Belgian Genetic Tests database

ANALYTE:
G6PC1

NAME:	glucose-6-phosphatase catalytic subunit 1
SYMBOL:	G6PC1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	G6PC1 GSD1a glycogen storage disease type I, von Gierke disease
XREF(S):	Orphanet HGNC OMIM Reactome SwissProt Ensembl Genatlas
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RELATED CONTENT

Related Genetic Tests

- Glycogen storage disease type 1a
- Hepatology (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Metabolic disorders including disorders of glycosylation, peroxisomal disorders, organic acidurias, glycogenosis disorders, neurotransmitter disorders (213 genes)
- Nephrogenetics / Nephropathy (gene panel)
- Nephrotic syndrome, Focal Segmental Glomerulosclerosis (FSGS) , Alport syndrome and podocytopathy (gene panel)
- Tubulopathy (gene panel)

Related Diseases

- Glycogen storage disease due to glucose-6-phosphatase deficiency type Ia

Related Gene Panels

- Hepatology panel - UGent
- Metabolic disorders (213 genes) - VUB
- Nephropathy panel - UGent
- Nephrotic syndrome, FSGS, Alport syndrome (76 genes) - IPG
- Panel Nephro-ULG-V1
- Tubulopathy/Nephrolithiasis (106 genes) - IPG

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