DISEASE:
Lafora disease

NAME:	Lafora disease
DESCRIPTION:	A rare, inherited, severe, progressive myoclonic epilepsy characterized by myoclonus and/or generalized seizures, visual hallucinations (partial occipital seizures), and progressive neurological decline.
ORPHACODE:	501
SYNONYMS:	EPM2 PME type 2 Progressive myoclonic epilepsy type 2 Progressive myoclonus epilepsy type 2
XREF(S):	Orphanet OMIM MeSH MedDRA ICD-10
ANALYTE(S):	EPM2A NHLRC1
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