal Tail Cyst

Imaging: Complex heterogeneous solid and cystic lesion of the epididymal tail with peripheral vascularity

Pathology: benign epididymal cyst with hemorrhage
References


- Statdx https://my.statdx.com/

- Radiopaedia http://radiopaedia.org/