

# Orexin Receptor 1 antibody

# Cat. No. GTX54847

Host	Rabbit
Clonality	Polyclonal
Isotype	IgG
Applications	WB, ICC/IF, IHC-P, IHC-Fr
Reactivity	Human, Mouse, Rat

References (1)
Package
50 μl

# Applications

# **Application Note**

\*Optimal dilutions/concentrations should be determined by the researcher.

Suggested dilution	Recommended dilution
WB	Assay dependent
ICC/IF	Assay dependent
IHC-P	Assay dependent
IHC-Fr	Assay dependent

Not tested in other applications.

Calculated MW 47 kDa. (Note)

Properties	
Form	Liquid
Buffer	PBS, 1% BSA
Preservative	0.05% Sodium azide
Storage	Store as concentrated solution. Centrifuge briefly prior to opening vial. For short-term storage (1-2 weeks), store at 4°C. For long-term storage, aliquot and store at -20°C or below. Avoid multiple freeze-thaw cycles.
Concentration	0.8 mg/ml (Please refer to the vial label for the specific concentration.)
Immunogen	Peptide (C)RNWKRPSEQLEAQH, corresponding to amino acid residues 256-269 (3rd intracellular loop) of rat OX1R (Accession: P56718).
Purification	Purified by antigen-affinity chromatography
Conjugation	Unconjugated



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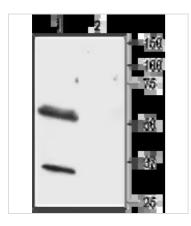


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#### Note

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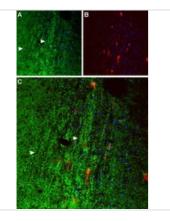
## DATA IMAGES



#### GTX54847 WB Image

WB analysis of rat brain lysate using GTX54847 Orexin Receptor 1 antibody preincubated with or without immunogen peptide.

Dilution: 1:500



## GTX54847 IHC-Fr Image

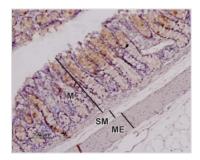
IHC-Fr analysis of mouse septum tissue using GTX54847 Orexin Receptor 1 antibody. Cell nuclei (blue) are visualized with Hoechst 33342.

Panel A: OX1R (green) appears in axonal processes (right-pointing triangles).

Panel B: Parvalbumin (red) appears in septal neurons.

Panel C: Merge of OX1R and parvalbumin suggests that orexinergic innervation covers the entire septal nucleus rather than restricted to individual neurons.

Dilution: 1:50



## GTX54847 IHC-P Image

IHC-P analysis of rat colon tissue using GTX54847 Orexin Receptor 1 antibody. Note that the stain (redbrown color) is highly specific for absorptive cells in the superior third of the intestinal glands.

Immunolabeling was detected using DAB as the chromogen and hematoxilin as the counterstain.

Dilution: 1:100



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