Belgian Genetic Tests database

DISEASE:
Microlissencephaly

NAME:	Microlissencephaly
DESCRIPTION:	Microlissencephaly describes a heterogenous group of a rare cortical malformations characterized by lissencephaly in combination with severe congenital microcephaly, presenting with spasticity, severe developmental delay, and seizures and with survival varying from days to years.
ORPHACODE:	1083
XREF(S):	Orphanet ICD-10 OMIM OMIM
ANALYTE(S):	NDE1 KATNB1
CREATED:	13 May 2019 - 01:02
CHANGED:	01 Aug 2021 - 06:46

Source URL: http://gentest.healthdata.be/disease/1602

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Related Laboratories

- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

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- nudE neurodevelopment protein 1