

# Autosomalna recesywna paraplegia spastyczna typu 28

**Kod Orpha: 101008 Kod OMIM: 609340**

## Opis choroby \*

### Definicja

Autosomal recessive spastic paraplegia type 28 is a pure form of hereditary spastic paraplegia characterized by a childhood or adolescent onset of slowly progressive, pure crural muscle spastic paraparesis which manifests with mild lower limb weakness, gait difficulties, extensor plantar responses, and hyperreflexia of lower extremities. Less common manifestations include cerebellar oculomotor disturbance with saccadic eye pursuit, pes cavus and scoliosis. Some patients also present pin and vibration sensory loss in distal legs.

### Dane

**Klasyfikacja**  
Choroba

**Synonimy**  
SPG28  
SPG28

**Kod ORPHA**  
101008

**Kod OMIM**  
609340

**Kod ICD10**  
G11.4

**Kod ICD11**  
8B44.01

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[\\*Źródło](#)

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## Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.

Dostępna na stronie [www.orphanet.pl](http://www.orphanet.pl)