Opis choroby *

Definicja

A rare ectodermal dysplasia syndrome characterized by the association of sparse, fine, dry, slow growing hair with variable dental abnormalities including oligodontia, peg-shaped incisors, and shell teeth. Mild intellectual disability, microcephaly, and dysmorphic facial features have also been reported.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
Kersey syndrome
Zespół Kersey'a

Kod ORPHA
3351

Kod OMIM
601453

Kod ICD10
Q82.4

Kod ICD11
LD27.0Y

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*Źródło

orphanet