

Mouse CCL7/MCP-3/MARC Antibody

Antigen Affinity-purified Polyclonal Goat IgG Catalog Number: AF-456-NA

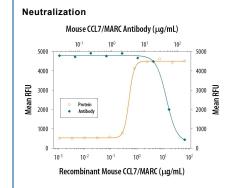
DESCRIPTION			
Species Reactivity	Mouse		
Specificity	Detects mouse CCL7/MCP-3/MARC in direct ELISAs and Western blots. In direct ELISAs, approximately 30% cross-reactivity with recombinant human (rh) MCP-1 is observed, approximately 15% cross-reactivity with rhEotaxin is observed, and less than 3% cross-reactivity with recombinant mouse Eotaxin, rhMARC, and rhMCP-2 is observed.		
Source	Polyclonal Goat IgG		
Purification	Antigen Affinity-purified		
Immunogen	E. coli-derived recombinant mouse CCL7/MCP-3/MARC Gln24-Pro97 Accession # Q03366		
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.		

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	0.1 μg/mL	Recombinant Mouse CCL7/MCP-3/MARC (Catalog # 456-MC)	
Neutralization	Measured by its ability to neutralize CCL7/MCP-3/MARC-induced chemotaxis in the BaF3 mouse pro-B cell line transfected with human CCR2A. The Neutralization Dose (ND $_{50}$) is typically 15-45 μ g/mL in the presence of		
	2 µg/mL Recombinant Mouse CCL7/MCP-3/MARC.		

DATA



Chemotaxis Induced by CCL7/MARC and Neutralization by Mouse CCL7/MARC Antibody. Recombinant Mouse CCL7/ MARC (Catalog # 456-MC) chemoattracts the BaF3 mouse pro-B cell line transfected with human CCR2A 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 CCL7/ MARC (2 µg/mL) is neutralized (green line) by increasing concentrations of Mouse CCL7/MARC Antigen Affinitypurified Polyclonal Antibody (Catalog # AF-456-NA). The ND₅₀ is typically 15-45 µg/mL.

PREPARATION AND STORAGE

 Reconstitution
 Reconstitute at 0.2 mg/mL in sterile PBS.

 Shipping
 The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.

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

Mouse MARC, a member of the β subfamily of chemokines, was initially identified as a transcript that is induced in a mouse mast cell line after Fc epsilon RI triggering by IgE plus antigen. Sequence comparisons suggest that MARC may be the mouse homologue of the human MCP-3 gene. Mouse MARC/MCP-3 expression has also been detected during murine experimental allergic encephalomyelitis in the spinal cord, and in LPS-stimulated murine WEHI -3 cells and Swiss 3T3 cells where MARC expression is glucocorticoid-attenuated. Except for one amino acid substitution, mouse MARC is identical to mouse FIC, the product of a growth factor-activated gene. The mouse MARC cDNA encodes a 97 amino acid residue precursor protein with a 23 amino acid residue signal peptide that is cleaved to yield a 74 amino acid residue mature protein. Mouse CCR2, a mouse chemokine receptor, has been shown to bind JE/MCP-1 with high affinity and MARC/MCP-3 with lower affinity. The *E. coli*-expressed mouse MARC/MCP-3 produced at R&D Systems has been shown to be a monocyte and T-lymphocyte chemoattractant.

References:

- 1. Kulmburg, P.A. et al. (1992) J. Exp. Med. 176:1773.
- 2. Thirion, S. et al. (1994) Biochem. Biophys. Res. Commun. 201:493.
- 3. Smith, J.B. and H.R. Herschman (1995) J. Biol. Chem. 270:16756.
- 4. Kurihara, T. and R. Bravo (1996) J. Biol. Chem. 271:11603.



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