DISEASE:
Subcortical band heterotopia

NAME:	Subcortical band heterotopia
DESCRIPTION:	A rare, non-syndromic cerebral malformation due to abnormal neuronal migration characterized by variable clinical manifestation depending on the location, size and thickness of subcortical bands. Clinical presentation ranges from mild cognitive deficit to developmental delay with severe intellectual disability, seizures and behavioral problems.
ORPHACODE:	99796
SYNONYMS:	Subcortical laminar heterotopia
XREF(S):	Orphanet OMIM OMIM OMIM ICD-10
ANALYTE(S):	DCX PAFAH1B1 EML1
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/disease/3810

RELATED CONTENT

Related Genetic Tests

- Lissencephaly / subcortical band heterotopia
- Lissencephaly, X-linked / subcortical band heterotopia
- Malformations of cortical development (235 genes)
- Subcortical band heterotopia, lissencephaly

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Brussel VUB

Related Analytes

- doublecortin
- EMAP like 1
- platelet activating factor acetylhydrolase 1b regulatory subunit 1