Cystic echinococcosis, review and illustration of non-hepatic manifestations

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Abstract
Cystic echinococcosis (CE) or hydatidosis (hydatid cysts), is an infection with a wide spectrum of manifestations, from asymptomatic infection to fatal disease. Ultrasound (US) allows screening, diagnosis, differential diagnosis, treatment guidance and follow-up of CE under many circumstances. Hydatid cysts are predominantly observed in the liver but many other organs can be involved. As part of a series of publications, herewith we present a review describing the characteristic imaging features of the broad variety of organs which can be involved

Keywords: cystic echinococcosis (CE); hydatidosis; ultrasound (US); non-liver; imaging

Introduction
Cystic echinococcosis (CE) or hydatidosis (hydatid cysts), is an infection caused by the larval stage (metacestode) of the cestode (tapeworm) Echinococcus granulosus. In humans, it may result in a wide spectrum of manifestations, from asymptomatic infection to fatal disease.

The clinical manifestations [1] and imaging features of modern imaging for liver CE has been recently described [2]. In brief, infection most commonly affects the liver without generating symptoms. The evolution of cysts can be defined into stages according to the WHO classification [3] which updated Gharbi’s work [4]; from active stages of simple cysts and daughter cysts, those with the pathognomonic ‘water-lily’ sign, or daughter cysts with solid or calcified components, to non-viable forms with increasing fibrosis and calcification [1]. As cysts grow, local compressive effects and infiltration of surrounding structures can precipitate symptoms and a broad range of local complications [2]; secondary infection, rupture into the biliary tree, the peritoneal cavity, or through the diaphragm to the thorax.
In this review, we want to describe the wide variety of organ involvement of CE and to demonstrate characteristic imaging features.

**Pancreatic involvement**

Infection of the pancreas by hydatid cysts is very rare, even in endemic areas, where full body imaging is necessary for accurate diagnosis, particularly in instances of atypical deposits. Even when surgery is not possible, chemotherapy as treatment or prophylaxis against recurrence can be effective [5,6]. Migration to the pancreas occurs predominantly via hematological spread or peri-pancreatic lymphatic invasion. Very rare cases are reported via retroperitoneal spread [6]. Within the pancreas, the head is most frequently involved (57%), followed by the body (24%) and the tail (19%) [7]. Pancreatic hydatid cysts can cause non-specific symptoms, commonly related to the location and size of the cyst, but are usually asymptomatic. Those in the head of the pancreas can cause obstructive jaundice or pancreatitis. Large cysts in the body or tail of pancreas, most commonly cause abdominal pain [8].

The imaging features reported for pancreatic hydatid cysts include contrast enhanced computer tomography (CE-CT) demonstrating a hypodense mass in the uncinate process with a suggestion of hyperdense/enhancing curvilinear densities, magnetic resonance imaging (MRI) showing a non-enhancing cystic mass, and endoscopic ultrasound which might demonstrate a cystic structure with curvilinear floating membranes consistent with the water lily sign, pathognomonic of hydatid disease [9].

**Splenic manifestations**

Splenic involvement is an uncommon manifestation of hydatid cysts, accounting for only 4% of all abdominal cases. Most cases of splenic involvement are seen in conjunction with hepatic or peritoneal HD, with splenic seeding from primary cyst rupture and/or contiguous cyst growth [10,11]. The clinical manifestations include non-specific left upper abdominal pain, right upper abdominal pain with dyspepsia or they may be asymptomatic [12].

The imaging findings of splenic hydatid cysts are similar to those of hydatid cysts seen in other abdominal viscerca and range from purely cystic lesions to completely solid or calcified masses (fig 1). A solitary or multi-locular cystic lesion is the most common finding. Heavily calcified masses are indicative of non-viable cysts. Multiple lesions have also been reported in the spleen [10,13]. A spleen-saving approach is likely an effective method for the removal of CE spleen cysts given that it is a quick procedure and preserves the function of the spleen. Total splenectomy procedures may be required for particularly large, centrally located cysts [14].

**Pelvic manifestation**

Pelvic echinococcosis can be either primary or secondary, but are extremely rare, with an incidence of only 0.2-2.25%. Secondary pelvic hydatid cysts are more common and occur following the cyst rupture of a primary hepatic, splenic or mesenteric lesion. If no other associated cyst in the abdomen is identified, a pelvic hydatid cyst is considered to be primary [15]. Pelvic lesions present with symptoms usually due to mass effect. These may be confused with obstructive uropathy (due to compression of the urinary tract), ovarian cysts, appendicitis or even pelvic sepsis (when they rupture). An ultrasound followed by a CECT scan is usually required for diagnosis. MRI in most instances is not required, as it offers little additional information over CT [15] (fig 2). Hydatid cysts in the uterus can mimic other conditions such as multi-cystic ovarian tumor, hemorrhagic ovarian cyst,
endometrioma, cystadenoma, leiomyoma, etc [16]. Primary ovary hydatid disease in postmenopausal women must be carefully differentiated from ovarian malignancy [17,18].

Retrovesical hydatid cysts may have a varied and non-specific presentation. Ultrasound and CECT are both excellent imaging modalities for the detection of urinary tract hydatid cysts. The treatment of choice in this context is principally a careful and complete surgical excision, which can be achieved either by open or laparoscopic approaches [19].

**Lung involvement**

The lung is the second most common organ involved in adults, but the most common infected organ in children. The majority of patients with even multi-locular lung involvement are asymptomatic, particularly in children and young patients. However, chest pain, cough, dyspnoea and hemoptysis may occur. A single cyst may be found in 50% of cases, two in 20% and multiple cysts in 30%. Cysts are usually located in the lower lobes and may grow up to 10 cm [20-22]. Most of the cysts are Gharbi 1 or 2 cysts, calcification and daughter cyst formation being rare in lung hydatidosis [21] (fig 3). The use of lung ultrasound so far is limited but promising [23-27].

Pulmonary hydatid cysts are usually diagnosed with plain radiography and then further characterized by CT. US can depict lesions peripherally located or with pleural contact. The “wall sign” describes a double-layered wall in univesicular cysts and a double-layered septum in cases of multi-vesicular cysts [22] (fig 4). It has been
reported to have a specificity of nearly 100% for diagnosing pulmonary hydatid cysts [3]. In peripheral pulmonary disease, ultrasound may be an appropriate alternative to CT for children, as an easily available and non-irradiating modality [21]. The majority of patients with even multi-locular lung involvement are asymptomatic, particularly in children and young patients [28-30]. However, chest pain, cough, dyspnea and hemoptysis may occur. Complications include rupture with spilling of larva fragments and protoscolices into the bronchial tree and pleural cavity, bacterial super-infection, abscess formation, empyema, pneumothorax and systemic spread. Concomitant liver involvement occurs in 20% of patients, and is more common in children than adults [27,28,30].

Kidney and urinary tract involvement

Hydatid cysts of the kidney are rare, occurring in 3% of cases [20]. They may present with acute renal colic and hydatiduria. Specific features of urinary hydatid cyst may result after a cyst ruptures into the collecting system [10,31]. Involvement of the kidney is mainly due to parasitic dissemination through the portal venous system and retroperitoneal lymphatics to reach the renal fossae [32]. Renal hydatid cysts are generally located in the upper or lower pole of the kidney cortex, usually solitary and can grow up to 10 cm without symptoms. Complications include infection and rupture (18% of cases) into the urinary tract [33] (fig 5). The clinical manifestations of renal hydatid cyst otherwise nonspecific; back pain, dysuria, and hematuria are reported most frequently. The US signs are similar to hepatic hydatid cysts [20]. In cases of multi-locular cysts, differentiation from simple renal cysts, cystic nephroma or renal cell carcinoma is required [34,35] (fig 6-8). A unilocular cyst with a calcified rim or a multilocular cystic lesion with heterogeneity on CT suggests hydatid disease [36].

Other organ involvement

Involvement of organs other than those described above are even less frequently encountered but significant morbidity and mortality occurs. Systemic infiltrations affecting the central nervous system, musculoskeletal system, subcutaneous cyst with skin involvement, heart and pericardium with pericardial effusion or tamponade, ocular cysts and appendix involvement [37,38] can occur. Cysts are mostly asymptomatic, so that patients often present with clinical evidence of mass or vague symptoms. This explains the common delay to diagnosis, and lack of data on correlation to the real moment of infection [39]. The diagnosis is confirmed by clinical evidence, medical history including epidemiology and radiological support. US is frequently considered the gold standard in imaging of the cyst, whilst CT scan and MRI often offer improved definition of anatomic relations [40].
Brain

Multiple intracranial lesions can result from the rupture of an initial single intracranial cyst (in cystic echinococcosis) or from dissemination of systemic disease of the lung, liver or heart (cystic and alveolar echinococcosis) [41]. The two main subtypes, cystic and alveolar echinococcosis, present differently and have distinct imaging features in the brain. Most cases of intracranial cystic echinococcosis present with features of a space-occupying lesion such as headache, vomiting or focal neurological signs [40]. On CT or MR imaging, cystic echinococcosis characteristically appears as a well-defined, thin-walled cerebrospinal fluid-signal cyst, which is usually solitary but may contain multiple daughter cysts. There is usually no surrounding oedema unless secondarily infected [41] (fig 9).

Orbit and eye

Head and neck involvement of echinococcosis is a rare entity and extension of the hydatid cyst into the intraorbital region or infiltrating into the surrounding orbital bone is even rarer [42]. Clinical and radiological findings are important but may not be sufficient in the preoperative diagnosis. Propositis may be seen with orbital or infratemporal hydatidosis, and although rare should be considered a differential diagnosis [43]. The role of a radiologist is crucial in expediting the diagnosis of orbital lesions with the help of characteristic imaging features on ultrasound, CT or MRI. These also help identify complications where formulation of an early and effective management strategy is vital for preserving vision [44] (fig 10).

Conclusion

Echinococcosis hybrid cystic lesions are frequently encountered in clinical practice in specific geographic areas of the world. Hydatidosis affects multiple organs and has diverse imaging presentations. Application of the WHO classification system outside of the liver is of practical benefit. Knowledge of the specific imaging features enables accurate diagnosis, facilitates appropriate management and helps avoid complications.

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References

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