

Peroxide Assay Kit

Cat. No. Kit-0676 **Lot. No.** (See product label)

SPECIFICATION

Product Overview	Peroxide Assay Kit is a quantitative colorimetric peroxide determination at 585 nm.
Description	Peroxide (e.g. hydrogen peroxide H₂O₂) 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. Peroxide Assay Kit is designed to measure peroxide concentration in biological samples without any pretreatment. The improved method utilizes the chromogenic Fe³⁺-xylene orange reaction, in which a purple complex is formed when Fe²⁺ provided in the reagent is oxidized to Fe³⁺ 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: H₂O₂ in biological samples (e.g. serum, citrate-plasma, urine, cell lysate, culture medium). Pharmacology: effects of drugs on peroxide metabolism.
Usage	For research use only (RUO)
Storage	Store the kit at -20°C.
Kit Components	Reagent A: 1 mL Reagent B: 50 mL Standard: 100 μ 3% stabilized H₂O₂. Kit shipped at room temperature. Storage conditions. The kit is shipped at room temperature.

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Store all reagents at 4°C. Shelf life: 6 months after receipt. Precautions: reagents are for research use only. Normal precautions for laboratory reagents should be exercised while using the reagents. Please refer to Material Safety Data Sheet for detailed information.

Detection method Colorimetric

Compatible Sample Types Cell Lysate, Culture Medium, Plasma (Citrate), Serum, Urine

Features & Benefits Sensitive and accurate: Enhanced color intensity using sorbitol. Detection range 0.2 μ M (7 ng/mL) to 30 μ M (1,020 ng/mL) H₂O₂ in 96-well plate assay. Simple and high-throughput: The procedure involves addition of a single detection 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.

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