

## Opis choroby \*

### Definicja

A rare genetic parenchymatous liver disease characterized by infantile or early childhood onset of recurrent episodes of acute liver failure precipitated by a febrile illness. During the life-threatening episodes, patients present with vomiting, lethargy, jaundice, as well as elevated levels of liver enzymes and coagulopathy. There is usually complete recovery between the episodes with conservative treatment.

### Dane

### Klasyfikacja

Choroba

#### Kod ORPHA

464724

#### Kod OMIM

618641

#### Kod ICD10

K72.0

#### Kod ICD11

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### \*Źródło

orphanet