

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	Synonimy
Grupa fenomenów	CNM CNM

Kod ORPHA	Kod OMIM	Kod ICD10
595	-	G71.2

Kod ICD11
8C72.01

*Źródło

orphanet