Opis choroby *

Definicja

A rare ectodermal dysplasia syndrome characterized by neonatal teeth, hypo- or oligodontia of the secondary dentition, flexural acanthosis nigricans, and sparse body and scalp hair (the latter being thin and slow-growing). There have been no further descriptions in the literature since 1995.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
69083

Kod OMIM
601345

Kod ICD10
Q82.4

Kod ICD11
LD27.0Y

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*Źródło

orphanet