产品名称: CHRM2 Mouse Monoclonal Antibody

产品货号: AMM85970



产品概述 (Summary)

产品名称 (Production Name) CHRM2 Mouse Monoclonal Antibody

描述 (Description) Mouse monoclonal Antibody

宿主 (Host) Mouse

应用 (Application)WB,IHC,ICC,FC种属反应性 (Reactivity)Human, Mouse

产品性能 (Performance)

個联物 (Conjugation) Unconjugated 修饰 (Modification) Unmodified 同种型 (Isotype) Mouse IgG1 充隆 (Clonality) Monoclonal Liquid

Store at 4°C short term. Aliquot and store at -20°C long term. Avoid 存放说明 (Storage)

freeze/thaw cycles.

储存溶液 (Buffer) Purified antibody in TBS with 0.05% sodium azide.

纯化方式 (Purification) Affinity Purification

免疫原信息 (Immunogen)

基因名 (Gene Name) CHRM2

别名 (Alternative Names) Muscarinic acetylcholine receptor M2, CHRM2

基因 ID (Gene ID) 1129.0

P08172.This antibody is generated from a mouse immunized with a 蛋白ID (SwissProt ID)

recombinant protein.

产品应用 (Application)

稀释比 (Dilution Ratio) WB 1:500-1:1000,IHC 1:100-1:500,ICC 1:25-1:50,FC 1:25-1:50

蛋白分子量 (Molecular Weight) 51.7kDa

研究背景 (Background)

The muscarinic acetylcholine receptor mediates various cellular responses, including inhibition of adenylate cyclase,

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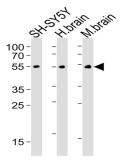


breakdown of phosphoinositides and modulation of potassium channels through the action of G proteins. Primary transducing effect is adenylate cyclase inhibition.

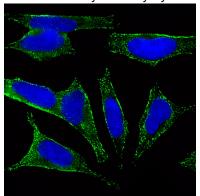
研究领域 (Research Area)

PI3K-Akt signaling pathway

图片 (Image Data)



Western blot analysis of lysates from SH-SY5Y cell line, human brain, mouse brain tissue(from left to right), using CHRM2 Antibody(Cat. #AMM85970). AMM85970 was diluted at 1:500 at each lane. A goat anti-mouse IgG H&L(HRP) at 1:3000 dilution was used as the secondary antibody. Lysates at 20µg per lane.



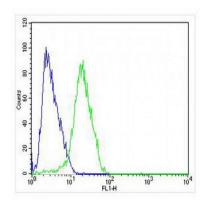
Fluorescent image of SH-SY5Y cells stained with CHRM2 Antibody (Cat#AMM85970). AMM85970 was diluted at 1:25 dilution. An Alexa Fluor® 488-conjugated goat anti-mouse IgG at 1:400 dilution was used as the secondary antibody (green). DAPI was used to stain the cell nuclear (blue).

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Overlay histogram showing SH-SY5Y cells stained with (green line). The cells were fixed with 4% paraformaldehyde (10 min) and then permeabilized with 90% methanol for 10 min. The cells were then icubated in 2% bovine serum albumin to block non-specific protein-protein interactions followed by the antibody (, 1:25 dilution) for 60 min at 37°C. The secondary antibody used was Alexa Fluor® 488 goat anti-mouse IgG (166821) at 1/200 dilution for 40 min at 37°C. Isotype control antibody (blue line) was mouse IgG1 (1µg/1x10^6 cells) used under the same conditions. Acquisition of >10, 000 events was performed.

注意事项 (Note)

For research use only.

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