DISEASE:
Mild hemophilia B

NAME:	Mild hemophilia B
DESCRIPTION:	A mild form of hemophilia B characterized by a small deficiency of factor IX (biological activity between 5 and 40 IU/dL) leading to abnormal bleeding as a result of minor injuries or following trauma, surgery or tooth extraction. Spontaneous hemorrhages do not occur. The condition may affect males and female carriers of disease-causing mutations.
ORPHACODE:	169799
SYNONYMS:	Mild congenital F9 deficiency Mild congenital factor IX deficiency
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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- Hemophilia B
- Hemophilia B
- Trombosis - Hemostasis (gene panel)

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- Centre de Génétique Médicale UCL
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- coagulation factor IX

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