Opis choroby *

Definicja

A rare, indolent primary cutaneous B-cell lymphoma characterized by a solitary or grouped erythematous plaques or tumors, preferentially located on the head, neck or trunk region, and composed of centroblasts and centrocytes arranged in a follicular, diffuse, or mixed growth pattern. The lesions are smooth and typically do not ulcerate. The neoplastic cells express pan B cell markers and Bcl-6, and typically lack Bcl-2.

Dane

Klasyfikacja
Choroba

Synonimy
PCFCL
PCFCL

Kod ORPHA
178540

Kod OMIM
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Kod ICD10
C82.6

Kod ICD11
2A80.3

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*Źródło

orphanet