

# Recombinant Protein Technical Manual Recombinant Human ICOS Ligand/ICOSL Protein (His Tag)(Active)

### Product Data:

**Product SKU:** RPES1050 **Size:** 100μg

Species: Human Expression host: HEK293 Cells

**RPES1050** 

**Uniprot:** NP\_056074.1

### **Protein Information:**

Molecular Mass: 28 kDa

AP Molecular Mass: 50-60 kDa

Tag: C-His

Bio-activity: Measured by its binding ability in a functional ELISA. Immobilized human B7-H2 at

1 μg/ml (100 μl/well) can bind human ICOS with a linear range of 1.6-200 ng/ml.

**Purity:** > 98 % as determined by reducing SDS-PAGE.

**Endotoxin:** < 1.0 EU per μg as determined by the LAL method.

**Storage:** Lyophilized proteins are stable for up to 12 months when stored at -20 to -80°C.

Reconstituted protein solution can be stored at 4-8°C for 2-7 days. Aliquots of

reconstituted samples are stable at < -20°C for 3 months.

**Shipping:** This product is provided as lyophilized powder which is shipped with ice packs.

**Formulation:** Lyophilized from sterile PBS, pH 7.4

**Reconstitution:** Please refer to the printed manual for detailed information.

**Application:** Functional ELISA

**Synonyms:** ICOS Ligand; B7 Homolog 2; B7-H2; B7-Like Protein Gl50; B7-Related Protein 1;

B7RP; CD275; ICOSLG; B7H2; B7RP1; ICOSL; KIAA0653

# Immunogen Information:

Sequence: Met 1-Ser 258

## **Background:**

Inducible co-stimulator ligand (ICOSL), also known as B7-H2, is a member of the B7 family of co-stimulatory molecules related to B7 and B7-2. It is a transmembrane glycoprotein with extracellular IgV and IgC domains, and binds to ICOS on activated T cells, thus delivers a positive costimulatory signal for optimal T cell function. The structural features of ICOSL are crucial for its costimulatory function. Present study shows that ICOSL displays a marked oligomerization potential, resembling more like B7 than B7-2. B7-H2-dependent signaling may play an active role in a proliferative response rather than in cytokine and chemokine production. The CD28/B7 and ICOS/B7-H2 pathways are both critical for costimulating T cell immune responses. Deficiency in either pathway results in defective T cell activation, cytokine production and germinal center formation.