

# Product Information

## CF®488A Annexin V and 7-AAD Apoptosis Kit

**Catalog Number:** 30060

**Unit Size:** 100 assays

### Kit Contents

Component	Size
99902: 5X Annexin V Binding Buffer	3 x 15 mL
99946: CF®488A Annexin V	1 x 500 µL
99947: 7-AAD	1 x 20 µL

### Storage and Handling

Store at 4°C, protected from light. Do not freeze. Product is stable for at least 6 months from date of receipt when stored as recommended. 7-AAD binds nucleic acids, handle with universal laboratory safety precautions.

### Spectral Properties

CF®488A Annexin V Ex/Em: 490/515 nm

7-AAD Ex/Em: 548/648 nm (with DNA)

### Product Description

The CF®488A Annexin V and 7-AAD Apoptosis Assay Kit provides a convenient method for quantifying apoptotic (green) and necrotic (far-red) cells within the same cell population by flow cytometry or fluorescence microscopy.

Fluorescent conjugates of Annexin V can be used to label apoptotic cells. The human anticoagulant Annexin V is a 35-36 kilodalton, Ca<sup>2+</sup>-dependent phospholipid-binding protein with high affinity for phosphatidylserine (PS). In normal viable cells, PS is located on the inner leaflet of the cytoplasmic membrane. However, in apoptotic cells, PS is translocated from the inner to the outer leaflet of the plasma membrane, where it is available for binding to fluorescently labeled Annexin V, which can be detected by fluorescence microscopy or flow cytometry. Our CF®488A dye is superior to fluorescein/FITC as it is brighter, not affected by pH, and has much better photostability.

7-AAD (7-aminoactinomycin D) is a membrane-impermeant DNA-binding dye that is excluded by live cells and early apoptotic cells, but stains necrotic and late apoptotic cells with compromised membrane integrity. 7-AAD can be excited by the 488, 532, or 546 nm laser lines, but emits fluorescence in the far-red region, well separated from the green fluorescence of CF®488A.

Biotium also offers the CF®488A Annexin V and PI Apoptosis Kit, which works by the same assay principle, but features the red nucleic acid dye PI instead of 7-AAD (see related products).

Please visit our website at [www.biotium.com](http://www.biotium.com) for information on our life science research products, including environmentally friendly EvaGreen® qPCR master mixes, fluorescent CF® dye antibody conjugates and reactive dyes, apoptosis reagents, fluorescent probes, and kits for cell biology research.

Texas Red is a registered trademark of Thermo Fisher Scientific. Cy is a registered trademark of GE Healthcare.

Materials from Biotium are sold for research use only, and are not intended for food, drug, household, or cosmetic use.

### General Assay Considerations

1. This protocol was optimized using Jurkat cells treated with staurosporine to induce apoptosis. Assay optimization may be required for use with other inducing agents or other cell types.
2. Annexin V requires calcium for binding; usually concentrations near 2.5 mM are used. Annexin V Binding Buffer contains calcium.
3. If you prefer not to wash your cells, staining can be performed in cell culture medium with serum instead of Annexin V Binding Buffer, but the concentration of Annexin V may require optimization.

### Assay Protocols

#### Staining protocol for flow cytometry

1. Prepare a positive control by inducing apoptosis in your cells by the desired method. Include an untreated cell sample as a negative control. Also include samples for single-stained controls if compensation is required.
2. Dilute 5X Annexin V Binding Buffer 1:5 with distilled water. Prepare approximately 1 mL of 1X Binding Buffer for each sample to be stained.
3. Harvest cells after treatment by centrifugation and wash with PBS.
4. Centrifuge cells again, discard supernatant and resuspend cells at 5x10<sup>6</sup> to 10<sup>7</sup> cells per mL in 1X Binding Buffer.
5. Aliquot cells into flow cytometry tubes at 100 µL/tube.
6. Prepare a working solution of 7-AAD by diluting 1:10 in 1X Binding Buffer.
7. Add 5 µL of CF®488A Annexin V and 1-2 µL of 7-AAD working solution to each tube. Also prepare single stained compensation controls.
8. Incubate at room temperature for 15-30 minutes in the dark. The incubation can be carried out on ice to arrest the apoptotic process if desired.
9. Add 400 µL 1X Binding Buffer to each tube and analyze the cells by flow cytometry within 30 minutes of staining.
10. Detect CF®488A Annexin V in the FITC channel, and 7-AAD in the PE-Cy®5 channel.

### Related Products

Catalog number	Product
30061	CF®488A Annexin V and PI Apoptosis Kit
30065	Apoptosis and Necrosis Quantitation Kit Plus
99902	5X Annexin V Binding Buffer
29003-29085	Annexin V CF® Dye Conjugates
29003R-29085R	Annexin V CF® Dye Conjugates, Azide-Free, Lyophilized
10402	NucView® 488 Caspase-3 Substrate, 1 mM in DMSO
10405	NucView® 405 Caspase-3 Substrate, 1 mM in DMSO
10406	NucView® 530 Caspase-3 Substrate, 1 mM in DMSO
30062	NucView® 488 and MitoView™ 633 Apoptosis Kit
30063	CF®488A TUNEL Assay Apoptosis Detection Kit
30064	CF®594 TUNEL Assay Apoptosis Detection Kit
32002-32013	Live-or-Dye™ Fixable Viability Staining Kits
32010	Live-or-Dye™ NucFix Red Staining Kit
30020	ATP-Glo™ Bioluminescent Cell Viability Assay
30068	ViaFluor® SE 405 Cell Proliferation Kit