Opis choroby *

Definicja

A rare idiopathic inflammatory myopathy characterized by diffuse destructive infiltration of CD68+ macrophages into the fascia rather than muscle fibers in muscle biopsies, proximal muscle weakness and myalgia with or without scaly dermatomyositis-like or atypical non-dermatomyositis-like skin lesions, elevation of creatine kinase levels and thickening of muscle fascia in muscle MRI.

Dane

Klasyfikacja
Choroba

Synonimy
IMAM
IMAM

Kod ORPHA
247718

Kod OMIM
-

Kod ICD10
G72.4

Kod ICD11
-

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*Źródło

orphanet