

## 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	Kod OMIM	Kod ICD10
527468	-	Q87.8
Kod ICD11		
-		

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### \*Źródło

orphanet