

Dual Apoptosis Assay Kit with NucView™ 488 Caspase-3 Substrate & Sulforhodamine 101-Annexin V (Texas Red®-Annexin V)

Catalog Number: 30030

Unit Size: 50 flow cytometry assays (200 µL assay volume).
Number of fluorescence microscopy staining assays may vary based on size of culture vessel and staining volume used.

Kit Contents

- NucView™ 488 Caspase-3 Substrate, 0.2 mM in DMSO, 250 µL
- Sulforhodamine 101-Annexin V (Texas Red®-Annexin V) 50 µg/mL in TE buffer, pH 7.5/0.1% BSA/0.1% Na₂S₂O₈, 250 µL
- 5X Annexin V Binding Buffer, 15 mL
- Caspase-3 Inhibitor Ac-DEVD-CHO, 2 mM in DMSO, 100 µL

Storage and Handling

Store at 4°C. When stored as directed the kit is stable for at least 6 months from the date it is received.

Product Description

NucView™ 488 and Sulforhodamine 101-Annexin V (Texas Red®-Annexin V) Dual Apoptosis Assay Kit allows simultaneous detection of caspase-3 activation and phosphatidylserine (PS) translocation in apoptotic cells by fluorescence microscopy, flow cytometry, or fluorescence plate reader.

Traditional fluorogenic caspase substrates require cell lysis and cannot be used to measure caspase activity in live cells; furthermore such assays measure only the average caspase activity in a cell population. Fluorescently-labeled caspase inhibitor assay (FLICA) reagents can enter live cells to detect caspase activity, but because FLICA probes are also irreversible caspase inhibitors, they cannot be used to follow caspase activity in real time. In contrast, NucView™ 488 Caspase-3 substrate detects caspase-3 activity within individual intact cells without interfering with caspase-3 activity. NucView™ 488 Caspase-3 substrate is a novel fluorogenic substrate consisting of a DNA dye and a DEVD substrate moiety specific for caspase-3. The substrate, which is initially non-fluorescent, crosses the cell membrane to enter the cytoplasm, where it is cleaved by caspase-3 to release a high-affinity DNA dye, which migrates to the cell nucleus to stain the nucleus with bright green fluorescence, allowing caspase-3 detection and visualization of apoptotic nuclear morphology in adherent or suspension cells.

During apoptosis, phosphatidylserine (PS) is translocated from the inner to the outer surface of the cell, allowing the dying cell to be engulfed by phagocytic cells. Annexin V is a 35 kD Ca²⁺-dependent phospholipid binding protein with a high affinity for PS. Fluorescent conjugates of annexin V can be used to label apoptotic cells by binding to PS exposed on the outer leaflet of the plasma membrane. The deep red fluorescence of Texas Red® (excitation/emission: 596/615 nm) is well separated from the green fluorescence of DNA-bound NucView™ 488.

References

- Cen H, et al. DEVD-NucView488: A novel class of enzyme substrates for real-time detection of caspase-3 activity in live cells. *FASEB J.* 22, 2243–2252 (2008).
- Martin, SJ, et al. Early redistribution of plasma membrane phosphatidylserine is a general feature of apoptosis regardless of the initiating stimulus: inhibition by overexpression of Bcl-2 and Abl. *J Exp Med.* 182(5):1545-56 (1995).

Assay Protocols

Controls:

We recommend that you performing the following controls:

1. Negative control with cells not induced to undergo apoptosis
2. Positive control with cells induced to undergo apoptosis
3. Inhibitor control with cells induced to undergo apoptosis and incubated with caspase-3 inhibitor prior to addition of NucView™ substrate

Assay Optimization:

The concentrations of NucView™ 488 substrate Ac-DEVD-CHO inhibitor provided in this protocol may require optimization depending on cell type. Optimal substrate concentration may vary between 1-10 µM. Use a final concentration of caspase-3 inhibitor Ac-DEVD-CHO that is at least 2-fold higher than the substrate concentration. Caspase-3 inhibitor Ac-DEVD-CHO is a reversible competitive inhibitor. In some cell types, caspase-3 inhibition may require the use of an irreversible caspase-3 inhibitor such as Z-DEVD-FMK, or may require addition of inhibitor before or during apoptosis induction.

For flow cytometry:

1. Induce apoptosis in cells by desired methods. Remember to include an untreated cell sample as a control.
2. Prepare 1X Binding Buffer by diluting 5X Annexin V Binding Buffer 1:5 with dH₂O.
3. Pellet cells and discard supernatant. For adherent cells, detach cells from culture plate with trypsin or other cell dissociation method prior to staining.
4. Resuspend cells in 1X Binding Buffer at a density of 10⁶ cells/mL.
5. Pipette 0.2 mL cell suspension into a flow cytometry test tube.
6. For inhibitor control sample, add 1 µL caspase-3 inhibitor Ac-DEVD-CHO directly to the cell suspension and mix well to obtain a final inhibitor concentration of 10 µM (see Assay Optimization).
7. Incubate cells with inhibitor at room temperature for at least 15 minutes.
8. Add 5 µL of 0.2 mM NucView™ 488 Caspase-3 substrate stock solution and 5 µL Sulforhodamine 101-Annexin V (Texas Red®-Annexin V) stock solution directly into cell suspension and mix gently. This yields a final NucView™ 488 Caspase-3 substrate concentration of 5 µM (see Assay Optimization).
9. Incubate cells at room temperature, protected from light, for 15-30 minutes.
10. Add 400 µL 1X Binding Buffer to each tube and measure fluorescence in FITC (excitation/emission 485/515 nm) and Texas Red® (excitation/emission 594/615) channels.

For fluorescence microscopy

1. Induce apoptosis in cells by desired methods. Remember to include an untreated cell sample as a control.
2. Prepare 1X Binding Buffer by diluting 5X Annexin V Binding Buffer 1:5 with dH₂O.
3. Wash cells twice with PBS.
4. For inhibitor control samples, add 1X Binding Buffer containing 10 µM caspase-3 inhibitor Ac-DEVD-CHO. Prepare a sufficient volume of buffer to cover cells.
5. Incubate cell with inhibitor at room temperature for 15-30 min.
6. Prepare assay solution by adding 5 µL of 0.2 mM NucView™ 488 Caspase-3 substrate stock solution and 5 µL Sulforhodamine 101-Annexin V (Texas Red®-Annexin V) stock solution per 100 µL 1X Binding Buffer. This yields a final NucView™ 488 Caspase-3 substrate concentration of 5 µM. For inhibitor control samples, include 10 µM inhibitor (see Assay Optimization). Prepare sufficient volume of solution to completely cover cells.
7. Incubate cells with assay solution at room temperature, protected from light, for 30 minutes.
8. Wash cells twice with 1X Binding Buffer.
9. Add a sufficient volume of 1X Binding Buffer to completely cover cells.
10. Observe fluorescence using FITC and Texas-Red® filter sets.

Optional: cells can be counterstained with Hoechst dye (catalog number 40046, 1 µM final concentration) to visualize cell nuclei. Formaldehyde fixation may be performed for long term preservation of cell staining. Annexin V binding to PS is calcium-dependent, therefore buffers used for fixation should contain 1.25 mM calcium chloride (CaCl₂).

Spectra

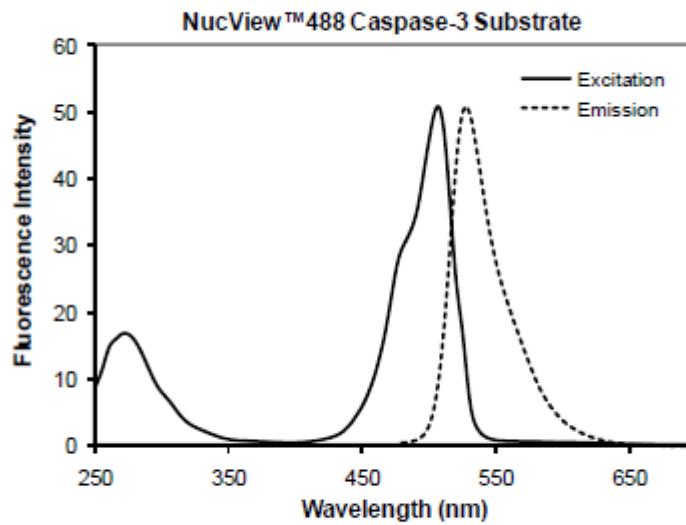


Figure 1. Excitation and emission spectra of enzymatically-cleaved NucView™488 caspase-3 substrate in the presence of dsDNA.

Related Products

Catalog number	Product
30067	Dual Apoptosis Assay Kit with NucView™ 488 caspase-3 substrate and CF594™-Annexin V
30065	Apoptosis & Necrosis Quantitation Kit Plus
30066	Apoptotic, Necrotic & Healthy Cells Quantitation Kit Plus
30060	CF™488A Annexin V and 7-AAD Apoptosis Kit
30061	CF™488A Annexin V and PI Apoptosis Kit
30029	NucView™ 488 Caspase-3 Assay Kit for live cells
30062	NucView™488 and MitoView™633 Apoptosis Kit
30001	JC-1 Mitochondrial Membrane Detection Kit
30063	CF™488A TUNEL Assay Apoptosis Detection Kit
30064	CF™594 TUNEL Assay Apoptosis Detection Kit