DISEASE:
Autosomal recessive spastic paraplegia type 67

NAME:	Autosomal recessive spastic paraplegia type 67
DESCRIPTION:	Autosomal recessive spastic paraplegia type 67 is an extremely rare, complex hereditary spastic paraplegia characterized by an infancy or childhood onset of global developmental delay and progressive spasticity with tremor in the distal limbs, increased deep tendon reflexes and extensor plantar responses, which may be associated with mild intellectual disability. Additional features include muscle wasting and cerebellar abnormalities.
ORPHACODE:	401820
SYNONYMS:	SPG67
XREF(S):	Orphanet ICD-10
ANALYTE(S):	PGAP1
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

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