

# Autosomalna dominująca paraplegia spastyczna typu 4

**Kod Orpha: 100985 Kod OMIM: 182601**

## Opis choroby \*

### Definicja

A rare form of hereditary spastic paraplegia with high intrafamilial clinical variability, characterized in most cases as a pure phenotype with an adult onset (mainly the 3rd to 5th decade of life, but that can present at any age) of progressive gait impairment due to bilateral lower-limb spasticity and weakness as well as very mild proximal weakness and urinary urgency. In some cases, a complex phenotype is also reported with additional manifestations including cognitive impairment, cerebellar ataxia, epilepsy and neuropathy. A faster disease progression is noted in patients with a later age of onset.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

SPG4

SPG4

#### Kod ORPHA

100985

#### Kod OMIM

182601

#### Kod ICD10

G11.4

#### Kod ICD11

8B44.00

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[\\*Źródło](#)

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## Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.

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