

Product Data Sheet

PD-1 Protein, Human (147a.a, HEK293, His)

Cat. No.:	HY-P70766
Synonyms:	Programmed cell death protein 1; hPD-1; PDCD1; CD279
Species:	Human
Source:	HEK293
Accession:	Q15116 (P21-Q167)
Gene ID:	5133
Molecular Weight:	Approximately 38.0 kDa

DDODEDTIEC	
PROPERTIES	
AA Sequence	PGWFLDSPDR PWNPPTFSPA LLVVTEGDNA TFTCSFSNTS ESFVLNWYRM SPSNQTDKLA AFPEDRSQPG QDCRFRVTQL PNGRDFHMSV VRARRNDSGT YLCGAISLAP KAQIKESLRA ELRVTERRAE VPTAHPSPSP RPAGQFQ
Biological Activity	10 $\mu g/mL$ (100 $\mu L/well) of immobilized Human PD-1-His can bind Anti-PD1 with an ED50 value of 3.1 \mu g/mL$
Appearance	Lyophilized powder.
Formulation	Lyophilized from a 0.2 µm filtered solution of PBS, pH 7.4.
Endotoxin Level	<1 EU/µg, determined by LAL method.
Reconsititution	It is not recommended to reconstitute to a concentration less than 100 μg/mL in ddH ₂ O. For long term storage it is recommended to add a carrier protein (0.1% BSA, 5% HSA, 10% FBS or 5% Trehalose).
Storage & Stability	Stored at -20°C for 2 years. After reconstitution, it is stable at 4°C for 1 week or -20°C for longer (with carrier protein). It is recommended to freeze aliquots at -20°C or -80°C for extended storage.
Shipping	Room temperature in continental US;may vary elsewhere.

DESCRIPTION

BackgroundPD-1 protein functions as an inhibitory receptor on antigen-activated T-cells, playing a crucial role in the induction and
maintenance of immune tolerance to self. Upon binding to its ligands CD274/PDCD1L1 and CD273/PDCD1LG2, PD-1 delivers
inhibitory signals and associates with CD3-TCR in the immunological synapse, directly impeding T-cell activation. This
inhibitory action is further executed through the recruitment of PTPN11/SHP-2, leading to the dephosphorylation of key TCR
proximal signaling molecules. Exploited by tumors to attenuate anti-tumor immunity, PD-1's interaction with
CD274/PDCD1L1 inhibits cytotoxic T lymphocytes (CTLs) effector function. Blockage of the PD-1-mediated pathway has

shown promise in reversing the exhausted T-cell phenotype and normalizing the anti-tumor response, providing a rationale for cancer immunotherapy.

Caution: Product has not been fully validated for medical applications. For research use only.

 Tel: 609-228-6898
 Fax: 609-228-5909
 E-mail: tech@MedChemExpress.com

 Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA