Belgian Genetic Tests database

ANALYTE:
SLC25A24

NAME:	solute carrier family 25 member 24
SYMBOL:	SLC25A24
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	APC1 DKFZp586G0123
XREF(S):	Orphanet Ensembl OMIM Genatlas HGNC SwissProt
CREATED:	13 May 2019 - 01:01
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Craniosynostosis (gene panel)
- Intellectual disability (virtual gene panel)
- Mitochondrial disorders (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Skeletal dysplasia (gene panel)

Related Diseases

- Gorlin-Chaudhry-Moss syndrome
- Progeroid syndrome, Petty type

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Craniosynostosis (32 genes) - KUL
- Intellectual disability (gene panel)
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- mitochondrial disease, nuclear based (343 genes) - VUB

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