## 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