Opis choroby *

Definicja

Autosomal recessive spastic paraplegia type 59 is a very rare, complex hereditary spastic paraplegia characterized by an early onset of progressive lower limb spasticity, tip-toe walking, scissor gait, hyperreflexia and clonus that may be associated with borderline intellectual disability. Nystagmus and pes equinovarus have also been reported.

Dane

Klasyfikacja
Choroba

Synonimy
SPG59
SPG59

Kod ORPHA
401795

Kod OMIM
-

Kod ICD10
G11.4

Kod ICD11
8B44.01

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*Źródło

orphanet