产品名称: C1INH Rabbit Polyclonal Antibody

产品货号: APRab07717



产品概述 (Summary)

产品名称 (Production Name) C1INH Rabbit Polyclonal Antibody

描述 (Description) Rabbit polyclonal Antibody

宿主 (Host) Rabbit

应用 (Application)WB,IHC,ICC/IF,ELISA种属反应性 (Reactivity)Human,Rat,Mouse

产品性能 (Performance)

偶联物 (Conjugation)Unconjugated修饰 (Modification)Unmodified

同种型 (Isotype) IgG

克隆 (Clonality) Polyclonal 形式 (Form) Liquid

Store at 4°C short term. Aliquot and store at -20°C long term. Avoid 存放说明 (Storage)

freeze/thaw cycles.

Liquid in PBS containing 50% glycerol, 0.5% protective protein and 0.02% 储存溶液 (Buffer)

New type preservative N.

纯化方式 (Purification) Affinity purification

免疫原信息 (Immunogen)

基因名 (Gene Name) SERPING1

SERPING1; C1IN; C1NH; Plasma protease C1 inhibitor; C1 Inh; C1Inh; C1 別名 (Alternative Names)

esterase inhibitor; C1-inhibiting factor; Serpin G1

基因 ID (Gene ID) 710.0

P05155.The antiserum was produced against synthesized peptide derived 蛋白ID (SwissProt ID)

from human SERPING1. AA range:342-391

产品应用 (Application)

稀释比 (Dilution Ratio) WB 1:500-1:2000,IHC 1:100-1:300,ICC/IF 1:50-1:200,ELISA 1:20000-1:40000

蛋白分子量 (Molecular Weight) 55kDa

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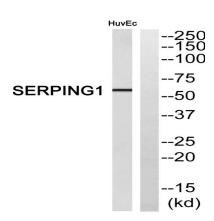
研究背景 (Background)

This gene encodes a highly glycosylated plasma protein involved in the regulation of the complement cascade. Its protein inhibits activated C1r and C1s of the first complement component and thus regulates complement activation. Deficiency of this protein is associated with hereditary angioneurotic oedema (HANE). Alternative splicing results in multiple transcript variants encoding the same isoform. [provided by RefSeq, Jul 2008], disease: Defects in SERPING1 are the cause of hereditary angioedema (HAE) [MIM:106100]; also called hereditary angioneurotic edema (HANE). HAE is an autosomal dominant disorder characterized by episodic local subcutaneous edema and submucosal edema involving the upper respiratory and gastrointestinal tracts. HAE due to C1 esterase inhibitor deficiency is comprised of two clinically indistinguishable forms. In HAE type 1, representing 85% of patients, serum levels of C1 esterase inhibitor are less than 35% of normal. In HAE type 2, the levels are normal or elevated, but the protein is non-functional, function: Activation of the C1 complex is under control of the C1-inhibitor. It forms a proteolytically inactive stoichiometric complex with the C1r or C1s proteases. May play a potentially crucial role in regulating important physiological pathways including complement activation, blood coagulation, fibrinolysis and the generation of kinins. Very efficient inhibitor of FXIIa. Inhibits chymotrypsin and kallikrein., online information:C1-inhibitor entry, online information:SERPING1 mutation db,polymorphism:Chymotrypsin uses Ala-465 as its reactive site in normal plasma protease C1 inhibitor, and His-466 as its reactive site in the variant His-466, PTM:Can be proteolytically cleaved by E.coli stcE., PTM: Highly glycosylated (49%), similarity: Belongs to the serpin family, subunit: Binds to E.coli stcE which allows localization of SERPING1 to cell membranes thus protecting the bacteria against complementmediated lysis. Interacts with MASP1.,

研究领域 (Research Area)

Complement and coagulation cascades;

图片 (Image Data)



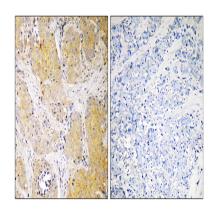
Western blot analysis of SERPING1 Antibody. The lane on the right is blocked with the SERPING1 peptide.

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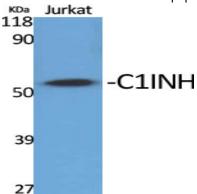
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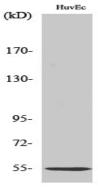




Immunohistochemistry analysis of paraffin-embedded human breast cancer, using SERPING1 Antibody. The picture on the right is blocked with the SERPING1 peptide.



Western Blot analysis of various cells using C1INH Polyclonal Antibody



Western Blot analysis of HuvEc cells using C1INH Polyclonal Antibody

注意事项 (Note)

For research use only .