

FA11 (heavy chain, Cleaved-Glu19) rabbit

pAb

Cat No.:ES19970

For research use only

Overview

| Product Name | FA11 (heavy chain, Cleaved-Glu19) rabbit pAb |
|--------------------------|--|
| Host species | Rabbit |
| Applications | WB; ELISA |
| Species Cross-Reactivity | Human; Mouse |
| Recommended dilutions | WB 1:1000-2000 ELISA 1:5000-20000 |
| Immunogen | Synthesized peptide derived from human FA11 |
| | (heavy chain, Cleaved-Glu19) |
| Specificity | This antibody detects endogenous levels of Human, |
| | Mouse FA11 (heavy chain, Cleaved-Glu19, protein |
| | was cleaved amino acid sequence between 18-19) |
| Formulation | Liquid in PBS containing 50% glycerol, 0.5% BSA and |
| | 0.02% sodium azide. |
| Storage | Store at -20 $^\circ\!{ m C}$. Avoid repeated freeze-thaw cycles. |
| Protein Name | FA11 (heavy chain, Cleaved-Glu19) |
| Gene Name | F11 |
| Cellular localization | Secreted. |
| Purification | The antibody was affinity-purified from rabbit |
| | antiserum by affinity-chromatography using |
| | epitope-specific immunogen. |
| Clonality | Polyclonal |
| Concentration | 1 mg/ml |
| Observed band | 40 66kD |
| Human Gene ID | 2160 |
| Human Swiss-Prot Number | P03951 |
| Alternative Names | Coagulation factor XI (FXI;EC 3.4.21.27;Plasma |
| | thromboplastin antecedent;PTA) [Cleaved into: |
| | Coagulation factor XIa heavy chain; Coagulation |
| | factor XIa light chain] |
| Background | This gene encodes coagulation factor XI of the blood |
| | coagulation cascade. This protein is present in |
| | plasma as a zymogen, which is a unique plasma |
| | coagulation enzyme because it exists as a |
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homodimer consisting of two identical polypeptide chains linked by disulfide bonds. During activation of the plasma factor XI, an internal peptide bond is cleaved by factor XIIa (or XII) in each of the two chains, resulting in activated factor XIa, a serine protease composed of two heavy and two light chains held together by disulfide bonds. This activated plasma factor XI triggers the middle phase of the intrisic pathway of blood coagulation by activating factor IX. Defects in this factor lead to Rosenthal syndrome, a blood coagulation abnormality. [provided by RefSeq, Jul 2008],



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