abcam

Product datasheet

Alexa Fluor® 790 Anti-GAPDH antibody [mAbcam 9484] - Loading Control ab184578

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Overview

Product name Alexa Fluor® 790 Anti-GAPDH antibody [mAbcam 9484] - Loading Control

Description Alexa Fluor® 790 Mouse monoclonal [mAbcam 9484] to GAPDH - Loading Control

Host species Mouse

Conjugation Alexa Fluor® 790. Ex: 782nm, Em: 805nm

Tested applications Suitable for: WB

Species reactivity Reacts with: Mouse, Human

Immunogen Full length native protein (purified). This information is proprietary to Abcam and/or its suppliers.

Positive control This antibody gave a positive signal in both HeLa and NIH3T3 whole cell lysates.

General notes According to our customer's feedback this antibody does not recognise meningococcal GapA1 or

GapA2 (GAPDH) recombinant proteins.

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outlicensing@thermofisher.com.

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

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Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C short term (1-2 weeks). Upon delivery aliquot. Store at -20°C or -

80°C. Avoid freeze / thaw cycle. Stable for 12 months at -20°C. Store In the Dark.

Storage buffer Preservative: 0.02% Sodium azide

Constituents: PBS, 1% BSA, 30% Glycerol (glycerin, glycerine)

Clonality Monoclonal

Clone number mAbcam 9484

Myeloma Sp2/0-Ag14

lsotype lgG2b **Light chain type** kappa

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab184578 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>(2)</u>	Use a concentration of 1 µg/ml. Detects a band of approximately 37 kDa (predicted molecular weight: 37 kDa).

Target

Function

Has both glyceraldehyde-3-phosphate dehydrogenase and nitrosylase activities, thereby playing a role in glycolysis and nuclear functions, respectively. Participates in nuclear events including transcription, RNA transport, DNA replication and apoptosis. Nuclear functions are probably due to the nitrosylase activity that mediates cysteine S-nitrosylation of nuclear target proteins such as SIRT1, HDAC2 and PRKDC (By similarity). Glyceraldehyde-3-phosphate dehydrogenase is a key enzyme in glycolysis that catalyzes the first step of the pathway by converting D-glyceraldehyde 3-phosphate (G3P) into 3-phospho-D-glyceroyl phosphate.

Pathway Carbohydrate degradation; glycolysis; pyruvate from D-glyceraldehyde 3-phosphate: step 1/5.

Sequence similaritiesBelongs to the glyceraldehyde-3-phosphate dehydrogenase family.

Post-translational S-nitrosylation of Cys-152 leads to interaction with SIAH1, followed by translocation to the

modifications nucleus.

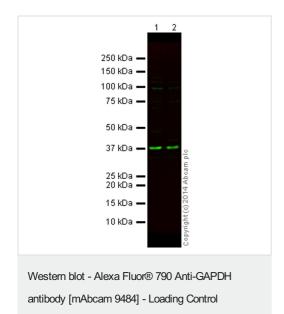
ISGylated.

Cytoplasm > cytosol. Nucleus. Cytoplasm > perinuclear region. Membrane. Translocates to the

nucleus following S-nitrosylation and interaction with SIAH1, which contains a nuclear localization

signal (By similarity). Postnuclear and Perinuclear regions.

Images



(ab184578)

All lanes : Alexa Fluor® 790 Anti-GAPDH antibody [mAbcam 9484] - Loading Control (ab184578) at 1 µg/ml

Lane 1 : HeLa (Human epithelial carcinoma cell line) Whole Cell Lysate

Lane 2: NIH 3T3 (Mouse embryonic fibroblast cell line) Whole Cell Lysate

Lysates/proteins at 10 µg per lane.

Predicted band size: 37 kDa **Observed band size:** 37 kDa

This blot was produced using a 4-12% Bis-tris gel under the MOPS buffer system. The gel was run at 200V for 50 minutes before being transferred onto a Nitrocellulose membrane at 30V for 70 minutes. The membrane was then blocked for an hour using 5% Milk before being incubated with ab184578 overnight at 4°C. Antibody binding was detected after washing to remove excess antibody and imaged using the Licor Odyssey CLx.

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

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