## Opis choroby *

Definicja

A very rare hepatic and biliary tract tumor characterized by a growth pattern ressembling that found in hepatocellular carcinomas and cholangiocarcinomas but presenting atypical histological and immunohistochemical features (such as trabecular, organoid, microcystic and/or blastemal-like architecture and inhibin A, cytokeratin 7 and/or cytokeratin 19 positivity) that do not allow a formal diagnosis of the more common aforementioned liver cancers. Patients may present abdominal distension and pain, a palpable abdominal mass and elevated liver enzymes.

Dane

**Klasyfikacja**
Choroba

Synonimy
Adenocarcinoma of the liver and IBT
Gruczolakorak wątroby i IBT

**Kod ORPHA**
424943

**Kod OMIM**
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**Kod ICD10**
C22.0

**Kod ICD11**
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*Źródło

orphanet