Belgian Genetic Tests database

ANALYTE:
GALT

NAME:	galactose-1-phosphate uridylyltransferase
SYMBOL:	GALT
VERSION OF ORPHANET:	2023-06-22 14:14:43
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/3773

RELATED CONTENT

Related Genetic Tests

- Cataract (gene panel)
- Cerebral palsy (gene panel)
- Congenital disorders of glycosylation (79 genes)
- Hepatology (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Leukodystrophy (gene panel)
- Metabolic disorders including disorders of glycosylation, peroxisomal disorders, organic acidurias, glycogenosis disorders, neurotransmitter disorders (213 genes)
- Movement Disorders (gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Tubulopathy (gene panel)

Related Diseases

- Classic galactosemia

Related Gene Panels

- Cataract - UGent

- Cerebral palsy (212 genes) - UZA
- Congenital disorders of glycosylation (79 genes) - KUL
- Hepatology panel - UGent
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Leukodystrophy - UGent
- Metabolic disorders (213 genes) - VUB
- Movement Disorders - ULG
- 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
