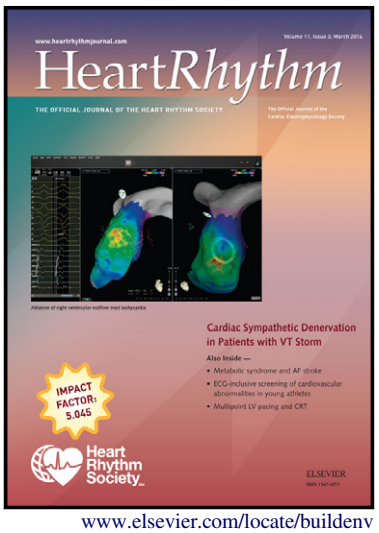

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Clinical and molecular characterization of a cardiac ryanodine receptor founder mutation causing catecholaminergic polymorphic ventricular tachycardia

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SHORT TITLE: RyR2 founder mutation causing CPVT

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