Opis choroby *

Definicja

A rare non-Langerhans cell histiocytosis characterized by multiple small yellowish-red or brown papules initially erupting predominantly in the head and neck region. The histopathological hallmark of these eventually self-healing lesions is a dermal proliferation of histiocytes with intracytoplasmic comma-shaped bodies, coated vesicles, and desmosome-like structures. Birbeck granules are absent. The disease typically occurs in young children.

Dane

Klasyfikacja
Choroba

Kod ORPHA
157997

Kod OMIM
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Kod ICD10
D76.3

Kod ICD11
EE81

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*Źródło

orphanet