Opis choroby *

Definicja

A rare genetic multiple congenital anomalies/dysmorphic syndrome characterized by congenital diaphragmatic hernia, short bowel, and asplenia. Dysmorphic facial features include long forehead, hypertelorism, upturned nares, and small mandible. Atresia of the duodenum has also been reported.

Dane

Klasyfikacja
Zespół wad wrodzonych

Kod ORPHA
527468

Kod OMIM
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Kod ICD10
Q87.8

Kod ICD11
-

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*Źródło

orphanet