DISEASE:
Autosomal recessive spastic paraplegia type 71

NAME:	Autosomal recessive spastic paraplegia type 71
DESCRIPTION:	A type of autosomal recessive pure hereditary spastic paraplegia characterized by infancy onset of crural spastic paraperesis with scissors gait, extensor plantar response, and increased tendon reflexes. Neuroimaging reveals a thin corpus callosum and electromyography and nerve conduction velocity studies are normal.
ORPHACODE:	401840
SYNONYMS:	SPG71
XREF(S):	Orphanet ICD-10
ANALYTE(S):	ZFR
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- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)

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