abcam

ab300526

Anti-EPCR/CD201 antibody [EPR25647-5] - BSA and Azide free

Recombinant | RabMAb

Key facts

Isotype	IgG
Host species	Rabbit
Storage buffer	pH: 7.2 - 7.4 Constituents: PBS
Form	Liquid
Clonality	Monoclonal
Immunogen	The exact immunogen used to generate this antibody is proprietary information.
Carrier free	Yes
Clone number	EPR25647-5
Purification technique	Affinity purification Protein A
Concentration	1.05 mg/mL The concentration of this product may be batch-dependent Batch concentration finder →

Reactivity data

WB

Tested

Species	Human
Dilution info	-

Notes	-

Not recommended

Species	Mouse, Rat
Dilution info	-
Notes	-

IHC-P

Not recommended

Species	Human, Mouse, Rat
Dilution info	-
Notes	-

ICC/IF

Tested

Species	Human
Dilution info	-
Notes	-

Not recommended

Species	Mouse, Rat
Dilution info	-
Notes	-

Flow Cyt

Tested

Species	Human
Dilution info	-
Notes	-

Not recommended

Species	Mouse, Rat
Dilution info	-
Notes	-

IP

Tested

Species	Human
Dilution info	-
Notes	-

Not recommended

Species	Mouse, Rat
Dilution info	-
Notes	-

Target data

See full target information PROCR

Function	Binds activated protein C. Enhances protein C activation by the thrombin-thrombomodulin complex; plays a role in the protein C pathway controlling blood coagulation.

Storage

Shipped at conditions	Blue Ice
Appropriate short-term storage conditions	+4°C
Appropriate long-term storage conditions	+4°C
Aliquoting information	Upon delivery aliquot

Storage information	Do Not Freeze

Notes

Patented technology
Our RabMAb® technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to RabMAb® patents.

What are the advantages of a recombinant monoclonal antibody?
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 batch production

For more information, read more on recombinant antibodies.

Conjugation ready
Our carrier-free antibodies are typically supplied in a PBS-only formulation, purified and free of BSA, sodium azide and glycerol. 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 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.

Compatibility
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.

Supplementary info

This supplementary information is collated from multiple sources and compiled automatically.

Activity summary	EPCR also known as CD201 stands for Endothelial Protein C Receptor. It is a glycoprotein with a molecular mass of approximately 46 kDa. EPCR is expressed mainly on endothelial cells which line the interior surface of blood vessels. This receptor plays a mechanical role in binding to protein C a zymogen that is converted to activated protein C (APC) by thrombin-thrombomodulin complex. By facilitating the activation of protein C EPCR contributes to the regulation of blood coagulation and inflammation processes.
Biological function summary	The EPCR is part of the protein C anticoagulant pathway where it acts as a cofactor. EPCR is vital in enhancing the activation of protein C on the endothelial surface reducing coagulation activity. It interacts closely with thrombomodulin and protein C to form this complex preventing excessive thrombus formation and contributing to vascular integrity. Apart from its anticoagulant role EPCR also modulates inflammatory responses supporting its function as a bridge between coagulation and inflammation.

Pathways	EPCR lies within the protein C anticoagulation pathway and the endothelial signaling pathways. Its primary partner in the anticoagulation pathway is APC which degrades activated factors Va and VIIIa providing a check against coagulation. Additionally EPCR's interaction with APC and thrombomodulin links this protein to the endothelial signaling pathway where it affects cell survival and inflammation regulation. This interaction is essential to maintain hemostatic balance and control vascular responses to inflammatory stimuli.
Associated diseases and disorders	EPCR associates with conditions like thrombosis and sepsis. In thrombosis inadequate expression or function of EPCR can lead to uncontrolled blood clotting due to insufficient activation of protein C. Sepsis a severe inflammatory response to infection shows abnormal modulation of the protein C pathway where EPCR's role becomes pivotal. Additionally APC which interacts directly with EPCR is critical in the pathological states of these diseases highlighting the therapeutic potential in targeting EPCR pathway.

Product promise

Tested

We have tested this species and application combination and it works. It is covered by our product promise.

Expected

We have not tested this specific species and application combination in-house, but expect it will work. It is covered by our product promise.

Predicted

This species and application combination has not been tested, but we predict it will work based on strong homology. However, this combination is not covered by our product promise.

Not recommended

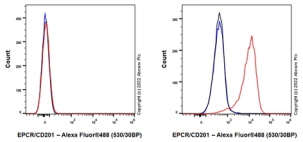
We do not recommend this combination. It is not covered by our product promise.

We are dedicated to supporting your work with high quality reagents and we are here for you every step of the way should you need us.

In the unlikely event of one of our products not working as expected, you are covered by our product promise.

Full details and terms and conditions can be found here:
Terms & Conditions.

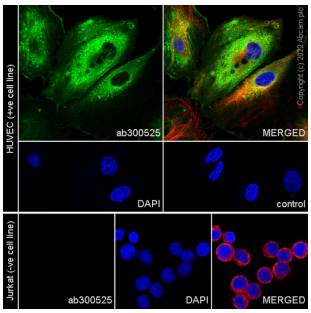
6 product images



Flow Cytometry - Anti-EPCR/CD201 antibody [EPR25647-5] - BSA and Azide free (ab300526)

EPCR/CD201 Flow Cytometry staining using rabbit Anti-EPCR/CD201 antibody

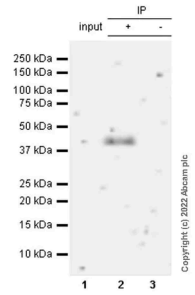
This data was developed using ab300525, the same antibody clone in a different buffer formulation.Flow cytometric analysis of Jurkat (human T cell leukemia T lymphocyte, Left) / HUVEC (human umbilical vein endothelial cell, Right) cells labelling EPCR/CD201 with ab300525 at 1/500 dilution (0.1ug) (Red) compared with a Rabbit monoclonal IgG (ab172730) (Black) isotype control and an unlabelled control (cells without incubation with primary antibody and secondary antibody) (Blue). Goat Anti-Rabbit IgG (Alexa Fluor® 488, ab150081) at 1/2000 dilution was used as the secondary antibody. Gated on viable cells.Negative control: Jurkat (PMID: 11703343).



Immunocytochemistry/ Immunofluorescence - Anti-EPCR/CD201 antibody [EPR25647-5] - BSA and Azide free (ab300526)

EPCR/CD201 Immunocytochemistry/ Immunofluorescence staining using rabbit Anti-EPCR/CD201 antibody

This data was developed using ab300525, the same antibody clone in a different buffer formulation.Immunofluorescent analysis of 4% Paraformaldehyde-fixed, 0.1% TritonX-100 permeabilized HUVEC (human umbilical vein endothelial cell) cells labelling EPCR/CD201 with ab300525 at 1/50 (9.8 ug/ml) dilution, followed by ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed antibody at 1/1000 2 ug/ml dilution (Green). Confocal image showing cytoplasmic and membranous staining in HUVEC cell lineNegative control: Jurkat (PMID: 11703343) is observed. ab195889 Anti-alpha Tubulin mouse monoclonal antibody - Microtubule Marker (Alexa Fluor® 594) was used to counterstain tubulin at 1/200 2.5 ug/ml dilution (Red). The Nuclear counterstain was DAPI (Blue). Secondary antibody only control: Secondary antibody is ab150081 Goat Anti-Rabbit IgG H&L (Alexa Fluor® 488) preadsorbed at 1/1000 2 ug/ml dilution.



Immunoprecipitation - Anti-EPCR/CD201 antibody [EPR25647-5] - BSA and Azide free (ab300526)

This data was developed using ab300526, the same antibody clone in a different buffer formulation. EPCR/CD201 was immunoprecipitated from 0.35 mg HUVEC (human umbilical vein endothelial cell) whole cell lysate 10 ug with ab300525 at 1/30 dilution (2ug in 0.35mg lysates). Western blot was performed on the immunoprecipitate using ab300525 at 1/1000 dilution. VeriBlot for IP secondary antibody (HRP)(ab131366) was used at 1/1000 dilution. Lane 1: HUVEC (human umbilical vein endothelial cell) whole cell lysate 10 ug Lane 2: ab300525 IP in HUVEC whole cell lysate Lane 3: Rabbit monoclonal IgG (ab172730) instead of ab300525 in HUVEC whole cell lysate. Blocking and dilution buffer and concentration: 5% NFDM/TBST. Exposure time: 3 minutes
Lane 1:
Immunoprecipitation - Anti-EPCR/CD201 antibody [EPR25647-5] (ab300525) at 1/30 dilution

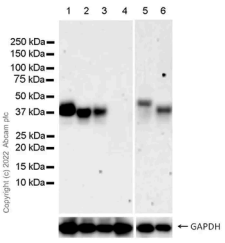
Lane 2:
Immunoprecipitation - Anti-EPCR/CD201 antibody [EPR25647-5] (ab300525) at 1/1000 dilution

All lanes:
HUVEC whole cell lysate at 10 µg

Secondary

All lanes:
Immunoprecipitation - VeriBlot for IP Detection Reagent (HRP) (ab131366) at 1/5000 dilution

Exposure time: 3min



Western blot - Anti-EPCR/CD201 antibody [EPR25647-5] - BSA and Azide free (ab300526)

EPCR/CD201 Western blot staining using rabbit Anti-EPCR/CD201 antibody

This data was developed using ab300525, the same antibody clone in a different buffer formulation.

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

Negative control: Jurkat (PMID: 7929370)

Exposure time: Lanes 1-4: 103 seconds; Lanes 5-6: 3 minutes

All lanes:
Western blot - Anti-EPCR/CD201 antibody [EPR25647-5] (ab300525) at 1/1000 dilution

Lane 1:
HUVEC (human umbilical vein endothelial cell), whole cell lysate at 10 µg

Lane 2:
Ea.Hy926 (human somatic cell hybrid endothelial), whole cell lysate at 10 µg

Lane 3:
RKO (human colon epithelial cell), whole cell lysate at 10 µg

Lane 4:
Jurkat (human T cell leukemia T lymphocyte), whole cell lysate at 10 µg

Lane 5:
Human placenta tissue lysate at 10 µg

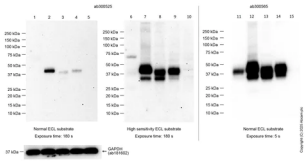
Lane 6:
Human lung tissue lysate at 10 µg

Secondary

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

Predicted band size: 27 kDa

Observed band size: 45 kDa



Western blot - Anti-EPCR/CD201 antibody [EPR25647-5] - BSA and Azide free (ab300526)

EPCR/CD201 Western blot staining using rabbit Anti-EPCR/CD201 antibody

This data was developed using ab300525, the same antibody clone in a different buffer formulation.
Exposure time: ab300525: 180 seconds; ab300565: 5 seconds

In Western blot, Anti-GAPDH antibody [EPR16891] - Loading Control (ab181602) staining at 1/1000000 dilution.

The band beneath the target band is supposed to be a soluble form (PMID: 19806250).

ab300525 is less sensitive than ab300565. ab300525 needs a high sensitivity ECL substrate.

Lanes 1 - 15:
Western blot - Anti-EPCR/CD201 antibody [EPR25647-5] (ab300525) at 1/1000 dilution

Lanes 11 - 15:
Western blot - Anti-EPCR/CD201 antibody [EPR25647-49] (ab300565) at 1/1000 dilution

Lanes 1, 6 and 11:
Human lung tissue lysate at 20 µg

Lanes 2, 7 and 12:
HUVEC (Human umbilical vein endothelial cell) whole cell lysate at 20 µg

Lanes 3, 8 and 13:
Ea.Hy926 (human somatic cell hybrid endothelial) whole cell lysate at 20 µg

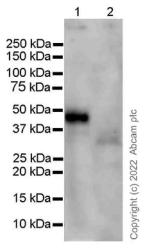
Lanes 4, 9 and 14:
RKO (human colorectal carcinoma cell) whole cell lysate at 20 µg

Lanes 5, 10 and 15:
Jurkat (Human T cell leukemia cell line from peripheral blood) whole cell lysate at 20 µg

Secondary

All lanes:
Goat Anti-Rabbit IgG (HRP) with minimal cross-reactivity with human IgG at 1/2000 dilution

Observed band size: 45 kDa



Western blot - Anti-EPCR/CD201 antibody [EPR25647-5] - BSA and Azide free (ab300526)

This data was developed using ab300525, the same antibody clone in a different buffer formulation.
Blocking and diluting buffer and concentration: 5% NFDM/TBST

This blot was developed using a high sensitivity ECL substrate. The high-sensitivity ECL substrate used allows for the detection of proteins in the mid-femtogram range.

All lanes:
Western blot - Anti-EPCR/CD201 antibody [EPR25647-5] (ab300525) at 1/1000 dilution

Lane 1:
Untreated HUVEC whole cell lysate at 10 µg

Lane 2:
HUVEC whole cell lysate treated with Protein Deglycosylation Mix II at 10 µg

Secondary

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

Developed using the ECL technique.

Predicted band size: 27 kDa

Observed band size: 45 kDa

Exposure time: 3min

Please note: All products are 'FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC OR THERAPEUTIC PROCEDURES'.