

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

*[Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.