

Human N-cadherin Sandwich ELISA Kit Datasheet

Please read it entirely before use

Catalogue Number: KE00328

Size: 96T

Sensitivity: 0.03 ng/mL

Range: 0.156 - 10 ng/mL

Usage: For the quantitative detection of human N-cadherin concentrations in serum, plasma and cell culture supernatant.

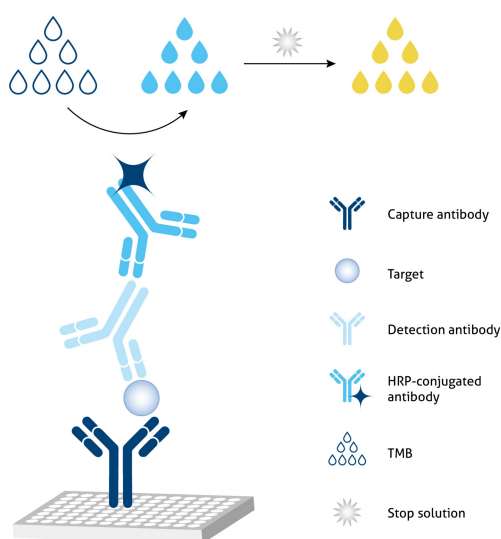
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1. Background

Cadherins are a family of transmembrane glycoproteins that mediate calcium-dependent cell-cell adhesion and play an important role in the maintenance of normal tissue architecture. N-cadherin (neural cadherin), also known as CDH2 (cadherin 2), is a 130-kDa transmembrane protein and a classical member of the cadherin superfamily which also include E-, P-, R-, and B-cadherins. Expression of N-cadherin has been reported on various cell types including neurons, endothelial cells and cardiac myocytes. N-cadherin has functions in early brain morphogenesis, synaptogenesis and synaptic plasticity. Soluble N-cadherin (sN-CAD) is found in biological fluids, including serum, seminal fluid samples, and urine. It has been shown that significantly higher amounts of serum sN-CAD are present in cancer patients than in persons with no evidence of disease.

2. Principle



Sandwich ELISA structure (HRP conjugated secondary antibody)

A capture antibody is pre-coated onto the bottom of wells which binds to analyte of interest. A detection antibody also binds to the analyte. Horseradish peroxidase (HRP)-conjugated secondary antibody binds to the detection antibody. 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) | 1 plate | 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 - 20 ng/bottle; lyophilized | 2 bottles | |
| Detection Antibody (100×) - 120 µL/vial* | 1 vial | |
| HRP-conjugated antibody (100×) - 120 µL/vial* | 1 vial | |
| Sample Diluent PT 3 - 30 mL/bottle. For human serum and plasma | 1 bottle | |
| Sample Diluent PT 1 - 30 mL/bottle. For cell culture supernatant | 1 bottle | |
| Detection Diluent - 30 mL/bottle | 1 bottle | |
| Wash Buffer Concentrate (20×) - 30 mL/bottle | 1 bottle | |
| Tetramethylbenzidine Substrate (TMB) - 12 mL/bottle | 1 bottle | |
| Stop Solution - 12 mL/bottle | 1 bottle | |
| Plate Cover Seals | 4 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.

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 100X Detection Antibody solution for a few seconds prior to use).

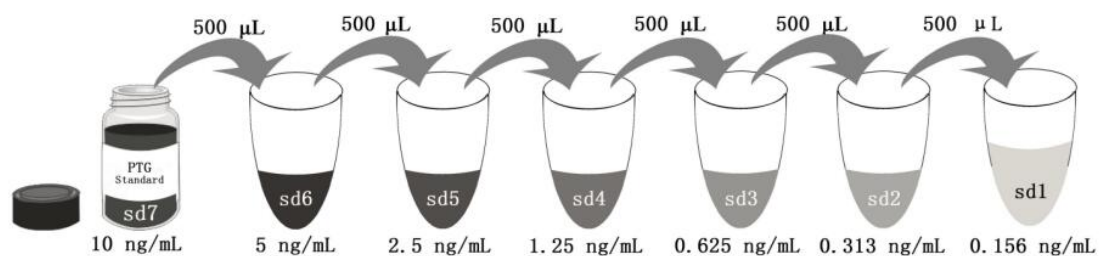
7.3 HRP-conjugated antibody(1X): Dilute 100X HRP-conjugated antibody 1:100 using Detection Diluent prior to assay. Suggested 1:100 dilution: 10 μ L 100X HRP-conjugated antibody + 990 μ L Detection Diluent (Centrifuge the 100X HRP-conjugated antibody solution for a few seconds prior to use).

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 or 1:8 is recommended for human serum and plasma; 1:2 is recommended for cell culture supernatant.

7.5 Standard Serial Dilution:

For human serum and plasma, add 2mL Sample Diluent PT 3 in protein standard. For cell culture supernatant, add 2mL Sample Diluent PT 1 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 1 or PT 3 | 2000 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 HRP-conjugated antibody 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 HRP-conjugated antibody 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).

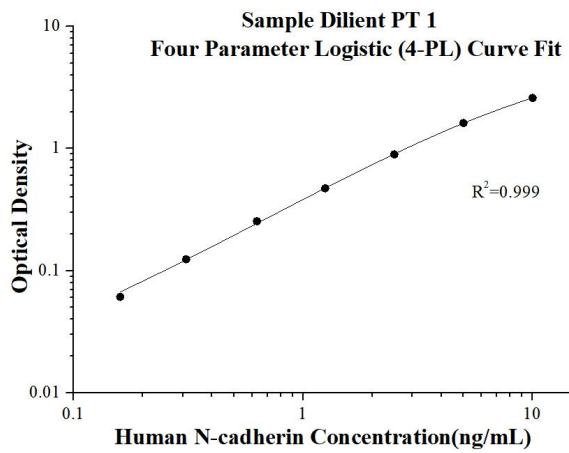
8.12 Data analysis: Calculate the average of the duplicate readings (OD value) for each standard and sample, and subtract the average of the zero standard absorbance. Construct a standard curve by plotting the mean absorbance for each standard on the y-axis against the concentration on the x-axis, use 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. | | | | |

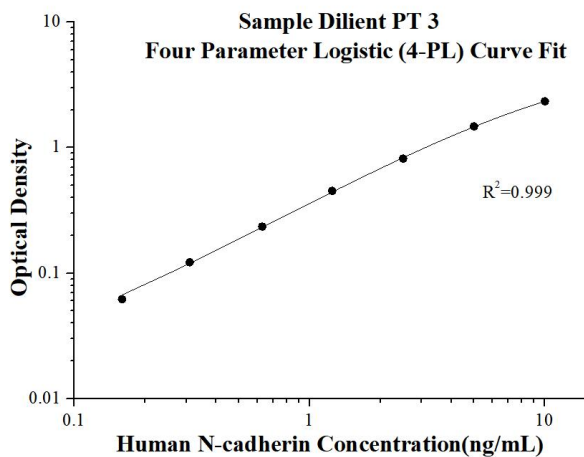
9. Validation Data

9.1 Standard curve

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



| (ng/mL) | O.D | Average | Corrected |
|---------|------------------|---------|-----------|
| 0 | 0.0754 0.0769 | 0.07615 | - |
| 0.156 | 0.1373 0.1377 | 0.1375 | 0.06135 |
| 0.313 | 0.1966 0.2029 | 0.19975 | 0.1236 |
| 0.625 | 0.3264 0.3334 | 0.3299 | 0.25375 |
| 1.25 | 0.5492 0.5485 | 0.54885 | 0.4727 |
| 2.5 | 0.9679 0.9771 | 0.9725 | 0.89635 |
| 5 | 1.6631 1.7262 | 1.69465 | 1.6185 |
| 10 | 2.6905 2.6548 | 2.67265 | 2.5965 |



| (ng/mL) | O.D | Average | Corrected |
|---------|------------------|---------|-----------|
| 0 | 0.0683 0.0693 | 0.0688 | - |
| 0.156 | 0.1301 0.1316 | 0.13085 | 0.06205 |
| 0.313 | 0.1898 0.1908 | 0.1903 | 0.1215 |
| 0.625 | 0.3049 0.3022 | 0.30355 | 0.23475 |
| 1.25 | 0.5309 0.5102 | 0.52055 | 0.45175 |
| 2.5 | 0.8937 0.8783 | 0.886 | 0.8172 |
| 5 | 1.565 1.5199 | 1.54245 | 1.47365 |
| 10 | 2.4241 2.3887 | 2.4064 | 2.3376 |

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 | | | | | Inter-assay Precision | | | | |
|-----------------------|----|--------------|------|------|-----------------------|----|--------------|------|------|
| Sample | n | Mean (ng/mL) | SD | CV% | Sample | n | Mean (ng/mL) | SD | CV% |
| 1 | 20 | 5.31 | 0.19 | 3.58 | 1 | 24 | 5.12 | 0.36 | 7.08 |
| 2 | 20 | 1.24 | 0.04 | 3.22 | 2 | 24 | 1.23 | 0.09 | 7.20 |
| 3 | 20 | 0.31 | 0.02 | 5.38 | 3 | 24 | 0.26 | 0.02 | 7.80 |

9.3 Recovery

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

| Sample Type | | Average% of Expected | Range (%) |
|--------------------------|------|----------------------|-----------|
| Human plasma | 1:8 | 102 | 91-117 |
| | 1:16 | 101 | 82-114 |
| Cell culture supernatant | 1:2 | 103 | 88-115 |
| | 1:4 | 102 | 89-115 |

9.4 Sample values

Human plasma - human plasma sample was evaluated for the presence of human N-cadherin in this assay.

| Sample Type | Mean (ng/mL) | Range (ng/mL) |
|---------------------|--------------|---------------|
| Human plasma (n=16) | 3.85 | 0.27-6.12 |

Cell culture supernatant - Hela cell were grown in DMEM with 10% fetal bovine serum. Aliquots of the cell culture supernatants were removed and assayed for levels of human N-cadherin, measured less than the lowest standard 0.156 ng/mL.

9.5 Sensitivity

The minimum detectable dose of human N-cadherin is 0.02 ng/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, human plasma samples were diluted with the appropriate **Sample Diluent** to produce samples with values within the dynamic range of the assay. Cell culture supernatant samples were spiked with high concentrations of human N-cadherin in various matrices and diluted with the appropriate **Sample Diluent** to produce samples with values within the dynamic range of the assay.

| | | Human plasma (Sample Diluent PT 3) | Cell culture supernatant (Sample Diluent PT 1) |
|------|----------------------|------------------------------------|--|
| 1:2 | Average% of Expected | 100 | 109 |
| | Range (%) | - | 105-117 |
| 1:4 | Average% of Expected | 110 | 111 |
| | Range (%) | 105-116 | 102-125 |
| 1:8 | Average% of Expected | 105 | 110 |
| | Range (%) | 94-117 | 104-120 |
| 1:16 | Average% of Expected | 111 | 111 |
| | Range (%) | 94-122 | 108-115 |

9.7 Specificity

This assay recognizes natural and recombinant human N-cadherin.

The following factors prepared at 50 ng/mL were assayed and exhibited no cross-reactivity or interference.

human:

NCAM-1/CD56

FGF R4/Fc Chimera

10. References

1. Soler AP, et al. N-cadherin involvement in cardiac myocyte interaction and myofibrillogenesis. *Dev Biol.* 162(1):9-17 (1994).
2. Navarro P, et al. Differential localization of VE- and N-cadherins in human endothelial cells: VE-cadherin competes with N-cadherin for junctional localization. *J Cell Biol.* 140(6):1475-84 (1998).
3. Puch S, et al. N-cadherin is developmentally regulated and functionally involved in early hematopoietic cell differentiation. *J Cell Sci.* 114(Pt 8):1567-77 (2001).
4. Derycke Let al. Soluble N-cadherin in human biological fluids. *Int J Cancer.* 119(12):2895-900 (2006).
5. Moya PR, et al. Rare missense neuronal cadherin gene (CDH2) variants in specific obsessive-compulsive disorder and Tourette disorder phenotypes. *Eur J Hum Genet.* 21(8):850-4 (2013).