

## 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

---

#### \*Źródło

orphanet