DISEASE:
Severe hemophilia B

NAME:	Severe hemophilia B
DESCRIPTION:	A severe form of hemophilia B characterized by a large deficiency of factor IX (biological activity <1 IU/dL) leading to frequent spontaneous hemorrhage and abnormal bleeding as a result of minor injuries or following trauma, surgery or tooth extraction. It primarily affects males but may also be observed in female carriers of disease-causing mutations.
ORPHACODE:	169793
SYNONYMS:	Severe congenital F9 deficiency Severe congenital factor IX deficiency
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	F9
CREATED:	13 May 2019 - 01:02
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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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