
DISEASE:
Acyl-CoA dehydrogenase 9 deficiency

NAME:	Acyl-CoA dehydrogenase 9 deficiency
DESCRIPTION:	A rare disorder characterized by neurological dysfunction, hepatic failure and cardiomyopathy due to a deficiency of complex I of the respiratory chain.
ORPHACODE:	99901
SYNONYMS:	ACAD9 deficiency
XREF(S):	<u>Orphanet</u> <u>OMIM</u> <u>ICD-10</u>
ANALYTE(S):	<u>ACAD9</u>
CREATED:	13 May 2019 - 01:02
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Source URL: <http://gentest.healthdata.be/disease/3786>

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