

Fluorescence-based apoptosis assay using CELENA[®] X

Key features

- High throughput screening of apoptosis using the CELENA[®] X high content imaging system.
- Optimized analysis pipeline to evaluate apoptosis.

INTRODUCTION

Apoptosis is a form of programmed cell death (PCD). It is a highly controlled cellular process that acts to eliminate unwanted and/or damaged cells during a cell's early growth and development. There are two pathways, intrinsic and extrinsic, that induce apoptosis. Despite being distinct pathways both the intrinsic and the extrinsic pathway require activation of caspase 3 and caspase 7. As a result, monitoring caspase 3/7 activity is used as a proxy for monitoring apoptosis. Here, we demonstrate the ability of the CELENA[®] X High Content Imaging System to monitor and quantify apoptosis using the CellEvent[™] Caspase-3/7 Green Detection Kit.

APPLICATION

Cell preparation

McCoy cells were counted using the LUNA-II[™] Automated Cell Counter, seeded at a density of 1×10^3 cells/100 μ L/well in a 96-well plate and grown overnight. Nuclei were counterstained with Hoechst 33342 (Life Technologies, H3570). Cells were washed twice using the cell culture media to remove excess dye before imaging.

Imaging and analysis

Apoptosis was evaluated by measuring the increase in Caspase-3/7 activity over time after inducing using CellEvent[™] Caspase-3/7 Green Detection Reagent (catalog number C10423; Life Technologies). The CellEvent reagent only becomes fluorescent when bound to DNA after cleavage by Caspase-3/7 which making it an ideal tool for monitoring apoptosis. Imaging and analysis were performed using the CELENA[®] X High Content Imaging System, CELENA[®] X Cell Analyzer software, and the CX Stage Top Incubator Pro [Tokai] (95 % humidity, 5 % CO₂, and 37 °C). Time lapse image acquisition was performed at T0 (time point) and then at 15-minute intervals for 2 hours. The cell culture media were exchanged with the fresh media containing 2 μ M CellEvent[™] Caspase-3/7 Green Detection Reagent and incubated for 30 minutes. Camptothecin was then added to cells with the final concentration of 0 μ M and 2.5 μ M to induce apoptosis. Live-cell imaging started immediately after adding Camptothecin. Images were captured using an Olympus 10X LWD high NA objective, a DAPI filter cube (Ex375/28, Em460/50) and an EGFP filter cube (Ex470/30, Em530/50).

Quantifying apoptotic events was done by measuring the area occupied by green fluorescence (Caspase-3/7 activity) occupied and the area occupied by blue fluorescence (Nuclei). Objects containing green and blue fluorescence were identified using the 'IdentifyPrimaryObjects' module. The area of fluorescence was measured using the 'MeasureImageAreaOccupied' module (Table 1). To visualize the data, the 'MaskImage' module was used to eliminate areas of low signal. The 'GrayToColor' module was used to convert the grayscale images to color images (Figure 1).

Order	Module	Use
1	<i>IdentifyPrimaryObjects</i>	Detects blue fluorescing Hoechst 33342
2	<i>IdentifyPrimaryObjects</i>	Detects green fluorescing CellEvent™ Green Detection Reagent
3	<i>MaskImage</i>	Eliminates cells with light levels below background
4	<i>MeasureImageAreaOccupied</i>	Measures area occupied by fluorescing CellEvent™ Green Detection reagent and Hoechst 33342
5	<i>GrayToColor</i>	Produces color images from gray scale

Table 1. Description of the pipeline used to assess apoptosis with CELENA® X Cell Analyzer

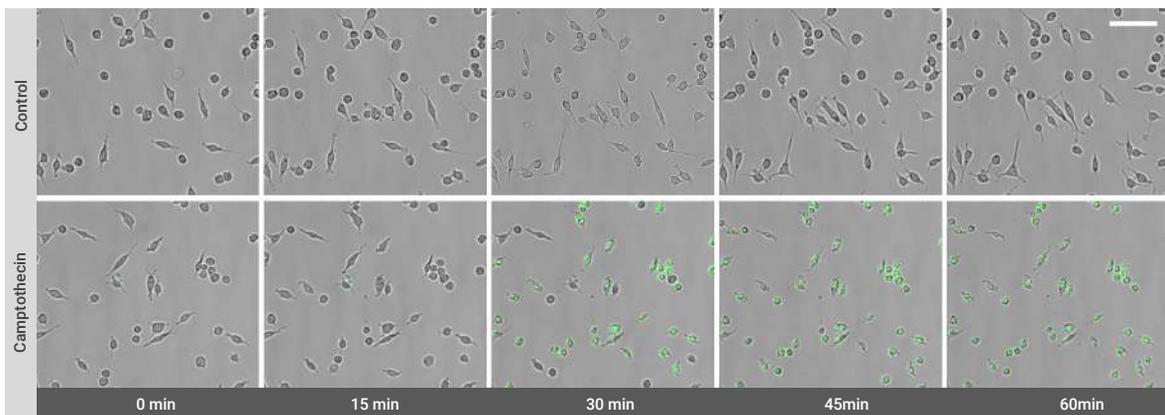


Figure 1. Time-lapse images acquired using CELENA® X system showing green fluorescence increasing over time (scale bar: 100 μ m).

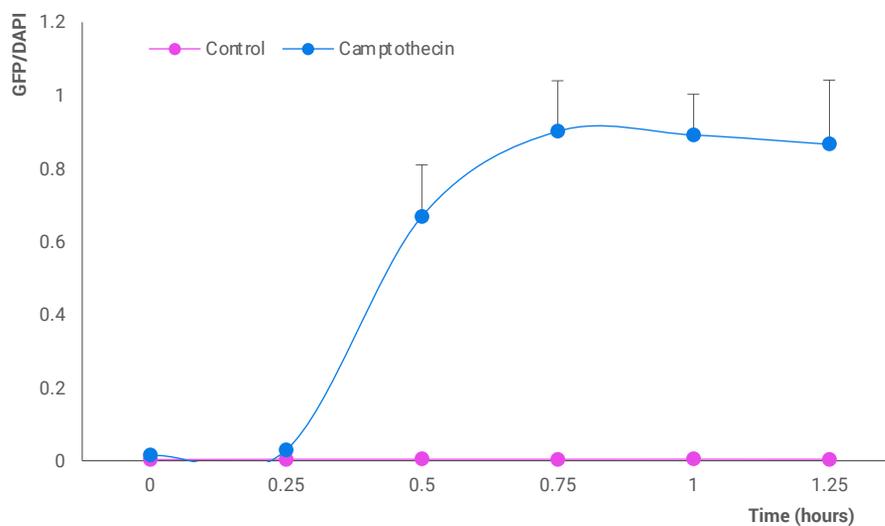


Figure 2. Quantification of apoptosis. The area occupied by green fluorescence (showing caspase activity) was divided by DAPI (showing Nuclei). Camptothecin treatment increases caspase activity over time.

CONCLUSION

In this study, we analyzed apoptotic events after Camptothecin treatment using the CELENA® X High Content Imaging System. Images were automatically taken and analyzed to identify objects, measure area, and produce images showing Caspase-3/7 activity. As shown in Figure 1 and 2, caspase activity was increased, and cells underwent apoptosis over time. Establishing the analysis pipeline was simple and straightforward. Overall, the CELENA® X High Content Imaging System, CELENA® X Cell Analyzer software, and CX Stage Top incubator proved to be an effective and efficient system for quantifying and monitoring apoptosis activity.

REFERENCES

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