Opis choroby *

Definicja

A rare autosomal dominant disorder characterized by aplasia, atresia or hypoplasia of the lacrimal and salivary glands leading to varying features since infancy such as recurrent eye infections, irritable eyes, epiphora, xerostomia, dental caries, dental erosion and oral inflammation.

Dane

Klasyfikacja
Choroba

Synonimy
ALSG
ALSG
Wrodzony brak punktów łzowych i gruczołów ślinowych
Congenital absence of lacrimal puncta and salivary glands

Kod ORPHA
86815

Kod OMIM
180920

Kod ICD10
Q38.4

Kod ICD11
LA14.10

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*Źródło

orphanet