abcam

Product datasheet

Anti-Factor VII antibody ab104564

画像数1

製品の概要

製品名 Anti-Factor VII antibody

製品の詳細 Sheep polyclonal to Factor VII

由来種 Sheep

特異性 ab104564 is specific for Factor VII as determined by immunodiffusion. Identity was established

with purified Factor VII.

 アプリケーション
 適用あり: IHC-P

 種交差性
 交差種: Human

免疫原 Human Factor VII purified from plasma.

ポジティブ・コントロール Human Liver tissue

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.

バッファー pH: 7.40

Constituents: 50% Glycerol (glycerin, glycerine), HEPES

精製度 Ammonium Sulphate Precipitation

ポリ/モノ ポリクローナル

アイソタイプ IgG

アプリケーション

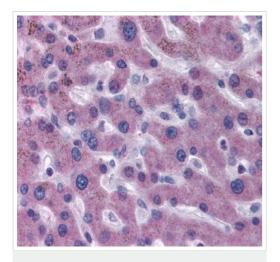
アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

| アプリケーション | Abreviews | 特記事項 |
|----------|-----------|--|
| IHC-P | | Use a concentration of 5 µg/ml. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol. |

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]. FA7D is a rare hereditary hemorrhagic disease. The clinical picture can be very severe, with the early occurrence of intracerebral hemorrhages or hemarthroses, or, in contrast, moderate with cutaneous-mucosal hemorrhages (epistaxis, menorrhagia) or hemorrhages provoked by a surgical intervention. Numerous subjects are completely asymptomatic despite a very low F7 level. |
| 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. |
| 細胞内局在 | Secreted. |

画像



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-Factor VII antibody (ab104564)

ab104564, at 5 ug/ml, staining Factor VII in formalin-fixed, paraffinembedded Human Liver by Immunohistochemistry, using a biotinylated secondary antibody, alkaline phosphatase-streptavidin and chromogen.

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

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