abcam

Product datasheet

Anti-Fibrinogen gamma chain antibody [EPR3085] ab133339



1 References 2 Images

Overview

Product name Anti-Fibrinogen gamma chain antibody [EPR3085]

Description Rabbit monoclonal [EPR3085] to Fibrinogen gamma chain

Host species Rabbit

Tested applications Suitable for: WB

Unsuitable for: ICC/IF,IHC-P or IP

Species reactivity Reacts with: Human

Immunogen Synthetic peptide. This information is proprietary to Abcam and/or its suppliers.

Positive control Human plasma lysate

General notesThis product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility

- Improved sensitivity and specificity

- Long-term security of supply

- Animal-free production

For more information see here.

Our RabMAb[®] technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to **RabMAb**[®] **patents**.

Mouse, Rat: We have preliminary internal testing data to indicate this antibody may not react with

these species. Please contact us for more information.

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at -20°C. Stable for 12 months at -20°C.

Storage buffer pH: 7.2

Preservative: 0.05% Sodium azide

Constituents: 0.1% BSA, 40% Glycerol (glycerin, glycerine), 9.85% Tris glycine, 50% Tissue

culture supernatant

Purity Tissue culture supernatant

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Clonality Monoclonal
Clone number EPR3085
Isotype IgG

Applications

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab133339 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		1/1000 - 1/10000. Predicted molecular weight: 52 kDa.

Application notes Is unsuitable for ICC/IF,IHC-P or IP.

Target

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.

Domain A 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

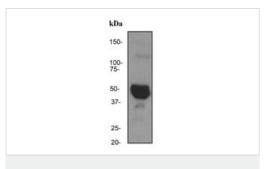
the formation of the soft clot. The soft clot is converted into the hard clot by factor XIIIA which catalyzes the epsilon-(gamma-glutamyl)lysine cross-linking between gamma chains (stronger)

and between alpha chains (weaker) of different monomers.

Sulfation of C-terminal tyrosines increases affinity for thrombin.

Cellular localization Secreted.

Images



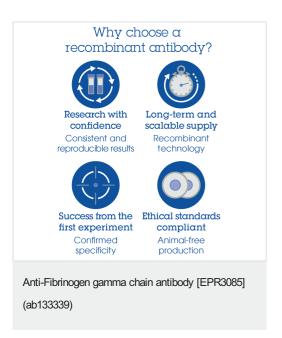
Western blot - Anti-Fibrinogen gamma chain antibody [EPR3085] (ab133339)

Anti-Fibrinogen gamma chain antibody [EPR3085] (ab133339) at 1/1000 dilution + Human plasma cell lysate at 10 μg

Secondary

Goat Anti-rabbit HRP at 1/2000 dilution

Predicted band size: 52 kDa



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

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