DISEASE:
Autosomal recessive spastic paraplegia type 59

NAME:	Autosomal recessive spastic paraplegia type 59
DESCRIPTION:	Autosomal recessive spastic paraplegia type 59 is a very rare, complex hereditary spastic paraplegia characterized by an early onset of progressive lower limb spasticity, tip-toe walking, scissor gait, hyperreflexia and clonus that may be associated with borderline intellectual disability. Nystagmus and pes equinovarus have also been reported.
ORPHACODE:	401795
SYNONYMS:	SPG59
XREF(S):	Orphanet ICD-10
ANALYTE(S):	USP8
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RELATED CONTENT

Related Genetic Tests

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Related Laboratories

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

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- ubiquitin specific peptidase 8

Related Gene Panels

- Spastic Paraplegia (89 genes) - IPG