Belgian Genetic Tests database

ANALYTE:
SLC12A1

NAME:	solute carrier family 12 member 1
SYMBOL:	SLC12A1
VERSION OF ORPHANET:	2021-08-01 04:46:43
SYNONYMS:	NKCC2
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
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Source URL: http://gentest.healthdata.be/analyte/3918

RELATED CONTENT

Related Genetic Tests

- Cardiomyopathy, hereditary (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Inherited Kidney Diseases (Gene Panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Tubulopathy (gene panel)

Related Diseases

- Antenatal Bartter syndrome

Related Gene Panels

- Cardiomyopathy, hereditary (208 genes) - VUB
- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1
- Tubulopathy/Nephrolithiasis (106 genes) - IPG