RD SYSTEMS a biotechne brand

Recombinant Human IL-6R alpha

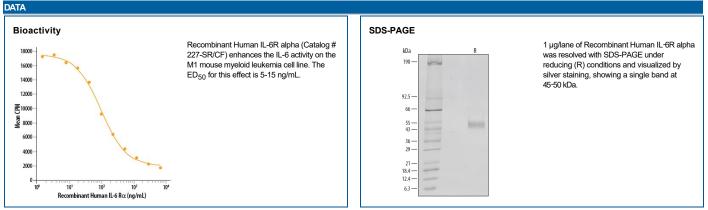
Catalog Number: 227-SR/CF

DESCRIPTION	
Source	<i>Spodoptera frugiperda, Sf</i> 21 (baculovirus)-derived human IL-6R alpha protein Leu20-Asp358
	Accession # P08887.1
N-terminal Sequence Analysis	Leu20
Predicted Molecular Mass	38 kDa

SPECIFICATIONS	
SDS-PAGE	45-50 kDa, reducing conditions
Activity	Measured by its ability to enhance the IL-6 activity on M1 mouse myeloid leukemia cells. Saito, T. <i>et al.</i> (1991) J. Immunol. 147 :168. The ED ₅₀ for this effect is 5-15 ng/mL.
Endotoxin Level	<1.0 EU per 1 µg of the protein by the LAL method.
Purity	>97%, by SDS-PAGE under reducing conditions and visualized by silver stain.
Formulation	Lyophilized from a 0.2 μm filtered solution in PBS. See Certificate of Analysis for details.

PREPARATION AND STORAGE	
Reconstitution	Reconstitute at 100 µg/mL in sterile PBS.
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.
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.

- 3 months, -20 to -70 °C under sterile conditions after reconstitution.



BACKGROUND

The multifunctional factor interleukin 6 (IL-6) exerts its activities through binding to a high-affinity receptor complex consisting of two membrane glycoproteins: an 80 kDa component receptor that binds IL-6 with low affinity (IL-6R α) and a signal-transducing component of 130 kDa (gp130) that does not bind IL-6 by itself, but is required for high-affinity binding of IL-6 by the complex. Both components of the receptor complex, IL-6R α and gp130 have been cloned, sequenced, and expressed (1-4).

A soluble form of the IL-6R α has been found in the urine of healthy adult humans (5). This soluble receptor apparently arises from proteolytic cleavage of membranebound IL-6R α . No naturally-occurring mRNA encoding a truncated form of the IL-6R α has been reported. Soluble forms of human and murine IL-6R α s have been constructed, however, by insertion of termination codons into the regions of the IL-6R α cDNAs encoding the external portions of the receptors and prior to the transmembrane domains. These soluble receptors have been expressed in COS-7 and CHO cells and have been shown to bind to IL-6 in solution and to augment the activity of IL-6 as a result of the binding of the IL-6/IL-6R α complex to membrane-bound gp130 (6, 7).

References:

- 1. Yamasaki *et al.* (1988) Science **241**:825.
- 2. Baumann et al. (1990) J. Biol. Chem. 265:19853.
- 3. Hibi et al. (1990) Cell 63:1149.
- 4. Schooltink et al. (1991) Eur. J. Biochem. 277:659.
- 5. Novick et al. (1989) J. Exp. Med. 170:1409.
- 6. Yasukawa et al. (1990) J. Biochem. 108:673.
- 7. Saito et al. (1991) J. Immunology 147:168.

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