Opis choroby *

Definicja

A rare syndromic intestinal malformation characterized by ulcer formation in the umbilical cord associated with congenital upper-intestinal atresia, typically presenting with intra-uterine hemorrhaging from the ulcer site and subsequent fetal bradycardia.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
3405

Kod OMIM
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Kod ICD10
Q43.8

Kod ICD11
LD2F.1Y

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*Źródło

orphanet