

**DESCRIPTION**

**Source** Chinese Hamster Ovary cell line, CHO-derived cynomolgus monkey SIRP beta 1/CD172b protein  
Glu30-Pro369, with a C-terminal 6-His tag  
Accession # XP005568593.1

**N-terminal Sequence Analysis** Glu30

**Predicted Molecular Mass** 38 kDa

**SPECIFICATIONS**

**SDS-PAGE** 53-68 kDa, under reducing conditions

**Activity** Measured by its binding ability in a functional ELISA.  
When Recombinant Cynomolgus Monkey SIRPβ1/CD172b is immobilized at 1 µg/mL, 100 µL/well, the concentration of **Recombinant Human SP-D** (Catalog # 1920-SP) that produces 50% of the optimal binding response is 0.15-0.9 µg/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.

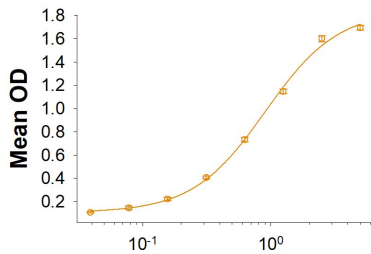
**Shipping** The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below.

**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.

**DATA**

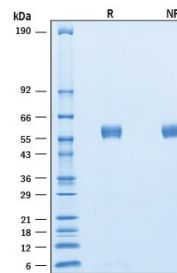
**Binding Activity**



When Recombinant Cynomolgus Monkey SIRPβ1/CD172b (Catalog # 10192-SB) is immobilized at 1 µg/mL, 100 µL/well, **Recombinant Human SP-D** (Catalog # 1920-SP) binds with an ED<sub>50</sub> of 0.15-0.9 µg/mL.

**Recombinant Human SP-D (µg/mL)**

**SDS-PAGE**



2 µg/lane of Recombinant Cynomolgus Monkey SIRP beta 1/CD172b His-tag was resolved with SDS-PAGE under reducing (R) and non-reducing (NR) conditions and visualized by Coomassie® blue staining, showing bands at 53-68 kDa.

## BACKGROUND

Signal-regulatory protein beta 1 (SIRP beta 1) is a disulfide-linked type I membrane glycoprotein that belongs to the SIRP/SHPS (CD172) family of the immunoglobulin (Ig) superfamily (1). Mature cynomolgus SIRP beta 1 consists of a 342 amino acid (aa) extracellular domain (ECD), a 26 aa transmembrane segment, and a short 4 aa cytoplasmic domain. Within the ECD, cynomolgus SIRP beta 1 shares 87% aa sequence identity with human SIRP beta 1. The SIRP family are paired receptors that have similar extracellular domains but differing C-terminal domains and functions (1). Members of this family are characterized by an extracellular region containing a V-set Ig domain containing a J-like sequence and two C1-set Ig domains (2). Positively charged residues within the transmembrane domain mediate interactions with DAP12 proteins which contain immunoreceptor tyrosine-based activation motifs (ITAMs) (3). Proteins in the SIRP family are typically expressed in cells of monocyte, macrophage or dendritic lineages (4). SIRP beta 1 has a relatively short cytoplasmic region and lacks the signaling motifs for association with phosphatases. However, formation of the SIRP beta 1/DAP12 complex in myeloid cells induce tyrosine phosphorylation, mitogen-activated protein kinase activation, and cellular activation (5, 6). Engagement of SIRP beta 1 by specific monoclonal antibodies promoted Fc $\gamma$  receptor-dependent or -independent phagocytosis in mouse peritoneal macrophages (7). Surfactant protein D (SP-D) has been shown to bind SIRP alpha and SIRP beta 1 in a calcium-dependent and sugar-specific manner on a distinct binding site from CD47 (8). Although the SIRP beta 1 extracellular regions share a high degree of homology with the SIRP alpha, SIRP beta 1 has been shown not to bind CD47 (9).

## References:

1. vanBeek, E.M. *et al.* (2005) *J. Immunol.* **175**:7781.
2. van den Berg, T. *et al.* (2008) *Trends in Immunology* **29**:203.
3. Liu, Y. *et al.* (2005) *Journal of Biological Chemistry* **280**:36132.
4. Matozaki, T. *et al.* (2009) *Trends in Cell Biology* **19**:72.
5. Dietrich, J. *et al.* (2000) *J Immunol.* **164**:9.
6. Brook, G. *et al.* (2004) *J Immunol.* **173**:2562.
7. Hayashi, A. *et al.* (2004) *J Biol Chem.* **279**:29450.
8. Fournier, B. *et al.* (2012) *J. Biol. Chem.* **287**:19386.
9. Seiffert, M. *et al.* (2001) *Blood* **97**:2741.