

# **Live-Dead Cell Staining Kit**

(Catalog #K501-100; 100 stainings; Store kit at -20°C)

#### I. Introduction:

Distinguishing between live and dead cells is very important for investigation of growth control and cell death. The *Live-Dead Cell Staining Kit* provides the ready-to-use reagents for convenient discrimination between live and dead cells. The kit utilizes Live-Dye<sup>TM</sup>, a cell-permeable green fluorescent dye (Ex/Em = 488/518 nm), to stain live cells. Dead cells can be easily stained by propidium iodide (PI), a cell non-permeable red fluorescent dye (Ex/Em = 488/615). Stained live and dead cells can be visualized by fluorescence microscopy using a band-pass filter (detects FITC and rhodamine). The kit provides sufficient reagents for 100 stainings using 24-well plate.

#### II. Kit Contents:

Component	K501-100	Part Number
Solution A (1 mM Live-Dye)	50 µl	K501-100-1
Solution B (2.5 mg/ml PI)	50 μl	K501-100-2
Staining Buffer	50 ml	K501-100-3

# III. Cell Staining Protocol:

- Prepare enough Staining Solution for your assay (0.5 ml per well in 24 well dish): mix 1 µl of Solution A and 1 µl of Solution B in 1 ml of Staining Buffer. Scale up accordingly for larger numbers of assays.
- 2. Collect cells (1 x 10<sup>6</sup> cells) by centrifugation at 500 X g for 5 min.
- 3. Resuspend to 0.5 ml Staining Solution
- 4. Incubate for 15 min at 37°C.
- 5. Place the cell suspension on a glass slide. Cover the cells with a glass coverslip. For analyzing adherent cells, grow cells directly on a coverslip. Following incubation with the Staining Solution, invert coverslip on a glass slide and visualize cells.
- 6. Observe cells immediately under a fluorescence microscope using a band-pass filter (detects fluorescein and rhodamine).
  - Healthy cells stain only the cell-permeable Live-Dye, fluorescing green. Dead cells can stain both the cell-permeable Live-Dye and the cell non-permeable PI (red), the overlay of green and red appears to be yellow-red.

# IV. Caution:

As the optimal staining conditions may vary among different cell types, we recommend that a suitable concentration of Solution A and B be determined individually.

Please note that PI is suspected to be highly carcinogenic, so careful handling of the reagent is required.

# V. Storage and Stability:

Store kit at -20°C. Protect from light. Store Staining Buffer at 4°C after opening. All reagents are stable for 1 year under proper storage conditions.

#### FOR RESEARCH USE ONLY! Not to be used on humans.

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