Case Report

Isolated primary hydatid cyst of the pancreas: Management challenges of a cystic masquerade

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Abdominal hydatid cyst disease mostly involves the liver. Involvement of the pancreas as an isolated primary organ is rare accounting for < 2% of all systemic echinococcosis cases. It mostly involves the head of the pancreas. Symptoms depend on the location, size, and associated complications; therefore, it can have varied presentations including acute pancreatitis. On imaging, it can mimic other common pancreatic cystic lesions like pseudocyst or cystic neoplasm. Accurate preoperative diagnosis is usually difficult and requires a very high index of suspicion even in endemic areas. Herein, a case of primary isolated hydatid cyst of the pancreas that was initially diagnosed and managed as acute pancreatic pseudocyst is reported.

Key Words: Echinococcosis; Pancreatic cyst; Pancreatic pseudocyst; Pancreatitis; Membranes

INTRODUCTION

Hydatid disease is a zoonotic parasitic disease caused by the echinococcal infection. Hydatid cysts in humans most commonly result from Echinococcus granulosus infection. The liver (50%–77%) is the most common site of infection, while the other organs are less involved; lungs (15%–47%), spleen (0.5%–8%), and kidney (2%–4%) [1].

Primary involvement of the pancreas by the hydatid disease is very rare. It accounts for < 2% of total systemic echinococcosis cases [2] with only 33 cases reported in literature since 2011 [3]. Pancreas being a rare site of involvement, primary isolated pancreatic hydatid cyst (PHC) may pose diagnostic and management challenges as it may mimic common pathologies like pseudocyst [4,5].

Herein, we report a case of a primary PHC that mimicked pancreatic pseudocyst and was managed laparoscopically.

CASE

A 20-year-old female presented with intermittent epigastric pain for 8 months with no history of nausea, vomiting, altered bowel habits, fever, jaundice, anorexia, weight loss, or a history of similar pain in the past. General physical examination was normal. Abdominal examination revealed a small vague non-tender lump in the epigastric region.

The patient was evaluated for the symptoms 6 months before at another center. All the routine hematological investigations were normal, but serum amylase (422 U/L, normal range 28–100 U/L) and serum lipase (1,048 U/L, normal range < 67 U/L) levels were raised. Ultrasonography showed a well-defined cystic lesion of 7.4 cm × 7.3 cm in the pancreatic head suggesting a pancreatic pseudocyst. The rest of the pancreas and gallbladder were normal. Contrast-enhanced computed tomography (CECT) (Fig. 1A, 1B) and magnetic resonance imaging (MRI), including magnetic resonance cholangiopancreatography (MRCP), (Fig. 1C, 1D) revealed a well-defined cystic lesion (8 cm × 8 cm, wall thickness 3 mm) in relation to the pancreatic head, bulky pancreatic body and tail, and prominent main pancreatic duct. No evidence of any septa or solid component...
was noted within the cyst. Pancreatic pseudocyst was suggested with a differential of choledochal cyst and cystic neoplasm. The patient was managed conservatively as a case of pancreatic pseudocyst with partial symptomatic relief.

Since there was no significant improvement in symptoms in the next few months, the patient consulted at our center. The pancreatic cystic lesion was re-evaluated with repeat cross-sectional imaging. CECT of the abdomen showed a well-defined thin-walled hypodense cystic lesion (5.3 cm × 6.7 cm × 7.6 cm) with multiple, thin, linear, and wavy high attenuating structures suggesting floating membranes within it (water Lilly sign) in the pancreatic head and a thin rim of pancreatic parenchyma surrounding it (Fig. 2A, 2B). MRI of the abdomen also showed similar findings (Fig. 2C); hyperintense cystic lesion with multiple thin T2 hypointense floating membranes in the pancreatic head. The rest of the pancreas, including the main pancreatic duct, was found to be normal on both of these imaging. The biliary ductal system was found to be normal throughout its course on these imaging. There was no evidence of a similar cystic lesion in any other abdominal organ or in the chest. Based on these imaging findings, a diagnosis of primary cystic echinococcosis of the pancreas was suggested which was classified as CE3a stage (cyst with liquid contents and detached endocyst) as per the WHO-International Working Group on Echinococcus (IWGE) classification [6,7].

Routine hematological investigations, liver function test, and serum amylase (55 U/L) were within the normal range. Echinococcal serology was non-reactive (immunoglobulin [Ig] G 6 U/mL, normal range < 10 U/mL; IgM 10 U/mL, normal range < 20 U/mL).

After the diagnosis of primary isolated PHC, the patient underwent laparoscopic partial cystopericystectomy and omentoplasty with perioperative albendazole therapy (15 mg/kg/day). The laparoscopic procedure was performed using four ports (Fig. 3) with the patient in the French position. Initial diagnostic laparoscopy revealed a large (~6 cm × 6 cm) thick wall cyst in the pancreatic head and uncinate process, splaying the...
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duodenal C loop, bulging through the transverse mesocolon, and lying in-close approximation with the right and middle colic vessels (Fig. 4A). The anterior aspect of the cyst was exposed by dividing the overlying gastrocolic omentum and mobilizing the proximal transverse colon (Fig. 4B). The lesser sac was not entered at this stage to prevent its contamination with cyst contents during subsequent steps of the procedure. The cyst was isolated by placing 10% Povidone-iodine (scolicidal agent) soaked roller gauze around it (Fig. 4B); cyst decompression was done with a laparoscopic aspiration needle (Fig. 4C); scolicidal agent (10% Povidone-iodine solution) was injected into the cyst cavity with a sterilization time of 15 minutes; the contents aspirated again; the cyst was opened carefully at the most prominent point with an ultrasonic energy device (Fig. 4D); cyst contents were evacuated with a 10 mm suction cannula (Fig. 4E); cyst cavity was rinsed with scolicidal agent (10% Povidone-iodine) again; and then partial cystopericystectomy (derooﬁng) was done using ultrasonic shears (Fig. 4F).

To prevent any spillage of cyst contents during the procedure, 2 suction cannulas were used during the initial opening and evacuation of the cyst cavity (Fig. 4D) and one lip of the cyst wall (at the opening site) was retracted upwards throughout the cyst evacuation process (Fig. 4E). Care was also taken during the “cystopericystectomy” step to protect adjoining luminal structures (e.g., duodenum, stomach, colon, pancreas) and vascular structures (e.g., right gastroepiploic vascular arcade, right colic, middle colic, and superior mesenteric vessels).

After partial cystopericystectomy, the cyst cavity was inspected for residual contents (Fig. 4G), irrigated again with normal saline and 10% Povidone-iodine, omentoplasty was done (Fig. 4H), and the cavity was drained with a closed suction drain (16F).

Excised cyst wall and membranes (Fig. 5) were sent for histopathological examination that showed a lamellated membrane with an inner, degenerated germinal layer comprising degenerated protoscolices with hooklets and calcified areas, along with chronic pancreatitis changes in the adjoining pancreas (Fig. 6). These findings conﬁrmed the diagnosis of PHC.

Fig. 3. Port positions (marked with ‘X’) used in the laparoscopic partial cystopericystectomy procedure. Two 10 mm ports (at the umbilicus [marked with ‘*’ for the camera, second for surgeon’s left working hand/suction/specimen extraction), two 5 mm ports (lower one for surgeon’s right working hand and upper one for assistant/surgeon).

Fig. 4. (A) Intraoperative image showing a large cystic lesion (*) in the head of the pancreas. (B) Cyst exposure after the division of overlying gastrocolic omentum. The cyst has been isolated from the surrounding structures by roller gauze soaked in scolicidal agent (10% Povidone-iodine). (C) Cyst content is being aspirated in a controlled manner. (D) Cyst is being opened with an ultrasonic energy device in a controlled manner with 2 suction cannulas placed close by to contain spillage. (E) Cyst contents (* membrane) are being evacuated with a 10 mm suction cannula. (F) Partial cystopericystectomy is being performed using ultrasonic energy device. (G) Opened up cyst after partial cystopericystectomy. (H) Omentoplasty.

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There was a pancreatic leak in the postoperative period with a small fluid collection in the residual cavity that was managed conservatively. The patient was discharged on the 9th post-operative day on albendazole (15 mg/kg/day) for 8 weeks. The patient is doing well on follow-up at 3 months.

**DISCUSSION**

Pancreatic involvement in hydatid disease is very rare with a reported incidence of 0.14% to 2% [8]. Most of the aspects of primary PHC, including its natural history, diagnosis, and management, are not clear so far [8]. The pancreas is thought to be involved primarily through hematogenous dissemination [3]. Pancreatic hydatid involvement is solitary in a majority of cases [1,3]. It may remain asymptomatic and detected as an incidental finding [9] or it may present with a variety of symptoms depending on its size, location, and associated complications. Common symptoms include epigastric/left upper quadrant pain and/or a lump, nausea, vomiting, and fever [1].

Acute pancreatitis is a rare presentation that may occur due to pancreatic ductal obstruction by external compression by the cyst or by internal occlusion by scoles after cyst-ductal communication [10]. Other rare complications include sinistral portal hypertension (by pancreatic tail lesions), rupture into the biliary tract, cholangitis, pancreatic fistula, and recurrent pancreatitis [1]. Necrotising pancreatitis has also been reported [10]. In the present case, abdominal pain was the primary symptom although the patient had acute pancreatitis (mild) initially. Associated chronic pancreatitis changes in the adjoining pancreas, as revealed on the histopathological examination and most probably due to chronic cyst compression, have not been reported so far.

Preoperative diagnosis of primary PHC is very difficult because symptoms are non-specific and characteristics radiological findings are not always present. As the cystic echinococcosis evolves, morphological changes in the cyst occur. Based on the morphological changes in imaging, cystic echinococcosis has been classified into five types or stages as per WHO-IWGE classification [6,7]. CL is a very initial or latent stage (unilocular cyst with no cyst wall or any pathognomonic signs). CE1 is an active stage with unilocular cyst having cyst wall and hydatid sand. CE2 is also an active stage with multivesicular cyst (rosette-like) with cyst wall. CE3 is a transitional cyst that has started to degenerate but may still produce a daughter cyst, thus it is a stage between an active cyst and inactive cyst; CE3a refers to a cyst with liquid contents and detached laminated membrane/endocyst and CEb refers to unilocular cyst with a predominantly solid component with daughter cysts. CE4 is an inactive phase and refers to a cyst with solid heterogenous de-

![Fig. 5. Evacuated hydatid cyst membranes.](image)

![Fig. 6. (A) Hydatid cyst wall showing lamellated membrane with inner degenerative germinal layer (H&E, low power 200×). (B) The degenerative protoscolices of echinococcus with hooklets (arrow) (H&E, high power 400×). (C) Pericyst (H&E, low power 200×). (D) Adjoining pancreas showing chronic pancreatitis changes with acinar atrophy and interstitial fibrosis (H&E, low power 200×).](image)
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None.

CONFLICT OF INTEREST

No potential conflict of interest relevant to this article was reported.

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Conceptualization: VG, RS, EJ. Data curation: All authors. Methodology: PKK, VG, RS, EJ, HW, JS. Visualization: PKK, VG, RS, EJ. Writing - original draft: PKK, VG. Writing - review & editing: All authors.

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