

## 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