Images in...

Apical hypertrophic cardiomyopathy in an Indian patient: straight from the book
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DESCRIPTION
A 55-year-old non-hypertensive Indian man presented with a history of exertional angina. His electrocardiogram showed left ventricular hypertrophy (with biggest R-wave of 47 mm in V4) and giant T-wave inversion (more than 10 mm deep) with deepest T-wave inversion of 12 mm in V4 (figure 1). Echocardiogram (ECHO) showed left ventricular hypertrophy with a septal thickness of 11.7 mm and a posterior wall thickness of 12.2 mm. There was striking apical hypertrophy with a thickness of 28.4 mm (Videos 1–3). The left ventricular cavity resembled an ‘ace of spades’ in diastole (figure 2). There was no evidence of left ventricular outflow tract obstruction or coarctation of the aorta.

The case has classical presentation, electrocardiographic and echocardiographic features of apical hypertrophic cardiomyopathy (HCM). An absolute apical thickness of more than 15 mm and a ratio of apical thickness to basal LV wall thickness of more than 1.3 define apical HCM on ECHO. Standard ECHO may miss this entity where a contrast ECHO may be useful but cardiac MR imaging (CMR) is best. 

CMR is superior to ECHO in detecting variants of HCM such as apical HCM, presence of focal hypertrophy, severe hypertrophy (>30 mm) and LV apical aneurysm. Patchy mid-wall-type delayed hyperenhancement of myocardium, on contrast-enhanced CMR, identifies fibrosis. It is associated with a higher risk of adverse LV remodelling and systolic dysfunction. Its association with ventricular tachycardia has been reported but its predictive value for sudden cardiac death is not clear.

The annual cardiovascular mortality is ~0.1%. New York Heart Association class II or more, age less than 41 years and presence of left atrial enlargement predict an increased incidence of myocardial infarction and atrial fibrillation.
Learning points

▸ The case stresses that secondary causes of angina should be carefully searched for in the initial evaluation of angina before performing the stress test.
▸ Careful electrocardiographic and echocardiographic examination may help in identifying rare aetiologies of angina like apical hypertrophic cardiomyopathy.
▸ Cardiac MR imaging provides useful diagnostic and prognostic information in patients with hypertrophic cardiomyopathy.

Figure 2  Apical four-chamber view of a transthoracic echocardiogram showing apical hypertrophy and an ‘ace of spades’-shaped cavity (black arrow in white box) in diastole.

Video 2  Parasternal short axis view of a transthoracic echocardiogram showing apical hypertrophy and complete obliteration of a cavity in systole.

Video 1  Apical four-chamber view of a transthoracic echocardiogram showing apical hypertrophy and an ‘ace of spades’-shaped cavity in diastole.

Video 3  Apical four-chamber view of a transthoracic echocardiogram with colour Doppler showing apical hypertrophy and an ‘ace of spades’-shaped cavity in diastole.

Competing interests  None.
Patient consent  Obtained.

REFERENCES
Singh S. Apical hypertrophic cardiomyopathy in an Indian patient: straight from the book. BMJ Case Reports 2012;10.1136/bcr-2012-006288, Published XXX

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