Belgian Genetic Tests database

ANALYTE:
CACNB2

NAME:	calcium voltage-gated channel auxiliary subunit beta 2
SYMBOL:	CACNB2
VERSION OF ORPHANET:	2023-06-22 14:14:43
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Cardiomyopathy, hereditary (gene panel)
- Cardiopathies, hereditary (gene panel)
- Heart / Cardio disorders / Cardiopathy (gene panel)
- Inherited cardiac arrhytmia (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders 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 Diseases

- Brugada syndrome

Related Gene Panels

- Brugada syndrome (16 genes) - VUB
- Cardiomyopathy, hereditary (208 genes) - VUB
- Cardiopathies, hereditary (102 genes) - KUL
- Inherited cardiac arrhytmia (25 genes) - IPG
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Primary Electrical disorders/Brugada syndrome (genepanel) - UZA

- Primary cardiac arrhythmias (113 genes) - VUB
- cardiopathy panel - UGent
