abcam

Product datasheet

Anti-CD168 antibody [EPR4054] ab124729





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Overview

Product name Anti-CD168 antibody [EPR4054]

Description Rabbit monoclonal [EPR4054] to CD168

Host species Rabbit

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

Unsuitable for: Flow Cyt or ICC/IF

Species reactivity Reacts with: Mouse, Rat, Human

Immunogen Synthetic peptide within Human CD168 aa 1-100. The exact sequence is proprietary.

Positive control Lysates of: MDA-MB-435, LnCaP, T47-D, RAW264.7, PC-12. Human testis tissue

General notes 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**.

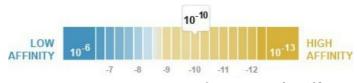
Properties

Form Liquid

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

Avoid freeze / thaw cycle. Stable for 12 months at -20°C.

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



Learn more about K_D

Storage buffer pH: 7.20

Preservative: 0.01% Sodium azide

Constituents: 40% Glycerol, 0.05% BSA, 59% PBS

Purity Protein A purified

Clonality Monoclonal
Clone number EPR4054

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab124729 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>(1)</u>	1/1000. Predicted molecular weight: 84 kDa. For unpurified use at 1/1000 - 1/10000.
		The mouse and rat recommendation is based on the WB results. This antibody may not be suitable for IHC with mouse or rat samples.
IP		1/10 - 1/100.
IHC-P		1/250. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol. For unpurified use at 1/50 - 1/100. Antigen retrieval is recommended

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

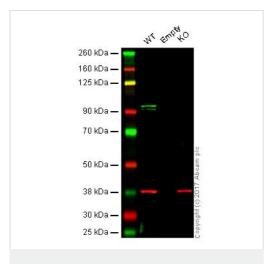
transformation and metastasis formation, and in regulating extracellular-regulated kinase (ERK)

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] (ab124729)

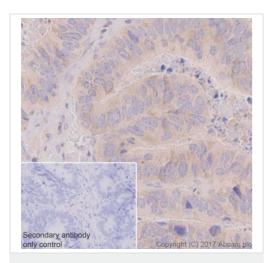
Lane 1: Wild type HAP1 whole cell lysate (20 µg)

Lane 2: Empty

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

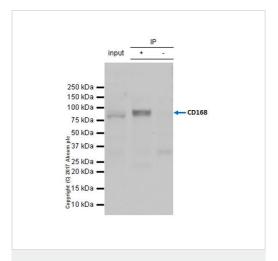
Lanes 1 - 3: Merged signal (red and green). Green - ab124729 observed at 90 kDa. Red - loading control, <u>ab9484</u>, observed at 37 kDa.

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.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD168 antibody
[EPR4054] (ab124729)

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.



Immunoprecipitation - Anti-CD168 antibody [EPR4054] (ab124729)

ab124729 (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

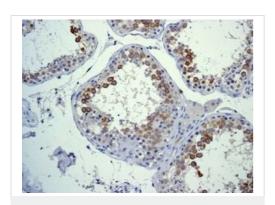
Lane 2 (+): ab124729 & MCF-7 (Human breast adenocarcinoma epithelial cell) whole cell lysate

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

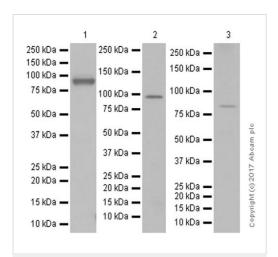
For western blotting, VeriBlot for IP Detection Reagent (HRP) (ab131366) was used for detection at 1:1000 dilution.

Blocking and diluting buffer: 5% NFDM/TBST.

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



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD168 antibody
[EPR4054] (ab124729)



Western blot - Anti-CD168 antibody [EPR4054] (ab124729)

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

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 virus-induced 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

5% NFDM/TBST

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

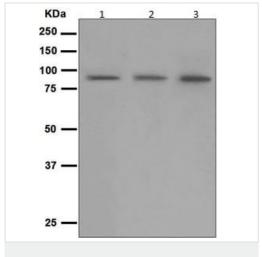
Lane 1: MDA-MB-435 cell lysate

Lane 2 : LnCaP cell lysate

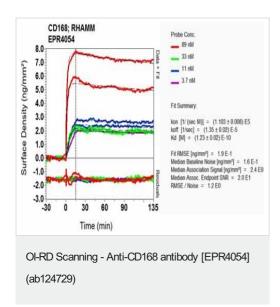
Lane 3 : T47-D cell lysate

Lysates/proteins at 10 µg per lane.

Predicted band size: 84 kDa



Western blot - Anti-CD168 antibody [EPR4054] (ab124729)



Equilibrium disassociation constant (K_D) Learn more about K_D

Click here to learn more about Kn



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