

Data sheet

XTT Cell Proliferation Assay Kit

Cat. No: CA031
1000 reactions

Introduction

The XTT Cell Proliferation Assay Kit is a colorimetric assay that detects the cellular metabolic activities. During the assay, the yellow tetrazolium salt XTT (sodium 2,3-bis(2-methoxy-4-nitro-5-sulfophenyl)-5-[(phenylamino)-carbonyl]-2H-tetrazolium) is reduced to a highly coloured formazan dye by dehydrogenase enzymes in metabolically active cells. This conversion only occurs in viable cells and thus, the amount of the formazan produced is proportional to viable cells in the sample. The formazan dye formed in the assay is soluble in aqueous solution and can be quantified by measuring the absorbance at wavelength 450 nm using a spectrophotometer. An electron coupling reagent, such as PMS (N-Methylphenazonium methyl sulfate), can significantly improve the efficiency of XTT reduction in cells.

Advantages/Features:

- Easy to use: there is no need for additional reagents and/or cell washing procedures.
- Rapid (no solubilisation step as in an MTT assay) results within 2-5 hours.
- Sensitivity: can be assayed even in low cell concentrations.
- Accuracy: dye absorbance is proportional to the number of cells in each well.
- No radioactivity
- Ideal for high throughput assays (no washing or other steps that can cause cell loss and variability)
- The entire assay can be performed directly in a microtiter plate.

Kit Contents

Components	1000 assays
XTT Reagent	2x25 mL
Activation Reagent	1 ml

Storage

Upon receipt, store the kit at -20°C. Protect XTT solution from light. Kit components are stable for at least 12 months from date of receipt when stored as recommended.

XTT Working Solution

Immediately before use, add 100 µl of the Activation Reagent to 5 mL of XTT reagent to make XTT Working Solution. Prepared XTT Working Solution should be added to cells within several minutes.

If sediment is present in the XTT Solution, heat the solution to 37°C and swirl gently until a clear solution is obtained

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Assay Procedure

1. Plate cells into 96-well tissue culture plates at a density of 10^4 - 10^5 cells/well in 100 μ l and the medium may contain up to 10% serum.
2. Carry out desired cell treatments.
3. Culture the cells in a CO₂ incubator at 37°C for 24-48 hours.
4. Add 50 μ l of the reconstituted XTT mixture to each well. Mix gently for one minute.
5. Incubate the cells for two hours (adherent culture) to four hours (suspension culture) at 37°C in a CO₂ incubator.

Note: *The optimal incubation time for this assay depends on experimental setup, such as: cell type, cell number, and treatment. Optimization of incubation time can be determined by reading one plate at various time points after addition of XTT solution.*

6. Shake the plate gently to evenly distribute the dye in the wells.
7. Measure the absorbance signal of the samples with a spectrophotometer at a wavelength of 450-500 nm. Measure background absorbance at a wavelength between 630-690 nm. Subtract background absorbance from signal absorbance to obtain normalized absorbance values:

$$A = A_{450-500\text{nm}}(\text{Test}) - A_{450-500\text{nm}}(\text{Blank}) - A_{630-690\text{nm}}(\text{Test})$$

8. Plot the absorbance values of your normalized data on the ordinate (Y-axis) and your experimental parameters (e.g., number of cells) on the abscissa (X-axis).

PRODUCT USE LIMITATION

This product is developed, designed and sold exclusively for research purposes and in vitro use only. The product was not tested for use in diagnostics or for drug development, nor is it suitable for administration to humans or animals. Please refer to www.canvaxbiotech.com for Material Safety Data Sheet of the product.

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