

Mito-Tracker Deep Red 633

Introduction

Mito-Tracker Deep Red 633 is a red-fluorescent probe that selectively labels mitochondria. Like some conventional mitochondria probes tetramethylrosamine and rhodamine 123, Mito-Tracker Deep Red 633 is potential-dependent. So Mito-Tracker Deep Red 633 stains live cells well but is not suitable for fixed cells. Meanwhile, Mito-Tracker Deep Red 633 can be used to monitor changes in mitochondria membrane potential during apoptosis. Mito-Tracker Deep Red 633 is easy to use that rapidly accumulates in mitochondria and can be imaged without washing.

Components and Storage

| Components | B8809-50 μg | |
|--|-------------|--|
| Mito-Tracker Deep Red 633 | 50 μg | |
| This product should be stored at -20°C away from light and moisture. Stable for 3 years. | | |

Properties

| Physical Appearance | Solid | Landon Company |
|---------------------|-----------------|---|
| Ex/Em | 622/648 | A Paragraphic and the state of |
| Solubility | Soluble in DMSO | Table Paris |

Protocol

1. **Preparation of the stock solution:** Dissolve 50 μg Mito-Tracker Deep Red 633 in 460 μL anhydrous DMSO to make a 200 μM stock solution. The stock solution should be stored at -20°C away from light. It is recommended to aliquot the stock solution into small volumes and avoid repeated freeze/thaw cycles.

*Note: Allow Mito-Tracker Deep Red 633 to warm to room temperature before opening.

2. Preparation of the working solution: Dilute the stock solution in a suitable buffer (for example, HBSS with Calcium and Magnesium) or growth medium to make a working solution. The recommended concentration of the working solution is 20-200 nM. To reduce non-specific staining, keep the concentration of working solution as low as possible. It is suggested to dilute Mito-Tracker Deep Red 633 when using it.

*Note: The optimal concentration of working solution varies depending on the type of cell.

3. Labeling of Mitochondria: For adherent cells, grow cells to reach the desired density. Remove the growth medium and add an appropriate working solution to cover the cells. Incubate at 37°C away from light for 15 min. Replace the working solution with a fresh, pre-warmed suitable buffer or growth medium. Then detect the fluorescence signal of cells by a microscope with a Cy5 filter set.

*Note: The optimal time for incubation varies depending on the type of cells. For suspension cells, harvest cells and perform similarly to the adherent cells.

Note

- 1. Fluorescent probes are easy to quench, please protect them from light when using.
- 2. For your safety and health, please wear lab coats and gloves during the experiment.
- 3. For research use only. Not to be used in clinical diagnostic or clinical trials.









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