Opis choroby *

Definicja

A rare genetic skin disease characterized by childhood onset of follicular keratotic papules slowly progressing to characteristic ''honeycomb'' atrophy on the cheeks, preauricular area, and forehead. Less frequently, the condition may affect also the upper lip, ears, or limbs. Additional features include facial erythema, milia, and follicular plugs.

Dane

Klasyfikacja
Choroba

Synonimy
Folliculitis ulerythematosa reticulate
Rogowacenie zapalne mieszków włosowych siatkowate

Kod ORPHA
79100

Kod OMIM
604093

Kod ICD10
L66.4

Kod ICD11
ED56

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*Źródło

orphanet