# DHR Probe

KP-06-004 250/500/1000 test



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All chemicals should be handled with care

➤ This kit is for R&D use only

## Introduction

Reactive Oxygen Species can be induced by some stress conditions like exposure to oxidant or drugs. This fact leads to oxidative stress.

ROS induce damage in DNA, protein and lipids, with important consequences in cells.

Cell permeant reagent Dihydrorhodamine 123 (DHR 123) is a fluorogenic dye that is useful for the detection of reactive oxygen species (ROS) such as peroxide and peroxynitrite. After cell uptake, DHR 123 is oxidized by ROS into a fluorescent compound.

It seems that neither the superoxide, the NO, nor the hydrogen peroxide by themselves, are capable of oxidizing DHR. These ROS, are thought to combine with other cellular components such as cytochrome c oxidase or Fe2+ in order to oxidize DHR 123 to its fluorescent derivative Rhodamine 123.

Rhodamine 123 can be detected by fluorimeter, flow cytometer or fluorescence microscope with a maximum excitation and emission spectra of 500 and 536 nm, respectively.

It can be also detected by absorbance spectroscopy at 500 nm ( $\varepsilon = 78,800 \text{ M}^{-1} \text{ cm}^{-1}$ )

## Materials

## BQCkit DHR Probe KP06004-250 tests contains:

Product	Quantity	Storage
DHR 123	1 vial	RT

## BQCkit DHR Probe KP06004-500 tests contains:

Product	Quantity	Storage
DHR 123	2 vials	RT

## BQCkit DHR Probe KP06004-1000 tests contains:

Product	Quantity	Storage
DHR 123	4 vials	RT

# Reagent Preparation

Dilute DHR probe (1000X) with PBS (not included). Use the required amount of DHR and PBS for your tests.

Example: 1 µL of DHR probe (1000X) with 999 µL of PBS.

# Assay Protocol

# Short protocol:

#### -Adherent cells-

Seed adherent cells at 25 x 10<sup>3</sup> per well one day before performing the assay.

#### -Suspension cells-

Grow suspension cells in sufficient amount. (In the step 5 you will need  $100 \times 10^3$  cells per group).



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#### -Adherent cells-

Remove the media and add 100 µL/well of PBS.

## -Suspension cells-

Collect cells and wash by centrifugation PBS.

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#### -Adherent cells-

Remove PBS stain cells by adding density of µL/well 100 previously DHR 123 Reagent Preparation).

## -Suspension cells-

and Resuspend cells at a  $1x10^{6}$ of cells/mL. Stain the cells diluted with the desired volume (see of previously diluted DHR 123 (see Reagent Preparation).

#### -Adherent cells-

-Suspension cells-

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5



Incubate at cells' optimal temperature in dark conditions. An time Of incubation minutes İS 15-60 enough.

Incubate at cells' optimal temperature in dark conditions. An incubation time of 15-60 minutes is enough.

# -Suspension cells-



Ex/Em= 500/536 nm

### -Adherent cells-

Remove media and PBS. Measure fluorescence immediately.

Wash cells by centrifugation. Resuspend cells add at least 100 µL of PBS, seed in 96-well microplate with 100,000 stained cells/well and measure fluorescence immediately.

FLOW Cytometer: For cytometer application, follow the protocol for suspension cells, avoiding point 5.

# Warranties and Limitation of Liability

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Buyer's exclusive remedy and Bioquochem's sole liability hereunder shall be limited to a refund of the purchase price, or the replacement of all material that does not meet our specifications.

Said refund or replacement is conditioned on buyer giving written notice to Bioquochem within 30 days after arrival of the material at its destination.

Expiration date: 1 year from the date of delivery

For further details, please refer to our website <a href="https://www.bqckit.com">www.bqckit.com</a>.