Opis choroby *

Definicja

A rare genetic skin disease characterized by infantile onset of diffuse alopecia, abnormal skin pigmentation (hypo- and hyperpigmented macules of the trunk and face and areas of reticular hypo- and hyperpigmentation of the extremities), palmoplantar keratoderma, and nail dystrophy. Patients develop recurrent spinocellular carcinomas later in life. Brittle teeth resulting in early loss of dentition have also been described.

Dane

Klasyfikacja
Choroba

Kod ORPHA
447961

Kod OMIM
618373

Kod ICD10
D04.8

Kod ICD11
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*Źródło

orphanet