DISEASE:
Congenital factor V deficiency

NAME:	Congenital factor V deficiency
DESCRIPTION:	Congenital factor V deficiency is an inherited bleeding disorder due to reduced plasma levels of factor V (FV) and characterized by mild to severe bleeding symptoms.
ORPHACODE:	326
SYNONYMS:	Owren disease Parahemophilia Proaccelerin deficiency
XREF(S):	Orphanet MedDRA ICD-10 OMIM
ANALYTE(S):	F5
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

RELATED CONTENT

Related Genetic Tests

- Factor V- cambridge, liverpool and hong kong variant (hot spot mutations - p.Arg334Thr, p.Arg306)
- Trombosis - Hemostasis (gene panel)

Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- coagulation factor V

Related Gene Panels

- Trombosis - Hemostasis (107 genes) - KUL