abcam

Product datasheet

Anti-DPP4 antibody [EPR18215] - BSA and Azide free ab232221



5 Images

Overview

Product name Anti-DPP4 antibody [EPR18215] - BSA and Azide free

Description Rabbit monoclonal [EPR18215] to DPP4 - BSA and Azide free

Host species Rabbit

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

Species reactivity Reacts with: Mouse, Rat

Immunogen Recombinant fragment. This information is proprietary to Abcam and/or its suppliers.

Positive control IHC-P: Mouse liver tissue.

General notes ab232221 is the carrier-free version of ab187048.

> 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**.

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 EPR18215

Isotype IgG

Applications

The Abpromise guarantee

Our Abpromise quarantee covers the use of ab232221 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. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
WB		Use at an assay dependent concentration. Detects a band of approximately 110 kDa (predicted molecular weight: 88 kDa).
IHC-Fr		Use at an assay dependent concentration.

Target

Function

Cell surface glycoprotein receptor involved in the costimulatory signal essential for T-cell receptor (TCR)-mediated T-cell activation. Acts as a positive regulator of T-cell coactivation, by binding at least ADA, CAV1, IGF2R, and PTPRC. Its binding to CAV1 and CARD11 induces T-cell proliferation and NF-kappa-B activation in a T-cell receptor/CD3-dependent manner. Its interaction with ADA also regulates lymphocyte-epithelial cell adhesion. In association with FAP is involved in the pericellular proteolysis of the extracellular matrix (ECM), the migration and invasion of endothelial cells into the ECM. May be involved in the promotion of lymphatic endothelial cells adhesion, migration and tube formation. When overexpressed, enhanced cell proliferation, a process inhibited by GPC3. Acts also as a serine exopeptidase with a dipeptidyl peptidase activity that regulates various physiological processes by cleaving peptides in the circulation, including many chemokines, mitogenic growth factors, neuropeptides and peptide hormones. Removes N-terminal dipeptides sequentially from polypeptides having unsubstituted N-termini provided that the penultimate residue is proline.

Tissue specificity

Expressed specifically in lymphatic vessels but not in blood vessels in the skin, small intestine, esophagus, ovary, breast and prostate glands. Not detected in lymphatic vessels in the lung, kidney, uterus, liver and stomach (at protein level). Expressed in the poorly differentiated crypt cells of the small intestine as well as in the mature villous cells. Expressed at very low levels in the

colon.

Sequence similarities

Belongs to the peptidase S9B family. DPPIV subfamily.

Domain

The extracellular cysteine-rich region is necessary for association with collagen, dimer formation and optimal dipeptidyl peptidase activity.

Post-translational modifications

The soluble form (Dipeptidyl peptidase 4 soluble form also named SDPP) derives from the membrane form (Dipeptidyl peptidase 4 membrane form also named MDPP) by proteolytic processing.

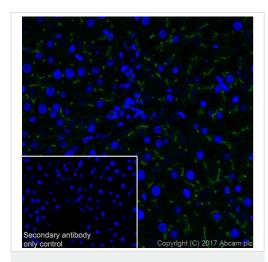
N- and O-Glycosylated.

Phosphorylated. Mannose 6-phosphate residues in the carbohydrate moiety are necessary for interaction with IGF2R in activated T-cells. Mannose 6-phosphorylation is induced during T-cell activation.

Cellular localization

Cell membrane. Apical cell membrane. Cell projection > invadopodium membrane. Cell projection > lamellipodium membrane. Cell junction. Membrane raft. Translocated to the apical membrane through the concerted action of N- and O-Glycans and its association with lipid microdomains containing cholesterol and sphingolipids. Redistributed to membrane rafts in T-cell in a interleukin-12-dependent activation. Its interaction with CAV1 is necessary for its translocation to membrane rafts. Colocalized with PTPRC in membrane rafts. Colocalized with FAP in invadopodia and lamellipodia of migratory activated endothelial cells in collagenous matrix. Colocalized with FAP on endothelial cells of capillary-like microvessels but not large vessels within invasive breast ductal carcinoma. Colocalized with ADA at the cell junction in lymphocyte-epithelial cell adhesion. Colocalized with IGF2R in internalized cytoplasmic vesicles adjacent to the cell surface and Secreted. Detected in the serum and the seminal fluid.

Images

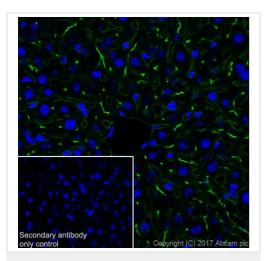


Immunohistochemistry (Frozen sections) - Anti-DPP4 antibody [EPR18215] - BSA and Azide free (ab232221) Immunohistochemical analysis of 4% PFA-fixed, 0.2 % Triton X-100 permeabilized rat liver tissue labeling CD26 with **ab187048** at 1/200 dilution, followed by Goat Anti-Rabbit lgG H&L (Alexa Fluor® 488) (**ab150077**) at 1/1000 dilution (green). Strong positive staining on the bile canaliculi and moderate positive staining in sinusoid endothelial cells is observed (PMID: 23361237). The nuclear counter stain is DAPI (blue)

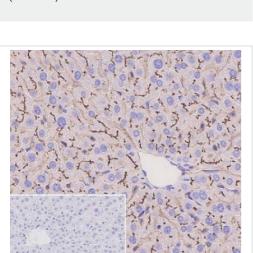
Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (Alexa Fluor® 488) (ab150077) at 1/1000 dilution.

Antigen retrieval: Heated Tris-EDTA (pH9.0) (ab94681).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab187048).



Immunohistochemistry (Frozen sections) - Anti-DPP4 antibody [EPR18215] - BSA and Azide free (ab232221)



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-DPP4 antibody

[EPR18215] - BSA and Azide free (ab232221)

Immunohistochemical analysis of 4% PFA-fixed, 0.2 % Triton X-100 permeabilized mouse liver tissue labeling CD26 with <u>ab187048</u> at 1/1000 dilution, followed by Goat Anti-Rabbit lgG H&L (Alexa Fluor® 488) (<u>ab150077</u>) at 1/1000 dilution (green). Strong positive staining on the bile canaliculi and moderate positive staining in sinusoid endothelial cells is observed (PMID: 23361237). The nuclear counter stain is DAPI (blue)

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit lgG H&L (Alexa Fluor® 488) (ab150077) at 1/1000 dilution.

Antigen retrieval: Heated Tris-EDTA (pH9.0) (ab94681).

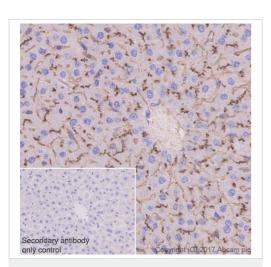
This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab187048).

Immunohistochemical analysis of paraffin-embedded rat liver tissue labeling CD26 with <u>ab187048</u> at 1/2000 dilution, followed by Goat Anti-Rabbit lgG H&L (HRP) Ready to use. Membranous staining on bile canaliculi of rat liver is observed (PMID: 23361237). Counter stained with hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) Ready to use.

Perform heat mediated antigen retrieval using <u>ab93684</u> (Tris/EDTA buffer, pH 9.0).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (<u>ab187048</u>).



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-DPP4 antibody

[EPR18215] - BSA and Azide free (ab232221)

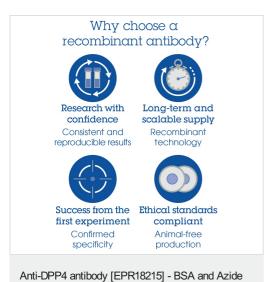
Immunohistochemical analysis of paraffin-embedded mouse liver tissue labeling CD26 with <u>ab187048</u> at 1/2000 dilution, followed by Goat Anti-Rabbit IgG H&L (HRP) Ready to use.

Membranous staining on bile canaliculi of mouse liver is observed (PMID: 23361237). Counter stained with hematoxylin.

Secondary antibody only control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) Ready to use.

Perform heat mediated antigen retrieval using <u>ab93684</u> (Tris/EDTA buffer, pH 9.0).

This data was developed using the same antibody clone in a different buffer formulation containing PBS, BSA, glycerol, and sodium azide (ab187048).



free (ab232221)

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