Opis choroby *

Definicja

A rare, pure or complex form of hereditary spastic paraplegia characterized by early adulthood onset of slowly progressive lower limb spasticity resulting in gait disturbances, hyperreflexia and extensor plantar responses, urinary urgency and/or incontinence, muscle weakness, decreased vibration sense and mild muscular atrophy in lower extremities. It may be associated with complicating signs, such as sensory neuropathy, ataxia (i.e. mild dysmetria, uncoordinated eye movement) and mild dysphagia.

Dane

Klasyfikacja
Choroba

Synonimy
SPG8
SPG8

Kod ORPHA
100989

Kod OMIM
603563

Kod ICD10
G11.4

Kod ICD11
8B44.00

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*Źródło

orphanet