

ANALYTE:
ACADL

NAME:	acyl-CoA dehydrogenase long chain
SYMBOL:	ACADL
VERSION OF ORPHANET:	2020-02-01 05:07:10
SYNONYMS:	ACAD4 LCAD
XREF(S):	Orphanet Genatlas HGNC OMIM Reactome SwissProt Ensembl
CREATED:	13 May 2019 - 01:01
CHANGED:	01 Feb 2020 - 12:07

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RELATED CONTENT

Related Genetic Tests

- 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)

Related Diseases

- Long chain acyl-CoA dehydrogenase deficiency

Related Gene Panels

- Neuromuscular disorders (548 genes) - ULB
- Neuromuscular disorders (232 genes) - KUL
- Neuromuscular disorders - UGent

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