

Datasheet for ABIN2859310 S100A8 ELISA Kit

Image



Overview

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Quantity:	96 tests
Target:	S100A8
Binding Specificity:	AA 2-89
Reactivity:	Mouse
Method Type:	Sandwich ELISA
Detection Range:	31.2-2000 pg/mL
Minimum Detection Limit:	31.2 pg/mL
Application:	ELISA

Product Details

Purpose:	Sandwich High Sensitivity ELISA kit for Quantitative Detection of Mouse S100A8
Brand:	PicoKine™
Sample Type:	Cell Culture Supernatant, Serum, Plasma (heparin), Plasma (EDTA)
Analytical Method:	Quantitative
Detection Method:	Colorimetric
Immunogen:	Expression system for standard: E.coli Immunogen sequence: P2-E89
Specificity:	E.coli, P2-E89
Cross-Reactivity (Details):	There is no detectable cross-reactivity with other relevant proteins.
Sensitivity:	<10pg/mL

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Product Details

Material not included:

Microplate reader in standard size. Automated plate washer. Adjustable pipettes and pipette tips. Multichannel pipettes are recommended in the condition of large amount of samples in the detection. Clean tubes and Eppendorf tubes. Washing buffer (neutral PBS or TBS). Preparation of 0.01M TBS: Add 1.2g Tris, 8.5g Nacl

Target Details

Target:	S100A8
Alternative Name:	S100A8 (S100A8 Products)
Background:	Protein Function: S100A8 is a calcium- and zinc-binding protein which plays a prominent role i
	the regulation of inflammatory processes and immune response. It can induce neutrophil
	chemotaxis and adhesion. Predominantly found as calprotectin (S100A8/A9) which has a wide
	plethora of intra- and extracellular functions. The intracellular functions include: facilitating
	leukocyte arachidonic acid trafficking and metabolism, modulation of the tubulin-dependent
	cytoskeleton during migration of phagocytes and activation of the neutrophilic NADPH-oxidas
	Activates NADPH- oxidase by facilitating the enzyme complex assembly at the cell membrane
	transferring arachidonic acid, an essential cofactor, to the enzyme complex and S100A8
	contributes to the enzyme assembly by directly binding to NCF2/P67PHOX. The extracellular
	functions involve proinfammatory, antimicrobial, oxidant-scavenging and apoptosis-inducing
	activities. Its proinflammatory activity includes recruitment of leukocytes, promotion of cytoki
	and chemokine production, and regulation of leukocyte adhesion and migration. Acts as an
	alarmin or a danger associated molecular pattern (DAMP) molecule and stimulates innate
	immune cells via binding to pattern recognition receptors such as Toll-like receptor 4 (TLR4)
	and receptor for advanced glycation endproducts (AGER). Binding to TLR4 and AGER activate
	the MAP-kinase and NF- kappa-B signaling pathways resulting in the amplification of the
	proinflammatory cascade. Has antimicrobial activity towards bacteria and fungi and exerts its
	antimicrobial activity probably via chelation of Zn(2+) which is essential for microbial growth.
	Can induce cell death via autophagy and apoptosis and this occurs through the cross-talk of
	mitochondria and lysosomes via reactive oxygen species (ROS) and the process involves
	BNIP3. Can regulate neutrophil number and apoptosis by an anti-apoptotic effect, regulates ce
	survival via ITGAM/ITGB and TLR4 and a signaling mechanism involving MEK-ERK. Its role as
	an oxidant scavenger has a protective role in preventing exaggerated tissue damage by
	scavenging oxidants. The iNOS-S100A8/A9 transnitrosylase complex is proposed to direct
	selective inflammatory stimulus-dependent S- nitrosylation of multiple targets such as GAPDF
	ANXA5, EZR, MSN and VIM by recognizing a [IL]-x-C-x-x-[DE] motif, S100A8 seems to contribut
	to S-nitrosylation site selectivity (By similarity)

	Background: S100 calcium-binding protein A8 (S100A8), also known as calgranulin A, is a
	protein that in humans is encoded by the S100A8 gene. It is mapped to chromosome 1q21.3
	based on an alignment of the S100A8 sequence with the genomic sequence. The proteins
	S100A8 and S100A9 form a heterodimer called calprotectin which is a major calcium- and zinc-
	binding protein in the cytosol of neutrophils, monocytes, and keratinocytes. This gene is a
	member of the S100 family of proteins containing 2 EF-hand calcium-binding motifs. S100
	proteins are localized in the cytoplasm and/or nucleus of a wide range of cells, and involved in
	the regulation of a number of cellular processes such as cell cycle progression and
	differentiation. And S100 genes include at least 13 members which are located as a cluster on
	chromosome 1q21. This protein may function in the inhibition of casein kinase and as a
	cytokine. Altered expression of this protein is associated with the disease cystic fibrosis.
	Synonyms: Protein S100-A8,Calgranulin-A,Chemotactic cytokine CP-10,Leukocyte L1 complex
	light chain,Migration inhibitory factor-related protein 8,MRP-8,p8,Pro-inflammatory S100
	cytokine,S100 calcium-binding protein A8,S100a8,Caga, Mrp8,
	Full Gene Name: Protein S100-A8
	Cellular Localisation: Secreted . Cytoplasm . Cytoplasm, cytoskeleton . Cell membrane,
	Peripheral membrane protein . Predominantly localized in the cytoplasm. Upon elevation of the
	intracellular calcium level, translocated from the cytoplasm to the cytoskeleton and the cell
	membrane. Upon neutrophil activation or endothelial adhesion of monocytes, is secreted via a
	microtubule-mediated, alternative pathway.
Gene ID:	20201
UniProt:	P27005

Pathways:

Transition Metal Ion Homeostasis, Positive Regulation of Endopeptidase Activity, S100 Proteins

Application Details

Application Notes:	Before using Kit, spin tubes and bring down all components to bottom of tube. Duplicate well assay was recommended for both standard and sample testing.
Plate:	Pre-coated
Protocol:	mouse S100A8 ELISA Kit was based on standard sandwich enzyme-linked immune-sorbent assay technology. A monoclonal antibody from rat specific for S100A8 has been precoated onto 96-well plates. Standards(E.coli, P2-E89) and test samples are added to the wells, a biotinylated detection polyclonal antibody from goat specific for S100A8 is added subsequently and then followed by washing with PBS or TBS buffer. Avidin-Biotin-Peroxidase Complex was

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Application Details

	added and unbound conjugates were washed away with PBS or TBS buffer. HRP substrate TMB was used to visualize HRP enzymatic reaction. TMB was catalyzed by HRP to produce a blue color product that changed into yellow after adding acidic stop solution. The density of yellow is proportional to the mouse S100A8 amount of sample captured in plate.
Assay Procedure:	Aliquot 0.1 mL per well of the 2000pg/mL,1000pg/mL, 500pg/mL, 250pg/mL, 125pg/mL, 62.5pg/mL, 31.2pg/mL mouse S100A8 standard solutions into the precoated 96-well plate. Add 0.1 mL of the sample diluent buffer into the control well (Zero well). Add 0.1 mL of each properly diluted sample of mouse cell culture supernates, serum or plasma(heparin, EDTA) to each empty well. See "Sample Dilution Guideline" above for details. It is recommended that each mouse S100A8 standard solution and each sample be measured in duplicate.
Assay Precision:	 Sample 1: n=16, Mean(pg/ml): 156, Standard deviation: 9.048, CV(%): 5.8 Sample 2: n=16, Mean(pg/ml): 724, Standard deviation: 34.03, CV(%): 4.7 Sample 3: n=16, Mean(pg/ml): 1345, Standard deviation: 55.15, CV(%): 4.1, Sample 1: n=24, Mean(pg/ml): 174, Standard deviation: 11.31, CV(%): 6.5 Sample 2: n=24, Mean(pg/ml): 617, Standard deviation: 33.32, CV(%): 5.4 Sample 3: n=24, Mean(pg/ml): 1436, Standard deviation: 112.01, CV(%): 7.8
Restrictions: Handling	For Research Use only
Handling Advice:	Avoid multiple freeze-thaw cycles.
Storage:	-20 °C,4 °C
Storage Comment:	Store at 4°C for 6 months, at -20°C for 12 months. Avoid multiple freeze-thaw cycles
Expiry Date:	12 months



ELISA

Image 1. Mouse S100A8 PicoKine ELISA Kit standard curve

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