

PRODUCT INFORMATION SHEET

Catalog number: 22542 Unit size: 200 Tests

Cell Meter™ VX500 fixable viability dye

 Component
 Storage
 Amount

 Cell Meter™ VX500 fixable viability dye
 Freeze (< -15 °C), Minimize light exposure</td>
 200 tests

OVERVIEW

Discrimination and exclusion of dead cells from live cells allows cleaner separation and identification of cell populations. Cell Meter™ fixable viability dyes are a large family of cell-impermeable fluorescent viability dyes that are optimized to match the major excitation lasers of common flow cytometers, such as 350, 405, 488, 633 and 647 nm. These dyes are impermeant to live cells but permeant to cells with compromised membranes. They irreversibly react with amine- and thiol-containing proteins and other cellular components. Since dead or fixed cells with a compromised membrane more readily react with Cell Meter™ fixable cell stains, thus stain brighter than live cells with an intact membrane, these dyes can be used to assess live vs. dead status of mammalian cells. There are a few factors to be considered when using these dyes, e.g., the titration of each dye to ensure that live cells have minimal to no staining. Cell Meter™ VX500 fixable cell stain is optimized to be excited with the violet laser at 405 nm with emission at 500 nm. Compared to other commercially similar viability dyes, this fixable viability dye is much more robust and stable.

AT A GLANCE

Protocol summary

- 1. Prepare samples in HHBS (0.5 mL/assay)
- 2. Add Cell Meter[™] VX500 to the cell suspension
- 3. Stain the cells at room temperature for 20 60 minutes
- 4. Wash the cells
- 5. Fix the cells (optional)
- Examine the sample with flow cytometer using 525/50 nm filter (Pacific Orange channel)

Important

Thaw at room temperature before starting the experiment.

KEY PARAMETERS

Flow cytometer

| Excitation | |
|-----------------------------|--|
| Emission | |
| Instrument specification(s) | |

405 nm laser 525/50 nm filter Pacific Orange channel

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.

Cell Meter™ VX500 stock solution (500X)

Add 200 μL of DMSO into the Cell MeterTM VX500 vial to make 500X stock solution.

Note The unused Cell Meter™ VX500 stock solution should be divided into single use aliquots and stored at -20 °C. Avoid repeated freeze/thaw cycles.

SAMPLE EXPERIMENTAL PROTOCOL

1. Treat cells as desired.

- Wash cells once with HHBS or the azide- and serum/protein-free buffer of your choice.
- Resuspend cells at 5 10 × 10 ⁶ /mL in HHBS or in the azide- and serum/protein-free buffer of your choice.
- Add 1 µL of 500X Cell Meter™ VX500 stock solution to 0.5 mL of cells/assay and mix it well.
- 5. Incubate at room temperature, 5% CO ₂ incubator for 20 60 minutes, protected from light.

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

- Wash cells twice and resuspend cells with HHBS or the buffer of your choice.
- 7. Fix cells as desired (optional).
- Analyze cells with a flow cytometer using 525/50 nm filter (Pacific Orange channel).

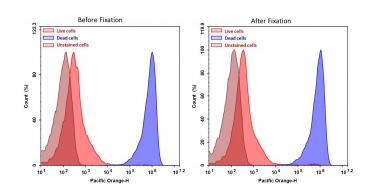


Figure 1. Detection of Jurkat cell viability by Cell Meter[™] fixable viability dye. Jurkat cells were treated and stained with Cell Meter[™] VX500 (Cat#22542), and then fixed in 3.7% formaldehyde and analyzed by flow cytometry. The dead cell population (Blue peak) is easily distinguished from the live cell population (Red peak) with Pacific Orange channel, and nearly identical results were obtained before and after fixation.

DISCLAIMER

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EXAMPLE DATA ANALYSIS AND FIGURES