DISEASE:
Autosomal dominant spastic paraplegia type 13

NAME:	Autosomal dominant spastic paraplegia type 13
DESCRIPTION:	A rare, pure or complex form of hereditary spastic paraplegia characterized by progressive spastic paraplegia with pyramidal signs in the upper and lower limbs, and decreased vibration sense.
ORPHACODE:	100994
SYNONYMS:	SPG13
XREF(S):	Orphanet OMIM MeSH ICD-10
ANALYTE(S):	HSPD1
CREATED:	13 May 2019 - 01:02
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- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)

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