DISEASE:
Hereditary combined deficiency of vitamin K-dependent clotting factors

NAME:	Hereditary combined deficiency of vitamin K-dependent clotting factors
DESCRIPTION:	Combined vitamin K-dependent clotting factors deficiency (VKCFD) is a congenital bleeding disorder resulting from variably decreased levels of coagulation factors II, VII, IX and X, as well as natural anticoagulants protein C, protein S and protein Z.
ORPHACODE:	98434
SYNONYMS:	Hereditary combined deficiency of factors II, VII, IX and X
XREF(S):	Orphanet OMIM OMIM ICD-10
ANALYTE(S):	GGCX VKORC1
CREATED:	13 May 2019 - 01:02
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RELATED CONTENT

Related Genetic Tests

- Deficiency of Vitamin K-Dependent Clotting Factors
- Pseudoxanthoma Elasticum with clotting deficiency
- Trombosis - Hemostasis (gene panel)

Related Laboratories

- Centrum Medische Genetica - UZ Gent
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- gamma-glutamyl carboxylase
- vitamin K epoxide reductase complex subunit 1

Related Gene Panels

- Deficiency of Vitamin K-Dependent Clotting Factors (2 genes) - UGent
- Trombosis - Hemostasis (107 genes) - KUL