

## Cytokines

### Human Recombinant MIP-1 beta (CCL4)

Macrophage inflammatory protein-1  
beta



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Catalog # 78090  
78090.1

5 µg  
25 µg

## Product Description

Macrophage inflammatory protein-1 beta (MIP-1 beta), also known as CCL4, is a member of CC family of chemokines and is most closely related to CCL3 or MIP-1 alpha. MIP-1 beta is predominantly synthesized by activated CD8+ T cells, specifically the perforin-low memory CD8+ T cells (Kamin-Lewis et al.). MIP-1 beta signals through CCR5 which is the major coreceptor required for entry of certain strains of HIV-1 into permissive cells and it is known that HIV-1-specific CD8+ T cell clones that exhibit cytotoxic T lymphocyte (CTL) activity secrete MIP-1 beta upon encounter with sensitized target cells (Menten et al.). MIP-1 beta attracts a variety of immune cells to sites of microbial infection. In vitro experiments show that human MIP-1 beta tends to attract CD4+ T lymphocytes, preferentially of the naive (CD45RA) phenotype (Schall et al.). In addition to its chemotactic functions, MIP-1 beta induces the release of proinflammatory cytokines, mast cell degranulation, and NK cell activation (Schall et al.). It has been shown that there is an increased expression of MIP-1 beta in CD8+ and CD4+ T cells on the site of inflammation in sarcoidosis patients (Barczyk et al.).

## Product Information

**Alternative Names:** ACT-2, Immune activation protein 2, LAG-1, Lymphocyte activation gene 1 protein, MIP-1b, Protein H400, SCYA2, SCYA4, Small-inducible cytokine A4, T-cell activation protein 2

**Accession Number:** P13236

**Amino Acid Sequence:** APMGSDPPTA CCFSYTARKL PRNFVVDYYE TSSLCSQPAV VFQTKRSKQV CADPSESWVQ EYVYDLELN

**Predicted Molecular Mass:** 7.6 kDa

**Species:** Human

**Cross Reactivity:** Mouse, Rat

**Formulation:** Lyophilized after dialysis against phosphate-buffered saline.

**Source:** E. coli

## Specifications

**Activity:** The specific activity is  $\geq 1 \times 10^4$  units/mg ( $EC_{50} \leq 0.1 \mu\text{g/mL}$ ) as determined by  $Ca^{2+}$  mobilization assay in CHO-K1/Gα15/hCCR5 cells (human Gα15 and human CCR5 stably expressed in CHO-K1 cells).

**Purity:**  $\geq 95\%$

**Endotoxin Level:** Measured by kinetic limulus amoebocyte lysate (LAL) analysis and is  $\leq 0.2$  EU/µg protein.

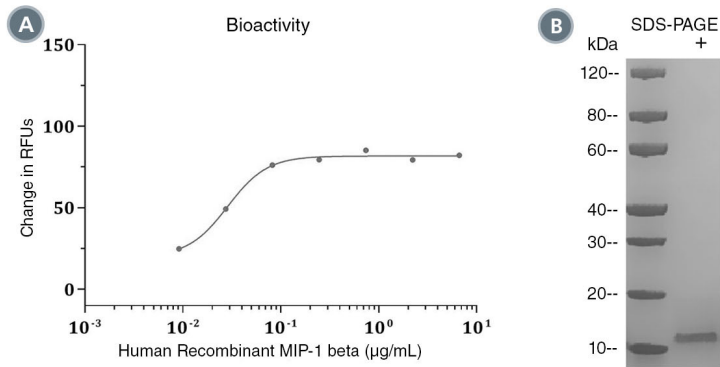
## Preparation and Storage

**Storage:** Store at  $-80^\circ\text{C}$ .

**Stability:** Stable as supplied for 12 months from date of receipt.

**Preparation:** Centrifuge vial before opening. Resuspend the product in sterile water or phosphate-buffered saline to at least 0.1 mg/mL by pipetting the solution down the sides of the vial. Do not vortex. Store at  $2 - 8^\circ\text{C}$  for up to 2 weeks or at  $-20^\circ\text{C}$  to  $-80^\circ\text{C}$  for up to 3 months. Avoid repeated freeze-thaw cycles.

## Data



(A) The biological activity of Human Recombinant MIP-1 beta (CCL4) was tested by its ability to mobilize Ca<sup>2+</sup> in CHO-K1/Gα15/hCCR5 cells (human Gα15 and human CCR5 stably expressed in CHO-K1 cells). Ca<sup>2+</sup> mobilization was measured using a fluorometric assay method. The EC<sub>50</sub> is defined as the effective concentration of the growth factor at which Ca<sup>2+</sup> mobilization is at 50% of maximum. The EC<sub>50</sub> in the example above is less than 0.1 µg/mL.

(B) 1 µg of Human Recombinant MIP-1 beta (CCL4) was resolved with SDS-PAGE under reducing (+) conditions and visualized by Coomassie Blue staining. Human Recombinant MIP-1 beta (CCL4) has a predicted molecular mass of 7.6 kDa.

## Related Products

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## References

- Barczyk A et al. (2014) Increased expression of CCL4/MIP-1β in CD8+ cells and CD4+ cells in sarcoidosis. *Int J Immunopathol Pharmacol* 27(2): 185–93.
- Kamin-Lewis R et al. (2001) Perforin-low memory CD8+ cells are the predominant T cells in normal humans that synthesize the beta-chemokine macrophage inflammatory protein-1beta. *Proc Natl Acad Sci U S A* 98(16): 9283–8.
- Menten P et al. (2002) Macrophage inflammatory protein-1. *Cytokine Growth Factor Rev* 13(6): 455–81.
- Schall TJ et al. (1993) Human macrophage inflammatory protein alpha (MIP-1 alpha) and MIP-1 beta chemokines attract distinct populations of lymphocytes. *J Exp Med* 177(6): 1821–6.

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