

Opis choroby *

Definicja

A rare intermediate form of junctional epidermolysis bullosa characterized by congenital blistering and erosions confined to intertriginous skin sites, the esophagus, groin, and perineum. Blistering is usually severe and lesions may heal with atrophic scarring and milia formation. Extracutaneous manifestations include nail dystrophy, enamel hypoplasia and dental caries, oral, esophageal and vaginal blisters and erosions.

Dane

Klasyfikacja

Choroba

Synonimy

JEB inversa

JEB-I

JEB inversa

JEB-I

Kod ORPHA

79405

Kod OMIM

226650

Kod ICD10

Q81.8

Kod ICD11

EC31

[*Źródło](#)

orphanet