Opis choroby *

Definicja

A pure form of hereditary spastic paraplegia characterized by a slowly progressive and relatively benign spastic paraplegia presenting in adulthood with spastic gait, lower limb hyperreflexia, extensor plantar responses, bladder dysfunction (urinary urgency and/or incontinence), and mild sensory and motor peripheral neuropathy.

Dane

Klasyfikacja
Choroba

Synonimy
SPG19
SPG19

Kod ORPHA
100999

Kod OMIM
607152

Kod ICD10
G11.4

Kod ICD11
8B44.00

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*Źródło

orphanet