## 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