

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	Synonimy
Choroba	CMT2 due to TFG mutation CMT2 spowodowana mutacją TFG

Kod ORPHA	Kod OMIM	Kod ICD10
435819	-	G60.0

Kod ICD11

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*Źródło

orphanet