DISEASE:
Very long chain acyl-CoA dehydrogenase deficiency

NAME:	Very long chain acyl-CoA dehydrogenase deficiency
DESCRIPTION:	Very long-chain acyl-CoA dehydrogenase (VLCAD) deficiency (VLCADD) is an inherited disorder of mitochondrial long-chain fatty acid oxidation with a variable presentation including: cardiomyopathy, hypoketotic hypoglycemia, liver disease, exercise intolerance and rhabdomyolysis.
ORPHACODE:	26793
SYNONYMS:	VLCAD deficiency VLCADD
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	ACADVL
CREATED:	13 May 2019 - 01:02
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RELATED CONTENT

Related Genetic Tests

- Cardiomyopathy, hereditary (gene panel)
- Neuromuscular disorders : congenital & distal myopathy, congenital muscle dystrophy / Limb-girdle muscular dystrophy / Rhabdomyolysis / Myopathy (with prominent contractures) / distal artrogryposis (gene panel)

Related Laboratories

- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

- acyl-CoA dehydrogenase very long chain

Related Gene Panels

- Cardiomyopathy, hereditary (208 genes) - VUB
- Neuromuscular disorders (166 genes) - VUB
