

## Opis choroby \*

### Definicja

A rare, pure or complex form of hereditary spastic paraplegia, with variable phenotype, typically characterized by childhood-onset of minimally progressive, bilateral, mainly symmetric lower limb spasticity and weakness, associated with *pes cavus*, scoliosis, sphincter disturbances and/or urinary bladder hyperactivity. Rare additional associated manifestations may include mild intellectual disability, axonal motor neuropathy, and seizures.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

Strümpell disease

Choroba Strümpella

#### Kod ORPHA

100984

#### Kod OMIM

182600

#### Kod ICD10

G11.4

#### Kod ICD11

8B44.00

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#### \*Źródło

orphanet