ANALYTE:
MYH3

NAME:	myosin heavy chain 3
SYMBOL:	MYH3
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	HEMHC MYHC-EMB MYHSE1 SMHCE muscle embryonic myosin heavy chain 3 myosin, skeletal, heavy chain, embryonic 1
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

- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Intellectual disability (virtual gene panel)
- 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)
- Neuromuscular disorders : congenital & distal myopathy, congenital muscle dystrophy / Limb-girdle muscular dystrophy / Rhabdomyolysis / Myopathy (with prominent contractures) / distal artrogryposis (gene panel)
- Skeletal dysplasia (gene panel)

Related Diseases

- Autosomal dominant multiple pterygium syndrome
- Autosomal recessive multiple pterygium syndrome
- Distal arthrogryposis type 1
- Freeman-Sheldon syndrome
- Sheldon-Hall syndrome
- Spondylocarpotarsal synostosis

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Intellectual disability (gene panel)
- Myopathy (332 genes) - IPG
- Myopathy (genepanel) - UZA
- Neuromuscular disorders (548 genes) - ULB
- Neuromuscular disorders (166 genes) - VUB
- Neuromuscular disorders (232 genes) - KUL
- Neuromuscular disorders - UGent
- Skeletal dysplasia (genepanel) - UZA
