Opis choroby *

Definicja

A syndromic genetic deafness characterized by erythrokeratoderma, hypotrichosis, nail dystrophy and sensorineural hearing loss. Erythema, recurrent skin infections and mucositis have also been associated.

Dane

Klasyfikacja
Choroba

Synonimy
Hypotrichosis-hearing loss syndrome
Hypotrichosis-hearing loss syndrome

Kod ORPHA
330029

Kod OMIM
-

Kod ICD10
H90.5

Kod ICD11
-

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*Źródło

orphanet