Opis choroby *

Definicja

A rare epidermolysis bullosa simplex characterized by the association of the typical trauma-induced blisters with additional features including hearing impairment, alopecia, hypo- or anodontia, and nail dystrophy. Occurrence of vitiliginous skin areas unrelated to the sites of the blisters has also been described.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
EBS with anodontia/hypodontia
Zespół Gamborga i Nielsena
Zespół Kallina
Kallin syndrome

Kod ORPHA
2325

Kod OMIM
-

Kod ICD10
Q81.0

Kod ICD11
-

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*Źródło

orphanet