DISEASE:
Bleeding disorder in hemophilia B carriers

NAME:	Bleeding disorder in hemophilia B carriers
DESCRIPTION:	A rare bleeding disorder in association with carrier mutations in the F9 gene (Xq27.1) encoding coagulation factor IX (FIX), with a biological activity of FIX ?40 IU/dL and characterized clinically by abnormal bleeding as a result of minor injuries or following trauma, surgery or tooth extraction. Spontaneous hemorrhages may occur occasionally. Heavy menstrual bleeding is the most frequent type of bleed in the carriers.
ORPHACODE:	177929
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	F9
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

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