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
-

Kod ICD10
M33.0

Kod ICD11
4A41.00

---

*Źródło

orphanet