Autosomalna dominująca choroba Charcota, Mariego i Tootha typu 2G
Kod Orpha: 99941 Kod OMIM: 614436

Opis choroby *

Definicja
A form of axonal Charcot-Marie-Tooth disease, a peripheral sensorimotor neuropathy with onset associated to development of foot deformity and walking difficulties between the 1st and the 8th decades, with a median range in the 2nd one. Weakness and sensory loss involve primarily the legs and ankles tendon reflexes are reduced. This disorder has a slowly progressive course.
Dane

Klasyfikacja
Choroba

Synonimy
CMT2G
CMT2G

Kod ORPHA
99941

Kod OMIM
614436

Kod ICD10
G60.0

Kod ICD11
8C20.1

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*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.