

## Opis choroby \*

### Definicja

A complex form of hereditary spastic paraplegia, characterized by an onset in childhood or adulthood of progressive spastic paraplegia (with spastic gait, spasticity, lower limb weakness, pes cavus and urinary urgency) associated with the additional manifestation of peripheral sensorimotor neuropathy.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

SPG36

SPG36

#### Kod ORPHA

320365

#### Kod OMIM

613096

#### Kod ICD10

G11.4

#### Kod ICD11

8B44.00

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#### \*Źródło

orphanet