DISEASE:
Leigh syndrome with cardiomyopathy

NAME:	Leigh syndrome with cardiomyopathy
ORPHACODE:	70474
SYNONYMS:	Cardiomyopathy with hypotonia due to cytochrome C oxidase deficiency Cardiomyopathy with myopathy due to COX deficiency Leigh disease with myopathy
XREF(S):	Orphanet OMIM OMIM ICD-10 OMIM
ANALYTE(S):	NDUFAF3 NDUFB8 SCO2 SURF1 NDUFS2
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RELATED CONTENT

Related Genetic Tests

- Leigh / NARP Syndrome
- Leigh syndrome
- Mitochondrial disorders (gene panel)

Related Laboratories

- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

- NADH:ubiquinone oxidoreductase complex assembly factor 3
- NADH:ubiquinone oxidoreductase subunit B8
- NADH:ubiquinone oxidoreductase core subunit S2
- synthesis of cytochrome C oxidase 2
- SURF1 cytochrome c oxidase assembly factor

Related Gene Panels

- Leigh syndrome (mtDNA / 37 genes) - VUB
- mitochondrial disease, nuclear based (343 genes) - VUB