Belgian Genetic Tests database

DISEASE:
Autosomal dominant spastic paraplegia type 9B

NAME:	Autosomal dominant spastic paraplegia type 9B
DESCRIPTION:	A rare predominantly pure hereditary spastic paraplegia characterized by juvenile or adult onset of slowly progressive spastic paraparesis, gait disturbances, and increased tendon reflexes. Additional variable manifestations include pes cavus, dysarthria, sensory impairment, and urinary symptoms. Cognition is normal.
ORPHACODE:	447757
SYNONYMS:	AD-SPG9B
XREF(S):	Orphanet ICD-10
ANALYTE(S):	ALDH18A1
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