Bahrain Medical Bulletin, Vol. 36, No. 2, June 2014

#### Leiomyoma of Tunica Albuginea

#### Kadim Zabar, CABS\* Akbar A Jalal, CABU\*\* Mohammed Matar Al Alawi, CABU\*\*\*

#### ABSTRACT

Leiomyoma of the tunica albuginea is an extremely rare benign tumor of the genitourinary tract. A thorough and proper examination and investigation are necessary for proper management and testis salvage surgery should be considered if feasible in such cases.

We present a fifty-six-year-old male who had benign smooth muscle tumor (leiomyoma) arising from tunica albuginea which was successfully treated with testis sparing excision of the tumor. This is the first case of leiomyoma of tunica albuginea which has been reported in the Kingdom of Bahrain.

Bahrain Med Bull 2014; 36(2):114-116

## INTRODUCTION

Leiomyomas are benign tumors which arise from smooth muscle cells and are often found as benign lesions arising in the uterus<sup>1,2</sup>. Leiomyomas are rarely found in the genitourinary tract outside the uterus; the renal capsule is the site of the majority of these tumors. Leiomyomas have been reported in the epididymis, spermatic cord, testis, tunica albuginea, bladder, prostate, scrotum and the glans penis<sup>3-5</sup>. Leiomyoma of the tunica albuginea is extremely rare.

The aim of this report is to present a case of a leiomyoma of the tunica albuginea treated successfully with testis sparing excision of the tumor.

## THE CASE

A fifty-six-year-old Bahraini male presented with a painless left hemi-scrotal swelling for more than a year; it was gradually increasing in size with occasional pain in left groin. The patient denied any history of genital trauma or infection.

On physical examination, swelling of his left scrotum was found; the testis could not be palpated because of tense hydrocele. The right testis, epididymis and spermatic cord were unremarkable. The patient's routine blood biochemistry and hematological workup were within normal limits.

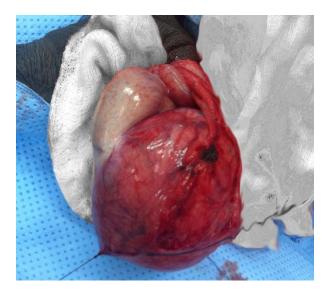
The scrotal ultrasound was suggestive of left-sided hydrocele with chronic inflammatory changes with normal testis. The right-sided testis, epididymis and hemiscrotum were normal.

<sup>\*</sup> Consultant Urologist

<sup>\*\*</sup> Specialist Urologist

\*\*\* Chief Resident Urologist Salmaniya Medical Complex Kingdom of Bahrain Email: drakbarjalal@gmail.com

The patient underwent elective hydrocele surgery. During exploration, it was found to have infective hydrocele with thickened tunica vaginalis with an extra testicular mass about 6x6x5 cm, arising from the lower part of the tunica albuginea and extending to the lower part of the epididymis; however, it was not involving the epididymis, see figure 1.



## Figure 1: Extra-testicular Mass Arising from the Tunica Albuginea

Intraoperative frozen section biopsy showed whorling bundles of smooth muscle cells of benign features.

As there was a cleavage line between the tumor and the testis, a decision was made to do testis sparing excision of the tumor. The tumor was completely excised sparing the testis and the epididymis, see figures 2 and 3. Postoperatively the patient recovered well. Follow-up tumor markers and the scrotal ultrasound were unremarkable.



Figure 2: Left Testis after Excision of the Tumor

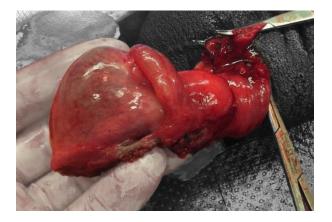


Figure 3: Left Testis and Epididymis after Excision of the Tumor

Final pathology revealed a well-circumscribed grey white mass measuring 6x6x5 cm and weighing 150 grams with whitish whorled appearance, see figures 4 and 5. Microscopically, the tumor was composed of interlacing and whorling bundles of smooth muscle cells. The tumor cells were spindled containing centrally located nucleolus and showing no mitotic activity or nuclear atypia, see figures 6 and 7. The mass was diagnosed as a leiomyoma.



Figure 4: Gross Appearance - Extratesticular mass 6x6x5 cm



Figure 5: Cut Section of the Tumor Showing Whitish Whorling Smooth Muscle, No Necrosis or Cystic Changes

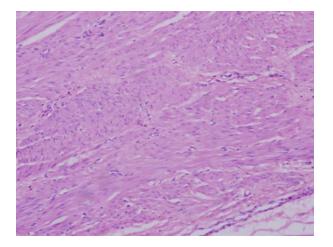


Figure 6: Microscopic Appearance – High Power Showing Fascicles of Benign Smooth Muscle, No Atypia

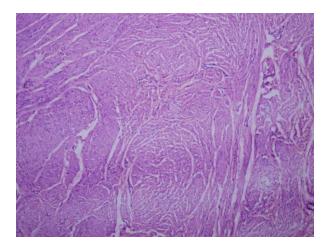


Figure 7: Microscopic Appearance – Low Power

# DISCUSSION

Leiomyomas are benign tumors derived embryologically from mesenchymal cells. Although rare in the genitourinary tract, leiomyomas can originate from any structure containing smooth muscle, most commonly the renal capsule. Leiomyoma of the tunica albuginea is extremely rare<sup>1-5</sup>.

Leiomyoma of the tunica albuginea is considered to be of benign behavior. It shows no invasive growth or metastasis<sup>6</sup>. Inflammatory hydrocele, multiloculated hematocele and a Sertoli cell tumor of the testis are potential differential diagnosis by sonography<sup>7,8</sup>. Tumors arising from the testicular tunics are rare and most cases are fibromas. The etiology of leiomyomas of the tunica albuginea is controversial. Leiomyomas could arise from the smooth muscle of blood vessels or totipotent teratoma<sup>9</sup>. Inflammatory myofibroblastic tumor (IMT) of the spermatic cord is a possible differential diagnosis<sup>10</sup>.

Scrotal mass must be properly evaluated and investigated to rule out the possibility of malignancy. Extratesticular masses are commonly benign and usually secondary to trauma, infection and inflammation or benign neoplasms; ultrasonographic study is needed to confirm the diagnosis. In our case, the extratesticular mass was missed by the physical examination and sonography due to the presence of large hydrocele.

Leiomyoma of the tunica albuginea is an extremely rare condition, only a few cases of this tumor have been reported<sup>11</sup>. It is advisable to do intraoperative histopathological examination of paratesticular tumors despite that three-quarters are non-malignant; the treatment of choice in non-malignant mass is simple extirpation. An Orchiectomy is not necessary; usually, a testis and epididymis sparing surgery can be achieved.

#### CONCLUSION

# Leiomyoma of the tunica albuginea is an extremely rare condition. An Orchiectomy is not necessary in such case and usually testis and epididymis salvage operation can be achieved and should be considered as treatment of choice as in our reported case.

**Author contribution:** All authors share equal effort contribution towards (1) substantial contributions to conception and design, acquisition, analysis and interpretation of data; (2) drafting the article and revising it critically for important intellectual content; and (3) final approval of the manuscript version to be published. Yes.

Potential conflicts of interest: None.

Competing interest: None. Sponsorship: None.

Submission date: 21 October 2013. Acceptance date: 5 April 2014.

Ethical approval: Approved by Surgical Department, SMC, Bahrain.

#### REFERENCES

- 1. Belis JA, Post GJ, Rochman SC, et al. Genitourinary Leiomyomas. Urology 1979; 13(4):424–9.
- 2. Robboy SJ, Bentley RC, Butnor K, et al. Pathology and Pathophysiology of Uterine Smooth-Muscle Tumors. Environ Health Perspect 2000; 108 Suppl 5:779–84.
- 3. Borri A, Nesi G, Bencini L, et al. Bizarre Leiomyoma of the Epididymis. A Case Report. Minerva Urol Nefrol 2000; 52(1):29-31.
- 4. Redman JF, Liang X, Ferguson MA, et al. Leiomyoma of the Glans Penis in a Child. J Urol 2000; 164(3 Pt 1):791.
- 5. Rosen Y, Ambiavagar PC, Vuletin JC, et al. Atypical Leiomyoma of Prostate. Urology 1980; 15(2):183–5.
- Lia-Beng T, Wei-Wuang H, Biing-Rorn C, et al. Bilateral Synchronous Leiomyomas of the Testicular Tunica Albuginea. A Case Report and Review of the Literature. Int Urol Nephrol 1996; 28(4):549-52.
- 7. Cunningham JJ. Sonographic Findings in Clinically Unsuspected Acute and Chronic Scrotal Hematoceles. AJR Am J Roentgenol 1983; 140(4):749-52.
- 8. Cunningham JJ. Echographic Findings in Sertoli Cell Tumor of the Testis. J Clin Ultrasound 1981; 9(6):341-2.
- Chiaramonte RM. Leiomyoma of Tunica Albuginea of Testis. Urology 1988; 31(4):344– 5.
- 10. Yee CH, To KF, Hou SM, et al. Inflammatory Myofibroblastic Tumor of Spermatic Cord in Undescended Testis. Urology 2009; 73(6): 1423.e9-12.
- Bremmer F, Kessel FJ, Behnes CL, et al. Leiomyoma of the Tunica Albuginea. A Case Report of a Rare Tumour of the Testis and Review of the Literature. Diagn Pathol 2012; 7:140.