Oculodental syndrome, Rutherfurd type

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Oculodental syndrome, Rutherfurd type. ORPHA:2709

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.