

## Opis choroby \*

### Definicja

A rare, pure or complex form of hereditary spastic paraplegia characterized by either a pure spastic paraplegia phenotype, usually presenting in the first or second decade of life, with spastic lower extremities, unsteady spastic gait, hyperreflexia and extensor plantar responses, or as a complicated phenotype with the additional manifestations of distal wasting, saccadic ocular movements, mild cerebellar ataxia and mild, distal, axonal neuropathy.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

SPG30

SPG30

#### Kod ORPHA

101010

#### Kod OMIM

610357

#### Kod ICD10

G11.4

#### Kod ICD11

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#### \*Źródło

orphanet