# Wrodzony Niedobór czynnika XII
# Kod Orpha: 330 Kod OMIM: 234000

## Opis choroby *

**Definicja**
A rare, autosomal recessive systemic dysfunction of the hemostatic pathway, that is due to a defect in the coagulation factor XII (FXII or Hageman factor), and is either asymptomatic or characterized by a prolonged activated partial thromboplastin time and an increased risk for thromboembolism. FXII deficiency is strongly associated with primary recurrent abortions.
**Dane**

**Klasyfikacja**
Choroba

**Synonimy**
Congenital Hageman factor deficiency
Wrodzony Niedobór czynnika Hagemana

**Kod ORPHA**
330

**Kod OMIM**
234000

**Kod ICD10**
D68.2

**Kod ICD11**
3B15

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*Źródło

orphanet

## Rozszerzony opis choroby

Brak opisu rozszerzonego dla tej choroby. Opracowanie w toku.