

Mouse CCL20/MIP-3α Antibody

Monoclonal Rat IgG₁ Clone # 114908 Catalog Number: MAB7601

DESCRIPTION	
Species Reactivity	Mouse
Specificity	Detects mouse CCL20/MIP-3a in ELISAs. Does not cross-react with recombinant human (rh) CCL19, rhCCL20, recombinant mouse (rm) CCL3, 4, 9, 19, rmCXCL2, or recombinant rat CCL20.
Source	Monoclonal Rat IgG ₁ Clone # 114908
Purification	Protein A or G purified from hybridoma culture supernatant
Immunogen	E. coli-derived recombinant mouse CCL20/MIP-3α Ala28-Met97 Accession # O89093
Endotoxin Level	<0.10 EU per 1 μg of the antibody by the LAL method.
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.

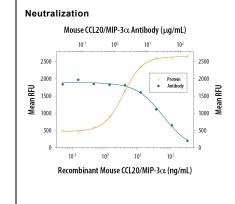
APPLICATIONS

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.

Mouse CCL20/MIP-3α Sandwich Immunoassay		Reagent
ELISA Capture	2-8 μg/mL	Mouse CCL20/MIP-3α Antibody (Catalog # MAB7601)
ELISA Detection	0.1-0.4 μg/mL	Mouse CCL20/MIP-3α Biotinylated Antibody (Catalog # BAF760)
Standard		Recombinant Mouse CCL20/MIP-3α (Catalog # 760-M3)
Neutralization	, ,	to neutralize CCL20/MIP-3α-induced chemotaxis in the BaF3 mouse pro-B cell line CCR6. The Neutralization Dose (ND ₅₀) is typically 3-15 μg/mL in the presence of 20 ng/mL

Recombinant Mouse CCL20/MIP-3a

DATA



Chemotaxis Induced by CCL20/MIP-3α and Neutralization by Mouse CCL20/MIP-3α Antibody. Recombinant Mouse CCL20/MIP-3α (Catalog # 760-M3) chemoattracts the BaF3 mouse pro-B cell line transfected with human CCR6 in a dosedependent manner (orange line). The amount of cells that migrated through to the lower chemotaxis chamber was measured by Resazurin (Catalog # AR002). Chemotaxis elicited by Recombinant Mouse CCL20/MIP-3a (20 ng/mL) is neutralized (green line) by increasing concentrations of Rat Anti-Mouse CCL20/MIP-3α Monoclonal Antibody (Catalog # MAB7601). The ND₅₀ is typically 3-15 μg/mL.

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.		

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BACKGROUND

MIP-3 α , also known as LARC (Liver and Activation-regulated Chemokine) and as Exodus, is one of many novel β chemokines identified through bioinformatics. Mouse MIP-3 α cDNA encodes a 97 amino acid residue precursor protein with a 27 aa residue putative signal peptide that is predicted to be cleaved to form the 70 amino acid (aa) residue mature secreted protein. MIP-3 α is distantly related to other β chemokines (20-28% aa sequence identity). Mouse MIP-3 α shares approximately 71% and 63% amino acid sequence homology with rat and human MIP-3 α , respectively.

MIP-3α has been shown to be expressed predominantly in lymph nodes, appendix, PBL, fetal liver, fetal lung, and epithelial cells of intestinal tissues. The expression of MIP-3α is strongly up-regulated by inflammatory signals and down-regulated by the anti-inflammatory cytokine IL-10. Synthetic or recombinant MIP-3α has been shown to be chemotactic for lymphocytes and dendritic cells, and inhibits proliferation of myeloid progenitors in colony formation assays. MIP-3α has now been shown to be a unique functional ligand for CCR-6 (previously referred to as GPR-CY4, CKR-L3, or STRL22 orphan receptor), a chemokine receptor that is selectively and highly expressed in human dendritic cells derived from CD34⁺ cord blood precursors.

References:

- 1. Baba, M. et al. (1997) J. Biol. Chem. 272:14893.
- 2. Hromas, R. et al. (1997) Blood 89:3315.
- 3. Greaves, D.R. et al. (1997) J. Exp. Med. 186: 857.
- 4. Tanaka, Y. et al. (1999) Eur. J. Immunol. 29:633.