Opis choroby *

Definicja

A complex form of hereditary spastic paraplegia, characterized by an onset in childhood or adulthood of progressive spastic paraplegia (with spastic gait, spasticity, lower limb weakness, pes cavus and urinary urgency) associated with the additional manifestation of peripheral sensorimotor neuropathy.

Dane

Klasyfikacja
Choroba

Synonimy
SPG36
SPG36

Kod ORPHA
320365

Kod OMIM
613096

Kod ICD10
G11.4

Kod ICD11
8B44.00

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*Źródło

orphanet