

## Biotinylated Recombinant Human PD-1 Fc Chimera Avi-tag

Catalog Number: AVI1086

DESCRIPTION					
Source	Chinese Hamster Ovary cell line, CHO-derived human PD-1 protein				
	Human PD-1 (Leu25-Thr168) Accession # Q15116.3	IEGRMD	Human IgG <sub>1</sub> (Pro100-Lys330)	HP, 3x (GGGS)	Avi-tag
N-terminal Sequence Analysis	Leu25				
Structure / Form	Disulfide-linked homodimer, biotinylated via Avi-tag				
Predicted Molecular Mass	45 kDa				

SPECIFICATIONS			
SDS-PAGE	62-88 kDa, under reducing conditions		
Activity	Measured by its binding ability in a functional ELISA. When Recombinant Human B7-H1/PD-L1 Fc Chimera (Catalog # 156-B7) is immobilized at 1 μg/mL (100 μL/well), the concentration of Biotinylated Recombinant Human PD-1 Fc Chimera Avi-tag (Catalog # AVI1086) that produces 50% of the optimal binding response is 10-90 ng/mL.		
Endotoxin Level	<0.10 EU per 1 $\mu$ g of the protein by the LAL method.		
Purity	>95%, by SDS-PAGE visualized with Silver Staining and quantitative densitometry by Coomassie® Blue Staining.		
Formulation	Lyophilized from a 0.2 µm filtered solution in PBS with Trehalose. See Certificate of Analysis for details.		

PREPARATION AND STORAGE				
Reconstitution	Reconstitute at 250 $\mu$ g/mL in PBS.			
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.			
Stability & Storage	<ul> <li>Use a manual defrost freezer and avoid repeated freeze-thaw cycles.</li> <li>12 months from date of receipt, -20 to -70 °C as supplied.</li> <li>1 month, 2 to 8 °C under sterile conditions after reconstitution.</li> <li>3 months, -20 to -70 °C under sterile conditions after reconstitution.</li> </ul>			



2 µg/lane of Biotinylated Recombinant Human PD-1 Fc Chimera Avi-tag (Catalog # AVI1086) was resolved with SDS-PAGE under reducing (R) and non-reducing (NR) conditions and visualized by Coomassie® Blue staining, showing bands at 62-88 kDa and 120-180 kDa, respectively.

Rev. 1/22/2020 Page 1 of 2



**Global** bio-techne.com info@bio-techne.com techsupport@bio-techne.com TEL +1 612 379 2956 USA TEL 800 343 7475 **Canada** TEL 855 668 8722 **China** TEL +86 (21) 52380373 **Europe | Middle East | Africa** TEL +44 (0)1235 529449



## Biotinylated Recombinant Human PD-1 Fc Chimera Avi-tag

Catalog Number: AVI1086

## BACKGROUND

Programmed Death-1 receptor (PD-1), also known as CD279, is type I transmembrane protein belonging to the CD28 family of immune regulatory receptors (1). Other members of this family include CD28, CTLA-4, ICOS, and BTLA (2-5). Mature mouse PD-1 consists of a 149 amino acid (aa) extracellular region (ECD) with one immunoglobulin-like V-type domain, a 21 aa transmembrane domain, and a 98 aa cytoplasmic region. The mouse PD-1 ECD shares 65% aa sequence identity with the human PD-1 ECD. The cytoplasmic tail contains two tyrosine residues that form the immunoreceptor tyrosine-based inhibitory motif (ITIM) and immunoreceptor tyrosine-based switch motif (ITSM) that are important for mediating PD-1 signaling. PD-1 acts as a monomeric receptor and interacts in a 1:1 stoichiometric ratio with its ligands PD-L1 (B7-H1) and PD-L2 (B7-DC) (6, 7). PD-1 is expressed on activated T cells, B cells, monocytes, and dendritic cells while PD-L1 expression is constitutive on the same cells and also on nonhematopoietic cells such as lung endothelial cells and hepatocytes (8, 9). Ligation of PD-L1 with PD-1 induces co-inhibitory signals on T cells promoting their apoptosis, anergy, and functional exhaustion (10). Thus, the PD-1:PD-L1 interaction is a key regulator of the threshold of immune response and peripheral immune tolerance (11). Finally, blockade of the PD-1: PD-L1 interaction by either antibodies or genetic manipulation accelerates tumor eradication and shows potential for improving cancer immunotherapy (12, 13).

## References:

- 1. Ishida, Y. et al. (1992) EMBO J. 11:3887.
- 2. Sharpe, A.H. and G. J. Freeman (2002) Nat. Rev. Immunol. 2:116.
- 3. Coyle, A. and J. Gutierrez-Ramos (2001) Nat. Immunol. 2:203.
- 4. Nishimura, H. and T. Honjo (2001) Trends Immunol. 22:265.
- 5. Watanabe, N et al. (2003) Nat. Immunol. 4:670.
- 6. Zhang, X. et al. (2004) Immunity 20:337.
- 7. Lázár-Molnár, E. et al. (2008) Proc. Natl. Acad. Sci. USA 105:10483.
- 8. Nishimura, H et al. (1996) Int. Immunol. 8:773.
- 9. Keir, M.E. et al. (2008) Annu. Rev. Immunol. 26:677.
- 10. Butte, M.J. et al. (2007) Immunity 27:111.
- 11. Okazaki, T. et al. (2013) Nat. Immunol. 14:1212.
- 12. Iwai, Y. et al. (2002) Proc. Natl. Acad. Sci. USA 99: 12293.
- 13. Nogrady, B. (2014) Nature 513:S10.

Rev. 1/22/2020 Page 2 of 2



Global bio-techne.com info@bio-techne.com techsupport@bio-techne.com TEL +1 612 379 2956 USA TEL 800 343 7475 Canada TEL 855 668 8722 China TEL +86 (21) 52380373 Europe | Middle East | Africa TEL +44 (0)1235 529449