Human Neudesin ELISA Kit

Instructions for Use | Cat. No.: ARE6652

I. INTRODUCTION

This gene encodes a neurotrophic factor that may play a role in neuron differentiation and development. A pseudogene of this gene is found on chromosome 12. Alternate splicing of this gene results in multiple transcript variants. It is mapped to 1q32.3.

II. ASSAY PRINCIPLES

AREX Biosciences Human Neudesin ELISA (Enzyme-Linked Immunosorbent Assay) kit is an in vitro enzyme-linked immunosorbent assay for the quantitative measurement of Human Neudesin in Cell Culture Supernates, Serum, Plasma. This assay employs an antibody specific for Human Neudesin coated on a 96-well plate. Standards and samples are pipetted into the wells and Neudesin present in a sample is bound to the wells by the immobilized antibody. The wells are washed and biotinylated anti-Human Neudesin antibody is added. After washing away unbound biotinylated antibody, HRP-conjugated streptavidin is pipetted to the wells. The wells are again washed, a TMB substrate solution is added to the wells and color develops in proportion to the amount of Neudesin bound. The Stop Solution changes the color from blue to yellow, and the intensity of the color is measured at 450 nm.

III. KIT COMPONENTS

Component	Volume
96-well Plate Coated with Anti-Human Neudesin Antibody	12 x 8 Strips
Human Neudesin Standard	10 ng x 2
Biotin-Labeled Detection Antibody (100X)	120 µl
Streptavidin-HRP (100X)	120 µl
Standard/Sample Diluent	30 ml
Detection Antibody Diluent	12 ml
Streptavidin-HRP Diluent	12 ml
Wash Buffer (20X)	30 ml
TMB Substrate Solution	12 ml
Stop Solution	12 ml
Plate Adhesive Strips	3 Strips
Technical Manual	1 Manual

IV. STORAGE AND STABILITY

All kit components are stable at 2 to 8 °C. Standard (recombinant protein) should be stored at -20 °C or -80 °C (recommended at -80 °C) after reconstitution. Opened Microplate Wells or reagents may be store for up to 1 month at 2 to 8 °C. Return unused wells to the pouch containing desiccant pack, reseal along entire edge.

Note: the kit can be used within one year if the whole kit is stored at -20 °C. Avoid repeated freeze-thaw cycles.

V. MATERIALS REQUIRED BUT NOT PROVIDED

1. Microplate reader capable of measuring absorbance at 450 nm.
2. Adjustable pipettes and pipette tips to deliver 2 µl to 1 ml volumes.
3. Adjustable 1-25 ml pipettes for reagent preparation.
4. 100 ml and 1 liter graduated cylinders.
5. Absorbent paper.
6. Distilled or deionized water.
7. Computer and software for ELISA data analysis.
8. Tubes to prepare standard or sample dilutions.

VI. HEALTH AND SAFETY PRECAUTIONS

1. Reagents provided in this kit may be harmful if ingested, inhaled or absorbed through the skin. Please carefully review the MSDS for each reagent before conducting the experiment.
2. Stop Solution contains 2 N Sulfuric Acid (H2SO4) and is an extremely corrosive agent. Please wear proper eye, hand and face protection when handling this material. When the experiment is finished, be sure to rinse the plate with copious amounts of running water to dilute the Stop Solution prior to disposing the plate.

VII. REAGENT PREPARATION

1. Sample Preparation

Store samples to be assayed within 24 hours at 2-8°C. For long-term storage, aliquot and freeze samples at -20°C. Avoid repeated freeze-thaw cycles.

Cell culture supernates: Remove particulates by centrifugation, assay immediately or aliquot and store samples at -20°C.

Serum: Allow the serum to clot in a serum separator tube (about 4 hours) at room temperature. Centrifuge at approximately 1000 X g for 15 minutes. Analyze the serum immediately or aliquot and store samples at -20°C.

Plasma: Collect plasma using heparin or EDTA as an anticoagulant. Centrifuge for 15 minutes at 1500 X g within 30 minutes of collection. Assay immediately or aliquot and store samples at -20°C.

Cell Lysates: Collect cells and rinse cells with PBS. Homogenize and lyse cells throughly in lysate solution. Centrifuge cell lysates at approximately 10000 X g for 5 minutes to remove debris. Aliquots of the cell lysates were removed and assayed.

Bone Tissue: Extract demineralized bone samples in 4 M Guanidine-HCl and protease inhibitors. Dissolve the final sample in 2 M Guanidine-HCl.

Tissue Homogenates: The preparation of tissue homogenates will vary depending upon tissue type. Rinse tissue with 1X PBS to remove excess blood, homogenized in 20 mL of 1X PBS and stored overnight at ⩽ -20 °C. After two freeze-thaw cycles were performed to break the cell membranes, the homogenates were centrifuged for 5 minutes at 5000 x g. The supernate was removed immediately and assayed. Alternatively, aliquot and store samples at ⩽ -20 °C.

Note: Some lysis buffer, such as RIPA can not be used. Some components will affect the binding.

Urine: Urinary samples should be cleared by centrifugation and then can be used directly without dilution. Storage at -20°C.

2. Human Neudesin Standard Preparation

Reconstitute the lyophilized Human Neudesin Standard by adding 1 ml of Standard/Sample Diluent to make the 10000 pg/ml standard stock solution. Allow solution to sit at room temperature for 5 minutes, then gently vortex to mix completely. Use within one hour of reconstituting. Two tubes of the standard (10 ng per tube) are included in each kit. Use one tube for each experiment.

Perform 2-fold serial dilutions of the top standards to make the standard curve within the range of this assay (156 pg/ml - 10000 pg/ml) as below. Standard/Sample Dilution Buffer serves as the zero standard (0 pg/ml).

Standard	Add	Into
10000 pg/ml	-	-
5000 pg/ml	500 µl of the Standard (10000 pg/ml)	500 µl of the Standard/Sample Diluent
2500 pg/ml	500 µl of the Standard (5000 pg/ml)	500 µl of the Standard/Sample Diluent
1250 pg/ml	500 µl of the Standard (2500 pg/ml)	500 µl of the Standard/Sample Diluent
625 pg/ml	500 µl of the Standard (1250 pg/ml)	500 µl of the Standard/Sample Diluent
312 pg/ml	500 µl of the Standard (625 pg/ml)	500 µl of the Standard/Sample Diluent
156 pg/ml	500 µl of the Standard (312 pg/ml)	500 µl of the Standard/Sample Diluent
0 pg/ml	1 ml of the Standard/Sample Diluent	-

Note: The standard solutions are best used within 2 hours. The 10000 pg/ml standard solution should be stored at 4°C for up to 12 hours, or at -20°C for up to 48 hours. Avoid repeated freeze-thaw cycles.

3. Biotin-Labeled Detection Antibody Working Solution Preparation

The Biotin-Labeled Detection Antibody should be diluted in 1:100 with the Detection Antibody Diluent and mixed thoroughly. The solution should be prepared no more than 2 hours prior to the experiment.

4. Streptavidin-HRP Working Solution Preparation

The Streptavidin-HRP should be diluted in 1:100 with the Streptavidin-HRP Diluent and mixed thoroughly. The solution should be prepared no more than 1 hour prior to the experiment.

5. Wash Buffer Working Solution Preparation

Pour entire contents (30 ml) of the Wash Buffer Concentrate into a clean 1,000 ml graduated cylinder. Bring final volume to 600 ml with glass-distilled or deionized water (1:20).

VIII. ASSAY PROCEDURE

The Streptavidin-HRP Working Solution and TMB Substrate Solution must be kept warm at 37°C for 30 minutes before use. When diluting samples and reagents, they must be mixed completely and evenly. Standard detection curve should be prepared for each experiment. The user will decide sample dilution fold by crude estimation of protein amount in samples.

1. Add 100 µl of each standard and sample into appropriate wells.

2. Cover well and incubate for 90 minutes at room temperature or overnight at 4°C with gentle shaking.

3. Remove the cover, discard the solution and wash plate 3 times with Wash Buffer Working Solution, and each time let Wash Buffer Working Solution stay in the wells for 1 - 2 minutes. Blot the plate onto paper towels or other absorbent material. Do NOT let the wells completely dry at any time.

4. Add 100 µl of Biotin-Labeled Detection Antibody Working Solution into each well and incubate the plate at 37°C for 60 minutes.

5. Wash plate 3 times with Wash Buffer Working Solution, and each time let Wash Buffer Working Solution stay in the wells for 1 - 2 minutes. Discard the Wash Buffer Working Solution and blot the plate onto paper towels or other absorbent material.

6. Add 100 µl of Streptavidin-HRP Working Solution into each well and incubate the plate at 37°C for 45 minutes.

7. Wash plate 5 times with Wash Buffer Working Solution, and each time let wash buffer stay in the wells for 1 - 2 minutes. Discard the wash buffer and blot the plate onto paper towels or other absorbent material.

8. Add 100 µl of TMB Substrate Solution into each well and incubate plate at 37°C in dark for 30 minutes.

9. Add 100 µl of Stop Solution into each well. The color changes into yellow immediately.

10. Read the O.D. absorbance at 450nm in a microplate reader within 30 minutes after adding the Stop Solution.

For calculation, (the relative O.D.450) = (the O.D.450 of each well) - (the O.D.450 of Zero well). The standard curve can be plotted as the relative O.D.450 of each standard solution (Y) vs. the respective concentration of the standard solution (X). The concentration of the samples can be interpolated from the standard curve.

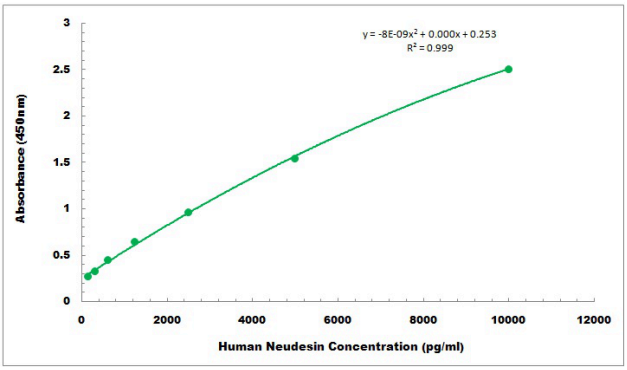
Note: If the samples measured were diluted, multiply the dilution factor to the concentrations from interpolation to obtain the concentration before dilution.

IX. ASSAY PROCEDURE SUMMARY

1. Prepare all reagents, samples and standards
2. Add 100 µl Standard or Sample
3. Wash plate 3 times with Wash Buffer Working Solution
4. Add 100 µl Biotin-Labeled Detection Antibody Working Solution
5. Wash plate 3 times with Wash Buffer Working Solution
6. Add 100 µl Streptavidin-HRP Working Solution
7. Wash plate 5 times with Wash Buffer Working Solution
8. Add 100 µl TMB Substrate Solution
9. Add 100 µl Stop Solution
10. Read the plate at 450nm

X. TYPICAL DATA

The standard curve is for demonstration only. A standard curve must be run with each assay.



XI. SENSITIVITY

The minimum detectable dose of Human Neudesin is typically less than 58 pg/ml.

XII. SPECIFICITY

Human Neudesin ELISA Kit allows for the detection and quantification of endogenous levels of natural and/or recombinant Human Neudesin proteins within the range of 156 pg/ml - 10000 pg/ml.

XIII. CROSS REACTIVITY

No detectable cross-reactivity with other relevant proteins.

XIV. TROUBLESHOOTING GUIDE

Problem	Possible Cause	Solution
High signal and background in all wells	Insufficient washing Too much Streptavidin-HRP Incubation time too long Development time too long	Increase number of washes Increase time of soaking between in wash Check dilution, titration Reduce incubation time Decrease the incubation time before the stop solution is added
No signal	Reagent added in incorrect order, or incorrectly prepared Standard has gone bad (If there is a signal in the sample wells) Assay was conducted from an incorrect starting point	Review protocol Check the condition of stored standard Reagents allows to come to 20 - 30 °C before performing assay
Too much signal-whole plate turned uniformly blue	Insufficient washing-unbound Streptavidin-HRP remaining Too much Streptavidin-HRP Plate sealer or reservoir reused, resulting in presence of residual Streptavidin-HRP	Increase number of washes Carefully Check dilution Use fresh plate sealer and reagent reservoir for each step
Standard curve achieved but poor discrimination between point	Plate not developed long enough Improper calculation of standard curve dilution	Increase substrate solution incubation time Check dilution, make new standard curve
No signal when a signal is expected, but standard curve looks fine	Sample matrix is masking detection	More diluted sample Recommended
Samples are reading too high, but standard curve is fine	Samples contain protein levels above assay range	Dilute samples and run Again
Edge effect	Uneven temperature around work surface	Avoid incubating plate in areas where environmental conditions vary Use plate sealer

XV. TECHNICAL SUPPORT

For troubleshooting, information or assistance, please contact us at info@arexbio.com.

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XVI. NOTES

For research use only. Not for diagnostic or therapeutic procedures.