Izolowany wrodzony niedorozwój języka/brak języka
Kod Orpha: 141152 Kod OMIM: 612776

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
-

*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.