Opis choroby *

Definicja

A rare, complex type of hereditary spastic paraplegia characterized by early-onset progressive spastic paraplegia presenting in infancy, associated with optic atrophy, fixation nystagmus, polyneuropathy occurring in late childhood/early adolescence leading to severe motor disability and progressive joint contractures and scoliosis.

Dane

Klasyfikacja
Choroba

Synonimy
SPOAN
SPOAN

Kod ORPHA
320406

Kod OMIM
609541

Kod ICD10
G11.4

Kod ICD11
8B44.01

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*Źródło

orphanet