Introduction

Persistent Mullerian duct syndrome (PMDS) is a rare form of disorder of sexual development (DSD), first described by Nilson in 1939 [1]. It is an autosomal recessive disorder in which genetically and phenotypically male subjects have persistence of Mullerian duct structures (uterus, fallopian tubes and vagina) due to a deficiency in the anti-Mullerian hormone (AMH) produced by Sertoli cells, or its type II receptor (AMHR-II), and can result from a deficiency of anti-Mullerian hormone (AMH) or a defect in the receptor gene [2,3]. The syndrome is rare, with only about 150 cases described in the literature to date [4] and only few cases of malignant transformation are reported. Similar to cryptorchidism, the gonads of PMDS patients are at an increased risk for malignant transformation. Embryonal carcinoma, seminoma, yolk sac tumor, and teratomas have been seen in patients having PMDS [5]. The overall incidence of malignant change is 15-18% [6].

We report a case of PMDS with seminoma testis.

Case Presentation

A 39 year old unmarried male presented with pain abdomen since 1 year. He had B/L undescended testes and had undergone left sided hernia repair with left orchidectomy 12 years back. On examination scrotum was empty, penis was well developed and no other abnormality was detected. CT scan showed 7cmx 5 cm mass behind the bladder. Few smaller similar density masses were found in small bowel mesentery (Figure 1). Biopsy from this mass was suggestive of seminoma testis. AFP level was normal. After 3 cycles of chemotherapy (BEP) excision of both masses done. Pathological examination showed uterus and fallopian tube with leigd cell hyperplasia in right testis.

Conclusion: Surgeons and pathologists should be aware of such a diagnosis in any patient who presents with undescended testis and mass behind urinary bladder in imaging.

Keywords: Persistent mullerian duct syndrome; Seminoma; Cryptorchidism

Discussion

PMDS is a rare form of male pseudo hermaphroditism. It is characterized by the persistence of mullerian duct structures and cryptorchid testis or testes in a 46, XY karyotypic male [6]. It may be
detected in infancy or adulthood. PMDS is thought to result from the synthesis or release of MIF or from the defective MIF receptor. The MIF gene is located on Chr 19p. From 7th week of gestation onwards sertoli cells of fetal testis releases MIF which causes regression of mullerian duct. Any defect in this normal pathway leads to persistence of mullerian duct structures. This remnant Mullerian structures lead to cryptorchidism by hindering the normal testicular descent [7]. The secretion and response of target organs to testosterone are not affected, so the wolffian duct structures and external genitalia progress to a normal male. PMDS is divided in to three types based on the location of testes and uterus. Type I (60-70%): Have intra-abdominal mullerian structures and testes in a position analogous to that of ovaries, type II (20-30%) also known as hernia uteri inguinal is have one testis in hernial sac or scrotum together with mullerian structures and type III (10%): have both testes located in the same hernia sac along with fallopian tubes and uterus (transverse testicular ectopia) [2].

Due to its rarity PMDS is rarely diagnosed before surgery. It is usually diagnosed postoperatively with the identification of mullerian duct structures on histopathological examination. In our case testicular seminoma was found in right testis with uterus and fallopian tube. Left testis was already removed during previous hernia surgery. Surgeons should be ready to face such surprises intraoperatively. The risk of malignancy in an ectopic testis in a case of PMDS is similar to that in a cryptorchid testis in a healthy male [8]. Germ cell tumors have been reported in the testis, whereas tumors of the Mullerian duct derivatives are very rare [9].

**Conclusion**

PMDS with seminoma is a rare entity. Because of its rarity it is difficult to diagnose preoperatively and hence it is difficult to predict intraoperatively. Surgeons and pathologists should be aware of this situation when dealing with cryptorchidism and malignancy in it. The risk of developing malignancy and other symptoms is higher in an intra-abdominal testis. When attempting for surgery for seminoma in such a scenario the mullerian remnants should also be removed.

**References**