CASE REPORT

Abdominoscrotal hydrocele with bilateral hydrenephrosis in an adult: Case report

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Abdominoscrotal hydrocele is a rare entity with unclear etiology which may be diagnosed with general examination and ultrasound imaging. During examination it may be misinterpreted as acute urinary retention of the bladder (globe-like) especially if associated with hydrenephrosis. It should be treated surgically. Here we present a case of left abdominoscrotal hydrocele with accompanying left grade 2 and right grade 1 hydrenephrosis.

KEY WORDS: Hydrocele; Hydrenephrosis; Ultrasound.

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INTRODUCTION

Abdominoscrotal hydrocele (ASH) is a rare entity with unclear etiology. It extends into the abdominal cavity through the inguinal canal (1, 2). Mostly observed in pediatric patients, ASH is scarcely described in adult population with only a few cases associated with one sided hydrenephrosis reported in the literature (3, 4). We present a case of ASH with bilateral hydrenephrosis.

CASE PRESENTATION

A 49-year-old male was admitted to our clinic with progressively increasing left scrotal swelling along a period of one year. During initial physical examination lower abdominal mass on the left and hydrocele of the left testis were observed (Figure 1). Hydrocele had simultaneous fluctuation with the lower abdominal swelling. In spite of clinical diagnosis based on physical findings and ultrasound imaging an urethral Foley catheter was inserted to rule out the presence of urinary retention and associated hydrenephrosis demonstrating an empty bladder. Bilateral hydrenephrosis occurred as a result of the compression of the bladder and the left ureter by the sac (Figure 2, 3).

Surgery was scheduled by inguinal approach in order to perform an high ligation of the processus vaginalis with complete excision of the abdominal component of the lesion. During the abdominal dissection of the sac we injured the peritoneum and it was repaired. We also mobilised the scrotal part of the sac in order to excise the

Figure 1.
tunica vaginalis of the testis and suture the edges of the remaining tunica vaginalis posterior to the testis. Mesh (Lichtenstein) method was used for inguinal hernia repair. Testis and cord were normal. The postoperative period was uneventful.

**DISCUSSION**

The etiology of ASH is not clear but there is general consensus on the need of surgical treatment. According to a Pubmed search there is only one case which Authors successfully managed by conservative approach (5). ASH is not a real benign condition, because it may be complicated with acute appendicitis, and it may lead to ureterohydronephrosis, testicular dismorphism and even leg edema in pediatric patients (6, 7). In our case at initial diagnosis we have observed bilateral hydronephrosis. ASH should be considered for the differential diagnosis of bilateral hydronephrosis. Mostly, it is one sided but in the literature bilateral ASH is also reported (8). Paratesticular malignant mesothelioma associated with ASH has also been reported in a 14-year-old boy (9). Ultrasound, magnetic resonance imaging (MRI) and computed tomography (CT) may be used for the diagnosis. During surgical treatment of ASH, dilated inguinal ring due to large sac should be repaired and mesh method should be used to avoid secondary herniation. Although open surgery is generally a preferred option, as we did, Bouhadba et al reported laparoscopic excision of ASH (10).

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