

Miopatia Brody'ego

Kod Orpha: 53347 Kod OMIM: 601003

Opis choroby *

Definicja

A rare genetic skeletal muscle disease characterized by childhood onset of exercise-induced progressive impairment of muscle relaxation, stiffness, cramps, and myalgia, predominantly in the arms, legs, and face (eyelids), and, biochemically, by a reduced sarcoplasmic reticulum Ca(2+)-ATPase activity. Symptoms improve after a few minutes of rest and may be exacerbated by cold. The term Brody syndrome refers to a clinically distinguishable subset of patients without *ATP2A1* mutations, with adolescence or adult onset and selective muscular involvement, in which myalgia is more common.

Dane

Klasyfikacja

Choroba

Kod ORPHA
53347

Kod OMIM
601003

Kod ICD10
G71.8

Kod ICD11
8C7Y

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.