Belgian Genetic Tests database

ANALYTE:
CLCN2

NAME:	chloride voltage-gated channel 2
SYMBOL:	CLCN2
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	CLC2 ClC-2 EJM6
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
- Hereditary Spastic Paraplegia (gene panel)
- Intellectual disability (virtual gene panel)
- Leukodystrophy (gene panel)
- Movement Disorders (gene panel)
- Nephropathies, hereditary (gene panel)
- Tubulopathy (gene panel)

Related Diseases

- Familial hyperaldosteronism type II
- Juvenile myoclonic epilepsy
- Leukoencephalopathy with mild cerebellar ataxia and white matter edema

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
- Hereditary Spastic Paraplegia & ataxia (genepanel) - UZA
- Intellectual disability (gene panel)
- Leukodystrophy - UGent
- Movement Disorders - ULG
- Nephropathies, hereditary (219 genes) - KUL
- Tubulopathy/Nephrolithiasis (106 genes) - IPG
