

Human CCL14a/HCC-1 Antibody

Antigen Affinity-purified Polyclonal Goat IgG Catalog Number: AF-324-PB

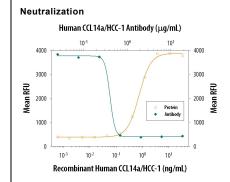
DESCRIPTION		
Species Reactivity	Human	
Specificity	Detects human CCL14a/HCC-1 in direct ELISAs and Western blots. In direct ELISAs and Western blots (non-reducing conditions), less than 5% cross-reactivity with recombinant human (rh) MCP-2, rhMCP-3, recombinant mouse (rm) C10, and rmMIP-1β is observed.	
Source	Polyclonal Goat IgG	
Purification	Antigen Affinity-purified	
Immunogen	E. coli-derived recombinant human CCL14a/HCC-1 Thr20-Asn93 Accession # Q16627	
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.

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	Recommended Concentration	Sample	
Western Blot	0.1 μg/mL	Recombinant Human CCL14a/HCC-1 aa 20-93 (Catalog # 324-HC)	
Neutralization	, ,	v to neutralize CCL14a/HCC-1-induced chemotaxis in the BaF3 mouse pro-B cell line n CCR1. The Neutralization Dose (ND ₅₀) is typically 0.15-0.75 µg/mL in the presence of	
	10 ng/mL Recombinar	nt Human CCL14a/HCC-1 aa 28-93.	

DATA



Chemotaxis Induced by CCL14a/HCC-1 and Neutralization by Human CCL14a/HCC-1 Antibody. Recombinant Human CCL14a/HCC-1 (Catalog # 1578-HC) chemoattracts the BaF3 mouse pro-B cell line transfected with human CCR1 in a dose-dependent 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 Human CCL14a/HCC-1 (10 ng/mL) is neutralized (green line) by increasing concentrations of Goat Anti-Human CCL14a/ HCC-1 Antigen Affinity-purified Polyclonal Antibody (Catalog # AF-324-PB). The ND₅₀ is typically 0.15-0.75 $\mu g/mL$.

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. *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

HCC-1 (Hemofiltrate CC Chemokine-1) was originally isolated from the hemofiltrate of human patients with chronic renal failure (1). It belongs to the CC chemokine superfamily and has been designated CCL14a. HCC-1/CCL14a cDNA encodes a 93 amino acid (aa) residue precursor with a 19 aa signal peptide that is cleaved to form the 74 aa secreted protein (aa 20-93). By alternative splicing, a second longer isoform named HCC-3/CCL14b, which includes sequences from exon 3, also exists (2). HCC-1/CCL14a is expressed constitutively in various normal tissues including spleen, liver, muscle, gut and bone marrow. It circulates at nanomolar concentrations in human plasma. Different post-translationally modified HCC-1/CCL14a, including O-glycosylated and N-terminally truncated variants of HCC-1/CCL14a, have been identified (3, 4). Whereas the 74 aa peptide is a weak CCR1 agonist, the proteolytically processed, truncated HCC-1/CCL14a (aa 28-93) is a highly potent agonist of CCR1, CCR5 and to a lesser extent, CCR3. HCC-1/CCL14a (aa 28-93) promotes chemotaxis of T lymphocytes, monocytes and eosinophils, and inhibits infection of M-tropic human immunodeficiency virus type 1. Activation of the HCC-1/CCL14a precursor to active peptide is mediated by the urokinase type plasminogen activator or plasmin (5).

References:

- 1. Schulz-Knappe, P. et al. (1996) J. Exp. Med. 183:295.
- 2. Forssmanns, U. et al. (2001) J. Leukocyte Biology 70:357.
- 3. Richter, R. et al. (2000) Biochemistry 39:10799.
- 4. Munch, J. et al. (2002) Antimicrob. Agents Chemother. 46:982.
- 5. Vakili, J. et al. (2001) J. Immunol. 167:3406.



