Primary Hydatid Cyst of the Diaphragm: A Case Report
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Abstract
Hydatid disease is a parasitic disease endemic in Romania which occurs frequently in liver and lungs, but it can also be present in almost any part of the body. We present the rare case of a 37-year-old male which was admitted in our service with a large hydatid cyst at the level of the right thoracic outlet and we were not able initially to establish its exact origin, liver or right lung. After imagistic investigations we decided for an abdominal approach considering that the cyst was liver related. Intraoperative findings showed a primary hydatid cyst of the diaphragm, a very rare entity that was successfully treated by the mentioned approach.

Keywords Echinococcus Granulosus; Hydatid Cyst; Diaphragm

Introduction
Hydatid disease (HD) is a parasitic disease caused by Echinococcus granulosus. The disease is endemic in Eastern Europe, including Romania which has a morbidity index of 5.6 to 100.000 people [1]. It occurs frequently in the liver (59-75%), but it can also be present in the lung (27%), kidney (3%), bone (1-4%) and diaphragm (1%), the last being generally associated with liver hydatidosis [2,3]. In this report we present a case with primary hydatid cyst of the diaphragm, with no other cysts present, which was successfully managed in our department.

Case Report
A 37-year-old male, with no history of surgery, was admitted in our department for thoracoabdominal pain for 3 days. At admission the patient was without fever and in good general condition, despite suffering of thoracoabdominal pain. Laboratory tests showed slightly elevated leucocytes and no eosinophilia was present. Chest X-ray performed at admittance revealed right-sided subpulmonic opacity with elevation of the right hemidiaphragm. The CT scan performed the next day showed a giant hydatid cyst extending from the abdominal cavity in to the right thoracic cavity but did not offer any information regarding the origin of the hydatid cyst (Figure 1). As it was considered that we deal with a liver cyst, we have decided for an abdominal approach.

An exploratory laparotomy was performed on the third day, during which time we discovered a giant hydatid cyst (Figure. 2) between the muscle fibers of the diaphragm, without liver or lung involvement, pushing down on the liver and compressing the inferior right pulmonary lobe. Careful dissection of the muscle fibers was carried out in order to avoid spillage in the abdominal cavity (Figure 3). Upon reaching the membrane of the cyst we decided to first evacuate the content with the help of a 16G needle. Next we irrigated the site with 90% alcohol solution in order to inactivate the remaining content, and finally we evacuated the cyst again using the same 16G needle.

Removal of the cyst’s membrane went without incidents, and closure of the defect was performed with individual silk suture points. Histological examination established the presence of muscle fibers in the pericyst and confirmed the diagnosis of hydatid cyst.

Postoperative recovery was uneventful and the patient was discharged on the fourth postoperative day and directed to a parasitological department for medical treatment.

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Discussion

The primary hydatid cyst of the diaphragmatic is a rare clinical entity defined as a cyst with no involvement of the pulmonary parenchyma and with no transmission from the abdomen to the thorax, whereas a secondary cyst is most likely caused by transdiaphragmatic migration from the posterior segments of the right hepatic lobe [4,5]. Diaphragmatic localization of the hydatid cyst is most likely caused by dissemination of the embryos through the arterial circulation [6]. Diagnosis is confirmed by the presence of muscle fibers in the pericyst during histological examination [7]. In the preoperative period careful topographic diagnosis between the diaphragm, lung, liver and abdominal localizations should be made either by use of CT or MRI examinations [8]. Although total excision of the cyst through the thoracotomy is considered an excellent approach [9], we prefer using the laparotomy in cases in which the CT examination doesn’t rule out liver involvement. Also, we have achieved good long term results without exciting completely the pericyst, simple removal of the membrane being sufficient. Suturing of the diaphragmatic defect is a must in order to avoid possible herniation and to ensure good pulmonary function.

Conclusion

The primary hydatid cyst of the diaphragmatic is a rare clinical entity and the diagnosis is challenging. Although total excision of the cyst through the thoracotomy is considered an excellent approach, we prefer using the laparotomy in cases in which the CT examination doesn’t rule out abdominal viscera involvement. A good diaphragmatic suture is mandatory to avoid possible herniation and respiratory dysfunction.

Conflict of interest

Authors have no conflict of interests to disclose.

References