DISEASE:
Spastic paraplegia type 2

NAME:	Spastic paraplegia type 2
DESCRIPTION:	A rare, X-linked leukodystrophy characterized primarily by spastic gait and autonomic dysfunction. When additional central nervous system (CNS) signs, such as intellectual deficit, ataxia, or extrapyramidal signs, are present, the syndrome is referred to as complicated SPG.
ORPHACODE:	99015
SYNONYMS:	SPG2 Spastic gait type 2 Spastic paraparesis type 2 X-linked spastic paraplegia type 2
XREF(S):	Orphanet ICD-10 MeSH OMIM
ANALYTE(S):	PLP1
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RELATED CONTENT

Related Genetic Tests

- 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

Related Analytes

- proteolipid protein 1

Related Gene Panels

- Hereditary Spastic Paraplegia & ataxia (genepanel) - UZA
- Spastic Paraplegia (89 genes) - IPG