

**DESCRIPTION**

<b>Source</b>	Chinese Hamster Ovary cell line, CHO-derived human CD47 protein			
	Human CD47 (Gln19-Pro139) Accession # Q08722	IEGRMD	Human IgG <sub>1</sub> (Pro100-Lys330)	Avi-tag
	N-terminus		C-terminus	
<b>N-terminal Sequence Analysis</b>	No results obtained: Gln19 predicted			
<b>Structure / Form</b>	Disulfide-linked homodimer, Biotinylated via Avi-tag			
<b>Predicted Molecular Mass</b>	42 kDa			

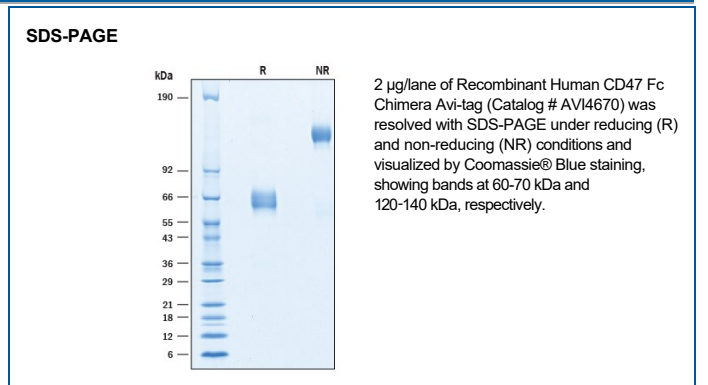
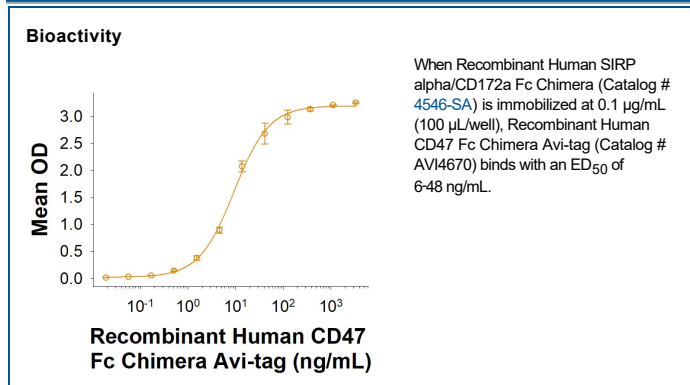
**SPECIFICATIONS**

<b>SDS-PAGE</b>	60-70 kDa, under reducing conditions
<b>Activity</b>	Measured by its binding ability in a functional ELISA. When Recombinant Human SIRP alpha/CD172a Fc Chimera (Catalog # 4546-SA) is immobilized at 0.1 µg/mL (100 µL/well), the concentration of Recombinant Human CD47 Fc Chimera Avi-tag (Catalog # AVI4670) that produces 50% of the optimal binding response is 6-48 ng/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 with Trehalose. See Certificate of Analysis for details.

**PREPARATION AND STORAGE**

<b>Reconstitution</b>	Reconstitute at 100 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>	Use a manual defrost freezer and avoid repeated freeze-thaw cycles. <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**

CD47, also known as Integrin-Associated Protein (IAP) and OA3, is a 40-60 kDa variably glycosylated atypical member of the immunoglobulin superfamily (1, 2). Human CD47 is an integral membrane protein that consists of a 123 amino acid (aa) extracellular domain (ECD) with a single Ig-like domain, five membrane-spanning regions with short intervening loops, and a 34 aa C-terminal cytoplasmic tail (3). Alternate splicing of human CD47 generates additional isoforms with deletions in the cytoplasmic tail (3). Within the N-terminal ECD, human CD47 shares 63% aa sequence identity with mouse and rat CD47. A portion of the N-terminal ECD can be shed from smooth muscle cells by MMP-2-mediated proteolysis (4). The ubiquitously expressed CD47 binds to SIRP family members on macrophages, neutrophils, and T cells (5, 6). These interactions prevent macrophage-mediated clearance of healthy CD47-expressing cells and promote immune cell transmigration across the vascular endothelium (5-8). The CD47-SIRP $\alpha$  interaction is species specific, and this lack of cross-species interaction has been implicated in xenotransplantation rejection (16). CD47 associates *in cis* with Fas on T cells and enhances Fas-mediated apoptosis; its ligation promotes T cell anergy and dampens Th1 immune responses (9-11). CD47 also associates *in cis* with Integrins  $\alpha 4\beta 1$ ,  $\alpha V\beta 3$ ,  $\alpha 2b\beta 3$ , and  $\alpha 2\beta 1$  which can positively or negatively modulate Integrin-mediated function (2, 12). In the vasculature, CD47 binding by Thrombospondin-1 inhibits the angiogenic and vasorelaxant effects of nitric oxide (2, 13, 14). On dendritic cells and myeloma cells, CD47 ligation by TSP-1 induces giant cell formation and osteoclast differentiation (15).

**References:**

1. Sarfati, M. *et al.* (2009) *Curr. Drug Targ.* **9**:842.
2. Isenberg, J.S. *et al.* (2008) *Arterioscler. Thromb. Vasc. Biol.* **28**:615.
3. Campbell, I.G. *et al.* (1992) *Cancer Res.* **52**:5416.
4. Maile, L.A. *et al.* (2008) *Mol. Endocrinol.* **22**:1226.
5. Oldenborg, P.-A. *et al.* (2000) *Science* **288**:2051.
6. Liu, Y. *et al.* (2002) *J. Biol. Chem.* **277**:10028.
7. Stefanidakis, M. *et al.* (2008) *Blood* **112**:1280.
8. de Vries, H.E. *et al.* (2002) *J. Immunol.* **168**:5832.
9. Manna, P.P. *et al.* (2005) *J. Biol. Chem.* **280**:29637.
10. Avice, M.-N. *et al.* (2001) *J. Immunol.* **167**:2459.
11. Bouguermouh, S. *et al.* (2008) *J. Immunol.* **180**:8073.
12. Barazi, H.O. *et al.* (2002) *J. Biol. Chem.* **277**:42859.
13. Isenberg, J.S. *et al.* (2009) *Nitric Oxide* **21**:52.
14. Isenberg, J.S. *et al.* (2009) *Matrix Biol.* **28**:110.
15. Kukreja, A. *et al.* (2009) *Blood* **114**:3413.
16. Wang H. *et al.* (2007) *Blood* **109**:836.