Autosomalna dominująca paraplegia spastyczna typu 42
Kod Orpha: 171863 Kod OMIM: 612539

Opis choroby *

Definicja
A pure form of hereditary spastic paraplegia characterized by slowly progressive spastic paraplegia of lower extremities with an age of onset ranging from childhood to adulthood and patients presenting with spastic gait, increased tendon reflexes in lower limbs, extensor plantar response, weakness and atrophy of lower limb muscles and, in rare cases, pes cavus. No abnormalities are noted on magnetic resonance imaging.
Dane

Klasyfikacja
Choroba

Synonimy
SPG42
SPG42

Kod ORPHA
171863

Kod OMIM
612539

Kod ICD10
G11.4

Kod ICD11
8B44.00

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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.