

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 inflammation, 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	Synonimy
Podtyp kliniczny	Granulomatous pyoderma gangrenosum Granulomatous pyoderma gangrenosum

Kod ORPHA 538872	Kod OMIM -	Kod ICD10 L88
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Kod ICD11
EB21

*Źródło

orphanet