DISEASE:
Congenital factor XI deficiency

NAME:	Congenital factor XI deficiency
DESCRIPTION:	A rare inherited bleeding disorder characterized by reduced levels and/or activity of factor XI (FXI) resulting in moderate bleeding symptoms, usually occurring after trauma or surgery.
ORPHACODE:	329
SYNONYMS:	Hemophilia C PTA deficiency Plasma thromboplastin antecedent deficiency Rosenthal factor deficiency Rosenthal syndrome
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	F11
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