

## Product datasheet

# Anti-PPP2R1A antibody [6G3] - BSA and Azide free ab255976

Recombinant

[1 References](#) [5 Images](#)

### Overview

<b>Product name</b>	Anti-PPP2R1A antibody [6G3] - BSA and Azide free
<b>Description</b>	Rat monoclonal [6G3] to PPP2R1A - BSA and Azide free
<b>Host species</b>	Rat
<b>Tested applications</b>	<b>Suitable for:</b> WB, ICC/IF <b>Unsuitable for:</b> IP
<b>Species reactivity</b>	<b>Reacts with:</b> Mouse, Rat, Human
<b>Immunogen</b>	Recombinant full length protein corresponding to Human PPP2R1A. Database link: <a href="#">P30153</a>
<b>Positive control</b>	WB: HeLa, Jurkat ,Ramos, HepG2, C6 and RAW 264.7 whole cell lysate. ICC/IF: HeLa, RAW 264.7 and C6 cells.
<b>General notes</b>	<p>ab255976 is the carrier-free version of <a href="#">ab24736</a>.</p> <p>This antibody clone is manufactured by Abcam. If you require a custom buffer formulation or conjugation for your experiments, please contact <a href="mailto:orders@abcam.com">orders@abcam.com</a>.</p> <p>Our <b>carrier-free</b> 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.</p> <p>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.</p> <p>Use our <b>conjugation kits</b> for antibody conjugates that are ready-to-use in as little as 20 minutes with &lt;1 minute hands-on-time and 100% antibody recovery: available for fluorescent dyes, HRP, biotin and gold.</p> <p>This product is compatible with the Maxpar<sup>®</sup> Antibody Labeling Kit from Fluidigm, without the need for antibody preparation. Maxpar<sup>®</sup> is a trademark of Fluidigm Canada Inc.</p> <p>This product is a recombinant monoclonal antibody, which offers several advantages including:</p> <ul style="list-style-type: none"> <li>- High batch-to-batch consistency and reproducibility</li> <li>- Improved sensitivity and specificity</li> </ul>

 [Run BLAST with](#)

 [Run BLAST with](#)

- Long-term security of supply
  - Animal-free production
- For more information [see here](#).

## Properties

<b>Form</b>	Liquid
<b>Storage instructions</b>	Shipped at 4°C. Store at +4°C. Do Not Freeze.
<b>Storage buffer</b>	pH: 7.2 Constituent: PBS
<b>Carrier free</b>	Yes
<b>Purity</b>	Ion Exchange Chromatography
<b>Clonality</b>	Monoclonal
<b>Clone number</b>	6G3
<b>Isotype</b>	IgG2a

## Applications

**The Abpromise guarantee** Our **Abpromise guarantee** covers the use of ab255976 in the following tested applications. The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

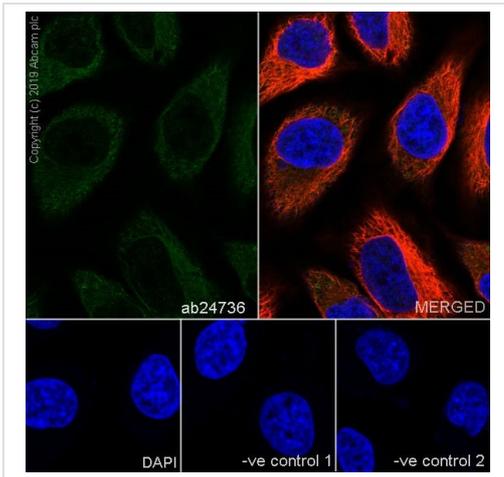
Application	Abreviews	Notes
WB		Use a concentration of 0.785 µg/ml. Predicted molecular weight: 65 kDa.
ICC/IF		Use a concentration of 7.85 µg/ml.

**Application notes** Is unsuitable for IP.

## Target

<b>Function</b>	The PR65 subunit of protein phosphatase 2A serves as a scaffolding molecule to coordinate the assembly of the catalytic subunit and a variable regulatory B subunit. Required for proper chromosome segregation and for centromeric localization of SGOL1 in mitosis.
<b>Sequence similarities</b>	Belongs to the phosphatase 2A regulatory subunit A family. Contains 15 HEAT repeats.
<b>Domain</b>	Each HEAT repeat appears to consist of two alpha helices joined by a hydrophilic region, the intrarepeat loop. The repeat units may be arranged laterally to form a rod-like structure.
<b>Cellular localization</b>	Cytoplasm. Chromosome > centromere. Centromeric localization requires the presence of BUB1.

## Images



Immunocytochemistry/ Immunofluorescence - Anti-PPP2R1A antibody [6G3] - BSA and Azide free (ab255976)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized HeLa (human cervix adenocarcinoma epithelial cell) cells labeling PPP2R1A with **ab24736** at 7.85µg/ml, followed by Goat Anti-Rat IgG (Alexa Fluor® 488) (**ab150157**) secondary antibody at 1/1000 dilution (green). Confocal image showing mainly showing cytoplasmic staining on HeLa cell line. The nuclear counterstain is DAPI (blue).

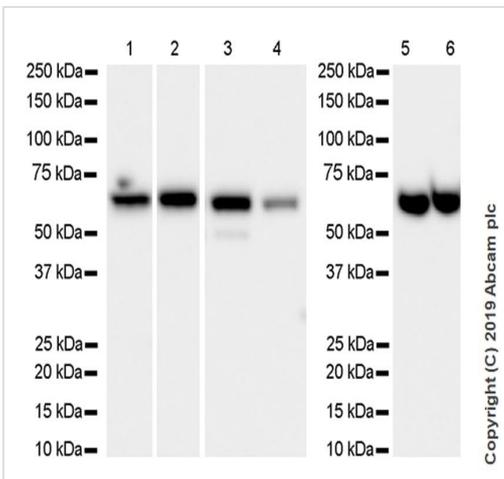
Tubulin is detected with **ab179504** Anti-beta IV Tubulin antibody - Microtubule Marker at 1/1000 dilution, followed by Goat Anti-Rabbit IgG (Alexa Fluor® 594) (**ab150080**) secondary antibody at 1/1000 dilution (red).

The negative controls are as follows:

**-ve control 1:** **ab24736** at 1/100 dilution, followed by Goat Anti-Rabbit IgG (Alexa Fluor® 594) (**ab150080**) secondary antibody at 1/1000 dilution.

**-ve control 2:** **ab179504** at 1/200 dilution, followed by **ab150157** AlexaFluor®488 Goat anti-rat secondary at 1/1000 dilution.

This image was produced using the same antibody clone but in a different formulation **ab24736**, PBS, sodium azide, glycerol and BSA.



Western blot - Anti-PPP2R1A antibody [6G3] - BSA and Azide free (ab255976)

**All lanes :** Anti-PPP2R1A antibody [6G3] (**ab24736**) at 0.785 µg/ml

**Lane 1 :** HeLa (human cervix adenocarcinoma epithelial cell), whole cell lysate

**Lane 2 :** Jurkat (human T cell leukemia T lymphocyte), whole cell lysate

**Lane 3 :** Ramos (Human Burkitt's lymphoma B lymphocyte), whole cell lysate

**Lane 4 :** HepG2 (human hepatocellular carcinoma epithelial cell), whole cell lysate

**Lane 5 :** C6 (rat glial tumor glial cell), whole cell lysate

**Lane 6 :** RAW264.7 (mouse Abelson murine leukemia virus-induced tumor macrophage), whole cell lysate

Lysates/proteins at 20 µg per lane.

### Secondary

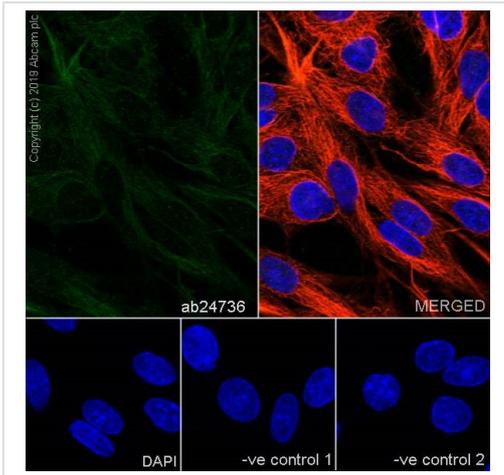
**All lanes :** Goat Anti-Rat IgG H&L (HRP) (**ab205720**) at 1/10000 dilution

**Predicted band size:** 65 kDa

**Exposure time:** 3 seconds

Blocking/Dilution buffer: 5% NFDM/TBST.

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Immunocytochemistry/ Immunofluorescence - Anti-PPP2R1A antibody [6G3] - BSA and Azide free (ab255976)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized C6 (rat glial tumor glial cell) cells labeling PPP2R1A with **ab24736** at 7.85µg/ml, followed by Goat Anti-Rat IgG (Alexa Fluor® 488) (**ab150157**) secondary antibody at 1/1000 dilution (green). Confocal image showing mainly showing cytoplasmic staining on C6 cell line. The nuclear counterstain is DAPI (blue).

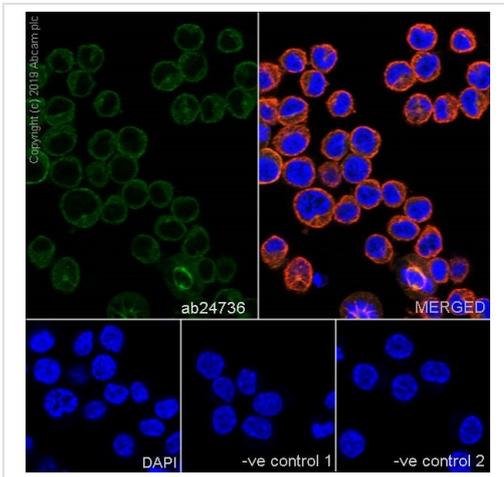
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Immunocytochemistry/ Immunofluorescence - Anti-PPP2R1A antibody [6G3] - BSA and Azide free (ab255976)

Immunofluorescent analysis of 4% paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized RAW 264.7 (mouse Abelson murine leukemia virus-induced tumor macrophage) cells labeling PPP2R1A with **ab24736** at 7.85µg/ml, followed by Goat Anti-Rat IgG (Alexa Fluor® 488) (**ab150157**) secondary antibody at 1/1000 dilution (green). Confocal image showing mainly showing cytoplasmic staining on RAW 264.7 cell line. The nuclear counterstain is DAPI (blue).

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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-PPP2R1A antibody [6G3] - BSA and Azide free (ab255976)

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