Opis choroby *

Definicja

A rare oblique facial cleft characterized by a congenital unilateral or bilateral defect beginning in the upper lip medial to the oral commissure and extending across the cheek as a groove ending between the middle and lateral third of the lower eyelid (resulting in coloboma). Bone involvement includes an alveolar cleft in the premolar region, extending across the maxilla lateral to the infraorbital nerve and up to the infraorbital rim and orbital floor. The malformation may be associated with Tessier number 3 and number 4 clefts, macrostomia, or anophthalmos.

Dane

Klasyfikacja
Wada morfologiczna

Kod ORPHA
141261

Kod OMIM
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Kod ICD10
Q18.8

Kod ICD11
LA51

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*Źródło

orphanet