

## Opis choroby \*

### Definicja

A rare genetic, orofacial clefting syndrome characterized by the association of bilateral microtia with severe to profound hearing impairment, and cleft palate.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych	Bilateral microtia-hearing loss-cleft palate syndrome

**Kod ORPHA**  
140963

**Kod OMIM**  
612290

**Kod ICD10**  
Q87.0

**Kod ICD11**  
LD2H.Y

---

### \*Źródło

orphanet