

DISEASE:**Acute infantile liver failure-multisystemic involvement syndrome**

NAME:	Acute infantile liver failure-multisystemic involvement syndrome
DESCRIPTION:	A rare, genetic, parenchymal hepatic disease characterized by acute liver failure, that occurs in the first year of life, which manifests with failure to thrive, hypotonia, moderate global developmental delay, seizures, abnormal liver function tests, microcytic anemia and elevated serum lactate. Other associated features include hepatosteatorrhea and fibrosis, abnormal brain morphology, and renal tubulopathy. Minor illness exacerbates deterioration of liver failure.
ORPHACODE:	370088
XREF(S):	<u>Orphanet</u> <u>ICD-10</u> <u>OMIM</u>
ANALYTE(S):	<u>LARS1</u>
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