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

Synonimy
Czeizel syndrome
Czeizel syndrome

Kod ORPHA
2736

Kod OMIM
258320

Kod ICD10
Q87.8

Kod ICD11
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*Źródło

orphanet