DISEASE:
Idiopathic ventricular fibrillation, non Brugada type

NAME:	Idiopathic ventricular fibrillation, non Brugada type
DESCRIPTION:	A rare, genetic, cardiac rhythm disease characterized by ventricular fibrillation in the absence of any structural or functional heart disease, or known repolarization abnormalities. The presence of J waves is associated with a higher risk of nocturnal ventricular fibrillation events and a higher risk of recurrence.
ORPHACODE:	228140
SYNONYMS:	Familial paroxysmal ventricular fibrillation, non Brugada type
XREF(S):	Orphanet OMIM OMIM ICD-10
ANALYTE(S):	SCN5A DPP6
CREATED:	13 May 2019 - 01:02
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RELATED CONTENT

Related Genetic Tests

- Cardiopathies, hereditary (gene panel)
- Primary Electrical disorders / Brugada syndrome / Long QT syndrome (LQT) / Short QT syndrome (SQT) / Arrhythmogenic right ventricular cardiomyopathy (ARVC) / Catecholaminergic polymorphic ventricular tachycardia (CPVT) (gene panel)
- Primary cardiac arrhythmias (Atrial fibrillation / Brugada syndome / Catech. polymorphic ventricular tachycardia / Early repolaristion syndrome / Ideopathic ventricular fibrillation / Long QT syndrome / Sick sinus syndrome / Short QT syndrome) (gene pane)

Related Laboratories

- Centrum Medische Genetica - UZ Antwerpen
- Centrum Medische Genetica - UZ Brussel VUB
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- dipeptidyl peptidase like 6
- sodium voltage-gated channel alpha subunit 5

Related Gene Panels

- Cardiopathies, hereditary (102 genes) - KUL
- Primary Electrical disorders/Brugada syndrome (genepanel) - UZA
- Primary cardiac arrhythmias (113 genes) - VUB
