abcam

Product datasheet

Anti-CD168 antibody [EPR4054] - BSA and Azide free ab271887



Recombinant

RabMAb

8 Images

Overview

Product name Anti-CD168 antibody [EPR4054] - BSA and Azide free

Description Rabbit monoclonal [EPR4054] to CD168 - BSA and Azide free

Host species Rabbit

Tested applications Suitable for: IHC-P, IP, WB

Unsuitable for: Flow Cyt or ICC/IF

Species reactivity Reacts with: Mouse, Rat, Human

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

Positive control IP: MCF-7. IHC-P: Human gastric carcinoma tissue. Human testis tissue.

General notes ab271887 is the carrier-free version of <u>ab124729</u>.

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 monoclonal antibodies. For details on our patents, please refer to **RabMAb**[®] **patents**.

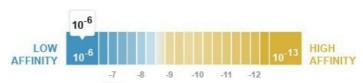
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Properties

Form Liquid

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

Dissociation constant (K_D) $K_D = 1.23 \times 10^{-6} M$



Learn more about K_D

Storage buffer pH: 7.2

Constituent: PBS

Carrier free Yes

Purity Protein A purified

ClonalityMonoclonalClone numberEPR4054

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab271887 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
IHC-P		Use at an assay dependent concentration. Antigen retrieval is recommended
IP		Use at an assay dependent concentration.
WB		Use at an assay dependent concentration. Predicted molecular weight: 84 kDa. The mouse and rat recommendation is based on the WB results. This antibody may not be suitable for IHC with mouse or rat samples.

Application notes Is unsuitable for Flow Cyt or ICC/IF.

Target

Function Involved in cell motility. When hyaluronan binds to HMMR, the phosphorylation of a number of

proteins, including the focal adhesion kinase occurs. May also be involved in cellular

transformation and metastasis formation, and in regulating extracellular-regulated kinase (ERK)

activity.

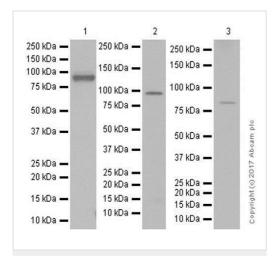
Tissue specificity

Expressed in breast cancer cell lines and in normal breast tissue.

Cellular localization

Cell surface. Cytoplasm.

Images



Western blot - Anti-CD168 antibody [EPR4054] - BSA and Azide free (ab271887)

All lanes : Anti-CD168 antibody [EPR4054] (ab124729) at 1/1000 dilution

Lane 1 : MDA-MB-435 (Human mammary gland ductal carcinoma melanocyte) whole cell lysates at 20 μg

Lane 2: RAW264.7 (Mouse Abelson murine leukemia virusinduced tumor macrophage) whole cell lysates at 15 μg

Lane 3 : PC-12 (Rat adrenal gland pheochromocytoma) whole cell lysates at 15 μg

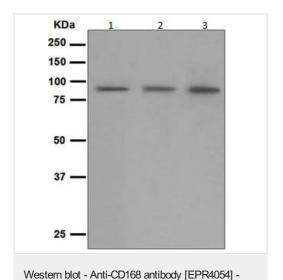
Secondary

All lanes : Goat Anti-Rabbit IgG H&L (HRP) (<u>ab97051</u>) at 1/20000 dilution

Predicted band size: 84 kDa

This data was developed using <u>ab124729</u>, the same antibody clone in a different buffer formulation:

5% NFDM/TBST



BSA and Azide free (ab271887)

All lanes: Anti-CD168 antibody [EPR4054] (ab124729)

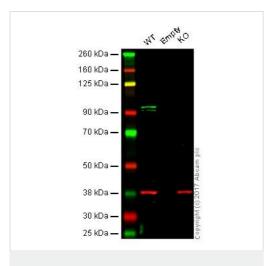
Lane 1: MDA-MB-435 cell lysate

Lane 2: LnCaP cell lysate

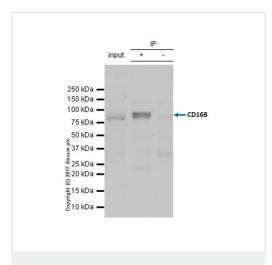
Lane 3: Sample: T47-D cell lysate

Lysates/proteins at 10 µg per lane.

Predicted band size: 84 kDa



Western blot - Anti-CD168 antibody [EPR4054] - BSA and Azide free (ab271887)



Immunoprecipitation - Anti-CD168 antibody [EPR4054] - BSA and Azide free (ab271887) This data was developed using <u>ab124729</u>, the same antibody clone in a different buffer formulation:

Lane 1: Wild type HAP1 whole cell lysate (20 µg)

Lane 2: Empty

Lane 3: CD168 knockout HAP1 whole cell lysate (20 µg)

Lanes 1 - 3: Merged signal (red and green). Green - <u>ab124729</u> observed at 90 kDa. Red - loading control, <u>ab9484</u>, observed at 37 kDa.

ab124729 was shown to specifically react with CD168 when CD168 knockout samples were used. Wild-type and CD168 knockout samples were subjected to SDS-PAGE. Ab124729 and ab9484 (Mouse anti-GAPDH loading control) were incubated overnight at 4°C at a 1/1000 dilution and 1/10000 dilution respectively. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preabsorbed ab216773 and Goat anti-Mouse IgG H&L (IRDye® 680RD) preabsorbed ab216776 secondary antibodies at 1/20000 dilution for 1 hour at room temperature before imaging.

<u>ab124729</u> (purified) at 1:100 dilution (2ug) immunoprecipitating CD168 in MCF-7 (Human breast adenocarcinoma epithelial cell) whole cell lysate.

Lane 1 (input): MCF-7 (Human breast adenocarcinoma epithelial cell) whole cell lysate 10ug

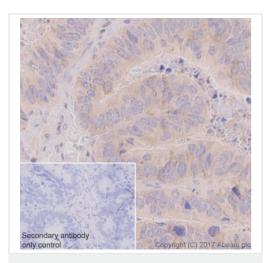
Lane 2 (+): <u>ab124729</u> & MCF-7 (Human breast adenocarcinoma epithelial cell) whole cell lysate

Lane 3 (-): Rabbit monoclonal IgG (<u>ab172730</u>) instead of <u>ab124729</u> in MCF-7 (Human breast adenocarcinoma epithelial cell) whole cell lysate

For western blotting, VeriBlot for IP Detection Reagent (HRP) (ab131366) was used for detection at 1:1000 dilution.

Blocking and diluting buffer: 5% NFDM/TBST.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab124729).

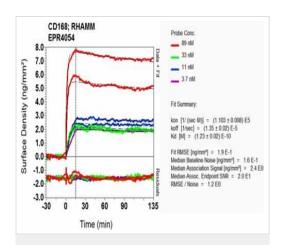


Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD168 antibody

[EPR4054] - BSA and Azide free (ab271887)

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) analysis of Human gastric carcinoma tissue sections labeling CD168 with purified ab124729 at 1:250 dilution (7.7 µg/ml). Heat mediated antigen retrieval was performed using EDTA Buffer, pH9.0. Tissue was counterstained with Hematoxylin. ImmunoHistoProbe one step HRP Polymer (ready to use) secondary antibody was used at 1:0 dilution. PBS instead of the primary antibody was used as the negative control.

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab124729).

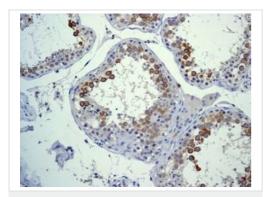


Ol-RD Scanning - Anti-CD168 antibody [EPR4054] - BSA and Azide free (ab271887)

Equilibrium disassociation constant (K_D) Learn more about K_D

Click here to learn more about KD

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab124729).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD168 antibody

[EPR4054] - BSA and Azide free (ab271887)

Immunohistochemical analysis of paraffin-embedded human testis tissue using unpurified <u>ab124729</u> at 1/50 - 1/100 dilution

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab124729).



Anti-CD168 antibody [EPR4054] - BSA and Azide free (ab271887)

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