A recurrent cyst hydatid abscess fistulised to the bronchial tree

Bronşiyal ağaca fistülize olan nüks kist hidatik absesi

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Abstract

Human cystic echinococcosis is a zoonosis transmitted by ingesting Echinococcus granulosus ova eliminated from infected dogs. A 41-year-old man was admitted to the hospital with symptoms of abdominal tenderness located in the epigastric region, fever and vomiting. We diagnosed a recurrent hydatid disease and after admission to our hospital he began to cough with greenyellowish colored sputum with abscess. Posteroanterior chest X-ray and right pleural US revealed no fluid collection but right lower lung atelectasis and diaphragma elevation. With these results we performed an operation for broncho-biliary fistula. He was discharged from the hospital on the ninth postoperative day. Careful assessment and early treatment of septic complications are essential in the successful treatment of this rare condition.

Keywords: Bronchobiliary fistula, echinococcosis

Introduction

Human cystic echinococcosis is a zoonosis transmitted by ingesting Echinococcus granulosus ova eliminated from infected dogs. A broncho-biliary fistula (BBF) is defined as an abnormal communication between the biliary system and the bronchial tree due to trauma, hemihepatectomy and necrotic hepatic infections, such as hydatid disease [1-3]. Several factors participate in promoting intrathoracic evolution of hydatid cysts of the hepatic dome.
1. The intrathoracic negative pressure tends to aspirate the hepatic hydatid cyst.
2. The mechanical compression maintained by the cyst on the diaphragm results in local ischemia, which can lead to muscle erosion.
3. Sepsis in the hepatic cyst can also contribute to muscle necrosis close to the cyst.
4. In the case of biliary fistulas, the caustic property of biliary secretions can lead to chemical erosion of the diaphragm and the bronchi.

For hydatid disease that fistulised to the thoracic cavity a classification had been established by Mestiri and colleagues, which was improved by Dévé (Table 1) [4]. According to this classification our case was a type I bronchobiliary fistula. In this report, we describe a case of bronchobiliary fistula in a man with recurrent hydatid disease abscess that was treated via laparotomy.

**Table 1. Classification established by Mestiri and colleagues, which improves on that proposed by Dévé.**

| Type I: Direct fistulas in the bronchial tree |
| IA: Small bronchial fistulas |
| IB: Major bronchial fistulas |
| Type II: Constitution of an intrapulmonary cavity |
| IIA: Small bronchial fistulas |
| IIB: Major bronchial fistulas |
| Type III: Constitution of an intrapleural pouch |
| IIIA: No bronchial fistulas |
| IIIB: With bronchial fistulas |
| IIIC: With pleuroparietal fistulas |
| Type IV: Rupture in the pleural cavity |
| IVA: Acute rupture: hydatic and biliary empyema |
| IVB: Secondary pleural hydatidosis |

**Case report**

A 41-year-old man was admitted to the hospital with symptoms of abdominal tenderness located in the epigastric region, fever and vomiting. Ultrasonography (US) and abdominal computerized tomography (CT) demonstrated a hydatid cyst recurrence at the dome of the liver nearly 12x10 cm in size (Figure 1a) and abscess with air-fluid level (Figure 1b). The patient underwent an operation for hydatic disease in another hospital 5 months ago. After this operation because of the persistence of symptoms 3 months later, an endoscopic retrograde cholangiopancreatography (ERCP) with sphincterotomy and puncture, aspiration, injection, and reaspiration (PAIR) was performed. Two months after these efforts, the patient presented to our hospital with the symptoms as described above. We hospitalised the patient and initiated systemic antibiotics. 2 days after admission he began to cough with greenyellowish colored sputum with abscess. The temperature of the patient was 38.1°C. No jaundice was detected, but there was mild tenderness in the right quadrant. Laboratory studies revealed increased levels of alkaline phosphatase, bilirubin, and white cell count.

Posteroanterior chest X-ray (Figure 2) and right pleural US revealed no fluid collection but right lower lung atelectasis and diaphragma elevation. With these results we performed an operation for BBF. The exploration findings was a cyst hydatid abscess pouch that had germinative membrane in it and this membrane penetrated to the pleural space (Figure 3). The cyst pouch was cleaned with scolicidal material and germinative membrane was excised, the diaphragmatic defect was primary repaired with tube thoracostomy. He was discharged from the hospital on the ninth postoperative day without any complication.
Figure 1a. Hydatid cyst recurrence at the dome of the liver nearly 12x10 cm in size and (b) abscess with air-fluid level.

Figure 2. Posteroanterior chest X-ray revealing right lower lung atelectasis and diaphragma elevation.

Figure 3. Peroperative findings of the cyst hydatid abscess pouch that had germinative membrane in it and this membrane had penetrated to the pleural space.

Discussion

Hydatid disease is endemic in Turkey, with high rates of infection (1/20,000-1/50,000) [1] one of the main complications of the hydatid cyst of the dome of the liver is its rupture into the bronchial tree [3, 5]. This occurs due to a combination of pressure erosion...
from an expanding hydatid cyst to the diaphragm and the destructive effect of superimposed infection. If enough adhesions precede the erosion, the cyst will rupture into the pulmonary parenchyma causing pneumonitis and BBF [3, 6].

In many series, bronchography is not considered to be a useful diagnostic tool; we also didn’t perform it to our patient. Today there are less invasive methods such as CT scan, ultrasonography and nuclear imaging that turned to be very important diagnostic tools [7, 8]. Management has traditionally been surgical resection, but recently embolization of the fistula [9] and stent placement to relieve common bile duct obstruction have been described [10]. Surgery still remains the treatment of choice in case of echinococcosis [11, 12]. BBF is a complicated surgical problem because it is characterised by a complex of pathological lesions involving organs in continuity in the thoracic and abdominal cavities as well as the interposed diaphragm. The characteristics of the primary cystic lesions, the site and seriousness of diaphragmatic and pulmonary secondaries, and the frequency of biliary complications affect our surgical approach to BBF [13].

Many authors are in favour of a transpleural approach. It gives a good exposure and the lung complications can be treated. If abdominal complications occur the incision can be extended to the abdomen or a separate abdominal incision made [14].

We do not agree with this opinion. Any radical treatment of the cyst attempted via standard thoracotomy is unsafe because of lack of control of the hepatic pedicle, inferior vena cava, and hepatic veins, and results in serious intraoperative risks. In our case we performed a surgical approach via laparotomy that would be less morbid for the patient. The BBF was type 1 according to Dévé’s classification and after the disconnection of the fistula we thought that the patient would benefit from this intervention.

Thoracoabdominal incision is rarely reported in the literature because it is thought to cause increased morbidity [15]. Thoracoabdominal approach (TA) provides a view of the entire complex of the BBF. The section of the diaphragm provides simultaneous control of both the diaphragmatic surfaces and facilitates the blunt dissection of the diaphragm adhering to the cyst and to the lung. TA gives good access to the hepatic pedicle and to the biliary tract, enabling haemostatic manoeuvres, intra-operative cholangiography and any required surgical procedure on the biliary tract. Since 1990 thoracic portion of TA has been performed among the fifth or seventh intercostal space because it provides better access to the principal and segmental hila of the right inferior lobe of the lung. The choice of the route of access and the surgical technique should not depend on the surgeon’s preference, but rather be tailored to the requirements posed by the site, nature, and extent of disease.

A risk of delaying treatment is rupture of the lung cyst with widespread pulmonary soiling and acute respiratory failure. Careful assessment and early treatment of septic complications are essential in successfully treating this rare condition.

References