

## **MITO-ID<sup>®</sup> Membrane Potential Detection Kit**

Catalog #: ENZ-51018

ENZ-51018-K100	100 assays
ENZ-51018-0025	25 assays

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Please read entire booklet before proceeding with the assay.



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## I. INTRODUCTION

Cell-based assays for evaluating the functional status of mitochondria are emerging as useful tools for elucidating the role of mitochondrial activity in drug-induced toxicity, the apoptosis cascade and other cellular and biochemical processes. The loss of the mitochondrial membrane potential (MMP) is often associated with the early stages of apoptosis. The collapse of MMP coincides with the opening of the mitochondrial permeability transition pores, leading to the release of cytochrome C into the cytosol, which in turn triggers other downstream events in the apoptotic cascade. Enzo Life Sciences' MITO-ID® Membrane Potential Detection Kit provides an improved approach for measuring changes in MMP, as demonstrated by a shift in the fluorescence emission maximum of a dual-emission MMP-sensitive dye from orange to green. Concomitantly, the kit monitors cell death as loss of plasma membrane integrity, as highlighted by the ability of a red-emitting, membrane-impermeable DNA intercalating dye to penetrate into the cell nucleus.

The MITO-ID® Membrane Potential Detection Kit has been optimized for measurement of MMP and cell viability by conventional fluorescence microscopy and flow cytometry. A related kit, MITO-ID® Membrane Potential Cytotoxicity Assay Kit (Cat. No. ENZ-51019-KP002) has been optimized for use in fluorescence microplate-based assays. The MITO-ID® MP Detection Reagent is a cationic carbocyanine dye that fluoresces either green or orange, depending upon MMP status. In energized cells, the MITO-ID® Membrane Potential dye exists as a green-fluorescent monomer in the cytosol, but more importantly also accumulates as orange-fluorescent aggregates in the mitochondria. However, in apoptotic and necrotic cells, this dye exits mitochondria and exists primarily as green-fluorescent monomers in the cytosol. A cell-impermeant DNA intercalator, the Necrosis Detection Reagent, similar to the red-emitting dye propidium iodide, is used to monitor late-stage apoptosis and necrotic cell death. Compared with the commonly used cationic carbocyanine dye JC-1,<sup>1-3</sup> the emission spectrum of the MITO-ID® Membrane Potential dye is slightly blue-shifted and thus more readily distinguishable from the Necrosis Detection Reagent, allowing easier compensation of spectral overlap. Additionally, the dye is more soluble, more photostable and approximately 10-25-fold more sensitive to MMP changes than the conventional JC-1 dye.

A control perturbation agent, CCCP, is provided as a positive control for monitoring loss in MMP. Potential applications for using the MITO-ID<sup>®</sup> Membrane Potential Detection Kit as a live-cell assay are for monitoring cellular events leading to MMP changes and/or apoptosis, as well as monitoring certain cell treatment conditions that lead to MMP change without eliciting apoptosis or alternatively lead to apoptosis without leading to MMP changes. Other cellular events that may alter MMP include drug-induced toxicity relating to perturbations in lipid metabolism, mitochondrial respiratory chain breakdown or reduced mitochondrial DNA content arising from anti-viral therapy. Simultaneously monitoring MMP and plasma membrane integrity can potentially provide useful mechanistic information for drug safety evaluation of novel compounds. For example, the assay can potentially be used to differentiate among compounds and rank their order of potency.



Avoid freeze / thaw cycles

## II. REAGENTS PROVIDED AND STORAGE

All reagents are shipped on dry ice. Upon receipt, the kit should be stored at -20°C, protected from light. **Avoid repeated freezing and thawing.**

Reagents provided in the ENZ-51018-K100 kit are sufficient for 100 assays using either live, adherent cells or cells in suspension. Reagents provided in the ENZ-51018-0025 kit are sufficient for 25 assays using either live, adherent cells or cells in suspension.

Reagent	ENZ-51018-K100	ENZ-51018-0025
MITO-ID <sup>®</sup> MP Detection Reagent	500µl	130µL
Necrosis Detection Reagent	600µl	150µL
CCCP Control (2mM)	100µl	100µL
10X Assay Buffer 1	25ml	7ml
50X Assay Buffer 2	5ml	1.5ml

### III. OTHER MATERIALS NEEDED

- Standard fluorescence microscope or a flow cytometer that is equipped with a 488nm laser for fluorescent dye excitation
- Phosphate buffered saline (PBS)
- 5ml tubes appropriate for holding cells during induction of apoptosis and necrosis
- Calibrated, adjustable precision pipets, preferably with disposable plastic tips
- Adjustable speed centrifuge with swinging buckets (for suspension cultures)
- Glass microscope slides (for microscope analysis only)
- Glass cover slips (for microscope analysis only)
- Deionized water
- Growth medium (e.g., Dulbecco's modified Eagle medium, D-MEM)

### IV. SAFETY WARNINGS AND PRECAUTIONS

- This product is for research use only and is not intended for diagnostic purposes.
- Some components of this kit may contain hazardous substances. They can be harmful if ingested or absorbed through the skin and may cause irritation to the eyes. The reagents of the kit should be treated as possible mutagens and should be handled with care and disposed of properly.
- Observe good laboratory practices. Gloves, lab coat, and protective eyewear should always be worn. Never pipet by mouth. Do not eat, drink or smoke in the laboratory areas. All blood components and biological materials should be treated as potentially hazardous and handled as such. They should be disposed of in accordance with established safety procedures.
- To avoid photobleaching, perform all manipulations in low light environments or protected from light by other means.



Handle with care



Protect from light

## V. METHODS AND PROCEDURES

**NOTE:** PLEASE READ THE ENTIRE PROCEDURE BEFORE STARTING. Allow all reagents to be used to warm to room temperature before proceeding. Upon thawing of solutions, gently hand-mix or vortex the reagents prior to use to ensure a homogenous solution.

### A. CELL PREPARATION

Cells should be maintained via standard tissue culture practices. A positive control for MMP status can be performed using the CCCP Control provided in the kit. It is recommended that treatment with the agent be performed using 2-10 $\mu$ M final concentration in order to observe changes in MMP and that the final percent DMSO in the assay not exceed 0.2%. Treated cells should be harvested and diluted to a working concentration of  $5.0 \times 10^5$  -  $1 \times 10^6$  per ml in media. Negative control cells should be treated with vehicle (DMSO, media or other solvent used to reconstitute or dilute an inducer or inhibitor) for an equal length of time under similar conditions.

### B. REAGENT PREPARATION

#### 1. Positive Control

CCCP (carbonyl cyanide 3-chlorophenylhydrazone) is a proton ionophore and uncoupler of oxidative phosphorylation in mitochondria. As such, it is useful for depolarizing mitochondria. Addition of CCCP should cause a dose-dependent reduction in mitochondrial orange fluorescence. Prepare this perturbation agent as a positive control for mitochondrial membrane potential loss. Before beginning the experiment, ensure that the vial of CCCP has equilibrated to room temperature. CCCP is supplied as a 2mM stock solution in DMSO.

#### 2. 1X Assay Solution

Allow the 10X Assay Buffer 1 and the 50X Assay Buffer 2 to warm to room temperature. Make sure that the reagents are free of any crystallization before use. Prepare enough 1X Assay Solution for the number of samples to be assayed. For each 10ml preparation of 1X Assay Solution, add 1ml 10X Assay Buffer, and 0.2ml 50X Assay Buffer 2 into 8.8ml deionized water. Mix well.

### 3. Dual Detection Reagent

**NOTE:** *The MITO-ID<sup>®</sup> MP Detection Reagent is light sensitive. Avoid direct exposure of the reagent to intense light. Aliquot and store unused reagent at –20°C, protected from light. Avoid repeated freeze/thaw cycles.*

Immediately prior to use, prepare sufficient amount of Dual Detection Reagent for the number of samples to be assayed as follows: To every 1ml of 1X Assay Solution (from step 2), add 10µl of MITO-ID<sup>®</sup> MP Detection Reagent and 4µl of Necrosis Detection Reagent. Mix well.

### C. LIVE CELL ANALYSIS BY FLUORESCENCE MICROSCOPY (SUSPENSION CELLS)

1. Cells should be cultured to a density not to exceed  $1 \times 10^6$  cells/mL. Make sure that cells are in the log phase of growth before starting an experiment.
2. Treat the cells with the compound of interest and the negative control cells with vehicle. Each cell line should be evaluated on an individual basis to determine the optimal cell density for monitoring MMP changes.
3. Transfer 0.5ml of cell suspension into a sterile centrifuge tube and centrifuge for 5 minutes at room temperature at 400 x *g*.
4. Remove the supernatant and carefully resuspend and wash the cells with 2ml of 1X Assay Solution (see **section B-2** for preparation).
5. Carefully remove the supernatant and suspend the cells in 500µl of the Dual Detection Reagent (prepared as per **section B-3**).
6. Protect the samples from light and incubate the cells at 37°C in a 5% CO<sub>2</sub> incubator for 15 min.

7. Observe the cells immediately following staining by fluorescence microscopy. The orange fluorescence emission associated with energized mitochondria with high MMP can be detected using a rhodamine filter set (excitation = 540nm, emission = 570nm). The green fluorescence emission associated with depolarized mitochondria with low MMP can be detected using a fluorescein (FITC) filter set (excitation = 485nm, emission = 530nm). The red fluorescence associated with nuclear staining of cells with compromised plasma membrane integrity can be detected using a 7-AAD or propidium iodide filter (excitation = 546nm, emission = 674nm).
8. Positive Control Samples: It is recommended that a set of control samples be pretreated with CCCP at a final concentration of 2 $\mu$ M in medium or buffer of choice for 30 mins. Once again, concentration of this inducer and duration of exposure may require adjustment, depending upon the cell line being evaluated.
9. Follow steps 2-7 for post-treatment.

#### **D. LIVE CELL ANALYSIS BY FLUORESCENCE MICROSCOPY (ADHERENT CELLS)**

1. Grow the cells directly onto glass slides or polystyrene tissue culture plates until ~80% confluent.
2. Treat the cells with the compound of interest and negative control cells with vehicle.
3. Carefully wash the cells twice with 1X Assay Solution (see **section B-2**, page 5, for preparation) in a volume sufficient for covering the cell monolayer.
4. Carefully remove the supernatant and dispense the Dual Detection Reagent (prepared as per **section B-3**, page 6) in a volume sufficient for covering the cell monolayer.
5. Protect the samples from light and incubate for 15 minutes at room temperature.
6. Flick the staining solution onto a paper towel and if necessary add a few drops of 1X Assay Solution to prevent the cells from drying out.

7. Cover the cells and observe under a fluorescence/confocal microscope. The orange fluorescence emission associated with energized mitochondria with high MMP can be detected using a rhodamine filter set (excitation = 540nm, emission = 570nm). The green fluorescence emission associated with depolarized mitochondria with low MMP can be detected using a fluorescein (FITC) filter set (excitation = 485nm, emission = 530nm). The red fluorescence associated with nuclear staining of cells with compromised plasma membrane integrity can be detected using a 7-AAD or propidium iodide filter (excitation = 546nm, emission = 674nm).
8. Positive Control Samples: It is recommended that a set of control samples be pretreated with CCCP at a final concentration of 2 $\mu$ M in medium or buffer of choice for 30 mins. Once again concentration of this inducer and duration of exposure may require adjustment, depending upon the cell line being evaluated.
9. Follow steps 3-7 for post-treatment.

#### **E. LIVE CELL ANALYSIS BY FLOW CYTOMETRY**

The protocol described in this manual assumes that the user is familiar with the basic principles and practices of flow cytometry and is able to run samples according to the operator's manual pertaining to the instrument being used.

1. Cells should be cultured to a density not to exceed 1 x 10<sup>6</sup> cells/mL during staining and analysis.
2. Treat the cells with the compound of interest and negative control cells with vehicle, per desired protocol. At the end of the treatment, trypsinize (adherent cells), or collect cells (suspension cells).
3. Centrifuge the cells for 5 min. at 400 x *g* at room temperature.
4. Carefully remove the supernatant by aspiration and gently resuspend and wash the pellet in 2ml 1X Assay Solution. (see **section B-2** for preparation)
5. Centrifuge the cells for 5 min. at 400 x *g* at room temperature.

6. Carefully remove the supernatant and gently resuspend the pellet in 500µl Dual Detection Reagent (prepared as per **section B-3**, page 6).
7. Protect the samples from light and incubate for 15 min. at room temperature.
8. Samples should be analyzed via flow cytometry using a 488nm laser with the FL 1 channel for the green fluorescent monomeric signal of the MITO-ID® Membrane Potential dye, the FL 2 channel for the orange fluorescent aggregated signal from the MITO-ID® Membrane Potential dye and the FL 3 channel for red fluorescent signal of the Necrosis Detection Reagent. It is recommended that compensation corrections be performed using single stained, induced cells to avoid any overlap between the MITO-ID® MP Detection Reagent and Necrosis Detection Reagent fluorescent signals.
9. Positive Control Samples: It is recommended that a set of control samples be pretreated with CCCP at a final concentration of 2µM for 30 mins. Once again, the concentration of this inducer and duration of exposure may require adjustment, depending upon the cell line being evaluated.
10. Follow steps 2-8 for post-treatment.

**Suggested Compensation Controls for Flow Cytometry (see Appendix B):**

- a. Unstained cells
- b. Cells stained with MITO-ID® MP Detection Reagent (without Necrosis Detection Reagent)
- c. Cells Stained with Necrosis Detection Reagent (without MITO-ID® MP Detection Reagent)

**Note:** Samples and controls should be kept on ice before the assay is run and analyzed via flow cytometry within one hour of staining.

## VI. APPENDICES

### A. FILTER SET SELECTION

The selection of optimal filter sets for fluorescence microscopy applications requires matching the optical filter specifications to the spectral characteristics of the dyes employed in the analysis. Please consult your instrument or filter set manufacturer for assistance in selecting optimal filter sets.

Filter sets suitable for Texas Red or tetramethylrhodamine (TRITC) are recommended for imaging the orange fluorescent signal of the MITO-ID® MP Detection Reagent, representing energized mitochondria. Filter sets suitable for fluorescein (FITC) are recommended for imaging the green fluorescent signal of the MITO-ID® MP Detection Reagent, representing de-energized mitochondria. Filter sets suitable for 7-AAD or propidium iodide (PI) are recommended for imaging the red fluorescent signal of the Necrosis Detection Reagent, representing compromised plasma membrane integrity.

### B. EXAMPLE OF FLOW CYTOMETRY COMPENSATION

Suggested controls for flow cytometry:

- Unstained cells (negative population)
  - Untreated cells stained with the MITO-ID® MP Detection Reagent only (highest orange and green signal)
  - CCCP-treated cells, stained with the MITO-ID® MP Detection Reagent only (lowest orange signal)
  - CCCP-treated cells, stained with Necrosis Detection Reagent only (highest red signal)
1. Generate a log FL1 (X-axis) versus log FL2 (Y-axis) dot plot. Run untreated cells stained with the MITO-ID® reagent only (compensation control 2). Set PMT for FL1 and FL2 channels so both green and orange signals will fall between the second and the third log decade scale of FL1 and FL2. Set up quadrant gate to register a dual positive cell population in the upper right quadrant (see **Figure 1**)
  2. Adjust compensation values for the MITO-ID® MP Detection Reagent and Necrosis Detection Reagents using the same untreated MITO-ID® stained control

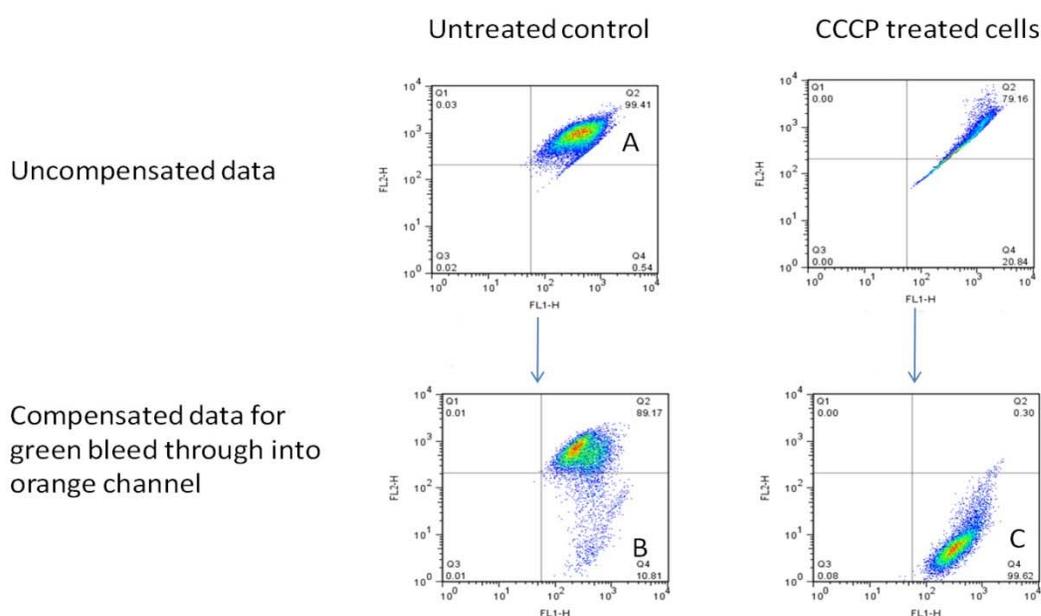
(compensation control 2) and PMT settings established as described above.

- The green MITO-ID® Membrane Potential dye signal fluoresces mostly in the FL1 channel, but bleeds over into the FL2 channel. This needs to be compensated (as shown in Figure 1) .
  - To compensate, subtract the FL1 bleed from FL2. The equation to use is  $FL2 - \dots \%FL1$  (The window “Compensation” in the Cells Quest or DiVa).
  - As the % compensation is increased, the green population is subtracted from the FL2 channel (double positive top right quadrant, A) and placed solely in single positive quadrant B.
  - Run Compensation control 3 to make sure that the overlap of the green signal into the orange channel is eliminated (see **Figure 1**).
  - Do not overcompensate – background of this sample, in terms of FL2 intensity, should be the same as that of a negative population (**Figure 1-C**)
  - If necessary, compensate for any bleed through from the orange channel into the green channel ( $FL1 - \dots \%FL2$ ).
3. Repeat the compensation adjustment for the Necrosis Detection Reagent using the single stained control (controls 3 and 4).
    - To compensate for red bleed through into orange use  $FL2 - \dots \%FL3$  and orange into red use  $FL3 - FL2\%$  (controls 3 and 4).
  4. Once compensation is complete do not change the voltages. Any further voltage changes will unbalance the compensations and the process will have to be repeated. The PMT settings for this assay may be quite low due to the strong fluorescence signals.
  5. Verify compensation values for each new experiment, or at least for each new cellular system tested.

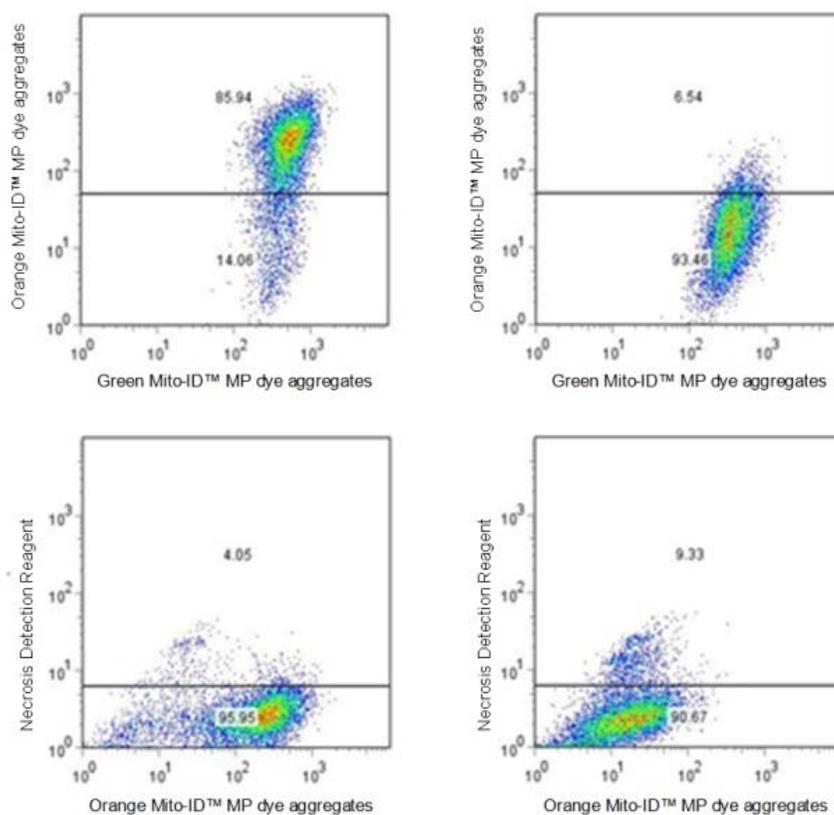
## C. RESULTS

Apoptosis is characterized by a diverse repertoire of biochemical and cellular signals that alter cellular morphology. Some of the downstream effects observed via fluorescence microscopy include cell shrinkage, loss of membrane symmetry, membrane blebbing and cytoplasmic condensation. Changes at the nuclear level are also evident, including the condensation and fragmentation of nuclear chromosomal DNA. By contrast, necrosis results from acute cellular damage or damage to an essential cellular system and is morphologically defined by early plasma membrane rupture and dilatation of organelles. MMP loss can be associated with a variety of forms of cell injury and death.

Typical results of flow cytometry-based analysis of cell populations using the MITO-ID<sup>®</sup> Membrane Potential Detection Kit are shown in Figure 2. The aggregated orange form of the MITO-ID<sup>®</sup> MP Detection Reagent has an emission maximum of 590nm, while the green monomeric form has emission maximum of 525nm, allowing 2-D flow cytometric analysis. Control cells show a high number of cells that display both orange and green fluorescence (Figure 2A). Treatment of cells with 2 $\mu$ M CCCP for 4 hours results in a dramatic shift in the fluorescence profile of the cell populations, with most cells displaying high green fluorescence but minimal orange fluorescence (Figure 2B).



**Figure 1:** Uncompensated data (top row) compared with compensated data (bottom row). This example demonstrates compensation for the green bleed through into the orange channel.



**Figure 2:** Monitoring mitochondrial dynamics in Jurkat cells by flow cytometry using MITO-ID<sup>®</sup> MP Detection Reagent. Jurkat cells were treated with DMSO (Left) or 2µM CCCP (Right) for 30 mins. Cells were labeled with MITO-ID<sup>®</sup> Detection Reagent for 15 minutes. Cells were analyzed on a FACS Calibur (Becton Dickinson, San Jose, CA) flow cytometer. A dot plot of orange fluorescence (FL2) versus green fluorescence (FL1) resolved live cells with intact MMP (Top Left) from de-energized cells that have lost MMP (Top Right). A dot plot of red (FL3) versus orange (FL2) fluorescence allowed to distinguish live (bottom left) and dead (bottom right) cells.

**REFERENCES**

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2. Cossarizza, Baccarani-Contri, Kalashnikova, and Franceschi (1993) "A new method for the cytofluorimetric analysis of mitochondrial membrane potential using the J-aggregate forming lipophilic cation 5,5',6,6'-tetrachloro-1,1',3,3' tetraethylbenzimidazolylcarbocyanine iodide (JC-1)." *Biochem. Biophys. Res. Commun.* 197 (1): 40-45.
3. Woollacott and Simpson. (2001) "High throughput fluorescence assays for the measurement of mitochondrial activity in intact human neuroblastoma cells." *J Biomol Screen* 6(6): 413-420.

## TROUBLESHOOTING GUIDE

Problem	Potential Cause	Suggestion
Fixed and/or permeabilized cells fail to stain with MITO-ID <sup>®</sup> MP Detection Reagent.	This dye is only suitable for live-cell staining.	Use the dye only for live-cell analysis.
Poor staining is observed.	The MITO-ID <sup>®</sup> MP Detection Reagent has been centrifuged.	Do not centrifuge the MITO-ID <sup>®</sup> MP Detection Reagent.
	Stained cells have been exposed to strong light.	Protect samples from exposure to strong light and analyze them immediately after staining.
	Kit reagent has degraded.	Verify that the reagents are not past their expiration dates before using them.
Control cells without treatment show low ratio of orange to green signal.	Control cells are not healthy. Extended storage of cells after staining may adversely affect their health.	Use healthy cells. Acquire data as soon as possible after staining the cells.
Cell staining is too strong.	The MITO-ID <sup>®</sup> MP Detection Reagent is too concentrated for the cell type being employed.	Dilute the staining solution further with the 1X Assay Solution.
Orange fluorescent MITO-ID <sup>®</sup> Membrane Potential dye signal increases upon treatment with the CCCP Control.	The orange emission filter in your instrument may be too wide and green fluorescent signal is being read, too.	Replace emission filter with a narrower band pass filter.

<b>Problem</b>	<b>Potential Cause</b>	<b>Suggestion</b>
Orange fluorescent MITO-ID <sup>®</sup> Membrane Potential dye intensity values are negative (< 0) on the microplate reader.	The fluorescence intensity value of the control sample treated with Dual Detection Reagent was set to zero instead of an actual fluorescence value.	Use 1X Assay Solution to set the zero value on the instrument.
Poor resolution observed between positive and negative populations.	Incomplete staining with the MITO-ID <sup>®</sup> MP Detection Reagent and/or the Necrosis Detection Reagent has occurred due to very low concentration of dye being employed for the number of cells or dye was incubated with the cells for an insufficient length of time.	Ensure that the positive and negative controls are stained adequately and if not, titer reagents to determine optimal dye-to-cell ratios. Make sure that the cells are stained for 30 minutes in a 37°C CO <sub>2</sub> incubator.
Accuracy of data from experiment to experiment is poor.	The concentration of the stained cell sample for data acquisition is too low.	Repeat experiments using a higher density of cells.
	Cells have not been properly resuspended prior to data acquisition.	Verify that cells have been properly resuspended prior to collecting data.



# Product Manual

## NOTES

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# Product Manual

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