

## Recombinant Protein Technical Manual

# Recombinant Mouse PD/PDCD1 Protein (Fc Tag)(Active)

**RPES0833** 

**Product SKU: RPES0833** Size: 10µg

**Expression host:** Human Cells **Species**: Mouse

Uniprot: Q02242

**Molecular Mass:** 43.3 kDa

AP Molecular Mass: 58-85 kDa

Tag: C-Fc

**Bio-activity:** Immobilized Human PD-L1-His(Cat: PKSH033557) at 2μg/ml(100 μl/well) can bind

Mouse PD-Fc.

> 95% as determined by reducing SDS-PAGE. **Purity:** 

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

Lyophilized protein should be stored at < -20°C, though stable at room Storage:

> temperature for 3 weeks. Reconstituted protein solution can be stored at 4-7°C for 2-7 days. Aliquots of reconstituted samples are stable at < -20°C for 3 months.

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

Formulation: Lyophilized from a 0.2 µm filtered solution of 20mM Tris,150mM NaCl,pH8.0.

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

**Application: Functional ELISA** 

Programmed cell death protein 1;PD;CD279;Pdcd1;mPD Synonyms:

### Immunogen Information:

Sequence: Leu25-Gln167

### Background:

Programmed Death (PD), firstly cloned from mouse T cell hybridoma 2B4.11, is one member of CD28/CTLA-4 superfamily. PD belongs to type I transmembrane protein and acts as an important immunosuppressive molecule. This family also include members of CD28, CTLA-4 and ICOS. The mouse Programmed Death protein, encoded by PD gene, comprises four parts including a putative 20 aa signal peptide, a 149 aa extracellular region, a 21 aa transmembrane domain and a 98 aa cytoplasmic region. The cytoplamsic tail of PD contains two structural motifs, an immunoreceptor tyrosine-based inhibitory motif (ITIM) and an immunoreceptor tyrosine-based switch motif (ITSM) formed by two tyrosine residues which make the difference in PD signal mediating. Mouse PD is expressed in thymus and shares about 69% aa sequence identity with human PD. Recently, programmed death (PD) with its ligands, programmed death ligand B7H1 (PD-L1) and B7DC (PD-L2), was found to regulate T-cell activation and tolerance, upon ligand binding, inhibiting T-cell effector functions in an antigen-specific manner. PD gene knocked out mice would induce some autoimmune diseases, which suggests that PD acts as a co-inhibitory molecule actively participating in maintaining peripheral tolerance. Thus, PD may be a useful target for the immunologic therapy of carcinoma, infection, autoimmune diseases as well as organ transplantation.