

Product datasheet

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

KO VALIDATED 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 notes | <p>This product is a recombinant monoclonal antibody, which offers several advantages including:</p> <ul style="list-style-type: none"> - High batch-to-batch consistency and reproducibility - Improved sensitivity and specificity - Long-term security of supply - Animal-free production <p>For more information see here.</p> <p>Our RabMAB[®] technology is a patented hybridoma-based technology for making rabbit monoclonal antibodies. For details on our patents, please refer to RabMAB[®] patents.</p> |

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 |

| | |
|---------------------|------------|
| Clonality | Monoclonal |
| Clone number | EPR17147 |
| Isotype | IgG |

Applications

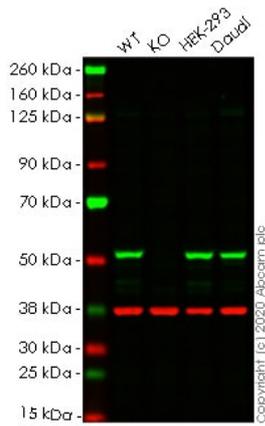
The Abpromise guarantee Our [Abpromise guarantee](#) covers the use of ab198500 in the following tested applications. The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

| 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 similarities | Belongs 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-293 cell lysate

Lane 4 : Daudi cell lysate

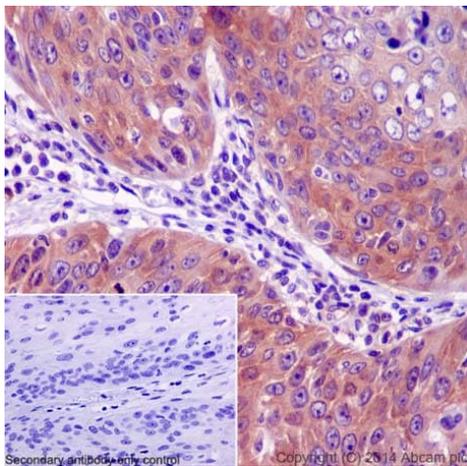
Lysates/proteins at 20 µg per lane.

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] - C-terminal 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.

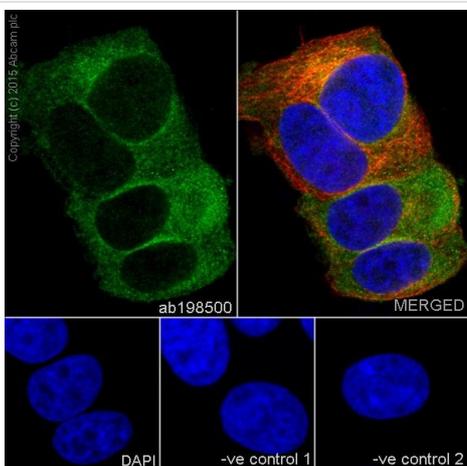


Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-PPP2R5E antibody [EPR17147] - C-terminal (ab198500)

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 is observed. 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.

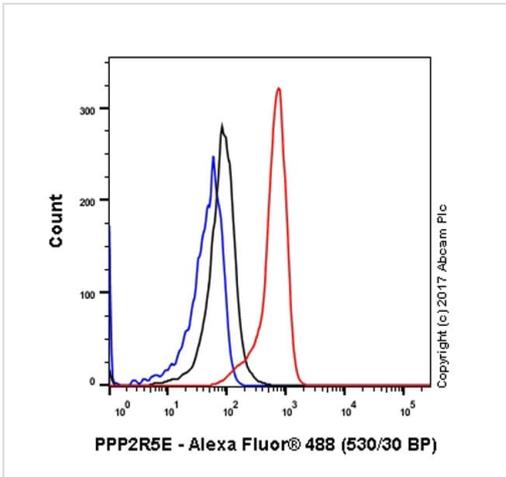


Immunocytochemistry/ Immunofluorescence - Anti-PPP2R5E antibody [EPR17147] - C-terminal (ab198500)

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).

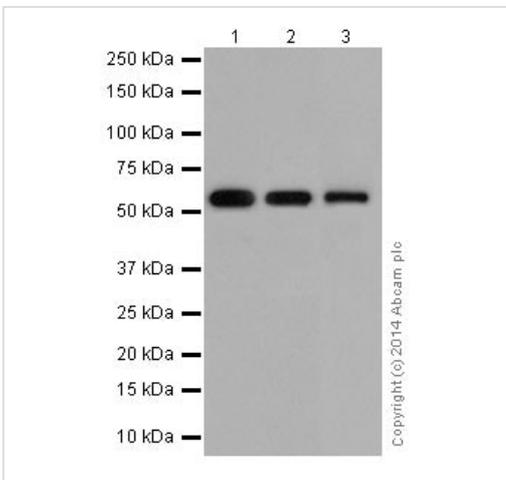
The negative controls are as follows;

1. ab198500 at 1/500 dilution followed by ab150120 (AlexaFluor®594 Goat anti-Mouse secondary) at 1/500 dilution.
2. ab7291 (anti-Tubulin mouse mAb) at 1/1000 dilution followed by ab150077 (Alexa Fluor®488 Goat Anti-Rabbit IgG 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 IgG (Alexa Fluor® 488) (ab150077) was used as the secondary antibody at a 1/2000 dilution. Black - Rabbit monoclonal IgG (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 IgG, (H+L), Peroxidase conjugate at 1/1000 dilution

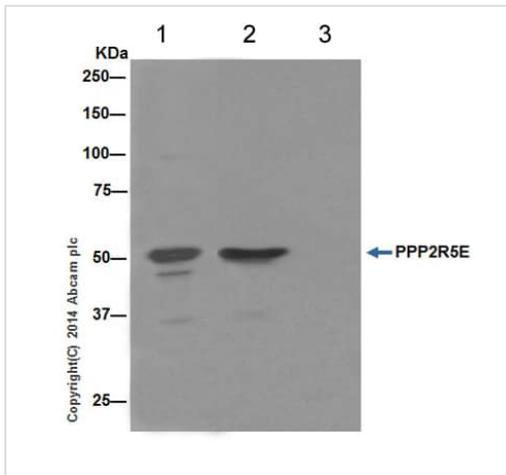
Developed using the ECL technique.

Predicted band size: 55 kDa

Observed band size: 55 kDa

Exposure time: 3 minutes

Blocking/Dilution buffer: 5% NFD/MTBST.



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

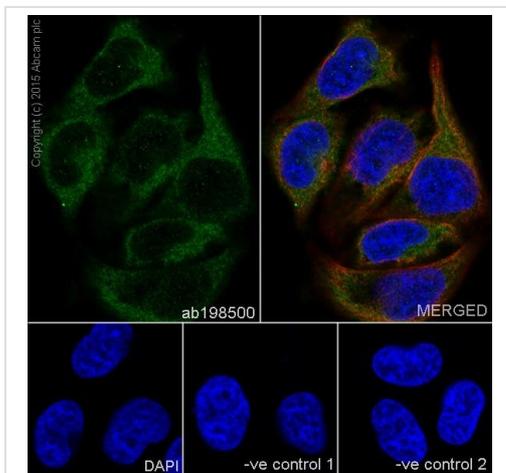
PPP2R5E was immunoprecipitated from 1mg of HeLa (Human cervix adenocarcinoma) whole cell extract with ab198500 at 1/175 dilution. Western blot was performed from the immunoprecipitate using ab198500 at 1/1000 dilution. Anti-Rabbit IgG (HRP), specific to the non-reduced form of IgG, was used as secondary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

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

Lane 2: HeLa whole cell extract

Lane 3: Rabbit monoclonal IgG (ab172730) instead of ab198500 in HeLa whole cell extract.

Blocking and dilution buffer and concentration: 5% NFD/MTBST.

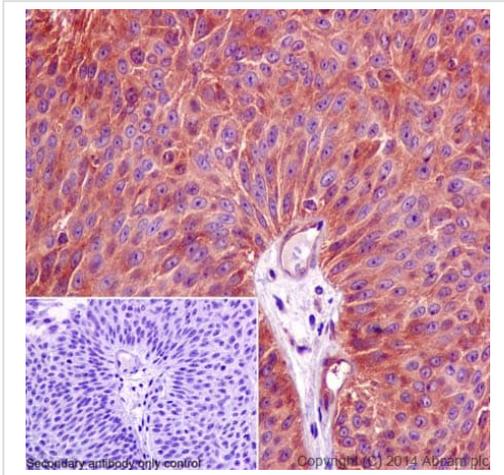


Immunocytochemistry/ Immunofluorescence - Anti-PPP2R5E antibody [EPR17147] - C-terminal (ab198500)

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 IgG (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;

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



Immunohistochemical analysis of paraffin-embedded Human transitional cell carcinoma of bladder 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 transitional cell carcinoma of bladder tissue is observed. 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.

Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-PPP2R5E antibody [EPR17147] - C-terminal (ab198500)

Why choose a recombinant antibody?

| | |
|---|---|
|  Research with confidence Consistent and reproducible results |  Long-term and scalable supply Recombinant technology |
|  Success from the first experiment Confirmed specificity |  Ethical standards compliant Animal-free production |

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

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