DISEASE:
Familial hypofibrinogenemia

NAME:	Familial hypofibrinogenemia
DESCRIPTION:	Familial hypofibrinogenemia is a coagulation disorder characterized by mild bleeding symptoms following trauma or surgery due to a reduced plasma fibrinogen concentration.
ORPHACODE:	101041
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	FGA FGB FGG
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

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