

A case report of arrhythmogenic right ventricular dysplasia

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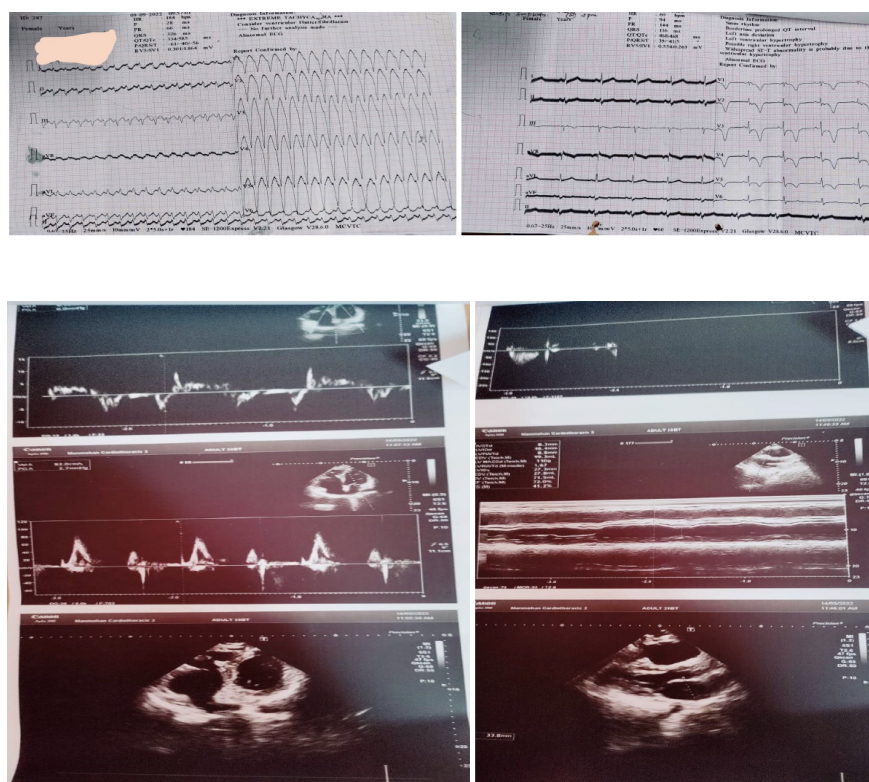
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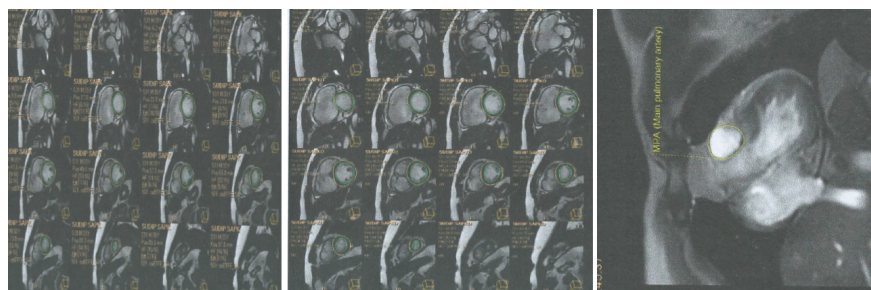
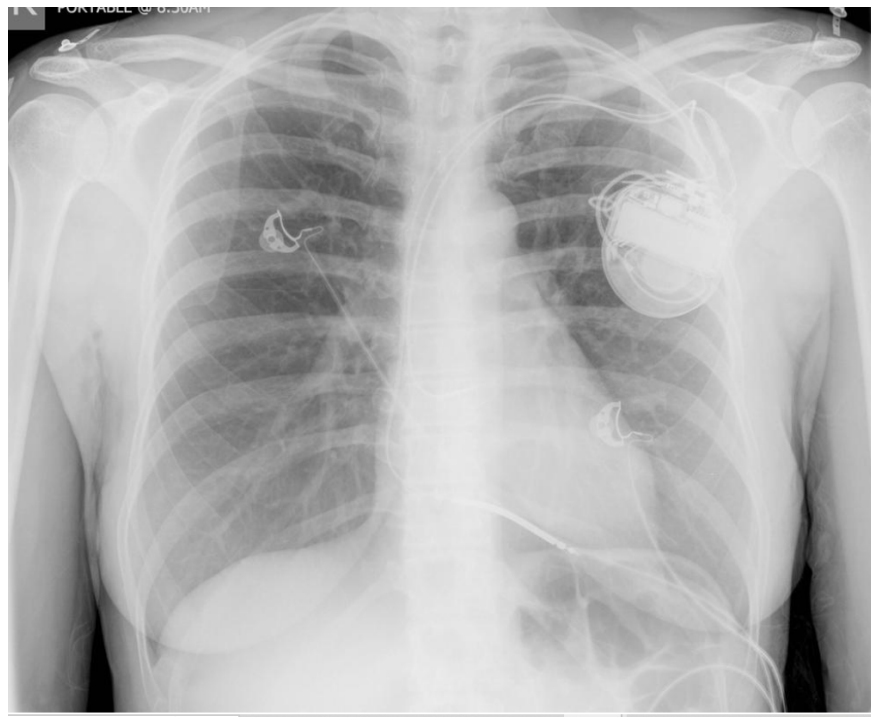
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Arrhythmogenic right ventricular tachycardia.docx available at <https://authorea.com/users/612442/articles/640332-a-case-report-of-arrhythmogenic-right-ventricular-dysplasia>





Category	Major criteria	Minor criteria
<i>Global and/or regional dysfunction and structural alterations</i>	1. Severe dilatation and reduction of right ventricular ejection fraction with no (or only mild) LV impairment 2. Localized right ventricular aneurysms (akinetic or dykinetic areas with diastolic bulging) 3. Severe segmental dilatation of the right ventricle	1. Mild global right ventricular dilatation and/or ejection fraction reduction with normal left ventricle 2. Mild segmental dilatation of the right ventricle 3. Regional right ventricular hypokinesia
<i>Tissue characterization of walls</i>	1. Fibrofatty replacement of myocardium on endomyocardial biopsy	
<i>Repolarisation abnormalities</i>		1. Inverted T waves in right precordial leads (V₂ and V₃) (people aged more than 12 yr; in absence of the right bundle branch block)
<i>Depolarisation/conduction abnormalities</i>	1. Epsilon waves or localized prolongation (> 110 ms) of the QRS complex in right precordial leads (V ₁ -V ₃)	1. Late potentials (signal-averaged ECG)
<i>Arrhythmias</i>		1. Left bundle branch block type ventricular tachycardia (sustained and non-sustained) (ECG, Holter, exercise testing). 2. Frequent ventricular extrasystoles (more than 1000/24 h) (Holter)
<i>Family history</i>	1. Familial disease confirmed at necropsy or surgery	1. Familial history of premature sudden death (<35 yr) due to suspected right ventricular dysplasia. Familial history (clinical diagnosis based on present criteria)
Dignosis of Arrhythmogenic Right Ventricular Dysplasia (ARVD)-Two major criteria or One major and Two minor criteria or Four minor criteria		

