Belgian Genetic Tests database

ANALYTE:
MYO18B

NAME:	myosin XVIIIB
SYMBOL:	MYO18B
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	BK125H2.1
XREF(S):	Orphanet HGNC OMIM Genatlas SwissProt Ensembl
CREATED:	13 May 2019 - 01:01
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Myopathy (gene panel)
- Myopathy (gene panel)
- Neuromuscular disorders (232 genes (= myopathy, metabolic myopathy, ion channel muscle diseases, muscular dystrophy, myotonic dystrophy, rhabdomyolysis, myasthenia)
- Neuromuscular disorders (548 genes)
- Neuromuscular disorders (gene panel)
- Skeletal dysplasia (gene panel)

Related Diseases

- Klippel-Feil anomaly-myopathy-facial dysmorphism syndrome

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Myopathy (332 genes) - IPG
- Myopathy (genepanel) - UZA
- Neuromuscular disorders (548 genes) - ULB
- Neuromuscular disorders (232 genes) - KUL
- Neuromuscular disorders - UGent
- Skeletal dysplasia (genepanel) - UZA

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