Isolated Left Ventricular Noncompaction in a 25 Year Old Male

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Abstract: Ventricular noncompaction (VNC) of the myocardium is a rare genetic cardiomyopathy caused by a disorder during endocardial morphogenesis and could be accompanied by life-threatening complications. The major clinical manifestations of VNC are heart failure, arrhythmias, and embolic events. The left ventricle is the most commonly reported affected site, but a few cases of right ventricular involvement have also been reported. We report the case of a young male who presented with features of heart failure. Trans thoracic echocardiography showed the characteristic trabeculations of left ventricular noncompaction with associated depressed left ventricular dysfunction.

Keywords: ventricular noncompaction, spongiform cardiomyopathy

1. Introduction

Ventricular noncompaction (VNC) of the myocardium is a rare genetic cardiomyopathy that is believed to arise from arrested endomyocardial development during embryogenesis [1]. It is characterised by the presence of deep intertrabecular recesses in the hypertrophied, frequently hypokinetic segment of ventricular myocardium[2,3]. Myocardial noncompaction can occur in isolation or in association with cardiac syndromes or disease[4]. Myocardial noncompaction can occur at any age; the clinical presentation is nonspecific and varies from no symptoms to conduction defects. thromboembolism, ventricular arrhythmias, severe heart failure, or sudden cardiac death[5,6]. Here we report the case of a young male who presented with features of heart failure

2. Case Report

A 25 year old male presented at our institutions emergency department with progressive shortness of breath associated with episodes of palpitations which were present on exertion but was relieved on resting. He did not have other cardiac symptoms like anginal pain, syncopal attacks, orthopnoea or paroxysmal nocturnal dyspnoea. He was a non-smoker, non-alcoholic, normotensive and non-diabetic. There was no history of sudden cardiac death in any of the family members.

Physical examination revealed only a 2/6 systolic ejection murmur which was loudest at the left midsternal border on cardiac auscultation. The laboratory results were within normal limits. Electrocardiogram (ECG) showed sinus rhythm with regular rate.

Trans thoracic echocardiography revealed a trabeculated, sponge-like appearance of the left ventricle, more affecting the inferior posterior wall. There were associated moderate tricuspid valve regurgitation, mild pericardial effusion with moderate left ventricular systolic dysfunction. The left ventricular ejection fraction was 32% only. Cardiac magnetic resonance imaging showed the same trabeculation pattern in the left ventricle. These findings let to the diagnosis of left ventricular noncompaction.

3. Discussion

VNC of the myocardium is a rare genetic cardiomyopathy. The left ventricle is more frequently involved with biventricular involvement rarely encountered. [7,8]. Oechslin et al.[7] demonstrated in a follow-up study of 34 patients that the most important clinical manifestations of VNC were heart failure (53%), ventricular tachycardia (41%), sudden cardiac death (35%), syncope (18%), and embolic events (24%).

Among all the presenting symptom of VNC, the most common reason for referral was heart failure, [7] with one-third of these patients having NYHA class III/IV at time of diagnosis [9]. Both systolic and diastolic dysfunction can develop. Systolic dysfunction and arrhythmias might be secondary to microcirculatory dysfunction and subepicardial hypoperfusion[10]

Echocardiography is crucial for the diagnosis of LVNC. Widely accepted diagnostic criteria [11] include the following:

- 1) An excessively thick myocardial wall structure in 2 differing layers—a thin, compacted epicardial layer and a thicker, noncompacted endocardial layer
- 2) A characteristic end-systolic ratio of >2:1 for noncompacted-to-compacted wall thickness
- Prominent multiple, chiefly intracavitary trabeculae, with color-Doppler echocardiographic evidence of communication between the deep intertrabecular recesses and the ventricular cavity
- 4) No cardiac abnormalities beyond the above, in isolated noncompaction cases.

Our patient presented with features of heart failure along associated palpitations with trans with thoracic echocardiography showing features consistent with left ventricular noncompaction. In patients with LVNC, treatment options vary on an individual basis, ranging from medical management in mild cases to heart transplantation in patients with refractory symptoms. Monitoring asymptomatic patients is encouraged, whether diagnosis is incidental or the consequence of familial screening, because of possible future complications in those individuals[5,6]. Our patient responded well to medical therapy.

4. Conclusion

VNC is an uncommon disorder accompanied by lifethreatening complications. Its unusual presentation highlights the importance of a complete and thorough workup of patients presenting with apparently noncardiac symptom(s). Data is limited on specific therapy for VNC, but it is recommended that medical management be tailored towards the clinical manifestations, and standard guidelines should be applied for patients with reduced LVEF, and heart failure with preserved systolic function [12], in the setting of VNC. Patients with VNC, who meet standard criteria for anticoagulation, should also be managed according to standard guidelines [13].

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