

## 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