DISEASE:
Autosomal recessive spastic paraplegia type 62

NAME:	Autosomal recessive spastic paraplegia type 62
DESCRIPTION:	A pure or complex form of hereditary spastic paraplegia characterized by an onset in the first decade of life of spastic paraperesis (more prominent in lower than upper extremities) and unsteady gait, as well as increased deep tendon reflexes, amyotrophy, cerebellar ataxia, and flexion contractures of the knees, in some.
ORPHACODE:	401785
SYNONYMS:	SPG62
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	ERLIN1
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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

- ER lipid raft associated 1

Related Gene Panels

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