Hernia uteri inguinalis with calcified fibroma of ovary in male : A case report

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Hernia uteri inguinalis (also known as Persistent Mullerian Duct Syndrome) is extremely rare, In this case, a 51 year-old man with the complaint of painful swelling of the right scortal sac and inguinal region was operated on under the suspicion of having right incarcerated hernia. A standard oblique inguinal incision was performed. The hernial sac contained with uterus, right utrine tube (sliding hernia type) and left ovarian tumour(intraperitoneal part). Hysterectomy with bilateral salpingooophorectomy and herniorrhaphy was made through the inguinal incision, To our knowledge, this is the first case reported of Hernia uteri inguinalis with calcified fibroma of ovary in male.

Keywords : Hernia uteri inguinalis, calcified fibroma of ovary, Persistent Mullerian Duct Syndrome.

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รายงานกรณีศึกษาผู้ป่วย 1 รายมีลักษณะทางเพศภายนอกเป็นชาย อายุ 51 ปี มา โรงพยาบาลด้วยอาการปวดบวมและมีก้อนที่ขาหนีบขวา ลักษณะเป็นใส้เลื่อนชนิดอุดดัน หลังจากทำ การผ่าตัดพบว่าเป็นใส้เลื่อนชนิด sliding ประกอบด้วยมดลูกและรังไข่ขวาบริเวณที่ผ่าตัดและเนื้องอก รังไข่ภายในช่องท้อง ได้ให้การรักษาโดยผ่าตัดมดลูก รังไข่และเย็บซ่อมผนังหน้าท้องทางแผลบริเวณ ขาหนีบขวา ผลการตรวจขึ้นเนื้อของรังไข่ซ้ายเป็น calcified fibroma ซึ่งเป็นผู้ป่วยรายแรกที่ได้รับ การรักษาและรายงาน

คำสำคัญ: Hernia uteri inguinalis, เนื้องอกรังไข่ชนิด calcified fibroma, Persistent Mullerain

Duct syndrome

Case Report

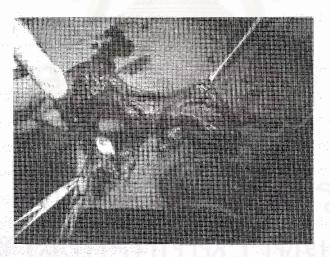
A 51 year-old man was referred to the emergency room complaining of painful swelling of the right scortal sac and the inguinal region 2 days prior to admit. He was classified as American Society of Anesthesiologist I. Past medical history was no contributory; he had no history or evidence of trauma or infertility.

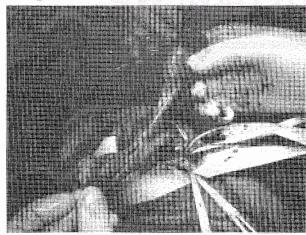
On physical examination, his vital signs were stable. The abdomen was soft and not tender. There was an obvious painful swelling of the right scortal sac and the inguinal region and normally male secondary sex characteristics, But the right scortum was not palpable. He was resuscitated with

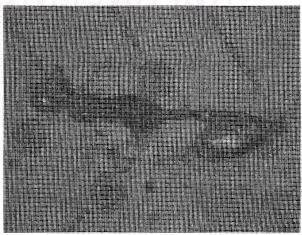
intravenous fluids being taken to the operating room. Since the preoperative diagnosis was right incarcerated hernia, the patient was given prophylactic antibiotics.

After standard oblique inguinal incision was performed, the hernial sac which was found to contain an immature uterus and right uterine tube, Atrophic testis, (sliding hernia type), The uterus was measuring 3.5x3x2 cm³. Its endometrium was 0.2 cm. Thick: myometrium 1 cm; the attached ovary 2x1x1 cm³. Left uterine tube with 6x3.5x3 cm³. ovarian tumor (intraperitoneal part) diameter, The uterous was excised and bilateral salpingoopherectomy and herniorrhaphy were made through the inguinal incision.

Gross pathology and intraoperative part as figure







The picture shows male secondary sex characteristic and hemia uter inguinalis with calcified mass of left ovary with testicular atrophy.

Postoperatively he made unevenful recovery and was discharged 3 days later, During his revisit to the Out-patient Department one month later, he was found healthy.

Surgical pathology report:

Microscopic finding: section of ovarain nodule composed of atrophic hyarinizes semiferous tubules with clusters of leydig's cells. Section of structure resembling fallopian tube show epididymis with dilated ducts.

Sections of separated mass composed totally of hyalinized collagenised tissue with extensive calcification. No ovarian tissue seen.

Diagnosis: Uterus with one adnexa

- weakly proliferative endometrium.
- unremarkable myometrium.
- attached adnexae composed of atrophic testis with dilated epididymis.

Separated mass

 totally calcified mass, probably calcified fibroma of ovary.

Discussion

Hernia uteri inguinalis (also known as Persistent Mullerian Duct Syndrome (PMDS)) which cause by impaired action of Mullerian Inhibiting Substance (MIS) in regreasithy the Mullerian duct. The Mullerian duct structures include the uterus, the fallopain tubes and the vagina. (1) An etiology of this

syndrome may be genetically inherited either by autosomal recessive genes with sex limitation or X-linked recessive mutation.⁽²⁾

Preoperative diagnosis of hernia uteri inguinalis is difficult. Various organs have been found in indirect inguinal sac, but the presence of uterus in inguinal hernia in male has only rarely been first described by Nilson 1939. (3) He described three main types of hernia uteri inguinalis, namely:-

Type I the uterus and adnexa of both sides are present in the hernia.

Type II the uterus and adnexa on oneside are contained in the hernia and the adnexa on the other side remain in the peritoneal cavity.

Type III a unicornuate uterus, or one horn of a bicornuate uterus, with its corresponding adnexa are contained in the hernia. (3)

The possibility of this type of intersex abnormality is not normally considered until laparotomy or repair of a hernia reveals the abnormality, Nilson, however, suggested that it should be possible to diagnose hernia uteri inguinalis preoperatively by bimanual examination. However, the rare condition is not known to the surgeon and it was not kept in mind then that diagnosis might be possible. The treatment is controversial with most clinicians recommending excising the persistent Mullerian structures but due to close location with the vas deferens and Mullerian structures, it is easy to injure the vas deferens. (2-5) Some clinician suggest leaving the persistent Mullerian structures intact because malignancy has not been reported and no such problems exist, in order to avoid malignancy orchidectomy should be performed in adults in case of undescended testis.

Conclusion

A case of Type II hemia uteri inguinalis (Nilson 1939) that occurred in a phenotypically normal 51 yearold male but, to our knowledge, this is the first case report of hemia uteri inguinalis with calcified fibroma of ovary in male.

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