

# Datasheet for ABIN116167

## anti-CCL4 antibody

# Publication



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Quantity:	50 μg	
Target:	CCL4	
Reactivity:	Human	
Host:	Rabbit	
Clonality:	Polyclonal	
Conjugate:	This CCL4 antibody is un-conjugated	
Application:	Western Blotting (WB), Immunohistochemistry (Paraffin-embedded Sections) (IHC (p)), Enzyme Immunoassay (EIA), Functional Studies (Func)	

### **Product Details**

Immunogen:	Highly pure (>98%) E.coli derived recombinant Human MIP-1 beta (CatNo PA107)
Specificity:	Recognizes Human Macrophage Inflammatory Protein-1 beta (MIP-1b). Other species not tested.
Purification:	Affinity Chromatography employing immobilized Human MIP-1beta matrix.

### **Target Details**

Target:	CCL4
Alternative Name:	MIP1 beta / CCL4 (CCL4 Products)
Background:	MIP1 alpha and MIP1 beta were originally co-purified from medium conditioned by an LPS- stimulated murine macrophage cell line. Human MIP1 beta refers to the products of several
	independently cloned cDNAs, including Act2, PAT 744, hH400, G26, HIMAP, HC21, and MAD 5a.

The predicted protein products of these cDNAs represent variants that are between 94 % - 98 % identical and these proteins are all approximately 75 % homologous to murine MIP1 beta. MIP1 beta also shares approximately 70 % amino acid identity with MIP1 alpha. MIP1 proteins are expressed primarily in T cells, B cells, and monocytes after antigen or mitogen stimulation. The MIP1 proteins have chemoattractant and adhesive effects on lymphocytes, with MIP1 alpha and MIP1 beta preferentially attracting CD8+ and CD4+ T cells, respectively. A signal transducing receptor designated the CC chemokine receptor 1 (CC CKR1) with seven transmembrane domains that binds MIP1 alpha, MIP1 beta, MCP1 and RANTES with varying affinities has been isolated. Synonyms: CCL-4, G-26 T-lymphocyte-secreted protein, HC21, LAG1, Lymphocyte activation gene 1 protein, MIP-1-beta, MIP1B, Macrophage inflammatory protein 1-beta, PAT744, Protein H400, SCYA4, SIS-gamma, Small-inducible cytokine A4, T-cell activation protein 2

Gene ID:

388372

NCBI Accession:

NP\_001001435

UniProt:

P13236

#### **Application Details**

**Application Notes:** 

Neutralization: To yield one-half maximal inhibition [ND50] of the biological activity ofhMIP-1 beta (50.00 ng/mL), a concentration of 1.75-3.00  $\mu$ g/mL of this antibody is required. ELISA: To detect hMIP-1 beta by direct ELISA (using 100l/well antibody solution) aconcentration of at least 0.5  $\mu$ g/mL of this antibody is required. This antigen affinitypurified antibody, in conjunction with compatible secondary reagents, allows the detection of 0.2-0.4 ng/well of recombinant hMIP-1 beta. Western Blot: To detect hMIP-1 beta by Western Blot analysis this antibody can be used at a concentration of 0.1-0.2  $\mu$ g/mL. Used in conjunction with compatible secondary reagents the detection limit for recombinant hMIP-1 beta is 1.5-3.0 ng/lane, under either reducing ornon-reducing conditions.

Other applications not tested.

Optimal dilutions are dependent on conditions and should be determined by the user.

Restrictions:

For Research Use only

#### Handling

Reconstitution:

Restore in sterile water to a concentration of 0.1-1.0 mg/mL.

Buffer:

PBS, pH 7.2 without preservatives

### Handling

Preservative:	Without preservative	
Handling Advice:	Avoid repeated freezing and thawing. Centrifuge vial prior to opening!	
Storage:	4 °C/-20 °C	
Storage Comment:	Prior to reconstitution store at 2-8 °C. Following reconstitution store the antibody at -20 °C.	
Publications		
Product cited in:	Schlaepfer, Audigé, Joller, Speck: "TLR7/8 triggering exerts opposing effects in acute versus	
	latent HIV infection." in: Journal of immunology (Baltimore, Md.: 1950), Vol. 176, Issue 5, pp.	
	2888-95, (2006) (PubMed).	