Case Report

First report of testis-sparing surgery for sertoliform cystadenoma: case presentation and review of literature

Razvan-George Rahota,1 Guillaume Ploussard,1 Jean-Romain Gautier,1 Christophe Almeras,1 Henri Ducoin,2 Christophe Tollon,1 Jacques Assoun,3 Guillaume Loison,1 Jean-Baptiste Beauval1 and Ambroise Salin1

Department of 1Urology, 2Pathology, and 3Radiology, La Croix du Sud Hospital, Quint Fonsegrives, France

Introduction: Sertoliform cystadenoma is a very rare, benign lesion of the rete-testis difficult to distinguish from other malignancies of the testicle.

Case presentation: We present the case of a 42-year-old male who presented with a right testicular mass, asymptomatic for 1 year. Clinical examination revealed a palpable, painless, and well-delimited right testicular superior pole nodule. Testicular ultrasound confirmed the nodule, whereas serum tumoral markers were normal. The patient underwent inguinal partial orchiectomy. Intraoperative excisional biopsy and frozen section pathology were performed, reporting undetermined tumoral origin with negative surgical margins. Ischemia time was 12 minutes. The final pathology report showed a Sertoliform cystadenoma of rete testis, with immunomorphology positive for AE1, CK7, and negative surgical margins.

Conclusion: To our knowledge, this is the first report of testicular sparing surgery for Sertoliform cystadenoma, a very rare benign tumor of rete testis. All previously reported cases were managed by radical inguinal orchidectomy.

Key words: excisional biopsy, frozen section, sertoliform cystadenoma, testicular-sparing surgery.

Keynote message
To our knowledge, this is the first report of testicular sparing surgery for Sertoliform cystadenoma, a very rare benign tumor of the rete testis. All previously reported cases were managed by radical inguinal orchidectomy. Partial inguinal orchectomy with frozen section pathology could be an alternative to radical surgery in selected cases.

Introduction
Testicular tumors account for 1% of male cancers and approximately 5% of urogenital malignancies, with an increasing incidence due to the widespread use of ultrasound examination.1 Paratesticular tumors, especially masses of the rete testis, are a rare finding and can be classified in non-neoplastic developmental pathologies (adenomatous hyperplasia, acquired or secondary cystic dysplasia), benign tumors (cystadenomas or Sertoliform cystadenoma), and malignant lesions such as adenocarcinoma of the rete testis.2,3 The first report in the literature of Sertoliform cystadenoma was made by Jones MA in 1997.4 All previously reported cases of sertoliform cystadenoma have been managed by radical surgery-inguinal orchidectomy.5,6 We present the first case of a Sertoliform cystadenoma in a 42-year-old male managed by conservative surgery.

Case report
A 42-year-old male with no prior medical history presented with a palpable mass in the right testicle, asymptomatic, and without any increase in size for 1 year. The patient is married, without children. At clinical examination, a firm, painless, well-delimited superior pole right testicular
lesion was objectified. Scrotal ultrasound revealed a 1.7/0.5 cm mass in the upper pole of the right testicle, with normal contralateral testicle and normal testicular volumes (Fig. 1). Serum tumor markers were collected and in normal range, as following: alpha-fetoprotein (0.5 ng/ml, N = 0.89–8.78), beta-human chorionic gonadotropin (0, N < 1.4 UI/L), and lactate dehydrogenase (10 UI/L, N = 120–280 UI/L). A staging chest–abdominal–pelvic contrast-enhanced CT scan was performed which did not show any evidence of visceral or lymph node metastasis.

Given the fact that the lesion was perceived as stable by the patient for 1 year, as described during the first consultation, we suspected that a benign etiology could be possible and offered the patient the option of testis-sparing surgery. The patient was informed of the risk of radical orchietomy in case of intraoperative aspect or frozen section pathology report highly suggestive for malignancy and salvage radical orchietomy if the final pathology report would be conclusive for malignancy. After informed consent was given, the patient underwent right partial orchiectomy by inguinal approach. We identified and dissected the elements of the spermatic cord, after which we separated the vas deferens from the testicular vessels. After clamping the testicular vessels, we incised the tunica albuginea and proceeded with the partial orchietomy within safety surgical margins. Frozen section assessment was demanded and revealed testicular tumor of indeterminate origin, with negative surgical margins. Thus, we closed the albuginea and reintegrated the ipsilateral testicle in the scrotum, at the same time performing an orchidopexy (Fig. 2). The total ischemia time was 12 min.

The final pathology report showed a Sertoliform cystadenoma of the rete testis, with the tubular architecture of variable sizes, without atypical cells and rare mitosis visible. The testicular parenchyma present on the specimen was without pathological characteristics and the surgical margins were negative. Immunohistochemistry was intense positive for AE1 and CK7, negative for Inhibin and EMA, without any signs of malignancy (Fig. 3).

The patient was discharged the same day, no immediate or long-term postoperative complications were reported. The 1-month postoperative scrotal ultrasound examination was normal, with no sign of local recurrence. Due to the fact that it is a rare histology, we performed also 2 and 6-months postoperative serum tumoral markers, which were all negative.

**Discussion**

Sertoliform cystadenoma is a rare benign tumor of the rete testis, being for the first time mentioned in the literature in 1997 by Jones et al. The largest case series in the literature was presented by Paluru et al. and is comprised of 15 cases; mean age reported was 46 years, mean tumor size 1.5 cm, mean follow-up of 8.1 years with no complications and no evidence of disease recurrence. All of the patients underwent radical inguinal orchietomy.

Several single case reports have been also identified in the literature. Sahnan et al. and Bremmer et al. describe the cases of a 19 and 66-year-old man, respectively, with a 6-month history of smooth, painless right testicular mass, stable over time, with normal serum tumoral markers in which right radical inguinal orchietomies were performed.7,8

The first pediatric case was reported by Kacar et al. in a 6-year-old boy who presented bilateral gynecomastia and left scrotal tumefaction. Serum tumoral markers were in the normal range but estradiol and testosterone levels were slightly increased (48.4 ng/mL and 20 ng/dL, respectively). No other adult patient reported in the literature displayed increased levels of sexual hormones. The patient underwent left inguinal orchietomy and the final pathology report revealed Sertoli cell proliferations, compatible with Sertoliform cystadenoma.

The most recent case of Sertoliform cystadenoma was presented by Lahouti et al. in a 39-year-old male with a right superior pole testicular mass, in which serum tumor markers were negative. The final pathology report revealed sertoliform cystadenoma with immunohistochemical examination intense positive for Inhibin, cytokeratin CAM5.2, PAX8.

Histologically, the most important differential diagnosis is made with rete testis adenocarcinoma. Sertoliform cystadenoma has been reported in patients between 19 and 84 years,
having variable size characteristics (a few mm to 4 cm) with a predominant tubular architecture, columnar to cuboidal tumor cells with eosinophilic cytoplasm, and basally located nucleus. Moreover, adenocarcinoma of rete testis is also a rare but aggressive finding, with an incidence peak around 70 years of age. The most important morphologic criteria required for diagnosis are the involvement of testicular mediastinum with the preservation of the parietal tunica vaginalis, in the absence of other lesions.\textsuperscript{5,10}

Although current recommendations are in favor of considering every scrotal mass as malignant until proven otherwise, we propose in selected cases where clinical history is in favor of benign lesion and patients desire maximum fertility preservation the alternative of organ sparing surgery. Testis-preserving surgery should always be accompanied by frozen section examination, due to the fact that it has a high concordance with the final pathology result.\textsuperscript{11} However, all patients must be informed of the risk of immediate or salvage radical orchidectomy if the frozen section or final pathology report are consistent with malignancy.

**Conclusions**

In conclusion, we report the first case of Sertoliform cystadenoma managed by conservative surgery—partial orchidectomy, without any early or late postoperative complications. Sertoliform cystadenoma remains a rare finding among intratesticular mass findings, with only 16 cases reported in the literature, all treated by radical inguinal orchidectomy.

**Funding source**

No funding was received for this study.

**Conflict of interest**

The authors disclose no conflicts of interest.

**Approval of the research protocol by an Institutional Reviewer Board**

This study was approved by the local ethics committee of the Croix du Sud hospital (register number 57/2021) and was conducted in line with the principles of the Declaration of Helsinki.

**Informed Consent**

Not applicable.
Registry and the Registration No. of the study/trial

Not applicable.

References

1 Albers P, Albrecht W, Algaba F et al. Update. Eur. Urol. 2015; 2015(68): 1054–68.
2 Emerson RE, Ulbright TM. Morphological approach to tumours of the testis and paratestis. J. Clin. Pathol. 2007; 60: 866–80.
3 Rosevear HM, Mishail A, Sheynkin Y, Wald M. Unusual scrotal pathology: an overview. Nat. Rev. Urol. 2009; 6: 491–500.
4 Jones MA, Young RH. Sertoliform rete cystadenoma: a report of two cases. J. Urol. Pathol. 1997; 7: 47–53.
5 Lahouti AH, Brodherson M, Larish Y, Unger PD. Sertoliform cystadenoma of the rete testis: report of a case and review of the literature. Int. J. Surg. Pathol. 2017; 25: 555–8.

Editorial Comment

Editorial Comment to First report of testis-sparing surgery for sertoliform cystadenoma: case presentation and review of literature

Orchiectomy is the gold standard for managing testicular tumors. According to the American Urological Association guideline, patients with a suspected malignant testicular lesion and a normal contralateral testis should undergo radical inguinal orchiectomy, instead of testis-sparing surgery (TSS). TSS may be offered as an alternative for treating testicular tumors in highly selected patients having tumors of diameter 2 cm or less who wish to preserve testicular function or whose testicular function is likely to be substantially impaired by orchiectomy. However, TSS is controversial for cases that are strongly suspected to be benign.

Sertoliform cystadenoma is a very rare benign tumor originating from the rete testis. Ultrasound findings show a cystic mass in many cases, but sometimes a solid mass is observed with an epididymal cyst. If malignancy cannot be completely ruled out after examination with various modalities, orchiectomy is still considered preferable. In particular, given the rarity of sertoliform cystadenoma, it may be difficult to definitely rule out malignancy preoperatively. However, when the mass is obviously benign and no malignant findings on rapid intraoperative pathology after ischemia of the testicular vessels exist, TSS with tumor enucleation with a wide enough margin may be considered. Preoperative evaluation of testicular function, including analysis of semen, serum testosterone levels, and gonadotropin levels, is advised when considering the possibility of conversion to orchiectomy.

In this case, the tumor was judged to be benign because there was no change over a long period, and the authors successfully performed TSS. If possible, preoperative testicular function should be evaluated to determine if TSS is truly necessary. Simultaneous testicular sperm extraction should be considered for patients who wish to have children and have azospermia in the preoperative semen analysis or who are expected to have decreased testicular function.

Teppei Takeshima M.D., Ph.D.
Department of Urology, Reproduction Center, Yokohama City University Medical Center, Kanagawa, Japan
teppei_t@yokohama-cu.ac.jp
DOI: 10.1002/jiu5.12378

Conflict of interest

The author declares no conflict of interest.

References

1 Stephenson A, Eggener SE, Bass EB et al. Diagnosis and treatment of early stage testicular cancer: AUA guideline. J. Urol. 2019; 202: 272–81.
2 Paluru S, Ulbright TM, Amin M, Montironi R, Epstein JI. The morphologic spectrum of sertoliform cystadenoma of the rete testis: a series of 15 cases. Am. J. Surg. Pathol. 2018; 42: 141–9.
3 Rahota R, Ploussard G, Gautier J et al. First report of testis-sparing surgery for sertoliform cystadenoma: case presentation and review of literature. IUJ Case Rep. 2021; 4: 425–8.
4 Tsutsumi S, Kawahara T, Takeshima T et al. Onco-testicular sperm extraction (onco-TESE) for bilateral testicular tumors: two case reports. J. Med. Case Rep. 2017; 11: 139.