Images in...

The ECG in arrhythmogenic right ventricular cardiomyopathy: ε-waves and anterior T-wave inversion

Constantinos O'Mahony, 1 Filippo Maria Cauti, 2 Antonios Pantazis 1

Correspondence to Dr Filippo Maria Cauti, filippocauti@hotmail.it

DESCRIPTION

Arrhythmogenic right ventricular cardiomyopathy (ARVC) is an uncommon inherited cardiomyopathy caused by mutations of desmosomal protein genes responsible for cardiomyocyte electrical integrity and intercellular adhesion. ARVC is clinically characterised by:

- (1) electrical instability manifested by abnormalities of the resting ECG and ventricular arrhythmias. Sudden arrhythmic cardiac death is a recognised complication.
- (2) right ventricular systolic impairment, dilatation and/or regional wall motion abnormalities. Left ventricular involvement is increasingly been recognised. The electrical manifestations tend to precede the structural abnormalities.

The ECG illustrates the typical findings in ARVC: (1) anterior T-wave inversion in the right precordial leads (V1–V3), present in 85% and (2) ϵ -waves, present in 33% of ARVC patients. ϵ -Waves are reproducible low-amplitude signals occurring after the end of QRS complex and before the T-wave in the right precordial leads.

This patient had a maternal first-degree relative who was diagnosed with the condition having presented with aborted sudden cardiac death. Both the proband and the patient were found to have a desmosomal protein gene mutation (plakophilin). The patient also had right ventricular dilation and regional wall motion abnormalities

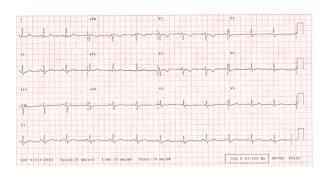


Figure 1 A standard 12-lead ECG showing T-wave inversion in leads V1 and V2.

with aneurysm formation and >1000 premature ventricular complexes in 24 h consistent with the diagnosis of ARVC.

ARVC is inherited in an autosomal dominant fashion, and first-degree relatives should be offered screening. The 12-lead ECG is an important diagnostic tool (figure 1). Double speed and amplitude traces are invaluable in the detection of ε -waves (figure 2).

This case illustrates the sensitivity of the ECG in detecting early electrical abnormalities in this condition.

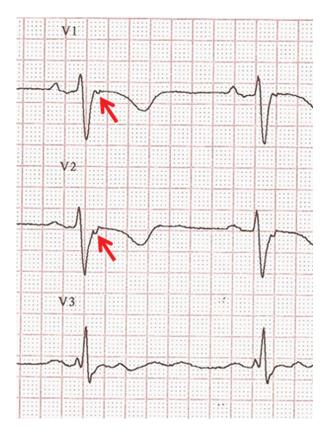


Figure 2 A double speed and amplitude 12-lead ECG (speed 50 mm/s, amplitude 20 mm/mV) demonstrate ϵ -waves (red arrows).

¹The Heart Hospital, University College of London, London, UK;

²Department of Cardiology, University of Rome "Sapienza", Via di Grottarossa, Rome, Italy

BMJ Case Reports

Competing interests None.

Patient consent Obtained.

 Nasir K, Bomma C, Tandri H, et al. Electrocardiographic features of arrhythmogenic right ventricular dysplasia/cardiomyopathy according to disease severity: a need to broaden diagnostic criteria. Circulation 2004;110:1527e34.

REFERENCES

 Marcus FI, McKenna WJ, Sherrill D, et al. Diagnosis of arrhythmogenic right ventricular cardiomyopathy/dysplasia: proposed modification of the task force criteria. Circulation 2010;121:1533–41.

This pdf has been created automatically from the final edited text and images.

Copyright 2011 BMJ Publishing Group. All rights reserved. For permission to reuse any of this content visit http://group.bmj.com/group/rights-licensing/permissions.

BMJ Case Report Fellows may re-use this article for personal use and teaching without any further permission.

Please cite this article as follows (you will need to access the article online to obtain the date of publication).

O'Mahony C, Cauti FM, Pantazis A. The ECG in arrhythmogenic right ventricular cardiomyopathy: ε-waves and anterior T-wave inversion. BMJ Case Reports 2011;10.1136/bcr.09.2010.3318, date of publication

Become a Fellow of BMJ Case Reports today and you can:

- ► Submit as many cases as you like
- ► Enjoy fast sympathetic peer review and rapid publication of accepted articles
- ► Access all the published articles
- ▶ Re-use any of the published material for personal use and teaching without further permission

For information on Institutional Fellowships contact consortiasales@bmjgroup.com

Visit casereports.bmj.com for more articles like this and to become a Fellow