

Human CD147 Sandwich ELISA Kit Datasheet

Please read it entirely before use

Catalogue Number: KE00135

Size: 5*96T

Sensitivity: 2.1 pg/mL

Range: 31.25-2000 pg/mL

Usage: For the quantitative detection of human CD147 concentrations in serum, plasma, cell culture supernatant, urine, saliva and human milk.

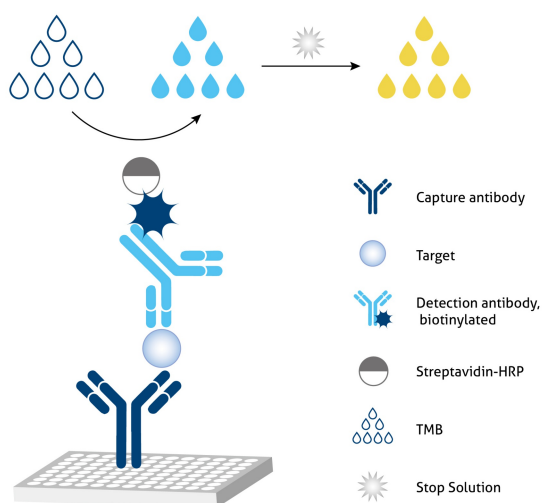
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1. Background

CD147, also called Basigin (BSG) or EMMPRIN, is a transmembrane glycoprotein that belongs to the immunoglobulin superfamily and is important in numerous physiological and pathological phenomena. It is involved in reproduction, neural function, inflammation and tumor invasion. CD147 is broadly distributed, and expressed at particularly high levels at the retinal pigment epithelium and neonatal blood-brain barrier, on tumour cells and on activated T cells. CD147 can stimulate adjacent fibroblasts to produce matrix metalloproteinases (MMPs). It has been considered as an objective and effective marker to predict invasion and prognosis in various cancers. Soluble CD147 (sCD147) has been uncovered in addition to membrane-bound CD147. sCD147 production is mediated by two different mechanisms: microvesicular secretion and proteolytic cleavage. The presence of sCD147 in body fluids is associated with several diseases such as systemic sclerosis, hepatocellular carcinoma, ankylosing spondylitis and coronary diseases.

2. Principle



Sandwich ELISA structure (Detection antibody labeled with biotin)

A capture antibody is pre-coated onto the bottom of wells which binds to analyte of interest. A detection antibody labeled with biotin also binds to the analyte. Streptavidin-HRP binds to the biotin. TMB acts as the HRP substrate and the solution color will change from colorless to blue. A stop solution containing sulfuric acid turns solution yellow. The color intensity is proportional to the quantity of bound protein which is measurable at 450 nm with the correction wavelength set at 630 nm.

3. Required Materials

- 3.1 A microplate reader capable of measuring absorbance at 450 nm with the correction wavelength set at 630 nm.
- 3.2 Calibrated, adjustable precision pipettes and disposable plastic tips. A manifold multi-channel pipette is recommended for large assays.
- 3.3 Plate washer: automated or manual.
- 3.4 Absorbent paper towels.
- 3.5 Glass or plastic tubes to prepare standard and sample dilutions.
- 3.6 Beakers and graduated cylinders.
- 3.7 Log-log or semi-log graph paper or computer and software for ELISA data analysis. A four-parameter logistic (4-PL) curve-fit is recommended.

4. Kit Components and Storage

Microplate - antibody coated 96-well microplate (8 well × 12 strips)	5 plates	Unopened Kit: Store at 2-8°C for 6 months or -20°C for 12 months. Opened Kit: All reagents stored at 2-8°C for 7 days. Please use a new standard for each assay.
Protein standard - 2000 pg/bottle; lyophilized	10 bottles	
Detection antibody, biotinylated (100×) - 600 μL/vial*	1 vial	
Streptavidin-horseradish peroxidase (HRP) (100×) - 600 μL/vial*	1 vial	
Sample Diluent PT 3 - 150 mL/bottle	1 bottle	
Detection Diluent - 150 mL/bottle	1 bottle	
Wash Buffer Concentrate (20×) - 150 mL/bottle	1 bottle	
Tetramethylbenzidine Substrate (TMB) - 60 mL/bottle	1 bottle	
Stop Solution - 60 mL/bottle	1 bottle	
Plate Cover Seals	15 pieces	

* Centrifugation immediately before use

5. Safety Notes

- 5.1 Avoid any skin and eye contact with Stop Solution and TMB. In case of contact, wash thoroughly with water.
- 5.2 Do not use the kit after the expiration date.
- 5.3 Do not mix or substitute reagents or materials from other kit lots or other sources.
- 5.4 Be sure to wear protective equipment such as gloves, masks and goggles during the experiment.
- 5.5 When using an automated plate washer, adding a 30 second soak period following the addition of Wash Buffer to improve assay precision

6. Sample Collection and Storage

- 6.1 Serum: Allow blood samples to clot for 30 minutes, followed by centrifugation for 15 minutes at 1000xg. Clear serum can be assayed immediately or aliquoted and stored at -20°C. Avoid repeated freeze-thaw cycles.
- 6.2 Plasma: Use EDTA, heparin, or citrate as an anticoagulant for plasma collection. Centrifuge for 15 minutes at 1000xg within 30 minutes of collection. The plasma can be assayed immediately or aliquoted and stored at -20°C. Avoid repeated freeze-thaw cycles.
- 6.3 Cell Culture Supernatant: Remove particulates by centrifugation for 5 minutes at 500xg and assay immediately or aliquot and store samples at ≤ -20°C. Avoid repeated freeze-thaw cycles.
- 6.4 Urine: Collect urine samples and centrifuge for 20 minutes at 1000xg. Collect the aqueous layer, assay immediately or aliquot and store samples at ≤ -20°C. Avoid repeated freeze-thaw cycles.
- 6.5 Saliva: Collect saliva samples and centrifuge for 5 minutes at 10,000xg. Collect the aqueous layer, assay immediately or aliquot and store samples at ≤ -20°C. Avoid repeated freeze-thaw cycles.
- 6.6 Human Milk: Collect milk samples and Centrifuge for 15 minutes at 1000xg at 2-8°C. Collect the aqueous fraction and repeat this process a total of 3 times. Assay immediately.

7. Regent Preparation

7.1 Wash Buffer (1X): If crystals have formed in the concentrate, warm to room temperature and mix gently until the crystals have completely dissolved. Add 30 mL of Wash Buffer Concentrate(20X) to 570 mL deionized or distilled water to prepare 1X Wash Buffer.

7.2 Detection Antibody (1X): Dilute 100X Detection Antibody 1:100 using Detection Diluent prior to assay. Suggested 1:100 dilution: 10 μ L 100X Detection Antibody + 990 μ L Detection Diluent (Centrifuge the 100 X Detection Antibody solution for a few seconds prior to use).

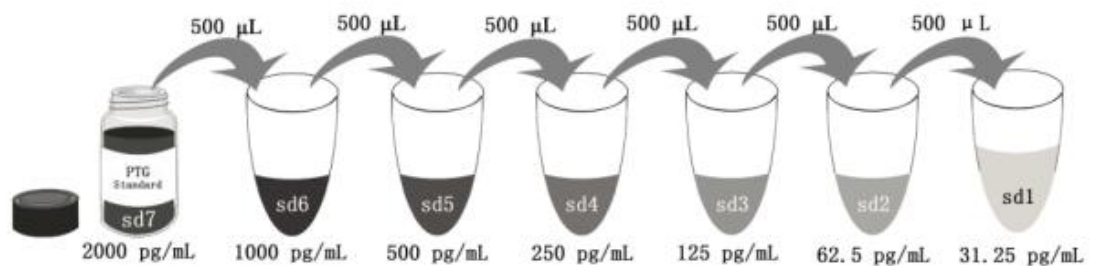
7.3 Streptavidin-HRP (1X): Dilute 100X Streptavidin-HRP 1:100 using Detection Diluent prior to assay. Suggested 1:100 dilution: 10 μ L 100X Streptavidin-HRP + 990 μ L Detection Diluent (Centrifuge the 100X Streptavidin-HRP solution for a few seconds prior to use).

7.4 Sample Dilution: Different samples should be diluted with corresponding Sample Diluent, samples may require further dilution if the readout values are higher than the highest standard OD reading. Variations in sample collection, processing and storage may affect the results of the measurement.

Recommended Dilution for different sample types: 1:4 is recommended for human serum and plasma; 1:4 is recommended for cell culture supernatant; 1:10 is recommended for urine; 1:2 is recommended for saliva; 1:40 is recommended for human milk.

7.5 Standard Serial Dilution:

Add 1 mL Sample Diluent PT 3 in protein standard.



Add # μ L of Standard diluted in the previous step	—	500 μ L	500 μ L	500 μ L	500 μ L	500 μ L	500 μ L
# μ L of Sample Diluent PT 3	1000 μ L	500 μ L	500 μ L	500 μ L	500 μ L	500 μ L	500 μ L
	"sd7"	"sd6"	"sd5"	"sd4"	"sd3"	"sd2"	"sd1"

8. Assay Procedure Summary

Bring all reagents to room temperature before use (Detection antibody and Streptavidin-HRP can be used immediately). To avoid cross-contamination, change pipette tips between additions of each standard level, between sample additions, and between reagent additions. Also, use separate reservoirs for each reagent.

8.1 Take out the required number of microplate strips and return excess strips to the foil pouch containing the drying reagent pack and reseal; store at 4°C immediately. Microplate strips should be used in one week.

8.2 Preset the layout of the microplate, including control group, standard group and sample group, add 100 µL of each standard and sample to the appropriate wells. (Make sure sample addition is uninterrupted and completed within 5 to 10 minutes, it is recommended to assay all standards, controls, and samples in duplicate).

8.3 Seal plate with cover seal, pressing it firmly onto top of microwells. Incubate the plate for 2 hours at 37°C.

8.4 Wash

1) Gently remove the cover seal. Discard the liquid from wells by aspirating or decanting. Remove any residual solution by tapping the plate a few times on fresh paper towels.

2) Wash 4 times with 1X Wash Buffer, using at least 350-400 µL per well. Following the last wash, firmly tap plates on fresh towels 10 times to remove residual Wash Buffer. Avoid getting any towel fibers in the wells or wells drying out completely.

8.5 Add 100 µL of 1X Detection Antibody solution (refer to Reagent Preparation 7.2) to each well. Seal plate with cover seal and incubate for 1 hour at 37°C.

8.6 Repeat wash step in 8.4.

8.7 Add 100 µL of 1X Streptavidin-HRP solution (refer to Reagent Preparation 7.3) to each well. Seal plate with cover seal and incubate the plate for 40 minutes at 37°C.

8.8 Repeat wash step in 8.4.

8.9 Signal development: Add 100 µL of TMB substrate solution to each well, protected from light. Incubate for 15 to 20 minutes. Substrate Solution should remain colorless until added to the plate.

8.10 Quenching color development: Add 100 µL of Stop Solution to each well in the same order as addition of the TMB substrate. Mix by tapping the side of the plate gently. NB: Avoid skin and eye contact with the Stop solution.

8.11 Read results: Immediately after adding Stop solution read the absorbance on a microplate reader at a wavelength of 450 nm. If possible, perform a double wavelength readout (450 nm and 630 nm).

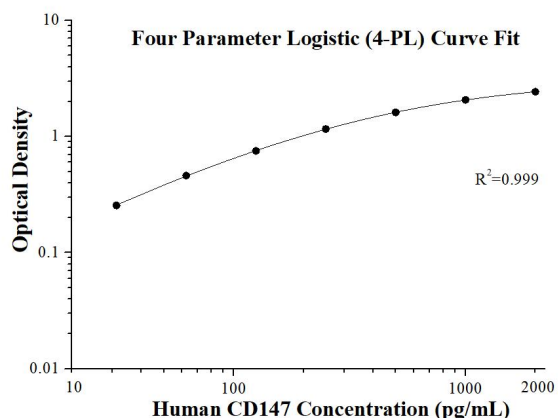
8.12 Data analysis: Calculate the average of the duplicate readings (OD value) for each standard and sample, and subtract the average of the zero standard absorbance. Construct a standard curve by plotting the mean absorbance for each standard on the y-axis against the concentration on the x-axis, use four-parameter logistic curve-fit (4-PL) analysis to do this. If the samples have been diluted, the OD readout from the standard curve must be multiplied by the dilution factor used.

Step	Reagent	Volume	Incubation	Wash	Notes
1	Standard and Samples	100 µL	120 min	4 times	Cover Wells incubate at 37°C
2	Diluent Antibody Solution	100 µL	60 min	4 times	Cover Wells incubate at 37°C
3	Diluent HRP Solution	100 µL	40 min	4 times	Cover Wells incubate at 37°C
4	TMB Substrate	100 µL	15-20 min	Do not wash	Incubate in the dark at 37°C
5	Stop Solution	100 µL	0 min	Do not wash	-
6	Read plate at 450 nm and 630 nm immediately after adding Stop solution. DO NOT exceed 5 minutes.				

9. Validation Data

9.1 Standard curve

These standard curves are provided for demonstration only. A standard curve should be generated for each set of samples assayed.



(pg/mL)	O.D	Average	Corrected
0	0.062 0.062	0.062	-
31.25	0.325 0.309	0.317	0.255
62.5	0.534 0.512	0.523	0.461
125	0.826 0.803	0.815	0.753
250	1.250 1.195	1.223	1.161
500	1.700 1.653	1.677	1.615
1000	2.168 2.095	2.095	2.070
2000	2.541 2.479	2.497	2.43

9.2 Precision

Intra-assay Precision (Precision within an assay) Three samples of known concentration were tested 20 times on one plate to assess intra-assay precision.

Inter-assay Precision (Precision between assays) Three samples of known concentration were tested in 24 separate assays to assess inter-assay precision.

Intra-assay Precision				
Sample	n	Mean (pg/mL)	SD	CV%
1	20	56.5	1.1	2.0
2	20	219.7	6.7	3.1
3	20	816.5	33.5	4.1

Inter-assay Precision				
Sample	n	Mean (pg/mL)	SD	CV%
1	24	56.2	2.5	4.4
2	24	221.9	9.6	4.3
3	24	865.6	58.0	6.7

9.3 Recovery

The recovery of human CD147 spiked to three different levels throughout the range of the assay in various matrices was evaluated.

Sample Type		Average% of Expected	Range (%)
Human plasma	1:16	106	93-121
	1:32	96	75-115
Human serum	1:16	95	76-117
	1:32	99	87-118
Cell culture supernatant	1:8	92	80-112
	1:16	93	80-115
Human milk	1:40	105	88-125
	1:80	102	95-112
Urine	1:20	102	78-121
	1:40	106	91-121
Saliva	1:4	98	84-118
	1:8	95	85-115

9.4 Sample values

Samples from healthy volunteers were evaluated for human CD147 in this assay. No medical histories were available for the donors used in this study.

Sample Type	Mean of Detectable (pg/mL)	Range (pg/mL)
Human serum (n=24)	3,146	1,946-5,924
Human milk (n=7)	5,824	3,636-9,293
Urine (n=7)	4,184	1,015-10,260
Saliva (n=7)	3,849	818-10,260

Human peripheral blood leucocytes cells (3.5×10^6 cells/mL) were cultured in DMEM supplemented with 8% fetal bovine serum, $5 \mu\text{M}$ β -mercaptoethanol, 2 mM L-glutamine, 100 U/mL penicillin, and 100 $\mu\text{g/mL}$ streptomycin sulfate. Cells were cultured unstimulated or stimulated with 10 $\mu\text{g/mL}$ PHA for 1 day, 3 days and 5 days. Aliquots of the cell culture supernatant was removed and assayed for levels of human CD147.

Condition	(pg/mL)
Unstimulated	189
Stimulated for 1d	135
Stimulated for 3d	697
Stimulated for 5d	3,645

9.5 Sensitivity

The minimum detectable dose of human CD147 is 2.1 pg/mL. This was determined by adding two standard deviations to the concentration corresponding to the mean O.D. of 20 zero standard replicates.

9.6 Linearity

To assess the linearity of the assay, samples were diluted with the appropriate **Sample Diluent** to produce samples with values within the dynamic range of the assay. (The human plasma, serum samples and cells culture supernatant was initially diluted 1:2, the human milk were initially diluted 1:20, the urine were initially diluted 1:5)

		Human plasma	Human serum	Cell culture supernatant	Human milk	Urine	Saliva
1:2	Average% of Expected	100	100	100	100	100	100
	Range (%)	-	-	-	-	-	-
1:4	Average% of Expected	96	103	91	101	98	99
	Range (%)	91-101	96-113	85-99	92-108	97-99	97-101
1:8	Average% of Expected	100	113	91	105	100	101
	Range (%)	90-111	106-125	77-106	93-117	98-103	97-106
1:16	Average% of Expected	105	120	100	102	108	107
	Range (%)	94-117	114-125	83-126	100-104	107-109	106-108

10. References

1. Muramatsu T, et al. Basigin (CD147): a multifunctional transmembrane protein involved in reproduction, neural function, inflammation and tumor invasion. *Histol Histopathol.* 18(3):981-7 (2003).
2. Toole BP, et al. Emmprin (CD147), a cell surface regulator of matrix metalloproteinase production and function. *Curr Top Dev Biol.* 54:371-89 (2003).
3. Kirk P, et al. CD147 is tightly associated with lactate transporters MCT1 and MCT4 and facilitates their cell surface expression. *EMBO J.* 19(15):3896-904 (2000).
4. Sun J, et al. Regulation of MMP-1 and MMP-2 production through CD147/extracellular matrix metalloproteinase inducer interactions. *Cancer Res.* 61(5):2276-81 (2001).
5. Iacono KT, et al. CD147 immunoglobulin superfamily receptor function and role in pathology. *Exp Mol Pathol.* 83(3):283-95 (2007).
6. Laopajon W, et al. Antibody biosensors for the measurement and characterization of soluble CD147 molecules. *Asian Pac J Allergy Immunol.* 36(3):191-200 (2018).