

## MCP-1 Antibody (Carboxy-terminal Antigen)



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## For Research Use Only. Not for Use in Diagnostic Procedures.

<b>Applications:</b> W, IP	Reactivity:	<b>Sensitivity:</b> Endogenous	<b>MW (kDa):</b> 13-15	Source/Isotype: Rabbit	UniProt ID: #P13500	Entrez-Gene Id: 6347
Product Usage Information	•	Application Western Blotting Immunoprecipitation	.5 ,5		<b>Dilution</b> 1:1000 1:50	30 //
Storage		Supplied in 10 mM sodium HEPES (pH 7.5), 150 mM NaCl, 100 $\mu$ g/ml BSA and 50% glycerol. Store at – 20°C. Do not aliquot the antibody.				
Specificity/Sensitivity		MCP-1 Antibody (Carboxy-terminal Antigen) recognizes endogenous levels of total MCP-1 protein.				
Source / Purification		Polyclonal antibodies are produced by immunizing animals with a synthetic peptide corresponding to residues near the carboxy terminus of human MCP-1 protein. Antibodies are purified by protein A and peptide affinity chromatography.				
Background		Monocyte chemoattractant protein-1 (MCP-1), also known as CCL2, monocyte chemotactic activating factor (MCAF) or glioma-derived chemotactic factor-2 (GDCF-2), is the product of the human $JE$ gene and a member of the family of C-C (or $\beta$ ) chemokines (1-4). The predicted molecular weight of MCP-1 protein is 11-13 kDa, but it may migrate at 20-30 kDa due to glycosylation. MCP-1 is secreted by a variety of cell types in response to pro-inflammatory stimuli and was originally described for its chemotactic activity on monocytes. This activity has led to studies demonstrating its role in diseases characterized by monocyte infiltrates such as psoriasis (5), rheumatoid arthritis (6) and atherosclerosis (7). MCP-1 may also contribute to tumor progression and angiogenesis (8). Signaling by MCP-1 is mediated by the G protein-coupled receptor CCR2 (9).				
Background References		<ol> <li>Matsushima, K. et al. (1989) J Exp Med 169, 1485-90.</li> <li>Furutani, Y. et al. (1989) Biochem Biophys Res Commun 159, 249-55.</li> <li>Robinson, E.A. et al. (1989) Proc Natl Acad Sci USA 86, 1850-4.</li> <li>Rollins, B.J. et al. (1988) Proc Natl Acad Sci USA 85, 3738-42.</li> <li>Gillitzer, R. et al. (1993) J Invest Dermatol 101, 127-31.</li> <li>Koch, A.E. et al. (1992) J Clin Invest 90, 772-9.</li> <li>Ylä-Herttuala, S. et al. (1991) Proc Natl Acad Sci USA 88, 5252-6.</li> <li>Salcedo, R. et al. (2000) Blood 96, 34-40.</li> <li>Charo, I.F. et al. (1994) Proc Natl Acad Sci USA 91, 2752-6.</li> </ol>				

**Species Reactivity** Species reactivity is determined by testing in at least one approved application (e.g., western blot).

Western Blot Buffer IMPORTANT: For western blots, incubate membrane with diluted primary antibody in 5% w/v nonfat

dry milk, 1X TBS, 0.1% Tween® 20 at 4°C with gentle shaking, overnight.

Applications Key W: Western Blotting IP: Immunoprecipitation

Cross-Reactivity Key H: Human

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