## Opis choroby \*

Definicja

A rare congenital laryngeal anomaly characterized by an abnormal dilation of the laryngeal saccule that is filled with air, maintains communication with the laryngeal lumen, and is either confined to the false vocal fold or extends upward, protruding through the thyrohyoid membrane to the neck. Symptoms may include cough, hoarseness, stridor, sore throat and uni- or bilateral swelling of the neck. Blockage of the laryngocele neck can result isn laryngomucocele, and forms laryngopyocele when infected.

Dane

## Klasyfikacja

Zespół wad wrodzonych

 Kod ORPHA
 Kod OMIM
 Kod ICD10

 2372
 Q31.3

Kod ICD11 LA71.1

\*Źródło

orphanet