

DESCRIPTION

Source Mouse myeloma cell line, NS0-derived rat SIRP alpha/CD172a protein
Lys32-Asn373, with a C-terminal 6-His tag
Accession # P97710

N-terminal Sequence Analysis Lys32

Predicted Molecular Mass 38.3 kDa

SPECIFICATIONS

SDS-PAGE 75-95 kDa, reducing conditions

Activity Measured by the ability of the immobilized protein to support the adhesion of mouse red blood cells.
The ED₅₀ for this effect is 1.2-4.8 μ g/mL.

Optimal dilutions should be determined by each laboratory for each application.

Endotoxin Level <0.10 EU per 1 μ g of the protein by the LAL method.

Purity >95%, by SDS-PAGE under reducing conditions and visualized by silver stain.

Formulation Lyophilized from a 0.2 μ m filtered solution in PBS. See Certificate of Analysis for details.

PREPARATION AND STORAGE

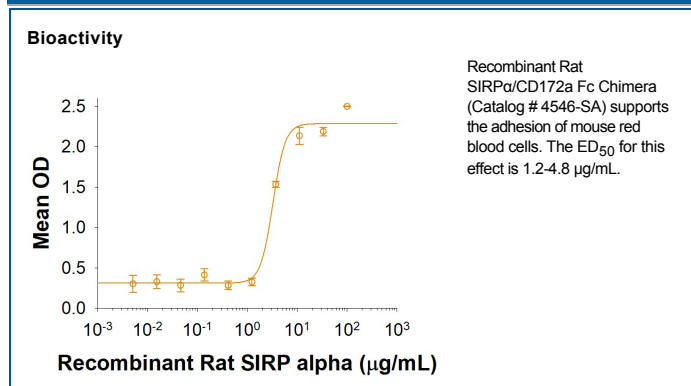
Reconstitution Reconstitute at 500 μ g/mL in PBS.

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



BACKGROUND

Signal regulatory protein alpha (SIRP α , designated CD172a), also called SHPS-1 (SHP substrate 1) and previously, MyD-1 (Myeloid/Dendritic-1), is a homodimeric, 100 - 120 kDa type I transmembrane glycoprotein that belongs to the SIRP/SHPS (CD172) family of the immunoglobulin superfamily (1 - 6). SIRPs are paired receptors, with similar extracellular domains but differing C-termini and functions (1, 2). The 509 amino acid (aa) rat SIRP α contains a 342 aa extracellular domain (ECD) with one V-type and two C1 type Ig domains and many potential N-glycosylation sites. It has a 113 aa cytoplasmic sequence with ITIM motifs that recruit tyrosine phosphatases SHP-1 and SHP-2 when phosphorylated (6). Rat SIRP α ECD shares 60% and 75% aa sequence identity with human and mouse SIRP α , respectively. Mouse and human SIRP α have at least 30 described polymorphisms, including the human SIRP α prominent variant BIT (Brain Ig like molecule with Tyrosine-based activation motifs, also called SIRP α_2 or PTPNS) (2). Less is known about rat SIRP α polymorphisms and family members. SIRP α is expressed mainly on myeloid cells, including macrophages, neutrophils, dendritic and Langerhans cells (3 - 7). It is also found on neurons, smooth muscle and endothelial cells (8 - 10). SIRP α shows adhesion to the ubiquitous CD47/IAP (integrin associated protein) (1, 2). Interaction between SIRP α and CD47 on red blood cells occurs in a species specific manner (17). Mouse and human SIRP α are allelic in nature, and variations in the V-type Ig-like domain likely impacts its binding to CD47 (11). SIRP α engagement generally produces a negative regulatory signal (4). Low SIRP α recognition of CD47, which occurs on aged erythrocytes or platelets or xenogenic cells, promotes clearance of CD47^{low} cells from circulation (12-14). SIRP α recognition of surfactants SP-A and SP-D in the lung can inhibit alveolar macrophage cytokine production (15). The CD47 integrin-SIRP α interaction is reported to promote macrophage fusion during osteoclastogenesis (16).

References:

1. Barclay, A.N. (2009) *Curr. Opin. Immunol.* **21**:47.
2. van Beek, E.M. *et al.* (2005) *J. Immunol.* **175**:7781.
3. Liu, Y. *et al.* (2005) *J. Biol. Chem.* **280**:36132.
4. Sano, S-I. *et al.* (1999) *Biochem. J.* **344**:667.
5. Lee, W.Y. *et al.* (2010) *J. Biol. Chem.* **285**:37953.
6. Fujioka, Y. *et al.* (1996) *Mol. Cell. Biol.* **16**:6887.
7. Miyashita, M. *et al.* (2004) *Mol. Biol. Cell* **15**:3950.
8. Wang, X.X. & K.H. Pfenninger (2005) *J. Cell Sci.* **119**:172.
9. Maile, L.A. *et al.* (2003) *Mol. Biol. Cell* **14**:3519.
10. Johansen, M.L. & E.J. Brown (2007) *J. Biol. Chem.* **282**:24219.
11. Takenaka, K. *et al.* (2007) *Nat. Immunol.* **8**:1313.
12. Ishikawa-Sekigami, T. *et al.* (2006) *Biochem. Biophys. Res. Commun.* **343**:1197.
13. Olsson, M. *et al.* (2005) *Blood* **105**:3577.
14. Ide, K. *et al.* (2007) *Proc. Natl. Acad. Sci. USA* **104**:5062.
15. Gardai, S.J. *et al.* (2003) *Cell* **115**:13.
16. Lundberg, P. *et al.* (2007) *Biochem. Biophys. Res. Commun.* **352**:444.
17. Subramanian *et al.* (2006) *Blood* **107**:2548.