

## 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

<b>Klasyfikacja</b>	<b>Synonimy</b>
Podtyp kliniczny	Granulomatous pyoderma gangrenosum Granulomatous pyoderma gangrenosum

<b>Kod ORPHA</b> 538872	<b>Kod OMIM</b> -	<b>Kod ICD10</b> L88
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**Kod ICD11**  
EB21

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### \*Źródło

orphanet