

Anti-CHRM2 Antibody

Rabbit polyclonal antibody to CHRM2 Catalog # AP61231

Specification

Anti-CHRM2 Antibody - Product Information

Application WB, IF/IC
Primary Accession P08172
Other Accession O9ERZ4

Reactivity
Human, Mouse, Rat, Pig, Bovine
Rabbit

Host Rabbit
Clonality Polyclonal
Calculated MW 51715

Anti-CHRM2 Antibody - Additional Information

Gene ID 1129

Other Names

Muscarinic acetylcholine receptor M2

Target/Specificity

KLH-conjugated synthetic peptide encompassing a sequence within the center region of human CHRM2. The exact sequence is proprietary.

Dilution

WB~~WB (1/500 - 1/1000), IF/IC (1/100 - 1/500) IF/IC~~N/A

Format

Liquid in 0.42% Potassium phosphate, 0.87% Sodium chloride, pH 7.3, 30% glycerol, and 0.09% (W/V) sodium azide.

Storage

Store at -20 °C. Stable for 12 months from date of receipt

Anti-CHRM2 Antibody - Protein Information

Name CHRM2

Function

The muscarinic acetylcholine receptor mediates various cellular responses, including inhibition of adenylate cyclase, breakdown of phosphoinositides and modulation of potassium channels through the action of G proteins. Primary transducing effect is adenylate cyclase inhibition. Signaling promotes phospholipase C activity, leading to the release of inositol trisphosphate (IP3); this then triggers calcium ion release into the cytosol.

Cellular Location



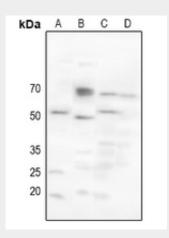
Cell membrane; Multi-pass membrane protein. Postsynaptic cell membrane; Multi-pass membrane protein. Note=Phosphorylation in response to agonist binding promotes receptor internalization {ECO:0000250|UniProtKB:P06199}

Anti-CHRM2 Antibody - Protocols

Provided below are standard protocols that you may find useful for product applications.

- Western Blot
- Blocking Peptides
- Dot Blot
- Immunohistochemistry
- Immunofluorescence
- <u>Immunoprecipitation</u>
- Flow Cytomety
- Cell Culture

Anti-CHRM2 Antibody - Images

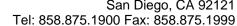


Western blot analysis of CHRM2 expression in HEK293T (A), mouse lung (B), mouse kidney (C), rat kidney (D) whole cell lysates.



Immunofluorescent analysis of CHRM2 staining in LOVO cells. Formalin-fixed cells were permeabilized with 0.1% Triton X-100 in TBS for 5-10 minutes and blocked with 3% BSA-PBS for 30 minutes at room temperature. Cells were probed with the primary antibody in 3% BSA-PBS and incubated overnight at 4~%C in a hidified chamber. Cells were washed with PBST and incubated







with a Alexa Fluor 594-conjugated secondary antibody (red) in PBS at room temperature in the dark.

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