DISEASE:
Autosomal dominant spastic paraplegia type 3

NAME:	Autosomal dominant spastic paraplegia type 3
DESCRIPTION:	A rare, pure or complex form of hereditary spastic paraplegia, with variable phenotype, typically characterized by childhood-onset of minimally progressive, bilateral, mainly symmetric lower limb spasticity and weakness, associated with pes cavus, scoliosis, sphincter disturbances and/or urinary bladder hyperactivity. Rare additional associated manifestations may include mild intellectual disability, axonal motor neuropathy, and seizures.
ORPHACODE:	100984
SYNONYMS:	Strümpell disease
XREF(S):	Orphanet ICD-10 MeSH OMIM
ANALYTE(S):	ATL1
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RELATED CONTENT

Related Genetic Tests

- Charcot-Marie-Tooth (other than type 1A) (gene panel, IPN panel)
- Hereditary Spastic Paraplegia (gene panel)
- Spastic Paraplegia (gene panel)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Antwerpen
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- atlastin GTPase 1

Related Gene Panels

- Inherited Peripheral Neuropathies gene panel (139 genes) - KUL
- Hereditary Spastic Paraplegia & ataxia (genepanel) - UZA
- Spastic Paraplegia (89 genes) - IPG