DISEASE:
Familial isolated arrhythmogenic ventricular dysplasia, left dominant form

NAME:	Familial isolated arrhythmogenic ventricular dysplasia, left dominant form
ORPHACODE:	293888
SYNONYMS:	Familial isolated arrhythmogenic ventricular cardiomyopathy, left dominant form
XREF(S):	Orphanet ICD-10 OMIM OMIM

ANALYTE(S):	PKP2 DSC2 TTN TMEM43 RYR2 TGFB3 DSG2 DSP JUP LDB3 LMNA CTNNA3
CREATED:	13 May 2019 - 01:02
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RELATED CONTENT

Related Genetic Tests

- Cardiomyopathy, hereditary (gene panel)
- Cardiomyopathy: hypertrophic cardiomyopathy, dilated cardiomyopathy, restrictive cardiomyopathy, left ventricular non-compaction cardiomyopathy, arrhythmogenic right ventricular cardiomyopathy (gene panel)
- 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)

Related Laboratories

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

Related Analytes

- catenin alpha 3
- desmocollin 2
- desmoglein 2
- desmoplakin
- junction plakoglobin
- LIM domain binding 3
- lamin A/C
- plakophilin 2

- ryanodine receptor 2
- transforming growth factor beta 3
- transmembrane protein 43
- titin

Related Gene Panels

- Cardiomyopathy (genepanel) - UZA
- Cardiomyopathy, hereditary (208 genes) - VUB
- Cardiopathies, hereditary (102 genes) - KUL
- Primary Electrical disorders/Brugada syndrome (genepanel) - UZA
