**Figure 1**

**a.** Overview of model inputs to *inSilicoCellModel* and output as pictures of cellular distributions colored according to where cells are in the cycle.

**b.** Adaptation from Drasdo-Hohme of how the model works.

**c.** Hierarchy of the model and what can be replaced.

**Figure 2**

**a.** Multiple growth rates in boundary and no boundary (show every curve)

**b.** Show density over time in both bounded and unbounded

**Figure 3**

Real Data from experiment plotted with points, simulation fits plotted with lines. PBS fitted from all simulations without drug effect, then initial density and cycle length are fixed and all drug simulations are fit with some drug effect. Fit done by minimizing L2.

**Figure 4**

**a.** Cartoon overview of how we simulate gene expression.

**b.** Illustrating how the scaling varies between 0 to 1 for each pathway.

**c.** Pick one dosage (10 um/mL) from **Figure 3** and plot each of the effective pathways vs time for that selection. Y-axis is % of pathway activity and all pathways can be plotted together. Show difference in pathways for drug effect.

**d.** Comparison to real gene expression data along each pathway.

**Figure 5**

**a** heatmap for 2 cell types w/ & w/o boundary (pick one)

**b.** Varying mean (normal) of drug effect w/o boundary in a simulation where you would typically observe a balance in the cellular populations.

**c.** Varying SD (normal) of drug effect w/o boundary where you would typically observe a balance in the cellular populations.

**d.** gene expression