## **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

<u>\*Źródło</u>

orphanet