

DISEASE:
Combined deficiency of factor V and factor VIII

NAME:	Combined deficiency of factor V and factor VIII
DESCRIPTION:	A rare inherited bleeding disorder due to the reduction in activity and antigen levels of both factor V (FV) and factor VIII (FVIII) and characterized by mild-to-moderate bleeding symptoms.
ORPHACODE:	35909
SYNONYMS:	F5F8D FV and FVIII combined deficiency
XREF(S):	Orphanet OMIM OMIM OMIM ICD-10
ANALYTE(S):	MCFD2 LMAN1
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- [Trombosis - Hemostasis \(gene panel\)](#)

Related Laboratories

- [Centrum Menselijke Erfelijkhed - KUL](#)

Related Analytes

- [lectin, mannose binding 1](#)
- [multiple coagulation factor deficiency 2, ER cargo receptor complex subunit](#)

Related Gene Panels

- [Trombosis - Hemostasis \(107 genes\) - KUL](#)

Source URL: <http://gentest.healthdata.be/disease/2952>