

## Opis choroby \*

### Definicja

Autosomal recessive spastic paraplegia type 23 (SPG23) is a rare, complex type of hereditary spastic paraplegia that presents in childhood with progressive spastic paraplegia, associated with peripheral neuropathy, skin pigment abnormalities (i.e. vitiligo, hyperpigmentation, diffuse lentiginosities), premature graying of hair, and characteristic facies (i.e. thin with "sharp" features). The SPG23 phenotype has been mapped to a locus on chromosome 1q24-q32.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Lison syndrome

Parapareza spastyczna - bielactwo nabyte -  
przedwczesne siwienie - charakterystyczna twarz

SPG23

Zespół Lisona

SPG23

Spastic paraparesis-vitiligo-premature graying-  
characteristic facies syndrome

#### Kod ORPHA

101003

#### Kod OMIM

270750

#### Kod ICD10

G11.4

#### Kod ICD11

8B44.01

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#### \*Źródło

orphanet