Opis choroby *

Definicja

A pure form of hereditary spastic paraplegia characterized by a childhood- to adulthood-onset of slowly progressive lower limb spasticity and hyperreflexia of lower extremities, extensor plantar reflexes, distal sensory impairment, variable urinary dysfunction and pes cavus.

Dane

Klasyfikacja
Choroba

Synonimy
SPG12
SPG12

Kod ORPHA
100993

Kod OMIM
604805

Kod ICD10
G11.4

Kod ICD11
8B44.00

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*Źródło

orphanet