

# ODE, RDE and SDE Models of Cell Cycle Dynamics and Clustering in Yeast

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## Abstract

Biologists have long observed periodic-like oxygen consumption oscillations in yeast populations under certain conditions and several unsatisfactory explanations for this phenomenon have been proposed.

We hypothesize that these oscillations could be caused by cell cycle synchronization or clustering. We develop some novel ordinary differential equation (ODE) models of the cell cycle. For these models, and for random and stochastic perturbations, we give both rigorous proofs and simulations showing that both positive and negative feedback are possible agents that can cause clustering of populations within the cell cycle. It occurs for a variety of models and a broad selection of parameter values in those models. These results suggest that the clustering phenomenon is robust and is likely to be observed in nature. Since there are necessarily an integer number of clusters, clustering would lead to periodic-like behavior with periods that are nearly integer divisors of the period of the cell cycle.

Related experiments have shown conclusively that cell cycle clustering occurs in some oscillating yeast cultures.

## 1 The Yeast Cell Cycle and Experimental Observations

Klevecz [13], McKnight [22], and others have observed marked oscillations in the dissolved  $O_2$  levels of various strains of *Saccharomyces cerevisiae* (yeast) under certain conditions. For instance in Figure 1 we show data from such an experiment for the yeast strain Cen.PK 113 [19]. It is seen in this experiment that dissolved  $O_2$  levels oscillate between about 5% to 55%, with a period of a little less than 4 hours. Since the doubling time for this culture was 7.8

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hours (as calculated from the dilution rate) it is natural to suppose that the doubling time and  $O_2$  oscillations are causally related.

Observations like these have been the subject of much speculation and analysis [3, 5, 8, 9, 10, 17, 22, 23]. Cultures exhibiting  $O_2$  oscillations, although planktonic, are very dense; the average distance between cells is on the order of 1 cell diameter [13]. This suggests the possibility of cell to cell signaling or media signaling in the cultures. Klevecz and others suggested that signaling is responsible for the  $O_2$  oscillations by setting up genomic synchronization. In experiments with the IFO0223 strain they showed that the compounds acetaldehyde and  $H_2S$  could be used to reset the phase of the oscillations [13, 14]. Other explanations of the oscillation involving various forms of signaling have been proposed. For instance, [10] suggested media signaling as a mechanism for “focusing” populations.

During aerobic growth, the cell cycle of *Saccharomyces cerevisiae* has four distinct phases, G1, S, G2 and M. The phase G1 begins at cell division and is characterized by growth of the cell. At the end of the G1 phase, which is thought to be triggered by a volume milestone, the cell enters the S phase. This is marked by the appearance of a bud and the beginning of replication of the DNA. Replication continues throughout the phase. Once replication is complete the cell enters a second growth phase G2. Most of the growth during this period takes place in the bud. The M phase is marked by narrowing or “necking” of the connection between the original cell and the bud and ends in cell division.

We hypothesize that the populations of cells in the S-phase can effect the growth rate of cell in the phase before the S-phase, either by positive or negative feedback.

We propose here that dissolved  $O_2$  oscillations in some strains could be due to a phenomenon that we will call *clustering*. In general we will define clustering loosely as significant groups of cells going through cell cycle milestones at approximately the same time, i.e. there is a weak form of temporal synchronization. The clustering could cause fluctuations in the  $O_2$  dilution levels by clusters passing in and out of high oxygen metabolism phases of the cell cycle, i.e. G1 phase.

In [19] it is shown for the Cen.PK 113 strain in experiments similar to those of [22] that bud index as well as cell density time-series (see again Figure 1) prove definitively that clustering occurs.

The goal of the present study is to show that various forms of signaling among cells, with either positive or negative feedback, robustly leads to clustering of cells within the cell cycle. We will consider several realizations of cell-cycle models, supposing that the population of cells in the S-phase effects the growth of cells in the later portion of the G1-phase of the cell cycle. We will denote this region by R.

We will prove mathematically in idealized models that clustering occurs and that the number of clusters formed depends on the widths of  $|S|$  and  $|R|$  of the S-phase and R. Of key importance is the necessity that if clustering occurs, then there are an integer number of clusters. This can provide the basis for a periodic-looking behavior with a period which is approximately an integer divisor of the cell-cycle length.

Models for which proofs are constructed include noisy or randomly perturbed models. We use simulations to demonstrate that the same type of behavior occurs in more detailed models.

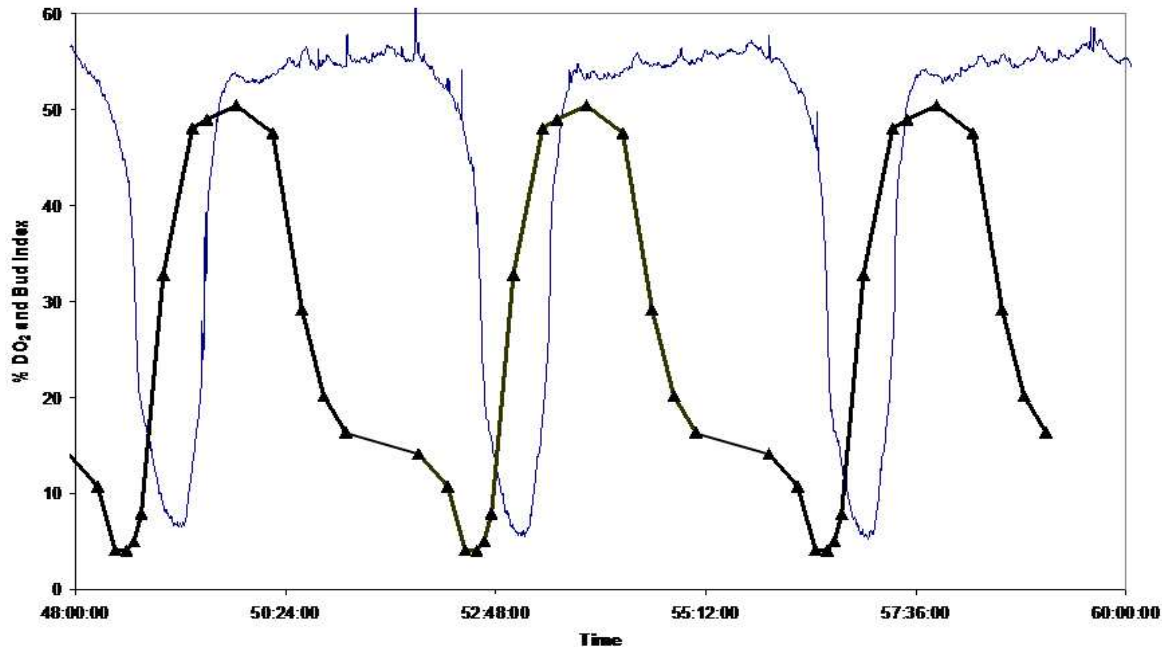


Figure 1: Dissolved  $O_2$  percentage and bud index percentage. The plot shows clearly that the bud index (percentage of cells with buds) are synchronized with the oscillation in the level of dissolved  $O_2$ . Moreover, time periods of sharp increases in bud index percentage show that a cluster of cell (consisting of almost 50% of the cells) is passing into S phase together. The doubling time of this culture is 7.8 hours. Two oscillations per 7.8 hour period also is clearly consistent with two clusters of cells.

In this paper we will focus primarily on models with the underlying assumption that the cell cycles of different generations are identical, or nearly so. Although this assumption is not justified in general yeast cultures, there is some evidence for it in the cultures under study. In conditions where the oscillations have been observed, the doubling time of the culture is extended significantly, by a factor of about 4. It is known that when the cell cycle is extended, most of the extension occurs in the G1 phase. This was confirmed by the authors in Cen.PK 113 experiments [19]. It is also known that much of the differences between the cell cycles of generations is due to differing lengths of the G1 phase (parent cells tend to divide with larger volumes than their daughter cells and so need less time to marshal sufficient resources to again undergo division). Thus by extending the doubling time, the G1 phases of all differing generations are stretched comparably, which decreases the relative differences of cell cycle lengths between generations. In related experiments on Cen.PK 113 [19], age distributions counts provide evidence that the assumption of equal cell cycle length for different generations is not too far off in that case.

In addition to models with the assumptions of identical or nearly identical cell cycles we also briefly treat models with small relative differences across generations and cases where even

larger differences could still lead to clustering. Elsewhere [20, 21], we undertake a careful study of a model with stratified generations under various assumptions and using biologically relevant values for all the parameters that can be determined. In all these models clustering emerges as a common and robust phenomenon.

In the following we begin with the most idealized models and add complexity to the models as we proceed.

## 2 Modeling of Cell Cycle Dynamics

### 2.1 General ODE model

We begin by defining a normalized logarithmic scale to represent the cell cycle. Customarily, the cell cycle is delineated by volume milestones. For a given cell, indexed by  $i$ , let  $v_i(t)$  denote its volume. It is usually assumed that the volume growth of a cell in a culture is proportional to its volume, i.e.

$$\frac{dv_i}{dt} = c_i(t)v_i$$

where the relative growth rate  $c_i(t)$  may depend on many factors, such as available resources, chemical composition of the culture substrate, etc. The growth rate may also be influenced by the state of the cell itself, thus the cell may react to environmental factors in differing way at different stages of its cycle. The environmental factors in turn may be influenced by the cells in the culture. Finally, the individual cells may have individual differences. We may include all of these factors into the rate  $c_i(t)$ .

Let  $V_{b,i}$  denote the volume of the cell at the beginning of its cell cycle and  $V_{d,i}$  its volume at division. Then consider the change of variables:

$$x_i = \frac{\ln(v_i/V_{b,i})}{\ln(V_{d,i}/V_{b,i})}.$$

We have then that  $x_i \in [0, 1)$  satisfies

$$\frac{dx_i}{dt} = c_i(t).$$

Next we may rescale time so that the average time span of a cell cycle in the culture is normalized to 1, i.e. scale by a factor  $\bar{c}(t)$  which is the average (over all cells) growth rate in a culture. In making this change of time we obtain that

$$\frac{dx_i}{dt} = \frac{c_i(t)}{\bar{c}(t)}.$$

If the variation between individual cells and their growth rate responses to the environment are not too great then the right hand side of the above equation is approximately 1. At this point, we distinguish between those influences on the growth rate that are common to all the cells in the culture and those due to individual differences. The common features we allow to depend on the state,  $x_i$  of the cell itself, as well as the conglomerate history of the whole

culture, which we denote by  $\bar{x}$ . We denote the common part of the growth rate as  $a$  and the individual part as  $g_i$ . We can thus write the equation as:

$$\frac{dx_i}{dt} = a(t, x_i, \bar{x}) + g_i(t). \quad (2.1)$$

The part of the growth rate due solely to individual differences in cells is contained in the term  $g_i$ . In most applications this term will be relatively small and “random” in the sense that differences are due to details of the cell process that are far too minute and complex to model. It contains both variations in the growth rate and differences in  $V_{b,i}$  and  $V_{d,i}$ . Thus we can actually view (2.3) as a Random Differential Equation (RDE) [1]. A reasonable approximation of this term in many circumstances is to replace  $g_i$  by a stochastic term:

$$\frac{dx_i}{dt} = a(t, x_i, \bar{x}) + \sqrt{\sigma} dW_i, \quad (2.2)$$

where  $dW_i$  represents an independent noise term. This Stochastic Differential Equation (SDE) must be interpreted in the usual way as an Ito integral equation. It is also reasonable under certain circumstances to consider (2.1) or (2.2) as perturbations of an Ordinary Differential Equation (ODE)

$$\frac{dx_i}{dt} = a(t, x_i, \bar{x}). \quad (2.3)$$

This model allows for variation of the growth on state within the cell cycle and the overall state of the system, but ignores differences between individual cells.

Thus far we have focused on the individual cell. Now we look at the culture as a whole. First of all, we note that in a culture, cells are constantly dividing, dying and perhaps being harvested. Thus the equations above must be applied to a changing set of cells, indexed by a changing finite set of positive integers  $S(t) \subset \mathbb{N}$ . If the  $i$ -th cell dies or is harvested, then  $i$  is dropped from  $S$ . As cell  $i$  divides, that index would be dropped from  $S$  and two new indices added. Alternatively, when a cell divides, one of the new cells could continue being denoted by  $i$ , with  $x_i$  resetting to 0 and the other (daughter cell) would be identified by a new index. This alternative would be appropriate in budding yeast where the mother and daughter cells are distinguishable. The conglomerate history  $\bar{x}$  would contain each  $x_i$  over its respective lifespan. Another way to keep track of this would be to let the index set  $S$  contain all indices for which cells are active at some time during an experiment and let  $x_i(t)$  assume some arbitrary value, such as  $-\infty$ , at times  $t$  outside the lifetime of the cell. With this convention the equations above could be coupled with stochastic processes that determine when harvest or death occur.

While these ODE models and perturbations described thus far are quite complex, they are not entirely intractable as we will see later. Now we turn our attention to partial differential equation approximations of the models.

## 2.2 General PDE model

Let  $U(x, t)$  denote the distribution of cells within the cell cycle as represented by  $x \in [0, 1]$ , i.e.

$$U(x, t) = \frac{1}{N} \sum_{i \in S(t)} \delta_{x_i(t)}(x)$$

where  $S(t) \subset S$  is the index set of cells that are living at time  $t$  and  $N$  might be taken as the average number of active cells. As is customary, we may approximate the point distribution by a more regular distribution  $u(x, t)$ .

If there is no death or harvesting, then by standard results, a distribution  $u(t, x)$  evolves locally under (2.2) by the Fokker-Plank equation, which takes the form:

$$\frac{\partial u}{\partial t} + \frac{\partial}{\partial x} (a(x, t, [u]) u) - \sigma \frac{\partial^2 u}{\partial x^2} = 0. \quad (2.4)$$

In this notation  $[u]$  denotes a functional dependence on  $u$  and possibly its history, for example via an integral operator. If the culture is well mixed and harvesting is via removal of bulk material, as in a bio-reactor, then all cells are equally likely to be harvested, regardless of state and the effect of harvesting on the density will be proportional to  $u(t, x)$ . Similarly, if death of a cell is equally likely at any stage of the cell cycle then the harvesting and death can be incorporated into the Fokker-Plank equation as a term  $-ku$  on the right hand side, i.e.:

$$\frac{\partial u}{\partial t} + \frac{\partial}{\partial x} (a(x, t, [u]) u) - \sigma \frac{\partial^2 u}{\partial x^2} = -ku. \quad (2.5)$$

To take cell division into account in the PDE model, we need to require the boundary conditions:

$$u(t, 0) = 2u(t, 1^-), \quad u_x(t, 0) = 2u_x(t, 1^-).$$

### 2.3 Periodic or Near Steady State

Next we suppose that the culture is growing in a bioreactor near a steady state or a periodic state so that the cell division is approximately balanced by harvesting. Suppose also that dependence of  $a$  on all variables besides  $x$  is small, or, that the dependence on  $t$  is averaged over one period, and that  $\gamma$  is small. Then equation (2.5) may be approximated by:

$$\frac{\partial u}{\partial t} + \frac{\partial}{\partial x} (a(x) u) = -ku.$$

If we make the change of variables:

$$u(t, x) = e^{-k \int_0^x \frac{1}{a(y)} dy} z(t, x),$$

then an easy calculation shows that  $z(t, x)$  satisfies:

$$\frac{\partial z}{\partial t} + \frac{\partial}{\partial x} (az) = 0. \quad (2.6)$$

Further, the balance between growth and harvesting implies:

$$k = \frac{\ln 2}{\int_0^1 \frac{1}{a}}.$$

This implies that the boundary conditions on  $z$  are

$$z(t, 0) = z(t, 1).$$

The ordinary differential equation that generates (2.6) is simply:

$$\frac{dx}{dt} = a(x),$$

on  $[0, 1)$ . If we reincorporate the other influences on the growth rate then the unperturbed and perturbed equations become:

$$\frac{dx_i}{dt} = a(t, x_i, [\bar{x}]), \quad (2.7)$$

and

$$\frac{dx_i}{dt} = a(t, x_i, [\bar{x}]) + g(t)_i. \quad (2.8)$$

Note that (2.6) is a conservative equation in the sense that  $\int z dx$  is preserved, thus in the ODE (2.7) and RDE (2.8) the total number of cells considered should not change. It is thus reasonable to take  $S$  to be a fixed set  $\{1, 2, \dots, N\}$  and let  $x_i(t)$  to be defined for all  $t \in [0, T]$ . To accommodate division we may reset  $x_i(t)$  to 0 each time it reaches 1. An interpretation of this model is the following: Given a living cell at time  $t$ , the *expected number* of cells descended from that cell at any later time  $t + n$  is exactly 1. Thus the original model with a changing index set  $S$  can be replaced by a fixed system, each variable  $x_i(t)$  representing the state of its expected descendant at time  $t$ .

With a fixed index set, this model is easily amenable to numerical investigation with currently available computer speeds. For instance, one can investigate the behavior of a conglomeration of 10,000 or more cells over several cell cycles on a desktop workstation. Further, under some additional assumptions on the dependence of  $a$  and  $g_i$ , we can investigate properties of the solutions of (2.7) and (2.8) rigorously as we show below.

## 2.4 Modeling of the growth term $a$

The standard biological assumption on  $a$  is that it does not depend on  $x_i$  i.e. it is independent of the cell's current state within the cycle. Rather it depends mostly on the nutrients available and other environmental factors. With these assumptions  $a = a(t, \bar{x})$  in the ODE models and  $a = a(t, [u])$  in the PDE models.

In recent studies it has been shown that observed behavior is consistent with a more sophisticated form  $a$ , in which the location of cells in cell cycle may influence the growth rate of themselves and other cells, i.e.

$$a = a(t, x_i, \bar{x}).$$

For example in budding yeast it has been hypothesized that cell in the S phase may promote or inhibit the growth of cells in the pre-budded G1 phase. In the PDE models we may represent such influence through integral operators:

$$a(t, x, [u]) = b(t, x) + \int_0^1 k(t, x, u(z), z) dz.$$

In the ODE model the continuous distribution  $u$  can be replaced by the point distribution  $U(t)$  in the integral. In an idealized version the dependence on  $U$  was taken to be in the form of counting the number of cells in a certain portion of the cell cycle (in this case the "S" phase of the yeast). This number can be assumed to influence the growth of cells just preceding that phase. This model is amenable to exact analysis and we explore it in the next sections.

### 3 Idealized Models and Clustering

#### 3.1 Idealized models

The S-phase of the cell can be represented by a subinterval  $S = [s, t] \subset [0, 1]$ . In our models the population of cells in the S-phase will effect the growth of cells in a preceding portion of the cell cycle which we denote by  $R = [r, s]$ , i.e. a portion of the G1-phase. We will consider a finite population of  $N$  cells, each of them travelling with the speed

$$\frac{dx_i}{dt} = \begin{cases} 1 & \text{if } x_i \notin R \\ 1 + F(\#\{\text{cells in } S\}) & \text{if } x_i \in R. \end{cases} \quad (3.1)$$

In this section we will consider two idealized (and discontinuous) forms of  $F$ . By the *advance model* we will mean that if the fraction of cells in  $S$  exceeds some threshold  $\tau$ , then all cells in  $R$  are instantaneously promoted to the beginning of  $S$ , from which they resume normal growth. This corresponds to a limit  $F \rightarrow \infty$  in (3.1) when  $\#\{\text{cells in } S\} \geq \tau$  and  $F = 0$  when  $\#\{\text{cells in } S\} < \tau$ .

By the *delay model* we mean that if the fraction of cells in  $S$  equals or exceeds some threshold  $\tau$ , then cells at  $s$  are delayed from proceeding into the S-phase, until the fraction of cells in  $S$  drops below the threshold. This corresponds to choice  $F = -1$  when  $x_i = s$  and  $\#\{\text{cells in } S\} \geq \tau$  and  $F = 0$  when  $\#\{\text{cells in } S\} < \tau$ . If more than  $\tau$  cells have accumulated at  $s$ , then all of those cells will enter  $S$  together when the fraction of cells drops below  $\tau$ .

These idealized models have an advantage of being analytically accessible, thus allowing an insight into more realistic models, that we later explore numerically.

#### 3.2 Clustering

Under either the advance or delay models, it is clear that some cells may become synchronized. Consider the advance model; whenever the fraction of cells in  $S$  exceeds the threshold  $\tau$ , then all cells in  $R$  are instantaneously synchronized at  $s$ . There being no mechanism in the model to subsequently differentiate them, they will remain synchronized from then on. Similarly, for the delay model, during a period when the threshold is exceeded, all cells arriving at  $s$  will be thereafter synchronized. We will call a group of synchronized cells a *cluster*. If the number of cells in a cluster is large enough to exceed the threshold of the model, we call it a *critical cluster*.

We will denote by  $\lfloor x \rfloor$  the largest integer smaller than  $x$  and by  $\lceil x \rceil$  the smallest integer larger than  $x$ .

Given an initial (discrete) distribution  $\bar{x}(0)$  or  $w(s) = \sum_{i=1}^N \delta_{x_i(0)}(s)$  the evolution of such distribution under either model will be called a *trajectory* starting at  $w(s)$  and denoted by  $\phi(t, w(s))$ . It is clear that different trajectories can merge.

A trajectory is called an *equilibrium* if it is stationary in the coordinate frame moving with the speed 1. In other words,  $\phi(t, w(s)) = w((x - t) \bmod 1)$  for all  $t \geq 0$ . A *periodic orbit* is a trajectory which is periodic in the moving frame. In such a case there exists  $T$  with

$\phi(T + t, w(x)) = \phi(t, w(\phi(x)))$  for all  $t \in [0, T]$ ; if  $T$  is the smallest number with this property, it is called a *period*.

### 3.3 Advance model

**Theorem 3.1** *Consider the advance model.*

- A.** *If the initial  $w(x)$  exceeds threshold on all intervals of length  $|S|$ , except possibly inside the interval  $R$ , then the trajectory converges to a periodic orbit.*
- B.** *Any initial  $w(x)$  that is below threshold on all intervals of length  $|S|$ , is an equilibrium point.*
- C.** *Every other initial condition converges to an equilibrium  $e$  with a finite number of critical clusters, separated by voids of length at least  $|R| + |S|$ .*

**Proof:** Let

$$q(t, x) = \int_{[x, x+|S|] \bmod 1} \phi(t, w(z)) dz. \quad (3.2)$$

The function  $q(t, x)$  is a form of local density of the discrete distribution of cells. Observe that if  $q(0, x) < \tau$  for all  $x$  then this is a fixed point of the dynamics. This corresponds to the case **B** above.

The case **A** corresponds to the case when  $q(0, x) \geq \tau$  for all  $x$ , except for  $x \in R$ . At  $t = 0$  all cells in  $R$  are advanced to the point  $s$ . Since the  $q(0, x) \geq \tau$  for all  $x$  the cells arriving at  $r$ , the start of the interval  $R$ , are immediately transferred to the point  $s$ . This continues indefinitely. This trajectory is a periodic orbit in the moving frame.

Now we do the analysis of the case **C**. Let  $q(i, x)$ ,  $i = 0, 1, \dots$  be the mass function after  $i$  passes through the cell cycle. We decompose the domain  $I = [0, 1]$  into intervals  $I_1^i \cup J_1^i \cup \dots \cup J_{n-1}^i \cup I_k^i$  where  $q(i, x) \geq \tau$  for all  $x \in I_j^i$  and  $q(i, x) < \tau$  for all  $x \in J_j^i$ . Set  $I_j^i := [d_j^i, c_j^i]$ . We compare the intervals  $I_j^i$  from iteration to iteration. There are several possibilities that can happen to  $I_j^i$  after a passage through a cell cycle:

1. The interval  $I_j^i$  will shorten by  $|R|$  and  $I_j^{i+1} = [d_j^i + |R|, c_j^i]$ .
2. The intervals  $I_j^i, \dots, I_{j-k}^i$  may merge, if all intervals in between fall within distance  $|R|$  i.e.  $c_j^i - c_{j-k}^i < |R|$ ;
3. If an interval  $I_{j-1}^i$  follows the interval  $I_j^i$  by a distance closer than  $|R| + |S|$  i.e.  $d_j^i - c_{j-1}^i \leq |R| + |S|$ , then the interval  $I_{j-1}^i$ , or a portion of that interval, will be promoted across  $R$ . The resulting distance between the two new intervals will be less than  $|S|$ . After each subsequent pass the distance between these two intervals will shorten by  $|R|$  and in finite number of steps they will be within  $|R|$  of each other and they will merge. The number of steps this takes is uniformly bounded above by  $l := \frac{|S|}{|R|} + 1$ . This process may result in a split of an interval  $I_j^i$  into two intervals with the total mass conserved

in the transaction. Notice that even though temporarily the number of intervals  $I_j^{i+1}$  may increase over the number  $I_j^i$ , after  $k$  subsequent passes through  $S$  the number of intervals  $I_j^{i+k}$  is less or equal to number of intervals  $I_j^i$ .

4. The interval  $I_j^i$  may shorten, if the preceding interval  $I_{j+1}^i$  has critical mass in  $S$  and therefore promotes part of the mass ahead of  $c_j^i$  across  $R$ . Since a mass ahead of  $c_j^i$  is lost,  $c_j^{i+1} < c_j^i$ . The difference between this case and case (3.) is that here  $|R| + |S| \leq d_j^i - c_{j-1}^i \leq |R| + |S| + |S|$ . Observe, that the transferred cells will not form a new interval  $I_j^{i+1}$ , since they do not have a sufficient mass.

In summary, the number of intervals  $I_j^i$  is greater or equal to the number of intervals  $I_j^{i+k}$  for every  $i$  and the length of each intervals is a non-increasing function of  $i$ . Finally, it is easy to see that the only interval  $I_j^i$  that will not change under the advance operator, is a singleton  $I_j^i = [c_j^i, c_j^i]$ . Since the population is finite, after a finite number of passes through  $S$ , each  $I_j$  is a singleton. If two such singletons follow each other closer then  $|R| + |S|$ , by the step 3. above, they will merge in finite number of steps. Eventually all remaining singletons  $I_j$  are followed by a empty void of length  $|R| + |S|$ .

□

**Corollary 3.2** *In the advance model no more than  $\lfloor (|R| + |S|)^{-1} \rfloor$  critical clusters can persist.*

**Proof:** By the previous Theorem each critical cluster has to be followed by gap of length at least  $|R| + |S|$ . □

### 3.4 Delay model

**Theorem 3.3** *In the delay model no more than  $\lceil |S|^{-1} \rceil$  critical clusters can persist.*

**Proof:** When two consecutive critical clusters pass through  $S$ , the second cluster must wait at  $s$  until the first cluster has passed  $t$ . Thus the distance between them must then be at least  $|S|$  and will remain at least  $|S|$  until the first cluster hits  $r$ . Then the first cluster may have to wait to enter  $S$ , possibly decreasing the distance between the two clusters in question.

Assume that the number  $n$  of persistent clusters is constant along some trajectory. Then either all these clusters are separated by a distance at least  $|S|$ , or there are  $n - 1$  clusters separated by a distance  $|S|$  and there is an additional cluster inside  $S$  whose distance to a waiting cluster at  $s$  is smaller than  $|S|$ .

This minimal spacing implies that

$$(n - 1)|S| \leq 1.$$

The result then follows. □

We have shown that a cluster cannot persistently follow a critical cluster closer than  $|S|$  (except while the critical cluster is being delayed at  $s$ ). Also, it is clear that as many as  $\lceil |S|^{-1} \rceil$  critical clusters may persist if the overall number  $N$  of cell satisfies:

$$N > \tau \lceil |S|^{-1} \rceil.$$

Recall the definition of  $q(t, x)$  in (3.2).

**Theorem 3.4** *In the delay model, if the cells are initially distributed so that the density  $q(x, 0)$  everywhere exceeds twice the threshold, then exactly  $\lceil |S|^{-1} \rceil$  clusters will develop and persist.*

**Proof:** Cells at  $s$  are initially delayed from entering  $S$  until the fraction of cells in  $S$  drops below the threshold. Since  $d(|S| - t) = \tau$ , this will occur at

$$t = |S| - \frac{\tau}{d}.$$

At this time the cells that have clustered at  $s$  number  $dt$  or  $d|S| - \tau$ . Since  $d|S|$  is assumed to be at least twice  $\tau$ , the threshold is again exceeded as the cluster enters  $S$ . The threshold will continue to be exceeded until this cluster leaves  $S$ . At this time a new cluster will have formed at  $s$  with volume  $d|S|$  which is by assumption above  $\tau$ . The second cluster is spaced exactly distance  $|S|$  behind the first cluster. At third cluster, etc. will form in the same way until the first cluster returns to  $s$ . At that time there is either a cluster in the interior of  $S$  or at  $t$ . In the former case the first cluster will wait less than time  $|S|$  to enter  $S$ , ahead of when the second cluster arrives. In the latter case it enters  $S$  immediately. This scenario will repeat itself indefinitely and the number of critical clusters thus produced is an easy calculation.  $\square$

## 4 Small Perturbations of the Advance and Delay Models

### 4.1 Small Perturbations

Next we consider *small perturbations* of the advance and delay models by which we mean that the progression of each individual cell is independently perturbed by a small term which is independent of the other cells.

$$\frac{dx_i}{dt} = \begin{cases} 1 + \epsilon \xi_i(x_i, t) & \text{if } x_i \notin R \\ 1 + F(\#\{\text{cells in } S\}) + \epsilon \xi_i(x_i, t) & \text{if } x_i \in R, \end{cases} \quad (4.1)$$

where  $|\xi_i(x_i, t)| \leq 1$  and  $\epsilon$  is small. We will take as our definition of small that  $\epsilon$  is smaller than  $\min(|S|, |R|)/6$ . The effect of the perturbation is to cause initially synchronized cells to drift from each other, but by no more than  $2\epsilon$  or  $\min(|S|, |R|)/3$  within a single cell cycle. Note that under these assumptions (4.1) is a random differential equation with bounded noise. For general results about RDE with bounded noise see [11].

Note that this model can viewed as a relaxation of the assumption that cells are indistinguishable. It also can be considered as incorporating random variations and noise.

With these perturbations, clusters will not remain precisely synchronized as in the unperturbed models, but rather will spread out as the cells proceed through the cell cycle. We now expand our definition of cluster to include any group of at least  $\tau$  cells that are within  $|S|/3$  of each other in the cell cycle.

We will assume below that the small perturbations act like noise in the ways we define below.

We will say that perturbations satisfy the *maximum principle* if

$$\max_x q(t_1, x) > \max_x q(t_2, x)$$

and

$$\min_x q(t_1, x) < \min_x q(t_2, x)$$

for any  $t_1 < t_2$  that are not separated by an activation or deactivation of delay or advance. We will say that the perturbations are *diffusive* if:

- the perturbations satisfy the maximum principle.
- synchronized cells will immediately de-synchronize.

## 4.2 Perturbed Delay Model

**Theorem 4.1** *In the delay model with small diffusive perturbations such that  $\epsilon < |S|/6$ , and an initial distribution of cells that satisfies  $q(0, x) > 2\tau + 2\epsilon$  for all  $x$ , the system will form either  $\lfloor |S|^{-1} \rfloor$ ,  $\lceil |S|^{-1} \rceil$ , or,  $\lceil |S|^{-1} \rceil + 1$  critical clusters within one cell-cycle period and these clusters will persist indefinitely.*

**Proof:** Denote  $d = \min q(0, x)$ . Cells at  $s$  are initially delayed from entering  $S$  until the fraction of cells in  $S$  drops below the threshold. This will occur at time  $t_0$  no smaller than

$$t_0 = \frac{|S| - \frac{\tau}{d}}{1 + \epsilon}.$$

At this time the cells that have clustered at  $s$  number at least  $t_0 d$  or  $(d|S| - \tau)/(1 + \epsilon)$ . This group of cells we will call cluster 1. By the assumptions, the threshold is again exceeded as the cluster enters and crosses  $S$ . While crossing  $S$  the cluster will de-synchronize, but all the cells will remain in  $S$  for at least  $t_1 = |S|/(1 + \epsilon)$  and no longer than  $|S|/(1 - \epsilon)$ . Thus while the first cluster crosses  $S$  at least  $t_1 d$  or  $d|S|/(1 + \epsilon)$  cell accumulate at  $s$ . This group we call the second cluster and from the assumption  $q(0, x) > 2\tau + 2\epsilon$  it is critical. Its crossing time  $t_2$  will again be bounded below by  $|S|/(1 + \epsilon)$  and above by  $|S|/(1 - \epsilon)$  and so this process of cluster formation will continue for as long as a density of cells at least  $d$  is arriving at  $s$ .

By a straight-forward calculation, the gap between any two adjacent clusters formed in this process is bounded below by:

$$|S| \frac{1 - \epsilon}{1 + \epsilon}$$

and obviously the gap is bounded above by  $|S|$ . If the first of two adjacent clusters travel at the minimal speed  $1 - \epsilon$  while outside of  $S$ , all the cells in the first cluster must arrive again at  $s$  no later than:

$$t^* = \frac{1}{1 - \epsilon} - \frac{|S|}{1 + \epsilon},$$

after cluster 2 leaves  $S$ . During the same time period the second of the adjacent clusters can travel at a rate at most  $1 + \epsilon$  and thus can travel a distance of no more than

$$d^* = t^*(1 + \epsilon) = \frac{1 + \epsilon}{1 - \epsilon} - |S| < 1.$$

Therefore the entirety of the first cluster must reach  $s$  before any of the second cluster does so. The gap at that time is at least

$$1 - d^* = |S| - \frac{2\epsilon}{1 + \epsilon} > \frac{2|S|}{3}.$$

Next consider the relative motion between the cells that began in  $S$  and the cells in cluster 1. Note that the initial gap will be at least  $|S|/2$ . Since  $\epsilon < |S|/4$ , the last of the cells will reach  $s$  before the first cluster returns.

When cluster 2 leaves  $S$ , it is easily shown that the gap between cluster 1 and 2 is no wider than

$$|S| \frac{1 + \epsilon}{1 - \epsilon} < |S|(1 + 2\epsilon).$$

At the same time the gap between cluster 1 and 3 is no more than the gap between cluster 1 and 2 plus  $|S|$ . That is,

$$2|S|(1 + \epsilon).$$

When cluster 3 reaches  $t$ , the gap between clusters 1 and 3 is no more than

$$2|S|(1 + \epsilon) + |S| \frac{1 + \epsilon}{1 - \epsilon} - |S| < 2(1 + 2\epsilon)|S|.$$

By induction one can show that the first cluster can be no more than

$$(j - 1)(1 + 2\epsilon)|S|$$

ahead of the  $j$ -th cluster when the  $j$  cluster is at  $t$ . From this we can show that at least  $\lfloor |S|^{-1} \rfloor$  clusters will form.

On the other hand, when cluster 2 leaves  $|S|$  the gap between clusters 1 and 2 must be at least

$$\frac{2}{1 + \epsilon}|S|$$

and cluster 1 must be at least

$$\frac{j - 1 - \epsilon}{1 + \epsilon}|S|$$

ahead of the  $j$ -cluster. It is clear then that  $j$  must be less than  $\lfloor |S|^{-1} \rfloor + 1$ .

Now to show that the clusters persist. Clearly, while cluster 1 is passing through the cycle the first time, there is always a critical cluster in  $S$ , at least until the cells that began in  $S$  again reach  $S$ . Note that the cells which were inside  $S$  when the first cluster was released from  $s$  are critical in number. We will call this group of cells the tail. When the tail begins to arrive back at  $S$ , its leading edge can be no more than

$$\frac{|S|(1 - |S|)}{1 - \epsilon}$$

ahead of cluster 1. At that moment there will either be (1) a critical in the interior of  $S$ , or, (2) there are critical clusters at both ends of  $S$ . Let us consider case (2) first. In this case the cluster at the beginning of  $S$  will be critical and as it enters  $S$  the cells from tail will be

delayed at  $s$ . Since the original width of the tail is less than  $|S|/2$ , its width when it reaches  $S$  will be less than  $2|S|/3$  and it will all reach  $S$  before the cluster ahead of it leaves  $S$ . Now two things can happen, both of which continue the process, either all or part of cluster 1 reaches  $S$  while the tail is delayed there, or it will be blocked by the tail. In the former situation part or all of cluster 1 will merge with the tail and be delayed and in the later the entirety of cluster will be delayed by the tail. Either way cluster 1 is delayed and will be inside  $S$  when cluster 2 arrives back. Now in case (1) all or part of the tail will reach  $S$  and be delayed while the previous cluster clears  $S$ . It will merge with other cells delayed at  $S$  and they will be a critical cluster when they are released. Thus when cluster 1 arrives at  $S$ , the last cluster including part or all of the tail will still be in  $S$ . Again in this case cluster 1 is delayed before reentering  $S$  and will be inside  $S$  when cluster 2 arrives. Now in either case cluster 1 is not only delayed, but resynchronized as it is delayed. The same happens for the subsequent clusters and the process continues.  $\square$

### 4.3 Perturbed Advance Model

In the perturbed advance model we make the following observation: *In the advance model with small random perturbations and  $|R| = |S|$ , an initial single critical cluster will have “forward leakage.”* Wherever a cluster begins in the cell cycle, when its edge reaches  $s$  it will have positive width. As some of the cells enter  $S$  the threshold will be exceeded, and the rest of the cells will be promoted to  $s$ . The cluster will then proceed around the cell cycle. When its edge reaches  $S$  again, its width will be larger and the leading edge flatter than during the previous cycle. Thus the front edge will reach deeper into  $S$  before the advance takes place. Thus the front edge of the cluster will grow wider and wider on each pass until some of the cells are able to pass through  $S$  completely before advance occurs. At this point these cells have essentially escaped from the cluster.

Note that the cells which leak forward may eventually slow down and drop back into the cluster, or, if they continue at a faster rate than the cluster will eventually be caught in the next cluster (or into the original cluster if it is the only one.) This implies that there may be steady states which have persistent clustering, but the cells in the clusters may not actually be synchronized in the strict sense.

## 5 More detailed models and simulations

### 5.1 Graduated but unstratified model.

Next we consider more realistic models. Rather than strict advance and delay we consider the possibility that the number of cells in  $S$  either act to slow down (inhibit) or speed up (accelerate) cell growth in the preceding region  $R$ . The population of cells in the S-phase is hypothesized to effect the growth of cells in a preceding phase  $R = [r, s]$  via a continuous function. This influence will be assumed to be a function of the number of cells in the  $S$  phase.

Thus we have the following equations of motion:

$$\frac{dx_i}{dt} = \begin{cases} 1 + \sqrt{\sigma}dW_i & \text{if } x_i \notin R \\ 1 + F(\#\{\text{cells in } S\}) + \sqrt{\sigma}dW_i & \text{if } x_i \in R. \end{cases} \quad (5.1)$$

Note that this form is quite general since we might consider any functional dependence  $F$  and  $R$  could be large or small.

We report simulations of this model where we will assume  $F$  to be linear function  $aS$ , with  $a < 0$  for the delay model and  $a > 0$  for the advance model. For example a nonlinear model would be obtained if  $F$  was a sigmoidal Hill function  $F(x) = \frac{A}{1+(x/\tau)^n}$ . In the delay case in the limit of Hill coefficient  $n \rightarrow \infty$  we obtain a strict delay model described in section 2.

In the simulations we also incorporate a small diffusive term (white noise) with variation  $\sigma$ . The program tracked the trajectories of 5000 individual cells, initially uniformly distributed, through 20 unperturbed cell cycles (20 units of time in equation 5.1). In each of Figures 1-4 we show (a) a histogram of the final distribution of cells within the cell cycle and (b) a time-series of the fraction of cells inside  $S$  over the final two periods.

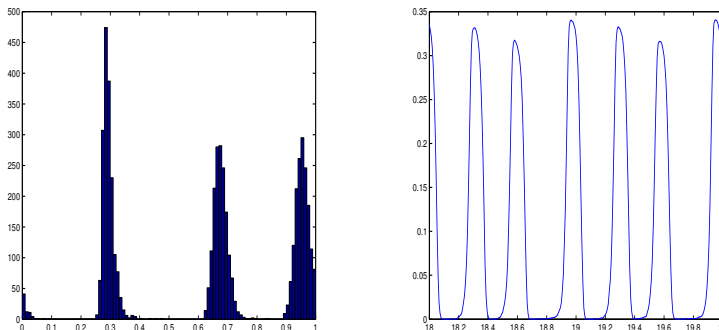


Figure 2: Simulations of the graduated model with acceleration. Here  $R = [.1, .2]$ ,  $S = [.2, .3]$ ,  $\sigma = .02$  and the linear promotion factor was 5. (a) Histogram of the final distribution of cells within the cell cycles. (b) Time-series of the final two time frames. One unit of time corresponds to one unperturbed cell cycle.

The figures clearly show clustering for all the specified parameter values. We note that increasing the noise too much could destroy the clustering effect. The amount of noise needed to do so was smaller for larger numbers of clusters.

We observe the following things from the simulations:

- The inhibition produces more clusters than acceleration.
- The number of clusters formed is inversely proportional to the width of  $S$  or  $|R| + |S|$ .
- Acceleration causes sharper clustering, i.e. the clusters appear to be completely separated.

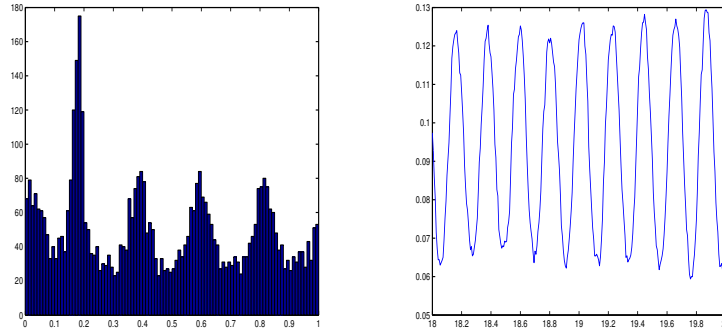


Figure 3: Simulations of the graduated model with inhibition. Here  $R = [.1, .2]$ ,  $S = [.2, .3]$ ,  $\sigma = .02$  and the linear inhibition factor was 5. (a) Histogram of the final distribution of cells. (b) Time-series of the final two time frames.

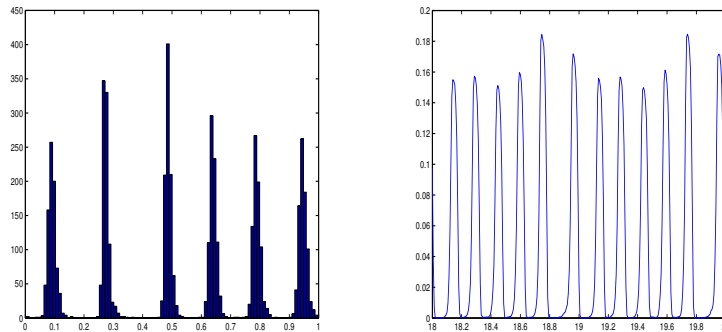


Figure 4: Simulations of the graduated model with acceleration. Here  $R = [.15, .2]$ ,  $S = [.2, .25]$ ,  $\sigma = .01$  and the linear promotion factor was 10. (a) Histogram of the final distribution of cells. (b) Time-series of the final two time frames.

The first two of these observations are consistent with the results concerning the idealized advance and delay models. In general these simulations with more realistic models confirm our conclusions from rigorous analysis of the idealized models.

## 5.2 Stratified Model

When yeast cells divide, one is the mother and the other the daughter. The mother carries a scar from the bud and the daughter does not. The number of daughters a cell has had can be determined by counting the bud scars. We will refer to a cell's *generation* as the number of daughters it has produced, starting the count from 0 for the daughters themselves in the first cell cycle.

It is known that after division the mother's volume is slightly larger than that of a daughter. Further, the later generations have slightly shorter times to budding and slightly shorter cell cycle times.

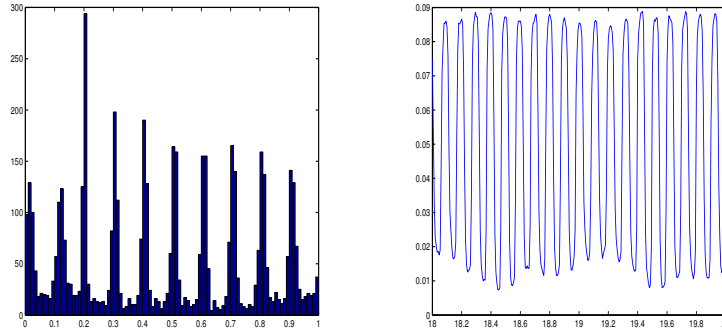


Figure 5: Simulations of the graduated model with inhibition. Here  $R = [.15, .2]$ ,  $S = [.2, .25]$ ,  $\sigma = .01$  and the linear inhibition factor was 10. (a) Histogram of the final distribution of cells. (b) Time-series of the final two time frames.

Leslie proposed a general stratified population model that takes into account the differences in generations that we adopt to yeast in [20]. As before, we will simplify the model by considering logarithmic coordinates, normalized by the cell cycle length of the zero-th generation. In these coordinates the cell cycle of the 0 generation is the unit interval  $I_0 = [0, 1]$ . We will denote the  $S$  phase and  $R$  phases of the 0 generation by  $R = [r_0, s_0]$  and  $S = [s_0, t_0]$ . The successive generations in these coordinates are represented by the intervals  $I_k = [0, D_k]$  with  $R_k = [r_k, s_k]$  and  $S_k = [s_k, t_k]$ .

For yeast under the conditions of the oscillation experiments, values of these parameters are not precisely known.

Note first that if  $D_k = 1$ ,  $R_k = R_0$  and  $S_k = S_0$  for all  $k \geq 1$ , then this model reduces to the simplified model of the previous sections. Next we note that if these conditions approximately hold then the above analysis can be applied in the same way as the small random perturbations.

If there are marked differences in the parameters between generations, then we still might be able to repeat a rigorous analysis. For example, suppose that  $D_k = 2/3$  for all  $k > 0$ . Then in the idealized models we could have 3 clusters in the 0-th generation and 2 clusters in the higher generations. In this scenario, at division parents would be synchronized with a cluster of daughters that was previously  $1/3$  of a cycle ahead of them. As another example, suppose that the ratio above is  $3/4$ , then the system could support 4 daughter clusters and 3 parent clusters. In the same way, rational ratios of cell cycles could produce a variety of clustering combinations.

In real systems, first of all, ratios close to the above idealized “resonances” could still lead to clustering. Secondly, we may suppose that only the first few generations matter since higher generations are represented in numbers that decay approximately geometrically. Biologically, the successive generations beyond the second have not been found to exhibit marked differences anyway.

Elsewhere we consider stratified Leslie Models in more detail, including careful numerical simulations [20], [21].

## 6 Discussion

The main conclusion we wish to emphasize is that clustering seems to be a very robust phenomenon, it occurs in all the models studied and for a large spectrum of parameter values. This latter point is quite important since the actual systems in question are so complex that many of the parameter values cannot be accurately obtained.

The observed synchronous behavior here is not driven by the cell cycle itself, but by the feedback mechanisms. However, the number of clusters must necessarily be an integer and so the oscillations produced by clustering would naturally appear with a period that is an integer fraction of the cell cycle period.

We observe two phenomena that possibly make inhibition a more reasonable explanation of yeast cell-cycle synchrony. First, inhibition allows for large numbers of clusters as seen in simulations with more realistic models. Second, inhibition is seen to initiate clustering more naturally, which we observe in analysis of the idealized models. However, either positive or negative feedback are possibly agents of clustering and conclusive evidence of either cause would need extensive biological modeling confirmed by experiment.

Given the robustness of cluster, we suspect that clustering occurs in many other biological systems with cell cycles. Specifically it could play a role in many types of microbiological systems with cell cycles and some type of signaling, including bacteria [4, 6, 7, 15, 16]

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