Cardiac Cysticercosis: A Rare Incidental Finding on Autopsy
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Abstract
Cardiac cysticercosis is an extremely rare condition, mostly found in postmortem cases as an incidental finding because it causes either no symptoms or conduction abnormalities in some. We report the case of a 40 years male who died of unknown cause. A cystic cavity was seen in heart which on histopathology found to be cysticercosis.

Keywords: Cysticercosis; Autopsy

Introduction
Cysticercosis refers to a parasitic infestation caused by encysted larval stage of the pork tapeworm, *Taenia solium*. In its life cycle, humans act as definitive host and they harbour the adult forms of the tape worm in their small intestine [1]. Cysticerci can be found anywhere in the body, their location and size determine the clinical presentation. Involvement of heart is very uncommon and mostly asymptomatic [2]. Infection occurs when the tapeworm larvae enter the body. Ingested eggs pass into the bloodstream and disseminate to various organs and form the cysts that characterize cysticercosis [3].

Case Report
Viscera samples of an adult male aged 40 years in a postmortem case were received. The cause of death was ‘unknown’ as per postmortem record. The various viscera received for histopathological analysis were brain, heart, lung, liver and kidneys. On gross examination, heart was enlarged, weighed 450 gm and measured 12x8x5 cm. Using the inflow outflow technique the heart

Figure 1: Cystic cavity on the anterior aspect of heart
Cysticercosis is an infection caused by the pork tapeworm, *Taenia solium*. It is most prevalent in Africa, South-East Asia and South America [3]. Infection occurs via ingestion of contaminated fruits, vegetables and uncooked pork [4]. Larval form disseminate from the intestine via the hepatoporal system to various organs of the body like subcutaneous tissues, skeletal muscles, lungs, brain, eyes, liver and occasionally the heart. Symptoms are variable according to the affected organs like convulsions, decreased visual acuity, arrhythmia, paraparesis, gait abnormality, muscular pseudohypertrophy etc. It is difficult to suspect and diagnose cardiac involvement clinically because of absence of any significant signs and symptoms in most of the cases. Therefore all the patients with disseminated cysticercosis should be screened with echocardiography for any myocardial involvement and carefully monitored during the course of therapy. With the advent and availability of improved imaging modalities to include echocardiography, CT, and MRI, more cases of cardiac cysticercosis are being diagnosed antemortem.

Discussion

Cardiac cysticercosis is predominantly a silent disease, yet cases in which heart function was adversely affected had been found. Pathological diagnosis of cardiac cysticercosis requires obtaining biopsies from heart which is technically difficult and ethically not justified. Demonstration of cysticerci in the autopsy specimen has been the rule to diagnose involvement of heart in disseminated cysticercosis [2]. Most commonly, it affects the central nervous system. Involvement of heart is extremely rare and is usually a postmortem incidental finding, as evidenced by multiple case reports and autopsy series [5,6]. Khelmiski reported the first case in 1962 of cardiac involvement which was detected on autopsy. There was associated skeletal muscle and brain involvement [7]. In India, Reddy and colleagues reported the first case of cysticercosis involving heart [8]. Cysticerci are randomly distributed throughout cardiac tissue, producing a variable inflammatory response ranging from minimal myocardial damage to granuloma formation and potentially fibrosis. Thus, patient is asymptomatic or has conduction abnormalities and arrhythmias [5,6,9]. Cysts may be single or multiple, ovoid or round, and range in size from 1.0 to 30.0 mm. They can be randomly distributed in the subpericardium, epicardium, subendocardium, or myocardium of the atria, ventricles, septum, or papillary muscles. Microscopically, a capsule may be visualized with the scolex, hooklets, and suckers underneath. The presence of calcification is variable [5,9].

Parasitic infections previously seen only in developing tropical settings can be currently diagnosed worldwide due to travel and population migration. Treatment of choice is prevention of cysticercosis with a better primary health care system, health education, proper sanitation, better food hygiene, access to safe water and clean water, and eradication of poverty.

References