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BIOXYTECH[®] H₂O₂-560™

Quantitative Hydrogen Peroxide Assay

For Research Use Only. Not For Use In Diagnostic Procedures.

Catalog No. 21024

INTENDED USE

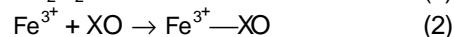
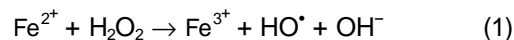
The BIOXYTECH[®] H₂O₂-560™ Assay is intended for the colorimetric quantitative determination of hydrogen peroxide (total hydroperoxides) in aqueous samples.

INTRODUCTION

The study of hydrogen peroxide in biological systems prompts the need for rapid and quantitative measurement of hydrogen peroxide. The researcher also requires a method of measuring hydrogen peroxide in laboratory reagents and detergents to prevent the unintended contribution of peroxidation by reagent use. The BIOXYTECH[®] H₂O₂-560™ Assay allows for simple colorimetric measurement of hydrogen peroxide.

PRINCIPLES OF THE PROCEDURE

The BIOXYTECH[®] H₂O₂-560™ Assay is based on the oxidation of ferrous ions (Fe²⁺) to ferric ions (Fe³⁺) by hydrogen peroxide under acidic conditions, equation (1). The ferric ion binds with the indicator dye xylenol orange (3,3'-bis[N,N-di(Carboxymethyl)-aminomethyl]-o-cresolsulfone-phthalein, sodium salt) to form a stable colored complex which can be measured at 560 nm, equation (2).



The Fe³⁺-xylenol orange complex has an extinction coefficient of $1.5 \times 10^4 \text{ M}^{-1}\text{cm}^{-1}$ at 560 nm when ferric ions are added in the absence of hydrogen peroxide. However, in the presence of sorbitol there is substantial chain oxidation of ferrous ion, increasing the color yield. This results in an apparent extinction coefficient of $2.67 \times 10^5 \text{ M}^{-1}\text{cm}^{-1}$, indicating that approximately 18 moles of Fe²⁺ are oxidized to Fe³⁺ for every mole of hydrogen peroxide present, increasing the sensitivity of the assay.

REAGENTS

Materials Provided

R1: 25 mM ammonium iron(II) sulfate, 2.5 M H₂SO₄, 1 mL.

R2: 100 mM sorbitol, 125 μM xylene orange in water, 100 mL.

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Warnings and Precautions

- **For Research Use Only. Not For Use In Diagnostic Procedures.**
- For *In Vitro* Use Only.
- Wear appropriate laboratory attire, including eye protection and disposable gloves.
- In case of accidental exposure to skin, eyes or mucous membranes, thoroughly rinse affected area with water.

Preparation

Prepare working reagent by mixing 1 volume of R1 with 100 volumes of R2. Prepare only the volume needed for the day's assays.

Reagent Storage

Store the reagents at 2-8°C. Reagents are stable until the expiration date on the label. The prepared working reagent is stable for 12 hours when stored at 2-8°C. Discard unused working reagent at the end of the day.

PROCEDURE

Materials Required But Not Provided

- 30% H₂O₂
- Deionized water
- Adjustable pipets with disposable tips
- Test tubes and spectrophotometric cuvettes or microplates
- Spectrophotometer or microplate reader capable of an absorbance reading in the 520-600 nm range
- 2.5 M H₂SO₄ (for optional sample blank)

Standard Preparation

1. Prepare fresh standards for each use.
2. Prepare a 25 mM standardized H₂O₂ stock solution by diluting 285 μL 30% H₂O₂ to 100 mL with DI H₂O.
3. Standardize using 43.6 M¹•cm⁻¹ as the molar extinction coefficient for H₂O₂ at 240 nm, *i.e.*, in a 1 cm cuvette, a 25 mM H₂O₂ solution will have an absorbance of 1.09 at 240 nm.
4. Dilute 25 mM H₂O₂ 1/250 with DI H₂O to give a 100 μM standard (highest standard).
5. Perform dilutions of the 100 μM standard to obtain a range of standards with appropriate concentrations for the samples to be assayed.

Assay Method

1. Add 1 volume of sample or standard to 10 volumes of working reagent.
2. Mix well and incubate at room temperature for 25-30 minutes.
3. Zero the spectrophotometer.
4. Measure the absorbance at 560 nm.

Stability of Final Reaction Material

The incubation step is necessary for the reaction to reach an endpoint. Once formed, the complex is stable for up to 12 hours. The absorbance should be measured the same day.

Assay Notes

- Excessive H_2O_2 can bleach color from the dye, resulting in lower measured absorbance. If the target concentration of the sample is unknown, perform a 1:100 dilution of the sample and repeat the assay. Absorbance values for the diluted sample that are the same or higher than the original reading indicate excessive H_2O_2 . Repeat the process of performing 1:100 dilutions and reassaying the sample until the dilution-corrected concentration is consistent.
- If endogenous iron is suspected, such as in biological samples, a sample blank should be assayed by replacing the working reagent with “blinking” reagent (1 volume 2.5 M H_2SO_4 mixed with 100 volumes R2). The measured absorbance should be subtracted from the sample absorbance.

RESULTS

Standard Curve

- Use a range of standards appropriate for the samples to measure.
- The maximum standard concentration to use is 100 μM . At concentrations above 100 μM the standard curve reaches a plateau.
- Plot absorbance *versus* standard concentration.
- The standard curve can be *approximated* by linear regression.
- Other curve fitting or graphical interpolation may be used.

Sample Values

- Subtract the blank absorbance values from sample absorbance values to obtain the net absorbance.
- If using linear regression or other mathematical curve fit, obtain concentration from the net absorbance using the equation of the standard curve.
- If using interpolation, obtain the concentration graphically from the plot.

LIMITATIONS OF THE PROCEDURE

- Endogenous iron and other transition metals may increase color development. Use the sample blanking procedure if needed.
- High concentrations of fructose, sorbitol, sucrose, glucose and formic acid in samples result in increased color yield.