

# Autosomalna dominująca paraplegia spastyczna typu 42

## Kod Orpha: 171863 Kod OMIM: 612539

### Opis choroby \*

#### Definicja

A pure form of hereditary spastic paraplegia characterized by slowly progressive spastic paraplegia of lower extremities with an age of onset ranging from childhood to adulthood and patients presenting with spastic gait, increased tendon reflexes in lower limbs, extensor plantar response, weakness and atrophy of lower limb muscles and, in rare cases, pes cavus. No abnormalities are noted on magnetic resonance imaging.

#### Dane

#### Klasyfikacja

Choroba

#### Synonimy

SPG42

SPG42

#### Kod ORPHA

171863

#### Kod OMIM

612539

#### Kod ICD10

G11.4

#### Kod ICD11

8B44.00

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[\\*Źródło](#)

orphanet

### Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.