Belgian Genetic Tests database

ANALYTE:
ANO5

NAME:	anoctamin 5
SYMBOL:	ANO5
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	GDD1
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Intellectual disability (virtual gene panel)
- Myopathy (gene panel)
- Myopathy (gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders 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)
- Osteogenesis Imperfecta (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)

Related Diseases

- Anoctamin-5-related limb-girdle muscular dystrophy R12
- Distal anoctaminopathy
- Gnathodiaphyseal dysplasia

Related Gene Panels

- Intellectual disability (gene panel)
- Myopathy (332 genes) - IPG
- Myopathy (genepanel) - UZA
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Neuromuscular disorders (548 genes) - ULB
- Neuromuscular disorders (166 genes) - VUB
- Neuromuscular disorders (232 genes) - KUL
- Neuromuscular disorders - UGent
- Osteogenesis Imperfecta (25 genes) - KUL
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
