Belgian Genetic Tests database

ANALYTE:
ALG9

NAME:	ALG9 alpha-1,2-mannosyltransferase
SYMBOL:	ALG9
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	dol-P-Man dependent alpha-1,2-mannosyltransferase dolichyl-P-Man:Man(6)GlcNAc(2)-PP-dolichol alpha-1,2-mannosyltransferase dolichyl-P-Man:Man(8)GlcNAc(2)-PP-dolichol alpha-1,2-mannosyltransferase
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Ciliopathy / polycystic kidney and liver diseases / ADTKD/ nephronophtisis / Bardet-Biedl syndromes and kidney cancers (gene panel)
- Congenital disorders of glycosylation (79 genes)
- Congenital malformation (gene panel - 1721 genes)
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy gene panel
- Hepatology (gene panel)
- Inherited Kidney Diseases (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)
- Nephropathies, hereditary (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Diseases

- ALG9-CDG
- Autosomal dominant polycystic kidney disease

Related Gene Panels

- Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophtisis, Bardet-Biedl syndromes and kidney cancers (146 genes) - IPG
- Cleft lip and palate / dysmorphic facial features / craniofacial anomalies (255 genes)) - UCL
- Congenital disorders of glycosylation (79 genes) - KUL
- Congenital malformation (1721 genes) - ULB
- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Hepatology panel - UGent
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Metabolic disorders (213 genes) - VUB
- 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
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
