Belgian Genetic Tests database

ANALYTE:
KCNJ10

NAME:	potassium inwardly rectifying channel subfamily J member 10
SYMBOL:	KCNJ10
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	Kir1.2 Kir4.1
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Ataxia (autosomic dominant and recessive / except expansion of triplets) (gene panel - 722 genes)
- Ataxia (gene panel)
- Ataxia Spasticity (gene panel)
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy (gene panel)
- Epilepsy gene panel
- Epilepsy, seizures (gene panel)
- Hearing loss (deafness), (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Tubulopathy (gene panel)

Related Diseases

- EAST syndrome
- Pendred syndrome

Related Gene Panels

- Ataxia (141 genes) - KUL
- Ataxia (348 genes) - ULB
- Ataxia Spasticity - UGent
- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Epilepsy, seizures (196 genes) - IPG
- Hearing loss (deafness) (genepanel) - UZA
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Panel Nephro-ULG-V1
- Rare epilepsy with developmental delay (> 240 genes) - UZA
- Tubulopathy/Nephrolithiasis (106 genes) - IPG
- test
