

Biotinylated Recombinant Human CD8α Fc Chimera Avi-tag

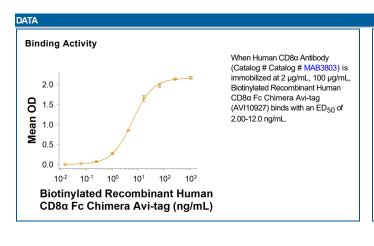
Catalog Number: AVI10927

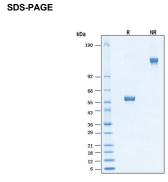
DESCRIPTION						
Source	Chinese Hamster Ovary cell line, Ch	ese Hamster Ovary cell line, CHO-derived human CD8 protein				
	Human CD8α (Ser22-Asp182) Accession # P01732.1	IEGRMD	Human IgG1 (Pro100-Lys330)	Avi-tag		
	N-terminus			C-terminus		

N-terminal Sequence Analysis	Ser22
Structure / Form	Biotinylated via Avi-tag.
Predicted Molecular	46 kDa

SPECIFICATIONS			
SDS-PAGE	55-61 kDa, under reducing conditions.		
Activity	Measured by its binding ability in a functional ELISA. When Human CD8α Antibody (Catalog # MAB3803) is immobilized at 2 μg/mL, 100 μg/mL, Biotinylated Recombinant Human CD8α Fc Chimera Avi-tag binds with an ED ₅₀ of 2.00-12.0 ng/mL.		
Endotoxin Level	<0.10 EU per 1 µg 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 500 μg/mL in PBS.		
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.		
Stability & Storage	Use a manual defrost freezer and avoid repeated freeze-thaw cycles. 12 months from date of receipt, -20 to -70 °C as supplied. 1 month, 2 to 8 °C under sterile conditions after reconstitution. 3 months, -20 to -70 °C under sterile conditions after reconstitution.		





Biotinylated Recombinant Human CD8α Fc Chimera Avi-tag Protein SDS-PAGE. 2 μg/lane of Biotinylated Recombinant Human CD8α Fc Chimera Avi-tag Protein (Catalog # AVI10927) was resolved with SDS-PAGE under reducing (R) and non-reducing (NR) conditions and visualized by Coomassie® Blue staining, showing bands at 55-61 kDa and 110-120 kDa, respectively.

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BACKGROUND

CD8, also known as Ly2 or Leu2, is a heterodimeric glycoprotein (alpha and beta subunits) that functions in conjunction with the T cell receptor in the recognition of MHC class I/peptide complexes (1, 2). CD8 alpha is expressed on double positive (CD4+ CD8+) thymocytes and mature CD8+ cytolytic T cells (CTL) (3-5), intraepithelial lymphocytes (IEL) (6), some γδ T cells (7), subsets of thymic and splenic dendritic cells (DC) (8), and peritoneal mast cells (9). It can form disulfide linked homodimers or heterodimers with CD8β (10). Thymic CD8+ DC express primarily αβ heterodimers, while splenic CD8+ DC primarily express αα homodimers (8). CD8α+ DC can present viral antigenic peptides in complex with MHC I and prime CTL responses (11). The approximately 35 kDa mature mouse CD8α consists of a 169 amino acid (aa) extracellular domain (ECD) with one Iglike domain, a 21 aa transmembrane segment, and a 30 aa cytoplasmic domain (12). Within the ECD, mouse CD8α shares 49% and 64% aa sequence identity with human and rat CD8α, respectively. Our Avi-tag Biotinylated human CD8α features biotinylation at a single site contained within the Avi-tag, a unique 15 amino acid peptide. Protein orientation will be uniform when bound to streptavidin-coated surface due to the precise control of bionylation and the rest of the protein is unchanged so there is no interference in the protein bioactivity.

References

- 1. Laugel, B. et al. (2011) J. Leukoc. Biol. 90:1089.
- 2. Cole, D.K. et al. (2012) Immunology 137:139.
- 3. Germain, R.N. (2002) Nat. Rev. Immunol. 2:309.
- 4. Ledbetter, J.A. et al. (1980) J. Exp. Med. 152:280.
- 5. Nakayama, K. et al. (1994) Science 263:1131.
- 6. Wang, J. and J.R. Klein (1994) Science 265:1860
- 7. MacDonald, H.R. et al. (1990) Eur. J. Immunol. 20:927.
- 8. Vremec, D. et al. (1992) J. Exp. Med. 176:47.
- 9. Lin, T.J. et al. (1998) J. Immunol. 161:6265.
- 10. Snow, P.M. and C. Terhorst (1983) J. Biol. Chem. 258:14675.
- 11. Belz, G.T. et al. (2004) J. Immunol. 172:1996.
- 12. Nakauchi, H. et al. (1985) Proc. Natl. Acad. Sci. USA 82:5126.