Opis choroby *

Definicja

A rare form of hereditary spastic paraplegia characterized by delayed walking, toe walking, unsteady and spastic gait, hyperreflexia of the lower limbs, and extensor plantar responses. Upper limbs spasticity and dystonia, subclinical axonal neuropathy, cognitive impairment and intellectual disability have also been associated.

Dane

Klasyfikacja
Choroba

Synonimy
SPG56
SPG56

Kod ORPHA
320411

Kod OMIM
615030

Kod ICD10
G11.4

Kod ICD11
8B44.01

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*Źródło

orphanet