Opis choroby *

Definicja

A rare syndrome characterized by gingival fibromatosis associated with progressive sensorineural hearing loss. It has been described in two families (with at least 16 affected members spanning five generations in one of the families, and five affected members spanning three generations in the other family). It is transmitted as an autosomal dominant trait.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
Gingival fibromatosis-progressive hearing loss syndrome
Zespół Jonesa
Jones syndrome

Kod ORPHA
2027

Kod OMIM
135550

Kod ICD10
H90.3

Kod ICD11
LD2H.Y

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*Źródło

orphanet