DISEASE:
Alpers-Huttenlocher syndrome

NAME:	Alpers-Huttenlocher syndrome
DESCRIPTION:	A cerebrohepatopathy and a rare and severe form of mitochondrial DNA (mtDNA) depletion syndrome characterized by the triad of progressive developmental regression, intractable seizures, and hepatic failure.
ORPHACODE:	726
SYNONYMS:	Alpers progressive sclerosing poliodystrophy Alpers syndrome Progressive neuronal degeneration of childhood with liver disease
XREF(S):	Orphanet ICD-10 OMIM MeSH MedDRA
ANALYTE(S):	POLG
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