CASE REPORT

Cardiac hydatid cyst in left ventricular free wall

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Summary

We report a rare case of a cardiac hydatid cyst that was incidentally found during routine work up for a redo-CABG and was picked up on echocardiography and confirmed by magnetic resonance imaging and, after successful removal, further confirmed by histopathology. The report emphasizes the importance of early and urgent surgery for such cardiac hydatid cysts whenever discovered to prevent fatal and unexpected death. Cardiac hydatidosis is a most infrequent type, in comparison with hydatidosis of the liver (65%) and lung (25%).

Learning points:

- Hydatidosis or cystic echinococcosis is caused by infection with the metacestode stage of the tapeworm *Echinococcus* (family Taeniidae). The adult tapeworm is usually found in dogs or other canines; the tapeworm eggs are expelled in the animal’s feces and humans become infected after ingestion of the eggs. The initial phase of primary infection is asymptomatic.
- Cardiac hydatidosis is extremely rare, more commonly the liver and lungs are affected.
- Morbidity from heart echinococcosis in men is three times higher than that in women. Solitary cysts occur in almost 60% of the cases; the most frequent location is the ventricular myocardium and they are usually subepicardially located, hence they rarely rupture in the pericardial space. The left ventricle is damaged twofold to threefold more frequently than the right one.
- The diagnosis of echinococcosis in heart can be divided into two steps: detection of the cyst and its identification as echinococcus. It is based on serological reactions, echocardiography, X-ray, computerized tomography, and/or magnetic resonance imaging.
- The most dangerous complication of cardiac echinococcosis is cyst perforation. After cyst perforation three quarters of the patients die from septic shock or embolic complications.
- It is very important to understand that chemotherapy may lead to cyst death, and destruction of its wall and result in cyst rupture. Therefore, no germicide must be administered before surgical removal.

Background

Cardiac hydatid cyst is a medical emergency. Rapid diagnosis must be performed using various imaging modalities with early surgical and pharmacological treatment of suspicious cystic masses, especially in endemic areas. Increased awareness is essential amongst cardiac physicians and diagnosticians (1).

Case presentation and investigations

A 67-year-old male patient who had undergone CABG in the year 2003 was admitted with complaints of chest heaviness on exertion for the preceding few weeks. The patient underwent a stress test, which was positive for reversible myocardial ischemia. Subsequently, a coronary angiogram was done, which showed a 90–95% stenosis distal to the left...
internal mammary artery-to-left anterior descending artery
graft anastomosis graft anastomosis. During the routine
work up for a redo-CABG, echocardiography showed the
presence of an intramural cystic mass \((2.3 \times 2.1 \text{ cm})\)
attached to the posterolateral wall of the left ventricular
cavity (Fig 1). On Doppler echocardiography no color flow
was observed within the cystic cavity. Cardiac magnetic
resonance imaging (MRI) was performed to further evaluate
the single cyst and rule out other possible differential
diagnosis, such as a simple post-infarction blood-filled
cyst, etc. The MRI confirmed the diagnosis, which implied
that the treatment of choice was urgent surgical cystectomy
with cardiopulmonary bypass. A serological test was
performed for specific echinococcus antibodies along with
routine hematology and biochemistry. Ultrasound, chest
X-ray, and remaining body scans were shown to be negative
for involvement of other organs in the echinococcus
infection. While the serology revealed equivocal results,
the cardiac MRI showed a \(2.2 \times 2.1 \text{ cm}\) well-encapsulated
lesion medial to the anterior papillary muscle, occupying
the left ventricular intracavitary space and inseparable from
the underlying endocardium. The lesion appeared hypo-to
iso-intense on T1WI images and had no post-contrast
enhancement. The patient was posted for early CABG and
left ventricular cyst removal (Figs 2 and 3).

**Treatment and follow-up**

After the redo-CABG, ventriculotomy was performed with a
cardiopulmonary bypass pump, and the intracavitary cystic
mass, identified to be similar in size on both echocardiogra-
phy and MRI, attached to the lateral wall of the left
ventricle was successfully removed. The patient had an
unremarkable post-operative course and was discharged after
a week staying on treatment with Albendazole 400 mg twice

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**Figure 1**
Echo images showing a left ventricular cavity cyst attached to the postero-
lateral wall in parasternal long- and short-axis views (arrow) respectively.

**Figure 2**
2D and 3D echo images showing left ventricular cavity cyst (arrow)
inseparable from the lateral wall in an apical four-chamber view.

**Figure 3**
Images showing no color flow in the cyst cavity on 2D echocardiography
(arrow) and cardiac MRI showing a left ventricular cavity cyst embedded in
the lateral wall (arrow) respectively.

**Figure 4**
Comparative images of 3D full-volume and cardiac MRI showing the left
ventricular cyst (arrow) embedded in the lateral wall.
daily per os for a period of 6 months, administered in cycles for of 3 weeks followed by 1 week of respite (in order to avoid toxicity), along with other drugs for coronary artery disease management. The patient was advised for timely follow-up to look for any reoccurrences in future (Figs 4 and 5).

Discussion

The incidence of left ventricular invasion by echinococcus is 55–60% as it has the maximum myocardial mass and abundant blood supply the incidence of involvement of the interventricular septum is 5–9% of cases. The right ventricle is involved in 15% of cases, while the right atrium is involved in 3–4% of cases (2). Distribution in pulmonary artery, left atrium, and pericardium is up to 7–8% (3). There are no age limits to the presentation and such cysts can cause obstruction in outflow tract, valves, and chambers of the heart, and can induce conduction disturbances such as atrioventricular nodal blocks, ventricular tachycardia and fibrillation, or cardiac tamponade, or can be completely asymptomatic (4). Pulmonary embolism, anaphylactic shock, and systemic metastasis are some more important and catastrophic complications of cardiac hydatid cysts. Left ventricular hydatid cyst are usually located subpericardially and rarely rupture into the pericardial space. The risks of surgery involve leakage of fluid from the cyst cavity leading to anaphylaxis and dissemination of infected scolices, which can be minimized by using scolicidal solutions such as iodine, hypertonic saline, methylene blue, or ethanol (5).

Declaration of interest

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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Patient consent

This is to confirm that written informed consent has been obtained from the patient for publication of the submitted article and accompanying images.

Author contribution statement

S Ohri and A Sachdeva were involved in obtaining the echocardiography images and in the clinical management of the patient under the supervision of S Shrivastava and M Bhatia – Head of Department of Cardiac MRI and CT – who was responsible for interpretation of the MRI used for establishing the diagnosis.

References

4. Di Bello R & Menendez H. 1963 Intracardiac rupture of hydatid cysts of the heart. A study based on three personal observations and 101 cases in the world literature. Circulation 27 366–374. (doi:10.1161/01.CIR.27.3.366)

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