Case Report—Inguinoscrotal ureteral hernia diagnosed on micturating cystourethrography

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Abstract

The presence of a ureter within an inguinal hernia is an extremely rare entity, usually discovered incidentally during herniorrhaphy and may pose a surgical risk. Early preoperative diagnosis is crucial to guide proper surgical approach and to preserve renal function.

Key words: Hernia; inguinal; ureteral

Introduction

Inguinal hernias may contain intraperitoneal or retroperitoneal structures, but the presence of a ureter within an inguinal hernia is rare.[1] Ureteral inguinoscrotal hernias are usually seen in obese elderly men and are generally discovered at herniorrhaphy.[2] We report the case of an inguinoscrotal hernia in an infant where a grossly dilated ureter was seen as its content.

Case Report

A 10-month-old male infant was referred for USG of the abdomen. The child had antenatally detected left-sided hydronephrosis, along with a history of groin swelling from the first week of life. The swelling was non-progressive in nature and decreased in size on lying down. There was no associated history of vomiting, abdominal distension, or excessive crying during urination.

On examination, a small soft swelling was palpable in the left groin, which was nontender, compressible, and reducible, with no local rise of temperature. An expansile cough reflux was present. A clinical diagnosis of left inguinal hernia was made.

USG abdomen revealed left-sided grade III hydroureteronephrosis with severe cortical thinning and a grossly dilated ureter throughout its extent. However, herniation of the ureter was not identified at the time of the USG examination. There was grade I hydronephrosis on the right side as well. A micturating cystourethrogram (MCU) was suggested to rule out a vesicoureteric reflux (VUR).

MCU during the filling [Figure 1A] and voiding [Figure 1B] phases showed a grossly dilated and tortuous left ureter in the left inguinoscrotal region with grade V VUR [Figure 1C]. A normal urethra was seen in the voiding phase [Figure 1B]. A radiological diagnosis of grade V VUR with ureter as a content of the left inguinal hernia was considered.

Dimercaptosuccinic acid (DMSA) scintigraphy was performed, which showed a nonfunctioning left kidney. The right kidney showed normal parenchymal uptake with no evidence of scarring.

Left inguinal herniotomy with ureterostomy was done. A hugely dilated ureter was present adjacent to the hernial sac, extending into the scrotal sac, suggestive of a paraperitoneal type of inguinoscrotal ureteral hernia. The postoperative period was uneventful. A DMSA scan repeated after one
month showed a nonfunctioning kidney (2%) on the left with a normally functioning kidney on the right. A left nephrectomy was planned for the patient but the patient was lost to follow-up.

Discussion

An inguinal hernia containing ureter is a rare entity.[1] Other sites where the ureter can herniate include femoral, sciatic, para-iliac, and thoracic.[3,4]

Patients with ureteral hernias rarely have any urinary symptoms and usually present with a mass in the groin.[2,5] It is important to identify the presence of the ureter prior to surgery in order to prevent injury to the ureter during hernia repair.[6] There are two types of ureteral inguinal hernias—paraperitoneal and extraperitoneal. Approximately 80% of ureteral hernias are paraperitoneal and 20% are extraperitoneal. The paraperitoneal type is associated with a peritoneal sac [Figure 2A], whereas an extraperitoneal herniation, which is rare, contains only a herniated ureter without a peritoneal sac [Figure 2B].

The paraperitoneal hernia is an indirect inguinal hernia that occurs due to traction on the underlying abdominal structures due to which the ureter gets pulled into the scrotum.[7] Abdominal organs like the urinary bladder, colon, etc., can make up the wall of the indirect hernial sac. The paraperitoneal hernia is not associated with other congenital renal or ureteric anomalies.[5,8]

Extraperitoneal ureteral herniation is associated with congenital anomalies of the urinary tract.[9] In this type of hernia, there may be an abnormal attachment of the ureter to the genitofemoral ligaments and Wolffian duct.[9] Retroperitoneal fat may be present in the hernia and the fat is often mistaken to be an indirect hernial sac. These hernias are smaller in size and are difficult to palpate. Unlike paraperitoneal hernia, extraperitoneal hernias present with urinary symptoms.[5] Crossed ectopia and nephrotosis are the most common associated congenital anomalies.[2,5]

Roach et al, described the CT scan findings, showing the path of the dilated ureter into the inguinal hernia.[10] There may be hydronephrosis in such cases, due to marked angulation of the ureter or due to direct ureteral compression because of inflammation of other herniated contents.[13] In our case, the hydronephrosis seen on USG was most likely due to vesicoureteric reflux. Inguinoscrotal ureteral hernias have also been described in postrenal transplant patients where hydronephrosis is seen due to a sharp stenosing bend of the ureter in the hernial sac, the ascending portion of the ureter remaining thin.[11]
So far, 140 cases have been described in the world literature, however, only one such case has been diagnosed on a preoperative urogram. The diagnosis of ureteral inguinoscrotal hernia is often missed because of the absence of urinary symptoms. Increased fatty tissue in the hernia should raise the suspicion of ureter within the hernial sac. CT scan clinches the diagnosis in patients with irreducible inguinal hernias presenting with urinary symptoms. Intravenous pyelography (IVP) may have a role when CT scan is not readily available. Surgical treatment usually involves replacement of the ureter into the retroperitoneum along with hernial repair.

With the widespread use of multiplanar imaging, it is likely that more such cases will be seen preoperatively.

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