

Product Datasheet

Muscarinic Acetylcholine Receptor M2/CHRM2 Antibody (JE38-35) NBP3-32617

Unit Size: 100 ul

Store at 4C short term. Aliquot and store at -20C long term. Avoid freeze-thaw cycles.

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NBP3-32617**Muscarinic Acetylcholine Receptor M2/CHRM2 Antibody (JE38-35)**

Product Information	
Unit Size	100 ul
Concentration	1 mg/ml
Storage	Store at 4C short term. Aliquot and store at -20C long term. Avoid freeze-thaw cycles.
Clonality	Monoclonal
Clone	JE38-35
Preservative	0.05% Sodium Azide
Isotype	IgG
Purity	Protein A purified
Buffer	1*TBS (pH7.4), 0.05% BSA and 40% Glycerol
Target Molecular Weight	52 kDa

Product Description	
Description	Novus Biologicals Rabbit Muscarinic Acetylcholine Receptor M2/CHRM2 Antibody (JE38-35) (NBP3-32617) is a recombinant monoclonal antibody validated for use in IHC, WB, Flow and ICC/IF. All Novus Biologicals antibodies are covered by our 100% guarantee.
Host	Rabbit
Gene ID	1129
Gene Symbol	CHRM2
Species	Human, Mouse, Rat
Immunogen	Synthetic peptide within Human Muscarinic Acetylcholine Receptor M2/CHRM2 aa 140-184 / 466. (Uniprot: P08172)

Product Application Details	
Applications	Western Blot, Immunohistochemistry-Paraffin, Flow Cytometry, Immunocytochemistry/ Immunofluorescence, Immunohistochemistry
Recommended Dilutions	Western Blot 1:1000, Flow Cytometry 1:500-1:1000, Immunohistochemistry, Immunocytochemistry/ Immunofluorescence 1:100, Immunohistochemistry-Paraffin 1:1000

Images

Western Blot: Muscarinic Acetylcholine Receptor M2/CHRM2 Antibody (JE38-35) [NBP3-32617] - Western blot analysis of Muscarinic Acetylcholine Receptor M2/CHRM2 on different lysates with Rabbit anti-Muscarinic Acetylcholine Receptor M2/CHRM2 antibody (NBP3-32617) at 1/1,000 dilution.

Lane 1: HepG2 cell lysate
 Lane 2: SH-SY5Y cell lysate
 Lane 3: U-87 MG cell lysate
 Lane 4: Neuro-2a cell lysate
 Lane 5: PC-12 cell lysate

Lysates/proteins at 20 ug/Lane.

Predicted band size: 52 kDa
 Observed band size: 52 kDa

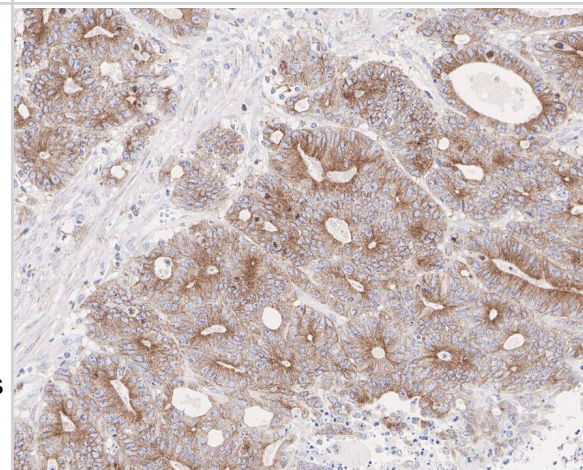
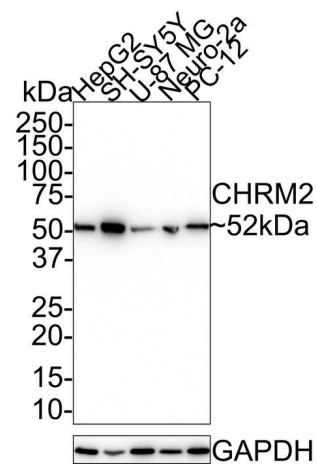
Exposure time: 1 minute 17 seconds;

4-20% SDS-PAGE gel.

Proteins were transferred to a PVDF membrane and blocked with 5% NFDM/TBST for 1 hour at room temperature. The primary antibody (NBP3-32617) at 1/1,000 dilution was used in 5% NFDM/TBST at 4 overnight. Goat Anti-Rabbit IgG - HRP Secondary Antibody at 1:100,000 dilution was used for 1 hour at room temperature.

Immunohistochemistry: Muscarinic Acetylcholine Receptor M2/CHRM2 Antibody (JE38-35) [NBP3-32617] - Immunohistochemical analysis of paraffin-embedded human colon carcinoma tissue with Rabbit anti-Muscarinic Acetylcholine Receptor M2/CHRM2 antibody (NBP3-32617) at 1/1,000 dilution.

The section was pre-treated using heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0) for 20 minutes. The tissues were blocked in 1% BSA for 20 minutes at room temperature, washed with ddH₂O and PBS, and then probed with the primary antibody (NBP3-32617) at 1/1,000 dilution for 1 hour at room temperature. The detection was performed using an HRP conjugated compact polymer system. DAB was used as the chromogen. Tissues were counterstained with hematoxylin and mounted with DPX.



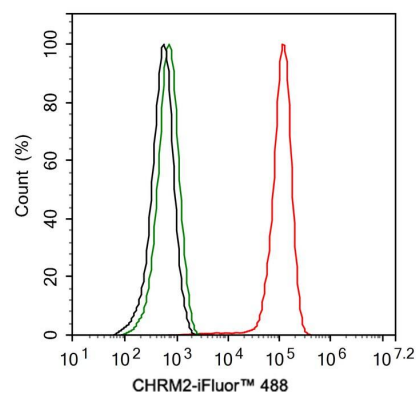
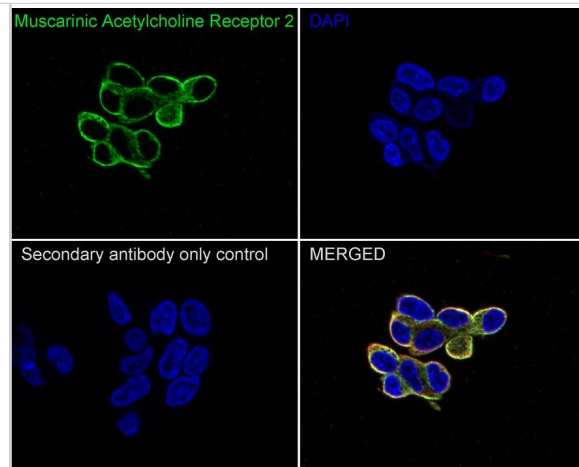
Immunocytochemistry/ Immunofluorescence: Muscarinic Acetylcholine Receptor M2/CHRM2 Antibody (JE38-35) [NBP3-32617] - Immunocytochemistry analysis of SH-SY5Y cells labeling Muscarinic Acetylcholine Receptor M2/CHRM2 with Rabbit anti-Muscarinic Acetylcholine Receptor M2/CHRM2 antibody (NBP3-32617) at 1/100 dilution.

Cells were fixed in 4% paraformaldehyde for 20 minutes at room temperature, permeabilized with 0.1% Triton X-100 in PBS for 5 minutes at room temperature, then blocked with 1% BSA in 10% negative goat serum for 1 hour at room temperature. Cells were then incubated with Rabbit anti-Muscarinic Acetylcholine Receptor M2/CHRM2 antibody (NBP3-32617) at 1/100 dilution in 1% BSA in PBST overnight at 4 °C. Goat Anti-Rabbit IgG H&L (iFluor™ 488) was used as the secondary antibody at 1/1,000 dilution. PBS instead of the primary antibody was used as the secondary antibody only control. Nuclear DNA was labelled in blue with DAPI.

Beta tubulin (red) was stained at 1/100 dilution overnight at +4 °C. Goat Anti-Mouse IgG H&L (iFluor™ 594) was used as the secondary antibody at 1/1,000 dilution.

Flow Cytometry: Muscarinic Acetylcholine Receptor M2/CHRM2 Antibody (JE38-35) [NBP3-32617] - Flow cytometric analysis of SH-SY5Y cells labeling Muscarinic Acetylcholine Receptor M2/CHRM2.

Cells were fixed and permeabilized. Then stained with the primary antibody (NBP3-32617, 1ug/ml) (red) compared with Rabbit IgG Isotype Control (green). After incubation of the primary antibody at +4 °C for an hour, the cells were stained with a iFluor™ 488 conjugate-Goat anti-Rabbit IgG Secondary antibody at 1/1,000 dilution for 30 minutes at +4 °C. Unlabelled sample was used as a control (cells without incubation with primary antibody; black).





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Products Related to NBP3-32617

NBP2-33376H	Blue Marker Antibody (6F4-F6) [HRP]
HAF008	Goat anti-Rabbit IgG Secondary Antibody [HRP]
NB7160	Goat anti-Rabbit IgG (H+L) Secondary Antibody [HRP]
NBP2-24891	Rabbit IgG Isotype Control

Limitations

This product is for research use only and is not approved for use in humans or in clinical diagnosis. Primary Antibodies are guaranteed for 1 year from date of receipt.

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