

ANALYTE:
CLCN5

NAME:	chloride voltage-gated channel 5
SYMBOL:	CLCN5
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	CLC5 CIC-5 DENTS Dent disease XLRH XRN hCIC-K2 hCIC-K2
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

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- [Genetic disorders of Calcium and Phosphate metabolism \(gene panel\)](#)
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- [Nephrocalcinosis and nephrolithiasis \(gene panel\)](#)
- [Nephrogenetics / Nephropathy \(gene panel\)](#)
- [Nephropathies, hereditary \(gene panel\)](#)
- [Nephrotic syndrome, Focal Segmental Glomerulosclerosis \(FSGS\) , Alport syndrome and podocytopathy \(gene panel\)](#)
- [Skeletal dysplasia \(gene panel\)](#)
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Related Diseases

- [Dent disease type 1](#)

Related Gene Panels

- [End-stage renal disease \(106 genes\) - IPG](#)
- [Genetic disorders of Calcium and Phosphate metabolism \(31 genes\) - KUL](#)
- [Intellectual disability \(gene panel\)](#)
- [Nephrocalcinosis and nephrolithiasis \(37 genes\) - IPG](#)
- [Nephropathies, hereditary \(219 genes\) - KUL](#)

- Nephropathy panel - UGent
- Nephrotic syndrome, FSGS, Alport syndrome (76 genes) - IPG
- Panel Nephro-ULG-V1
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
- Tubulopathy/Nephrolithiasis (106 genes) - IPG

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