

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

*Źródło

orphanet