

## Opis choroby \*

### Definicja

Autosomal recessive spastic paraplegia type 32 (SPG32) is a rare, complex type of hereditary spastic paraplegia characterized by a slowly progressive spastic paraplegia (with walking difficulties appearing at onset at 6-7 years of age) associated with mild intellectual disability. Brain imaging reveals thin corpus callosum, cortical and cerebellar atrophy, and pontine dysraphia. The SPG32 phenotype has been mapped to a locus on chromosome 14q12-q21.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

SPG32

SPG32

#### Kod ORPHA

171622

#### Kod OMIM

611252

#### Kod ICD10

G11.4

#### Kod ICD11

8B44.01

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#### \*Źródło

orphanet