Case Report

Bilateral Giant Hydatid Pulmonary Cysts in a Pediatric Patient: A Rare Case Report

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Abstract

Introduction

Pulmonary hydatid cysts, resulting from the Echinococcus tapeworm larva, primarily affect the lungs. The global incidence is elevated in regions with prevalent livestock farming due to the ingestion of Echinococcus eggs. Bilateral involvement is an infrequent presentation, carrying inherent respiratory risks. Accurate and prompt diagnosis necessitates a comprehensive approach involving imaging studies and serological tests to uncover this parasitic affliction.

Case presentation

In a 15-year-old male from an underprivileged rural background, bilateral giant hydatid cysts manifested. The patient's exposure to this parasitic infection was linked to his rural setting and contact with pets. The case unfolded with respiratory symptoms, prompting an investigation that revealed substantial bilateral cystic lesions. Successful surgical intervention and diligent postoperative care resulted in a smooth and uncomplicated recovery for the patient.

Conclusion

While giant bilateral hydatid cyst cases are rare, their documentation provides invaluable insights into the understanding and management of this complex condition. The prognosis hinges on various factors, underlining the importance of a multidisciplinary post-surgical approach for optimal patient outcomes.

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1. Introduction

A hydatid cyst, arising from the larval stage of the Echinococcus tapeworm, represents a parasitic infection manifesting particularly in the pulmonary form when cysts develop within the lungs [1]. The global incidence of this condition exhibits variations, with higher prevalence noted in regions where livestock farming is prevalent [1,2]. The fundamental causative factor lies in the ingestion of Echinococcus eggs, typically
occurring through the consumption of contaminated food, water or direct contact with animals harboring the infection [1].

Several risk factors contribute to the likelihood of contracting pulmonary hydatid cysts, including proximity to dogs, involvement in livestock handling, and residing in areas endemic to the disease [1-3]. While bilateral involvement is a relatively uncommon manifestation, its occurrence introduces heightened risks due to potential complications, particularly the compromise of respiratory function [4,5].

The clinical presentation of pulmonary hydatid cysts is diverse, encompassing symptoms such as cough, chest pain, and various respiratory manifestations [1,2]. Timely and accurate diagnosis is crucial and typically involves a comprehensive approach. Imaging studies, such as chest X-rays or CT scans, play a pivotal role in visualizing cystic structures [6,7]. Serological tests aid in confirming the presence of the Echinococcus infection, complementing the clinical evaluation for a comprehensive diagnostic assessment [8,9].

The significance of reporting a case involving bilateral giant pulmonary hydatid cysts cannot be overstated. Such cases are exceedingly rare and underscore the importance of understanding the potential complications associated with this unique presentation. Emphasizing the urgency of early diagnosis and intervention is imperative, given the heightened risk of respiratory compromise in bilateral cases.

2. Case Presentation

2.1. Patient information

A 15-year-old male patient reported a gradual emergence of dyspnea and cough persisting for a month. His past medical and surgical history was unremarkable. Coming from an underprivileged family in a rural setting, he had been exposed to pets.

2.2. Clinical findings

A physical examination revealed decreased bilateral air entry, although vital signs remained normal.

2.4. Therapeutic intervention

A chest x-ray depicted bilateral giant round homogeneous opacities in both the upper and middle zones of the right lung and the middle and lower zones of the left lung (Figure 1). A subsequent chest CT scan revealed two homogeneous cystic lesions, exhibiting peripheral enhancement and occupying the majority of both hemithoraces. These lesions measured 16x14 cm on the right and 14x12 cm on the left, indicative of bilateral giant hydatid cysts (Figure 2). The patient’s complete blood count highlighted eosinophilia, with an eosinophilic count of 700 cells/microliter. Both abdominal ultrasound and brain CT scan results were normal.

The patient received a preoperative dose of 400 mg intravenous ciprofloxacin, as per the hospital’s protocol.

Figure 1: Plane chest X-ray (posteroanterior view) showing bilateral large homogenous well-defined opacity.

Figure 2: computed tomography scan of the chest (coronal view) showing two giant hydatid cysts on both sides of the chest.

The surgery was performed under general anesthesia. Positioned laterally on the left with a single lumen endotracheal intubation, a 5 cm utility incision was made in the fifth intercostal space, located in the posterior axillary line. Additionally, a 1 cm incision was made in the sixth intercostal space, anterior axillary line, to accommodate the camera. For the right-side cyst, a controlled evacuation was executed post-isolation with packs soaked in hypertonic saline. The germinal membrane including the daughter cysts and pericysts was removed, the fistulae were closed, and the residual cavity was rinsed with hypertonic saline and closed by 2-0 vicryl (Figure 3). A chest drain was inserted. The same procedure was repeated for the left-side cyst within the same session (Figure 4).

The patient’s recovery was smooth and uncomplicated. Postoperative care included a prescription of oral albendazole 400 mg twice daily and pain management with paracetamol 1gm thrice daily, ketorolac 30 mg thrice daily, and pethidine 50 mg
as needed subcutaneously. Intravenous ciprofloxacin 400 mg was administered twice daily for 48 hours post-surgery. The left-side chest drain was removed on the third postoperative day, and the right-side drain on the fourth. The patient was discharged in good health on the fifth postoperative day.

2.4. Follow-up

At a follow-up visit six weeks post-surgery, the patient displayed no significant respiratory issues. He was instructed to continue taking albendazole for six months and to regularly monitor his liver enzymes.

4. Discussion

Pulmonary hydatid cysts pose a significant threat to health, yielding a spectrum of symptoms ranging from respiratory distress, persistent cough, and chest pain to potentially life-threatening complications, including cyst rupture, infection, or anaphylaxis [1,2]. The geographical prevalence of the disease is notably higher in regions where livestock farming is commonplace, such as parts of Asia, Africa, South America, the Middle East, and certain areas in Europe [10,11].

The diagnostic landscape of pulmonary hydatid cysts is marked by challenges attributed to the nonspecific nature of symptoms and the gradual progression of the disease [1,2,9]. Clinical suspicion arises particularly in individuals hailing from endemic regions with a history of animal contact [12]. Instrumental in the diagnostic process is imaging studies, with chest X-rays and CT scans playing a pivotal role in visualizing the characteristic cystic structures [6,7,13,14].

Guidelines for managing pulmonary hydatid cysts primarily advocate for surgical intervention, as endorsed by the World Health Organization (WHO) [15-19]. The overarching objective of surgery is to excise the cyst intact, thereby mitigating the risk of complications [20,21]. Complementary to surgical measures, medical treatment involving anthelmintic drugs like albendazole assumes the role of adjunctive therapy, aiding in reducing the likelihood of recurrence and addressing any residual cystic structures [22,23].

Despite the potential rarity of giant bilateral hydatid cyst cases in the existing literature, the importance of disseminating such cases cannot be overstated. Reports of these unique presentations contribute invaluable insights into evolving management strategies and outcomes, fostering a deeper understanding of this complex condition.

Prognosis and the success of surgical interventions hinge on multifactorial considerations, including the size and location of the cysts, the overall health of the patient, and the surgical approach adopted. When executed with precision, surgery often culminates in a favorable outcome, alleviating symptoms and improving the patient's quality of life [13,21]. However, the presence of complications, such as cyst rupture during surgical
5. Conclusion

The successful management of pulmonary hydatid cysts necessitates a collaborative and multidisciplinary approach, seamlessly integrating surgical expertise with ongoing medical treatment to optimize patient outcomes.

Declarations

Conflicts of interest: The author(s) have no conflicts of interest to disclose.

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