

# Mathematical Modeling in Biology

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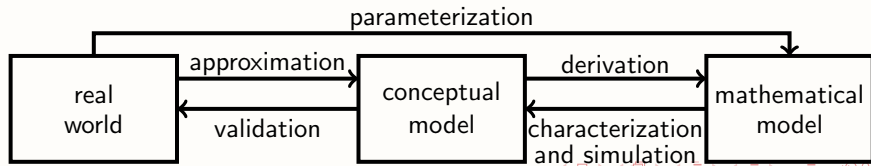
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# General Principles of Modeling

- ▶ Modeling contrasted with mathematics
- ▶ What are mathematical models?
- ▶ Models as functions of parameters

## Mathematics vs Modeling

- ▶ Mathematics
  - Assumptions define the setting.
  - Conclusions follow from mathematical logic.
  - Focus is on proof.
- ▶ Modeling
  - Assumptions define a *conceptual model* of a real setting.
  - Conclusions *for the model* follow from mathematical logic.
  - Conclusions *for the setting* are only as good as the conceptual model.
  - Focus is on checking results against known outcomes.



## What is a Mathematical Model?

- ▶ A **mathematical model** is a self-contained collection of one or more variables together with a set of rules (usually formulas and equations) that prescribe the values of those variables.
  - Models serve as an approximate quantitative description of some actual or hypothetical real-world scenario.
  - Models are created in the hope that the behavior they predict will capture enough of the features of that scenario to be useful.
  - The value of a model depends on the setting to which it is applied and the questions it is used to address.
- ▶ A **mechanistic model** is a mathematical model based on assumptions about the scientific principles that underlie the phenomena being modeled.

## Models as Functions of Parameters

How do we view the logistic growth model

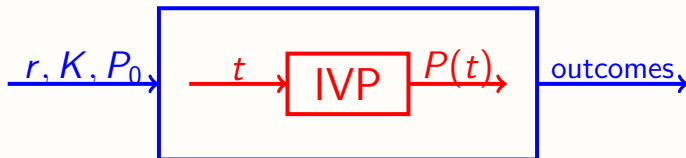
$$\frac{dP}{dt} = rP \left( 1 - \frac{P}{K} \right), \quad P(0) = P_0 > 0, \quad r, K > 0?$$

► **Narrow** view:

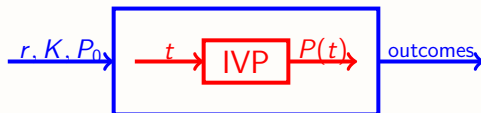
Initial value problem for  $P(t)$ , with parameters  $r$ ,  $K$ , and  $P_0$ .

► **Broad** view:

Function that maps parameters  $r$ ,  $K$ , and  $P_0$  to outcomes.



## Models as Functions of Parameters



- ▶ **Narrow** view: Math problem with fixed parameters.
  - The narrow view is used to determine the outcomes.
    - Narrow view questions are trivial: “Given  $K = 10$ ,  $R = 1$ , and  $P_0 = 1$ , when does the population reach  $P = 5$ ?”
- ▶ **Broad** view: Math problem with outcomes as functions of parameters.
  - The important questions are in the broad view.
    - Do solutions with any initial condition always approach  $K$ ?
    - At what point is the population growth the fastest?

## An Example – the SEIR Epidemic Model

1. Assumptions and mathematical derivation
  - 1.1 Class structure
  - 1.2 Processes
  - 1.3 Differential equations
  - 1.4 Basic reproductive number
2. Designing an investigation
  - 2.1 Asking questions
  - 2.2 Choosing study parameters and outcomes
3. Overview of Methods
4. Addressing questions and reporting results
  - 4.1 Parameter studies
  - 4.2 Using simulations
  - 4.3 Simulations can suggest conjectures

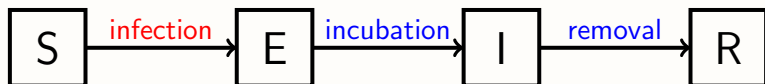
## 1.1. Class structure

- ▶ Individuals in a population are divided into classes. These vary among different epidemiological models. For SEIR:
  - **S**: *Susceptible* – can be infected
  - **E**: *Exposed* – infected but not infectious
  - **I**: *Infectious* – can transmit the disease to susceptibles
  - **R**: *Removed* – no longer infectious
- ▶ Sometimes the names are misleading.
  - '*Exposed*' should be '*Latent*' (already infected, not merely exposed)
  - *Removed* includes people who are still sick and may include people who are deceased
- ▶ Models are designated by the class structure: SIR, SIS, SEIR, SEAIR, SEAIRHD etc



## 1.2 Processes

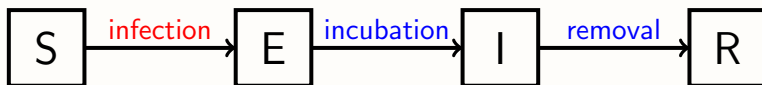
- ▶ Processes move individuals from one class to another.
  - Example: Basic SEIR model



- Rate of change of  $S$  is  $-\text{infection}$
- Rate of change of  $E$  is  $\text{infection} - \text{incubation}$
- Rate of change of  $I$  is  $\text{incubation} - \text{removal}$
- Rate of change of  $R$  is  $\text{removal}$

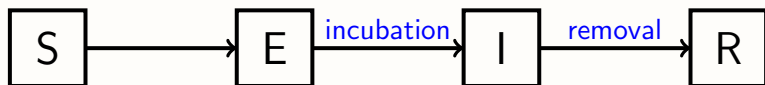
## 1.2 Processes

- ▶ Processes are either **transmissions** or **transitions**.



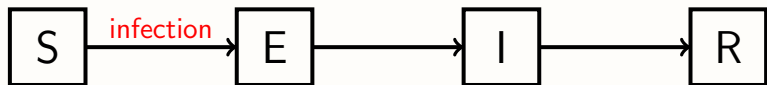
- **Transmissions** require interaction with another class.
  - Susceptibles are infected by Infectives.
- **Transitions** happen without any interaction.
  - Incubation of Latents and removal of Infectives happen spontaneously (but perhaps in phases).

## 1.2 Transitions



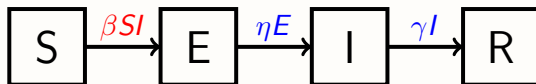
- ▶ Transition rates are proportional to the **leaving** class (assuming one-phase transitions)
  - incubation rate = constant \*  $E = \eta E$
  - removal rate = constant \*  $I = \gamma I$
- ▶ **Rate constants are reciprocals of average time in class.**
  - Average removal time 10 days  $\rightarrow \gamma = 0.1$

## 1.2 Transmissions



- ▶ Transmission rates are proportional to the **leaving** class size
  - infection rate = force of infection \*  $S = \lambda S$
- ▶ The force of infection is (usually) proportional to the sum of the **transmitting** classes (just I for SEIR)
  - force of infection = constant \*  $I = \beta I$
- ▶ The infection rate is  $\beta I * S = \beta SI$

## 1.3 Differential equation model



$$S' = -\beta SI$$

$$E' = \beta SI - \eta E$$

$$I' = \eta E - \gamma I$$

$$R' = \gamma I$$

- ▶ Let  $N = S + E + I + R$ . Then  $N' = 0$ , so  $N$  is constant. (Without loss of generality, we can take  $N = 1$ .)
- ▶ The model is an **autonomous dynamical system** (rates of change depend only on the state of the system).

## 1.4 Basic reproductive number

- ▶ **Basic reproductive number  $\mathcal{R}_0$ :**  
the average number of secondary infections caused by one primary infective in a fully susceptible population.
  - $\mathcal{R}_0 > 1$  is needed to start an epidemic.
- ▶ The total number is the average rate times the average time.
- ▶ Calculation of average transmission rate:
  - Recall that the **transmission rate** is  $\beta SI$
  - Transmission rate **per infective**:  $\beta S$
  - Rate per infective in a **fully-susceptible population**:  $\beta N$

## 1.4 Basic reproductive number

- ▶ **Basic reproductive number  $\mathcal{R}_0$ :**  
average transmission rate per infective in a fully susceptible population multiplied by average time in the Infectious class.
- ▶ Average transmission rate per infective in a fully susceptible population:  $\beta N$
- ▶ Average time in infectious class:  $1/\gamma$  (reciprocal of  $\gamma$ )
- ▶ Average number is average rate times average time:

$$\mathcal{R}_0 = \beta N \cdot \frac{1}{\gamma} = \frac{\beta N}{\gamma}.$$

- ▶ Other diseases (like COVID-19) can be more complicated.

## 2.1 Asking questions

- ▶ Models must be designed to answer specific questions.
  - If we want to know the impact of COVID-19 on health care resources, we need to modify the SEIR model to track hospitalizations and/or ICU patients.
  - If we want to know the impact of mitigation strategies on COVID-19, we need to build mitigation into the model.
    - My COVID-19 model is SEAIHRD with  $\lambda$  impacted by testing and a contact factor.
- ▶ Some common question types:
  - Is a specific claim supported by modeling or not?
  - What effect does parameter  $x$  have on outcome  $y$ ?



## 2.2 Choosing study parameters

- ▶ Model parameters are not always the best study parameters, especially when they are hard to measure.
  - Transition times are more fundamental than transmission rate constants. (Take  $\gamma = 1/T_i$ )
  - The transmission parameter  $\beta$  is dependent on population size, while the basic reproductive number  $\mathcal{R}_0$  is a fundamental disease property. (Take  $\beta = \gamma\mathcal{R}_0/N$ .)
- ▶ Some parameters are more fundamental than others.
  - The effect of the disease duration  $T_i$ , given fixed  $\mathcal{R}_0$ , is simply to change the time scale for the results.
  - Changing a model to dimensionless form (see Ledder, *Mechanistic Modeling*, Section 5) eliminates scale parameters.

## 2.2 Choosing outcomes

- ▶ Maximum number of new infections?
- ▶ Maximum number of hospitalizations per million?  
(compared to an average of 2800 hospital beds per million in the US)
  - Serves as a measure of the stress on the health care system
- ▶ Percent deaths? (0.1% in the US is 325,000 people)
  - Serves as a measure of the human cost
- ▶ Final fraction of susceptibles?
  - Serves as a measure of the risk of a new outbreak
- ▶ Times for any of these events?

### 3. Overview of methods

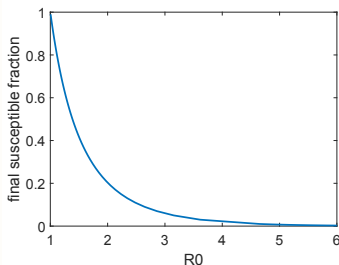
- ▶ Solution – seldom useful
  - Very few differential equation problems can be solved using methods of calculus. Even when possible, the results are often less useful than those obtained using other methods.
- ▶ Characterization – using hand computation to obtain properties of a model
  - We can often determine long-term behavior using mostly algebra.
  - Results are often general (parameters left unspecified).
- ▶ Simulation – using numerical computation to obtain results for individual scenarios
  - Results are never general (parameters must be specified).
  - Can still be used to address general questions.

## 4. Addressing questions and reporting results

- ▶ Answers to math questions are often numbers or formulas. **Modeling questions require verbal answers**, supplemented with visual aids.
  - ‘The graph goes up and then comes down’ is merely a *description*. An *explanation* connects to the real world scenario and **offers a reason** for the observed results.
- ▶ Graphs must be informative and not misleading.
  - No negative values for populations or parameters.
  - Axes must be labeled.
  - Sometimes multiple curves on the same axes are more informative than multiple graphs.
  - Measured data should be plotted as points; simulation results should be plotted as dot-to-dot ‘curves’.

## 4.1 Parameter studies

- ▶ Sensitivity analysis can determine how important a parameter is, but it does not determine the parameter's effect.
- ▶ Parameter studies determine the quantitative effect of a parameter on one or more outcomes.
  - SEIR example: Assume no initial immunity or mitigation. How does the fraction of people who don't get the disease depend on the basic reproductive number? (Homework problems 2–3)



## 4.2 Using simulations

Modeling with simulations requires two program components:

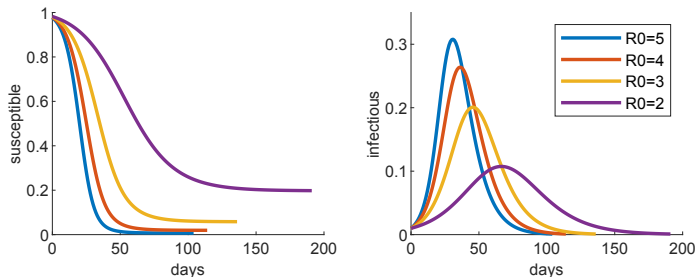
1. A **function** that
  - Accepts input values for
    - Disease parameters, like  $\gamma$  and  $\beta$ ,
    - Scenario parameters, like initial and terminal conditions,and
  - Returns the time history of the class counts.
2. A **driver script** that
  - Organizes an experiment,
  - Utilizes the function to obtain outcomes, and
  - Displays the results.

## 4.2 SIR\_paramstudy.R

This driver script plots various outcomes as a function of a parameter. Script elements:

1. Define function `sir_sim(beta,gamma,I0,V,target)`.
2. Define default scenario values.
3. Prescribe range and count of study parameter values.
4. Set up data structures.
5. Run loop:
  - Collect study parameter value.
  - Compute derived parameter values.
  - Use function `sir_sim` to collect results.
  - Add results to a data structure.
6. Create plots.

## 4.3 Simulations can suggest conjectures



We can prove these conjectures (**Homework problem 3**)

1. The fraction of susceptibles is continually decreasing, but is bounded away from 0.
2. The epidemic ends with  $I = 0$  and  $S = s_\infty$  for some  $s_\infty > 0$ .



## Homework

Problems 1–3 are essential. Problem 4 is ‘extra credit.’

1. Use characterization to find a fatal flaw in a commonly-used model.
2. Use an R program to produce the parameter study plot on the 4.1 slide. You will just need to make a few changes to `SIR_paramstudy.R`.
3. Use mathematics to prove the conjectures on the 4.3 slide.
4. Work through a model research study that could have been done by undergraduates (but without any prompting other than a general question).