

Opis choroby *

Definicja

A rare, genetic, otorhinolaryngological malformation characterized by congenital impatency of the nasolacrimal drainage 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

*Źródło

orphanet