Persistent Mullerian Duct Syndrome with Transverse Testicular Ectopia

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INTRODUCTION

The Mullerian duct regresses in male due to a glycoprotein secreted by the developing testes called Mullerian inhibiting factor (MIF). Failure to regress may be due to lack of MIF or defective MIF receptor, resulting in various disorders of regression, and one such disorder is persistent Mullerian duct syndrome (PMDS). PMDS is a rare form of male pseudohermaphroditism characterized by the presence of uterus and fallopian tubes in phenotypically and genotypically normal male (46, XY), with testosterone production and responsiveness.

The association between PMDS and transverse testicular ectopia (TTE) is even more uncommon.

CASE REPORT

A 25-year-old male presented with absent left testis since birth and right sided inguinal swelling for 5 years, and pain in swelling for 2 days. On physical examination, the left scrotum was empty, and right side revealed indirect inguinal hernia with right testis at the root of the scrotum. The penis was well developed. The patient was phenotypically male, exhibiting normal secondary sexual characteristics. The past history indicated primary infertility.

Semen analysis revealed azoospermia. Ultrasonography showed right inguinal hernia with undescended testis in the right inguinal canal and an absent testis on the left side.

Surgical exploration revealed indirect sac with the right testis. While retrieving the testis, we noticed a complex mass resembling uterus-like structure, and gonad (left) with the fimbria-like structure [Figure 1].

Both the gonads had vas deferens and vascular supply. Orchiopexy was done on the right side. The uterus like structure was removed along with left gonad as it was atrophic and sent for histopathology. Right inguinal hernioplasty was done.

Histopathological study proved it to be uterus [Figure 2] and left gonad like structure proved to be testis which was atrophic without spermatogenesis [Figure 3].

Chromosomal analysis revealed normal male genotype – 46 XY [Figure 4].

ABSTRACT

Persistent Mullerian duct syndrome (PMDS) is a rare form of male pseudohermaphroditism characterized by the presence of Mullerian duct structures in a normal male with 46, XY karyotype. Transverse testicular ectopia (TTE) is rare form of testicular ectopia in which two testes are located on one inguinal side. The opposite scrotum is empty. PMDS with TTE is rare. We report a case of PMDS with TTE discovered during surgery for a right inguinal hernia in a 25-year-old male.

Key words: Mullerian inhibiting factor, persistent Mullerian duct syndrome, transverse testicular ectopia
DISCUSSION

Embryologically, between 7th and 8th week of gestation, masculinization occurs in a male fetus. Testosterone secreted by the Leydig cells in the testis helps in the development of Wolffian duct into vas deferens, epididymis and seminal vesicle. MIF secreted by Sertoli cells in the testis, acts locally to suppress the Mullerian ducts and causes their regression by 8th and 10th weeks of gestation. PMDS results from failure of synthesis of MIF, or failure of end organs to respond to MIF.[2] MIF gene has been mapped to chromosome 19. Receptor for MIF is expressed in Sertoli cells, granulosa cells and Mullerian duct.[3] MIF does not have direct role in the descent of testes. The mechanical effect of persistent Mullerian duct structures might produce cryptorchidism by preventing normal testicular descent.[4]

Male pseudohermaphroditism results from the defective formation or action of androgens or MIF. PMDS represents a small fraction of this broad spectrum of male pseudohermaphroditism, characterized by the presence of well-developed or rudimentary uterus, cervix, vagina and fallopian tube in normal 46 XY male. Patients with PMDS have normal androgen production, male external genitalia and normal penile development. Testis show features of atrophy due to cryptorchidism. Infertility is common but fertility has been reported in a few.[5]

Normally the testis is located in the scrotum at birth. Failure of the testis to descend into the scrotum at birth results in undescended testis. Ectopic testis have been reported at various sites including superficial inguinal pouch, suprapubic, femoral, perineal region and the base of penis. Migration of the testis to the opposite side where both testis pass through the same inguinal canal is known as TTE.

PMDS occurs as:
- Partially descended testicles (80–90% of cases) with unilateral cryptorchidism and contralateral inguinal
hernia. The term “hernia uteri inguinalis” is used when uterus is found in the hernia sac
• Sometimes the contralateral testis is found in the hernia sac due to abnormal mobility of the Mullerian derivatives, known as TTE
• Bilateral cryptorchidism with uterus fixed in pelvis and both testes embedded in round ligaments.[4,6]

The undescended gonads have a high risk of malignancy.[1,7]

A rare form of clear cell adenocarcinoma of the Mullerian duct in PMDS[8] has been reported. Diagnosis is usually made incidentally during surgery for an inguinal hernia or exploration for cryptorchidism. TTE is suspected preoperatively in patients with unilateral inguinal hernia and contralateral nonpalpable testis.[4,9] Laparoscopy, ultrasonography, and MIF detection using bioassay techniques is helpful in diagnosing the syndrome.[1] The diagnosis is confirmed with testicular biopsies and chromosomal studies.[6,9]

Removal of these structures may damage the vas deferens which lies in close proximity. Hysterectomy is indicated when the Mullerian structures limit intrascrotal placement of testis. Orchidectomy is indicated when testes cannot be mobilized to a palpable location. Otherwise orchiopexy is done and patient kept on long-term follow-up.[6,9]

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Conflicts of interest

There are no conflicts of interest.

REFERENCES