Case Report

Asymptomatic Presentation in Arrhythmogenic Right Ventricular Cardiomyopathy: a Case Report.

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Abstract

Arrhythmogenic right ventricular cardiomyopathy (ARVC) is an inherited cardiac condition leading to one of the common cause for sudden cardiac death in young individuals. It is characterized by fibrofatty replacement of the cardiac myocytes predominantly the right ventricle. As the presentation can be varied and very nonspecific, it often leads to delay in diagnosis and specific treatment. Here, we present a 30 year male who was asymptomatic and detected to have electrocardiograph (ECG) and echocardiographic abnormalities suggestive of ARVC. Further, a cardiac imaging with cardiac MRI confirmed the diagnosis of ARVC highlighting the importance of multimodality imaging.

Keywords: Arrhythmogenic Right ventricular Cardiomyopathy; Asymptomatic Presentation.

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Case

A 30 year old male teacher by occupation presented for a routine health examination at our institute. He had no symptoms of breathlessness, chest pain, dizziness or palpitation. His clinical examination was unremarkable with normal vital parameters. There was no significant history of cardiac disease or any sudden cardiac death in the family. He had been active and no any comorbidities till date. His routine electrocardiograph (ECG) showed repolarization abnormalities in form of T wave in version in leads V1-V4 and lead III (major). (Image 1)

His echocardiogram revealed dilated right atrium (RA) and right ventricle (RV) with dyskinetic RV apex along with dilated right ventricle outflow tract (RVOT):(parasternal long axis (PLAX) dimension-51mm, parasternal short axis dimension (PSAX)-53mm, moderate tricuspid regurgitation with mild pulmonary hypertension Tricuspid annular plane systolic excursion (TAPSE)-19mm, RV systolic tissue doppler velocity (RV s TDI) 9.9mm and normal Left Ventricle (LV) dimension and systolic function (major). He further underwent a cardiac MRI (plain and gadolinium enhanced) which showed a markedly dilated right ventricle and markedly hypokinetic free walls of the right ventricle with associated thinning of the RV wall and patchy fatty areas noted in the right ventricular wall. Late gadolinium enhancement scan showed patchy enhancement of the right ventricle free wall. Right ventricle apex and mid free wall showed hypertrabeculated appearance. (Major) These findings with compared to the Task Force criteria (2010) for Arrhythmogenic right ventricular cardiomyopathy (ARVC) fulfilled for the definite diagnosis of ARVC with 2 major (both Echo and cardiac MRI/ECG). The index patient was totally asymptomatic and hence was offered no active treatment but with close follow up with monitoring of any new symptoms. However, all first degree relatives of the index case have been advised for a screening echocardiographic examination.

Image 1: ECG showing T wave inversion in V2-V4.

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Discussion

In 1978 Frank and Fontain first described ARVD (arrhythmogenic right ventricular dysplasia) in six patients presenting with ventricular tachycardia (VT). It was defined as “total or partial replacement of right ventricular muscle with fibro-fatty tissues associated with left bundle branch (LBBB) morphology arrhythmias”. True incidence of this disease is not known but the prevalence is approximately 0.02–0.1% in the general population.

The diagnosis of ARVC is made in 80% of cases in patients below the age of 40 years. Men are more commonly affected. Hence, it should be suspected in young patients who have history of resuscitated cardiac arrest or aborted sudden cardiac death, syncope and ventricular arrhythmias in the past. The clinical presentation can be varied from asymptomatic to life threatening arrhythmias. This case highlights the role of basic 2D echocardiography to identify suspected ARVC case along with multimodality imaging with cardiac MRI to make a definite diagnosis of ARVC.

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Conclusion

ARVC is an important cause of sudden cardiac death in young individuals. It is an in herited condition characterized by fibrofatty replacement of the right ventricle. Diagnosis should always be suspected in young patients who have history of resuscitated cardiac arrest or aborted sudden cardiac death, syncope and ventricular arrhythmias in the past. The clinical presentation can be varied from asymptomatic to life threatening arrhythmias. This case highlights the role of basic 2D echocardiography to identify suspected ARVC case along with multimodality imaging with cardiac MRI to make a definite diagnosis of ARVC.