Opis choroby *

Definicja

A rare developmental defect during embryogenesis characterized by muscular hypertrophy, adenoid hyperplasia, or vascular malformation that results in an enlarged, often protruding, tongue. Complications include difficulty in swallowing, breathing and mastication, drooling, dental and skeletal deformities, such as malocclusion, open bite, asymmetry in maxillary and mandibular arches. It may be isolated or associated with genetic syndromes.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
2430

Kod OMIM
153630

Kod ICD10
Q38.2

Kod ICD11
LA31.0

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*Źródło

orphanet