Bowel Occlusion Secondary to Retrovesical Hydatid Cyst: Exceptional Complication for Atypical Localization

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Abstract

Background: Hydatid pathology is endemic in our region. Although hepatic and pulmonary sites are the most common, the parasite can implant in any part of the body. We will report an exceptional case of retro-vesical hydatid cyst complicated by bowel obstruction.

Case Presentation: 45-year-old patient, admitted to the emergency department for an occlusive syndrome, the clinical examination found, distended tympanic but flexible abdomen, rectal examination: empty bulb. The biological assessment was without particularity. An abdominopelvic CT showed a cystic mass in the Douglas pouch with caliber disparity of bowel and intestinal distension upstream evoking a retro-vesical hydatid cyst. The patient was operated; a monobloc resection of hydatid cyst was performed. The postoperative course was simple. The biological analysis confirmed the diagnosis of retro-vesical hydatid cyst.

Conclusion: The retrovesical localization of hydatid cyst is rare, often asymptomatic, the clinical signs occur at a stage where the volume of the cyst is quite important. The diagnosis is often made by ultrasound coupled with CT. The hydatid serology has a great value of diagnostic orientation. The treatment is surgical based on a total perkystectomy. In an endemic area, any pelvic cystic mass must evoke a hydatid cyst.

Introduction

Hydatid pathology is endemic in our region. Although hepatic and pulmonary sites are the most common, the parasite can implant in any part of the body. Pelvic retrovesical location represents less than 1% of cases in the literature [1]. We will report an exceptional case of secondary acute intestinal obstruction to a retro-vesical hydatid cyst, by trying to meet the various challenges identified by this pathology.

Case Presentation

A 45-year-old patient, with no particular pathological history, admitted to the emergency department for an occlusive syndrome dating back to 3 days before his hospitalization. The clinical examination found a patient in fairly good general condition, temperature 37°C, distended abdomen tympanic but flexible, digital rectal examination: ampoule rectal empty. The biological assessment revealed leukocytosis at 12000/mm³, a slight functional renal failure. X-ray of the abdomen without preparation showed intestinal hydro-aerial levels. An abdominopelvic CT showed an intestinal distension with caliber disparity regarding a cystic mass in the Douglas pouch, hypodense of 7 cm in diameter, well limited evoking a retro-vesical hydatid cyst. Another hydatid cyst type 1 of 3 cm in diameter was located in the spleen (Figures 1-3). The liver was normal. The chest X-ray did not show a thoracic location. The patient was admitted to the operating room, an umbilical laparotomy was performed. After exposure and retro-grade emptying, exploration found a cystic mass occupying the Douglas pouch with intimate adhesions with the ileum responsible for caliber disparity with intestinal distention upstream, no other localization was found apart the hydatid cyst of the spleen. Hydatid cyst monobloc resection was performed after its release from its bladder and parietal attachments and after aspiration of its contents to prevent an intraoperative rupture responsible for recurrence (Figure 4 and 5). Splenic hydatid cyst was respected. The patient discharged at D+3 with no postoperative complications. The biological analysis of the cyst fluid as well as the hydatid serology confirmed the diagnosis of retro-vesical hydatid cyst. The patient received postoperative Albendazole for 3 months and follow up done 6 months after surgery with no complications.
Discussion

Retrovesical hydatid cysts are considered to be an “aberrantly” or “ectopic” localization and result from parasite development in sub- and retrovesical fat. They can be divided into two types: those with intraperitoneal development, and those with peritoneal development [2,3]. The mode of contamination is not well understood. It is most often secondary to a rupture of hepatic hydatid cyst in the peritoneal cavity with daughter cysts who continue their development. A secondary endothelium excludes them from the peritoneal cavity. However cases of primitive pelvic hydatid cyst as is the case of our patient have been reported. They are probably secondary to hematogenous contamination after passing through the liver and lung filter by the parasite. This form can be retained only in the absence of other hepatic or pulmonary localization [4-6]. This affection has a slow and silent evolution which explains why the signs as is the case of our patient who did not complain of any functional symptomatology before the installation of the occlusive syndrome. Bowel obstruction may be secondary to extrinsic compression or development of fleshy adhesions with the gastrointestinal tract. Diagnosis is often carried by imaging. Abdominal-pelvic CT is the gold standard for occlusive complication. Intestinal distention may interfere with ultrasound scanning [1,7]. The echographic classification of liver hydatid cyst is also valid for pelvic hydatid cyst [8]. Differential diagnosis may occur with ovarian cyst, ovarian tumor, seminal vesicle cyst, large ectopic ureterocele, posterior bladder diverticulum [7,9]. In this case, the use of biology with serology is very useful. Magnetic resonance imaging allows the analysis of pelvic reports of the hydatid cyst inaccessible to CT. It also allows in case of doubt to make the differential diagnosis with perirectal, and vesigial tumors of nerve or bone [4]. Surgical treatment remains the only therapeutic option. The monobloc resection of the cyst after protection of the operative field should be carried out if possible, otherwise resection of the protruding dome is recommended in case of close adhesion with the neighboring structures. Medical treatment based on Albendazole can be prescribed postoperatively on a case by case basis. There is no consensus on the indications and/or duration of treatment [7,10].

Conclusion

The retro-vesical localization of hydatid cyst is rare, often asymptomatic, the clinical signs occur at a stage where the volume of the cyst is quite important. The diagnosis is often made by ultrasound coupled with CT. The hydatid serology has a great value of diagnostic orientation. The treatment is surgical based on a total perkystectomy. In an endemic area, any pelvic cystic mass must evoke a hydatid cyst. Big effort must be expended to combat this affection which is a public health problem in our country.

References

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