

Opis choroby *

Definicja

A complex hereditary spastic paraplegia characterized by mild to severe lower limb spasticity, hyperreflexia, extensor plantar responses, impaired vibration sensation, *pes cavus*, and significant wasting and weakness of the small hand muscles. Temporal lobe epilepsy and cognitive dysfunction have been also reported.

Dane

Klasyfikacja

Choroba

Synonimy

SPG38

SPG38

Kod ORPHA

171617

Kod OMIM

612335

Kod ICD10

G11.4

Kod ICD11

8B44.00

*Źródło

orphanet