Opis choroby *

Definicja

A rare larynx anomaly characterized by complete absence of the laryngeal lumen resulting in congenital upper airway obstruction syndrome which, without fetal or neonatal intervention, is incompatible with life. Fetal sonography shows a dilated trachea, hyperechoic lungs, pleural effusion, minimal fetal abdominal ascites or hydrops, and amniotic fluid abnormalities.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
1202

Kod OMIM
150300

Kod ICD10
Q31.8

Kod ICD11
LA71.Y

---

*Źródło

orphanet