Product datasheet

Anti-Fibrinogen Alpha Chain/FGA Antibody

Catalog Number: A00816-3



Building C21, 3rd to 5th Floors, Optics Valley Biopharmaceutical Accelerator, East Lake High-Tech Development Zone, Wuhan.

Web: www.boster.com Phone: 027-67845390/1/2 Email: boster@boster.com

Basic Inform	nation	
Product Name	Anti-Fibrinogen Alpha Chain/FGA Antibody	
Gene Name	FGA	
Source	Rabbit	
Clonality	Polyclonal	
Isotype	IgG	
Species Reactivity	human	
Tested Application	WB, IHC, FCM, ELISA	
Contents	500 ug/ml antibody with PBS, 0.02% NaN3, 1 mg/ml BSA and 50% glycerol.	
Immunogen	E. coli-derived human Fibrinogen alpha chain/FGA recombinant protein (Position: E139-Q784). Human FGA shares 61.6% and 62.5% amino acid (aa) sequence identity with mouse and rat FGA, respectively.	
Concentration	500 ug/ml	
Purification	Immunogen affinity purified.	
Observed MW	95 kDa	
Dilution Ratios	Western blot (WB): Immunohistochemistry (IHC): Flow Cytometry (Fixed): Enzyme linked immunosorbent assay (ELISA): (Boiling the paraffin sections in 10mM citrate buffe mins is required for the staining of formalin/paraffil must be determined by end user.	

Storage

12 months from date of receipt, -20°C as supplied. 6 months 2 to 8°C after reconstitution. Avoid repeated freezing and thawing.

Background Information

Fibrinogen alpha chain is a protein that in humans is encoded by the FGA gene. This gene encodes the alpha subunit of the coagulation factor fibrinogen, which is a component of the blood clot. Following vascular injury, the encoded preproprotein is proteolytically processed by thrombin during the conversion of fibrinogen to fibrin. Mutations in this gene lead to several disorders, including dysfibrinogenemia, hypofibrinogenemia, afibrinogenemia and renal amyloidosis. Alternative splicing results in multiple transcript variants, at least one of which encodes an isoform that undergoes proteolytic processing.

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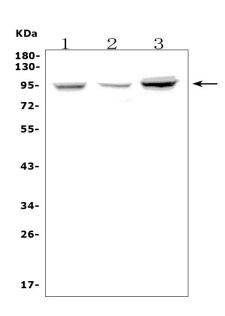
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Reference

Anti-Fibrinogen Alpha Chain/FGA Antibody被引用在1文献中。

Selected Validation Data



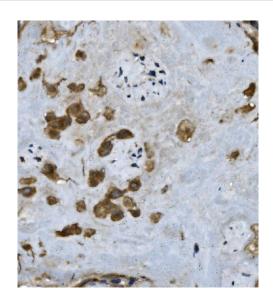
Western blot analysis of Fibrinogen Alpha Chain/FGA using anti-Fibrinogen Alpha Chain/FGA antibody (A00816-3). The sample well of each lane was loaded with 30 ug of sample under reducing conditions.

Lane 1: human T-47D whole cell lysates,

Lane 2: human Caco-2 whole cell lysates,

Lane 3: human SW620 whole cell lysates.

After electrophoresis, proteins were transferred to a membrane. Then the membrane was incubated with rabbit anti-Fibrinogen Alpha Chain/FGA antigen affinity purified polyclonal antibody (A00816-3) at a dilution of 1:1000 and probed with a goat anti-rabbit IgG-HRP secondary antibody (Catalog # BA1054). The signal is developed using ECL Plus Western Blotting Substrate (Catalog # AR1197). A specific band was detected for Fibrinogen Alpha Chain/FGA at approximately 95 kDa. The expected band size for Fibrinogen Alpha Chain/FGA is at 95 kDa.



IHC analysis of Fibrinogen Alpha Chain/FGA using anti-Fibrinogen Alpha Chain/FGA antibody (A00816-3).

Fibrinogen Alpha Chain/FGA was detected in a paraffin-embedded section of human placenta tissue. Biotinylated goat anti-rabbit IgG was used as secondary antibody. The tissue section was incubated with rabbit anti-Fibrinogen Alpha Chain/FGA Antibody (A00816-3) at a dilution of 1:200 and developed using Strepavidin-Biotin-Complex (SABC) (Catalog # SA1022) with DAB (Catalog # AR1027) as the chromogen.

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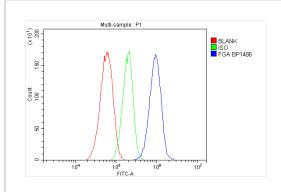
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Flow Cytometry analysis of HepG2 cells using anti-Fibrinogen Alpha Chain/FGA antibody (A00816-3).

Overlay histogram showing HepG2 cells stained with A00816-3 (Blue line). To facilitate intracellular staining, cells were fixed with 4% paraformaldehyde and permeabilized with permeabilization buffer. The cells were blocked with 10% normal goat serum. And then incubated with rabbit anti-Fibrinogen Alpha Chain/FGA Antibody (A00816-3) at 1:100 dilution for 30 min at 20°C. DyLight® 488 conjugated goat anti-rabbit IgG (BA1127) was used as secondary antibody at 1:100 dilution for 30 minutes at 20°C. Isotype control antibody (Green line) was rabbit IgG at 1:100 dilution used under the same conditions. Unlabelled sample without incubation with primary antibody and secondary antibody (Red line) was used as a blank control.