DISEASE:
Cytosolic phospholipase-A2 alpha deficiency associated bleeding disorder

NAME:	Cytosolic phospholipase-A2 alpha deficiency associated bleeding disorder
DESCRIPTION:	A rare genetic hematologic and intestinal disease characterized by childhood onset of bleeding tendency with epistaxis, gum bleeding, gastrointestinal bleeding, hematuria, and menorrhagia due to impaired platelet aggregation and secretion, as well as recurrent gastrointestinal ulcera. Mildly reduced levels of coagulation factor XI have been reported in addition.
ORPHACODE:	477787
SYNONYMS:	PLA2G4A-related platelet dysfunction Platelet dysfunction due to cytosolic phospholipase-A2 alpha deficiency
XREF(S):	Orphanet OMIM ICD-10
ANALYTE(S):	PLA2G4A
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