Huge Isolated Retrovesical Hydatid Cyst Affecting a Little Girl

Azzawi M Hadi*

Department of surgery, Tikrit medical college, Iraq

*Corresponding author: Azzawi M Hadi, Department of surgery, Tikrit medical college, Iraq. Email: azzawihadi@gmail.com

Submission: April 23, 2018; Published: July 03, 2018

Abstract:

A 8 year old girl presented with pain abdomen & frequency. Ultrasound & CT scan of abdomen showed hydatid cyst Retrovesical area. Surgery was performed and pathology of the excised mass confirmed the diagnosis. The rarity of the case of primary retrovesical hydatid cyst in such early age and such a huge size was the urging cause to present it.

Keywords: Hydatid cyst; Retrovesical; Echinococcus granulosus

Case Description

A 8-year-old girl presented with 4 months abdominal pain which was mild but continues, partially responding to analgesics, she also had frequency and nocturia, not associated with dysuria, nor hematuria. She had no remarkable systemic manifestations, nor a significant evidence in the past medical, and surgical history. She completed her vaccination schedule in time. She was born and lives in rural area.

On examination she was very well, and no finding can be detected other than mild lower abdominal tenderness, with palpable soft mass. Urinalysis was normal, as well her blood and biochemical indexes. Abdominal ultrasound requested, the report said that she had normal liver abdominal organs, with a cystic mass posterior to the bladder of 7x6cm, which contain echo free fluid, gives a deferential diagnosis of mesenteric cyst, big ovarian cyst, congenital diverticulum. Axial CT scan confirmed these findings as shown in (Figure 1).

Figure 1: CT scan findings, 7x6cm retrovesical cyst
Surgical exploration done by lower abdominal pfannenstiel incision, the mass was in midline, and was located in posterior to the bladder; pushing the posterior bladder wall anteriorly. All other abdominal organs were normal as well the chest views of the CT were normal. Surgery was performed to excise the mass. The cyst contents aspirated, which was clear fluid, and there was an endocyst of hydatid cyst that have been proved by the microscopic examination which revealed a laminated membrane, (Figure 2). Post operatively she had a smooth recovery, and sent home on albendazole tab 200mg daily in two divided doses for 3 months. The girl followed for one year by abdominal ultrasound, that shows no evidence of recurrence.

Discussion

Echinococcus granulosus the parasite that causes hydatid disease is endemic in many areas of the world. Hydatid cysts can develop in almost any part of the body [1]. In most cases, the first and most important locations for this parasite are the liver and the lungs. Hydatid disease of the urinary tract is uncommon, accounting for only 2-3% of all hydatid cases [2]. If hydatid cysts located at some the unusual sites, it may cause a diagnostic uncertainty. Peritoneal cavity hydatid cysts or pelvic hydatid cysts are usually result from rupture of a primary liver focus that occur spontaneously or during surgical management [1]. Although the most common site of echinococcal involvement is the liver, yet cystic echinococcus infestation can occur in any part of the body and should be considered in the differential diagnosis of any cystic masses [3]. Isolated retrovesical hydatid cyst, however, is very rare site with a few cases reported in literature only [4-6]. Unusually retrovesical hydatid cyst can be a cause for retention of urine [7,8]. It can presented as an extraperitoneal cyst, as well in the retrovesical area [9].

The diagnosis is based on clinical findings, imaging techniques and serology. Ultrasound is useful as a first line investigation in evaluation of cysts in the abdomen. Computed tomography and MRI are indicated in disseminated disease, lesions located out of the abdomen, sub-diaphragmatic sites, and pre-operative evaluation [10,11]. Abdominal ultrasound & native CT scan regarded helpful in diagnosis of abdominal hydatid disease and prove the suspicion of the clinical presentation. Although there was a unilocular cyst in our case, the multilocular appearance of the cyst supported by the density of the cyst contents indicated that the cyst was alive & viable [12,13]. Sagittal fast spin-echo T2-weighted MR image clearly identifies small cystic lesions representing daughter cysts [14]. The Dèvè’s classical theory states that fissuring or rupture of a primary hepatic, splenic or mesenteric cyst may seed its contents to the abdominal cavity, leading to a new lesions, while the primary cyst may heal and disappear, leaving behind a scar that may not detected by any diagnostic tools [9,11,15]. Haematogenous or lymphatic dissemination is another theory tried to explain the absence of the primary visceral lesion, and the absent evidence of peritoneal seeding [9,11,15]. Other hypothesis said that the larva may remained in the rectal ampulla and migrated through the haemorrhoidal vessels to reach a perirectal or retrovesical location [16].

Surgical treatment of Symptomatic large hydatid cysts by cysto-pericystectomy still the gold standard procedure [17]. Preoperative and postoperative use of albendazole have been found to decreases the viability of cysts at the time of surgery and significantly reduces the chances of cyst recurrence. Thus, albendazole is regarded as an effective adjuvant therapy in the management of hepatic hydatid cyst [18]. The cyst's growth rate is of about 10mm per year [19], but it is surprising that the cyst in this case reaches such a size at this age.

Conclusion

Hydatid cyst should be regarded as one of the differential diagnoses in patients with an isolated retrovesical cysts discovered during routine assessment for other conditions, such as patients presented with nonspecific symptoms of lower urinary tract, because prophylactic measures need to be considered in the operation to take spillage prevention in consideration.

References


