ANALYTE:
SLC4A4

NAME:	solute carrier family 4 member 4
SYMBOL:	SLC4A4
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	HNBC1 NBC1 NBC2 hhNMC pNBC
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/analyte/3912

RELATED CONTENT

Related Genetic Tests

- 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

- Autosomal recessive proximal renal tubular acidosis

Related Gene Panels

- 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

- Tubulopathy/Nephrolithiasis (106 genes) - IPG
