Belgian Genetic Tests database

DISEASE:
Kufor-Rakeb syndrome

NAME:	Kufor-Rakeb syndrome
DESCRIPTION:	Kufor-Rakeb syndrome (KRS) is a rare genetic neurodegenerative disorder characterized by juvenile Parkinsonism, pyramidal degeneration (dystonia), supranuclear palsy, and cognitive impairment.
ORPHACODE:	306674
SYNONYMS:	PARK9
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	ATP13A2
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Source URL: http://gentest.healthdata.be/disease/2053

RELATED CONTENT

Related Genetic Tests

- Neurodegeneration (gene panel)
- Neurodegeneration with Brain Iron Accumulation (gene panel)

Related Laboratories

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

Related Analytes

- ATPase cation transporting 13A2

Related Gene Panels

- Neurodegeneration (99 genes) - IPG
- Neurodegeneration with Brain Iron Accumulation (NBIA) - UGent