Opis choroby *

Definicja

A rare group of inherited neuromuscular disorders characterized by clinical features of a congenital myopathy and centrally placed nuclei on muscle biopsy. The clinical picture and other histologic features varies according to gene involved and mode of inheritance.

Dane

Klasyfikacja
Grupa fenomenów

Synonimy
CNM
CNM

Kod ORPHA
595

Kod OMIM
-

Kod ICD10
G71.2

Kod ICD11
8C72.01

---

*Źródło

orphanet