Belgian Genetic Tests database

ANALYTE:
SLC7A9

NAME:	solute carrier family 7 member 9
SYMBOL:	SLC7A9
VERSION OF ORPHANET:	2023-06-22 14:14:43
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Inherited Kidney Diseases (Gene Panel)
- Metabolic disorders including disorders of glycosylation, peroxisomal disorders, organic acidurias, glycogenosis disorders, neurotransmitter disorders (213 genes)
- Nephrocalcinosis and nephrolithiasis (gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Tubulopathy (gene panel)

Related Diseases

- Cystinuria type B

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Metabolic disorders (213 genes) - VUB
- 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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