DISEASE:
Autosomal recessive spastic paraplegia type 9B

NAME:	Autosomal recessive spastic paraplegia type 9B
DESCRIPTION:	A rare complex hereditary spastic paraplegia characterized by early onset of slowly progressive spastic para- or tetraparesis, increased tendon reflexes, positive Babinski sign, global developmental delay, cognitive impairment, and pseudobulbar palsy. Additional manifestations include dysmorphic facial features, tremor, short stature, and urinary incontinence.
ORPHACODE:	447760
SYNONYMS:	AR-SPG9B
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	ALDH18A1
CREATED:	13 May 2019 - 01:02
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