

Human VCAM-1 / CD106 ELISA Instructions

Cat: EH0067

CONTENT

	CAT	Volume
① CP (Coated Plate)	EH0067CP	96 well
2 S (Standard)	EH0067S,S1~S7,S0	9 vial
3 DA (Detect Antibody)	EH0067DA	6 ml/bottle
4 SD (Sample Diluent)	ESD01	12 ml/bottle
5 SH (Streptavidin-HRP)	ESH04	12 ml/bottle
(3 AB (Assay Buffer 1×)	EAB01	12 ml/bottle
7 TS (TMB Substrate)	ETS01	12 ml/bottle
8 SS (Stop Solution)	ESS01	12 ml/bottle
9 WB (Wash Buffer 10×)	EWB01	50 ml/bottle
SF (Sealer Film)	ESF01	6 pieces

NOTE: After the kit is opened, the stabilization period of each content is 30 days.

SAMPLE DILUTION

Samples such as serum、plasma require at least a 40-fold dilution into Sample Diluent. A suggested 40-fold dilution is 5 μ l of sample + 195 μ l of Sample Diluent

REAGENT PREPARATION

Washing Buffer (1×) Preparation

Pour entire contents (50 ml) of the Washing Buffer Concentrate (10×) into a clean 500 ml graduated cylinder. Bring to final volume of 500 ml with glass-distilled or deionized water. Transfer to a clean wash bottle and store at 2 to 25°C.

Standard Curve Preparation:

 ${\rm S1}$ to ${\rm S7}$ and ${\rm S0}$ is ready to use for serum and plasma.

Other sample type, prepare the standard curve with whatever buffer (SPB, Sample Prepared Buffer) is used to prepare the sample, such as cell culture supernatant, tissue grinding liquid, cell lysate, etc. Urine sample use AB (Assay Buffer) prepare standard curve.

The human VCAM-1 Standard EH0067S 300000 pg/ml 30 μ l + 270 μ l SPB serves as the high standard (30000 pg/ml). Pipette 200 μ l of SPB into each tube. Use the high standard to produce a 1:2 dilution series. Mix each tube thoroughly before the next transfer. SPB serves as the zero standard (0 pg/ml).



ASSAY PROCEDURE

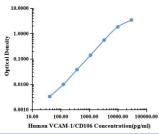
Bring all reagents and samples to room temperature before use.

- 1 Prepare all reagents and working standards as directed in the previous sections.
- 2 Remove excess CP (Coated Plate) strips from the plate frame, return them to the foil pouch and reseal.
- 3 Add 50 µl of AB (Assay Buffer) to each well.
- 4 Add 50 μl of Standard or sample per well. Ensure reagent addition is uninterrupted and completed within 15 minutes.
- \bigcirc Add 50 μ l of DA (Detect Antibody) to each well.
- **6** Cover with an **SF** (Sealer Film). Incubate at room temperature (18 to 25°C) for 1 hour on a microplate **shaker** set at 500 rpm.
- $\ensuremath{\mathfrak{T}}$ Aspirate each well and wash, repeating the process four times. Wash by filling each well with WB (Washing Buffer 300 μ l). Complete removal of liquid at each step is essential to good performance. After the last wash, remove any remaining WB (Washing Buffer) by aspirating or decanting. Invert the plate and blot it against clean paper towels.
- 8 Add 100 μ l of SH (Streptavidin-HRP) to each well
- **9** Cover with a new **SF** (Sealer Film). Incubate at room temperature (18 to 25°C) for 30 min on a microplate **shaker** set at 500 rpm.
- 10 Repeat aspiration/wash as in step 7.
- 11) Add 100 μ l of TS (TMB Substrate) to each well. Incubate for 5-30 minutes at room temperature.
- 12 Add 100 ul of SS (Stop Solution) to each well.
- (B) Determine the optical density within 30 minutes, using microplate **reader** set to 450 nm corrected with 570 nm or 630 nm.



TYPICAL DATA

Human VCAM-1/CD106 Typical Standard



pg/ml	О.	O.D.		Corrected
0.00	0.0090	0.0093	0.0092	
41.15	0.0130	0.0118	0.0124	0.0033
123.46	0.0190	0.0189	0.0190	0.0098
370.37	0.0495	0.0436	0.0466	0.0374
1111.11	0.1538	0.1445	0.1492	0.1400
3333.33	0.5925	0.5271	0.5598	0.5507
10000.00	1.9090	1.7580	1.8335	1.8244
30000.00	3.4470	3.4170	3.4320	3.4229

SENSITIVITY

The minimum detectable dose (MDD) of human VCAM-1 is typically less than 7.07 pg/ml.

The MDD was determined by adding two standard deviations to the mean optical density value of ten zero standard replicates and calculating the corresponding concentration.

PRECISION

Intra-assay Precision (Precision within an assay) Three samples of known concentration were tested twenty times on one plate to assess intra-assay precision.

Inter-assay Precision (Precision between assays)

	Intra-assay Precision		Inter-assay Precision			
Sample Number	S1	S2	S3	S1	S2	S3
	22	22	22	6	6	6
Average (pg/ml)	646.6	3231.2	10895.8	635.0	3119.9	10903.3
Standard Deviation	27.9	158.1	698.2	47.7	184.0	787.2
Coefficient of Variation (%)	4.3	4.9	6.4	7.5	5.9	7.2

RECOVERY

The spike recovery was evaluated by spiking 3 levels of human VCAM-1 into health human serum sample. The un-spiked serum was used as blank in this experiment.

The recovery ranged from 91% to 117% with an overall mean recovery of 105%.

LINEARITY

To assess the linearity of the assay, five samples were spiked with high concentration of VCAM-1 in human serum and diluted with Sample Diluent to produce samples with values within the dynamic range of the assay.

The linearity ranged from 97% to 115% with an overall mean recovery of 103%.

SAMPLE VALUES

Serum/Plasma – Thirty samples from apparently healthy volunteers were evaluated for the presence of human VCAM-1 in this assay. No medical histories were available for the donors.

Sample Matrix	Sample Evaluated	Range (ng/ml)	Detectable %	Mean of Detectable (ng/ml)
Serum	30	212.34-306.21	100	255.91

n.d. = non-detectable. Samples measured below the sensitivity are considered to be non-detectable.