

Fluorimetric Hydrogen Peroxide Assay Kit (Near Infrared Fluorescence)

Product Information

Cat

Kit-0985

Common Name

H2O2

Cat.No.

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Product Overview

The Fluorimetric Hydrogen Peroxide Assay Kit uses our unique IR peroxidase substrate to quantify hydrogen peroxide in solutions and cell extracts. IR generates the fluorescence that is pH-independent from pH 4 to 10. It is a superior alternative to ADHP for the detections that require low pH where ADHP has reduced fluorescence. In addition, IR generates a product that has maximum absorption at 647 nm with maximum emission at 670 nm. This near infrared fluorescence minimizes the assay background that is often caused by the autofluorescence of biological samples. It can also be used to detect a variety of oxidase activities through enzyme-coupled reactions.

This Fluorimetric Hydrogen Peroxide Assay Kit is an optimized "mix and read" assay that is compatible with HTS liquid handling instruments. It provides a sensitive, one-step fluorometric assay to detect as little as 3 picomoles of H2O2 in a 100 μ L assay volume (30 nM). The assay can be performed in a convenient 96-well or 384-well microtiter-plate format and easily adapted to automation without a separation step. Its signal can be easily read by either a fluorescence microplate reader at Ex/Em = ~640/680 nm or an absorbance microplate reader at ~650 nm. Due to its long emission wavelength, this kit has low interference from biological samples.

Description

Hydrogen peroxide (H2O2) is a reactive oxygen metabolic by-product that serves as a key regulator for a number of oxidative stress-related states. It is involved in a number of biological events that have been linked to asthma, atherosclerosis, diabetic vasculopathy, osteoporosis, a number of neurodegenerative diseases and Down's syndrome. Perhaps the most intriguing aspect

Fluorimetric Hydrogen Peroxide Assay Kit (Near Infrared Fluorescence)

of H₂O₂ biology is the recent report that antibodies have the capacity to convert molecular oxygen into hydrogen peroxide to contribute to the normal recognition and destruction processes of the immune system. Measurement of this reactive species will help to determine how oxidative stress modulates a variety of intracellular pathways.

Storage

Keep in freezer and avoid exposure to light.

Size

500 Tests

Kit Components

Component A: IR Peroxidase Substrate 1 vial

Component B: H₂O₂ 1 vial (3% stabilized solution, 200 μL)

Component C: Assay Buffer 1 bottle (100 mL)

Component D: Horseradish Peroxidase 1 vial (20 units)

Component E: DMSO 1 vial (0.5 mL)

Features & Benefits

Broad Application: Can be used for quantifying hydrogen peroxide in solutions, in cell extracts and in live cells, and for detecting a variety of oxidase activities through enzyme-coupled reactions.

Sensitive: Detect as low as 30 picomoles of H₂O₂ in solution.

Continuous: Easily adapted to automation without a separation step.

Convenient: Formulated to have minimal hands-on time. No wash is required.

Non-Radioactive: No special requirements for waste treatment.

Preparation

1. Prepare stock solutions:

1.1 100X IR peroxidase substrate stock solution: Add 250 μL of DMSO (Component E) into the vial of IR Substrate (Component A). The stock solution should be used promptly; any remaining solution should be aliquoted and refrozen at -20°C.

Note: Avoid repeated freeze-thaw cycles and protect from light.

1.2 20 U/mL Peroxidase stock solution: Add 1 mL of Assay Buffer (Component C) into the vial of

Fluorimetric Hydrogen Peroxide Assay Kit (Near Infrared Fluorescence)

Horseradish Peroxidase (Component D).

Note: The unused HRP solution should be divided into single use aliquots and stored at -20°C.

1.3 20 mM H₂O₂ stock solution: Add 22.7 uL of 3% H₂O₂ (0.88 M, Component B) into 977 uL of Assay Buffer (Component C).

Note: The diluted H₂O₂ solution is not stable. The unused portion should be discarded.

2. Prepare H₂O₂ reaction mixture: Prepare the H₂O₂ reaction mixture according to the following table and keep from light:

Table 1 H₂O₂ Reaction mixture for one 96-well plate (2X)

IR Peroxidase Substrate Stock Solution (100X, from Step 1.1) 50 uL

20 U/ml Peroxidase Stock Solution (from Step 1.2) 200 uL

Assay Buffer (Component C) 4.75 mL

Total volume 5 mL

3. Prepare serial dilutions of H₂O₂ standard (0 to 10 μM):

Warning 1: IR Peroxidase Substrate (Component A) is unstable in the presence of thiols such as DTT and β- mercaptoethanol. If the final concentration of the thiols is higher than 10 uM, it would significantly decrease the assay dynamic range.

Warning 2: NADH and glutathione (reduced form of GSH) may interfere with the assay.

3.1 Add 1 μL of 20 mM H₂O₂ solution (from Step 1.3) into 1999 μL of Assay Buffer (Component C) to get a 10 μM H₂O₂ standard.

3.2 Take 200 μL of 10 μM H₂O₂ standard to perform 1:3 serial dilutions to get 3, 1, 0.3, 0.1, 0.03, 0.01 and 0 μM serial dilutions of H₂O₂ standard.

3.3 Add serial dilutions of H₂O₂ standard and H₂O₂-containing test samples into a solid black 96-well microplate as described in Tables 2 and 3.

Table 2 Layout of H₂O₂ standards and test samples in a solid black 96-well microplate

BL BL TS TS

HS1 HS1

HS2 HS2

HS3 HS3

HS4 HS4

HS5 HS5

Fluorimetric Hydrogen Peroxide Assay Kit (Near Infrared Fluorescence)

HS6 HS6

HS7 HS7

Note: HS= H₂O₂ Standards; BL=Blank Control; TS=Test Samples

Table 3 Reagent composition for each well

H₂O₂ Standard: Serial dilutions*: 50 μL

Blank Control: Assay Buffer (Component C): 50 μL

Test Sample: 50 μL

*Note: Add the serially diluted H₂O₂ standards from 0.01 μM to 10 μM into wells from HS1 to HS7 in duplicate.

Assay Protocol

4. Run H₂O₂ assay in supernatants reaction:

4.1 Add 50 μL of H₂O₂ reaction mixture (from Step 2) into each well of H₂O₂ standard, blank control, and test samples (see Step 3.3) to make the total H₂O₂ assay volume of 100 μL/well.

Note: For a 384-well plate, add 25 μL of sample and 25 μL of H₂O₂ reaction mixture in each well.

4.2 Incubate the reaction at room temperature for 15 to 30 minutes, protected from light.

4.3 Monitor the fluorescence increase with a fluorescence plate reader at Ex/Em = 640/680 nm

Note 1: IR peroxidase substrate is easy to be self-oxidized, so read the fluorescence as soon as the H₂O₂ reaction mixture was added to increase the signal to noise ratio.

Note 2: 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 650 nm. The absorption detection has lower sensitivity compared to the fluorescence reading.

5. Run H₂O₂ assay for cells:

The Fluorimetric Hydrogen Peroxide Assay Kit can be used to measure the release of H₂O₂ from cells. The following is a suggested protocol that can be modified to meet the specific research needs.

5.1 The H₂O₂ reaction mixture should be prepared as Step 2 except that the Assay Buffer (Component C) should be replaced with the media that is used in your cell culture system.

Suggested media including (a) Krebs Ringers Phosphate Buffer (KRPB); (b). Hanks Balanced Salt Solution (HBSS); or (c) Serum-free media.

Fluorimetric Hydrogen Peroxide Assay Kit (Near Infrared Fluorescence)

5.2 Prepare cells in a 96-well plate (50 - 100 μ L/well), and activate the cells as desired.

Note: The negative controls (media alone and non-activated cells) are included for measuring background fluorescence.

5.3 Add 50 μ L of H₂O₂ reaction mixture (from Step 5.1) into each well of cells and H₂O₂ standards (from Step 3.3).

Note: For a 384-well plate, add 25 μ L of cells and 25 μ L of H₂O₂ reaction mixture into each well.

5.4 Incubate the reaction at room temperature for 15 to 30 minutes, protected from light.

5.5 Monitor the fluorescence increase with a fluorescence plate reader at Ex/Em = 640/680 nm.

Analysis

The fluorescence in blank wells (with the assay buffer only) is used as a control, and is subtracted from the values for those wells with the H₂O₂ reactions.

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.
