ANALYTE:
PYGM

NAME:	glycogen phosphorylase, muscle associated
SYMBOL:	PYGM
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	GSD5 McArdle syndrome glycogen phosphorylase, muscle form glycogen storage disease type V myophosphorylase
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Hepatology (gene panel)
- Mc Ardle disease, glycogene storage disease type V
- Metabolic disorders including disorders of glycosylation, peroxisomal disorders, organic acidurias, glycogenosis disorders, neurotransmitter disorders (213 genes)
- Myopathy (gene panel)
- Myopathy (gene panel)
- Nephrogenetics / Nephropathy (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)

Related Diseases

- Glycogen storage disease due to muscle glycogen phosphorylase deficiency

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Hepatology panel - UGent
- Metabolic disorders (213 genes) - VUB

- Myopathy (332 genes) - IPG
- Myopathy (genepanel) - UZA
- Nephropathy panel - UGent
- Neuromuscular disorders (548 genes) - ULB
- Neuromuscular disorders (166 genes) - VUB
- Neuromuscular disorders (232 genes) - KUL
- Neuromuscular disorders - UGent
