DISEASE:
Autosomal dominant spastic paraplegia type 12

NAME:	Autosomal dominant spastic paraplegia type 12
DESCRIPTION:	A pure form of hereditary spastic paraplegia characterized by a childhood- to adulthood-onset of slowly progressive lower limb spasticity and hyperreflexia of lower extremities, extensor plantar reflexes, distal sensory impairment, variable urinary dysfunction and pes cavus.
ORPHACODE:	100993
SYNONYMS:	SPG12
XREF(S):	Orphanet OMIM MeSH ICD-10
ANALYTE(S):	UBAP1 RTN2
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

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

- reticulon 2
- ubiquitin associated protein 1

Related Gene Panels

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