Opis choroby *

Definicja

Oculodental syndrome, Rutherfurd type is a rare genetic disorder that is primarily characterized by the classical triad of gingival fibromatosis, non-eruption of tooth and corneal dystrophy (bilateral corneal vascularization and opacity). Abnormally shaped teeth have also been reported. The syndrome is transmitted as an autosomal dominant trait.

Dane

Klasyfikacja
Zespół wad wrodzonych

Synonimy
Gingival hypertrophy-corneal dystrophy
Przerost dziąseł - dystrofia rogówki
Zespół Rutherfurda
Rutherfurd syndrome

Kod ORPHA
2709

Kod OMIM
180900

Kod ICD10
Q87.8

Kod ICD11
9A70.Y

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*Źródło

orphanet