INTRODUCTION

Cardiac hydatid cysts are uncommonly encountered entity of hydatid disease, even in regions where hydatid cysts are endemic. Most patients are adults and asymptomatic. Presentation may be with non-specific symptoms such as dyspnea, chest pain, palpitations, cough, and sometimes with life-threatening events. We present a rare case of a 9-year-old girl who was diagnosed with a cardiac hydatid cyst in the right ventricle, and underwent surgical excision.

Since time immemorial, human Echinococcosis has represented a serious disease burden of global concern. It is a parasitic disease caused by the infection of larval stages of tapeworms belong to genus Echinococcus through the ingestion of food or water contaminated with their eggs, or by a direct contact with the animal hosts. The development of the larvae of Echinococcus Granulosus in humans (as accidental intermediate hosts) leads to Cystic Echinococcosis (CE) (Hydatidosis). It is a hyperendemic disease in different regions of the world, significantly in rural areas. The most frequently involved organs by hydatidosis are liver (more than 65% of cases) and lungs (both comprise at least 90% of cases). Cardiac hydatid disease is rare (0.5–2% of all locations), and this may be due to the filtration of the embryos through the liver and lungs, and to the fact that cardiac rhythmic contractions prevent the cysts to implant within the cardiac wall. Clinical presentation of cardiac hydatid cysts is quite variable and depends on the age, size, location, and integrity of the cyst. In the majority of cases, patients are asymptomatic. Most common reported symptoms are dyspnea, chest pain, palpitations, and cough. On the other hand, it may present with life-threatening events such as anaphylactic shock, pulmonary embolism, acute coronary syndrome, or malignant arrhythmias due to subendocardial rupture which leads to the release of cyst contents into the blood stream. In endemic geographic areas, the most important issue regarding diagnosis is clinical suspicion. Transthoracic echocardiography (TTE) is the main diagnostic tool for cardiac hydatid disease. Computed tomography (CT) and magnetic resonance imaging (MRI) may provide additional diagnostic information. Surgery is the treatment of choice in all cases to achieve higher survival rates with lower recurrence.
2 | CASE PRESENTATION

A nine-year-old girl from northern province in Syria presented to our hospital with recent developing exertional dyspnea. In her past medical history, she had undergone surgical resection of liver hydatid cysts four years ago. TTE showed a 50 × 55 mm heterogeneous cystic mass localized in the free wall of the right ventricle outlet causing moderate obstruction (Max gradient across right ventricular outflow tract was 40 mm Hg) (Figure 1). Routine laboratory tests were normal. Depending on TTE findings, the patient’s demographic origin, and her surgical history, the diagnosis of right ventricle hydatid cyst was established. After a 5-day oral Albendazole course, surgery was performed through median sternotomy, and cardiopulmonary bypass (CPB) was prepared. The cyst was apparent within the wall of the right ventricle outlet (Figure 2). The operating field was isolated by medical gauze saturated with hypertonic saline solution. Without arresting the heart, a small incision on right ventricular outlet was carried out. The contents of the cyst were aspirated, and the cyst cavity was injected with 30% saline solution. Then, the cyst was dissected and enucleated (Figure 3). The residual cavity of the cyst (Figure 4) was closed by direct continuous suturing. The patient convalescence in the intensive care unit was uneventful and was discharged home six days later on albendazole for 12 weeks. The patient was followed up for 2 years. Serial examinations by TTE showed no cardiac recurrence, and the patient was asymptomatic.

3 | DISCUSSION

CE is highly prevalent in southern and northern provinces of Syria; however, epidemiologic studies on human echinococcosis are still not available.11 Our patient was from northeastern province in Syria, which is notably endemic for CE. In rare cases, the larvae reach the heart mainly through the coronary arteries and develop into hydatid cysts. Young adults are the most frequently affected, but very rare cases of cardiac hydatid cysts in children have been reported. Since it receives the most blood supply, the left ventricular wall is most frequently involved (55%–60%), whereas the right ventricle is less.

FIGURE 1  TTE image of the cyst (The Star) in the right ventricle in four chamber view

FIGURE 2  Intraoperative image showing the swelling in the right ventricle outlet (The Star) which refers to the cyst

FIGURE 3  Image of the excised cyst
involved (10%–15%). Our case is a rare presentation of cardiac hydatid cyst within the right ventricle in a child. It has been reported that in cases of right ventricle hydatid cysts, intracavitary rupture is more frequent and may lead to pulmonary embolization, pulmonary hypertension, and death.2,3,9,10 To avoid these complications, it is crucial to make an urgent diagnosis and investigations. High index of suspicion was the key of diagnosis in our case (the geographic origin and surgical history). There are multiple laboratory tests, such as Casoni’s intradermal test, the Weinberg reaction, and peripheral blood eosinophil count; however, they are not reliable since they carry false-negative results.7 Electrocardiogram and chest radiograph are not specific.7 TTE is the preferred diagnostic modality in detecting the cysts and their location and size.3,7 Other highly useful modalities include CT and MRI, which can be informative for a better surgical approach.3,5,7 The surgical excision combined with Albendazole therapy is still the gold standard for treatment of cardiac hydatid cysts.1–10 It is possible to excise the superficially or epicardially located cysts without the use of CPB; however, it is more recommended to use CPB as it shows higher safety and less risk of cyst contents leakage during surgery.7,8 In order to reduce the cyst tension and avoid an accidental rupture, the entire cyst contents must be aspirated. Moreover, the cyst must be injected with protoscolicidal agent to create an osmotic gradient that kills the protoscolices to avoid any possible recurrence in case of their spillage or locally invasion during the operation. For this purpose, packing the operative field with gauze and sponges soaked in a scolicidal agent is performed. We used hypertonic saline solution due to its effectiveness and relatively non-toxicity.8,13 Albendazole (400 mg twice daily) should be prescribed at least 4 days preoperatively and continued for 12 weeks postoperatively.2,3,8,9 The effectiveness of other therapeutic techniques such as puncture, aspiration, injection of a scolicidal agent, reaspiration, and radiofrequency ablation in the treatment of cardiac hydatid cysts is still unclear.8

4 | CONCLUSION

Cardiac hydatid cyst is very uncommon disease, rarely affecting the right ventricle in children. The clinical course varies widely, and the presentation may be insidious with persistent potential risk of life-threatening events. CE must be suspected in endemic areas, diagnosed with appropriate imaging techniques, and treated appropriately.

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CONFLICT OF INTEREST

The authors have no conflict of interest.

AUTHOR CONTRIBUTIONS

Alwaleed Al-Dairy planned and performed the work leading to the report. Wrote and reviewed successive versions and participated in their revisions. Rahim Abo Kasem wrote and reviewed the successive versions and participated in their revisions.

ETHICAL APPROVAL

The manuscript was approved by ethics committee at Damascus University.

CONSENT FOR PUBLICATION

Consent for publication was granted by the patient’s mother.

DATA AVAILABILITY STATEMENT

The data that support the findings of this study are available from the corresponding author, [A.A], upon reasonable request.

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