abcam

Product datasheet

Anti-PPP2R5E antibody [EPR17147] - C-terminal ab198500



Recombinant

RabMAb

9 Images

Overview

Product name Anti-PPP2R5E antibody [EPR17147] - C-terminal

Description Rabbit monoclonal [EPR17147] to PPP2R5E - C-terminal

Host species Rabbit

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

Species reactivity Reacts with: Human

Predicted to work with: Mouse, Rat

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

Positive control WB: HeLa, HEK-293T and Daudi cell lysates; Human skeletal muscle lysates. IHC: Human cervix

carcinoma and bladder transitional cell carcinoma tissues. ICC/IF: HeLa cells. Flow Cyt (intra):

HeLa cell lysate. IP: HeLa cell lysate.

General notesThis product is a recombinant monoclonal antibody, which offers several advantages including:

- High batch-to-batch consistency and reproducibility

Improved sensitivity and specificityLong-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 long

term. Avoid freeze / thaw cycle.

Storage buffer pH: 7.2

Preservative: 0.01% Sodium azide

Constituents: 40% Glycerol (glycerin, glycerine), 59% PBS, 0.05% BSA

Purity Protein A purified

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ClonalityMonoclonalClone numberEPR17147

Isotype IgG

Applications

The Abpromise guarantee Our **Abpromise guarantee** covers the use of ab198500 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		1/5000. Detects a band of approximately 55 kDa (predicted molecular weight: 55 kDa).
IP		1/75.
ICC/IF		1/500.
IHC-P		1/500. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.
Flow Cyt (Intra)		1/2500.

Target

Function The B regulatory subunit might modulate substrate selectivity and catalytic activity, and also might

direct the localization of the catalytic enzyme to a particular subcellular compartment.

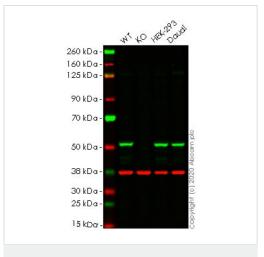
Sequence similaritiesBelongs to the phosphatase 2A regulatory subunit B56 family.

Post-translational modifications

Phosphorylated on serine residues.

Cellular localization Cytoplasm.

Images



Western blot - Anti-PPP2R5E antibody [EPR17147] - C-terminal (ab198500)

All lanes : Anti-PPP2R5E antibody [EPR17147] - C-terminal (ab198500) at 1/5000 dilution

Lane 1: Wild-type HeLa cell lysate

Lane 2: PPP2R5E knockout HeLa cell lysate

Lane 3: HEK-293T cell lysate

Lane 4: Daudi cell lysate

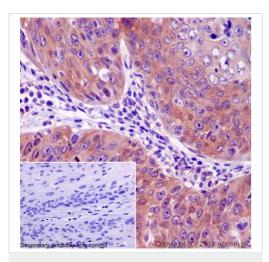
Lysates/proteins at 20 µg per lane.

Performed under reducing conditions.

Predicted band size: 55 kDa

Lanes 1-4: Merged signal (red and green). Green - ab198500 observed at 55 kDa. Red - loading control **ab8245** observed at 37 kDa.

ab198500 Recombinant Anti-PPP2R5E antibody [EPR17147] - Cterminal was shown to specifically react with PPP2R5E in wild-type HeLa cells. Loss of signal was observed when knockout cell line ab265637 (knockout cell lysate ab258135) was used. Wild-type and PPP2R5E knockout samples were subjected to SDS-PAGE. ab198500 and Anti-GAPDH antibody [6C5] - Loading Control (ab8245) were incubated overnight at 4°C at 1 in 5000 dilution and 1 in 20000 dilution respectively. Blots were developed with Goat anti-Rabbit IgG H&L (IRDye® 800CW) preadsorbed (ab216773) and Goat anti-Mouse IgG H&L (IRDye® 680RD) preadsorbed (ab216776) secondary antibodies at 1 in 20000 dilution for 1 hour at room temperature before imaging.



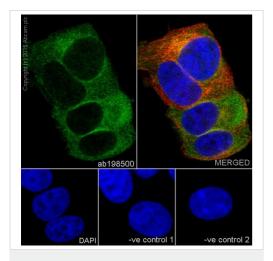
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-PPP2R5E antibody

[EPR17147] - C-terminal (ab198500)

Immunohistochemical analysis of paraffin-embedded Human cervix carcinoma tissue labeling PPP2R5E with ab198500 at 1/500 dilution followed by Goat Anti-Rabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Cytoplasm staining on Human cervix carcinoma tissue isobserved. Counter-stained with Hematoxylin.

Negative control: Using PBS instead of primary antibody.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol.

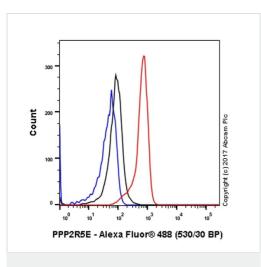


Immunocytochemistry/ Immunofluorescence - Anti-PPP2R5E antibody [EPR17147] - C-terminal (ab198500)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized MCF-7 (Human breast carcinoma) cells labeling PPP2R5E with ab198500 at 1/500 dilution, followed by Goat anti-rabbit IgG (Alexa Fluor® 488) (ab150077) secondary antibody at 1/500 dilution (green). Cytoplasm staining on MCF7 cell line is observed. The nuclear counter stain is DAPI (blue). Tubulin is detected with ab7291 (anti-Tubulin mouse mAb) at 1/1000 dilution and ab150120 (AlexaFluor®594 Goat anti-Mouse secondary) at 1/500 dilution (red).

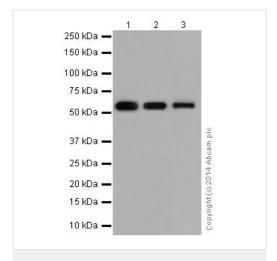
The negative controls are as follows;

- 1. ab198500 at 1/500 dilution followed by <u>ab150120</u> (AlexaFluor®594 Goat anti-Mouse secondary) at 1/500 dilution.
- 2. <u>ab7291</u> (anti-Tubulin mouse mAb) at 1/1000 dilution followed by <u>ab150077</u> (Alexa Fluor®488 Goat Anti-Rabbit lgG H&L) at 1/500 dilution.



Flow Cytometry (Intracellular) - Anti-PPP2R5E antibody [EPR17147] - C-terminal (ab198500)

Intracellular Flow Cytometry analysis of HeLa (Human cervix adenocarcinoma epithelial cell) cells labeling PPP2R5E (red) with purified ab198500 at a 1/2500 dilution (1ug/mL). Cells were fixed with 4% paraformaldehyde and permeabilized with 90% methanol. A goat anti rabbit lgG (Alexa Fluorr® 488) (ab150077) was used as the secondary antibody at a 1/2000 dilution. Black - Rabbit monoclonal lgG (Black) (ab172730). Blue (unlabeled control) - Cell without incubation with primary antibody and secondary antibody (Blue).



Western blot - Anti-PPP2R5E antibody [EPR17147] - C-terminal (ab198500)

All lanes : Anti-PPP2R5E antibody [EPR17147] - C-terminal (ab198500) at 1/5000 dilution

Lane 1: HeLa cell lysate at 10 μg **Lane 2**: 293 cell lysate at 10 μg

Lane 3: Human skeletal muscle lysate at 20 µg

Secondary

All lanes : Goat Anti-Rabbit lgG, (H+L), Peroxidase conjugate at 1/1000 dilution

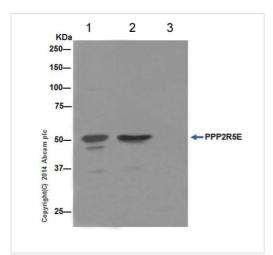
Developed using the ECL technique.

Predicted band size: 55 kDa

Observed band size: 55 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFDM/TBST.



Immunoprecipitation - Anti-PPP2R5E antibody [EPR17147] - C-terminal (ab198500)

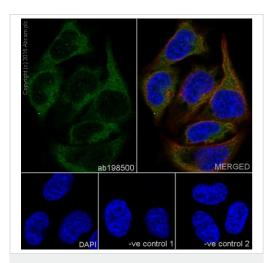
PPP2R5E was immunoprecipitated from 1mg of HeLa (Human cervix adenocarcinoma) whole cell extract with ab198500 at 1/175 dilution. Western blot was performed from the immunoprecipitate using ab198500 at 1/1000 dilution. Anti-Rabbit lgG (HRP), specific to the non-reduced form of lgG, was used as secondary antibody at 1/1500 dilution.

Lane 1: HeLa whole cell extract 10µg (Input).

Lane 2: HeLa whole cell extract

Lane 3: Rabbit monoclonal $\lg G$ ($\underline{ab172730}$) instead of ab198500 in HeLa whole cell extract.

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

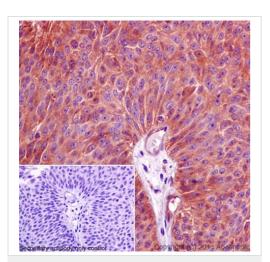


Immunocytochemistry/ Immunofluorescence - Anti-PPP2R5E antibody [EPR17147] - C-terminal (ab198500)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized HeLa (Human cervix adenocarcinoma) cells labeling PPP2R5E with ab198500 at 1/500 dilution, followed by Goat anti-rabbit lgG (Alexa Fluor® 488) (ab150077) secondary antibody at 1/500 dilution (green). Cytoplasm staining on HeLa cell line is observed. The nuclear counter stain is DAPI (blue). Tubulin is detected with ab7291 (anti-Tubulin mouse mAb) at 1/1000 dilution and ab150120 (AlexaFluor®594 Goat anti-Mouse secondary) at 1/500 dilution (red).

The negative controls are as follows;

ab198500 at 1/500 dilution followed by <u>ab150120</u>
 (AlexaFluor®594 Goat anti-Mouse secondary) at 1/500 dilution.
 <u>ab7291</u> (anti-Tubulin mouse mAb) at 1/1000 dilution followed by <u>ab150077</u> (Alexa Fluor®488 Goat Anti-Rabbit lgG H&L) at 1/500 dilution.



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-PPP2R5E antibody

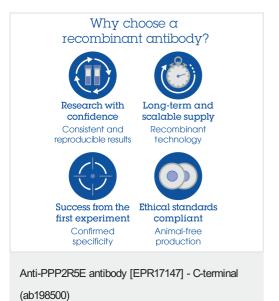
[EPR17147] - C-terminal (ab198500)

Immunohistochemical analysis of paraffin-embedded
Human transitional cell carcinoma of bladder tissue labeling
PPP2R5E with ab198500 at 1/500 dilution followed by Goat AntiRabbit IgG H&L (HRP) (ab97051) at 1/500 dilution. Cytoplasm
staining on Human transitional cell carcinoma of bladder tissue
isobserved. Counter-stained with Hematoxylin.

Negative control: Using PBS instead of primary antibody.

Perform heat mediated antigen retrieval with Tris/EDTA buffer pH

9.0 before commencing with IHC staining protocol.



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