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

Kod OMIM

Kod ICD10

401785

615681

G11.4

Kod ICD11 8B44.01

*Źródło

orphanet