Belgian Genetic Tests database

ANALYTE:
KCNJ5

NAME:	potassium inwardly rectifying channel subfamily J member 5
SYMBOL:	KCNJ5
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	CIR G protein-activated inward rectifier potassium channel 4 GIRK4 KATP1 Kir3.4 LQT13
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Cardiopathies, hereditary (gene panel)
- Heart / Cardio disorders / Cardiopathy (gene panel)
- Inherited cardiac arrhytmia (gene panel)
- Nephropathies, 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)
- Tubulopathy (gene panel)

Related Diseases

- Andersen-Tawil syndrome
- Familial atrial fibrillation
- Familial hyperaldosteronism type III
- NON RARE IN EUROPE: Aldosterone-producing adenoma
- Romano-Ward syndrome

Related Gene Panels

- Cardiopathies, hereditary (102 genes) - KUL
- Inherited cardiac arrhytmia (25 genes) - IPG
- Long QT (14 genes) - VUB
- Nephropathies, hereditary (219 genes) - KUL

- Primary Electrical disorders/Brugada syndrome (genepanel) - UZA
- Primary cardiac arrhythmias (113 genes) - VUB
- Tubulopathy/Nephrolithiasis (106 genes) - IPG
- cardiopathy panel - UGent

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