

**DESCRIPTION**

<b>Source</b>	Human embryonic kidney cell, HEK293-derived		
	Cynomolgus Monkey PD-L1 (Phe19-Thr239) Accession # XP_005581836	IEGRMD	Human IgG <sub>1</sub> (Pro100-Lys330)
	N-terminus		C-terminus
<b>N-terminal Sequence Analysis</b>	Phe19		
<b>Structure / Form</b>	Disulfide-linked homodimer		
<b>Predicted Molecular Mass</b>	52 kDa		

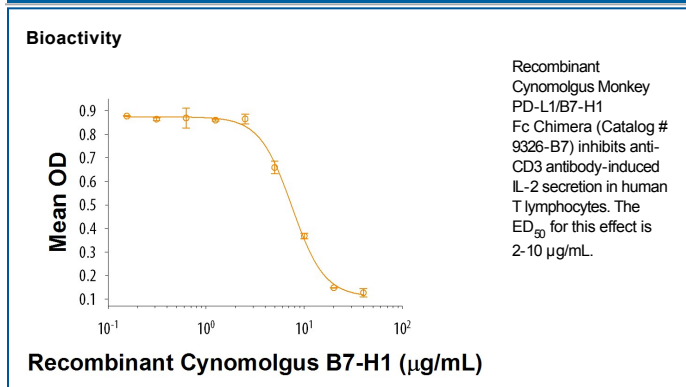
**SPECIFICATIONS**

<b>SDS-PAGE</b>	58-75 kDa, reducing conditions
<b>Activity</b>	Measured by its ability to inhibit anti-CD3 antibody induced IL-2 secretion in human T lymphocytes. The ED <sub>50</sub> for this effect is 2-10 µg/mL.
<b>Endotoxin Level</b>	<0.10 EU per 1 µg of the protein by the LAL method.
<b>Purity</b>	>95%, by SDS-PAGE visualized with Silver Staining and quantitative densitometry by Coomassie® Blue Staining.
<b>Formulation</b>	Lyophilized from a 0.2 µm filtered solution in PBS. See Certificate of Analysis for details.

**PREPARATION AND STORAGE**

<b>Reconstitution</b>	Reconstitute at 200 µg/mL in PBS.
<b>Shipping</b>	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.
<b>Stability &amp; Storage</b>	<p><b>Use a manual defrost freezer and avoid repeated freeze-thaw cycles.</b></p> <ul style="list-style-type: none"> <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>

**DATA**



**BACKGROUND**

B7-H1, also known as PD-L1 and CD274, is an approximately 65 kDa transmembrane glycoprotein in the B7 family of immune regulatory molecules (1, 2). Mature cynomolgus B7-H1 consists of a 220 amino acid (aa) extracellular domain (ECD) with two immunoglobulin-like domains, a 21 aa transmembrane segment, and a 30 aa cytoplasmic domain. Within the ECD, cynomolgus B7-H1 shares 92%, 72%, and 72% aa sequence identity with human, mouse, and rat B7-H1, respectively. In addition, cynomolgus B7-H1 shares 98%, 94%, 94%, 88%, 78%, 98%, 95%, 94%, and 94% aa sequence identity with rhesus macaque, chimpanzee, sumatran orangutan, white-tufted-ear marmoset, Garnett's greater bushbaby, olive baboon, green monkey, western lowland gorilla, and northern white-cheeked gibbon B7-H1, respectively. B7-H1 is expressed on inflammatory-activated immune cells including macrophages, T cells, and B cells (3-6), keratinocytes (7, 8), endothelial and intestinal epithelial cells (8, 10), as well as a variety of carcinomas and melanoma (10, 11). B7-H1 is a B7 ligand and binds to B7-1/CD80 and PD-1 receptors on T cells (6, 7, 11-14). It suppresses T cell activation and proliferation (4, 7, 13, 15) and induces the apoptosis of activated T cells (10). It plays a role in the development of immune tolerance by promoting T cell anergy (6, 13) and enhancing regulatory T cell development (15). B7-H1 favors the development of anti-inflammatory IL-10 and IL-22 producing dendritic cells (4, 9) and inhibits the development of Th17 cells (15). In cancer, B7-H1 provides resistance to T cell mediated lysis, enhances EMT, and enhances the tumorigenic function of Th22 cells (5, 8, 11, 14). B7-H1/PD-1 coinhibitory pathway was exploited therapeutically resulting in remarkable outcomes with 20-90% response in various types of cancer (16).

**References:**

1. Ceeraz, S. *et al.* (2013) Trends Immunol. **34**:556.
2. Dong, H. *et al.* (1999) Nat. Med. **5**:1365.
3. Tamura, H. *et al.* (2001) Blood **97**:1809.
4. Chen, L. *et al.* (2007) J. Immunol. **178**:6634.
5. Kuang, D.-M. *et al.* (2014) J. Clin. Invest. **124**:4657.
6. Tsushima, F. *et al.* (2007) Blood **110**:180.
7. Mazanet, M.M. and C.C.W. Hughes (2002) J. Immunol. **169**:3581.
8. Cao, Y. *et al.* (2010) Cancer Res. **71**:1235.
9. Scandiuizzi, L. *et al.* (2014) Cell Rep. **6**:625.
10. Dong, H. *et al.* (2002) Nat. Med. **8**:793.
11. Azuma, T. *et al.* (2008) Blood **111**:3635.
12. Butte, M.J. *et al.* (2008) Mol. Immunol. **45**:3567.
13. Park, J.-J. *et al.* (2010) Blood **116**:1291.
14. Ritprajak, P. *et al.* (2010) J. Immunol. **184**:4918.
15. Herold, M. *et al.* (2015) J. Immunol. **195**:3584.
16. Bardhan K. *et al.* (2016); Front Immunol. **7**:550.