

# Live or Dead™ Fixable Dead Cell Staining Kit \*Blue Fluorescence with 405 nm Excitation\*

Catalog number: 22500 Unit size: 200 Tests

Component	Storage	Amount	
Component A: Stain It™ Violet 450	Freeze (<-15 °C), Minimize light exposure	1 vial	
Component B: DMSO	Freeze (<-15 °C)	1 vial (200 μL)	

#### **OVERVIEW**

Our Live or Dead™ Fixable Dead Cell Staining Kits are a set of tools for labeling cells for fluorescence microscopic investigations of cellular functions. The effective labeling of cells provides a powerful method for studying cellular events in a spatial and temporal context. This particular kit is designed to uniformly label fixed mammalian cells in blue fluorescence for flow cytometry applications with violet laser excitation. The kit uses a proprietary blue fluorescent dye that is more fluorescent upon bonding to cellular components. The fluorescent dye used in the kit is well excited with the violet laser (405 nm excitation) to fluorescence at 460 nm. The kit provides all the essential components with an optimized cell-labeling protocol. It is an excellent tool for preserving of fluorescent images of particular cells, and can also be used for fluorescence flow cytometry applications.

#### AT A GLANCE

#### **Protocol summary**

- 1. Prepare samples in HHBS (0.5 mL/assay)
- 2. Replace with HHBS
- 3. Add Stain It™ Violet 450 to the cell suspension
- 4. Stain the cells at room temperature or 37°C for 20 60 minutes
- 5. Wash the cells
- 6. Fix the cells (optional)
- Examine the sample with flow cytometer and/or fluorescence microscope using the appropriate Excitation/Emission filter

**Important** Thaw all the components at room temperature before starting the experiment.

#### **KEY PARAMETERS**

Instrument: Fluorescence microscope

Excitation: 410 nm Emission: 450 nm

Recommended plate: Black wall/clear bottom

Instrument: Flow cytometer Instrument specification(s): see Table 1

### PREPARATION OF STOCK SOLUTIONS

Unless otherwise noted, all unused stock solutions should be divided into single-use aliquots and stored at -20 °C after preparation. Avoid repeated freeze-thaw cycles.

1. Stain It™ Violet 450 stock solution (500X):

Add 200 µL DMSO (Component B) into the vial of Stain It™ Violet 450 (Component A) to make 500X Stain It™ Violet 450 stock solution.

## PREPARATION OF CELL SAMPLES

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

#### SAMPLE EXPERIMENTAL PROTOCOL

**Table 1.** Fluorescence spectra properties and suggested excitation laser for flow cytometry analysis

Cat. #	Description	Ex (nm)	Em (nm)	Excitation
				Source
22500	Blue Fluorescence with 405 nm Excitation	410	450	405 nm
22501	Green Fluorescence with 405 nm	408	512	405 nm
	Excitation			
22502	Orange Fluorescence with 405 nm	398	550	405 nm
	Excitation			
22599	Red Fluorescence Optimized for Flow	523	617	488 nm
	Cytometry			
22600	Blue Fluorescence	353	442	335 nm
22601	Green Fluorescence	498	521	488 nm
22602	Orange Fluorescence	547	573	561 nm or
				488 nm
22603	Red Fluorescence	583	603	561 nm
22604	Deep Red Fluorescence	649	660	633 nm
22605	Near Infrared Fluorescence	749	775	633 nm

- Prepare cells using 1X Hanks and 20 mM Hepes buffer (HHBS) or sodium azidefree and serum/protein-free buffer of your choice.
- Wash cells once with HHBS or the azide- and serum/protein-free buffer of your choice.
- 3. Resuspend cells at 5  $10 \times 10^6/mL$  in HHBS or in the azide- and serum/protein-free buffer of your choice.
- Add 1 µL of 500X Stain It<sup>™</sup> Violet 450 stock solution to 0.5 mL of cells/assay and mix it well.
- Incubate at room temperature or 37°C, 5% CO<sub>2</sub> incubator for 20 60 minutes, protected from light.

**Note** The optimal stain concentrations and incubation time should be experimentally determined for different cell lines.

- 6. Wash cells twice and resuspend cells with HHBS or the buffer of your choice.
- 7. Fix cells as desired (optional).
- 8. Analyze cells with flow cytometer and/or fluorescence microscope using the appropriate Excitation/Emission filter (see Table 1).

### **EXAMPLE DATA ANALYSIS AND FIGURES**

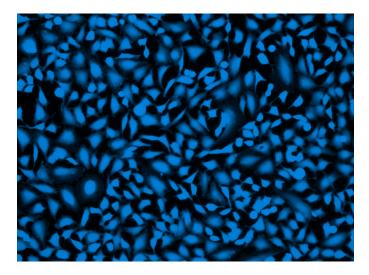


Figure 1. Image of Hela cells fixed with formaldehyde and stained with Live or Dead™ Fixable Dead Cell Staining kit \*Blue Fluorescence with 405 nm Excitation\* in a Costa black wall/clear bottom 96-well plate.

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