

## Cell Meter™ Colorimetric MTT Cell Proliferation Kit

Catalog number: 22768, 22769  
Unit size: 1000 Tests, 5000 Tests

Component	Storage	Amount (Cat No. 22768)	Amount (Cat No. 22769)
Component A: MTT Reagent A	Freeze (< -15 °C), Minimize light exposure	40 mL	2 X 100 mL
Component B: MTT Reagent B	Room temperature (10-25 °C)	100 mL	500 mL

### OVERVIEW

Cell Meter™ assay kits are a set of tools for monitoring cell viability. A variety of parameters can be used to monitor cell viability. Cell Meter™ Colorimetric MTT Cell Proliferation Kit uses MTT (3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide) to quantify the number of live cells. The water-soluble MTT produces water-insoluble purple formazan in metabolically active cells. The amount of formazan produced is directly proportional to the number of living cells. The absorbance increase is measured around 560 nm. Unlike other commercial MTT assays that require the tedious solubilization step to dissolve the water-insoluble formazan product, Cell Meter™ Colorimetric MTT Cell Proliferation Kit is a simple mix and read format with the minimal hands-one time. Our proprietary formulation eliminates the time-consuming solubilization step. It is the most robust MTT test for determining the number of viable cells in cell proliferation and cytotoxicity assays. The detection sensitivity is 10 times higher than the traditional MTT assays.

### AT A GLANCE

#### Protocol summary

1. Prepare cells in a 96-well plate (100 µL/well)
2. Add MTT working solution (140 µL/well)
3. Incubate at 37 °C for 2-4 hours
4. Read absorbance at 560 nm

#### Important

Thaw all the kit components at room temperature before use.

### KEY PARAMETERS

#### Absorbance microplate reader

Absorbance 562 nm  
Recommended plate Clear bottom

### PREPARATION OF WORKING SOLUTION

#### MTT working solution

Prepare the amount of MTT working solution needed by mixing 4 mL of MTT Reagent A (Component A) with 10 mL of MTT Reagent B (Component B) (2:5, v/v ratio of MTT Reagent A: B). Mix well.

**Note** 14 mL MTT working solution is enough for 100 tests in a 96-well plate. Prepare enough MTT working solution right before the experiment, use promptly.

### SAMPLE EXPERIMENTAL PROTOCOL

1. Plate 1000 to 40,000 cells/well in a tissue culture microplate with clear bottom.
2. Add test compounds into the cells and incubate for a desired period of time (such as 24, 48 or 96 hours) in a 37 °C, 5% CO<sub>2</sub> incubator. For blank wells (medium without the cells), add the same amount of test compounds. The suggested total volume is 100 µL for a 96-well plate, and 25 µL for a 384-well plate.

**Note** Each cell line should be evaluated on an individual basis to

determine the optimal cell density for proliferation or cytotoxicity induction. For proliferation assays, use fewer cells; for cytotoxicity assays, use more cells to start with.

3. Add 140 µL/well (96-well plate) or 35 µL/well (384-well plate) of MTT working solution to each well.
4. Incubate the plate at 37 °C for 2-4 hours, protect from light.

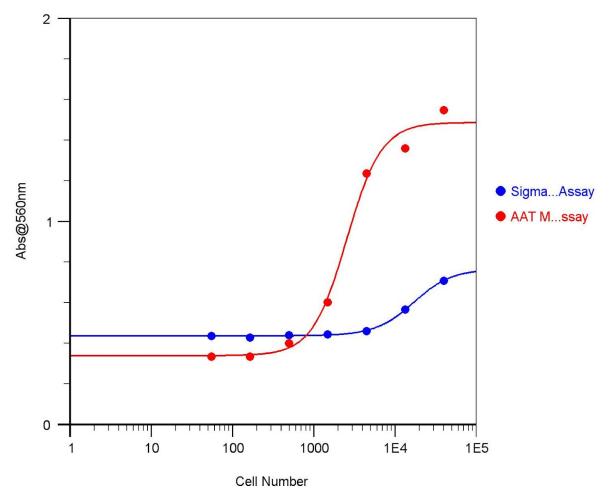
**Note** The incubation time could be from 2 hours to overnight depending on the individual cell type and cell concentration used. Optimize the incubation time for each experiment.

5. Monitor the absorbance increase with an absorbance plate reader at OD = 562 nm.

### EXAMPLE DATA ANALYSIS AND FIGURES

The reading (Abs@560nm) obtained from the blank standard well is used as a negative control. Subtract this value from the other standards' readings to obtain the base-line corrected values. Then, plot the standards' readings to obtain a standard curve and equation. This equation can be used to calculate Cell Number samples. We recommend using the Online Four Parameter Logistics Calculator which can be found at:

<https://www.aatbio.com/tools/four-parameter-logistic-4pl-curve-regression-online-calculator>



Cell numbers were determined with Cell Meter™ Colorimetric MTT Cell proliferation Kit. HeLa cells at 0 to 40,000 cells/well/100 µL were added in a clear bottom 96-well plate for overnight. The absorbance was measured at 560 nm using a SpectraMax reader (Molecular Devices). 500 cells/well was detected compare to ~5,000 cells/well with Sigma's MTT assay kit.

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**Figure 1.** Cell numbers were determined with Cell Meter™ Colorimetric MTT Cell proliferation Kit. HeLa cells at 0 to 40,000 cells/well/100 µL were added in a clear bottom 96-well plate for overnight. The absorbance was measured at 560 nm using a SpectraMax reader (Molecular Devices). 500 cells/well was detected compare to ~5,000 cells/well with Sigma's MTT assay kit.

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