CASE SUMMARY
A 25-year-old woman presented to the cardiology department with a history of shortness of breath and palpitations. A general cardiac examination showed regular pulse with a heart rate of 74 bpm and normal blood pressure of 120/80 mm/Hg. Heart sounds were normal. Blood work showed a raised eosinophil count. Chest X-ray showed mild cardiomegaly. The ECG was normal.

IMAGING FINDINGS
Transthoracic echocardiography revealed a cystic mass within the heart. Cardiac MR imaging (MRI) was performed with ECG gating to further characterize this mass. Bright blood cine SSFP gradient 4 chamber and short axis views revealed a large, well-marginated, round mass in the lateral wall myocardium of the left ventricle (Figure 1A) which demonstrated markedly hyperintense signal brighter than blood (Figure 1B). The mass had thin regular walls and was seen bulging into the LV cavity in the vicinity of the mitral valve. On double inversion recovery black blood T2-weighted MRI, homogeneous hyperintense fluid signal intensity in the mass indicated it to be a cyst and a surrounding thin hypointense rim was appreciated (Figure 2). There was no enhancement of the mass while the normal myocardium was seen enhancing on dynamic post-gadolinium inversion recovery MRI (Figure 3).

DIAGNOSIS
 Intramyocardial hydatid cyst involving the left ventricle

DISCUSSION
Cardiac echinococcosis is rare and constitutes less than 2% of all cases of hydatid disease.1 Parasite Echinococcus granulosus is the culprit pathogen that causes hydatidosis in the intermediary hosts, human beings. Human infestation by the larval stage of the parasite can lead to hydatid cyst formation in any organ, pleural cavity, or peritoneal cavity, as the larvae gain access to the portal circulation after ingestion of contaminated food. Hydatidosis commonly develops in the liver and lungs in humans. Imaging is the backbone for detection of echinococcal cysts anywhere in the human body, and especially within the heart.

Cardiac hydatid cysts are most unusual and tend to remain asymptomatic until the cyst becomes large. The incidence ranges between .5 and 2%, even in endemic areas.2 The larvae usually gain access to the heart through the coronary circulation, though direct spread by rupture from primary lung or hepatic hydatid cyst is known to occur.

The favored site for growth of hydatid cyst in the heart is the left ventricle free wall. With its substantial myocardial mass and varying pressures, the LV provides optimal conditions for development and growth of the parasite.3 Other sites in the heart where hydatid cyst may be found are

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the right ventricular wall (15%) left atrial wall (8%), interventricular septum (5-9%), pericardium (8%) and pulmonary atrium (7%).

Complications of cardiac hydatidosis differ with location of the cyst within the heart. Hydatidosis involving the IVS can cause conduction disturbances and A-V block. Hydatid cysts in the left ventricle myocardium tend to be located in the sub-pericardial region and therefore, are likely to rupture into the pericardium leading to inflammatory pericarditis. Cardiac tamponade can develop as a further complication. In the right ventricle, echinococcal cysts are usually sub-endocardial and prone to intracavitary rupture which can result in anaphylactic shock and sudden death; or the cyst debris can cause life-threatening pulmonary or aortic embolism. Intra-cavitary rupture results in carotid vessel occlusion by parasitic emboli and hemispheric stroke may occur.

Ninety percent of patients remain asymptomatic as hydatid cysts are slow growing, and a period of 5 to 10 years may elapse before the lesion attains a size large enough to cause cardiovascular symptoms. Ten percent of patients with larger cysts present with dyspnea, arrhythmias or precordial chest pain that may be confused with angina.

Imaging forms the crux of diagnosis. On TTE the characteristic appearance is of a spheroidal cystic mass with well-defined edges and presence of internal trabeculae corresponding to daughter membranes. On MDCT cysts are identified by their fluid attenuation and internal septations. Cyst wall calcification and daughter cysts can be seen.

Cardiac MRI is superior in diagnosis and detection of typical morphologic features of hydatidosis due to its excellent soft-tissue resolution. On MRI the cyst appears as a unilocular or multilocular, smoothly marginated, round mass that has hypointense signal on T1-weighted images and hyperintense signal on black blood DIR T2W images. The cysts are mostly single with an average size of 3-5 cm. The cyst wall may be thin or thick. When daughter cysts are present a multilocular mass is seen. A pericyst is typically seen in cardiac hydatidosis which appears as a hypointense peripheral ring around the lesion on T2-weighted black blood images. This hypointensity results from a dense fibrous capsule formed by the reactive host tissue. At times, the cyst may undergo degeneration and appear solid and this solid-appearing intracardiac mass proves difficult to differentiate from cardiac myxomas. Membranes may detach from the cyst wall and on SSFP MR Sequences, are visualized floating within the cyst. Dynamic gadolinium first-pass MRI serves to make this differentiation possible as cardiac hydatid cysts do not enhance, differentiating them from homogeneously enhancing myxomas.

Similarly the complications of cardiac hydatidosis are well-depicted on MRI. Cyst seen protruding into the cardiac cavity on MR is a sign of impending intracavitary rupture. Intracavitary rupture of cyst can be diagnosed on bright blood SSFP gradient images as hypointense filling defects within the atrium or ventricle. Rupture of cardiac hydatid into the pericardium produces pericarditis that manifests as thickening and enhancement of pericardium on gadolinium enhanced MR.

A very large Echinococcal cyst located in the ventricle cavity may prevent co-aptation of valve leaflets causing...
valvular dysfunction. Phase-contrast MRI is useful as it quantifies regurgitation across the valve. Mechanical outflow obstruction by the cyst can also be well demonstrated by MR morphologically and functionally.9

The treatment of choice for cardiac echinococcosis is cardiopulmonary bypass surgery and extirpation of the cyst. Prior to cystectomy, the cyst is punctured carefully (taking precautions to avoid spillage of contents), aspirated, and then resected. The residual fibrous cavity is plicated with sutures and closed resulting in cyst removal and cure.10

CONCLUSION
Cardiac MRI is an excellent modality for detecting and characterizing intracardiac hydatidosis. The modality confirms TTE findings, demonstrates the cyst, identifies associated complications and assists in planning surgery. Early diagnosis leads to quick referral for surgery, enabling prompt surgical cystectomy.

REFERENCES

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