## **Product Data Sheet**

## **Purified anti-human IL-4**

**Catalog # / Size:** 3104010 / 500 μg

Clone: MP4-25D2 Isotype: Rat IgG1, κ

Immunogen: CHO-expressed, recombinant human IL-

4

Reactivity: Human

**Preparation:** The antibody was purified by affinity

chromatography.

**Formulation:** Phosphate-buffered solution, pH 7.2,

containing 0.09% sodium azide.

Concentration: 0.5

## **Applications:**

**Applications:** Other

Recommended

Usage:

Each lot of this antibody is quality control tested by ELISA assay. Purified MP4-25D2 has been tested by blocking fluorochrome conjugated MP4-25D2 for intracellular cytokine staining. In order to obtain complete blocking results, a saturated amount of purified antibody (≤ 5.0 microg/million cells) should be used for incubation with target cells, prior to staining with fluorochrome conjugated antibody. It is recommended that the reagent be titrated for optimal performance for each application.

Application Notes:

**ELISA Detection**<sup>1,3</sup> **or ELISPOT Detection**<sup>4,5</sup>: The biotinylated MP4-25D2 antibody is useful as a detection antibody for a sandwich ELISA or ELISPOT assay, when used in conjunction with purified 8D4-8 antibody (Cat. No. 500702/500707) as the capture antibody.

**Flow Cytometry**<sup>6,9</sup>: The fluorochrome-labeled MP4-25D2 antibody is useful for intracellular immunofluorescent staining and flow cytometric analysis to identify IL-4 -producing cells within mixed cell populations.

**Neutralization**<sup>1-3</sup>: The LEAF<sup>TM</sup> purified antibody (Endotoxin <0.1 EU/ $\mu$ g, Azide-Free, 0.2  $\mu$ m filtered) is recommended for neutralization of human IL-4 bioactivity (Cat. No. 500815). The MP4-25D2 antibody can neutralize the bioactivity of natural or recombinant IL-4.

Application References:

- 1. Chretien I, et al. 1989. J. Immunol. Methods 117:67. (ELISA Detection, Neut)
- 2. Ramanathan L, et al. 1993. Biochem. 32:3549. (Neut)
- 3. Abrams J, *et al.* 1992. *Immunol. Rev.* 127:5. (ELISA Detection, Neut) 4. Mahanty S, *et al.* 1992. *J. Immunol.* 148:3567. (ELISPOT Detection)
- 5. Klinman D, et al. 1994. Curr. Prot. Immunol. John Wiley and Sons New York. Unit

6.19. (ELISPOT Detection)

- 6. Prussin C, et al. 1995. J. Immunol. Methods 188:117. (ICFC)
- 7. Raqib R, et al. 1995. Infect. Immun. 63:289. 8. Andersson J, et al. 1994. Immunology 83:16.
- 9. Iwamoto S, et al. 2007. J. Immunol. 179:1449. (ICFC) PubMed
- 10. Kubota M, et al. 1997. J. Immunol. 158:5321.
- 11. Dzhagalov I, et al. 2007. J. Immunol. 178:2113. PubMed
- 12. Kroneke MA, et al. 2012. J. Immunol. 188:3734. PubMed

**Description:** IL-4 is a pleiotropic cytokine that is produced by activated T cells, mast cells, and

basophils. IL-4 elicits many different biological responses but has two dominant

functions. The first is regulating differentiation of naı̈ve CD4<sup>+</sup> T cell to the Th2 type. Th2 cells produce IL-4, IL-5, IL-10, and IL-13, which tend to favor a humoral immune response while suppressing a cell-mediated immune response controlled by Th1 cells. The second is regulating IgE and IgG1 production by B cells.

## Antigen References:

- 1. Fitzgerald K, *et al.* Eds. 2001. The Cytokine FactsBook. Academic Press San Diego.
- 2. Boulay J, et al. 1992. Curr. Opin. Immunol. 4:294.
- 3. Dullens H, et al. 1991. In vivo 5:567.
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