Opis choroby *

Definicja

A rare, congenital, esophageal malformation characterized by the presence of an abnormal connection between the esophagus and the trachea (typically occurring in the lower cervical or upper thoracic area and taking an oblique path upward to trachea), without concomitant esophageal atresia. Depending on the size of the lumen, presentation varies from neonatal episodes of choking and cyanosis on feeding to subtle symptoms of wheezing and recurrent respiratory infections in childhood or early adulthood.

Dane

Klasyfikacja
Wada morfologiczna

Synonimy
H-type tracheoesophageal fistula
H-type tracheoesophageal fistula

Kod ORPHA
454750

Kod OMIM
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Kod ICD10
Q39.2

Kod ICD11
LB12.2

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*Źródło

orphanet