

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