

OPRM1 Antibody (Center)
Affinity Purified Rabbit Polyclonal Antibody (Pab)
Catalog # AP9130C

Specification

OPRM1 Antibody (Center) - Product Information

Application	WB, IHC-P, FC,E
Primary Accession	P35372
Other Accession	P33535 , Q95247 , P42866 , Q95M54
Reactivity	Human, Mouse
Predicted	Monkey, Pig, Rat
Host	Rabbit
Clonality	Polyclonal
Isotype	Rabbit Ig
Calculated MW	44779
Antigen Region	161-187

OPRM1 Antibody (Center) - Additional Information

Gene ID 4988

Other Names

Mu-type opioid receptor, M-OR-1, MOR-1,
Mu opiate receptor, Mu opioid receptor,
MOP, hMOP, OPRM1, MOR1

Target/Specificity

This OPRM1 antibody is generated from rabbits immunized with a KLH conjugated synthetic peptide between 161-187 amino acids from the Central region of human OPRM1.

Dilution

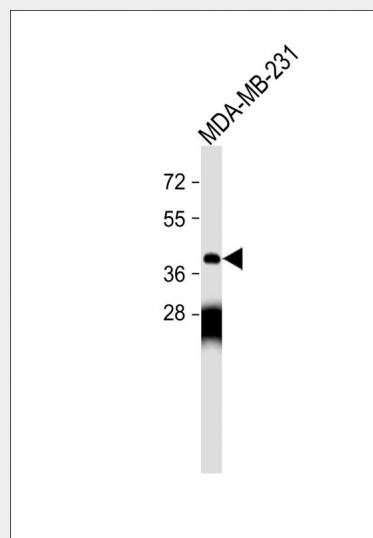
WB~~1:1000
IHC-P~~1:10~50
FC~~1:10~50

Format

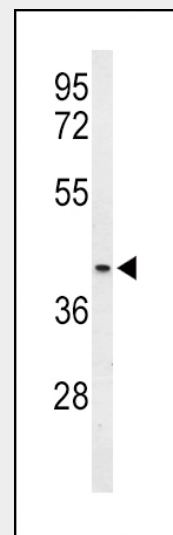
Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.

Storage

Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw



Anti-OPRM1 Antibody (Center) at 1:500 dilution + MDA-MB-231 whole cell lysate
Lysates/proteins at 20 µg per lane.
Secondary Goat Anti-Rabbit IgG, (H+L),
Peroxidase conjugated at 1/10000 dilution.
Predicted band size : 44 kDa
Blocking/Dilution buffer: 5% NFDM/TBST.



Western blot analysis of OPRM1 Antibody (Center) (Cat. #AP9130c) in mouse heart tissue lysates (35ug/lane). OPRM1 (arrow) was detected using the purified Pab.

cycles.

Precautions

OPRM1 Antibody (Center) is for research use only and not for use in diagnostic or therapeutic procedures.

OPRM1 Antibody (Center) - Protein Information

Name OPRM1

Synonyms MOR1

Function

Receptor for endogenous opioids such as beta-endorphin and endomorphin. Receptor for natural and synthetic opioids including morphine, heroin, DAMGO, fentanyl, etorphine, buprenorphin and methadone (PubMed:<a href="http://www.uniprot.org/citations/7905839"

target="_blank">7905839 ,

PubMed:<a href="http://www.uniprot.org/citations/7957926"

target="_blank">7957926 ,

PubMed:<a href="http://www.uniprot.org/citations/7891175"

target="_blank">7891175 ,

PubMed:<a href="http://www.uniprot.org/citations/12589820"

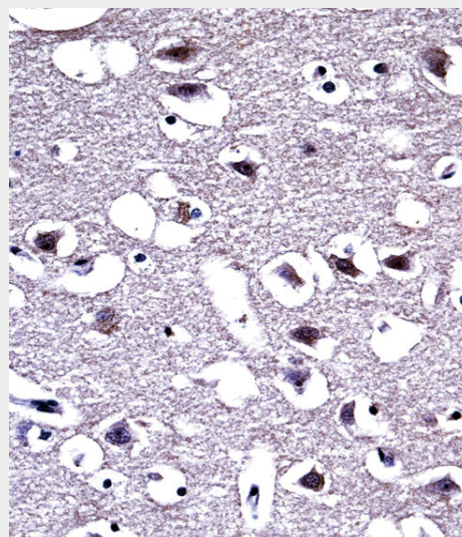
target="_blank">12589820 ,

PubMed:<a href="http://www.uniprot.org/citations/9689128"

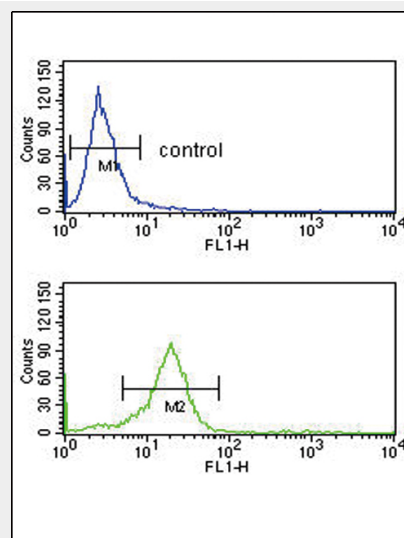
target="_blank">9689128). Agonist binding to the receptor induces coupling to an inactive GDP-bound heterotrimeric G-protein complex and subsequent exchange of GDP for GTP in the G-protein alpha subunit leading to dissociation of the G-protein complex with the free GTP-bound G-protein alpha and the G-protein beta-gamma dimer activating downstream cellular effectors (PubMed:<a href="http://www.uniprot.org/citations/7905839"

target="_blank">7905839). The

agonist- and cell type-specific activity is predominantly coupled to pertussis toxin-sensitive G(i) and G(o) G alpha proteins, GNAI1, GNAI2, GNAI3 and GNAO1 isoforms Alpha-1 and Alpha-2, and to a lesser extent to pertussis toxin-insensitive G alpha proteins GNAZ and GNA15 (PubMed:12068084). They mediate an array of downstream cellular responses, including inhibition of adenylate



OPRM1 Antibody (Center) (AP9130c) immunohistochemistry analysis in formalin fixed and paraffin embedded human brain tissue followed by peroxidase conjugation of the secondary antibody and DAB staining. This data demonstrates the use of OPRM1 Antibody (Center) for immunohistochemistry. Clinical relevance has not been evaluated.



OPRM1 Antibody (Center) (Cat. #AP9130c) flow cytometry analysis of K562 cells (bottom histogram) compared to a negative control cell (top histogram). FITC-conjugated goat-anti-rabbit secondary antibodies were used for the analysis.

OPRM1 Antibody (Center) - Background

OPRM1 is the mu opioid receptor, which is the primary site of action for the most commonly

cyclase activity and both N-type and L-type calcium channels, activation of inward rectifying potassium channels, mitogen-activated protein kinase (MAPK), phospholipase C (PLC), phosphoinositide/protein kinase (PKC), phosphoinositide 3-kinase (PI3K) and regulation of NF-kappa-B. Also couples to adenylate cyclase stimulatory G alpha proteins. The selective temporal coupling to G- proteins and subsequent signaling can be regulated by RGSZ proteins, such as RGS9, RGS17 and RGS4. Phosphorylation by members of the GPRK subfamily of Ser/Thr protein kinases and association with beta-arrestins is involved in short-term receptor desensitization. Beta- arrestins associate with the GPRK-phosphorylated receptor and uncouple it from the G-protein thus terminating signal transduction. The phosphorylated receptor is internalized through endocytosis via clathrin-coated pits which involves beta-arrestins. The activation of the ERK pathway occurs either in a G-protein-dependent or a beta-arrestin-dependent manner and is regulated by agonist-specific receptor phosphorylation. Acts as a class A G-protein coupled receptor (GPCR) which dissociates from beta-arrestin at or near the plasma membrane and undergoes rapid recycling. Receptor down-regulation pathways are varying with the agonist and occur dependent or independent of G- protein coupling. Endogenous ligands induce rapid desensitization, endocytosis and recycling whereas morphine induces only low desensitization and endocytosis. Heterooligomerization with other GPCRs can modulate agonist binding, signaling and trafficking properties. Involved in neurogenesis. Isoform 12 couples to GNAS and is proposed to be involved in excitatory effects (PubMed:20525224). Isoform 16 and isoform 17 do not bind agonists but may act through oligomerization with binding-competent OPRM1 isoforms and reduce their ligand binding activity (PubMed:16580639).

Cellular Location

Cell membrane; Multi-pass membrane protein. Cell projection, axon

used opioids, including morphine, heroin, fentanyl, and methadone. It is also the primary receptor for endogenous opioid peptides beta-endorphin (see POMC, MIM 176830) and the enkephalins (see PENK, MIM 131330). The OPRM1 receptor is a member of the G protein-coupled receptor family (Bond et al., 1998 [PubMed 9689128]). There are at least 3 types of opioid receptors, mu, kappa (OPRK1; MIM 165196), and delta, each with a distinct pharmacologic profile.

OPRM1 Antibody (Center) - References

- Bare,L.A., et.al., FEBS Lett. 354 (2), 213-216 (1994)
Wang,J.B., et.al., FEBS Lett. 338 (2), 217-222 (1994)

{ECO:0000250|UniProtKB:P97266}.
Perikaryon
{ECO:0000250|UniProtKB:P97266}. Cell
projection, dendrite
{ECO:0000250|UniProtKB:P97266}.
Endosome
{ECO:0000250|UniProtKB:P97266}. Note=Is
rapidly internalized after agonist binding.
{ECO:0000250|UniProtKB:P97266}

Tissue Location

Expressed in brain. Isoform 16 and isoform
17 are detected in brain.

OPRM1 Antibody (Center) - Protocols

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

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)