

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

Pemphigoid pyoderma gangrenosum

Pemphigoid pyoderma gangrenosum

Kod ORPHA

538869

Kod OMIM

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Kod ICD10

L88

Kod ICD11

EB21

*Źródło

orphanet