Opis choroby *

Definicja

A rare, genetic, otorhinolaryngological malformation characterized by congenital impatency of the nasolacrimal draingage system in various members of a family. Presentation is not specific and may include a uni- or bilateral medial canthal mass, dacryocystitis, nasal obstruction, periorbital cellulitis, and epiphora. Dacryocystocele and lacrimal puncta agenesis may be associated.

Dane

Klasyfikacja
Wada morfologiczna

Kod ORPHA
451612

Kod OMIM
149700

Kod ICD10
Q10.5

Kod ICD11
LA14.14

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*Źródło

orphanet