Opis choroby *

Definicja

A rare larynx anomaly characterized by a partial or complete narrowing of the upper airway extending from just below the vocal folds to the lower border of the cricoid cartilage. Clinical presentation is variable and includes recurrent, croup-like, upper respiratory infections, stridor, dyspnea, barking cough, and in most severe cases acute airway compromise at delivery. It may be an isolated finding, or associated with other congenital anomalies and syndromes.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
141121

Kod OMIM
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Kod ICD10
Q31.1

Kod ICD11
LA71.3

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*Źródło

orphanet