

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

*Źródło

orphanet