Opis choroby *

Definicja

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.

Dane

Klasyfikacja
Choroba

Synonimy
AD-SPG9B
AD-SPG9B

Kod ORPHA
447757

Kod OMIM
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Kod ICD10
G11.4

Kod ICD11
8B44.00

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*Źródło

orphanet