

# Catalog NO.: FY-EH3062S EasyPro Human NFE2L2(Nuclear Factor, Erythroid Derived 2, Like 2) ELISA Kit

For the quantitative determination of Human NFE2L2 concentrations in serum, plasma, and other biological fluids.

This package insert must be read in its entirety before using this product.

For research use only. Not for use in diagnostic procedures.

#### PRINCIPLE OF THE ASSAY

This ELISA kit uses the Sandwich-ELISA principle. The micro ELISA plate provided in this kit has been pre-coated with an antibody specific to Human NFE2L2. Samples (or Standards) and and Horseradish Peroxidase (HRP) linked antibody specific for Human NFE2L2 are added to the micro ELISA plate wells. Human NFE2L2 in samples (or standards) combines with the coated antibody and HRP linked detection antibody special to Human NFE2L2. Excess conjugate and unbound sample or standard are washed from the plate. The enzyme-substrate reaction is terminated by the addition of stop solution and the color turns yellow. The optical density (OD) is measured spectrophotometrically at a wavelength of 450 nm. The OD value is proportional to the concentration of Human NFE2L2. You can calculate the concentration of Human NFE2L2 in the samples by comparing the OD of the samples to the standard curve.

#### PERFORMANCE

- ★ Sensitivity:0.1 ng/mL.
- ★ Detection Range:0.16-10ng/mL.
- ★ No significant cross-reactivity or interference between Human NFE2L2 and analogues was observed.
- ★ Repeatability: Coefficient of variation is 10%.

## PRECAUTIONS BEFORE THE EXPERIMENT

- 1. The detection range of the kit is different from the concentration range of the substance to be measured in the sample. It is recommended to estimate the concentration by looking up the reference before the experiment, and determine the actual concentration by pre-experiment. If the concentration is too high or too low, dilute or concentrate the sample appropriately.
- 2. The newly opened ELISA plate may contain water fog-like substances, which is a normal phenomenon and will not have any impact on the experimental results.
- 3. Standard working solution and HRP Conjugate working solution should not be reused.
- 4. The micro plate reader requires a filter with a wavelength of 450±3nm and a detector capable of detecting 450±3nm, and the optical density value is between 0 and 3.5.
- 5. Avoid cross-contamination, change the pipette head between each standard concentration addition, between sample addition, and between reagent addition. In addition, separate storage bottles should be used for each reagent.
- 6. In order to obtain accurate experimental results, it is necessary to ensure that the sealing plate membrane seals the enzyme label plate during incubation.
- 7. The substrate solution should remain colorless and free from light until it is added to the plate. When added to the plate, the substrate solution should change from colorless to a gradient of blue.
- 8. The stop solution should be added to the Micro ELISA Plate in the same order as the substrate solution. After the stop solution is added, the color of the solution in the well will change from blue to yellow. Green wells indicate that the stop solution is not sufficiently mixed with the substrate solution. If the color is uneven, shake the plate gently to mix the solution evenly.

## REAGENTS&MATERIALS PROVIDED

Item	Specifications	Storage
Micro ELISA Plate (Dismountable)	96T: 8 wells ×12 strips	-20°C, 6 months
	48T: 8 wells ×6 strips	
	24T: 8 wells ×3 strips	
Reference Standard	96T: 2 vials	
	48T: 1 vial	
	24T: 1 vial	
Concentrated HRP Conjugate (100×)	96T: 1 vial, 60μL	-20°C, 6 months
	48T: 1 vial, 30μL	
	24T: 1 vial, 30μL	
Reference Standard & Sample	1:-1 20 I	
Diluent	1 vial, 20 mL	
HRP Conjugate Diluent	1 vial, 13mL	2-8°C, 6 months
Concentrated Wash Buffer (25×)	1 vial, 30 mL	
Substrate Reagent	1 vial, 10 mL	2-8°C (Protect from light)
Stop Solution	1 vial, 10 mL	2-8°C, 6 months
Plate Sealer	5 pieces	
Product Description	1 copy	RT
Certificate of Analysis	1 copy	

## OTHER SUPPLIES REQUIRED

Microplate reader with 450nm wavelength filter
Incubator capable of maintaining 37°C
Single or multi-channel pipettes with high precision
disposable pipette tips
EP tubes
Container for Wash Solution
Squirt bottle, manifold dispenser
Deionized or distilled water
Absorbent paper for blotting the microplate

#### SAMPLE COLLECTION&STORAGE

**Serum:** Allow blood samples to clot for 2 hours at room temperature or overnight at 2-8°C before centrifugation for 15 min at 2000×g at 2-8°C. Supernatants should be taken for assay testing immediately or stored at -20°C or -80°C for later use. Avoid repeated freeze-thaw cycles.

**Plasma:** Collect plasma using EDTA or heparin as an anticoagulant. Centrifuge samples for 15 min at 2000×g at 2-8°C within 30 min of collection. Supernatants should be taken for assay testing immediately or stored at -20°C or -80°C for later use. Avoid repeated freeze-thaw cycles.

Cell culture supernatant: Remove particulates by centrifugation and assay immediately or aliquot and store samples at  $\leq$  -20 °C. Avoid repeated freeze-thaw cycles.

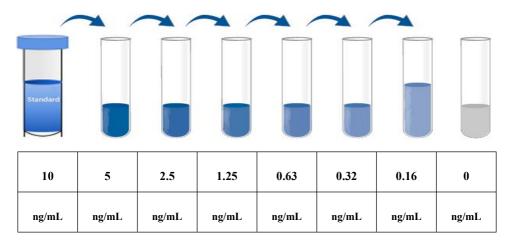
Other Types of samples: Please check our official website for the collection and processing method(https://www.feiyuebio.cn/index/e31/66.html).

## REAGENT PREPARATION

- 1. Allow all reagents to return to room temperature (18-25°C) before use.
- 2. Wash Buffer: Dilute 20 mL of Concentrated Wash Buffer with 480 mL of deionized or distilled water to prepare 500 mL of Wash Buffer.

#### 3. Standard working solution:

- ①Centrifuge the Reference Standard at 10000×g for 1 min.
- ②Add 1mL of Reference Standard & Sample Diluent, let it stand for 10min and then mix gently, that is 10ng/mL of standard working solution.
- ③Prepare 7 EP tubes, add  $500\mu$ L standard & sample diluent to each tube, take  $500\mu$ L from 10ng/mL to make 5ng/mL, and so on, the last tube is blank (no more liquid needs to be drawn from the penultimate tube). Use it now!



4.HRP Conjugate working solution: Calculate the amount of HRP Conjugate required (50 $\mu$ L/ well). The actual preparation should be 100-200 $\mu$ L. Centrifuge the concentration tube before use and dilute 100× concentrated HRP Conjugate to 1× working solution (e.g.  $10\mu$ L + 990 $\mu$ L HRP Conjugate diluent).

#### ASSAY PROTOCOL

- 1. The Micro ELISA Plate slats to be used were removed from the plate frame and the remaining slats were returned to the aluminum foil bag containing the desiccants and then resealed for storage.
- 2. Determine wells for diluted standard, blank and sample. Add 50μL each dilution of standard, blank and sample into the appropriate wells (It is recommended that all samples and standards be assayed in duplicate). Immediately add 50μL of HRP Conjugate working solution to each well. Cover the plate with a new sealer. Incubate for 60 min at 37°C. Note: solutions should be added to the bottom of the micro ELISA plate well, avoid touching the inside wall and causing foaming as much as possible.
- 3. Decant the solution from each well add 300µL of wash buffer to each well. Soak for 0.5 min and aspirate or decant the solution from each well and pat it dry against clean absorbent paper. Repeat this wash step 3 times. Note: a microplate washer can be used in this step and other wash steps. Make the tested strips in use immediately after the wash step. Do not allow wells to be dry.
- 4. Add 100μL of Substrate Reagent to each well. Cover the plate with a new sealer. Incubate for about 15 min at 37°C. Protect the plate from light. Note: the reaction time can be shortened or extended according to the actual color change, but not more than 30 min. Preheat the Microplate Reader for about 15 min before OD measurement.
- 5. Add 50μL of Stop Solution to each well. Note: adding the stop solution should be done in the same order as the substrate solution.
- 6. Determine the optical density (OD value) of each well at once with a micro plate reader set to 450 nm.

## **SUMMARY FOR PROCEDURE**

Respectively add standard and sample 50µL/ well

Add 50µL of HRP Conjugate 50µL/ well and incubated at 37°C for 60 min

Wash the plate three times

Add substrate 100µL/ well and incubate at 37°C for 15 minutes (avoid light and not exceed 30 minutes)

Add stop solution 50µL/ well

#### **EXAMPLE DATA**

For each test, a standard curve must be set for each plate. The standard curves below are for example purposes only.

ng/mL	Corrected OD	Typtical Standard Curve
0	0	
0.16	0.042	10
0.32	0.089	
0.63	0.213	OD 450nm
1.25	0.451	0.1
2.5	0.958	0.01
5	1.687	0.1 1 10 100 Human NFE2L2 Concentration(ng/mL)
10	2.362	

## **CALCULATION OF RESULTS**

Calculate the average OD value of the standard and sample, then subtract the OD value of the zero concentration standard.

With the standard product concentration as the horizontal coordinate and OD value as the vertical coordinate, the standard curve is formed by using the nonlinear four parameter logistic(4-PL) and the sample concentration is calculated by the curve.Please refer to our official website for the specific fitting method(https://www.feiyuebio.cn/index/e31/65.html).

If samples have been diluted, the concentration read from the standard curve must be multiplied by the dilution factor.

#### **SENSITIVITY**

The minimum detection concentration of the Human NFE2L2 detected by this kit is usually less than the 0.1 ng/mL. The sensitivity was determined by adding two standard deviations to the mean OD value of twenty zero standard replicates and calculating the corresponding concentration.

#### **PRECISION**

Mean coefficient of variation for Intra-Assay and Inter-Assay: 3 samples with low, middle and high level concentration were tested for repeat multiple times, respectively. The results showed that the coefficient of variation of the kits was less than 10%, which met the precision quality control standard.

#### **SPECIFICITY**

This method has high sensitivity and specificity for the detection of Human NFE2L2. No significant cross-reactivity was observed between the Human NFE2L2 and its analogues.

## **RECOVERY**

The known concentration of Human NFE2L2 was added to different samples respectively, and the recovery experiment was conducted. The results showed that the recovery range and average recovery rate of the kit were 80-120%, which met the recovery quality control standard.

## **LINEARITY**

Dilute linear experiments were performed on samples with Human NFE2L2 to evaluate the linearity of the kit. The results showed that the kit linear range (%) was 80-120%, which met the linear quality control standard.

#### **DECLARATION**

- 1. Limited by current conditions and scientific technology, we can't conduct comprehensive identification and analysis of all the raw material provided. There might be some qualitative and technical risks for users using the kit.
- 2. This assay is designed to eliminate interference by factors present in biological samples. Until all factors have been tested in the ELISA immunoassay, the possibility of interference cannot be excluded.
- 3. The final experimental results will be closely related to the validity of products, operational skills of the operators, the experimental environments and so on We are only responsible for the kit itself, but not for the samples consumed during the assay. The users should calculate the possible amount of the samples used in the whole test. Please reserve sufficient samples in advance.
- 4. To get the best results, please only use the reagents supplied by the manufacturer and strictly comply with the instructions.
- 5. Incorrect results may occur because of incorrect operations during the reagents preparation and loading, as well as incorrect parameter settings of the Micro plate reader. Please read the instructions carefully and adjust the instrument prior to the experiment.
- 6. Even the same operator might get different results in two separate experiments. In order to get reproducible results, the operation of every step in the assay should be controlled.
- 7. Every kit has strictly passed QC test. However, results from end users might be inconsistent with our data due to some variables such as transportation conditions, different lab equipment, and so on. Intra assay variance among kits from different batches might arise from the above reasons too.
- 8. Kits from different manufacturers or other methods for testing the same analyte could bring out inconsistent results, since we haven't compared our products with those from other manufacturers.
- 9. The kit is designed for research use only, we will not be responsible for any issues if the kit is applied in clinical diagnosis or any other related procedures.

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## **PLATE LAYOUT**

Use this plate layout to record standards and samples assayed.



If you have any questions, please contact us.

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