

Product name: Cell Tracker CM-DiD

Product number: RA20008

Basic Information

Product Name	Cell Tracker CM-DiD
Size	200 μL
Storage	Store at 4 °C, away from light
Shipping	Shipped with ice pack
Validity	12 months

Experimental procedures

The volume of working solution should be optimized according to different cell lines and experimental systems. It is recommended to start exploring the optimal conditions from 10 times the recommended volume.

Suspension cell staining

- (1) Add an appropriate volume of staining working solution to resuspend the cells to a density of 1×10^6 cells/mL, and then add the staining stock solution at a ratio of 1:200.
- (2) Incubate the cells at 37°C for 2 to 20 minutes. Different cells have different optimal culture times. You can use 20 minutes as the initial incubation time and then optimize the system.
- (3) After incubation, centrifuge at 1000-1500 rpm for 5 min. Pour off the supernatant and slowly add 37°C preheated culture medium to resuspend the cells. Repeat twice.

Adherent cell staining

- (1) Prepare the staining working solution: add 5 μ L of the staining stock solution to every 1 mL of culture medium and vortex to mix.
- (2) Culture the adherent cells on a sterile coverslip. After the culture is completed, remove the coverslip and aspirate away excess culture medium, but keep the surface moist.
- (3) Add 100 μ L of dye working solution to one corner of the coverslip and gently shake to allow the dye to evenly cover all cells.
- (4) Incubate the cells at 37°C for 2 to 20 minutes. The optimal incubation time varies for different cells. You can use 20 minutes as the initial incubation time, and then optimize the system to obtain a uniform labeling effect.
- (5) Aspirate the dye working solution, wash the coverslip 2-3 times with culture medium, cover all cells with prewarmed culture medium each time, incubate for 5-10 min, and then aspirate the culture medium. However, keep the surface moist.

Cell Tracker CM-DiD excitation/emission wavelength: 644/663 nm

Note: This reagent is for scientific research use only!