Opis choroby *

Definicja

A rare congenital malformation characterized by an interruption in the continuity of the esophagus, with or without persistent communication with the trachea. The clinical presentation varies according to the anatomy, and can lead to the inability to swallow or, in the most severe cases, respiratory distress.

Dane

Klasyfikacja
Wada morfologiczna

Synonimy
CEA
Congenital esophageal atresia
EA/TEF
Esophageal atresia with or without trachea-esophageal fistula
Oesophageal atresia

Kod ORPHA
1199

Kod OMIM
189960

Kod ICD10
Q39.1

Kod ICD11
LB12.1

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*Źródło

orphanet