

# Factor XIIIa Rabbit mAb

货号: B30634

产品信息

|       |  |
|-------|--|
| 反应    | Human  |
| 宿主    | Rabbit   |
| 克隆性   | Monoclonal   |
| 预测反应  |  |
| 应用    | WB IHC IF/ICC FC   |
| 推荐浓度  | <b>WB:</b> 1:500 - 1:2000<br><b>IHC:</b> 1:50 - 1:200<br><b>IF/ICC:</b> 1:50 - 1:200<br><b>FC:</b> 1:20 - 1:50 |
| 理论分子量 | 83kDa  |
| 实测分子量 | 83kDa  |
| 形式    | Liquid   |
| 保存条件  | Store at -20°C. Avoid freeze / thaw cycles.<br>Buffer: PBS with 0.75% BSA,50% glycerol,pH7.3.                  |
| 偶联物   | Unconjugated   |
| 阳性对照  | Jurkat,SH-SY5Y,BxPC-3,Mouse testis,Rat brain,Rat testis  |
| 细胞定位  | Cytoplasm,Secreted   |
| 纯化    | Affinity purification  |

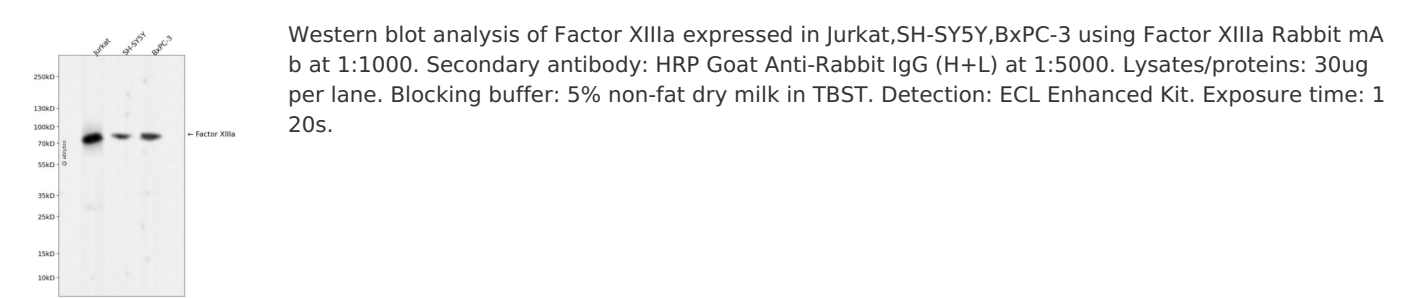
抗原信息

|      |  |
|------|--|
| 抗原信息 | Recombinant fusion protein corresponding to Human Factor XIIIa.  |
| 序列   | LEQASLHFFVTARINETRDVLAKQKSTVLTPIEIIKVRGTQVVGSDMTVTVEFTNPLKETLRNVVWVHLDGPGVTRPMKKMF<br>REIRPNSTVQWEEVCRPWVSGHRKLIASMSSDSL RHVYGELDVQIQRRPSM |

靶点信息

|       |   |
|-------|---|
| 研究背景  | This gene encodes the coagulation factor XIII A subunit. Coagulation factor XIII is the last zymogen to become activated in the blood coagulation cascade. Plasma factor XIII is a heterotetramer composed of 2 A subunits and 2 B subunits. The A subunits have catalytic function, and the B subunits do not have enzymatic activity and may serve as plasma carrier molecules. Platelet factor XIII is comprised only of 2 A subunits, which are identical to those of plasma origin. Upon cleavage of the activation peptide by thrombin and in the presence of calcium ion, the plasma factor XIII dissociates its B subunits and yields the same active enzyme, factor XIIIa, as platelet factor XIII. This enzyme acts as a transglutaminase to catalyze the formation of gamma-glutamyl-epsilon-lysine crosslinking between fibrin molecules, thus stabilizing the fibrin clot. It also crosslinks alpha-2-plasmin inhibitor, or fibronectin, to the alpha chains of fibrin. Factor XIII deficiency is classified into two categories: type I deficiency, characterized by the lack of both the A and B subunits; and type II deficiency, characterized by the lack of the A subunit alone. These defects can result in a lifelong bleeding tendency, defective wound healing, and habitual abortion. |
| 基因ID  | 2162  |
| 基因名   | F13A1   |
| Swiss | P00488  |
| 别名    | F13A1;F13A  |

产品验证



实验步骤

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