# abcam

# Product datasheet

# Anti-Factor VIIa antibody ab61412

### 1 References

#### 製品の概要

製品名 Anti-Factor VIIa antibody

製品の詳細 Rabbit polyclonal to Factor VIIa

由来種 Rabbit

特異性 ab61412 will also cross react with Factor VII, the inactive form of the protein.

アプリケーション 適用あり: ELISA, WB

種交差性 交差種: Human

免疫原 Full length native purified Human Factor VIIa

特記事項 The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

#### 製品の特性

製品の状態 Liquid

保存方法 Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

パップァー Constituents: 50% Water, 50% Glycerol (glycerin, glycerine)

精製度 Ion Exchange Chromatography

特記事項(精製) Salt fractionation followed by ion exchange chromatography

**ポリ/モノ** ポリクローナル

アイソタイプ lgG

#### アプリケーション

The Abpromise guarantee Abpromise保証は、次のテスト済みアプリケーションにおけるab61412の使用に適用されます

アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
ELISA		Use at an assay dependent concentration.
WB		Use at an assay dependent concentration.

# ターゲット情報

機能 Initiates the extrinsic pathway of blood coagulation. Serine protease that circulates in the blood in

a zymogen form. Factor VII is converted to factor VIIa by factor Xa, factor XIIa, factor IXa, or thrombin by minor proteolysis. In the presence of tissue factor and calcium ions, factor VIIa then converts factor X to factor Xa by limited proteolysis. Factor VIIa will also convert factor IX to factor

IXa in the presence of tissue factor and calcium.

組織特異性 Plasma.

関連疾患 Defects in F7 are the cause of factor VII deficiency (FA7D) [MIM:227500]. A hemorrhagic disease

with variable presentation. The clinical picture can be very severe, with the early occurrence of intracerebral hemorrhages or repeated hemarthroses, or, in contrast, moderate with cutaneous-mucosal hemorrhages (epistaxis, menorrhagia) or hemorrhages provoked by a surgical

intervention. Finally, numerous subjects are completely asymptomatic despite very low factor VII

levels.

**配列類似性** Belongs to the peptidase S1 family.

Contains 2 EGF-like domains.

Contains 1 Gla (gamma-carboxy-glutamate) domain.

Contains 1 peptidase S1 domain.

翻訳後修飾 The vitamin K-dependent, enzymatic carboxylation of some glutamate residues allows the

modified protein to bind calcium.

The iron and 2-oxoglutarate dependent 3-hydroxylation of aspartate and asparagine is (R)

stereospecific within EGF domains.

O- and N-glycosylated. N-glycosylation at Asn-205 occurs cotranslationally and is mediated by STT3A-containing complexes, while glycosylation at Asn-382 is post-translational and is mediated STT3B-containing complexes before folding. O-fucosylated by POFUT1 on a conserved serine or threonine residue found in the consensus sequence C2-X(4,5)-[S/T]-C3 of

EGF domains, where C2 and C3 are the second and third conserved cysteines.

細胞内局在 Secreted.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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