

Session 4: Phenotype & Diffusion

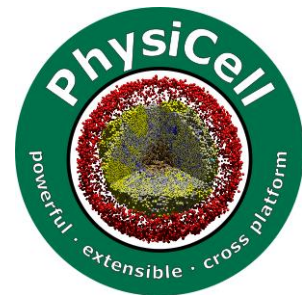


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Agenda:

- Cell Phenotype Continued
 - Cell Cycle
 - Cell Death
 - Cell Secretion and Uptake

Key parts of a PhysiCell model (2)

- **Cell agents (individual players):**
 - Which cell type? (the cell agent is initialized based on a cell definition)
 - State variables:
 - ♦ position
 - ♦ mechanical pressure
 - ♦ interaction list (optional)
 - Phenotype **(the script)**
 - ♦ Cell cycle
 - ♦ Volume
 - ♦ Death
 - ♦ Motility
 - ♦ Mechanics
 - ♦ Substrate uptake & release
 - Custom variables
 - Custom functions that act upon the phenotype, variables, and state **(script)**

A note about time steps

- PhysiCell is designed to account for the multiple time scales inherent to these problems, and has 3 time scales:

▪ $\Delta t_{\text{diffusion}}$	diffusion, secretion, and uptake	(default: 0.01 min)
▪ $\Delta t_{\text{mechanics}}$	cell movement	(default: 0.1 min)
▪ Δt_{cell}	phenotype and volume changes	(default: 6 min)
- This allows some efficiency improvements: not all functions need to be evaluated at each time step.
- See the PhysiCell method paper. (Oddly, not in the User Guide (yet).)

Cell phenotype

- One of the most critical data elements in a PhysiCell Cell is ***phenotype***
- Hierarchically organize key behavioral elements:
 - Phenotype
 - ♦ **cycle**: advancement through a cell cycle model
 - ♦ **death**: one or more types of cell death
 - ♦ **volume**: cell's volume regulation
 - ♦ **geometry**: cell's radius and surface area
 - ♦ **mechanics**: adhesion and resistance to deformation ("repulsion")
 - ♦ **motility**: active motion (other than "passive" mechanics)
 - ♦ **secretion**: both release and uptake of chemical substrates. Interfaces with BioFVM
 - ♦ **intracellular**: used for interacting with intracellular models. See Sessions 10-11.
 - ♦ **molecular**: a place to store internalized substrates

Documentation: User Guide, Sec. 10

Phenotype: Cycle

- Each agent's **phenotype** had a **cycle** with:

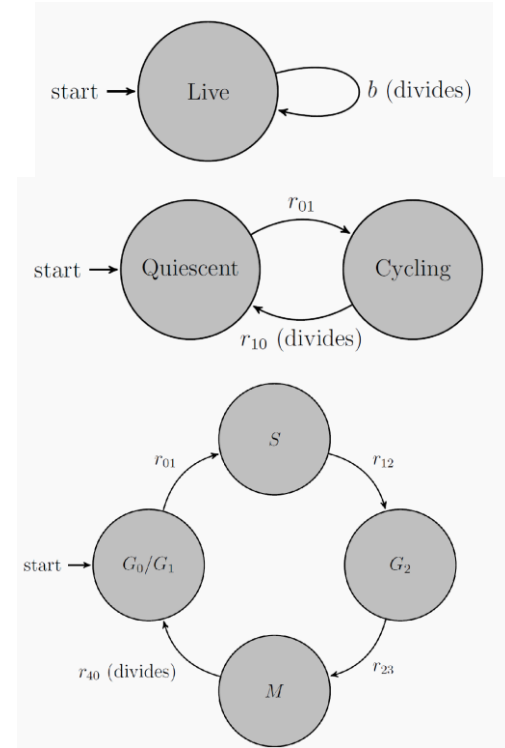
- Cycle model**

- ♦ A directional graph: *nodes* are cycle **phases** $\{P_i\}$ and *edges* are **transition rates** $\{r_{ij}\}$
- ♦ r_{ij} is the transition rate from phase P_i to phase P_j
- ♦ One of the transitions must be marked as a *division transition*
- ♦ Users can attach **arrest condition** functions to these transitions (e.g., size checks)

- Cycle data**

- ♦ stores the cell's current transition rates

- Documentation:** User Guide, Sec. 11.1



Phenotype: Cycle

Cell Cycles available in PhysiCell

- Live
- Ki-67 Basic
- Ki-67 Advanced
- Flow Cytometry
- Flow Cytometry Separated
- Cycling-Quiescent

Cell definition: cycle

```
<cycle code="5" name="live"> <!-- pick a code to match PhysiCell_constants.h -->
    <!-- use phase_transition_rates OR phase_durations -->
    <phase_transition_rates units="1/min">
        <rate start_index="0" end_index="0" fixed_duration="false">0.002</rate>
    </phase_transition_rates>

    <!-- use phase_transition_rates OR phase_durations -->

    <phase_durations units="min">
        <duration index="0" fixed_duration="false">500.0</duration>
    </phase_durations>

-->

</cycle>
```

- For some problems, it's easier to work in terms of transition rates. Use the "phase_transition_rates" code for these.
 - In this example, the "live" cell cycle (with a single phase) transitions at a rate of 0.002 1/min.
- Sometimes, it's easier to work in terms of how long a cell spends in a phase. Use "phase_durations" for these.
 - In this example, the "live" cell cycle (with a single phase) lasts (on average) 500 minutes.

cycle app demo

- <https://nanohub.org/resources/trcycle>



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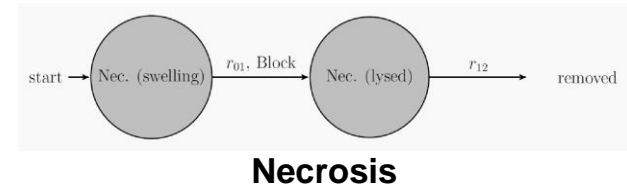
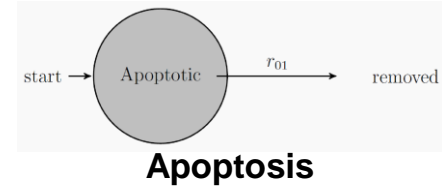
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Phenotype: Death

- **Death** has one or more death models:
 - A specialized cycle model with a *removal* transition rate
 - Extra parameters to help govern cell volume
 - Each death model has an associated death rate
 - Also stores an easy Boolean **dead** to easily check if the cell is alive.
- PhysiCell has built-in apoptosis and necrosis death models



Documentation: User Guide, Sec. 11.2

Cell definition: death

```
<death>

<model code="100" name="apoptosis">
  <death_rate units="1/min">0</death_rate>
  <!-- use phase_transition_rates OR phase_durations -->
  <phase_durations units="min">
    <duration index="0" fixed_duration="true">516</duration>
  </phase_durations>
  <parameters>
    <unlysed_fluid_change_rate units="1/min">0.05</unlysed_fluid_change_rate>
    <lysed_fluid_change_rate units="1/min">0</lysed_fluid_change_rate>
    <cytoplasmic_biomass_change_rate units="1/min">1.66667e-02</cytoplasmic_biomass_change_rate>
    <nuclear_biomass_change_rate units="1/min">5.83333e-03</nuclear_biomass_change_rate>
    <calcification_rate units="1/min">0</calcification_rate>
    <relative_rupture_volume units="dimensionless">2.0</relative_rupture_volume>
  </parameters>
</model>

<model code="101" name="necrosis">
  <death_rate units="1/min">0.0</death_rate>
  <!-- necrosis uses phase_duration[0] = 0 so that it always immediately
       tries to transition and instead checks volume against the rupture volume -->
  <phase_durations units="min">
    <duration index="0" fixed_duration="true">0</duration>
    <duration index="1" fixed_duration="true">86400</duration>
  </phase_durations>
  <parameters>
    <unlysed_fluid_change_rate units="1/min">0.05</unlysed_fluid_change_rate>
    <lysed_fluid_change_rate units="1/min">0</lysed_fluid_change_rate>
    <cytoplasmic_biomass_change_rate units="1/min">1.66667e-02</cytoplasmic_biomass_change_rate>
    <nuclear_biomass_change_rate units="1/min">5.83333e-03</nuclear_biomass_change_rate>
    <calcification_rate units="1/min">0</calcification_rate>
    <relative_rupture_volume units="dimensionless">2.0</relative_rupture_volume>
  </parameters>
</model>

</death>
```

- Use death_rate to determine the rate of *starting* each mode of death.
- Use the phase_durations and parameters to control how cells progress through each death model.

death app demo

- <https://nanohub.org/resources/trdeath>



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Phenotype: Secretion

- **Secretion** stores parameters for secretion, uptake, and generalized export of diffusing substrates

$$\frac{\partial \rho}{\partial t} = \nabla \cdot (D \nabla \rho) - \lambda \cdot \rho + \sum_i \delta(\mathbf{x} - \mathbf{x}_i) V_i (\mathbf{S}_i \cdot (\rho_i^* - \rho) - \mathbf{U}_i \cdot \rho + \mathbf{E}_i)$$

PhysiCell automatically tracks the mass of substrates removed from the tissue (added to cells) or added to tissue (removed from cells).

Documentation: User Guide Sec. 11.7

Important Parameters

- Differentiate between net export vs secretion rate
- Secretion rate is dependent upon Volume

Cell definition: Secretion

```
<secretion>
  <substrate name="chemical_A">
    <secretion_rate units="l/min">0</secretion_rate>
    <secretion_target units="substrate density">1</secretion_target>
    <uptake_rate units="l/min">0</uptake_rate>
    <net_export_rate units="total substrate/min">0</net_export_rate>
  </substrate>

  <substrate name="chemical_B">
    <secretion_rate units="l/min">0</secretion_rate>
    <secretion_target units="substrate density">1</secretion_target>
    <uptake_rate units="l/min">0</uptake_rate>
    <net_export_rate units="total substrate/min">0</net_export_rate>
  </substrate>

  <substrate name="chemical_C">
    <secretion_rate units="l/min">0</secretion_rate>
    <secretion_target units="substrate density">1</secretion_target>
    <uptake_rate units="l/min">0</uptake_rate>
    <net_export_rate units="total substrate/min">0</net_export_rate>
  </substrate>
</secretion>
```



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secretion app demo

- <https://nanohub.org/resources/trsecretion>

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