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

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*Źródło

orphanet