Belgian Genetic Tests database

ANALYTE:
KCNJ1

NAME:	potassium inwardly rectifying channel subfamily J member 1
SYMBOL:	KCNJ1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	ATP-sensitive inward rectifier potassium channel 1 Kir1.1 ROMK1
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy gene panel
- Inherited Kidney Diseases (Gene Panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Tubulopathy (gene panel)

Related Diseases

- Antenatal Bartter syndrome
- Bartter syndrome type 2

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1
- Tubulopathy/Nephrolithiasis (106 genes) - IPG

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