

Autosomalna dominująca mioglobinuria

Kod Orpha: 99846 Kod OMIM: 160010

Opis choroby *

Definicja

A rare metabolic myopathy characterized by episodic myalgia with myoglobinuria which is induced by fever, viral or bacterial infection, prolonged exercise or alcohol abuse, and could, on occasion, lead to acute renal failure. Between episodes, patients may be asymptomatic or could present elevated creatine kinase levels and mild muscle weakness. There have been no further descriptions in the literature since 1997.

Dane

Klasyfikacja

Choroba

Kod ORPHA

99846

Kod OMIM

160010

Kod ICD10

R82.1

Kod ICD11

5C5Y

[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.