

Product Data Sheet

PerCP/Cy5.5 anti-mouse IL-2

Catalog # / Size: 503821 / 25 µg

503822 / 100 µg

Clone: JES6-5H4 **Isotype:** Rat IgG2b, κ

Immunogen: E. coli-expressed, recombinant mouse IL-2

Reactivity: Mouse

Preparation: The antibody was purified by affinity chromatography, and conjugated with

PerCP/Cy5.5 under optimal conditions. The solution is free of unconjugated

PerCP/Cy5.5 and unconjugated antibody.

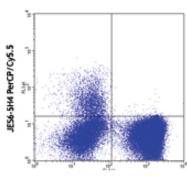
Formulation: Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and

0.2% (w/v) BSA (origin USA).

Concentration: 0.2 mg/ml

Storage: The antibody solution should be stored undiluted at 4°C and protected from

prolonged exposure to light. Do not freeze.



B220 (RA3-6B2) APC

PMA+ionomycin-stimulated (6 hours) C57BL/6 mouse splenocytes intracellular stained with B220 (RA3-6B2) APC and JES6-5H4 PerCP/Cv5.5

Applications:

Applications: ICFC - Quality tested

Recommended Usage: Each lot of this antibody is quality control tested by intracellular immunofluorescent staining with flow cytometric analysis. For immunofluorescent staining, the suggested use of this reagent is 5 µl per million cells or 5 µl per 100 µl of whole blood. It is recommended that the reagent be titrated for optimal performance for each application.

* PerCP/Cy5.5 has a maximum absorption of 482 nm and 564 nm and a maximum emission of 690 nm.

Application Notes: ELISA Detection¹⁻³ or ELISPOT Detection⁴⁻⁶: The biotinylated JES6-5H4 antibody is useful as a detection antibody for a sandwich ELISA or ELISPOT assay, when used in conjunction with the purified JES6-1A12 antibody (Cat. No. 503702/503704) as capture antibody and recombinant mouse IL-2 (Cat. No. 575409) as the standard.

Flow Cytometry⁸⁻¹⁰: The fluorochrome-labeled JES6-5H4 antibody is useful for intracellular immunofluorescent

staining and flow cytometric analysis to identify IL-2 -producing cells within mixed cell populations.

Neutralization¹,7: The LEAF™ purified antibody (Endotoxin in vivo and *in vitro* (Cat. No. 503812) is recommended for

Additional reported applications (for the relevant formats) include: immunoprecipitation¹, immunohistochemical staining of paraformaldehyde-fixed, saponin-treated frozen tissue sections², *in vivo* capture⁷, and

immunocytochemistry.

Note: For testing mouse IL-2 in serum, plasma or supernatant, BioLegend's ELISA Max™ Sets (Cat. No. 431001 to 431006) are specially developed and recommended.

Cy3, Cy5, Cy5.5 and Cy7 are subject to proprietary rights of GE Healthcare Bio-Sciences Corp. and Carnegie Mellon University and made and sold under license from GE Healthcare Bio-Sciences Corp. Sale of this product is licensed for research use only.

- Application References: 1. Abrams J, et al. 1992. Immunol. Rev. 127:5.
 - 2. Sander B, et al. 1993. J. Immunol. Meth. 166:201.
 - 3. Abrams J. 1995. Curr. Prot. Immunol. John Wiley and Sons New York. Unit 6.20.
 - 4. Klinman D, et al. 1994. Curr. Prot. Immunol. John Wiley and Sons New York. Unit 6.19. 5. Mo X, et al. 1995. J. Virol. 69:1288.

 - 6. Karulin A, et al. 2000. J. Immunol. 164:1862.
 - 7. Finkelman F, et al. 2003. Curr. Prot. Immunol. John Wiley & Sons New York. Unit 6.28. 8. Ko SY, et al. 2005. J. Immunol. 175:3309. PubMed

 - 9. Kang SS and Allen PM. 2005. J. Immunol. 174:5382.

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 Lawson BR, et al. 2007. J. Immunol. 178:5366.
 Aldhamen YA, et al. 2012. J. Immunol. 189:1349. PubMed.
 Yamamoto M, et al. 2013. Cytokine. 61:831. PubMed.

Description: IL-2 is a potent lymphoid cell growth factor which exerts its biological activity primarily on T cells. Additionally, IL-2 has been found to stimulate growth and differentiation of B cells, NK cells, LAK cells, monocytes, and oligodendrocytes.

- Antigen References: 1. Fitzgerald K, et al. Eds. 2001. The Cytokine FactsBook. Academic Press San Diego.

 - Taniguchi T, et al. 1993. Cell 73:5.
 Nistico G. 1993. Prog. Neurobiol. 40:463.
 - 4. Waldmann T, et al. 1993. Ann. NY Acad. Sci. 685:603.



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Related Products: Product

Cell Staining Buffer PerCP/Cy5.5 Rat IgG2b, κ Isotype Ctrl

Clone

RTK4530

Application FC, ICC, ICFC FC, ICFC



