

Opis choroby *

Definicja

Oculodental syndrome, Rutherford 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	Synonimy
Zespół wad wrodzonych	Gingival hypertrophy-corneal dystrophy Przerost dziąseł - dystrofia rogówki Zespół Rutherforda Rutherford syndrome

Kod ORPHA 2709	Kod OMIM 180900	Kod ICD10 Q87.8
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Kod ICD11
9A70.Y

*Źródło

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