Opis choroby *

Definicja

A rare, axonal hereditary motor and sensory neuropathy characterized by adult onset of slowly progressive distal muscle weakness and atrophy, decreased deep tendon reflexes of lower limbs, and mild distal sensory loss leading to gait difficulties in most patients.

Dane

Klasyfikacja
Choroba

Synonimy
CMT2 due to TFG mutation
CMT2 spowodowana mutacją TFG

Kod ORPHA
435819

Kod OMIM
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Kod ICD10
G60.0

Kod ICD11
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*Źródło

orphanet