A 31-Year-Old Female with an Acute Episode of Cough and Hemoptysis

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WHAT IS YOUR DIAGNOSIS?

A 31 year-old female was admitted to the hospital with an acute episode of cough and hemoptysis, along with low grade fever. She was nonsmoker without any previous history of respiratory symptoms. Laboratory findings showed WBC count 12,200, NEUT: 74%, LYM: 21%, and EOS: 5%.

Figures 1 and 2. Show the CXR and chest CT of the patient.
Diagnosis: Ruptured hydatid cyst and diffuse pneumonitis

Chest X-ray (Fig.1) shows a homogeneous opacity in lower half of the left hemithorax with a wavy appearance in the upper border, silhouetting left heart border. There are also alveolar shadows in right paracardiac region. Chest CT (Fig.2) reveals cavitary consolidation in left upper lobe (LUL) with internal floating membrane mimicking ‘water lily’. Scattered bilateral alveolar opacities are also seen.

At bronchoscopy, a shiny membranous tissue was seen in the lumen of LUL bronchus, suggesting a ruptured hydatid cyst (Fig. 3). Because of pulmonary infiltration, TBLB was done, and pathological findings included lymphocytic and neutrophilic infiltration in the interstitium and alveolar space due to pneumonitis. Smear of bronchoalveolar lavage did not show hooklets and culture was negative for bacteriology. Serologic tests for hydatid cyst were positive. Surgery was performed, and a large hydatid cyst was resected (Fig 4). Parenchymal infiltration resolved subsequently.

DISCUSSION

Hydatid cyst disease is a parasitic infection and one of the important public health problems in endemic areas like Iran. Liver is a common site of hydatidosis. Pulmonary involvement is less common, and comprises about 25% of patients with hydatid cyst disease (1, 2). Chest x-ray is the most valuable diagnostic method for pulmonary hydatid cyst. The typical findings for uncomplicated hydatid cyst are homogeneous round masses with smooth borders that may cause parenchymal atelectasis and pleural reaction (3). Occasionally, in ruptured cysts, ‘Cumbo-sign’ and ‘onion peel’ may be seen when air invades between the pericyst and the endocyst (4). Calcification of hepatic cysts is common, while pulmonary cysts are rarely calcified (3).

Some conditions such as benign tumors, carcinomas, metastasis, inflammatory masses, fluid-filled cysts and abscess can mimic the radiologic features of pulmonary
hydatid cyst. CT scan reveals the cystic nature of the masses, and lesions are localized for surgery guidance.

‘Inverse crescent sign’ is a new terminology for the separation of membranes from the posterior aspect of the cyst by air dissection without any anterior extension (5).

Although transient and local parenchymal infiltration after pulmonary hydatid cyst rupture is common, presentation with diffuse pulmonary infiltration is rare. Bacterial super-infection of the cyst is the most serious complication commonly seen after cyst rupture; but in our patient bronchoalveolar lavage for microorganisms was negative which highly suggests a chemical pneumonitis caused by hydatid cyst rupture. In endemic areas like Iran, familiarity with rare and unusual imaging manifestations of pulmonary hydatidosis, such as diffuse alveolar shadows and cavitary consolidations is essential for early diagnosis.

REFERENCES