

Colorimetric TUNEL Apoptosis Assay

Cat. No. 8088

50 tests

Introduction

The ScienCell™ Colorimetric TUNEL Apoptosis Assay is used for detection of apoptosis (programmed cell death) in individual cells based on the terminal deoxynucleotidyl transferase (TdT)-mediated dUTP-biotin nick-end-labeling (TUNEL) technology. In brief, the 3'-OH end of the DNA strand breaks in apoptotic cells is labeled with biotinylated nucleotides using the enzyme TdT. Streptavidin conjugated horseradish peroxidase (HRP) is then bound to the biotinylated nucleotides, and visualized using the peroxidase substrate, 3,3'-diaminobenzidine (DAB). The nuclei of apoptotic cells should be observed dark brown under light microscope.

Kit Components

Cat. No.	# of vials	Name	Quantity	Storage
8088a	1	Equilibrium Buffer	5 ml	-20°C
8088b	1	TdT Solution	250 µl	-20°C
8088c	1	Biotin-dUTP Solution	2.5 ml	-20°C
8088d	1	Stock Streptavidin-HRP (100×)	50 µl	4°C, dark
8088e	1	Stock DAB Substrate (20×)	250 µl	4°C

Materials to be Supplied by the User

PBS
Tris-HCl, pH 7.5, 50 mM
BSA
Deoxyribonuclease I (optional)
Formaldehyde
Triton X-100
H₂O₂
Hematoxylin (optional)

Quality Control

The ScienCell™ Colorimetric TUNEL Apoptosis Assay is applied to Human Astrocytes (HAs) treated with and without DNase I, which serve as our positive and negative controls, respectively. Dark brown nucleus can only be observed in cells treated with DNase I (Figure 1).

The ScienCell™ Colorimetric TUNEL Apoptosis Assay is also applied to Rat Cortical Neurons cultured with ScienCell™ Neuronal Medium for 3 days, and apoptotic cells can be identified with dark brown nucleus (Figure 2).

Procedures (96-well plate)

A. Cell culture

1. Seed cells in 96-well culture plate in culture medium with or without test compounds. Culture the cells in a CO₂ humidified incubator at 37°C for the desired period of time.

B. Pretreatment of cells

1. Rinse cells three times with PBS, and fix cells by incubating with freshly prepared 4% formaldehyde in PBS for 10 minutes at room temperature.
2. Wash cells three times with PBS, for 2 minutes each time.
3. Permeabilize cells by incubating with 0.2% Triton X-100 in PBS for 15 minutes at room temperature.
4. Wash cells three times with PBS, for 2 minutes each time.

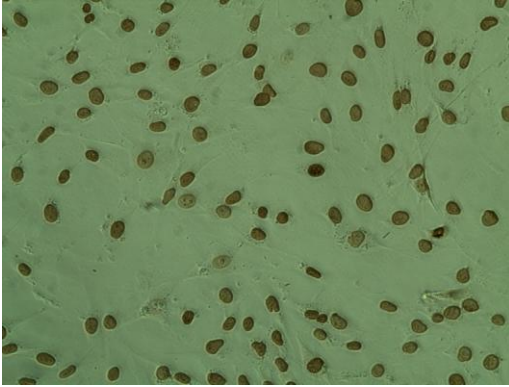
C. Setup of positive controls (optional)

1. Incubate control cells cultured without any test compounds with Deoxyribonuclease I (400 Kunits/ml in 50 mM Tris, pH 7.5 with 1 mg/ml BSA) for 10 minutes at room temperature to induce cleavage of genomic DNA.

D. TUNEL staining of cells

1. Pre incubate cells with 100 μ l/well of Equilibrium Buffer for 10 minutes at room temperature.
2. Prepare TUNEL reaction mixture just before use based on the number of samples to be assessed. For each well of 96-well plate, mix 5 μ l of TdT Solution with 45 μ l of Biotin-dUTP Solution.
3. Incubate cells in TUNEL reaction mixture (50 μ l per well) for 60 minutes at 37°C, protected from light.
4. Wash cells three times with PBS, for 2 minutes each time.
5. Block the endogenous peroxidase activity by incubating cells with 100 μ l/well of 0.3% H₂O₂ in PBS for 5 minutes at room temperature.
6. Wash cells three times with PBS, for 2 minutes each time.
7. Prepare working streptavidin-HRP by diluting Stock Streptavidin-HRP 100 \times with PBS. Add 100 μ l of working streptavidin-HRP to each well of 96-well plate, and incubate for 30 minutes at room temperature.
8. Wash cells three times with PBS, for 2 minutes each time.
9. Prepare working DAB substrate by diluting Stock DAB Substrate 20 \times with PBS containing 0.3% H₂O₂. Add 100 μ l of working DAB substrate to each well of 96-well plate, and incubate for about 10 minutes at room temperature in the dark, or until a light brown background develop.
10. Wash cells three times with PBS, for 2 minutes each time.
11. Counter stain cells with hematoxylin if needed.
12. Observe staining with a light microscope.

A.



B.

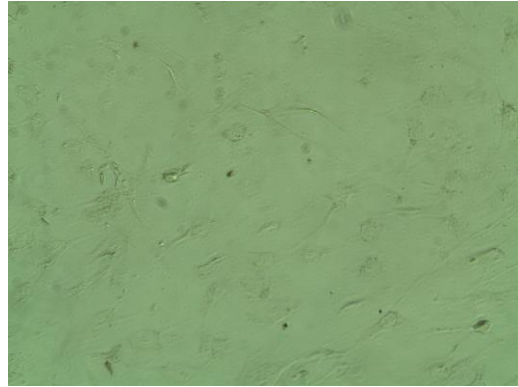


Figure 1. The ScienCell™ Colorimetric TUNEL Apoptosis Assay is applied to human astrocytes treated with (A) and without (B) DNase I, and data shows that dark brown nuclei can only be observed in those cells treated with DNase I.

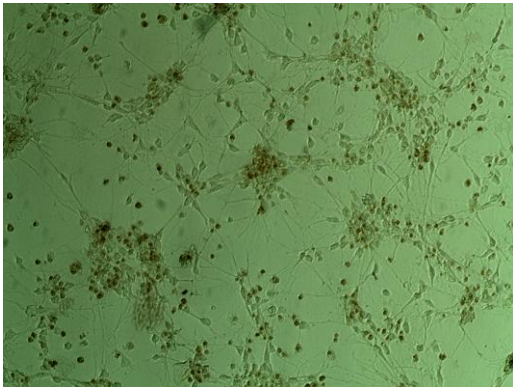


Figure 2. The ScienCell™ Colorimetric TUNEL Apoptosis Assay is applied to rat cortical neurons cultured with ScienCell™ Neuronal Medium for 3 days, and apoptotic cells can be identified with dark brown nucleus.