ANALYTE:
CLDN16

NAME:	claudin 16
SYMBOL:	CLDN16
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	HOMG3 PCLN1 hypomagnesemia 3, with hypercalciuria and nephrocalcinosis paracellin-1
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
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RELATED CONTENT

Related Genetic Tests

- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy gene panel
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability & Epilepsy (gene panel)
- Nephrocalcinosis and nephrolithiasis (gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Tubulopathy (gene panel)

Related Diseases

- Primary hypomagnesemia with hypercalciuria and nephrocalcinosis without severe ocular involvement

Related Gene Panels

- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Intellectual disability & Epilepsy - UGent
- Nephrocalcinosis and nephrolithiasis (37 genes) - IPG
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1
- Tubulopathy/Nephrolithiasis (106 genes) - IPG

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