

## Opis choroby \*

### Definicja

A pure form of hereditary spastic paraplegia characterized by a childhood- to adulthood-onset of slowly progressive spastic gait, extensor plantar responses, brisk tendon reflexes in arms and legs, decreased vibration sense at ankles and urinary dysfunction. Ankle clonus is also reported in some patients.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

SPG37

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#### Kod ORPHA

171612

#### Kod OMIM

611945

#### Kod ICD10

G11.4

#### Kod ICD11

8B44.00

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#### \*Źródło

orphanet