

Annexin V-FITC Early Apoptosis Detection Kit

✓ 1 Kit
(100 assays)



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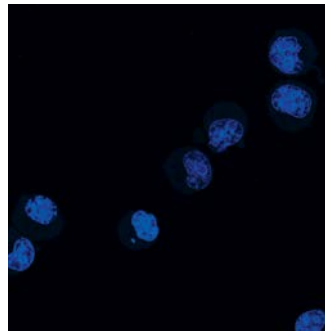
For Research Use Only. Not For Use In Diagnostic Procedures.

Products Included	Product #	Quantity
Annexin V-FITC Conjugate	4894	100 µl
10X Annexin V Binding Buffer	11732	2 x 1.5 ml
Propidium Iodide (PI) Solution	11733	1 x 1.3 ml

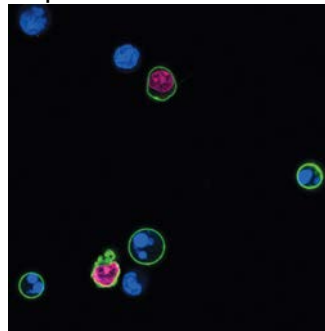
Description: The Annexin V-FITC Early Apoptosis Detection Kit enables researchers to identify early apoptotic cells within a cell population. Annexin V-FITC conjugated protein binds to cell surfaces expressing phosphatidylserine, an early apoptosis marker. Cells stained with propidium iodide (PI), a non-cell-permeable DNA dye, indicate necrotic cells. Cells stained with both PI and annexin V-FITC demonstrate later stage apoptosis and early necrosis. This kit provides enough reagent to perform 100 assays, based on a 250 µl assay volume.

Background: Annexin V belongs to the annexin family of proteins consisting of over 160 members. All annexin family members share the same characteristic of Ca²⁺-dependent binding to negatively charged phospholipid surfaces (1). During early apoptosis, phosphatidylserine (PS) is translocated from the cytosolic side of the plasma membrane to the cellular surface. This translocation exposes PS to the extracellular environment with the plasma membrane left intact (2). Annexin V protein demonstrates high affinity, specificity, and sensitivity for PS and can be used as a marker of early apoptosis. In order to rule out "leaky" necrotic cells, annexin V must be used in tandem with reagents that determine the integrity of the cell membrane, such as propidium iodide (2).

Untreated



Camptothecin-treated



Entrez Gene ID # 308
UniProt ID # P08758

Storage: The components within the Annexin V-FITC Early Apoptosis Staining Kit are stable for 1 year at 4°C. *Protect from light.*

Directions for Use:

1. Dilute the 10X Annexin V Binding Buffer with dH₂O to 1X and keep on ice.
2. Collect cells of interest by centrifugation. Wash cells with ice-cold culture medium or PBS.
3. Resuspend cells at 10⁵ to 10⁶ cells/ml with 1X Annexin V Binding Buffer. Aliquot 96 µl cell suspension in an assay tube.
4. Add 1 µl Annexin V-FITC Conjugate and 12.5 µl Propidium Iodide (PI) Solution to each 96 µl cell suspension. Allow cells to incubate 10 min on ice in the dark.
5. Dilute the cell suspension to a final volume of 250 µl/assay with ice-cold, 1X Annexin V Binding Buffer. Analyze immediately.

Background References:

- (1) Boersma, H.H. et al. (2005) *J Nucl Med* 46, 2035-50.
- (2) Vermes, I. et al. (1995) *J Immunol Methods* 184, 39-51.

◀ Confocal immunofluorescent analysis of live Jurkat cells, untreated (upper) or camptothecin-treated (lower), using Annexin V-FITC Conjugate (green). Red = Propidium Iodide (fluorescent DNA dye). Blue pseudocolor = DRAQ5® #4084 (fluorescent DNA dye).

◀ Flow cytometric analysis of Jurkat cells untreated (left) or treated with camptothecin (10µM, 4 hr; right) using Annexin V-FITC Early Apoptosis Detection Kit.

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The Annexin V-FITC conjugate is a product manufactured for Cell Signaling Technology by TAU Technologies BV.

