abcam

Product datasheet

Anti-RhoGDI antibody [EPR3773] - BSA and Azide free ab248423



Recombinant

RabMAb

3 Images

Overview

Product name Anti-RhoGDI antibody [EPR3773] - BSA and Azide free

Description Rabbit monoclonal [EPR3773] to RhoGDI - BSA and Azide free

Host species Rabbit

Tested applications Suitable for: WB, IHC-P

Unsuitable for: Flow Cyt or IP

Species reactivity Reacts with: Mouse, Human

Predicted to work with: Rat

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

Positive control WB: HEK293T, Jurkat and HeLa cell lysates.

General notes ab248423 is the carrier-free version of **ab133248**.

Our <u>carrier-free</u> antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. The carrier-free buffer and high concentration allow for increased conjugation efficiency.

This conjugation-ready format is designed for use with fluorochromes, metal isotopes, oligonucleotides, and enzymes, which makes them ideal for antibody labelling, functional and cell-based assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our <u>conjugation kits</u> for antibody conjugates that are ready-to-use in as little as 20 minutes with <1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.

This product is compatible with the Maxpar[®] Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar[®] is a trademark of Fluidigm Canada Inc.

This 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

Properties

Form Liquid

Storage instructions Shipped at 4°C. Store at +4°C. Do Not Freeze.

Storage buffer pH: 7.2

Constituent: PBS

Carrier free Yes

Purity Protein A purified

Clonality Monoclonal
Clone number EPR3773

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab248423 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		Use at an assay dependent concentration. Predicted molecular weight: 23 kDa.
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.

Application notes Is unsuitable for Flow Cyt or IP.

Target

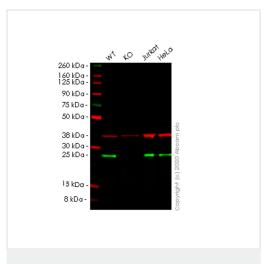
Function Regulates the GDP/GTP exchange reaction of the Rho proteins by inhibiting the dissociation of

GDP from them, and the subsequent binding of GTP to them.

Sequence similarities Belongs to the Rho GDI family.

Cellular localization Cytoplasm.

Images



Western blot - Anti-RhoGDI antibody [EPR3773] - BSA and Azide free (ab248423)

All lanes : Anti-RhoGDI antibody [EPR3773] (<u>ab133248</u>) at 1/1000 dilution

Lane 1: Wild-type HEK293T cell lysate

Lane 2: ARHGDIA knockout HEK293T cell lysate

Lane 3 : Jurkat cell lysate

Lane 4 : HeLa cell lysate

Lysates/proteins at 20 µg per lane.

Secondary

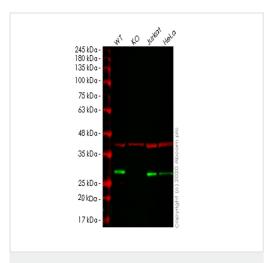
All lanes : Goat anti-Rabbit lgG H&L (IRDye® 800CW) preadsorbed (ab216773) at 1/10000 dilution

Predicted band size: 23 kDa Observed band size: 23 kDa

This data was developed using the same antibody clone in a different buffer formulation (<u>ab133248</u>).

Lanes 1-4: Merged signal (red and green). Green - <u>ab133248</u> observed at 23 kDa. Red - loading control <u>ab8245</u> observed at 36 kDa.

<u>ab133248</u> Anti-RhoGDI antibody [EPR3773] was shown to specifically react with RhoGDI in wild-type HEK293T cells. Loss of signal was observed when knockout cell line <u>ab266446</u> (knockout cell lysate <u>ab257355</u>) was used. Wild-type and RhoGDI knockout samples were subjected to SDS-PAGE. <u>ab133248</u> and Anti-GAPDH antibody [6C5] - Loading Control (<u>ab8245</u>) were incubated at room temperature for 2.5 hours at 1 in 1000 dilution and 1 in 20000 dilution respectively. Blots were developed with Goat anti-Rabbit lgG H&L (IRDye[®] 800CW) preadsorbed (<u>ab216773</u>) and Goat anti-Mouse lgG H&L (IRDye[®] 680RD) preadsorbed (<u>ab216776</u>) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.



Western blot - Anti-RhoGDI antibody [EPR3773] - BSA and Azide free (ab248423)

All lanes : Anti-RhoGDI antibody [EPR3773] (<u>ab133248</u>) at 1/1000 dilution

Lane 1: Wild-type HEK293T cell lysate

Lane 2: ARHGDIA knockout HEK293T cell lysate

Lane 3 : Jurkat cell lysate

Lane 4 : HeLa cell lysate

Lysates/proteins at 20 µg per lane.

Secondary

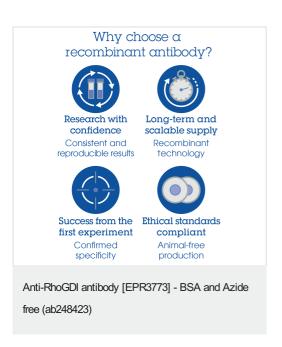
All lanes : Goat anti-Rabbit lgG H&L (IRDye® 800CW) preadsorbed (**ab216773**) at 1/10000 dilution

Predicted band size: 23 kDa **Observed band size:** 27 kDa

This data was developed using the same antibody clone in a different buffer formulation (<u>ab133248</u>).

Lanes 1-4: Merged signal (red and green). Green - <u>ab133248</u> observed at 27 kDa. Red - loading control <u>ab8245</u> observed at 36 kDa.

<u>ab133248</u> Anti-RhoGDI antibody [EPR3773] was shown to specifically react with RhoGDI in wild-type HEK293T cells. Loss of signal was observed when knockout cell line <u>ab266447</u> (knockout cell lysate <u>ab257356</u>) was used. Wild-type and RhoGDI knockout samples were subjected to SDS-PAGE. <u>ab133248</u> and Anti-GAPDH antibody [6C5] - Loading Control (<u>ab8245</u>) were incubated overnight at 4°C at 1 in 1000 dilution and 1 in 20000 dilution respectively. Blots were developed with Goat anti-Rabbit lgG H&L (IRDye® 800CW) preadsorbed (<u>ab216773</u>) and Goat anti-Mouse lgG H&L (IRDye® 680RD) preadsorbed (<u>ab216776</u>) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.



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

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