# Apoptosis (anti-PS) Biocytometry Kit



## **Assessment of Early Stage Apoptosis**

The Apoptosis (anti-PS) Biocytometry Kit offers a reliable method for quantification of early-stage apoptotic cells with exposure of phosphatidylserine (PS) on the outer leaflet of the cell membrane. The bioparticle system utilizes the well-characterized Annexin V moiety to identify target cells. Necrotic cells are selectively bypassed, enabling effective separation of apoptotic response to tested compounds from other events causing secondary necrosis, such as poor handling, pH imbalance, or temperature fluctulations.

### Sample types

compatible with many sample types:

- PBMCs
- cryopreserved samples
- primary cell cultures
- organoids
- immortalized cell lines

### Instrument requirements

requires only standard equipment:

- fixed-rotor centrifuge
- multimode plate reader (>1000 amol)
  - vortex
- 30°C incubator
- thermoshaker

### Workflow

- 1. add bioparticles & mix: 15 min
- 2. incubate: 5 hr, walkaway
- 3. add substrate & measure: 1 min

### **Performance Assessment**



#### Figure 1. Apoptosis quantification in Jurkat and HaCaT cells following treatment with apoptogenic agents.

(a) Time-course analysis using HaCaT and Jurkat cell monocultures. Cells were seeded at a density of 2,500 cells/well in RPMI and treated with 0.25 mM DTT to induce apoptosis. Incubation proceeded at 37°C in a 5% CO2 atmosphere for intervals of 2, 3, 4, or 5 hours. Apoptosis was quantified using Apoptosis (anti-PS) Biocytometry Kit following established preparation protocol. Signals were normalized to the number of cells, established prior to the biocytometric analysis. Analysis reveals differential sensitivity of HaCaT and Jurkat cell lines to DTT treatment. (b) Concentration-response curves for Jurkat cell line and camptothecin. Cells were seeded at a density of 2,500 cells/well in RPMI supplemented with 10% FBS and treated with varying concentrations of camptothecin (0 - 10  $\mu$ M). Incubation proceeded at 37°C in a 5% CO2 atmosphere for intervals of 2, 3, 4, or 5 hours. Apoptosis was quantified using Apoptosis (anti-PS) Biocytometry Kit following established preparation protocol. Signals were normalized to the number of cells, established preparation protocol. Signals were normalized to the number of cells, established preparation protocol. Signals were normalized to the number of cells, established preparation protocol. Signals were normalized to the number of cells, established preparation protocol. Signals were normalized to the number of cells, established prior to the biocytometric analysis. Notable onset of apoptosis is observed at the 3-hour mark.

# **Technical Specifications**<sup>1</sup>

### Sample

The Apoptosis (anti-PS) kit is designed to process samples of suspended cells and cell clusters. All intact cells in the sample will be processed.

100 µ
10,00
1

### **Target cells**

The Apoptosis (anti-PS) kit identifies target cells that are defined by presence of phosphatidylserine on the outer leaflet of the membrane. Such phenotype is a common predictor of early stage of apoptosis.

target antigen target phenotype necrotic cells phosphatidylserine early-stage apoptosis not quantified

<sup>1</sup> all values provided are approximate and for reference only. Specifications for samples and target cells may differ based on the application. For comprehensive guidelines on assays, workflows, and updates on new product releases, please visit our website at www.samplinghuman.com. For inquiries regarding custom solutions, high-throughput workflows, or licensing options, feel free to reach out to us at info@samplinghuman.com.