Opis choroby *

Definicja

A rare biliary tract disease characterized by congenital absence of the gallbladder and cystic duct. The majority of patients are asymptomatic. Possible clinical manifestations include abdominal pain and tenderness in the right upper quadrant, nausea, vomiting, fatty food intolerance, and jaundice. Frequency of choledocholithiasis is increased significantly.

Dane

Klasyfikacja
Wada morfologiczna

Kod ORPHA
440987

Kod OMIM
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Kod ICD10
Q44.0

Kod ICD11
LB20.10

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*Źródło

orphanet