

## Opis choroby \*

### Definicja

Autosomal recessive spastic paraplegia type 59 is a very rare, complex hereditary spastic paraplegia characterized by an early onset of progressive lower limb spasticity, tip-toe walking, scissor gait, hyperreflexia and clonus that may be associated with borderline intellectual disability. Nystagmus and pes equinovarus have also been reported.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

SPG59

SPG59

#### Kod ORPHA

401795

#### Kod OMIM

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#### Kod ICD10

G11.4

#### Kod ICD11

8B44.01

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#### \*Źródło

orphanet