Belgian Genetic Tests database

DISEASE:
CDKL5-deficiency disorder

NAME:	CDKL5-deficiency disorder
DESCRIPTION:	A rare genetic neurodevelopmental disorder characterized by early-onset drug-resistant seizures and severe neurodevelopmental impairment with major motor development delay.
ORPHACODE:	505652
SYNONYMS:	CDD
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	CDKL5
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/disease/2593

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