

# ANTI-HUMAN INTERLEUKIN-2 SOLUBLE RECEPTOR $\beta$ (IL-2 sR $\beta$ ) Developed in Goat, Affinity Isolated Antibody

Product Number I6277

## **ProductInformation**

#### **Product Description**

Anti-Human Interleukin-2 Soluble Receptor Beta (IL-2 sR $\beta$ ) is developed in goat using a recombinant human IL-2 sR $\beta$ ), expressed in Sf21 cells as immunogen. The antibody is purified using IL-2 R $\beta$  affinity chromatography.

Anti-Human IL-2 sR $\beta$  may be used to neutralize human cell surface IL-2 R $\beta$  mediated-bioactivity. For best results using cells expressing the high affinity IL-2 receptors, the use of both anti-IL-2 R $\alpha$  (Product No. I6152) and anti-IL-2 R $\beta$  is recommended. By ELISA and immunoblotting the antibody shows < 15% cross-reactivity with recombinant human IL-3 sR $\alpha$ , < 5% cross-reactivity with recombinant human IL-1 RII, IL-2 sR $\gamma$ , IL-5 sR $\alpha$ , and < 2% cross-reactivity with recombinant human IL-5 sR $\beta$ . In addition, the antibody shows no cross-reactivity with other cytokines tested.\*

Anti-Human IL-2 sR $\beta$  may be used for neutralization of the biological activity mediated by IL-2 R $\beta$  and for the detection of IL-2 R $\beta$  by immunoblotting and ELISA.

The biological effects of IL-2R signals are much more complex than simply mediating T-cell growth. Depending on the set of conditions, IL-2R signals may also promote cell survival, effector function, and apoptosis. These sometimes contradictory effects underscore the fact that a diversity of intracellular signaling pathways are potentially activated by IL-2R. There are at least 3 components of the IL-2 receptor, IL-2 R $\alpha$ , IL-2  $\beta$ R, and IL-2 R $\gamma$  chains. The IL-2 R $\gamma$  chain is shared by IL-2, IL-4 and IL-7. The low affinity  $\alpha$ chain is a 55 kD polypeptide. It is incapable of transmitting intracellular signals due to its short cytoplasmic tail. However, it can bind IL-2 rapidly to the cell membrane. The  $\beta$  chain (75 kD) and the  $\gamma$  chain (64 kD) form a complex that can bind IL-2 with high affinity and slow dissociation and can mediate signal transduction.

Cells known to express the β-chain include: activated CD56<sup>+</sup> (NK) cells plus CD8<sup>+</sup> and CD4<sup>+</sup> T cells,<sup>3,4</sup> resting NK cells and, perhaps, CD8<sup>+</sup> T cells,<sup>3,4</sup> activated and

resting B cells, <sup>5</sup> mature thymocytes, <sup>6</sup> embryonic fibroblasts, <sup>7</sup> resting monocytes <sup>8</sup> and neutrophils. <sup>9</sup>

#### Reagents

The product is supplied lyophilized from a  $0.2 \mu m$  filtered solution in phosphate buffered saline. Endotoxin level is < 10 ng per mg antibody as determined by the LAL method.

#### **Preparation Instructions**

To one vial of lyophilized powder, add 1 ml of 0.2 µm-filtered PBS to produce a 0.1 mg/ml stock solution of antibody. If aseptic technique is used, no further filtration should be needed for use in cell culture environments.

#### Storage/Stability

Prior to reconstitution, store at -20°C. Reconstituted product may be stored at 2-8°C for at least one month. For prolonged storage, freeze in working aliquots at -20°C. Avoid repeated freezing and thawing.

### **Procedure**

Anti-Human IL-2 sR $\beta$  is tested for its ability to neutralize human cell surface IL-2 R $\beta$  mediated IL-2 bioactivity in a  $^3$ H-thymidine incorporation assay using MO7e cells.  $^{10}$  The ND $_{50}$  of the antibody is defined as the concentration of antibody resulting in a one-half maximal inhibition of the cell surface IL-2 R $\beta$  mediated recombinant human IL-2 response on a responsive cell line.

#### **Product Profile**

For neutralization, a working concentration of 10-30  $\mu$ g/ml of Anti-Human IL-2 sR $\beta$  will block 50% of the bioactivity due to 30 ng/ml recombinant human IL-2 in a  $^3$ H-thymidine incorporation assay using 10 $^5$ /ml MO7e cells.

For Indirect Immunoblotting, a working concentration of 1-2  $\mu$ g/ml is determined using recombinant human IL-2 sR $\beta$  at 1 ng/lane under non-reducing conditions and 0.5 ng/ml under reducing conditions.

For Indirect ELISA, a working concentration of 0.5 - 1  $\mu$ g/ml is determined to detect recombinant IL-2 sR $\beta$  to a limit of 0.15 ng/well.

Note: In order to obtain best results in different techniques and preparations we recommend determining optimal working dilutions by titration test.

- References
  1. Noguchi, M., et al., Science, **262**, 1877 (1993)
- 2. Russel, S.M., et al., Science, 262, 1880 (1993).
- 3. Nakamura, Y., et al., Nature, **369**, 330 (1994).
- 4. Uchiyama, T., et al., J. Immunol., 126, 1393 (1981).
- F. Taniquahi T and Minami V Call **72** 5 (1902)
- 5. Taniguchi, T. and Minami, Y., Cell, **73**, 5 (1993).
- Vanham, G., et al., Clin. Immunol. Immunopathol., 71, 60 (1994).
- Caligiuri, M.A., et al., J. Exp. Med., 171, 1509 (1990).
- 8. Nakanishi, K., et al., Proc. Natl. Acad. Sci. USA, **89**, 3551 (1992).
- 9. Leclercq, G., et al., Int. Immunol., **7**, 843 (1995).
- 10. Hicks, C., et al., Growth Factors, 5, 201 (1991).
- 11. Plaisance, S., et al., Int. Immunol., 4, 739 (1992).
- 12. Benveniste, E.N., et al., J. Neuroimmunol., **17**, 301 (1988).
- 13. Espinoza-Delgado, I., et al., J. Leukoc. Biol., **57**, 13 (1995).
- 14. Waldmann, T.A., Science, 232, 727 (1986).
- 15. Herrmann, F., et al., J. Exp. Med., **162**, 1111 (1985).

- 16. Weidmann, E., et al., Cancer Res., **52**, 5963 (1992).
- \* rhANG, rhAR, rmB7-2, rhBTC, rhβ-NGF, rhBDNF, rmC10, rhCD4, rhCD8, rhCD28, rhCNTF, rrCNTF, rhEGF, rhENA-78, rhEpo, rhFGF acidic, rhFGF basic, rhFGF-4, rhFGF-5, rhFGF-6, rhFGF-7, rhFGF-9, rhG-CSF, rmG-CSF, rhGM-CSF, rhGM-CSF Rα, rmGM-CSF, rhGRO $\alpha$ , rhGRO $\beta$ , rhGRO $\gamma$ , rhHB-EGF, rhHRG- $\alpha$ , rhHGF, rhI-309, rhIFN- $\gamma$ , rhIGF-I, rhIGF-I R, rhlL- $1\alpha$ , rhlL-1 RI, rmlL- $1\alpha$ , rhlL- $1\beta$ , rmlL- $1\beta$ , rhlL-1ra, rmIL-1ra, rhIL-2, rmIL-2, rhIL-3, rmIL-3, rhIL-4, rmIL-4, rhIL-5, rmIL-5, rhIL-6, rhIL-6 sR, rmIL-6, rhIL-7, rhIL-7 R, rmIL-7, rhIL-8, rhIL-9, rmIL-9, rhIL-10, rhIL-10 sR, rmIL-10, rhIL-11, rhIL-12, rmIL-12, rhIL-13, rmIL-13, rhIL-15, rhIP-10, rhJAK-1, rmJAK-1, rmJE, rhLIF, rmLIF, rhM-CSF, rmM-CSF, rhMCP-1, rhMCP-1 R, rhMCP-2, rhMCP-3, rhMidkine, rhMIP-1 $\alpha$ , rmMIP-1 $\alpha$ , rhMIP-1 $\beta$ , rmMIP-1 $\beta$ , rmMIP-2, rhNT-3, rhNT-4, rhOSM, rhPD-ECGF, hPDGF, pPDGF. rhPDGF-AA, rhPDGF-AB, rhPDGF-BB, rhPDGF Rα, rhPIGF, rhPTN, rhRANTES, rhSCF, rmSCF, rhsqp130, rhSLPI, rhSTAT-1, rmSTAT-4, hTfR, rhTGF-α, rhTGF-β1, rhTGF-β2, rhTGF-β3, raTGF-β5, rhLAP (TGF- $\beta$ 1), rhLatent TGF- $\beta$ 1, rhTGF- $\beta$  sRII, rhTGF- $\beta$  sRIII, rhTNF- $\alpha$ , rmTNF- $\alpha$ , rhTNF- $\beta$ , rhsTNF RI, rhsTNF RII, rhTPO, rhVEGF

lpg 4/99