

Autosomalna dominująca paraplegia spastyczna typu 32

Kod Orpha: 171622 Kod OMIM: 611252

Opis choroby *

Definicja

Autosomal recessive spastic paraplegia type 32 (SPG32) is a rare, complex type of hereditary spastic paraplegia characterized by a slowly progressive spastic paraplegia (with walking difficulties appearing at onset at 6-7 years of age) associated with mild intellectual disability. Brain imaging reveals thin corpus callosum, cortical and cerebellar atrophy, and pontine dysraphia. The SPG32 phenotype has been mapped to a locus on chromosome 14q12-q21.

Dane

Klasyfikacja

Choroba

Synonimy

SPG32

SPG32

Kod ORPHA

171622

Kod OMIM

611252

Kod ICD10

G11.4

Kod ICD11

8B44.01

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.