DISEASE:
Congenital factor II deficiency

NAME:	Congenital factor II deficiency
DESCRIPTION:	A rare inherited bleeding disorder due to reduced activity of factor II (FII, prothrombin) and characterized by mucocutaneous and soft tissue bleeding symptoms.
ORPHACODE:	325
SYNONYMS:	Dysprothrombinemia Hypoprothrombinemia Prothrombin deficiency
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	F2
CREATED:	13 May 2019 - 01:02
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