

## Opis choroby \*

### Definicja

Hemifacial hyperplasia is a rare morphological anomaly of the maxillofacial region characterized by unilateral overgrowth of all facial structures (bone, soft tissues, teeth), called true hemifacial hypertrophy, or overgrowth of one or more but not all facial structures, called partial hemifacial hypertrophy. It may be isolated or related to some syndromes (e.g. Beckwith-Wiedemann, Proteus, Klippel-Trenaunay-Weber, McCune-Albright syndrome, Neurofibromatosis type 1). It may be associated with airway obstruction, sensorineural hearing loss or swallowing difficulties.

### Dane

<b>Klasyfikacja</b>	<b>Synonimy</b>
Zespół wad wrodzonych	Hemifacial hypertrophy

<b>Kod ORPHA</b>	<b>Kod OMIM</b>	<b>Kod ICD10</b>
141145	133900	Q67.4

**Kod ICD11**  
LA52

---

### \*Źródło

orphanet