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