

## 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