DISEASE:
Familial LCAT deficiency

NAME:	Familial LCAT deficiency
DESCRIPTION:	Familial LCAT (lecithin-cholesterol acyltransferase) deficiency (FLD) is a form of lecithin-cholesterol acyltransferase deficiency (LCAT; see this term) characterized clinically by corneal opacities, hemolytic anemia, and renal failure, and biochemically by severely decreased HDL cholesterol and complete deficiency of the LCAT enzyme.
ORPHACODE:	79293
SYNONYMS:	Complete LCAT deficiency FLD Norum disease
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	LCAT
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