Biotinylated Recombinant Human Siglec-2/CD22 Fc Chimera Avi-tag

Catalog Number: AVI1968

DESCRIPTION Source Chinese Hamster Ovary cell line, CHO-derived human Siglec-2/CD22 protein Human Siglec-2/CD22 Human IgG₁ (Asp20-Arg687) **IEGRMD** Avi-tag (Pro100-Lys330) Accession # CAA42006.1

N-terminus C-terminus

N-terminal Sequence Asp20 **Analysis**

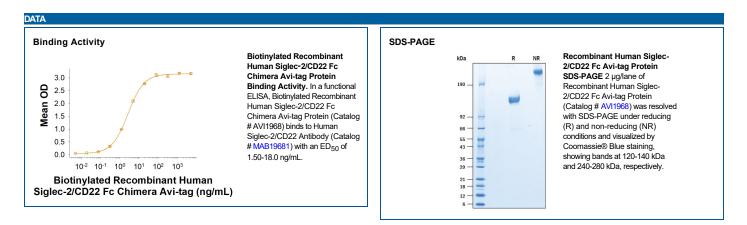
Structure / Form Disulfide-linked homodimer, biotinylated via Avi-tag

Predicted Molecular 104 kDa

Mass

SPECIFICATIONS	
SDS-PAGE	120-140 kDa, under reducing conditions.
Activity	Measured by the ability of the immobilized protein to support the adhesion of human red blood cells. Kelm, S. <i>et al.</i> (1994) Current Biology 4 :965. The ED ₅₀ for this effect is 0.1-0.9 μg/mL.
	Measured by its binding ability in a functional ELISA. Biotinylated Recombinant Human Siglec-2/CD22 Fc Chimera Avi-tag (Catalog # AVI1968) binds to Human Siglec-2/CD22 Antibody (Catalog # MAB19681) with an ED ₅₀ of 1.50-18.0 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.



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BACKGROUND

Sialic acid-binding immunoglobulin-like lectin 2 (Siglec-2), also known as B-cell receptor CD22 or B-lymphocyte cell adhesion molecule (BL-CAM), is a I-type (Ig-type) lectin belonging to the sialoadhesin subclass of the immunoglobulin superfamily (1). Fourteen human and nine mouse Siglecs have been characterized and are divided into 2 families: CD33 related and evolutionarily conserved (2, 3). The extracellular domain (ECD) of Siglecs are characterized by an N-terminal Ig-like V-type domain, which mediates sialic acid binding, followed by varying numbers of Ig-like C2-type domains (1-3). The predominant form of human Siglec-2 contains a N-terminal Iglike V-type domain, six Ig-like C2-type domains, a transmembrane region and a cytoplasmic tail with six tyrosine residues and four immunoreceptor tyrosine-based inhibition motifs (ITIMs) (1-3). A variant form of Siglec-2 missing two Ig-like C2-type domains along with a truncated cytoplasmic tail has also been identified (4). The mature ECD of human Siglec-2 shares 59% and 58% amino acid sequence identity with mouse and rat Siglec-2, respectively. Siglec-2 is an adhesion molecule that preferentially binds alpha 2,6- linked sialic acid on the same (cis) or adjacent (trans) cells (5). Besides its role as an adhesion molecule, Siglec-2 is a coreceptor that physically interacts with B-cell receptor (BCR), negatively regulating BCR signals by recruiting tyrosine phosphatase SHP-1 to its ITIMs. Phosphorylated Siglec-2 can also interact with other intracellular effector proteins such as Syk, PLC gamma, Pl3 kinase and Grb-2, suggesting it may play a role in positive signaling (2). Another function of Siglec-2 is that it mediates the anti-phagocytic effect of α2,6-linked sialic acid, and inhibition of Siglec-2 promotes the clearance of myelin debris, amyloidβ oligomers and α-synuclein fibrils in vivo (6). Siglec-2 also plays a role in autoimmunity and has great potential for Siglec-2-based immunotherapeutics for the treatment of autoimmune diseases such as systemic lupus erythematosus (SLE) (7). Our Avi-tag Biotinylated Recombinant Human Siglec-2 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 biotinylation and the rest of the protein is unchanged so there is no interference in the protein's bioactivity.

- 1. Sato, S. et al. (1996) Immunity. 5:551.
- 2. Crocker, P.R. and A. Varki (2001) Trends Immunol. 22:337.
- 3. Macauley, M.S. et al. (2014) Nature Rev Imm. 14:653.
- 4. Stamenkovic, I. and B. Seed (1990) Nature 345:74.
- Collins, B.E. et al. (2004) Proc. Natl. Acad. Sci. 101:6104.
- 6. Pluvinage, JV. et al. (2019) Nature. 568:7751.
- 7. Clark, E.A. et al. (2018) Front Immunol. 9:2235.