Belgian Genetic Tests database

DISEASE:
Autosomal recessive spastic paraplegia type 69

NAME:	Autosomal recessive spastic paraplegia type 69
DESCRIPTION:	A rare, complex hereditary spastic paraplegia disorder characterized by infantile onset of progressive lower limb spasticity, global developmental delay, hyperreflexia, clonus and extensor plantar reflexes, associated with dysarthria, intellectual disability, cataracts and hearing impairment.
ORPHACODE:	401830
SYNONYMS:	SPG69
XREF(S):	Orphanet ICD-10
ANALYTE(S):	RAB3GAP2
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Source URL: http://gentest.healthdata.be/disease/2264

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Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)

Related Analytes

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Related Gene Panels

- Spastic Paraplegia (89 genes) - IPG