CytoFix™ Red Mitochondrial Stain

Catalog number: 23200
Unit size: 500 Tests

Component	Storage	Amount (Cat No. 23200)
CytoFix™ MitoRed	Freeze (< -15 °C), Minimize light exposure	1 vial (100 µL)

OVERVIEW

CytoFix™ Red mitochondrial stain is a dye that selectively stains mitochondria independent of mitochondrial membrane potential gradient. Due to this functionality, CytoFix™ Red mitochondrial stain is well retained in mitochondria even after fixation. The dye permeates intact live cells and gets trapped in live cells. Its key features include high staining efficiency, long retention after fixation with minimal hands on time. CytoFix™ Red mitochondrial stain can be used with GFP expressed cells without overlapping the fluorescence of GFP, making it useful for multiplexing analysis. It can be used for both suspension and adherent cells and readily adapted for a wide variety of fluorescence platforms.

AT A GLANCE

1. Prepare cells in growth medium
2. Incubate cells with CytoFix™ MitoRed working solution for 20-30 minutes at 37 °C
3. Remove CytoFix™ MitoRed working solution
4. Fix cells with formaldehyde (Optional)
5. Analyze under fluorescence microscope with Cy3/TRITC filter set

KEY PARAMETERS

Fluorescence microscope

Emission	Cy3/TRITC filter set
Excitation	Cy3/TRITC filter set
Recommended plate	Black wall/clear bottom
Instrument specification(s)	Cy3/TRITC filter set

CELL PREPARATION

For guidelines on cell sample preparation, please visit https://www.aatbio.com/resources/guides/cell-sample-preparation.html

PREPARATION OF WORKING SOLUTION

Add 20 µL of stock solution into 10 mL of Hanks and 20 mM Hepes buffer (HHBS) or buffer of your choice or cell culture medium, and mix well.

Note 20 µL stock solution is enough for one 96-well plate assay. The staining conditions may be modified according to the particular cell type and the permeability of the cells or tissues to the probe.

Note Unused CytoFix™ MitoRed stock solution can be aliquoted and stored at ≤ -20 °C with smaller aliquots. Protect from light and avoid repeated freeze-thaw cycles.

SAMPLE EXPERIMENTAL PROTOCOL

1. Prepare cells in growth medium.
2. Add 100 µL/well (96-well plate) or 50 µL/well (384-well plate) of CytoFix™ MitoRed working solution in the cell plate.

   Note The optimal concentration of the cell membrane probe varies depending on the specific application.

   Note Serum in the culture growth medium may interfere with the staining. Staining in the absence of culture growth medium should improve the staining intensity.
3. Incubate the cells at 37 °C for 20-30 minutes, protected from light.
4. Remove working solution in each well. Wash cells with HHBS or buffer of your choice. (Optional)
5. **Optional:** Fix the cells with 4 % solution of paraformaldehyde for 20-30 minutes at room temperature. Wash twice to get rid of fixation solution.
6. Observe the fluorescence signal in cells using fluorescence microscope with a Cy3/TRITC filter set.

EXAMPLE DATA ANALYSIS AND FIGURES

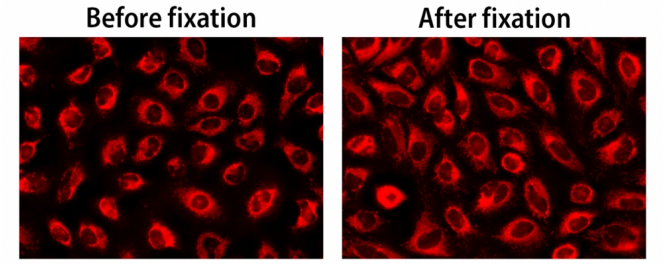


Figure 1. The fluorescence images of HeLa cells stained with CytoFix™ MitoRed in a 96-well black-wall clear-bottom plate. Image was acquired before (Left) and after (Right) fixation with 4% formaldehyde solution for 20 minutes at RT. The cells were imaged using fluorescence microscope with a Cy3/TRITC filter.

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