

Opis choroby *

Definicja

A rare autoimmune bullous skin disease characterized by acquired, subepidermal tense bullae occurring on normal or inflamed skin and that is typically widespread (occurring in the flexor regions of the proximal arms and legs, in the armpits, groin and the abdomen) and often associated with pruritus. The evolution is typically chronic with spontaneous exacerbations and remission.

Dane

Klasyfikacja

Choroba

Kod ORPHA

703

Kod OMIM

-

Kod ICD10

L12.0

Kod ICD11

EB41.0

[*Źródło](#)

orphanet