DISEASE:
Multiple acyl-CoA dehydrogenase deficiency, mild type

NAME:	Multiple acyl-CoA dehydrogenase deficiency, mild type
ORPHACODE:	394532
SYNONYMS:	Glutaric aciduria type 2, mild type MAD deficiency, mild type MADD, mild type
XREF(S):	Orphanet OMIM OMIM ICD-10
ANALYTE(S):	SLC25A32 FLAD1 ETFA ETFB ETFDH
CREATED:	13 May 2019 - 01:02
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RELATED CONTENT

Related Genetic Tests

- 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

- electron transfer flavoprotein subunit alpha
- electron transfer flavoprotein subunit beta
- electron transfer flavoprotein dehydrogenase
- flavin adenine dinucleotide synthetase 1
- solute carrier family 25 member 32

Related Gene Panels

- Neuromuscular disorders (166 genes) - VUB