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ICAM1 ELISA Kit





Overview

Quantity:	96 tests
Target:	ICAM1
Reactivity:	Human
Method Type:	Sandwich ELISA
Detection Range:	62.5-4000 pg/mL
Minimum Detection Limit:	62.5 pg/mL
Application:	ELISA

Product Details

Sample Type:	Cell Culture Supernatant, Serum, Plasma (heparin), Plasma (citrate), Plasma (EDTA)
Analytical Method:	Quantitative
Detection Method:	Colorimetric
Specificity:	Natural and recombinant Human sICAM-1 Ligand
Sensitivity:	31 pg/mL
Material not included:	Microplate reader.Pipettes and pipette tips.

Target Details

Target:	ICAM1	

· EP tube Deionized or distilled water.

Alternative Name:	sICAM-1 (ICAM1 Products)
Target Type:	Viral Protein
Background:	Intercellular Adhesion Molecule 1 (ICAM-1), also known as CD54, is a nearly ubiquitous
	transmembrane glycoprotein that plays a key role in leukocyte migration and activation (1, 2).
	Human ICAM-1 contains five Ig-like domains in its extracellular domain (ECD) and associates
	into non-covalently linked dimers (3, 4). Soluble forms of monomeric and dimeric ICAM-1
	(sICAM-1) can be generated via proteolytic cleavage by cathepsin G, elastase, MMP-9, MMP-
	14/MT1-MMP, and TACE/ADAM17 (5 - 8). In the mouse, alternate splicing generates isoforms
	that lack particular lg-like domains and are differentially sensitive to proteolysis (5). Within the
	ECD, human ICAM-1 shares 53 % amino acid sequence identity with mouse and rat ICAM-1. The
	principal binding partners of ICAM-1 are the leukocyte integrins LFA-1 (CD11a/CD18) and Mac-
	1 (CD11b/CD18) (9 - 11). The multivalency of dimeric ICAM-1 increases its strength of
	interaction with LFA-1 (9, 10). ICAM-1 also binds several non-integrin ligands including
	CD43/sialophorin, fibrinogen, hyaluronan, rhinoviruses, and Plasmodium falciparum-infected
	erythrocytes (12 - 16). At sites of inflammation, ICAM-1 is upregulated on endothelial and
	epithelial cells where it mediates the adhesion and paracellular migration of leukocytes
	expressing activated LFA-1 and Mac-1 (17 - 20). ICAM-1 ligation prolongs antigen presentation
	by dendritic cells and promotes T cell proliferation and cytokine release (21 - 23). ICAM-1
	activation also participates in angiogenesis, wound healing, and bone metabolism (24 -
	26). Soluble ICAM-1 has been reported in serum, cerebrospinal fluid, urine, and bronchoalveolar
	lavage fluid (2, 27 - 31). Elevated levels of sICAM-1 in these fluids are associated with
	cardiovascular disease, type 2 diabetes, organ transplant dysfunction, oxidant stress,
	abdominal fat mass, hypertension, liver disease, and certain malignancies (32 - 40). sICAM-1
	promotes angiogenesis and serves as an indicator of vascular endothelial cell activation or
	damage (41, 42). It also functions as an inhibitor of transmembrane ICAM-1 mediated activities
	such as monocyte adhesion to activated endothelial cells and sensitivity of tumor cells to NK
	cell-mediated lysis (7, 8).
Pathways:	Cellular Response to Molecule of Bacterial Origin, Regulation of Actin Filament Polymerization,
	Carbohydrate Homeostasis, Regulation of Leukocyte Mediated Immunity, Thromboxane A2
	Receptor Signaling
Application Details	
Application Notes:	Detection Wavelength: 450 nm

Sample Volume:

 $20\,\mu L$

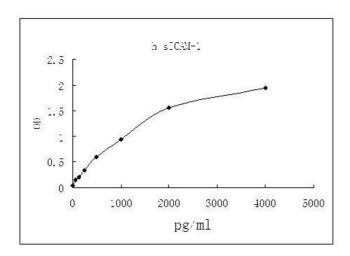
Application Details

Assay Time:	3 h
Plate:	Pre-coated
Restrictions:	For Research Use only

Handling

Storage: 4 °C

Images



ELISA

Image 1.