Opis choroby *

Definicja

A pure or complex form of hereditary spastic paraplegia characterized by an onset in the first decade of life of spastic paraperesis (more prominent in lower than upper extremities) and unsteady gait, as well as increased deep tendon reflexes, amyotrophy, cerebellar ataxia, and flexion contractures of the knees, in some.

Dane

Klasyfikacja
Choroba

Synonimy
SPG62
SPG62

Kod ORPHA
401785

Kod OMIM
615681

Kod ICD10
G11.4

Kod ICD11
8B44.01

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*Źródło

orphanet