



Figure 2 : Dilated and hyperechoic zones within the head and body of the epididymis.

The tuberculin skin test was negative. Urinalysis showed aseptic leukocyturia. The urine PCR test for tuberculosis on three successive days was negative, the urine culture in Ziehl-Neelsen's medium was negative, and the IFG test was negative. The search for biological markers (alpha-fetoprotein, total HCG, and lactate dehydrogenase) was negative, and HIV serology was negative. The diagnosis was made only after performing a surgical epididymal biopsy, which demonstrated epithelioid and giantocellular granulomatous epididymitis, with the presence of caseous necrosis compatible with tuberculosis. Pus culture came back positive. Our patient underwent an imaging assessment, which did not reveal any other secondary location of tuberculosis. The patient was put on antituberculosis treatment.

DISCUSSION

Genital tuberculosis can occur at any age, but it primarily affects men between the ages of 30 and 50 years of age [3]. The first organ most commonly affected is the epididymis, followed by the seminal vesicle, the prostate, the testis, and the vas deferens [3]. A recent study indicated that isolated epididymal tuberculosis may be the first or only manifestation of early genitourinary tuberculosis [3]. The epididymal-testicular disease can occur either through the reflux of infected urine into the genital tract or through direct blood or lymphatic transmission [4]. The latter explains the increasing occurrences of isolated genital tuberculosis, which is similar to that of our patient. Sexual transmission has also been reported in the literature [5]. The weakening of the immune system increases susceptibility to tuberculosis. Presently, HIV stands as the primary driver behind tuberculosis development. Besides reactivating dormant infections, HIV-induced immunosuppression can hasten the advancement of new infections or reinfections. Between 20 and 50% of HIV-positive individuals globally experience active tuberculosis [6]. Testicular involvement can manifest as testicular cancer, resulting in unnecessary orchiectomy when unrecognized [2]. Testicular involvement is a severe form of genital tuberculosis [4]. Clinically, epididymotesticular tuberculosis may manifest as scrotal ulcer, scrotal swelling, single, bipolar, or multiple epididymal nodules, testicular induration, and disappearance of the epididymotesticular groove [7]. This atypical clinical presentation is responsible for a significant delay in diagnosis with the risk of several complications (hypogonadism, infertility, or sexual transmission) [2].

Lee et al. conducted a study on 29 patients with testicular tuberculosis, demonstrating that only 17.2% were initially considered to have tuberculosis [5]. Our patient presented with nontender epididymal-testicular induration with weight loss and night sweats.

Ultrasound remains the best exam to explore the scrotum, testis, epididymis, and vas deferens [4]. In most cases, epididymal tuberculosis can appear as hypoechoic lesions, sometimes hyperechoic, or mixed lesions. Sometimes, we can find an enlargement of the whole epididymis or collections that correspond to tuberculosis abscesses similar to what our patient had on ultrasound. The testes can also be heterogeneous and enlarged. In some cases, testicular tuberculosis can appear as patches of hyperechoic lesions, resembling miliary testicular tuberculosis, as was observed in our patient.

Diagnosis of isolated genital involvement is still challenging today and relies on clinical, biological, radiological, and bacteriological evidence. All available laboratory tests are either nonspecific or have not yet been evaluated. Intradermal injection of tuberculin (purified protein) has an estimated sensitivity of 60% to 88% in urogenital tuberculosis [8]. New immunological tests, such as QuantiFERON-TB Gold, have an estimated sensitivity between 81% and 89% and specificity between 96% and 100% for both pulmonary and extra-pulmonary tuberculosis [9]. There is no specific assessment for genital tuberculosis. For our patient, the gamma interferon test was negative.

Investigations conducted on genital secretions have been poorly evaluated in isolated cases such as our case [1]. Sperm culture is rarely performed and little evaluated; therefore, it is not recommended in current practice [1]. The culture of pus and peritesticular fluid is a reliable test. However, it takes a long time (up to 42 days) to get the results [2]. However, it has the advantage of being able to test for sensitivity and resistance to the four standard antituberculosis drugs [2].

Unfortunately, biopsy and orchiectomy remain the only means to make a rapid diagnosis up until now. This was the only way to obtain the diagnosis of genital tuberculosis in our case. CT and IVU keep their essential place in exploring the urinary tract [3]. CT IVU revealed no anomaly in our patient. This is suggestive of isolated genital disease of Tuberculosis. Treatment is based on antituberculosis agents [1].

CONCLUSION

There is an increasing incidence of Isolated genital tuberculosis, which is a form of urogenital tuberculosis. There is still no codified approach to diagnosing this condition due to the absence of specific and sensitive investigations. Currently, biopsy and orchiectomy remain the only means of confirming this condition. We recommend the serial semen culture, reactional hydrocele fluid culture, or intracrotal pus culture to identify *Mycobacterium tuberculosis* when possible.

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AUTHORS' CONTRIBUTIONS

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COMPETING INTERESTS

The authors declare no competing interests in this case.

PATIENT CONSENT

Written informed consent was obtained from the patient.

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