Opis choroby *

Definicja

A pure form of hereditary spastic paraplegia characterized by adult onset of crural spastic paraparesis, hyperreflexia, extensor plantar responses, proximal muscle weakness, mild muscle atrophy, decreased vibration sensation at ankles, and mild urinary dysfunction. Foot deformities have been reported to eventually occur in some patients. No abnormalities are noted on brain magnetic resonance imaging and peripheral nerve conduction velocity studies.

Dane

Klasyfikacja
Choroba

Synonimy
SPG73
SPG73

Kod ORPHA
444099

Kod OMIM
616282

Kod ICD10
G11.4

Kod ICD11
8B44.00

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*Źródło

orphanet