Opis choroby *

Definicja

Autosomal recessive spastic paraplegia type 25 (SPG25) is a rare, complex type of hereditary spastic paraplegia characterized by adult-onset spastic paraplegia associated with spinal pain that radiates to the upper or lower limbs and is related to disk herniation (with minor spondylosis), as well as mild sensorimotor neuropathy. The SPG25 phenotype has been mapped to a locus on chromosome 6q23-q24.1.

Dane

Klasyfikacja
Choroba

Synonimy
Autosomal recessive spastic paraplegia-disc herniation syndrome
Autosomalna recesywna paraplegia spastyczna - przepuklina dysku
SPG25
SPG25

Kod ORPHA
101005

Kod OMIM
608220

Kod ICD10
G11.4

Kod ICD11
8B44.01

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*Źródło

orphanet