DISEASE:
Autosomal recessive spastic paraplegia type 70

NAME:	Autosomal recessive spastic paraplegia type 70
DESCRIPTION:	Autosomal recessive spastic paraplegia type 70 is a very rare, complex subtype of hereditary spastic paraplegia that presents in infancy with delayed motor development (i.e. crawling, walking) and is characterized by lower limb spasticity, increased deep tendon reflexes, extensor plantar responses, impaired vibratory sensation at ankles, amyotrophy and borderline intellectual disability. Additional signs may include gait disturbances, Achilles tendon contractures, scoliosis and cerebellar abnormalities.
ORPHACODE:	401835
SYNONYMS:	SPG70
XREF(S):	Orphanet ICD-10
ANALYTE(S):	MARS1
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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

- methionyl-tRNA synthetase 1

Related Gene Panels

- Spastic Paraplegia (89 genes) - IPG