abcam

Product datasheet

Anti-CD2 antibody [EPR6451] - BSA and Azide free ab248400

Recombinant RabMAb

7 Images

Overview

Product name Anti-CD2 antibody [EPR6451] - BSA and Azide free

Description Rabbit monoclonal [EPR6451] to CD2 - BSA and Azide free

Host species Rabbit

Tested applications Suitable for: WB, IHC-P, IP, Flow Cyt (Intra)

Species reactivity Reacts with: Human

Immunogen Synthetic peptide within Human CD2 aa 200-350. The exact sequence is proprietary.

Database link: P06729

Positive control WB: Human Jurkat cell lysates. IHC: Human tonsil, colon and spleen tissue. IP: Jurkat lysate. Flow

cyto: Jurkat (Human T cell leukemia T lymphocyte) cells

General notes ab248400 is the carrier-free version of ab131276.

> Our carrier-free 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 cellbased assays, flow-based assays (e.g. mass cytometry) and Multiplex Imaging applications.

Use our conjugation kits 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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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 +4°C. Do Not Freeze.

Storage buffer pH: 7.2

Constituent: PBS

Carrier free Yes

Purity Protein A purified

Clonality Monoclonal
Clone number EPR6451

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab248400 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: 39 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.
IP		Use at an assay dependent concentration.
Flow Cyt (Intra)		Use at an assay dependent concentration.

Target

Function CD2 interacts with lymphocyte function-associated antigen (LFA-3) and CD48/BCM1 to mediate

adhesion between T-cells and other cell types. CD2 is implicated in the triggering of T-cells, the

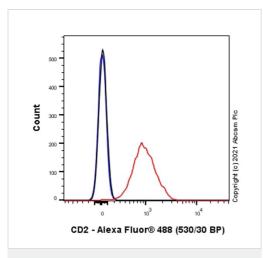
cytoplasmic domain is implicated in the signaling function.

Sequence similarities Contains 1 lg-like C2-type (immunoglobulin-like) domain.

Contains 1 lg-like V-type (immunoglobulin-like) domain.

Cellular localization Membrane.

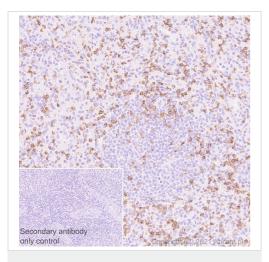
Images



Flow Cytometry (Intracellular) - Anti-CD2 antibody [EPR6451] - BSA and Azide free (ab248400)

This data was developed using <u>ab131276</u>, the same antibody clone in a different buffer formulation.

Intracellular Flow Cytometry analysis of Jurkat (Human T cell leukemia T lymphocyte) cells labeling CD2 with purified **ab131276** at 1:20 dilution (10 µg/ml) (Red). Cells were fixed with 4% Paraformaldehyde and permeabilised with 90% Methanol. A Goat anti rabbit lgG (Alexa Fluor® 488, **ab150081**) (1/2000 dilution) was used as the secondary antibody. Rabbit monoclonal lgG (Black) was used as a isotype control. Cell without incubation with primary antibody and secondary antibody (Blue) were used as unlabeled control.



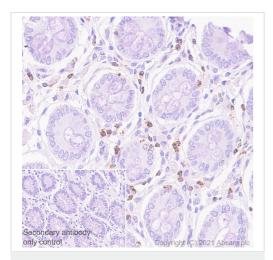
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD2 antibody [EPR6451] - BSA and Azide free (ab248400)

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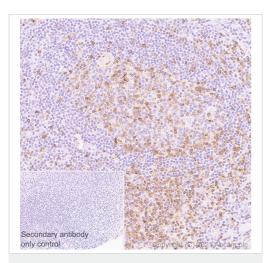
Immunohistochemical analysis of Paraffin-embedded sections
Human spleen tissue labelling CD2 with <u>ab131276</u> at 1/1000
dilution, followed by a ready to use secondary Rabbit specific IHC
polymer detection kit HRP/DAB (<u>ab209101</u>). Staining on Human
spleen tissue is observed. Counter stained with Haematoxylin.
Secondary antibody only control: Used PBS instead of primary
antibody, secondary antibody is ready to use Rabbit specific IHC
polymer detection kit HRP/DAB (<u>ab209101</u>).

Heat mediated antigen retrieval using Bond™ Epitope Retrieval Solution 2 (pH 9.0).

The immunostaining was performed on a Leica Biosystems BOND® RX instrument.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-CD2 antibody [EPR6451] - BSA and Azide free (ab248400)



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This data was developed using <u>ab131276</u>, the same antibody clone in a different buffer formulation.

Immunohistochemical analysis of Paraffin-embedded sections
Human colon tissue labelling CD2 with <u>ab131276</u> at 1/1000
dilution, followed by a ready to use secondary Rabbit specific IHC
polymer detection kit HRP/DAB (<u>ab209101</u>). Staining on Human
colon tissue is observed. Counter stained with Haematoxylin.
Secondary antibody only control: Used PBS instead of primary
antibody, secondary antibody is ready to use Rabbit specific IHC
polymer detection kit HRP/DAB (<u>ab209101</u>).

Heat mediated antigen retrieval using Bond™ Epitope Retrieval Solution 2 (pH 9.0).

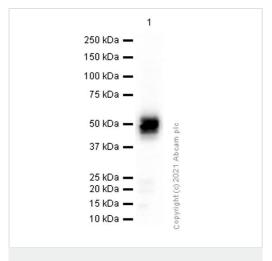
The immunostaining was performed on a Leica Biosystems BOND® RX instrument.

This data was developed using <u>ab131276</u>, the same antibody clone in a different buffer formulation.

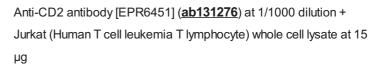
Immunohistochemical analysis of Paraffin-embedded sections
Human tonsil tissue labelling CD2 with <u>ab131276</u> at 1/1000
dilution, followed by a ready to use secondary Rabbit specific IHC
polymer detection kit HRP/DAB (<u>ab209101</u>). Staining on Human
tonsil tissue is observed. Counter stained with Haematoxylin.
Secondary antibody only control: Used PBS instead of primary
antibody, secondary antibody is ready to use Rabbit specific IHC
polymer detection kit HRP/DAB (<u>ab209101</u>).

Heat mediated antigen retrieval using Bond™ Epitope Retrieval Solution 2 (pH 9.0).

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Western blot - Anti-CD2 antibody [EPR6451] - BSA and Azide free (ab248400)



Secondary

Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/20000 dilution

Predicted band size: 39 kDa **Observed band size:** 45-58 kDa

This data was developed using <u>ab131276</u>, the same antibody clone in a different buffer formulation.

Blocking and diluting buffer and concentration: 5% NFDM/TBST.

The molecular weight observed is consistent with what has been described in the literature (PMID: 10510361).

Immunoprecipitation - Anti-CD2 antibody [EPR6451] - BSA and Azide free (ab248400)

This data was developed using <u>ab131276</u>, the same antibody clone in a different buffer formulation.

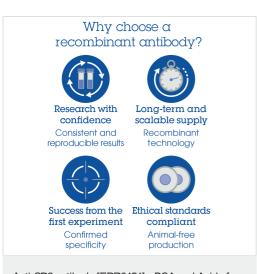
CD2 was immunoprecipitated from 0.35 mg Jurkat (Human T cell leukemia T lymphocyte) whole cell lysate 10 μ g with <u>ab131276</u> at 1/50 dilution (2 μ g). VeriBlot for IP Detection Reagent (HRP) (<u>ab131366</u>) was used at 1/5000 dilution.

Lane 1: Jurkat (Human T cell leukemia T lymphocyte) whole cell lysate 10 μg

Lane 2: abab131276 IP in Jurkat whole cell lysate

Lane 3: Rabbit monoclonal lgG ($\underline{ab172730}$) instead of $\underline{ab131276}$ in Jurkat whole cell lysate

Blocking and dilution buffer and concentration: 5% NFDM/TBST.



Anti-CD2 antibody [EPR6451] - BSA and Azide free (ab248400)

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