Belgian Genetic Tests database

DISEASE:
CLN2 disease

NAME:	CLN2 disease
ORPHACODE:	228349
SYNONYMS:	Classic late infantile NCL Classic late infantile neuronal ceroid lipofuscinosis
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	TPP1
CREATED:	13 May 2019 - 01:02
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Source URL: http://gentest.healthdata.be/disease/1532

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