

Opis choroby *

Definicja

A rare hematologic disease characterized by the transfer of maternal alloantibodies against red blood cell antigens of the Kell family to a fetus positive for this antigen across the placental barrier, causing suppression of erythropoiesis with reticulocytopenia and anemia, as well as alloimmune hemolysis. Severe anemia may lead to hydrops fetalis. Significant hyperbilirubinemia is rare in this condition.

Dane

Klasyfikacja

Choroba

Synonimy

Anti-K HDN

Matczyna alloimmunizacja anty-Kell

Maternal anti-Kell alloimmunization

Kod ORPHA

275944

Kod OMIM

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Kod ICD10

P55.8

Kod ICD11

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*Źródło

orphanet