abcam

Product datasheet

Anti-Fibrinogen gamma chain antibody [5A6] abl 19948

★★★★★ 1 Abreviews 8 References 3 Images

Overview

Product name Anti-Fibrinogen gamma chain antibody [5A6]

Description Mouse monoclonal [5A6] to Fibrinogen gamma chain

Host species Mouse

Tested applications
Suitable for: WB, ELISA, ICC/IF
Species reactivity
Reacts with: Mouse, Human

Immunogen Recombinant fragment, corresponding to amino acids 210-437 of Human Fibrinogen gamma

chain

Positive control Recombinant Fibrinogen gamma chain protein, NIH3T3 cells

General notesThis product was changed from ascites to supernatant. Lot no's high than GR240839-12 are from

Tissue Culture Supernatant

The Life Science industry has been in the grips of a reproducibility crisis for a number of years. Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Store at -20°C long term.

Storage buffer Preservative: 0.05% Sodium azide

Constituent: PBS

Purity Protein G purified

Purification notes Purified from tissue culture supernatant.

Clonality Monoclonal

Clone number 5A6

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Isotype IgG2b

Applications

The Abpromise guarantee

Our <u>Abpromise guarantee</u> covers the use of ab119948 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
WB	*** <u>*</u>	1/500 - 1/2000. Predicted molecular weight: 52 kDa.
ELISA		1/10000.
ICC/IF		1/200 - 1/1000.

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Function Fibrinogen has a double function: yielding monomers that polymerize into fibrin and acting as a

cofactor in platelet aggregation.

Involvement in disease Defects in FGG are a cause of thrombophilia.

Defects in FGG are a cause of congenital afibrinogenemia (CAFBN) [MIM:202400]. It is a rare

autosomal recessive disorder characterized by complete absence of detectable fibrinogen.

Sequence similarities

Contains 1 fibrinogen C-terminal domain.

DomainA long coiled coil structure formed by 3 polypeptide chains connects the central nodule to the C-

terminal domains (distal nodules). The long C-terminal ends of the alpha chains fold back,

contributing a fourth strand to the coiled coil structure.

Post-translational modifications

Conversion of fibrinogen to fibrin is triggered by thrombin, which cleaves fibrinopeptides A and B from alpha and beta chains, and thus exposes the N-terminal polymerization sites responsible for

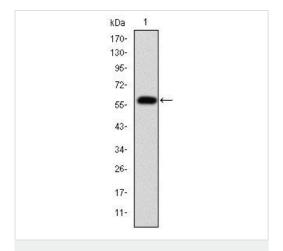
the formation of the soft clot. The soft clot is converted into the hard clot by factor XIIIA which catalyzes the epsilon-(gamma-glutamyl)lysine cross-linking between gamma chains (stronger)

and between alpha chains (weaker) of different monomers.

Sulfation of C-terminal tyrosines increases affinity for thrombin.

Cellular localization Secreted.

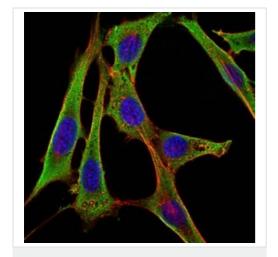
Images



Western blot - Anti-Fibrinogen gamma chain antibody [5A6] (ab119948)

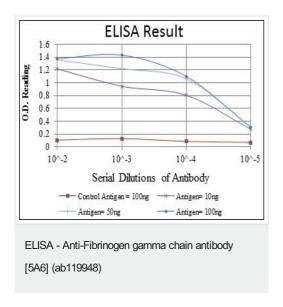
Anti-Fibrinogen gamma chain antibody [5A6] (ab119948) at 1/500 dilution + Recombinant Fibrinogen gamma chain protein

Predicted band size: 52 kDa



Immunocytochemistry/ Immunofluorescence - Anti-Fibrinogen gamma chain antibody [5A6] (ab119948)

Immunofluorescence analysis of NIH3T3 cells using ab119948 at a dilution of 1/200 (green). Blue: DRAQ5 fluorescent DNA dye. Red: Actin filaments have been labeled with Alexa Fluor-555 phalloidin.



ELISA, using ab119948 at a dilution of 1/10000.

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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