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