Opis choroby *

Definicja

Autosomal recessive spastic paraplegia type 35 is a rare form of hereditary spastic paraplegia characterized by childhood (exceptionally adolescent) onset of a complex phenotype presenting with lower limb (followed by upper limb) spasticity with hyperreflexia and extensor plantar responses, with additional manifestations including progressive dysarthria, dystonia, mild cognitive decline, extrapyramidal features, optic atrophy and seizures. White matter abnormalities and brain iron accumulation have also been observed on brain magnetic resonance imaging.

Dane

Klasyfikacja
Choroba

Synonimy
SPG35
SPG35

Kod ORPHA
171629

Kod OMIM
612319

Kod ICD10
G11.4

Kod ICD11
8B44.01

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*Źródło

orphanet