

Przejęciowa rodzinna hiperbilirubinemia noworodków

Kod Orpha: 2312 Kod OMIM: 237900

Opis choroby *

Definicja

A rare genetic hepatic disease characterized by very high serum bilirubin levels in a newborn, clinically presenting as jaundice during the first few days of life. The condition is usually self-resolving, although in some cases it can lead to kernicterus with corresponding symptoms (including lethargy, high-pitched crying, hypotonia, missing reflexes, vomiting, or seizures, among others), which may result in chronic disability and even death.

Dane

Klasyfikacja

Choroba

Synonimy

Lucey-Driscoll syndrome

Zespół Lucey i Driscoll

Kod ORPHA

2312

Kod OMIM

237900

Kod ICD10

P59.8

Kod ICD11

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[*Źródło](#)

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.