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Elite™ NADH Assay Kit (Red Fluorescence)

CATALOG NUMBER: CA-N205, 400 assays

Description

Nicotinamide adenine dinucleotide (NAD+) and nicotinamide adenine dinucleotide phosphate (NADP+) are two important cofactors found in cells. NADH is the reduced form of NAD+, the oxidized form of NADH. NAD forms NADP with the addition of a phosphate group to the 2' position of the adenyl nucleotide through an ester linkage.

The traditional NAD/NADH and NADP/NADPH assays are done by monitoring the changes in NADH or NADPH absorption at 340 nm. The short UV wavelength of the traditional NAD/NADH and NADP/NADPH assays makes these methods to suffer low sensitivity and high interference. Due to the weak absorption of NAD and NADH, the UV absorption method requires large sample sizes, making the same NAD and NADH measurement unpractical if the availability of samples is limited.

NADH + Elite™ NADH Sensor → **NAD + Red Fluorescent Product**

This Elite™ Fluorimetric NADH Assay Kit provides a convenient method for the detection of NADH. The enzymes in the system specifically recognize NADH in an enzyme recycling reaction. In addition, this assay has very low background since it is run in the red visible range that significantly reduces the interference resulted from biological samples. The Elite™ Fluorimetric NADH Assay Kit provides a sensitive, one-step assay to detect as little as 100 pico-moles of NADH in a 100 µl assay volume (1 µM). The assay can be performed in a convenient 96-well or 384-well microtiter-plate format. Its signal can be easily read by either a fluorescence microplate reader at Ex/Em = 540/590 nm or an absorbance microplate reader at ~576 nm.

Features

- **Broad application**: NADH detection in solution or cell extracts.
- **Continuous**: Easily adapted to automation without a separation step.
- **Convenient**: Formulated to have minimal hands-on time. No wash required.
- **Non-radioactive**: No special requirement for waste disposal.

Kit Components

• **Component A:** NADH Recycling Enzyme Mixture	2 bottles (lyophilized powder)
• **Component B:** NADH Assay Buffer	1 bottle (20 ml)
• **Component C:** NADH Standard	1 vial (142 µg)
• **Component G**: Lysis Buffer	1 bottle (10 ml)

Storage

Keep in freezer (-20 °C) and avoid exposure to light.
All reagents are stable for at least 6 months after receipt when stored properly at the recommended conditions.

Materials Required (but not supplied)

- 96-well black microplates with black bottom
- Fluorescence microplate reader

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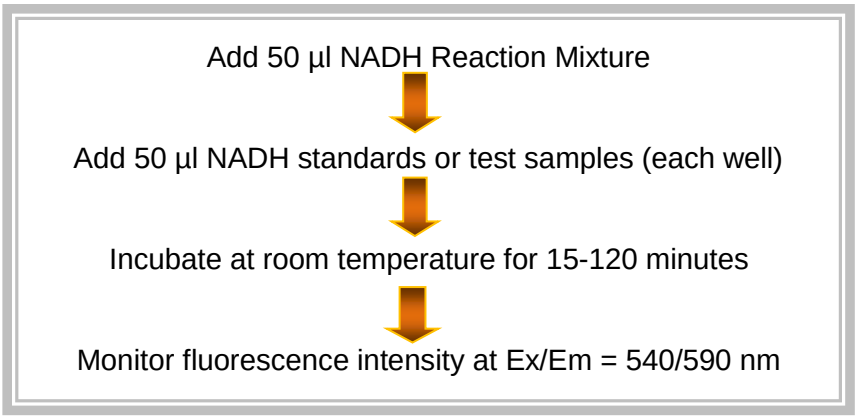
Assay Protocol (for 96-Well Plate)

Summary of the Assay



1. Prepare NADH stock solution:

Add 200 µl of PBS buffer into the vial of NADH standard (**Component C**) to have 1 mM (1 nmol/ µl) NADH stock solution.
Note: The unused NADH stock solution should be divided into single use aliquots and stored at -20 °C.

2. Prepare NADH reaction mixture:

Add 10 ml of Elite™ NADH Assay Buffer (**Component B**) to the bottle of NADH Recycling Enzyme Mixture (**Component A**), and mix well.

Note: This NADH reaction mixture is enough for two 96-well plates or four 384-well plates. The unused NADH reaction mixture should be divided into single use aliquots and stored at -20 °C.

3. Prepare serially diluted NADH standards (0 to 100 µM):

3.1. Add 50 µl of 1 mM NADH stock solution (from Step 1) into 450 µl PBS buffer (pH 7.4) to generate 100 µM (100 pmols/ µl) NADH standard solution.

Note: Diluted NADH standard solution is unstable, and should be used within 4 hours.

3.2. Take 200 µl of 100 µM NADH standard solution to perform 1:3 serial dilutions to get 30, 10, 3, 1, 0.3, 0.1, and 0 µM serially dilutions of NADH standard.

3.3. Add serial dilutions of NADH standards and NADH containing test samples into a solid black 96-well microplate as described in Tables 1 and 2.

Note: Prepare cells or tissue samples as desired or as described in appendix A. Lysis Buffer (**Component G**) can be used for lysing the cells.

Table 1. Layout of NADH standards and test samples in a solid black 96-well microplate

BL	BL	TS	TS	…	…						
NS1	NS1	…	…								
NS2	NS2										
NS3	NS3										
NS4	NS4										
NS5	NS5										
NS6	NS6										
NS7	NS7										

Note: NS= NADH Standards; BL=Blank Control; TS=Test Samples.

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Table 2. Reagent composition for each well

NADH Standard	Blank Control	Test Sample
Serial Dilutions: 50 µl	PBS: 50 µl	50 µl

Note: Add the serially diluted NADH standards from 0.1 µM to 100 µM into wells from NS1 to NS7 in duplicate.

4. Run NADH assay in supernatants reaction:

4.1. Add 50 µl of NADH reaction mixture (from Step 2) into each well of NADH standard, blank control, and test samples (from Step 3.3) to make the total NADH assay volume of 100 µl/well.

4.2. Incubate the reaction at room temperature for 15 minutes to 2 hours, protected from light.

4.3. Monitor the fluorescence increase with a fluorescence plate reader at Ex/Em = 530-570/590-600 nm (optimal at Ex/Em=540/590 nm, cutoff=570 nm).

Note: The contents of the plate can also be transferred to a white clear bottom plate and read by an absorbance microplate reader at the wavelength of 576 ± 5 nm. The absorption detection has lower sensitivity compared to fluorescence reading.

Data Analysis

The fluorescence in blank wells (with the PBS buffer only) is used as a control, and is subtracted from the values for those wells with the NADH reactions. A NADH standard curve is shown in Figure 1.

Note: The fluorescence background increases with time, thus it is important to subtract the fluorescence intensity value of the blank wells for each data point.

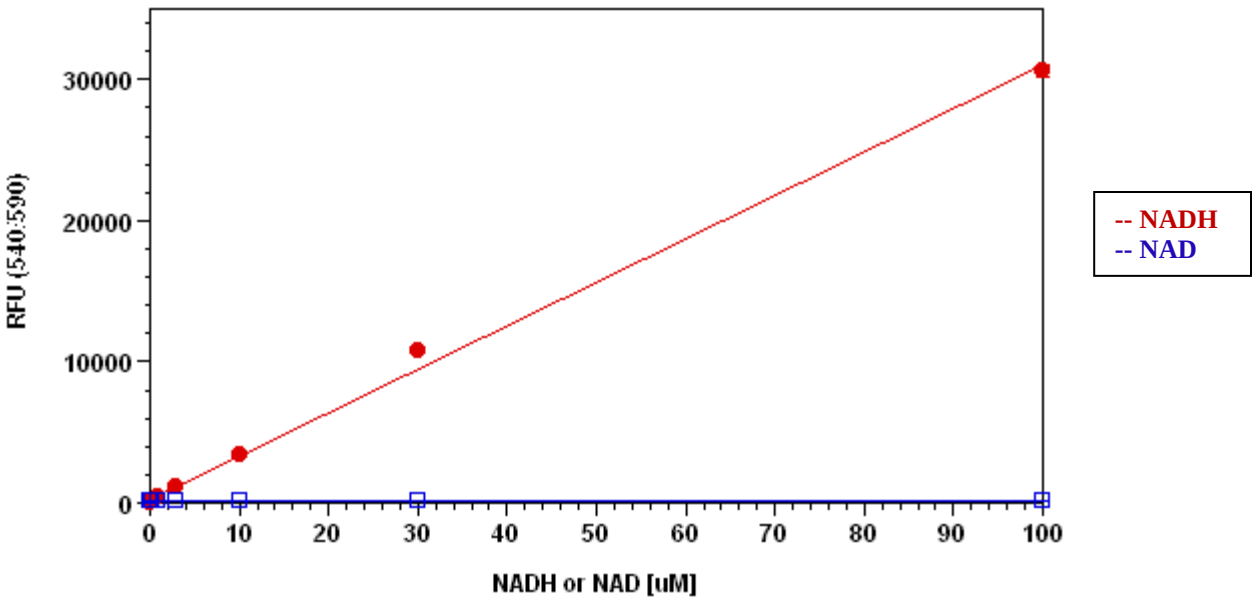

Figure 1. NADH dose response was measured with Elite™ NADH Assay Kit in a 96-well black plate using a NOVOStar microplate reader (BMG Labtech). As low as 1 µM (10 nmol/well) of NADH can be detected with 1 hour incubation (n=3) while there is no response from NAD.

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Appendix A

Sample Lysis Preparation for NADH/NAD or NADPH/NADP Assay

1. Lysis of bacterial cells

   Collecting bacterial cells by centrifugation ((10,000 x g, 0°C, 15 min). Use about 100 to 10 million cells/mL lysis buffer, leave at room temperature for 15 minutes. And then centrifuge at 2500 rpm for 5 minutes, use the supernatant for the assay.

2. Lysis of plant cells

   Homogenize the leave with the lysis buffer at 200 mg/mL, and then centrifuge at 2500 rpm for 5-10 minutes, use the supernatant for the assay.

3. Lysis of mammalian cells

   Simply remove medium from the plates (wells), use about 100 uL lysis buffer per 1-5 million cells (or 100uL/ well in a 96-well cell culture plate), and leave at room temperature for 15 minutes. You can use the cell lysate directly or simply centrifuge at 1500 rpm for 5 minutes, use the supernatant for the assay.

4. Lysis of tissues

   Weight ~20 mg tissue, wash with cold PBS, homogenize with 400 µl of lysis buffer in a micro-centrifuge tube, and then centrifuge at 2500 rpm for 5-10 minutes, use the supernatant for the assay.

   Note: Lysis buffer is supplied as **Component G** (Cat# CA-N015g).