



VitroView™ In Situ TUNEL DAB Apoptosis Detection Kit
(50 assays)
SKU#: VB-4005D

Description

DNA fragmentation represents a characteristic hallmark of apoptosis. Terminal deoxynucleotidyl transferase dUTP nick end labeling (TUNEL) is designed to detect apoptotic cells that undergo extensive DNA degradation during the late stages of apoptosis. The method is based on the ability of terminal deoxynucleotidyl transferase (TdT) to label blunt ends of double-stranded DNA breaks independent of a template. The results are visualized with 3,3-diaminobenzidine (DAB).

Kit Contents

1. Proteinase K stock solution (20×) -----0.25 ml
2. 1×Proteinase K working buffer -----5 ml
3. TdT equilibration buffer -----6 ml
4. TdT enzyme -----15 µl
5. Biotinylated dUTP -----22 µl
6. RTU Streptavidin-HRP -----5 ml
7. DAB stock solution (40×) -----0.5 ml
8. Stable H₂O₂ solution (40×)----- 0.5 ml
9. DAB buffer -----15 ml
10. TUNEL positive FFPE slides -----2 slides

Storage

1. RTU Streptavidin-HRP and TUNEL positive FFPE slides: Stored at 4 °C.
2. Others: Stored at -20 °C.

Protocol

1. Preparation of Slides

A. Frozen Sections

- 1) Snap frozen fresh tissues in liquid nitrogen or isopentane pre-cooled in liquid nitrogen, embedded in OCT compound in cryomolds. Store the frozen tissue block at -80°C until ready for sectioning.
- 2) Transfer the frozen tissue block to a cryotome cryostat (e.g. -20°C) prior to sectioning and allow the temperature of the frozen tissue block to equilibrate to the temperature of the cryotome cryostat.
- 3) Section the frozen tissue block into a desired thickness (typically 5-10 µm) using the cryotome.
- 4) Place the tissue sections onto glass slides suitable for immunohistochemistry (e.g. Superfrost)
- 5) Sections can be stored in a sealed slide box at -80°C for later use.
- 6) Before staining, warm slides at room temperature for 30 minutes and fix slides by 2% neutralized formalin for 30 min. Rinse slides with PBS then transfer to a Coplin jar containing ice-cold 70% ethanol for 1 h. Slides may be stored overnight in 70% ethanol at 4°C.
- 7) Wash in PBS.
- 8) Follow procedure for pretreatment as required with TdT reaction buffer.

B. Paraffin Sections

- 1) Deparaffinize sections in xylene, 3×5min.

- 2) Hydrate with 100% ethanol, 2×2min.
- 3) Hydrate with 95% ethanol, 2×2min.
- 4) Rinse in distilled water.
- 5) Prepare 1ml of 1× proteinase K working solution by mixing 50µl of 20× proteinase K solution with 950 µl of 1×Proteinase K working buffer.
- 6) Prepare carefully blot away excess water and pipette 75-100 µl of 1×proteinase K solutions to cover sections. Incubate 8~15 min at room temperature.
- 7) Following proteinase K treatment, wash slides 3 × 5 min with ddH₂O.
- 8) (Optional) Inactivate endogenous peroxidases by covering sections with 2% hydrogen peroxide for 5 min at room temperature. Wash slides 3 × 5 min with ddH₂O.
- 9) Follow procedure for pretreatment as required with TdT reaction buffer.

C. Adherent Cells

- 1) Adherent cells may be cultured on glass chamber slides. Wash in PBS.
- 2) Fix slides by 2% neutralized formalin for 15 min. Rinse slides with PBS then transfer to a Coplin jar containing ice-cold 70% ethanol for 1 h. Slides may be stored overnight in 70% ethanol at 4°C
- 3) Wash in PBS. Inactivate endogenous peroxidases by covering sections with 2% hydrogen peroxide for 5 min at room temperature. Wash slides 3 × 5 min with ddH₂O.
- 4) Follow procedure for pretreatment as required with TdT reaction buffer.

D. Suspension cells

- 1) Harvest cells and wash them twice in PBS using centrifugation (400g for 5 min) to remove residual protein.
- 2) Adjust the cell concentration to 4–5×10⁶ cells per mL in PBS.
- 3) Attach cells to slides using either the adhesion *or* centrifugation method
 - a. Adhesion method
 - a) Clean and label the slides.
 - b) Wash slides in PBS for 5 minutes at RT.
 - c) Place slides in a humidified box to prevent them from drying.
 - d) Place 20 to 50 µL of the cell suspension (at least enough to cover the well) in each well of the adhesion slides and let cells adhere at room temperature (RT) for 20 min.
 - e) Fix slides by 2% neutralized formalin for 15 min. Rinse slides with PBS then transfer to a Coplin jar containing ice-cold 70% ethanol for 1 h. Slides may be stored overnight in 70% ethanol at 4°C .
 - f) Inactivate endogenous peroxidases by covering sections with 2% hydrogen peroxide for 5 min at room temperature. Wash slides 3 × 5 min with ddH₂O.
 - g) Wash in PBS.
 - h) Follow procedure for pretreatment as required with TdT reaction buffer.
 - b. Centrifugation method
 - a) Assemble the CytoSpin centrifuge's sample chamber, filter card, slide, and racks according to the manufacturer's instructions.
 - b) Load 100 µL of cells in each sample chamber.
 - c) Centrifuge the slides at 600 rpm for 2 to 4 min.
 - d) Remove the slides from the rack and place them on a staining rack.
 - e) Fix slides by 2% neutralized formalin for 15 min. Rinse slides with PBS then transfer to a Coplin jar containing ice-cold 70% ethanol for 1 h. Slides may be stored overnight in 70% ethanol at 4°C
 - f) Wash in PBS. Inactivate endogenous peroxidases by covering sections with 2% hydrogen peroxide for 5 min at room temperature. Wash slides 3 × 5 min with ddH₂O.
 - g) Follow procedure for pretreatment as required with TdT reaction buffer.

2. TUNEL Reaction

- 1) Carefully blot away excess water then cover sections with TdT equilibration buffer for 10 min at room temperature.

2) Preparation of TdT reaction buffer:

	2 samples	4 samples	10 samples
TdT equilibration buffer	24 µl	48 µl	94 µl
TdT enzyme	0.5 µl	1 µl	2.0µl
Biotinylated dUTP	0.8 µl	1.6µl	3.2 µl

Mix well. Prepare fresh from stock solution prior to use.

- 3) Remove TdT equilibration buffer and cover sections with 10-12 µl of TdT reaction buffer. Incubate slides in a humidified chamber for 30 min at 37°C. In order to conserve reagents a reduced volume of TdT buffer may be carefully covered with a glass coverslip during the incubation. Take care to avoid trapping air bubbles which may lead to staining artifacts.
- 4) Stop reaction by incubating slides 2 × 10 min in 1×SSC.

3. Visualized with DAB

- 1) Rinse slides in PBS then block nonspecific binding by covering tissue sections with 2% BSA solution for 10 min at room temperature.
- 2) Rinse in PBS for 3×2min.
- 3) Detection: incubate sections with 3-4 drops of RTU streptavidin-HRP for 30 minutes at room temperature.
- 4) Rinse in PBS for 3×2min.
- 5) Chromogen/Substrate: incubate sections with 100-150µl (2-3 drops) of DAB solution for 2-8 minutes. Monitor signal development under a microscope.
Note: DAB solution is made by mixture of 20 µl of DAB stock solution and 20 µl of stable H₂O₂ solution with 1ml of DAB buffer.
- 6) Rinse in distilled water 2×2 min.
- 7) Counterstain: For using Hematoxylin Nuclear Counterstaining Kit (CAT#: VB-6004), incubate sections with 3 drops of RTU hematoxylin solution for 1-2 minutes. Rinse in tap water 2×2 min.
- 8) Dehydrate through 75% ethanol for 2 min, 95% ethanol for 2 min, and 100% ethanol for 2x3min. Clear in xylene for 2×5min.
- 9) Coverslip with mounting medium.

Negative Control: As a negative control, omit the TdT enzyme from the TdT reaction buffer.

Note: This product is intended for research purposes only. This product is **not** intended to be used for therapeutic or diagnostic purposes in humans or animals.

Precautions: Handle with care. Avoid contact with eyes, skin and clothing. Do not ingest. Wear gloves.