

Coenzyme II NADP (H) Content Assay Kit (MTT Chromogenic Method)

Note: Take two or three different samples for prediction before test.

Operation Equipment: Spectrophotometer/microplate reader

Cat No: NA0170

Size:100T/48S

Components:

Acid Extract solution : Liquid 50 mL×1. Storage at 2-8°C.

Alkaline Extract solution: Liquid 50 mL×1. Storage at 2-8°C.

Reagent I : Liquid 10 mL×1. Storage at 2-8°C.

Reagent II : Powder×1. Storage at -20°C . Add 4mL of distilled water before using to dissolve it.

Reagents can be stored at 2-8°C for 4 weeks after dissolution.

Reagent III : Liquid 4 mL×1. Storage at 2-8°C.

Reagent IV: Liquid 4 mL×1. Storage at 2-8°C.

Reagent V : Liquid 25 mL×1. Storage at 2-8°C.

NADP standard: Powder×1. Storage at -20°C .Add 1.27 mL of distilled water before use to obtain a standard of 5 µmol/mL, which can be stored at -20°C for 2 weeks.

NADPH standard: Powder×1. Storage at -20°C .Add 1.2 mL of distilled water before use to obtain a standard of 5 µmol/mL, which can be stored at -20°C for 2 weeks.

Product Description:

Coenzyme II NADP(H) is widely present in animals, plants, microorganisms and cultured cells. The determination of NADP⁺ and NADPH content can calculate the content of NADP (NADPH + NADP⁺) and the ratio of NADPH/NADP⁺, and its changes are related to the pentose phosphate pathway and biosynthesis and Antioxidative responses are closely related. The NADPH/NADP⁺ ratio is not only one of the main markers of cellular redox state, but also plays an important regulatory role in the PPP pathway, biosynthesis and antioxidant metabolism.

The NADP⁺ and NADPH in the samples were extracted with acidic and basic extraction solutions, respectively. Under the action of 1-mPMS, WST-1 can react with NADPH to produce water-soluble formazan, which has a characteristic absorption peak at 450nm, while NADP⁺ can be reduced to NADPH by 6-phosphate glucose dehydrogenase, which is further detected by WST-1.

Reagents and Equipment Required but Not Provided:

Spectrophotometer/microplate reader, centrifuge, water bath, mortar/homogenizer, sonicator, adjustable pipette, micro glass cuvettes/96 well plates, ice, and distilled water.

Sample Preparation :

1. Serum

Extract NADP⁺: Take 0.1 mL of serum (slurry), add 0.5 mL of acidic extract, boil for 5 minutes (cover tightly to prevent water loss), and after cooling in an ice bath, centrifuge at 10,000g at 4°C for

10 minutes; take 200 μ L of supernatant, add an equal volume of alkaline extract ; Mix well, centrifuge at 10,000g at 4°C for 10 min, take the supernatant, and store on ice for testing.

Extract NADPH: Take 0.1 mL of serum (slurry), add 0.5 mL of alkaline extract, boil for 5 minutes (cover tightly to prevent water loss), after cooling in an ice bath, centrifuge at 10,000g at 4°C for 10 minutes, take 200 μ L of supernatant, and add an equal volume of acidic extract ; Mix well, centrifuge at 10000g at 4°C for 10min, take the supernatant and store on ice for testing.

2. Tissue

Extract NADP⁺: Weigh about 0.1g of tissue, add 0.5mL of acidic extract, grind in ice bath, boil for 5min (cover tightly to prevent water loss), after cooling in ice bath, centrifuge at 10,000g at 4°C for 10min, take 200 μ L of supernatant, add an equal volume of alkaline extract ; Mix wel, centrifuged at 10,000 g at 4°C for 10 min, and the supernatant was taken and stored on ice for testing.

Extract NADPH: Weigh about 0.1 g of tissue, add 0.5 mL of alkaline extraction solution, grind in an ice bath, boil for 5 min (cover tightly to prevent water loss), after cooling in an ice bath, centrifuge at 10,000 g at 4°C for 10 min, take 200 μ L of supernatant, and add an equal volume of acidic extract ; Mix well, centrifuged at 10,000 g at 4°C for 10 min, and the supernatant was taken and stored on ice for testing.

3. Bacteria or cells

Extract NADP⁺: Collect 5 million cells or bacteria, add 0.5mL of acidic extract, ultrasonically disrupt for 1min (intensity 20% or 200W, ultrasonic for 2s, stop for 1s), boil for 5min (cover tightly to prevent water loss), cool in ice bath, 10000g centrifuge at 4°C for 10min, take 200uL of the supernatant into another new centrifuge tube, add an equal volume of alkaline extract to neutralize, mix well, centrifuge at 10,000g at 4°C for 10min, take the supernatant and store it on ice for testing .

Extract NADPH: Collect 5 million cells or bacteria, add 0.5mL alkaline extract, ultrasonically disrupt for 1min (intensity 20% or 200W, ultrasonic for 2s, stop for 1s), boil for 5min (cover tightly to prevent water loss), cool in an ice bath, Centrifuge at 10,000g at 4°C for 10min, take 200uL of the supernatant into another new centrifuge tube, add an equal volume of acidic extract to neutralize, mix well, centrifuge at 10,000g at 4°C for 10min, take the supernatant, and store it on ice for testing .

Determination procedure:

1. Preheat the spectrophotometer /microplate reader more than 30 minutes, adjust the wavelength to 450 nm, the spectrophotometer set zero with distilled water.
2. NADP⁺ standard: diluted with distilled water to a standard solution of 5,2.5,1.25, 0.625, 0.3125, 0.15625, 0nmol/mL (0nmol/mL is a blank tube).
3. NADPH standard: diluted with distilled water to a standard solution of 10,5, 2.5, 1.25, 0.625, 0.3125, 0.15625,0nmol/mL (0nmol/mL is a blank tube).
4. Adding sample table (add samples in the 1.5mL brown EP tube in sequence according to the following table):

Reagent (μ L)	Control tube (A ₁)	Test tube (A ₂)	Standard tube
Sample/ Standard	20	20	20
Reagent V	200	-	-

Reagent I	80	80	80
Reagent II	30	30	30
Reagent III	30	30	30
Reagent IV	30	30	30
Mix well and let stand for 1h at room temperature in the dark			
Reagent V		200	200

Mix well, take 200 μL into a micro glass cuvettes / 96-well plate, measure at 450nm, read the absorbance value, NADP^+ is marked as: $\Delta A \text{ NADP}^+ = A_2 - A_1$, NADPH is marked as $\Delta A \text{ NADPH} = A_2' - A_1'$, NADP standard tube is marked as $\Delta A_s = A_s - A_b$. The NADPH standard tube is marked as $\Delta A_s' = A_s' - A_b$. (The standard curve only needs to be done 1-2 times).

Calculation:

1. Standard curve drawing:

(1) Drawing of NADP^+ standard curve: According to the concentration of the standard tube (x_1 , nmol/mL) and the absorbance ΔA_s (y_1 , ΔA_s), establish a standard curve. From the standard curve, plug ΔA into the equation to get x_1 (nmol/mL).

(2) Drawing of the NADPH standard curve: According to the concentration of the standard tube (x_2 , nmol/mL) and the absorbance $\Delta A_s'$ (y_2 , $\Delta A_s'$), establish a standard curve. From the standard curve, plug ΔA into the equation to get x_2 (nmol/mL).

2. Calculation of NADP^+ and NADPH content:

① Calculation of NADP^+ content

(1) Calculated by liquid volume:

$$\text{NADP}^+\text{content (nmol/mL)} = x_1 \times (V_e + V_{se}) \div V \text{ serum} = 11 \times x_1$$

(2) Calculated by sample protein concentration

$$\text{NADP}^+ \text{ (nmol/mg prot)} = x_1 \times V_e \div (V_e \times C_{pr}) = x_1 \div C_{pr}$$

(3) Calculate content according to the fresh weight of the sample

$$\text{NADP}^+ \text{ (nmol/g fresh weight)} = x_1 \times V_e \div W = x_1 \div W$$

(4) Calculated by the number of cells:

$$\text{NADP}^+ \text{ content (nmol/10}^4 \text{ cell)} = x_1 \times V_e \div 500 = 0.002 \times x_1$$

② Calculation of NADPH content

(1) Calculated by liquid volume:

$$\text{NADPH content (nmol/mL)} = x_2 \times (V_e + V_s) \div V_s = 11 \times x_2$$

(2) Calculated by sample protein concentration

$$\text{NADPH (nmol/mg prot)} = x_2 \times V_e \div (V_e \times C_{pr}) = x_2 \div C_{pr}$$

(3) Calculate the content according to the fresh weight of the sample

$$\text{NADPH (nmol/g fresh weight)} = x_2 \times V_e \div W = x_2 \div W$$

(4) Calculated the content by the number of cells:

$$\text{NADPH (nmol/10}^4 \text{ cell)} = x_2 \times V_e \div 500 = 0.002 \times x_2$$

V_e : volume of added extract, 1 mL;

V_{se} : volume of serum (plasma), 0.1 mL;

C_{pr} : sample protein concentration, mg/mL;

W: sample mass, g;

500: the total number of bacteria or cells, 5 million.

Note:

1. If the number of samples for one-time determination is large, reagents I, II and III can be prepared into a mixed solution in proportion.
2. Avoid light during the reaction.
3. Since each measuring tube needs to set up a control tube, 100 tubes of this kit can measure 48 NADP⁺ or NADPH.
4. If the measured absorbance value exceeds the linear range absorbance value, you can increase the sample volume or dilute the sample before measuring. Simultaneously modify the calculation formula.

Experimental example:

1. Determination of NADP⁺: Weigh 0.1g, extract according to the extraction steps and then operate according to the determination steps. After measuring the absorbance value in the glass cuvette, calculate $\Delta A = A_2 - A_1$, $\Delta A = A_2 - A_1 = 0.075 - 0.069 = 0.006$, standard curve $y_1 = 0.2081x - 0.0144$, according to the standard curve, $x_1 = 0.098$, and the NADP⁺ content is:
NADP⁺ (nmol/g mass) = $x_1 \div W = 0.98$ nmol/g mass.
Determination of NADPH: Weigh 0.1g of holly leaves, extract according to the extraction steps, and then operate according to the determination steps. After measuring the absorbance value in a glass cuvette, calculate $\Delta A = A_2 - A_1$, $\Delta A = A_2 - A_1 = 0.155 - 0.110 = 0.045$, standard curve $y_2 = 0.0478x - 0.0188$, according to the standard curve, $x_2 = 1.335$, NADPH content:
NADPH (nmol/g mass) = $x_2 \div W = 13.35$ nmol/g mass.
2. Determination of NADP⁺: Weigh 0.1g of mouse liver, extract it according to the extraction steps, and then operate according to the determination steps. After measuring the absorbance value in a glass cuvette, calculate $\Delta A = A_2 - A_1$, $\Delta A = A_2 - A_1 = 0.194 - 0.146 = 0.048$, standard The curve $y_1 = 0.2081x - 0.0144$, according to the standard curve, $x_1 = 0.3$, NADP⁺ content:
NADP⁺ (nmol/g mass) = $x_1 \div W = 2.999$ nmol/g mass.
Determination of NADPH: Weigh 0.1g of mouse liver, extract it according to the extraction steps, and then operate according to the determination steps. After measuring the absorbance value in a glass cuvette, calculate $\Delta A = A_2 - A_1$, $\Delta A = A_2 - A_1 = 0.170 - 0.141 = 0.029$, standard curve $y_2 = 0.0478x - 0.0188$, according to the standard curve $x_2 = 1$, NADPH content:
NADPH (nmol/g mass) = $x_2 \div W = 10$ nmol/g mass.
3. Determination of NADP⁺: Take 0.1 mL of bovine serum, extract it according to the extraction steps, and operate according to the determination steps. After measuring the absorbance value in the glass cuvette, calculate $\Delta A = A_2 - A_1$, $\Delta A = A_2 - A_1 = 0.077 - 0.067 = 0.010$, standard curve $y_1 = 0.2081x - 0.0144$, according to the standard curve $x_1 = 0.117$, NADP⁺ content:
NADP⁺ (nmol/mL) = $11 \times x_1 = 1.29$ nmol/mL.
Determination of NADPH: Take 0.1 mL of bovine serum, extract it according to the extraction steps, and then operate according to the determination steps. After measuring the absorbance value in the glass cuvette, calculate $\Delta A = A_2 - A_1$, $\Delta A = A_2 - A_1 = 0.084 - 0.072 = 0.012$, standard curve $y_2 = 0.0478x - 0.0188$, according to the standard curve $x_2 = 0.644$, NADPH content: NADPH (nmol/mL) = $11 \times x_2 = 7.088$ nmol/mL.