

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

*Źródło

orphanet