Opis choroby *

Definicja

A rare skin disorder characterized by localized absence of skin that is usually located on the scalp but can occur anywhere on the body including the face, trunk and extremities. Aplasia cutis congenita (ACC) may occasionally be associated with other anomalies.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
1114

Kod OMIM
107600

Kod ICD10
Q84.8

Kod ICD11
LC60

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*Źródło

orphanet