Opis choroby *

Definicja

A rare subtype of pyoderma gangrenosum disease characterized by rapidly progressive, single or multiple, painful, aseptic ulcers which present overhanging, violaceous and undermined borders, surrounding induration and erythema, and granulation tissue (occasionally necrotic tissue and/or a purulent exudate) at the base, mainly affecting the legs (but other body surfaces may also be involved), leading to chronic ulcerations and often regressing with cribriform mutilating scars. The disease presents a chronic relapsing course and systemic features (e.g. fever, malaise, arthralgia, myalgia) may be associated.

Dane

Klasyfikacja
Podtyp kliniczny

Synonimy
Ulcerative pyoderma gangrenosum
Ulcerative pyoderma gangrenosum

Kod ORPHA
538863

Kod OMIM
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Kod ICD10
L88

Kod ICD11
EB21

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*Źródło

orphanet