Opis choroby *

Definicja

A rare acquired skin disease characterized by benign proliferation of mature plasma cells with a typical triad of cutaneous lesions, polyclonal hypergammaglobulinemia, and superficial lymphadenopathy, without an apparent underlying cause. The skin lesions consist of multiple round-to-oval, red-to-dark-brown macules, papules, and plaques most commonly found on the trunk, but also the face, neck, and axillae.

Dane

Klasyfikacja
Choroba

Kod ORPHA
451602

Kod OMIM
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Kod ICD10
L98.6

Kod ICD11
EK91.2

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*Źródło

orphanet