

# Recombinant Human IL-26/AK155 Dimer

Catalog Number: 1870-IL

DESCRIPTION	
Source	E. coli-derived Lys22-Gln171, with an N-terminal Met Accession # Q9NPH9.1
N-terminal Sequence Analysis	Met
Structure / Form	Disulfide-linked homodimer and a small amount of monomer (no more than 10%)
Predicted Molecular Mass	17.7 kDa (monomer)
SPECIFICATIONS	
Activity	Measured by its ability to induce IL-10 secretion in COLO 205 human colorectal adenocarcinoma cells. Hor, S. <i>et al.</i> (2004) J. Biol. Chem. <b>279</b> :33343.  The ED <sub>50</sub> for this effect is typically 0.04-0.2 μg/mL.
Endotoxin Level	<1.0 EU per 1 µg of the protein by the LAL method.
Purity	>95%, by SDS-PAGE under reducing conditions and visualized by silver stain.
Formulation	Supplied as a 0.2 µm filtered solution in NaH <sub>2</sub> PO <sub>4</sub> , NaCl and Glycerol with BSA as a carrier protein. See Certificate of Analysis for details.

### PREPARATION AND STORAGE

Shipping The product is shipped with dry ice or equivalent. 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 opening.

### **BACKGROUND**

IL-26 was originally cloned from herpesvirus saimiri (HVS)-transformed T-cells and named AK155 (1). It is a member of the IL-10 family of class II cytokines that signal via heterodimeric receptor complexes composed of two type I transmembrane receptor subunits (2). The human IL-26 gene has been mapped to chromosome 12q15. It encodes a 171 amino acid polypeptide with a 21 amino acid signal peptide. In addition to HVS-transformed T cells, IL-26 is also expressed in other virus transformed T cell lines, fresh peripheral mononuclear cells, activated NK cells and T cells. A mouse homologue of human IL-26 has not been identified. IL-26 binds with high-affinity to the heterodimeric complex consisting of the ligand-binding IL-20 R $\alpha$  and non ligand-binding IL-10 R $\beta$  (3). Activation of the receptor complex results in rapid phosphorylation of STAT1 and STAT3. Although the IL-26 receptor complex is highly specific for IL-26 and is not activated by other class II cytokines, the individual subunits of the IL-26 receptor complex are components in receptor complexes for other class II cytokines (1). IL-20 R $\alpha$  can form dimers with IL-20 R $\beta$  to function as signaling receptors for IL-19, IL-20, and IL-24. IL-10 R $\beta$  can complex with IL-10 R $\alpha$ , IL-22 R $\alpha$ , and IL-28 R $\alpha$  to transduce signals for IL-10, IL-22, and the three novel IFNs (IL-28A, IL-28B and IL-29), respectively. The physiological functions of IL-26 remain to be determined.

IL-26 was reported to be a homodimer in solution (1). *E. coli*-derived IL-26 produced by R&D Systems contains disulfide-linked homodimers and monomers. Both the pure monomeric and dimeric IL-26 can bind directly to IL-20 Rα and induce STAT activation in COLO-205 cells (4). Besides IL-20 Rα, R&D Systems IL-26 preparations have also been shown to bind IL-20 Rβ. The significance of this receptor-ligand interaction has not been investigated.

## References:

- 1. Knappe. A. et al. (2000) J. Virology 74:3881.
- 2. Renauld, J.-C. (2003) Nature Reviews Immunology 3:667.
- 3. Sheikh, F. et al. (2004) J. Immunol. 172:2006.
- 4. Unpublished data (2003) R&D Systems, Inc.

