



**MONOCLONAL ANTI-HUMAN INTERLEUKIN-2  
SOLUBLE RECEPTOR  $\gamma$  (IL-2 sR $\gamma$ )  
CLONE 38024.11  
Purified Mouse Immunoglobulin**

Product Number **I5902**

**Product Description**

Monoclonal Anti-Human Interleukin-2 Soluble Receptor gamma (IL-2 sR $\gamma$ ) (mouse IgG1 isotype) is derived from the 38024.11 hybridoma produced by the fusion of mouse myeloma cells and splenocytes from a mouse immunized with a mouse cell line transfected with human IL-2 sR $\gamma$ . The antibody is purified from ascites fluid using protein G chromatography.

Monoclonal Anti-Human IL-2 sR $\gamma$  will be used to block the biological activity mediated by IL-2. By ELISA, the antibody does not cross-react with recombinant human IL-2 R $\alpha$ , IL-2 R $\beta$ , IL-1 RI, IL-1 RII, IL-4 R, IL-6 R, IL-7 R and IL-10 R.

Monoclonal Anti-Human IL-2 sR $\gamma$  may be used for neutralization of the biological activity mediated by IL-2 R $\gamma$  and the detection of IL-2 R $\gamma$  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 R $\beta$ , and IL-2 R $\gamma$  chains. The IL-2 R $\gamma$  chain is shared by IL-2, IL-4 and IL-7.<sup>1,2</sup> 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 gamma-chain include monocytes,<sup>3,4</sup> neutrophils,<sup>5</sup> thymocytes,<sup>6</sup> CD4<sup>+</sup> and CD8<sup>+</sup> T cells, NK cells and B cells.<sup>7</sup>

**Reagents**

## Product Information

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  $\mu$ m-filtered PBS to produce a 0.5 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  $-20^{\circ}\text{C}$ . Reconstituted product may be stored at  $2-8^{\circ}\text{C}$  for at least one month. For prolonged storage, freeze in working aliquots at  $-20^{\circ}\text{C}$ . Avoid repeated freezing and thawing.

**Procedure**

Anti-Human IL-2 sR $\gamma$  is tested for its ability to neutralize human cell surface IL-2 R $\gamma$  mediated IL-2 bioactivity in a  $^3\text{H}$ -thymidine incorporation assay using MO7e cells.<sup>8</sup> The ND<sub>50</sub> of the antibody is defined as the concentration of antibody resulting in a one-half maximal inhibition of the cell surface IL-2 R $\gamma$  mediated recombinant human IL-2 response on a responsive cell line.

**Product Profile**

For neutralization, a working concentration of 2 - 6  $\mu\text{g/ml}$  of Monoclonal Anti- IL-2 sR $\gamma$  will block 50% of the bioactivity due to 10 ng/ml recombinant human IL-2 in a  $^3\text{H}$ -thymidine incorporation assay using  $10^5/\text{ml}$  MO7e cells.

For Indirect Immunoblotting, a working concentration of 1-2  $\mu\text{g/ml}$  is determined using recombinant human IL-2 R $\gamma$  at 100 ng/lane under non-reducing conditions.

For Indirect ELISA, a working concentration of 0.5 - 1  $\mu\text{g/ml}$  is determined to detect recombinant IL-2 R $\gamma$  to a limit of 3 ng/well.

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

**References**

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