Human IL-18 Rβ/IL-1 R7 Antibody

Antigen Affinity-purified Polyclonal Goat IgG Catalog Number: AF118

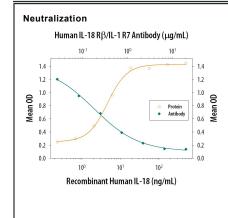
| DESCRIPTION | |
|--------------------|--|
| Species Reactivity | Human |
| Specificity | Detects human IL-18 Rβ in direct ELISAs and Western blots. In direct ELISAs, less than 1% cross-reactivity with recombinant mouse IL-18 Rβ (AcPL), recombinant human (rh) IL-18 R, rhIL-1 R1, rhIL-1 RII, and rhIL-1 Rrp2 is observed. |
| Source | Polyclonal Goat IgG |
| Purification | Antigen Affinity-purified |
| Immunogen | Mouse myeloma cell line NS0-derived recombinant human IL-18 Rβ/IL-1 R7 Phe20-Arg356 Accession # 095256 |
| Endotoxin Level | <0.10 EU per 1 µg of the antibody by the LAL method. |
| Formulation | Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose. See Certificate of Analysis for details. *Small pack size (-SP) is supplied as a 0.2 µm filtered solution in PBS. |

APPLICATIONS

Please Note: Optimal dilutions should be determined by each laboratory for each application. General Protocols are available in the Technical Information section on our website.

| | Recommended Concentration | Sample |
|----------------|---|--|
| Western Blot | 0.1 μg/mL | Recombinant Human IL-18 Rβ/IL-1 R7 Fc Chimera (Catalog # 118-AP) |
| Neutralization | | lity to neutralize IL-18/IL-1F4-induced IFN-γ secretion in the KG-1 human acute myelogenous Novick, D. <i>et al.</i> (1999) Immunity 10(1) :127. The Neutralization Dose (ND ₅₀) is typically |
| | 0.3–1.2 μ g/mL in the TNF– α . | e presence of 40 ng/mL Recombinant Human IL-18/IL-1F4 and 20 ng/mL Recombinant Human |

DATA



IFN-γ secretion Induced by IL-18/IL-1F4 and Neutralization by Human IL-18 Rβ/IL-1 R7 Antibody. In the presence of Recombinant Human TNF- α (20 ng/mL, Catalog # 210-TA), Recombinant Human IL-18/IL-1F4 stimulates IFN-y secretion in the KG-1 human acute myelogenous leukemia cell line in a dosedependent manner (orange line), as measured by the Human IFN-y Quantikine ELISA Kit (Catalog # DIF50). Under these conditions, IFN-γ secretion elicited by Recombinant Human IL-18/IL-1F4 (40 ng/mL) is neutralized (green line) by increasing concentrations of Human IL-18 Rβ/IL-1 R7 Antigen Affinity-purified Polyclonal Antibody (Catalog # AF118). The ND_{50} is typically 0.3-1.2 μ g/mL.

| Reconstitution | Reconstitute at 0.2 mg/mL in sterile PBS. | |
|---------------------|--|--|
| Shipping | The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below. *Small pack size (-SP) is shipped with polar packs. Upon receipt, store it immediately at -20 to -70 °C | |
| Stability & Storage | Use a manual defrost freezer and avoid repeated freeze-thaw cycles. 12 months from date of receipt, -20 to -70 °C as supplied. 1 month, 2 to 8 °C under sterile conditions after reconstitution. 6 months, -20 to -70 °C under sterile conditions after reconstitution. | |





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BACKGROUND

IL-18, originally described as an interferon- γ inducing factor (IGIF), is a member of the IL-1 family of cytokines that has multiple immunoregulatory functions. It has potent IFN- γ inducing activities and plays a key role in the activation of T helper type 1 (Th1) responses. The functional IL-18 receptor complex consists of two components, the IL-18 Rα (IL-1 R5) and IL-18 Rβ (also termed IL-1 R7 and AcPL) subunits. Both subunits are members of the IL-1 receptor superfamily. Although IL-18 Rα by itself binds IL-18 with low-affinity and IL-18 Rβ does not bind IL-18 *in vitro*, co-expression of IL-18 Rα and IL-18 Rβ is required for high-affinity binding and IL-18 responsiveness. Human IL-18 Rβ cDNA encodes a 599 amino acid (aa) residue precursor type I membrane protein with a 14 aa signal peptide, a 342 aa extracellular region containing three immunoglobulin-like domains, a single transmembrane domain and a 222 aa cytoplasmic domain. Human and mouse IL-18 Rβ share 65% aa sequence identity. The expression of IL-18 Rβ parallels that of IL-18 Rα and is detected in numerous tissues including lung, spleen, leukocytes and colon.

References:

- 1. Born, T.L. et al. (1998) J. Biol. Chem. 273:29445.
- 2. Okamura, H. et al. (2000) in Cytokine Reference, Vol. 2:1605, Academic Press.

SYSTEMS

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