DISEASE:
Congenital factor VII deficiency

NAME:	Congenital factor VII deficiency
DESCRIPTION:	A rare, genetic, congenital vitamin K-dependant coagulation factor deficiency disorder characterized by decreased levels or absence of coagulation factor VII (FVII), resulting in bleeding diathesis of variable severity.
ORPHACODE:	327
SYNONYMS:	Congenital proconvertin deficiency Hypoproconvertinemia
XREF(S):	Orphanet MedDRA ICD-10 OMIM
ANALYTE(S):	F7
CREATED:	13 May 2019 - 01:02
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