Opis choroby *

Definicja

A pure form of hereditary spastic paraplegia characterized by a childhood- to adulthood-onset of slowly progressive spastic gait, extensor plantar responses, brisk tendon reflexes in arms and legs, decreased vibration sense at ankles and urinary dysfunction. Ankle clonus is also reported in some patients.

Dane

Klasyfikacja
Choroba

Synonimy
SPG37
SPG37

Kod ORPHA
171612

Kod OMIM
611945

Kod ICD10
G11.4

Kod ICD11
8B44.00

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*Źródło

orphanet