# abcam

# Product datasheet

# Mouse IgG1 monoclonal [R312-MouseIgG1]-Isotype control - BSA and Azide free ab281291

Recombinant

# 1 References 9 Images

#### Overview

Product name Mouse IgG1 monoclonal [R312-MouseIgG1]-Isotype control - BSA and Azide free

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

**Immunogen** The details of the immunogen for this antibody are not available.

**General notes** ab281291 is the carrier-free version of <u>ab280974</u>. This format is designed for use in antibody

labeling, including fluorochromes, metal isotopes, oligonucleotides, enzymes.

Our **carrier-free formats** are supplied in a buffer free of BSA, sodium azide and glycerol for

higher conjugation efficiency.

Use our <u>conjugation kits</u> 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.

ab281291 is compatible with the Maxpar® Antibody Labeling Kit from Fluidigm.

Maxpar® is a trademark of Fluidigm Canada Inc.

This antibody clone is manufactured by Abcam. If you require a custom buffer formulation or conjugation for your experiments, please contact **orders@abcam.com**.

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 production

For more information see here.

### **Properties**

Form Liquid

**Storage instructions** Shipped at 4°C. Store at +4°C. Do Not Freeze.

Storage buffer Constituent: 100% PBS

Carrier free Yes

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Purity Protein A purified

**Clonality** Monoclonal

Clone number R312-MouselgG1

**Isotype** IgG1

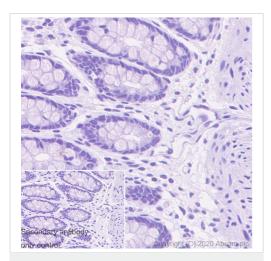
#### **Applications**

The Abpromise guarantee Our Abpromise guarantee covers the use of ab281291 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)		Use at an assay dependent concentration.
IHC-P		Use at an assay dependent concentration. Perform heat mediated antigen retrieval with citrate buffer pH 6 before commencing with IHC staining protocol.
ICC/IF		Use at an assay dependent concentration.

### **Images**



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Mouse IgG1 monoclonal [R312-MouseIgG1]-Isotype control - BSA and Azide free (ab281291)

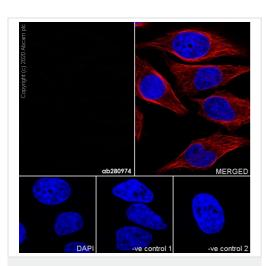
This data was developed using <u>ab280974</u>, the same antibody clone in a different buffer formulation.

Immunohistochemical analysis of paraffin-embedded human colon tissue labeling Mouse IgG1 with <u>ab280974</u> at 1/100 dilution followed by ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). The section was incubated with <u>ab280974</u> for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with Hematoxylin.

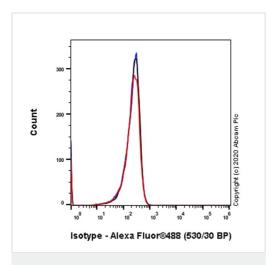
No staining on the human colon.

Secondary antibody only control: Secondary antibody is ready to use LeicaDS9800 (Bond™ Polymer Refine Detection).

Heat mediated antigen retrieval with Citrate buffer (pH 6.0, epitope retrieval solution 1) for 20 mins.



Immunocytochemistry/ Immunofluorescence Mouse IgG1 monoclonal [R312-MouseIgG1]-Isotype
control - BSA and Azide free (ab281291)



Flow Cytometry (Intracellular) - Mouse IgG1 monoclonal [R312-MouseIgG1]-Isotype control - BSA and Azide free (ab281291)

Immunofluorescent analysis of 4% Paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized HeLa cells labelling Mouse IgG1 with **ab280974** at 1/20 dilution, followed by **ab150113** Goat Anti-Mouse IgG H&L (Alexa Fluor<sup>®</sup> 488) antibody at 1/1000 dilution (Green). **ab179513** Anti-beta Tubulin rabbit monoclonal antibody was used to counterstain tubulin at 1/200 dilution followed by **ab150080** Goat Anti-Rabbit IgG H&L (Alexa Fluor<sup>®</sup> 594) at 1/1000 dilution (Red). The nuclear counterstain was DAPI (Blue).

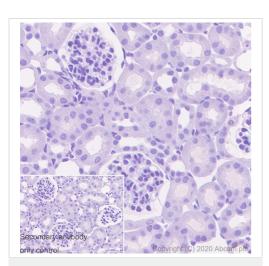
Confocal image showing **no staining** in HeLa cells.

**Negative control 1:** <u>ab280974</u> at a 1/20 dilution followed by ab150080 at a 1/1000 dilution.

**Negative control 2:** <u>ab179513</u> at a 1/200 dilution followed by <u>ab150113</u> at a 1/1000 dilution.

This data was developed using <u>ab280974</u>, the same antibody clone in a different buffer formulation.

Flow cytometric analysis of 4% paraformaldehyde fixed 90% methanol permeabilized HeLa (Human cervix adenocarcinoma epithelial cell) cells labelling Mouse IgG1 with <u>ab280974</u> at 1/500 dilution (0.1µg) (Red) compared with a Mouse IgG1, kappa monoclonal [MOPC-21] - isotype control <u>ab18443</u> (Black) and an unlabelled control (cells without incubation with primary antibody and secondary antibody) (Blue). A Goat anti mouse IgG (Alexa Fluor<sup>®</sup> 488, <u>ab150113</u>) at 1/2000 dilution was used as the secondary antibody.



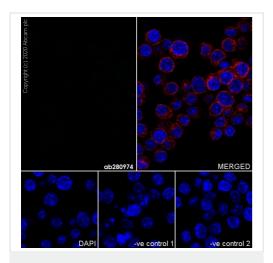
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Mouse IgG1 monoclonal [R312-MouseIgG1]-Isotype control - BSA and Azide free (ab281291)

Immunohistochemical analysis of paraffin-embedded mouse kidney tissue labeling Mouse IgG1 with <u>ab280974</u> at 1/100 dilution followed by a ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). The section was incubated with <u>ab280974</u> for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with Hematoxvlin.

No staining on the mouse kidney.

Secondary antibody only control: Secondary antibody is ready to use LeicaDS9800 (Bond™ Polymer Refine Detection).

Heat mediated antigen retrieval with Citrate buffer (pH 6.0, epitope retrieval solution 1) for 20 mins.



Immunocytochemistry/ Immunofluorescence Mouse IgG1 monoclonal [R312-MouseIgG1]-Isotype
control - BSA and Azide free (ab281291)

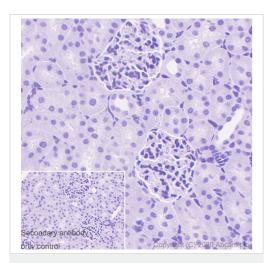
This data was developed using <u>ab280974</u>, the same antibody clone in a different buffer formulation.

Immunofluorescent analysis of 4% Paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized RAW 264.7 cells labelling Mouse IgG1 with <a href="mailto:ab280974">ab280974</a> at 1/20 dilution, followed by <a href="mailto:ab150113">ab150113</a> Goat Anti-Mouse IgG H&L (Alexa Fluor<sup>®</sup> 488) antibody at 1/1000 dilution (Green). <a href="mailto:ab179513">ab179513</a> Anti-beta Tubulin rabbit monoclonal antibody was used to counterstain tubulin at 1/200 dilution followed by <a href="mailto:ab150080">ab150080</a> Goat Anti-Rabbit IgG H&L (Alexa Fluor<sup>®</sup> 594) at 1/1000 dilution (Red). The nuclear counterstain was DAPI (Blue).

Confocal image showing **no staining** in RAW 264.7 cells.

**Negative control 1:** <u>ab280974</u> at a 1/20 dilution followed by <u>ab150080</u> at a 1/1000 dilution.

**Negative control 2:** <u>ab179513</u> at a 1/200 dilution followed by <u>ab150113</u> at a 1/1000 dilution.



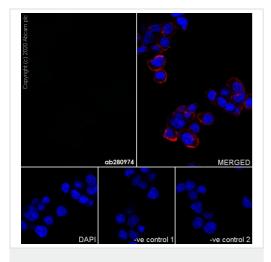
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Mouse IgG1 monoclonal [R312-MouseIgG1]-Isotype control - BSA and Azide free (ab281291)

Immunohistochemical analysis of paraffin-embedded rat kidney tissue labeling Mouse IgG1 with <u>ab280974</u> at 1/100 dilution followed by ready to use LeicaDS9800 (Bond™ Polymer Refine Detection). The section was incubated with <u>ab280974</u> for 30 mins at room temperature. The immunostaining was performed on a Leica Biosystems BOND® RX instrument. Counterstained with Hematoxylin.

No staining on the rat kidney.

Secondary antibody only control: Secondary antibody is ready to use LeicaDS9800 (Bond™ Polymer Refine Detection).

Heat mediated antigen retrieval with Citrate buffer (pH 6.0, epitope retrieval solution 1) for 20 mins.



Immunocytochemistry/ Immunofluorescence Mouse IgG1 monoclonal [R312-MouseIgG1]-Isotype
control - BSA and Azide free (ab281291)

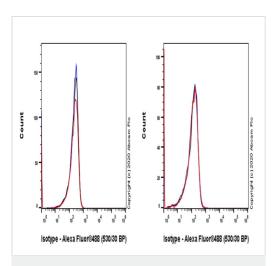
This data was developed using <u>ab280974</u>, the same antibody clone in a different buffer formulation.

Immunofluorescent analysis of 4% Paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized PC-12 cells labelling Mouse IgG1 with <a href="mailto:ab280974">ab280974</a> at 1/20 dilution, followed by <a href="mailto:ab150113">ab150113</a> Goat Anti-Mouse IgG H&L (Alexa Fluor<sup>®</sup> 488) antibody at 1/1000 dilution (Green). <a href="mailto:ab179513">ab179513</a> Anti-beta Tubulin rabbit monoclonal antibody was used to counterstain tubulin at 1/200 dilution followed by <a href="mailto:ab150080">ab150080</a> Goat Anti-Rabbit IgG H&L (Alexa Fluor<sup>®</sup> 594) at 1/1000 dilution (Red). The nuclear counterstain was DAPI (Blue).

Confocal image showing **no staining** in PC-12 cells.

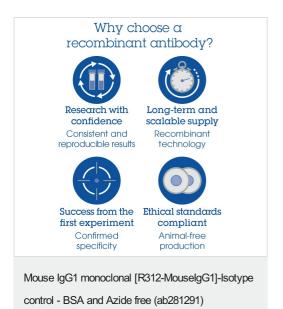
**Negative control 1:** <u>ab280974</u> at a 1/20 dilution followed by <u>ab150080</u> at a 1/1000 dilution.

**Negative control 2:** <u>ab179513</u> at a 1/200 dilution followed by <u>ab150113</u> at a 1/1000 dilution.



Flow Cytometry (Intracellular) - Mouse IgG1 monoclonal [R312-MouseIgG1]-Isotype control - BSA and Azide free (ab281291)

Flow cytometric analysis of 4% paraformaldehyde fixed 90% methanol permeabilized RAW 264.7 (Human cervix adenocarcinoma epithelial cell, Left) / PC-12 (Rat adrenal gland pheochromocytoma, Right) cells labelling Mouse IgG1 with ab280974 at 1/500 dilution (0.1µg) (Red) compared with a Mouse IgG1, kappa monoclonal [MOPC-21] - isotype control ab18443 (Black) and an unlabelled control (cells without incubation with primary antibody and secondary antibody) (Blue). A Goat antimouse IgG (Alexa Fluor® 488, ab150113) at 1/2000 dilution was used as the secondary antibody.



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

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