Opis choroby *

Definicja

A rare subtype of pyoderma gangrenosum disease characterized by a solitary, erythematous, ulcerated plaque, which lacks the violaceous border typically present in classic pyoderma gangrenosum, usually affecting individuals who are otherwise healthy. Histologically, the lesion presents a central layer containing neutrophilic inflamation, surrounded by a palisade of histiocytes, which are rimmed by a lymphocytic infiltrate. In comparison with the other variants of pyoderma gangrenosum, this subtype usually shows a good response to less aggressive treatments and underlying systemic disorders are less frequently associated. It is considered the most benign and uncommon clinical variant of pyoderma gangrenosum.

Dane

Klasyfikacja
Podtyp kliniczny

Synonimy
Granulomatous pyoderma gangrenosum
Granulomatous pyoderma gangrenosum

Kod ORPHA
538872

Kod OMIM
-

Kod ICD10
L88

Kod ICD11
EB21

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*Źródło

orphanet