DISEASE:
Congenital plasminogen activator inhibitor type 1 deficiency

NAME:	Congenital plasminogen activator inhibitor type 1 deficiency
DESCRIPTION:	A rare hemorrhagic disorder due to a constitutional haemostatic factors defect characterized by premature lysis of hemostatic clots and a moderate bleeding tendency.
ORPHACODE:	465
SYNONYMS:	Congenital PAI-1 deficiency
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	SERPINE1
CREATED:	13 May 2019 - 01:02
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