DISEASE:
Smith-Lemli-Opitz syndrome

NAME:	Smith-Lemli-Opitz syndrome
DESCRIPTION:	Smith-Lemli-Opitz syndrome (SLOS) is characterized by multiple congenital anomalies, intellectual deficit, and behavioral problems.
ORPHACODE:	818
SYNONYMS:	7-dehydrocholesterol reductase deficiency RSH syndrome SLOS
XREF(S):	Orphanet MeSH ICD-10 OMIM
ANALYTE(S):	DHCR7
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- Short Stature (gene panel)
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Related Laboratories

- Centre de Génétique Médicale UCL
- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
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Related Analytes

- 7-dehydrocholesterol reductase

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