Belgian Genetic Tests database

ANALYTE:
ACADVL

NAME:	acyl-CoA dehydrogenase very long chain
SYMBOL:	ACADVL
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	ACAD6 LCACD VLCAD
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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Source URL: http://gentest.healthdata.be/analyte/3185

RELATED CONTENT

Related Genetic Tests

- Ataxia (autosomic dominant and recessive / except expansion of triplets) (gene panel - 722 genes)
- Cardiomyopathy, hereditary (gene panel)
- Congenital malformation (gene panel - 1721 genes)
- Heart / Cardio disorders / Cardiopathy (gene panel)
- Hepatology (gene panel)
- Hypertrophic cardiomyopathy (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)

Related Diseases

- Very long chain acyl-CoA dehydrogenase deficiency

Related Gene Panels

- Ataxia (348 genes) - ULB
- Cardiomyopathy, hereditary (208 genes) - VUB

- Congenital malformation (1721 genes) - ULB
- Hepatology panel - UGent
- Hypertrophic cardiomyopathy (75 genes) - IPG
- 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
- cardiopathy panel - UGent
