

## Opis choroby \*

### Definicja

A rare head and neck malformation characterized by congenital partial (hypoglossia) or total (aglossia) absence of the tongue. Patients present feeding and respiratory difficulties, as well as delayed speech development and slurred speech. Taste perception is not severely compromised. Associated features include a characteristic facies due to mandibular transverse arch deficiency, oligodontia, and malocclusion, among others.

### Dane

### Klasyfikacja

Wada morfologiczna

#### Kod ORPHA

141152

#### Kod OMIM

612776

#### Kod ICD10

Q38.3

#### Kod ICD11

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### \*Źródło

orphanet