Opis choroby *

Definicja

A rare, pure or complex form of hereditary spastic paraplegia characterized by progressive spastic paraplegia with pyramidal signs in the upper and lower limbs, and decreased vibration sense.

Dane

Klasyfikacja
Choroba

Synonimy
SPG13
SPG13

Kod ORPHA
100994

Kod OMIM
605280

Kod ICD10
G11.4

Kod ICD11
8B44.00

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*Źródło

orphanet