

Product datasheet

Anti-PRPF4 antibody [EPR17207] - C-terminal ab198998

Recombinant RabMAb

9 Images

Overview

| | |
|----------------------------|--|
| Product name | Anti-PRPF4 antibody [EPR17207] - C-terminal |
| Description | Rabbit monoclonal [EPR17207] to PRPF4 - C-terminal |
| Host species | Rabbit |
| Tested applications | Suitable for: Flow Cyt (Intra), IHC-P, WB |
| Species reactivity | Reacts with: Mouse, Rat, Human |
| Immunogen | Synthetic peptide. This information is proprietary to Abcam and/or its suppliers. |
| Positive control | WB: Raji, HepG2, 293, C6, Raw264.7, NIH/3T3 and HeLa whole cell lysate, Mouse brain kidney and spleen tissue lysates. IHC: Human transitional cell carcinoma of bladder tissue, Human kidney tissue, Rat cerebral cortex tissue, and Mouse cardiac muscle tissue. ICC/I: MCF-7 and HeLa cells. Flow Cyt (intra): HeLa cells |
| 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: 59% PBS, 40% Glycerol (glycerin, glycerine), 0.05% BSA |
| Purity | Protein A purified |
| Clonality | Monoclonal |
| Clone number | EPR17207 |

Isotype

IgG

Applications

The Abpromise guarantee

Our **Abpromise guarantee** covers the use of ab198998 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 |
|------------------|-----------|---|
| Flow Cyt (Intra) | | 1/70. |
| IHC-P | | 1/250. Perform heat mediated antigen retrieval with Tris/EDTA buffer pH 9.0 before commencing with IHC staining protocol. |
| WB | | 1/1000. Detects a band of approximately 58 kDa (predicted molecular weight: 58 kDa). |

Target

Function

Participates in pre-mRNA splicing. Part of the U4/U5/U6 tri-snRNP complex, one of the building blocks of the spliceosome.

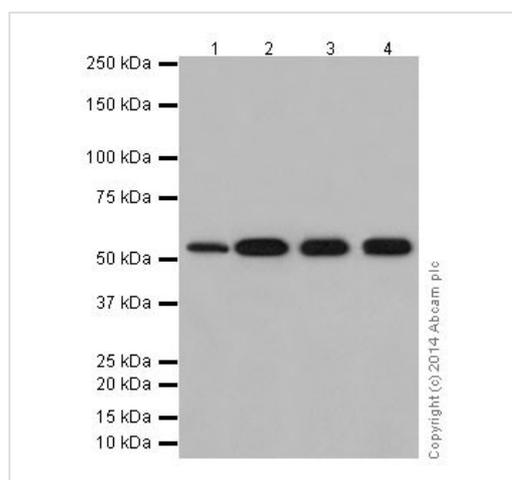
Sequence similarities

Contains 7 WD repeats.

Cellular localization

Nucleus speckle. Colocalizes with spliceosomal snRNPs.

Images



Western blot - Anti-PRPF4 antibody [EPR17207] - C-terminal (ab198998)

All lanes : Anti-PRPF4 antibody [EPR17207] - C-terminal (ab198998) at 1/20000 dilution

Lane 1 : Raji (Human Burkitt's lymphoma) whole cell lysate

Lane 2 : HepG2 (Human liver hepatocellular carcinoma) whole cell lysate

Lane 3 : 293 (Human epithelial cells from embryonic kidney) whole cell lysate

Lane 4 : HeLa (Human epithelial cells from cervix adenocarcinoma) whole cell lysate

Lysates/proteins at 10 µg per lane.

Secondary

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

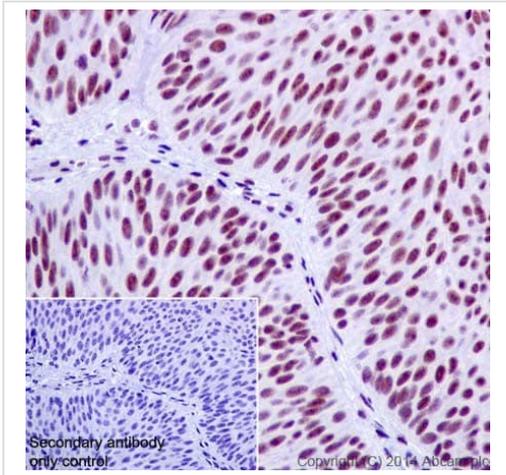
Developed using the ECL technique.

Predicted band size: 58 kDa

Observed band size: 58 kDa

Exposure time: 3 minutes

Blocking and diluting buffer was 5% NFDM/TBST

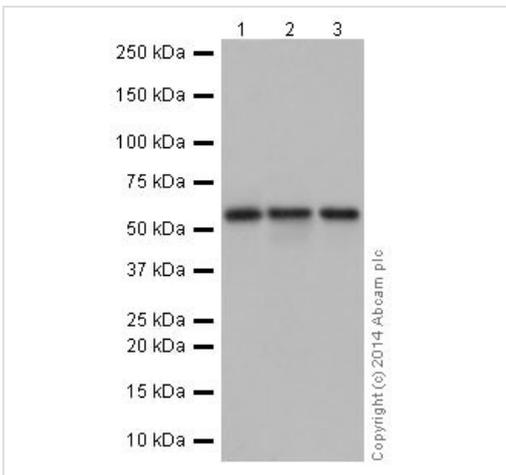


Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-PRPF4 antibody [EPR17207] - C-terminal (ab198998)

Immunohistochemical analysis of paraffin-embedded Human transitional cell carcinoma of bladder tissue labeling PRPF4 with ab198998 at 1/500 followed by Goat Anti-Rabbit IgG H&L (HRP) (**ab97051**) at 1/500. Counter stained with Hematoxylin. Nuclear staining on Human transitional cell carcinoma of bladder tissue was observed (Subcellular location - Nucleus speckle [UniProt]).

Negative control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) (**ab97051**) at 1/500 dilution.

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



Western blot - Anti-PRPF4 antibody [EPR17207] - C-terminal (ab198998)

All lanes : Anti-PRPF4 antibody [EPR17207] - C-terminal (ab198998) at 1/1000 dilution

Lane 1 : Mouse brain tissue lysate

Lane 2 : Mouse kidney tissue lysate

Lane 3 : Mouse spleen tissue lysate

Lysates/proteins at 10 µg per lane.

Secondary

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

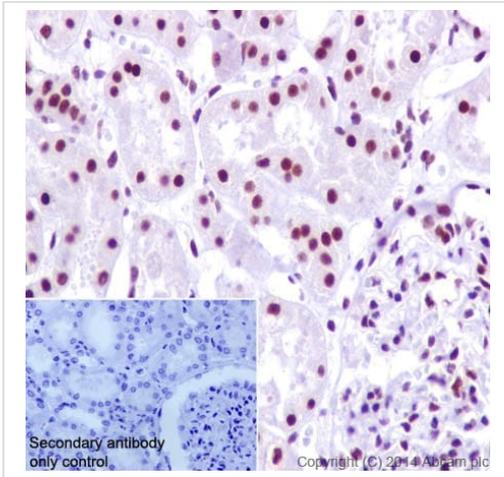
Developed using the ECL technique.

Predicted band size: 58 kDa

Observed band size: 58 kDa

Exposure time: 3 minutes

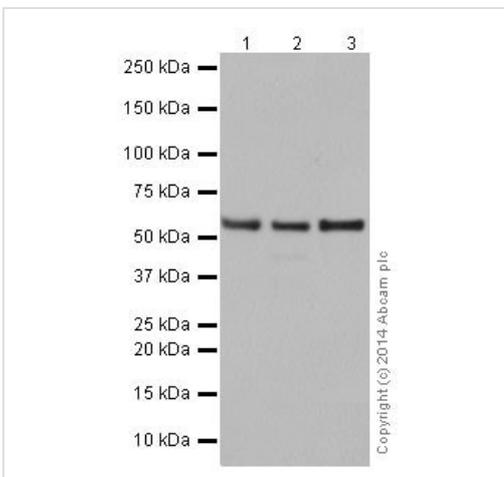
Blocking and diluting buffer was 5% NFDM/TBST



Immunohistochemical analysis of paraffin-embedded Human kidney tissue labeling PRPF4 with ab198998 at 1/500 followed by Goat Anti-Rabbit IgG H&L (HRP) (**ab97051**) at 1/500. Counter stained with Hematoxylin. Nuclear staining on Human kidney tissue was observed (Subcellular location - Nucleus speckle [UniProt]). Negative control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) (**ab97051**) at 1/500 dilution.

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-PRPF4 antibody [EPR17207] - C-terminal (ab198998)



Western blot - Anti-PRPF4 antibody [EPR17207] - C-terminal (ab198998)

All lanes : Anti-PRPF4 antibody [EPR17207] - C-terminal (ab198998) at 1/1000 dilution

Lane 1 : C6 (Rat glial tumor) whole cell lysate

Lane 2 : Raw264.7 (Mouse macrophages transformed with Abelson murine leukemia virus) whole cell lysate

Lane 3 : NIH/3T3 (Mouse embryo fibroblast) whole cell lysate

Lysates/proteins at 10 µg per lane.

Secondary

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

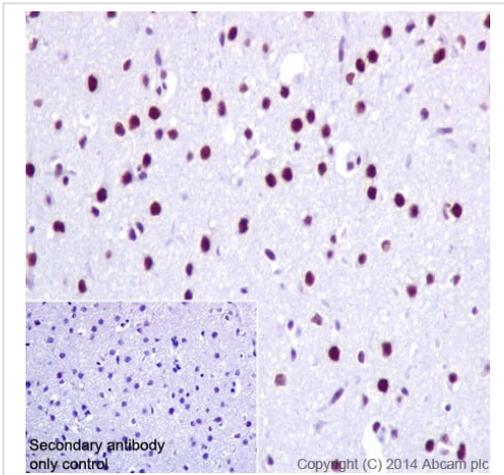
Developed using the ECL technique.

Predicted band size: 58 kDa

Observed band size: 58 kDa

Exposure time: 15 seconds

Blocking and diluting buffer was 5% NFDM/TBST

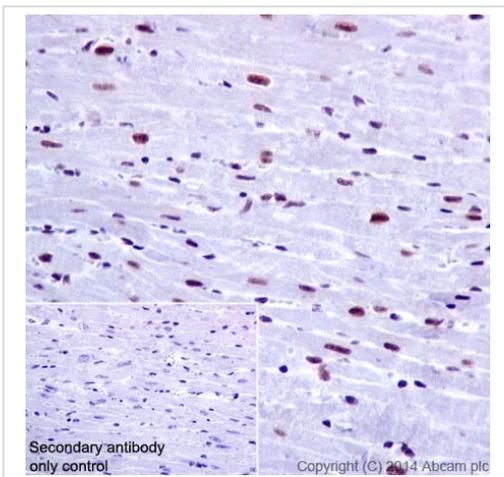


Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-PRPF4 antibody [EPR17207] - C-terminal (ab198998)

Immunohistochemical analysis of paraffin-embedded Rat cerebral cortex tissue labeling PRPF4 with ab198998 at 1/500 followed by Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/500. Counter stained with Hematoxylin. Nuclear staining on Rat cerebral cortex tissue was observed (Subcellular location - Nucleus speckle [UniProt]).

Negative control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/500 dilution.

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

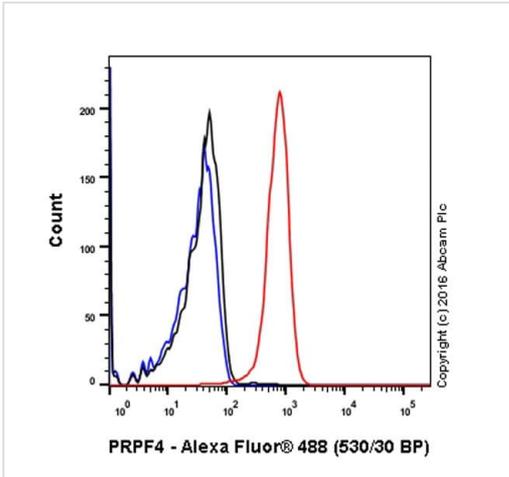


Immunohistochemistry (Formalin/PFA-fixed paraffin-embedded sections) - Anti-PRPF4 antibody [EPR17207] - C-terminal (ab198998)

Immunohistochemical analysis of paraffin-embedded Mouse cardiac muscle tissue labeling PRPF4 with ab198998 at 1/500 followed by Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/500. Counter stained with Hematoxylin. Nuclear staining on Mouse cardiac muscle tissue was observed (Subcellular location - Nucleus speckle [UniProt]).

Negative control: Used PBS instead of primary antibody, secondary antibody is Goat Anti-Rabbit IgG H&L (HRP) ([ab97051](#)) at 1/500 dilution.

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



Intracellular Flow Cytometry analysis of HeLa cells labelling PRPF4 (red) with purified ab198998 at dilution of 1/70. The secondary antibody used was Alexa Fluor[®] 488 goat-anti-rabbit IgG (1/2000). Cells were fixed with 4% paraformaldehyde and permeabilised with 90% methanol. Isotype control antibody was Rabbit monoclonal IgG (black). The blue line shows cells without incubation with primary antibody and secondary antibody.

Flow Cytometry (Intracellular) - Anti-PRPF4 antibody [EPR17207] - C-terminal (ab198998)

Why choose a recombinant antibody?

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

Anti-PRPF4 antibody [EPR17207] - C-terminal (ab198998)

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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