

Opis choroby *

Definicja

A rare idiopathic inflammatory myopathy (IIM) characterized by evocative skin lesions, muscle involvement with symmetrical proximal muscle weakness, and specific histological features. The clinical subtypes are defined by the presence of myositis-specific antibodies (anti-Mi2, anti-NXP2, anti-TIF1- γ , anti-MDA5, or anti-SAE antibodies) and are associated with specific clinical phenotypes and prognosis.

Dane

Klasyfikacja

Choroba

Synonimy

Adult dermatomyositis

DM

Kod ORPHA

221

Kod OMIM

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Kod ICD10

M33.0

Kod ICD11

4A41.00

*Źródło

orphanet