



### EnzyChrom™ NADP/NADPH Assay Kit

Pyridine nucleotides play an important role in metabolism and, thus, there is continual interest in monitoring their concentration levels. Quantitative determination of NADP<sup>+</sup>/NADPH has applications in research pertaining to energy transformation and redox state of cells or tissue.

Simple, direct and automation-ready procedures for measuring NADP<sup>+</sup>/NADPH concentration are very desirable. BioAssay Systems' EnzyChrom™ NADP<sup>+</sup>/NADPH assay kit is based on a glucose dehydrogenase cycling reaction, in which a tetrazolium dye (MTT) is reduced by NADPH in the presence of phenazine methosulfate (PMS). The intensity of the reduced product color, measured at 565 nm, is proportionate to the NADP<sup>+</sup>/NADPH concentration in the sample. Our assay is a convenient method to measure NADP, NADPH and their ratio.

#### KEY FEATURES

**Sensitive and accurate.** Detection limit 0.1 μM, linearity up to 10 μM NADP<sup>+</sup>/NADPH in 96-well plate assay.

**Convenient.** The procedure involves adding a single working reagent, and reading the optical density at time zero and 30 min at room temperature. No 37°C heater is required.

**High-throughput.** Can be readily automated as a high-throughput 96-well plate assay for thousands of samples per day.

#### APPLICATIONS

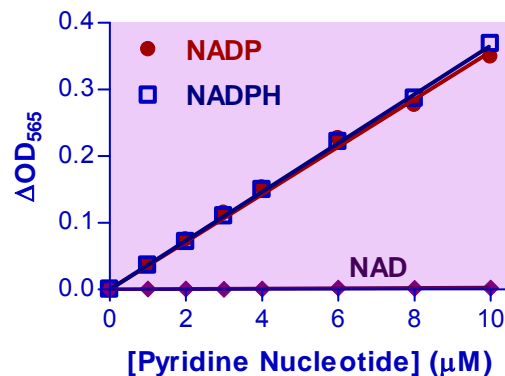
**Direct Assays:** NADP<sup>+</sup>/NADPH concentrations and ratios in cell or tissue extracts.

#### PRODUCT INFORMATION:

EnzyChrom™ NADP/NADPH Assay Kit ECNP-100

Each kit is sufficient for 100 assays in 96-well plate. Kit includes:

- 1 x 10 mL Assay Buffer
- 1 x 2 mL Glucose
- 1 x 2 mL PMS Solution
- 1 x 2 mL MTT Solution
- 1 x 120 μL Enzyme (GDH)
- 1 x 0.5 mL NADP<sup>+</sup> Standard
- 1 x 0.5 mL NADPH Standard
- 1 x 12 mL NAD(P) Extraction Buffer
- 1 x 12 mL NAD(P)H Extraction Buffer



Standard Curves in 96-well plate assay

#### REFERENCES:

- [1]. Zhao, Z, Hu, X and Ross CW (1987). Comparison of Tissue Preparation Methods for Assay of Nicotinamide Coenzymes. *Plant Physiol.* 84: 987-988.
- [2]. Matsumura, H. and Miyachi S (1980) Cycling assay for nicotinamide adenine dinucleotides. *Methods Enzymol.* 69: 465-470.
- [3]. Vilcheze, C et al. (2005). Altered NADH/NAD<sup>+</sup> Ratio Mediates Coresistance to Isoniazid and Ethionamide in Mycobacteria. *Antimicrobial Agents and Chemotherapy.* 49(2): 708-720.