Cystic echinococcosis: Late rupture and complication of a stable pulmonary cyst

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CASE REPORT

Cystic echinococcosis is observed worldwide. The majority of cases encountered in Canada occur among immigrants from endemic countries (1). Pulmonary involvement occurs in approximately 25% of cases. The liver is the most commonly affected organ, followed by the lungs (2). Although observation alone may be appropriate (3) for certain echinococcal cysts (generally hepatic), traditional management of hydatid cysts includes an invasive surgical approach with adjunctive chemotherapy. More recently, laparoscopic, percutaneous and parenchymal-sparing interventions have been introduced; however, the evidence to inform guidelines regarding the type of surgical and parenchymal-sparing interventions have been introduced; how-ever, the evidence to inform guidelines regarding the type of surgical intervention other than clinical and radiographic follow-up was deemed necessary. Between 1998 and 2002, he continued to be completely asymptomatic and, in fact, did not attend his follow-up appointments from 2003 onward.

The patient returned in 2007 only at the insistence of his family physician. He stated that in the intervening five years, he had remained completely asymptomatic, as he was during his current visit. A repeat CT scan of the thorax in 2007 (Figure 1C) showed no change in the size of the cyst, now measuring 5 cm × 4 cm × 3.5 cm. He was seen once again in 2008 and was asymptomatic, with chest radiography (Figure 1D) showing no change from previous imaging. He desired to discontinue regular follow-up.

He returned at his own request one year later, in August 2009, with a one-month history of shortness of breath, cough, right-sided pleuritic chest pain, fever, chills, night sweats and a 4.5 kg weight loss. He had already been prescribed two courses of antibiotics (cefuroxime and clarithromycin), with no significant improvement.

A chest examination revealed dullness to percussion and absent breath sounds over the right lower lung field. His white blood cell count was 9.13 × 10^9/L with 25% eosinophils. The chest radiograph showed right pleural effusion. A CT scan of the thorax confirmed that the previously well-circumscribed bronchogenic cyst had ruptured (Figures 2A and B). Analysis of the pleural fluid revealed that it was an exudate with a pH of 7.48 and cell count of 120 × 10^3/L (86% neutrophils and 2% eosinophils). Pleural fluid culture was negative.

Despite initial therapy with chest tube drainage and broad spectrum intravenous antimicrobial therapy, the patient continued to complain of cough, chest pain, shortness of breath and generalized fatigue. Pleural aspiration was performed and the patient required thoracotomy, decortication and resection of the ruptured cyst. Can Respir J 2011;18(5):258-260.

Key Words: Albendazole; Praziquantel; Pulmonary echinococcus

CT measurements of the cyst in 2000 showed some enlargement – it now measured at 5 cm × 4 cm × 3 cm. The patient was followed at least annually until 2002. He continued to be completely asymptomatic and, in fact, did not attend his follow-up appointments from 2003 onward.

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effusion was decreased, but was not completely eliminated despite several attempts to reposition and flush the chest tube. Surgical treatment was recommended. Five days after his admission to the respirology service, the patient underwent a right thoracotomy with decortication and resection of the ruptured cyst. Final pathology showed Echinococcus organisms (Figure 3). Medical therapy with albendazole and praziquantel was initiated. The patient tolerated these medications well throughout the three-month course. He improved rapidly and, when seen at follow-up three months and one year after discontinuation of medical therapy, he was asymptomatic with a virtually normal chest x-ray (Figure 4).

**DISCUSSION**

The present case illustrates that even long-standing and asymptomatic echinococcal cysts may rupture, and require combined surgical and medical management.

![Figure 1](image1.png) **Figure 1** A Chest radiograph on initial presentation in 1998. B Computed tomography scan of the chest on initial presentation in 1998. C Chest radiograph after 10 years of observation in 2008. D Computed tomography scan of the chest after nine years of observation in 2007

![Figure 2](image2.png) **Figure 2** Chest radiograph (A) and computed tomography scan (B) of the chest on presentation in 2009
When feasible, surgery (4,5) has traditionally been the treatment of choice for pulmonary hydatid disease. While percutaneous methods such as puncture, aspiration, injection and reaspiration (PAIR) are used with increasing frequency in the management of hepatic echinococcal disease, their use in pulmonary hydatid disease has been more limited. Gabal et al (6), described the use of a modified PAIR technique on five pulmonary hydatid cysts. Complications were limited to one pneumothorax. All cysts decreased in size, but none resolved completely. Recently, centres experienced in the management of cystic echinococcosis have suggested that observation alone of certain hydatid disease is sufficient (3), although this recommendation is probably more applicable to hepatic than pulmonary cysts, which demonstrate low rates of disease progression and complications (7). However, this ‘watch and wait’ approach has not been formally evaluated. Aribas et al (8) found high rates of morbidity (16.3%) and mortality (2.3%) in patients who presented with complications of pulmonary hydatid disease and subsequently suggested early surgical intervention.

The most recent update on pulmonary cystic echinococcosis (5) confirmed the previous recommendation (identifying surgery as the main therapeutic approach) and points out that the optimum timing of surgery in asymptomatic patients is unknown, that most patients are operated on after many years of infection and that nearly 50% present with complications – presumably, rupture and infection.

Patients who cannot undergo surgery for various reasons are treated medically. The most extensively studied drug for medical treatment of echinococcal cysts is albendazole. It is recommended as first-line chemotherapy in the treatment of cystic echinococcal disease (9). There is limited evidence to suggest that, in the setting of cyst rupture, the use of combination therapy with albendazole and praziquantel may reduce the risk of postoperative echinococcal recurrence (10).

Our case demonstrates that medical therapy with albendazole and praziquantel, in conjunction with surgical drainage, is successful in the treatment of echinococcal empyema. Furthermore, although Canada is not a country in which echinococcal disease is common, it clearly needs to be suspected in all immigrants arriving from endemic areas, whose chest radiograph shows bronchogenic cysts suspected of being due to Echinococcus organisms.

Finally, although the natural inclination in a stable asymptomatic patient with benign pulmonary lesion is to observe – rather than operate – this may not be the best approach in the case of echinococcal cysts, which may rupture spontaneously even after many years of clinical stability. Early surgery may be the best way to proceed to prevent eventual rupture of the cyst. Although there is no consensus as to the best surgical approach, current opinion (5) is that excision of the cyst using the parenchymal sparing technique (11) is the preferred approach. The surgery is safe, without any perioperative mortality, and with less than 9% morbidity, the most common being atelectasis, pleural effusion, prolonged air leak, hemoptyis and wound infection.

REFERENCES