




Incidental finding of cardiac hydatid cyst during autopsy

Nishant Purbey¹ , Amit Patil¹ , Shreekant Bharti² , Keshav Chandra¹ ,
Shashank Ranjan¹ 

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ABSTRACT

Hydatidosis or echinococcosis is an endemic parasitic disease caused by the ingestion of eggs of echinococcal species worldwide. In India, the annual incidence varies from 1 to 200 per one 100,000 hab., with the highest prevalence reported in the Indian states of Andhra Pradesh and Tamil Nadu. The dog is the definitive host, while humans, sheep, and cattle are intermediate hosts. The disease usually involves the liver and lungs, with the kidney and other organs rare involvement. Cardiac hydatidosis is still further rare, seen in 0.2% to 2% of the patients who remain asymptomatic until the development of its complications. Sudden deaths in cardiac echinococcosis are mostly attributed to cardiac arrhythmias, coronary artery diseases, valvular diseases, cardiomyopathies, pericarditis, and cardiac tamponade. We, herein, report a rare case of cardiac hydatid cyst incidentally found during the autopsy of a 26-year-old male who died due to electrical injuries. A single greyish-white cystic mass measuring 1.5cm X 1.2cm was detected on the left anterior ventricular wall 4 cm above the apex and was confirmed microscopically as a hydatid cyst. The cause of death was attributed to external injury.

Keywords

Echinococcus; Electric Injuries; Parasitic diseases; Autopsy; Forensic Pathology

INTRODUCTION

Hydatid disease, also known as echinococcosis, is a parasitic infection caused by the larval stage of *Echinococcus granulosus* tapeworms. The disease is widespread worldwide but endemic in the Middle East, South America and Mediterranean countries.¹ It is endemic in India, with annual incidence varying from 1 to 200 per 100,000 population,² with the highest prevalence reported from Andhra Pradesh and Tamil Nadu due to of poor hygiene and cattle and sheep rearing population. It is transmitted by ingesting food, water, or soil contaminated with stool of infected cattle.³⁻⁵ The alimentary tract is the most frequent route via which *Echinococcus granulosus* is

transmitted. Airborne bronchial venule penetration to the heart and systemic circulation is yet to be proven. It is also possible that parasites could bypass the portal filter and enter organs and tissues other than the liver and lungs through a lymphatic or venous shunt. Considering the muscle produces lactic acid, it is possible that *Echinococcus granulosus* eggs could hatch in soft tissues as well as the gastrointestinal tract in cases of involvement of the skeletal or cardiac muscles.⁶ For *Echinococcosis granulosus*, the dog is the definitive host; sheep, cattle, goats, and pigs are the usual intermediate hosts.⁷ Hydatid disease in humans most commonly occurs in the liver (55-70%)

¹ All India Institute of Medical Sciences, Department of Forensic Medicine and Toxicology, Patna, Bihar, India

² All India Institute of Medical Sciences, Department of Pathology, Patna, Bihar, India



and lungs (18-35%), with rare involvement of kidneys and other organs.^{8,9} Cardiac involvement in echinococcosis is sporadic, usually asymptomatic until the development of complications.⁹ We report a rare case of cardiac hydatidosis that was incidentally found during a routine autopsy of a man who died of external injury.

CASE REPORT

A 26-year-old male was found collapsed on an air-filling machine while working after an electrical shock. The autopsy was conducted the next day. On external examination, the corpse was of a young adult male, average built, 161 cm in length. At autopsy, there was multi-visceral congestion without any internal hemorrhage. Internal organs were unremarkable except for the heart (weight 380 g, mean reference range 327 g), with a soft greyish-white nodule on the left ventricle anterolateral aspect measuring 1.5cm X 1.2cm and was 4 cm above the apex (Figure 1). Externally, the cystic swelling was fixed and extended into the left myocardial ventricle wall. The remaining heart examination, on sectioning, was unremarkable.

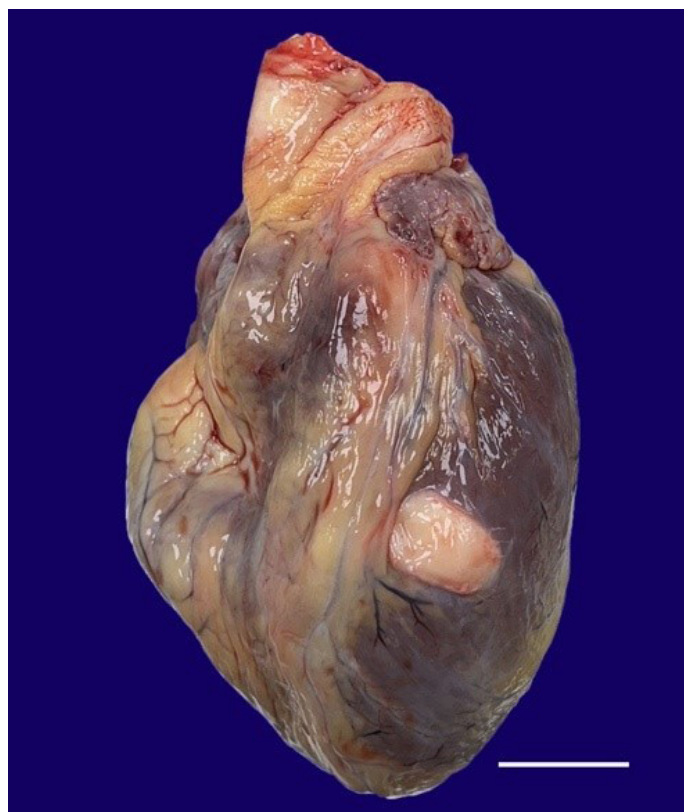


Figure 1. Gross view of heart with the cyst in the anterior ventricular wall (scale bar = 5cm).

Microscopic examination using H&E staining of the paraffin sections showed a cyst overlying the myocardial surface (Figure 2A, 2B). The presence of a tortuous linear lamellate structure showing hyalinized amorphous eosinophilic wall and tiny brood capsules on one side of its surface was depicted. The cyst was formed and walled off by a thick fibrous capsule, thus confirming the diagnosis of a cardiac hydatid cyst. Histological findings of the ventricles and coronary arteries were unremarkable.

DISCUSSION

Cardiac involvement by *Echinococcus granulosus* is seen in 0.2% to 2% of the patients who remain asymptomatic for a longer duration and can display nonspecific symptoms such as chest pain, cough, and palpitations.¹⁰⁻¹³ Existing 37 literature on cardiac hydatidosis suggests sample size varies from 2 to 62.¹⁴ The cysts usually grow slowly (1–5 cm per year) without causing symptoms, with probably only 10% of patients, especially those with large hydatid cysts, having clinical manifestations. The signs and symptoms of cardiac hydatid cysts are highly variable and directly related to the location and size of the cysts.^{15,16}

Diagnosing cardiac hydatidosis in the early phase is difficult due to the long latency from exposure to the infection and the manifestation of the disease. Mostly, they are asymptomatic but can present clinically with chest pain, dyspnea, palpitations, arrhythmias, and AV nodal block. T wave inversion and premature ventricular beats are electrocardiographic findings.¹⁷⁻¹⁹

Chest radiograph findings are nonspecific except for the detection of cardiomegaly, which depends upon the size and site of the lesion.²⁰ Echocardiography is the diagnostic method of choice for cardiac hydatidosis. It is an efficient, easy-to-perform, informative, and sensitive noninvasive technique to localize and detect cysts in living individuals. The most commonly used serology is the echinococcus indirect hemagglutination (EIHA) test and enzyme-linked immunosorbent assay (ELISA). However, a confirmatory diagnostic test for cardiac hydatidosis is the CT scan.²¹ Patients were managed surgically and pharmacologically, and their outcomes were good, so mortality was negligible. In the present case, the cyst was located on the anterior surface of the left ventricular free

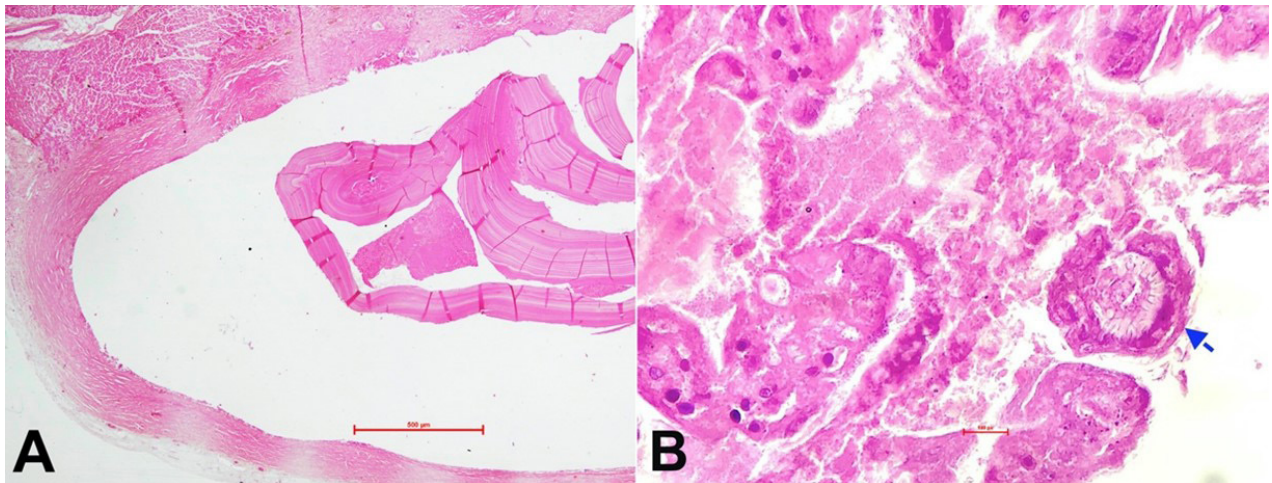


Figure 2. A – Superficial myocardium showing a cystic cavity with parasitic structure within the cyst (H&E, 10x); **B** – Hydatid cyst showing predominantly acellular wall and a protoscolex with refractile hooklets inside (Blue arrow) (H&E, 40x).

wall, one of the most common cardiac sites affected by the hydatid cyst owing to its thickness and rich blood supply, followed by the interventricular septum.^{22,23} The discovery of a hydatid cyst at autopsy requires confirmation by histopathology, an essential criterion standard for identifying the hydatid nature of the cyst. Microscopically, the wall of the hydatid cyst has 3 structural components: an outer acellular laminated membrane, the germinal membrane, and the protoscolices (Figure 2A, 2B). The hydatid cyst may be surrounded by a fibrous capsule or granulation tissue, including inflammatory infiltrate.²⁴ In our case, histological examination confirmed the hydatid nature of the cyst in the heart.

Sudden deaths caused by hydatid diseases are rarely reported in the literature. In these, death is either attributed to the cystic rupture, embolization of hydatid material to the cerebral circulation or to the pulmonary artery, and anaphylaxis.²⁵⁻²⁶ However, in cases of cardiac hydatidosis, sudden deaths are mostly attributed to major causes such as cardiac arrhythmias, coronary artery diseases, valvular diseases, and cardiomyopathies.^{27,28}

In the present case, though a cardiac hydatid cyst was confirmed as an incidental finding, the cause of death was attributed to an external cause.

CONCLUSION

Cardiac echinococcosis is a rare disease that may remain asymptomatic until the development of

complications and may be lethal in the absence of early diagnosis. In the present case, a rare incidental finding of a cardiac hydatid cyst was observed at autopsy. This was confirmed histologically. Cardiac hydatidosis can cause sudden death due to cardiac arrhythmias; however, that was not the cause of death in our case.

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Correspondence

Nishant Purbey

All India Institute of Medical Sciences, Department of Forensic Medicine and Toxicology
2ND FLOOR, Admin Building, AIIMS, Phulwari sharif, 801507, Patna, Bihar, India

Phone +917783814686

nishantpurbey@gmail.com