Opis choroby *

Definicja

A rare form of idiopathic inflammatory myopathy characterized by acute or subacute, severe, symmetrical, proximal muscle weakness usually associated with muscle-specific antibodies (anti-HMGCR or anti-SRP). Histopathological characteristics include myocyte necrosis and regeneration without significant inflammation, and C5b-9 deposition on non-necrotic myofibers.

Dane

Klasyfikacja
Choroba

Synonimy
Anti-HMG-CoA myopathy
NAM
IMNM
Miopatia anty-SRP
Miopatia anty-HMG-CoA
Autoimmunologiczne martwicze zapalenie mięsni
Miopatia immunologiczna z martwicą miocytów
Anti-SRP myopathy
Autoimmune necrotizing myositis
IMNM
Immune myopathy with myocyte necrosis
NAM

Kod ORPHA
206569

Kod OMIM
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Kod ICD10
G72.4

Kod ICD11
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*Źródło

orphanet