

Zombie UV387™ Fixable Viability Kit

Catalog# / Size	423125 / 100 tests 423126 / 500 tests
Regulatory Status	RUO
Other Names	Fixable Dye, Fixable Viability Dye, Dead, Apoptosis
Description	Zombie UV387™ is an amine-reactive fluorescent dye that is non-permeant to live cells but permeant to cells with compromised membranes. Thus, it can be used to assess live vs. dead status of mammalian cells. Zombie UV387™ is a polar, water-soluble dye, making it suitable for use in multi-color detection.

Product Details

Preparation	Zombie UV387™ Fixable Viability Kit is composed of lyophilized Zombie UV387™ dye and anhydrous DMSO. For reconstitution, pre-warm the kit to room temperature; add 100 µL of DMSO to one vial of Zombie UV387™ dye and mix until fully dissolved. 100 tests = 1 vial of Zombie UV387™ + DMSO, 500 tests = 5 vials of Zombie UV387™ + DMSO.
Storage & Handling	Store kit at -20°C upon receipt. Do not open vials until needed. Once the DMSO is added to the Zombie UV387™ dye, use immediately, or store at -20°C in a dry place and protected from light, preferably in a desiccator or in a container with desiccant for no more than one month.
Application	FC - Quality tested
Recommended Usage	Each lot of this product is quality control tested by immunofluorescent staining with flow cytometric analysis. For flow cytometry, the suggested dilution is 1:100 - 1:1000 for 1 - 10 million cells. For immunofluorescence microscopy, the suggested dilution is 1:1000. It is recommended that the reagent be titrated for optimal performance for each application, as optimal dosage varies with cell type.
Excitation Laser	Ultraviolet Laser (355 nm)
Application Notes	Zombie UV387™ dye is excited by the UV laser (355 nm) and has fluorescence emission maximum at 387 nm. If using in a multi-color panel design, filter optimization may be required depending on other fluorophores used. Zombie UV387™ has similar spectra to Spark UV™ 387.

Standard Cell Staining Protocol:

1. Prior to reconstitution, spin down the vial of lyophilized reagent in a microcentrifuge to ensure the reagent is at the bottom of the vial.
2. For reconstitution, pre-warm the kit to room temperature; add 100 µL of DMSO to one vial of Zombie UV387™ dye and mix until fully dissolved.
3. Wash cells with PBS buffer (no Tris buffer and protein free).
4. Dilute Zombie UV387™ dye at 1:100 - 1000 in PBS. Resuspend 1-10 x 10⁶ cells in diluted 100 µL Zombie UV387™ solution. To minimize background staining of live cells, titrate the amount of dye and/or number of cells per 100 µL for optimal performance. Different cell types can have a wide degree of variability in staining based on cell size and degree of cell death.

1. **Note:** Don't use Tris buffer as a diluent and be sure that the PBS does not contain any other protein like BSA or FBS.
2. **Note:** The amount of dye used can also influence the ability to detect apoptotic as well as live and dead cells.

5. Incubate the cells at room temperature (or 4°C), in the dark, for 15 - 30 minutes.
6. Wash one time with 2 mL BioLegend's Cell Staining Buffer (Cat. No. 420201) or equivalent buffer containing serum or BSA.
7. Continue performing antibody staining procedure as desired.
8. Cells can be fixed with paraformaldehyde or methanol prior to permeabilization or can be analyzed without fixation.

No-wash Sequential Staining Protocol:

1. Wash cells with PBS buffer (no Tris buffer and protein free).
2. For reconstitution, pre-warm the kit to room temperature; add 100 µL of DMSO to one vial of Zombie UV387™ dye and mix until fully dissolved.

3. Determine the total μL volume of antibody cocktail previously titrated and optimized for the assay that will be added to each vial/well of cells based on a final volume of 100 μL . Subtract that antibody volume from the 100 μL total staining volume intended for the assay. In the remaining volume, dilute Zombie UV387™ dye at 1:100 - 1000 in PBS as determined by prior optimization at that volume. For example, if you are adding 20 μL of antibody cocktail for a 100 μL total staining volume, use 80 μL of Zombie UV387™ solution. Resuspend $1 - 10 \times 10^6$ cells in the appropriate volume of Zombie UV387™ solution. Different cell types can have a wide degree of variability in staining based on cell size and degree of cell death.

1. **Note:** Don't use Tris buffer as a diluent and be sure that the PBS does not contain any other protein like BSA or FBS.
2. **Note:** The amount of dye used can also influence the ability to detect apoptotic as well as live and dead cells.

4. Incubate for 10 - 15 minutes at RT (or 4°C), protected from light. Without washing the cells, add the cell surface antibody cocktail and incubate for another 15 - 20 minutes.

5. Add 1 - 2 mL Cell Staining Buffer (Cat. No. 420201) or equivalent buffer containing BSA or serum. Centrifuge to pellet.

6. Continue with normal fixation and permeabilization procedure. If planning to skip fixation and analyze cells live, complete an additional wash step to minimize any unnecessary background of the live cells.

1. **Notes:** If the cell type in use cannot tolerate a protein-free environment, then titrate the Zombie UV387™ dye in the presence of the same amount of BSA/serum as will be present in the antibody staining procedure. A higher amount of Zombie UV387™ may be required since the BSA/serum will react with and bind up some proportion of the Zombie UV387™.

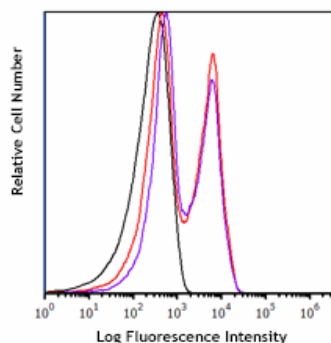
Antigen Details

Biology Area	Apoptosis/Tumor Suppressors/Cell Death, Cell Biology, Neuroscience
Gene ID	NA

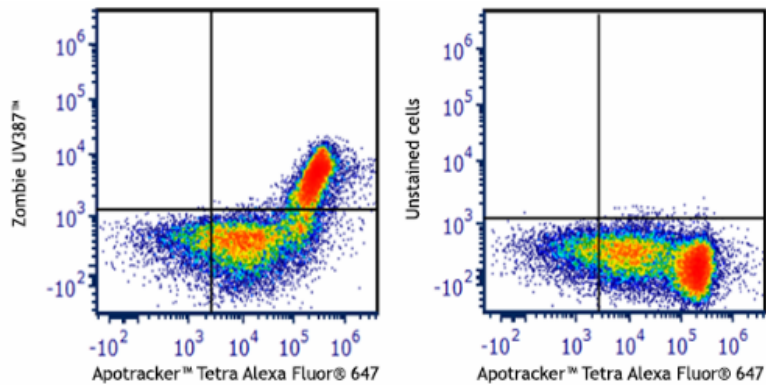
Related Protocols

- [Cell Surface Flow Cytometry Staining Protocol](#)

Product Data



Heat-killed (mixed 1:1 with live) C57BL/6 mouse splenocytes were stained with Zombie UV387™ and analyzed before fixation (purple) or after fixation (red). Cells alone, without Zombie UV387™ staining, are indicated in black.



Heat-killed (mixed 1:1 with live) C57BL/6 mouse splenocytes were stained with Apotracker™ Tetra Alexa Fluor® 647 and Zombie UV387™ (left) or stained with Apotracker™ Tetra Alexa Fluor® 647 only (right).

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