

## Opis choroby \*

### Definicja

A rare multiple congenital anomalies/dysmorphic syndrome characterized by the association of omphalocele and cleft palate. Other reported features include cleft lip, bifid uvula, bilateral talipes equinovarus, bicornuate uterus, and hydrocephalus internus. The condition is lethal in infancy.

### Dane

#### Klasyfikacja

Zespół wad wrodzonych  
Czeizel syndrome  
Czeizel syndrome

#### Synonimy

#### Kod ORPHA

2736

#### Kod OMIM

258320

#### Kod ICD10

Q87.8

#### Kod ICD11

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#### \*Źródło

orphanet