Biotinylated Mouse PD-1 / PDCD1 Protein, His,Avitag™ (MALS verified)

Catalog # PD1-M82E3



Synonym

PDCD1, PD1, CD279, SLEB2

Source

Biotinylated Mouse PD-1 Protein, His, Avitag(PD1-M82E3) is expressed from human 293 cells (HEK293). It contains AA Leu 25 - Gln 167 (Accession # Q02242-1).

Predicted N-terminus: Leu 25

Molecular Characterization

PD-1(Leu 25 - Gln 167) Q02242 -1 Poly-his Avi

This protein carries a polyhistidine tag at the C-terminus, followed by an Avi tag (AvitagTM).

The protein has a calculated MW of 19.7 kDa. The protein migrates as 37-42 kDa when calibrated against <u>Star Ribbon Pre-stained Protein Marker</u> under reducing (R) condition (SDS-PAGE) due to glycosylation.

Labeling

Biotinylation of this product is performed using AvitagTM technology. Briefly, the single lysine residue in the Avitag is enzymatically labeled with biotin.

Protein Ratio

Passed as determined by the HABA assay / binding ELISA.

Purity

>95% as determined by SDS-PAGE.

>90% as determined by SEC-MALS.

Formulation

Lyophilized from 0.22 μm filtered solution in PBS, pH7.4 with trehalose as protectant.

Contact us for customized product form or formulation.

Reconstitution

Please see Certificate of Analysis for specific instructions.

For best performance, we strongly recommend you to follow the reconstitution protocol provided in the CoA.

Storage

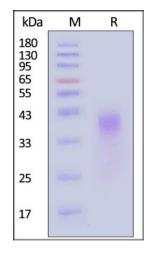
For long term storage, the product should be stored at lyophilized state at -20°C or lower.

Please avoid repeated freeze-thaw cycles.

This product is stable after storage at:

- -20°C to -70°C for 12 months in lyophilized state;
- -70°C for 3 months under sterile conditions after reconstitution.

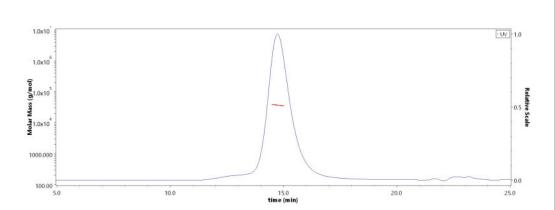
SDS-PAGE



Biotinylated Mouse PD-1 Protein, His, Avitag on SDS-PAGE under reducing (R) condition. The gel was stained with Coomassie Blue. The purity of the protein is greater than 95% (With <u>Star Ribbon Pre-stained Protein Marker</u>).

Bioactivity-ELISA

SEC-MALS



The purity of Biotinylated Mouse PD-1 Protein, His, Avitag (Cat. No. PD1-M82E3) is more than 90% and the molecular weight of this protein is around 30-40 kDa verified by SEC-MALS.

Report

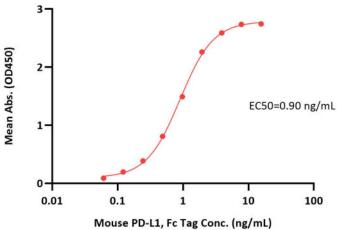


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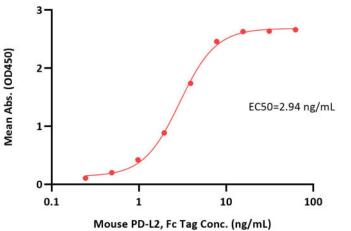


Biotinylated Mouse PD-1 / PDCD1 Protein, His, Avitag ELISA 0.1 μ g of Biotinylated Mouse PD-1 / PDCD1 Protein, His, Avitag per well



Immobilized Biotinylated Mouse PD-1 / PDCD1 Protein, His,Avitag (Cat. No. PD1-M82E3) at 1 μ g/mL (100 μ L/well) on streptavidin precoated (STN-N5116) can bind Mouse PD-L1, Fc Tag (Cat. No. PD1-M5251) with a linear range of 0.1-2 ng/mL (QC tested).

Biotinylated Mouse PD-1 / PDCD1 Protein, His, Avitag ELISA 0.1 μ g of Biotinylated Mouse PD-1 / PDCD1 Protein, His, Avitag per well



Immobilized Biotinylated Mouse PD-1 / PDCD1 Protein, His,Avitag (Cat. No. PD1-M82E3) at 1 μ g/mL (100 μ L/well)on streptavidin precoated (STN-N5116) can bind Mouse PD-L2, Fc Tag (Cat. No. PD2-M5254) with a linear range of 0.1-4 ng/mL (Routinely tested).

Background

Programmed cell death protein 1 (PD-1) is also known as CD279 and PDCD1, is a type I membrane protein and is a member of the extended CD28/CTLA-4 family of T cell regulators. PDCD1 is expressed on the surface of activated T cells, B cells, macrophages, myeloid cells and a subset of thymocytes. PD-1 has two ligands, PD-L1 and PD-L2, which are members of the B7 family. PD-L1 is expressed on almost all murine tumor cell lines, including PA1 myeloma, P815 mastocytoma, and B16 melanoma upon treatment with IFN-γ. PD-L2 expression is more restricted and is expressed mainly by DCs and a few tumor lines. PD1 inhibits the T-cell proliferation and production of related cytokines including IL-1, IL-4, IL-10 and IFN-γ by suppressing the activation and transduction of P13K/AKT pathway. In addition, coligation of PD1 inhibits BCR-mediating signal by dephosphorylating key signal transducer. In vitro, treatment of anti-CD3 stimulated T cells with PD-L1-Ig results in reduced T cell proliferation and IFN-γ secretion. Monoclonal antibodies targeting PD-1 that boost the immune system are being developed for the treatment of cancer.

