

## 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	Kod OMIM	Kod ICD10
2430	153630	Q38.2
Kod ICD11		
LA31.0		

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### \*Źródło

orphanet