

MESENTERIC HYDATID CYST: AN UNUSUAL PRESENTATION

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Hydatidosis is a common zoonosis that affects a large number of humans and animals, especially in poorly developed countries. Mesenteric hydatid cysts are rare intra abdominal lesions with variable clinical symptoms and signs that make preoperative diagnosis difficult. The treatment is surgical excision followed by mebendazole for four months. We report this unusual case where preoperative diagnosis of right tubo-ovarian mass was made and the diagnosis of mesenteric hydatid cyst could be made only on laparotomy and microscopic examination after surgery.

Keywords: Hydatidosis; Laparotomy; Mesenteric Hydatid cyst; Tubo ovarian mass

Hydatid disease is a zoonotic infection caused by larval stages of cestode *Echinococcus granulosus* (family taeniidae) and rarely by *Echinococcus multilocularis*. Dogs and other carnivores are definitive hosts whereas sheep and ruminants are intermediate hosts. Man becomes an accidental host by consuming vegetables and water contaminated with hydatid ova. Although liver (75%) and lung (15%) are the most commonly involved organs, the disease can be seen anywhere in the body (10%) [1,2]. Bickers [3] after reviewing 532 cases of hydatid disease from an endemic area over a 20 year period recorded only 12 instances of hydatid cyst in pelvis.

Case Report

A 35-year-old lady para two live two presented with pain lower abdomen for two years. The pain was dull in nature. There was no change in her bowel and bladder habits. She was para two live two and was using barrier contraceptive. Her menstrual cycle was regular. On physical examination vitals were stable. Her abdominal examination was unremarkable. On per speculum examination cervix and vagina was healthy. Per vaginum examination revealed normal size anteverted uterus deviated to left side. The mobility of the uterus was restricted. A large cystic mass measuring about

6×6 cm was palpable in the right fornix which was adherent to the uterus. Left adnexa were normal. On per rectal examination there was no nodularity in the pouch of Douglas and rectal mucosa was free. The routine laboratory investigations were within normal limits. X-ray chest and ultrasound (USG) upper abdomen was normal. USG lower abdomen and pelvis showed normal size uterus with a large multicystic septated mass in the right adnexa measuring 11×7 cm. Left adnexa was normal. Her CA-125 was normal. Diagnosis of right adnexal cyst was made and she was planned for exploratory laparotomy. Intra operatively a cyst measuring 10×6 cm was seen arising from the mesentery and was

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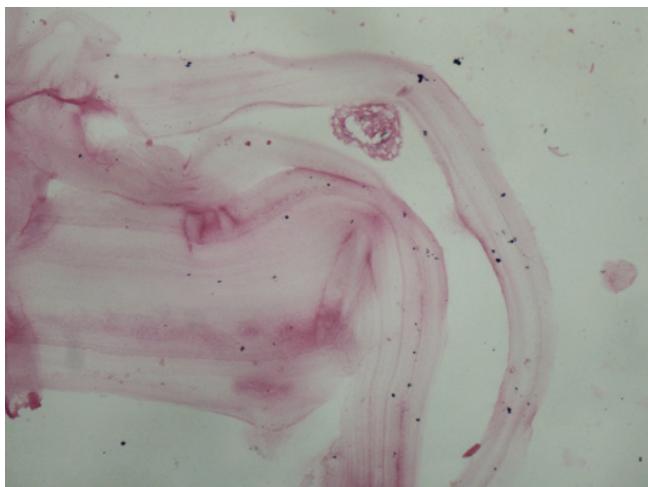


Fig. 1. Photomicrograph showing acellular laminated ectocyst and a single scolex (H&E, $\times 200$).

found to be badly adherent to the right ovary, fallopian tube and posterior surface of uterus and it could not be separated from the uterus and ovary. Removal of mesenteric cyst and right salpingo-oophorectomy was planned but total abdominal hysterectomy was also done as patient insisted on it. The whole specimen was sent for histopathological examination. Cut sections of mesenteric cysts showed glistening white multiple daughter cysts and microscopic examination revealed scolices with hooklets of *Echinococcus granulosus* (Fig. 1).

Cervix, right ovary and fallopian tube were unremarkable. Microscopic examination of uterus showed simple endometrial hyperplasia and myometrium was unremarkable. On taking history retrospectively it was found that infection was probably accidentally acquired in our patient, there was no history of pets nor a visit to an endemic area. Patient recovered well and was discharged on the tenth postoperative day with albendazole (10 mg/kg body-weight/day) for six months.

Discussion

Mesenteric hydatid cyst is the most unreported site so far as literature goes. There are rare lesions with prevalence of 1:100,000 in adults and 1:20,000 in children [4]. Hydatid disease is endemic in cattle grazing areas particularly Australia, New Zealand, Middle East, India, South America, and Turkey [5]. Most common locations of hydatid cyst are liver and lungs but unusual locations such as brain, heart, pericardium, kidney, intraperitoneum, retroperitoneum, bone, soft tissue and breast have been reported in literature. Extra

hepatic localization frequently occurs after hepatic involvement. Primary peritoneal echinococcosis is very rare and has been reported to occur in 2% of all abdominal hydatid diseases [6]. Clinical manifestations vary with the site and size of the cyst which ranges between a few centimeters and ten centimeters and symptoms are due to space occupying effect within organs or systemic reactions due to perforation. Fortunately they behave mostly as benign tumors, while malignancy accounts for 2% of cases [4].

The correct preoperative diagnosis of hydatidosis is difficult due to its rarity, lack of specific symptoms, diverse imaging appearances and striking resemblance between hydatid cyst and malignant disease of the related organ. Singh et al. [7] reported a case of primary hydatidosis of female genital tract for which pre operative diagnosis of malignant ovarian cyst was made and total abdominal hysterectomy with bilateral salpingo-oophorectomy was done. The diagnosis of hydatid cyst adherent to uterus and adnexa was made after histopathological examination. Okumuş et al. [8] also reported a case in which the primary involvement was of the uterus but the diagnosis was confirmed only by microscopic study after surgery. However the diagnosis may be possible by clinical finding, imaging studies and laboratory tests including echinococcal titers. The imaging modality of choice is abdominal ultrasonography. Computed tomography scan can reveal that the cyst is not arising from other organs like kidney, pancreas or ovary [9]. Surgery along with postoperative albendazole or mebendazole therapy is the treatment of choice for primary abdominal hydatid disease. It has been suggested that preoperative course of albendazole should be considered in order to sterilize the cyst, decrease the chances of anaphylaxis, decrease the tension in the cyst wall (thus reducing the risk of spillage during surgery) and to reduce the recurrence rate postoperatively [10,11].

Thus, we conclude that isolated mesenteric cyst is a unique manifestation of hydatid disease. Infection was probably accidentally acquired in our patient, there was no history of pets nor a visit to an endemic area. Owing to its rarity in area other than endemic areas and variable imaging appearances preoperative diagnosis of peritoneal/mesenteric hydatid disease is a clinical challenge so a suspicion of this disease is justified in all patients presenting with cystic mass especially in endemic areas.

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