DISEASE:
Lennox-Gastaut syndrome

NAME:	Lennox-Gastaut syndrome
DESCRIPTION:	A rare, severe early-onset developmental epileptic encephalopathy characterized by the triad of intellectual impairment, multiple seizure types, and typical electroencephalography (EEG) abnormalities.
ORPHACODE:	2382
XREF(S):	Orphanet OMIM OMIM OMIM MedDRA MeSH ICD-10 OMIM

ANALYTE(S):	CUX2 SCN1A MAPK10 CHD2 DNM1 GABRB3 CACNA1A CACNA1A
CREATED:	13 May 2019 - 01:02
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