

QuantiChrom™ Peroxide Assay Kit

Peroxide (e.g. hydrogen peroxide H_2O_2) is one of the key reactive oxygen species formed under oxidative stress conditions. High levels of peroxide formation have been linked to pathological conditions such as ageing, asthma, diabetes, atherosclerosis, cataract, inflammatory arthritis and neurodegenerative diseases.

Simple, direct and automation-ready procedures for quantitative determination of peroxide find wide applications in research and drug discovery. BioAssay Systems' peroxide assay kit is designed to measure peroxide concentration in biological samples without any pretreatment. The improved method utilizes the chromogenic Fe^{3+} -xylenol orange reaction, in which a purple complex is formed when Fe^{2+} provided in the reagent is oxidized to Fe^{3+} by peroxides present in the sample. The intensity of the color, measured at 540-610nm, is an accurate measure of the peroxide level in the sample. The optimized formulation substantially reduces interference by substances in the raw samples.

APPLICATIONS:

Direct Assays: H_2O_2 in biological samples (e.g. serum, citrate-plasma, urine, cell lysate, culture medium).

Pharmacology: effects of drugs on peroxide metabolism.

KEY FEATURES:

Sensitive and accurate. Enhanced color intensity using sorbitol. Detection range 0.4 μ M (14 ng/mL) to 100 μ M (3,400 ng/mL) H₂O₂ in 96-well plate assay. Simple and high-throughput. The procedure involves

Simple and high-throughput. The procedure involves addition of a single working reagent and incubation for 30 min. Can be readily automated as a high-throughput assay in 96-well plates for thousands of samples per day.

Potential interference reduced. For each sample, a background control is run under identical conditions with the omission of the Fe²⁺. This procedure corrects for any interference from endogenous iron present in biological samples.

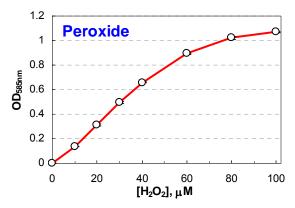
PRODUCT INFORMATION:

QuantiChrom[™] Peroxide Assay Kit

DIOX-250

Each kit is sufficient for 250 assays in 96-well plate. Kit includes:

- 1 x 1 mL Reagent A
- 1 x 1 mL Reagent B (for background control)
- 1 x 50 mL Reagent C
- 1 x 1 mL 35% stabilized H₂O₂Standard



Standard Curve in 96-well plate assay

REFERENCES:

- [1]. Karageuzyan KG (2005). Oxidative stress in the molecular mechanism of pathogenesis at different diseased states of organism in clinics and experiment. Curr Drug Targets Inflamm Allergy. 4(1):85-98.
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- [3]. Banerjee D, Madhusoodanan UK, Nayak S, Jacob J. (2003). Urinary hydrogen peroxide: a probable marker of oxidative stress in malignancy. Clin Chim Acta. 334: 205-9.