

## Opis choroby \*

### Definicja

A complex, hereditary, spastic paraplegia characterized by delayed motor development, spasticity, and inability to walk, later progressing to quadriplegia, motor aphasia, bowel and bladder dysfunction. Patients also present with vision problems and mild intellectual disability. The disease affects only males.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

SPG16

SPG16

#### Kod ORPHA

100997

#### Kod OMIM

300266

#### Kod ICD10

G11.4

#### Kod ICD11

8B44.02

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#### \*Źródło

orphanet