abcam

Product datasheet

Mouse IgG2b monoclonal [R311-MouseIgG2b] - Isotype control - BSA and Azide free ab281638

リコンピナント

画像数8

製品の概要

製品名

アプリケーション

免疫原

特記事項

Mouse IgG2b monoclonal [R311-MouseIgG2b] - Isotype control - BSA and Azide free

適用あり: ICC/IF, IHC-P, Flow Cyt

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

ab281638 is the carrier-free version of <u>ab281590</u>. This format is designed for use in antibody labeling, including fluorochromes, metal isotopes, oligonucleotides, enzymes.

Our <u>carrier-free formats</u> 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.

ab281638 is compatible with the Maxpar® Antibody Labeling Kit from Fluidigm.

Maxpar® is a trademark of Fluidigm Canada Inc.

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.

製品の特性

製品の状態 Liquid

保存方法 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.

パッファー Constituent: 100% PBS

キャリア・フリー はい

精製度 Protein A purified

ポリ/モノ モノクローナル

1

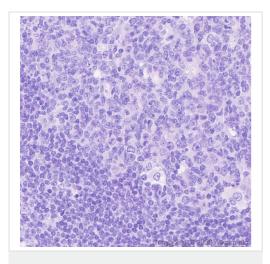
lgG2b

アプリケーション

The Abpromise guarantee <u>Abpromise保証は、</u>次のテスト済みアプリケーションにおけるab281638の使用に適用されます アプリケーションノートには、推奨の開始希釈率がありますが、適切な希釈率につきましてはご検討ください。

アプリケーション	Abreviews	特記事項
ICC/IF		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.
Flow Cyt		Use at an assay dependent concentration.

画像



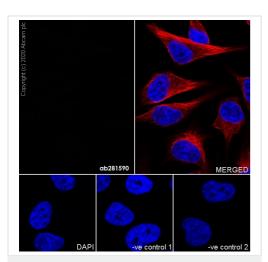
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Mouse IgG2b monoclonal [R311-MouseIgG2b] - Isotype control - BSA and Azide free (ab281638)

This data was developed using <u>ab281590</u>, the same antibody clone in a different buffer formulation.

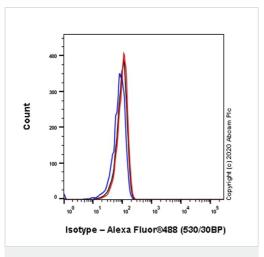
Immunohistochemical analysis of paraffin-embedded Human tonsil tissue labeling Mouse IgG2b with <u>ab281590</u> at 1/200 dilution followed by a ready to use Goat Anti-Mouse IgG H&L (HRP polymer) (<u>ab214879</u>). **Negative control:** No staining on human tonsil. Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is ready to use Goat Anti-Mouse IgG H&L (HRP polymer) (ab214879).

Heat mediated antigen retrieval using <u>ab93678</u> (citrate buffer, pH 6.0).



Immunocytochemistry/ Immunofluorescence Mouse IgG2b monoclonal [R311-MouseIgG2b] Isotype control - BSA and Azide free (ab281638)



Flow Cytometry - Mouse IgG2b monoclonal [R311-MouseIgG2b] - Isotype control - BSA and Azide free (ab281638)

This data was developed using <u>ab281590</u>, the same antibody clone in a different buffer formulation.

Immunofluorescent analysis of 4% Paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized HeLa cells labelling Mouse IgG2b with ab281590 at 1/20 dilution, followed by ab150113 Goat Anti-Mouse IgG H&L (Alexa Fluor[®] 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[®] 594) at 1/1000 dilution (Red). The nuclear counterstain was DAPI (Blue).

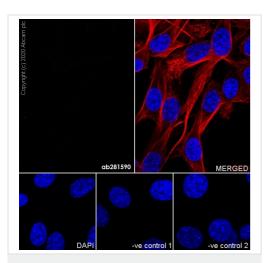
Confocal image showing **no staining** in HeLa cells.

Negative control 1: <u>ab281590</u> at a 1/20 dilution followed by <u>ab150113</u> at a 1/1000 dilution.

Negative control 2: <u>ab179513</u> at a 1/200 dilution followed by <u>ab150080</u> at a 1/1000 dilution.

This data was developed using <u>ab281590</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 IgG2b with <u>ab281590</u> at 1/500 dilution (0.1µg) (Red) compared with a Mouse (E7Q5L) mAb IgG2b Isotype Control (Black) and unlabelled control (cells without incubation with primary antibody and secondary antibody)(Blue). A Goat anti mouse IgG (Alexa Fluor[®] 488, <u>ab150113</u>) at 1/2000 dilution was used as the secondary antibody.



Immunocytochemistry/ Immunofluorescence Mouse IgG2b monoclonal [R311-MouseIgG2b] Isotype control - BSA and Azide free (ab281638)

ab281590 MERGED

DAPI -ve control 1 -ve control 2

Immunocytochemistry/ Immunofluorescence -Mouse IgG2b monoclonal [R311-MouseIgG2b] -Isotype control - BSA and Azide free (ab281638)

This data was developed using <u>ab281590</u>, the same antibody clone in a different buffer formulation.

Immunofluorescent analysis of 4% Paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized NIH/3T3 cells labelling Mouse IgG2b with ab281590 at 1/20 dilution, followed by ab150113 Goat Anti-Mouse IgG H&L (Alexa Fluor[®] 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[®] 594) at 1/1000 dilution (Red). The nuclear counterstain was DAPI (Blue).

Confocal image showing **no staining** in NIH/3T3 cells.

Negative control 1: <u>ab281590</u> at a 1/20 dilution followed by <u>ab150113</u> at a 1/1000 dilution.

Negative control 2: <u>ab179513</u> at a 1/200 dilution followed by <u>ab150080</u> at a 1/1000 dilution.

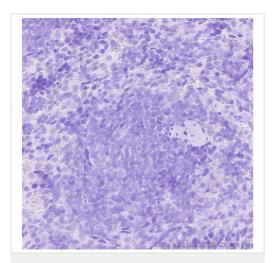
This data was developed using <u>ab281590</u>, the same antibody clone in a different buffer formulation.

Immunofluorescent analysis of 4% Paraformaldehyde-fixed, 0.1% Triton X-100 permeabilized PC-12 cells labelling Mouse IgG2b with ab281590 at 1/20 dilution, followed by ab150113 Goat Anti-Mouse IgG H&L (Alexa Fluor[®] 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[®] 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>ab281590</u> at a 1/20 dilution followed by <u>ab150113</u> at a 1/1000 dilution.

Negative control 2: <u>ab179513</u> at a 1/200 dilution followed by <u>ab150080</u> at a 1/1000 dilution.



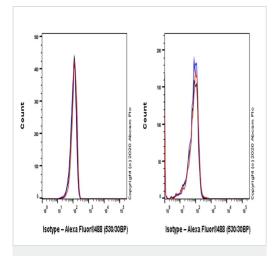
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Mouse IgG2b monoclonal [R311-MouseIgG2b] - Isotype control - BSA and Azide free (ab281638)

This data was developed using <u>ab281590</u>, the same antibody clone in a different buffer formulation.

Immunohistochemical analysis of paraffinembedded Rat spleen tissue labeling Mouse IgG2b with <u>ab281590</u> at 1/200 dilution followed by a ready to use Goat Anti-Mouse IgG H&L (HRP polymer) (<u>ab214879</u>). **Negative control:** No staining on rat spleen. Counterstained with Hematoxylin.

Secondary antibody only control: Secondary antibody is ready to use Goat Anti-Mouse IgG H&L (HRP polymer) (ab214879).

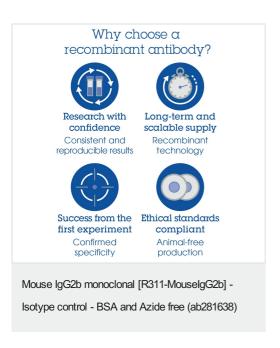
Heat mediated antigen retrieval using <u>ab93678</u> (citrate buffer, pH 6.0).



Flow Cytometry - Mouse IgG2b monoclonal [R311-MouseIgG2b] - Isotype control - BSA and Azide free (ab281638)

This data was developed using <u>ab281590</u>, the same antibody clone in a different buffer formulation.

Flow cytometric analysis of 4% paraformaldehyde fixed 90% methanol permeabilized NIH/3T3 (Mouse embryonic fibroblast, Left) / PC-12 (Rat adrenal gland pheochromocytoma, Right) cells labelling Mouse IgG2b with <u>ab281590</u> at 1/500 dilution (0.1µg) (Red) compared with a Mouse (E7Q5L) mAb IgG2b Isotype Control (Black) and unlabelled control (cells without incubation with primary antibody and secondary antibody)(Blue). A Goat anti mouse IgG (Alexa Fluor® 488, <u>ab150113</u>) at 1/2000 dilution was used as the secondary antibody.



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