Opis choroby *

Definicja

A pure form of hereditary spastic paraplegia characterized by onset in adolescence or early adulthood of slowly progressive spastic paraplegia, proximal muscle weakness of the lower extremities and small hand muscles, hyperreflexia, spastic gait and mild urinary compromise.

Dane

Klasyfikacja
Choroba

Synonimy
SPG41
SPG41

Kod ORPHA
320355

Kod OMIM
613364

Kod ICD10
G11.4

Kod ICD11
8B44.00

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*Źródło

orphanet