ANALYTE:
ALG1

NAME:	ALG1 chitobiosyldiphosphodolichol beta-mannosyltransferase
SYMBOL:	ALG1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	CDG1K HMAT1 HMT-1 Mat-1
XREF(S):	Orphanet Genatlas HGNC OMIM Reactome SwissProt Ensembl
CREATED:	13 May 2019 - 01:01
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RELATED CONTENT

Related Genetic Tests

- Congenital disorders of glycosylation (79 genes)
- Congenital malformation (gene panel - 1721 genes)
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy gene panel
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Metabolic disorders including disorders of glycosylation, peroxisomal disorders, organic acidurias, glycogenosis disorders, neurotransmitter disorders (213 genes)
- Nephrogenetics / Nephropathy (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel

Related Diseases

- ALG1-CDG

Related Gene Panels

- Congenital disorders of glycosylation (79 genes) - KUL
- Congenital malformation (1721 genes) - ULB
- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Intellectual disability & Epilepsy - UGent

- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Metabolic disorders (213 genes) - VUB
- Nephropathy panel - UGent
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
