Przejściowa 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
-

*Źródło

orphanet

Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.