## **PRODUCT** INFORMATION



SIRPa Extracellular Domain (human, recombinant)

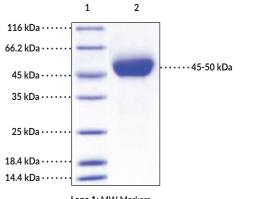
Item No. 32082

### **Overview and Properties**

Synonyms:	CD172A, CD172 Antigen-like Family Member A, MyD-1 Antigen, SHPS-1, SHP Substrate 1, Signal Regulatory Protein α
Source:	Active recombinant human C-terminal His-tagged SIRP $\alpha$ expressed in HEK293 cells
Amino Acids:	31-370
Uniprot No.:	P78324-2
Molecular Weight:	39 kDa
Storage:	-80°C (as supplied)
Stability:	≥1 year
Purity:	≥98% estimated by SDS-PAGE
Supplied in:	Lyophilized from sterile PBS, pH 7.4
Endotoxin Testing:	<1.0 EU/µg, determined by the LAL endotoxin assay
Bioactivity:	See figures for details
Information represents	the product experience. Detek experies and tical results are provided an each continents of and rise

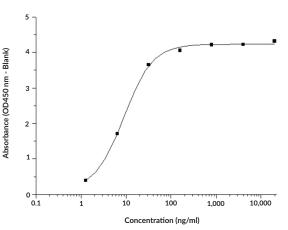
Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.





Lane 1: MW Markers Lane 2: SIRPa Extracellular Domain

SDS-PAGE Analysis of SIRPa Extracellular Domain. This protein has a calculated molecular weight of 39 kDa. It has a apparent molecular weight of approximately 45-50 kDa by SDS-PAGE under reducing conditions due to glycosylation.



SIRPa binding in functional ELISA. Immobilized human SIRPa at 10 µg/ml (100 µl/well) can bind human CD47-Fc. The EC<sub>50</sub> value of human CD47-Fc is 10.1-23.5 ng/ml.

WARNING THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

#### SAFETY DATA

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

Buyer agrees to purchase the material subject to Cayman's Terms and Conditions. Complete Terms and Conditions including Warranty and Limitation of Liability information can be found on our website.

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#### CAYMAN CHEMICAL

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# **PRODUCT** INFORMATION



#### Description

Signal regulatory protein  $\alpha$  (SIRP $\alpha$ ) is an inhibitory transmembrane receptor encoded by SIRPA in humans.<sup>1,2</sup> It is composed of three extracellular immunoglobulin-like (Ig-like) domains, a single transmembrane region, and a cytoplasmic domain that contains four tyrosine residues with immunoreceptor tyrosine-based inhibitory motifs (ITIMs) that are essential to SIRP $\alpha$  interaction with the Src homology region 2 (SH2) domain-containing phosphatases SHP-1 and SHP-2.<sup>2</sup> SIRP $\alpha$  is expressed on monocytes, macrophages, granulocytes, subsets of dendritic cells in lymphoid tissues, some bone marrow progenitor cells, and neurons. Binding of CD47 (Item No. 32085), a receptor expressed on healthy cells but often overexpressed in cancer cells, to SIRPa induces phosphorylation of the SIRPa ITIMs, coupling to SHP-1 or SHP-2, and repression of immune cell activation and phagocytosis.<sup>1,2</sup> Knockout of *Sirpa* increases dopaminergic neuronal loss in the substantia nigra pars compacta and enhances striatal microglia activation in a mouse model of MPTP-induced Parkinson's disease.<sup>3</sup> Sirpa<sup>-/-</sup> mice exhibit increased age-related podocyte injury and proteinuria compared with wild-type mice.<sup>4</sup> SIRPA expression is decreased in patients with focal segmental glomerular sclerosis and in mouse models of renal injury induced by puromycin aminonucleoside (Item No. 15509), doxorubicin (Item No. 15007), or streptozotocin (Item No. 13104). Cayman's SIRPa Extracellular Domain (human, recombinant) protein can be used for ELISA. This protein consists of 351 amino acids, has a calculated molecular weight of 39 kDa, and a predicted N-terminus of Glu31 after signal peptide cleavage. By SDS-PAGE, under reducing conditions, the apparent molecular mass of this protein is 45 to 50 kDa due to glycosylation.

#### References

- 1. Veillette, A. and Chen, J. SIRPα-CD47 immune checkpoint blockade in anticancer therapy. *Trends Immunol.* **39(3)**, 173-184 (2018).
- Barclay, A.N. and Van den Berg, T.K. The interaction between signal regulatory protein alpha (SIRPα) and CD47: Structure, function, and therapeutic target. Annu. Rev. Immunol. 32, 25-50 (2014).
- Wang, J., Ding, X., Wu, X., et al. SIRPα deficiency accelerates the pathologic process in models of Parkinson disease. Glia 67(12), 2343-2359 (2019).
- Li, L., Liu, Y., Li, S., *et al.* Signal regulatory protein α protects podocytes through promotion of autophagic activity. JCI Insight 5(9), e124747 (2019).

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