DISEASE:
Spastic paraplegia type 7

NAME:	Spastic paraplegia type 7
DESCRIPTION:	A form of hereditary spastic ataxia characterized by an onset usually in adulthood (but ranging from 10-72 years) of progressive bilateral lower limb weakness and spasticity and sometimes predominant cerebellar ataxia. In addition to frequent sphincter dysfunction and decreased vibratory sense at the ankles, manifestations may include optical neuropathy, nystagmus, blepharoptosis, ophthalmoplegia, decreased hearing, scoliosis, pes cavus, motor and sensory neuropathy, muscle atrophy, parkinsonism, and dystonia.
ORPHACODE:	99013
SYNONYMS:	SPG7
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	SPG7
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RELATED CONTENT

Related Genetic Tests

- Spastic Paraplegia (gene panel)

Related Laboratories

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

Related Analytes

- SPG7 matrix AAA peptidase subunit, paraplegin

Related Gene Panels

- Spastic Paraplegia (89 genes) - IPG