Opis choroby *

Definicja

A form of axonal Charcot-Marie-Tooth disease, a peripheral sensorimotor neuropathy, characterized by distal weakness primarily and predominantly occurring in the upper limbs and tendon reflexes absent or reduced in the arms and decreased in the legs. Progression is slow.

Dane

Klasyfikacja
Choroba

Synonimy
CMT2D
CMT2D

Kod ORPHA
99938

Kod OMIM
601472

Kod ICD10
G60.0

Kod ICD11
8C20.1

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*Źródło

orphanet