Opis choroby *

Definicja

A rare subtype of pyoderma gangrenosum disease characterized by grouped vesicles that rapidly spread and coalesce to form large bullae, which evolve into ulcerations that have an erythematous peripheral halo and central necrosis, mainly affecting the upper limbs and face. Lymphoproliferative diseases are frequently associated, thus prognosis is often compromised.

Dane

Klasyfikacja
Podtyp kliniczny

Synonimy
Phemphigoid pyoderma gangrenosum
Phemphigoid pyoderma gangrenosum

Kod ORPHA
538869

Kod OMIM
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Kod ICD10
L88

Kod ICD11
EB21

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*Źródło

orphanet