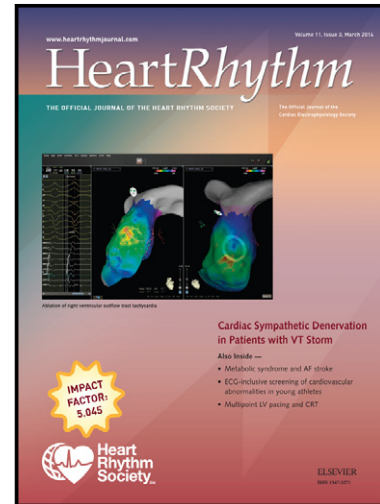


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Clinical and molecular characterization of a cardiac ryanodine receptor founder mutation causing catecholaminergic polymorphic ventricular tachycardia

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Clinical and molecular characterization of a cardiac ryanodine receptor founder mutation causing catecholaminergic polymorphic ventricular tachycardia

SHORT TITLE: RyR2 founder mutation causing CPVT

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CONTRIBUTIONS: FW and CBC contributed equally to this work. FW, PB, PMRH, and JB, performed clinical investigation, study design, and clinical data collection and interpretation. CP performed genealogy investigation. CBC, OC, CA, and AI performed genetic testing and genetic data analysis. CBC, PBA, GJP and FSS, performed functional experimental design, data collection,

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