Opis choroby *

Definicja

A rare genetic disease characterized by congenital cataract, neonatal hepatic failure and cholestatic jaundice, and global developmental delay. Neonatal death due to progressive liver failure has been reported.

Dane

Klasyfikacja
Choroba

Kod ORPHA
521432

Kod OMIM
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Kod ICD10
K83.1

Kod ICD11
-

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*Źródło

orphanet