[Primary Antibody]

human Fibrinogen Rabbit pAb

- DATASHEET -Host: Rabbit Isotype: IgG Clonality: Polyclonal GenelD: 2243 SWISS: P02671 Target: human Fibrinogen Purification: affinity purified by Protein A Concentration: 1mg/ml Storage: 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol. Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles. Background: Fibrinogen is the main protein of blood coagulation system. It is a large protein and it consists of two identical subunits that contain three polypeptide chains: alpha, beta and gamma. All chains are connected with each other by a number of disulfide bonds. Fibrinopeptides A (1 to 16 amino acids) and B (1 to 17 amino acids) are released by thrombin from the N terminal parts of alpha and beta chains, respectively. In this way fibrinogen is converted into fibrin, which by means of polymerization forms a fibrin clot. Fibrinogen clotting underlies pathogenesis of MI, thromboembolism and thromboses of arteries and veins, since fibrin is the main substrate for thrombus formation. Fibrinogen activation is also involved in pathogenesis of inflammation, tumor growth and many other diseases. The normal fibrinogen concentration in plasma is about 3 mg/ml. The elevated level of fibrinogen in patient's blood is regarded as an independent risk factor for cardiovascular diseases. An increase in blood fibrinogen concentration was shown to be a strong predictor of coronary heart disease (Sonel A. et al, and Rapold H.J. et al). All these facts make fibrinogen an important parameter in the diagnosis of cardiovascular diseases.

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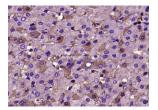
Applications: IHC-P (1:100-500) IHC-F (1:100-500) IF (1:100-500) Flow-Cyt (1µg /test)

Reactivity: Human, Rat (predicted: Mouse)

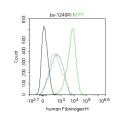
Predicted MW.: 95/340 kDa

Subcellular Location:

- VALIDATION IMAGES



Paraformaldehyde-fixed, paraffin embedded (human liver tissue); Antigen retrieval by boiling in sodium citrate buffer (pH6.0) for 15min; Block endogenous peroxidase by 3% hydrogen peroxide for 20 minutes; Blocking buffer (normal goat serum) at 37°C for 30min; Antibody incubation with (human Fibrinogen) Polyclonal Antibody, Unconjugated (bs-1240R) at 1:400 overnight at 4°C, followed by operating according to SP Kit(Rabbit) (sp-0023) instructionsand DAB staining.



Blank control (black line) :HepG2. Primary Antibody (green line): Rabbit Anti-human Fibrinogen antibody (bs-1240R) Dilution:1ug/Test; Secondary Antibody (white blue line) : Goat anti-rabbit IgG-AF488 Dilution: 0.5ug/Test. Isotype control (orange line) : Normal Rabbit IgG Protocol The cells were fixed with 4% PFA (10min at room temperature) and then permeabilized with 90% ice-cold methanol for 20 min at -20°C, The cells were then incubated in 5%BSA to block non-specific protein-protein interactions for 30 min at room temperature .Cells stained with Primary Antibody for 30 min at room temperature. The secondary antibody used for 40 min at room temperature. Acquisition of 20,000 events was performed.

- SELECTED CITATIONS -

- [IF=25.841] Eszter Á. Tóth. et al. Formation of a protein corona on the surface of extracellular vesicles in blood plasma. J Extracell Vesicles. 2021 Sep;10(11):e12140 Other ;Human. 34520123
- [IF=8.7] Xue Sun. et al. An injectable shape-adaptive hydrogel system for subconjunctival injuries: In situ and permanently releases rapamycin to prevent fibrosis via promoting autophagy. MATER TODAY BIO. 2025 Feb;30:101380 IF,WB ;Human. 39790484
- [IF=6.1] Shengjun Cheng. et al. Development of substrate-independent heparin coating to mitigate surface-induced thrombogenesis: efficacy and mechanism. J MATER CHEM B. 2024 Sep;: IF ;Human. 39352074
- [IF=5.4] Chen Xu. et al.Oral functional protein Z: Mitigation of thrombosis via thrombin inhibition to prevent cardiovascular disease.COLLOIDS AND SURFACES B: BIOINTERFACES.2025 Feb 20:251:114569. Turbidity assay ;Human. 39999698
- [IF=2.7] Ya Liu. et al. Validation for the function of protein C in mouse models. PEERJ. 2024 Apr;12:e17261 IHC ;Mouse. 38680896