

## Opis choroby \*

### Definicja

Autosomal recessive spastic paraplegia type 25 (SPG25) is a rare, complex type of hereditary spastic paraplegia characterized by adult-onset spastic paraplegia associated with spinal pain that radiates to the upper or lower limbs and is related to disk herniation (with minor spondylosis), as well as mild sensorimotor neuropathy. The SPG25 phenotype has been mapped to a locus on chromosome 6q23-q24.1.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Autosomal recessive spastic paraplegia-disc  
herniation syndrome

Autosomalna recesywna paraplegia spastyczna -  
przepuklina dysku

SPG25

SPG25

#### Kod ORPHA

101005

#### Kod OMIM

608220

#### Kod ICD10

G11.4

#### Kod ICD11

8B44.01

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#### \*Źródło

orphanet