## Background

Compound screening to assess cytotoxicity and to identify cancer targets is commonly performed using a highcontent imager, which is expensive, is overly complicated, and provides poor brightfi eld images making labelfree assays diffi cult. In addition the limited well coverage offered by these systems prevents their use in colony based assays.

## Synopsis

The Celigo Imaging Cytometer enables rapid, high quality, in situ, whole-well brightfi eld imaging for accurate label-free cell and colony analysis. In addition, three channel fl uorescence can be combined with brightfi eld for quantitative analysis of multiplexed assays. The Celigo is easy to use and offers a fl ow cytometry-like gating interface for optimal analysis of many different cell and colony types in multi-well plates (1536-well to 6-well) and T-fl asks (T-25, T75).

# Cancer Research Benefits

## **Tumor Sphere Analysis**

- Non-destructive quantification of live spheres for correlation with malignant cell behavior
- Rapid whole-well imaging and analysis of suspension spheres (analyze entire 12-well plates in <15 min)
- Analysis of sphere number, size, shape, and growth kinetics over time

## **Proliferation Assay**

- Accurate, whole-well brightfi eld imaging and segmentation provides counts of every cell in every well
- Label-free cell counting allows multiple reads of the same sample and requires no reagents
- Accurate normalization of wells using the actual number of cells in each well

## **Cell Cycle Analysis**

- Cell cycle analysis of adherent cells using assays developed for flow
- Scatter plot representation of DNA content/synthesis to identify each phase of cell cycle
- Easy-to-use interface that streamlines the application from image acquisition to data reporting

## Label Free Growth Tracking

- Determine growth characteristics of cells in situ, directly where they are grown
- Report growth curves, cell counts, confl uency, doubling time, and doubling rate for each well
- Quickly analyze cells growing in T-fl asks (T-75 fl asks can be analyzed in <15 min)

## Morphology Based Screening

- Development of cell-based assays relying on cell shape and morphology
- Rapid and label-free brightfi eld image acquisition and analysis (384-well plates can be analyzed in <5 min)

## **Multiparameter Gating**

- Interactive, real-time classifi cation of cell populations
- Unlimited number of histogram and scatter plots
- Combination of gates using a full set of boolean operators



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## **Tumor Sphere Analysis**



**Multiparameter Gating** 



