

# **Human Leptin R Antibody**

Recombinant Monoclonal Mouse IgG<sub>2B</sub> Clone # 52263R Catalog Number: MAB867R

DESCRIPTION			
Species Reactivity	Human		
Specificity	Detects human Leptin R in direct ELISAs.		
Source	Recombinant Monoclonal Mouse IgG <sub>2B</sub> Clone # 52263R		
Purification	Protein A or G purified from ascites		
Immunogen	Mouse myeloma cell line NS0-derived recombinant human Leptin R Phe22-Asp839 Accession # P48357		
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose. See Certificate of Analysis for details. *Small pack size (-SP) is supplied either lyophilized or as a 0.2 µm filtered solution in PBS.		

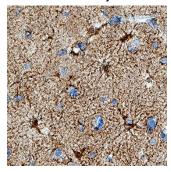
### **APPLICATIONS**

Please Note: Optimal dilutions should be determined by each laboratory for each application. General Protocols are available in the Technical Information section on our website.

	Recommended Concentration	Sample
Western Blot	1 μg/mL	Recombinant Human Leptin R Fc Chimera (Catalog # 389-LR)
Immunohistochemistry	0.5-25 μg/mL	See Below

### DATA

### Immunohistochemistry



Leptin R in Human Brain (Cortex). Leptin R was detected in immersion fixed paraffin-embedded sections of human brain (cortex) using Mouse Anti-Human Leptin R Monoclonal Antibody (Catalog # MAB867R) at 0.5 µg/mL for 1 hour at room temperature followed by incubation with the Anti-Mouse IgG VisUCyte™ HRP Polymer Antibody (Catalog # VC001). Before incubation with the primary antibody, tissue was subjected to heat-induced epitope retrieval using Antigen Retrieval Reagent-Basic (Catalog # CTS013). Tissue was stained using DAB (brown) and counterstained with hematoxylin (blue). Specific staining was localized to cell bodies and processes. View our protocol for IHC Staining with VisUCyte HRP Polymer **Detection Reagents.** 

PREPARATION AND STORAGE			
Reconstitution	Reconstitute at 0.5 mg/mL in sterile PBS.		
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below. *Small pack size (-SP) is shipped with polar packs. Upon receipt, store it immediately at -20 to -70 °C		
Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw cycles.  12 months from date of receipt, -20 to -70 °C as supplied.  1 month, 2 to 8 °C under sterile conditions after reconstitution.  6 months, -20 to -70 °C under sterile conditions after reconstitution.		

Rev. 2/17/2020 Page 1 of 2





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#### BACKGROUND

Leptin receptor (OB-R), also named B219, is a type I cytokine receptor family protein with significant amino acid sequence identity with gp130, G-CSF receptor, and the LIF receptor. Multiple isoforms of human and mouse OB-R, including a long form (OB-R<sub>L</sub>) with a large cytoplasmic domain capable of signal-transduction, and several receptor isoforms with short cytoplasmic domains (OB-R<sub>s</sub>) lacking signal-transducing capabilities, have been identified. The extracellular domains of the short and long forms of OB-R are identical. An OB-R transcript, lacking a transmembrane domain and potentially encoding a soluble form of the receptor, has also been described. OB-R<sub>L</sub> transcripts were reported to be expressed predominantly in regions of the hypothalamus previously thought to be important in body weight regulation. Expression of OB-R<sub>s</sub> transcripts have been found in multiple tissues, including the choroid plexus, lung, kidney, and primitive hematopoietic cell populations. OB-R has been shown to be encoded by the mouse diabetes (*db*) and rat fatty (*fa*) genes. Rodents homozygous for the *db* or *fa* mutations have been known to exhibit an obesity phenotype. Human OB-R long form encodes a 1165 amino acid (aa) precursor protein with a 22 aa signal peptide, an 819 aa extracellular domain, a 21 aa transmembrane domain and a 303 aa cytoplasmic domain. The extracellular domain of OB-R contain two hemopoietin receptor domains, a fibronectin type III domain and the WSXWS domain. Recombinant soluble OB-R has been shown to bind Leptin with high affinity and is a potent Leptin antagonist.

#### References:

- 1. Tartaglia, L.A. et al. (1995) Cell 83:1263.
- 2. Cioffi, J.A. et al. (1996) Nature Medicine 2:585.
- 3. Tartaglia, L.A. (1997) J. Biol. Chem. 272:6093.

Rev. 2/17/2020 Page 2 of 2

