

## Opis choroby \*

### Definicja

A form of axonal Charcot-Marie-Tooth disease, a peripheral sensorimotor neuropathy, characterized by a relatively late onset, pupillary abnormalities and deafness, in most patients, associated with distal weakness and muscle atrophy.

### Dane

#### Klasyfikacja

Choroba

#### Synonimy

CMT2J

CMT2J

#### Kod ORPHA

99943

#### Kod OMIM

607736

#### Kod ICD10

G60.0

#### Kod ICD11

8C20.1

---

#### \*Źródło

orphanet