

# 1 Epidemic Modelling Intro



Mathematics  
and Statistics

$$\int_M d\omega = \int_{\partial M} \omega$$

# Mathematics 4MB3/6MB3 Mathematical Biology

Instructor: David Earn

Lecture 1  
Epidemic Modelling Intro  
Monday 8 January 2018

# Where to find course information

- The course web page:  
<http://www.math.mcmaster.ca/earn/4MB3>
- Click on “Course information”.
- Download pdf or read online.
- Let's [have a look now...](#)

# Who is NOT available at these times?

- Monday 10:30-11:30
- Monday 12:30-1:30
- Wednesday 10:30-11:30
- Wednesday 12:30-1:30
- Friday 12:30-1:30
- Friday 2:30-3:20

# Group formation

**Most work in this course will be done in groups.**

- Attempt to form a group of 4 students (you and 3 others) **no later than Thursday night this week.**
- After you have done your best to form a group of four, **exactly one** member of your group must **e-mail the instructor no later than Thursday night this week:**
  - Include “Math 4MB3” and your proposed group name in the subject line.
  - **Copy your message to all members of your proposed group so I have everyone’s e-mail in the thread.**
- If you were unable to form a group, then e-mail the instructor explaining what you did to try to form a group, and describe your skills/preferences. (*This is a last resort – please try your best to form a group.*)
- *Instructor may change groups based on survey results.*

# Online Surveys

*You will be required to fill in online surveys during this course. Doing so in a timely manner contributes to your participation mark.*

The first online survey has been posted:

- Go to the [Surveys page](#) on the [course web site](#).
- Follow the link for [Background and Group formation Survey](#).
- Complete the survey **no later than 11:59pm this Thursday (11 Jan 2018)**.
- It should take only  $\sim 5$  minutes.
- Note that *surveys sometimes fail to save*.
  - Type long answers into a file first and paste them into the survey. Then you won't get as frustrated if it fails to save.

# Software

- **ASAP**, install the software discussed on the [Software page](#) on the [course web site](#):

- L<sup>A</sup>T<sub>E</sub>X



- R



- RStudio



- XPPAUT

- Emacs



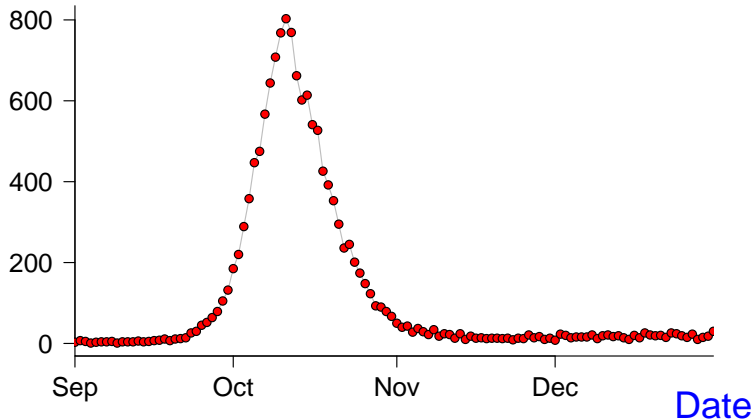
- If you have installation problems, please contact [Ken Moyle](#) <[moylek@mcmaster.ca](mailto:moylek@mcmaster.ca)>, who created the [Software page](#).
- **Note:** the [Software page](#) also contains some info about spell-checking and counting words in L<sup>A</sup>T<sub>E</sub>X documents.

# Epidemic Modelling



# Pneumonia & Influenza Mortality, Philadelphia, 1918

## P&I Deaths



# Modelling challenge

Develop a model that helps us understand the [graph on the previous slide](#), based on mechanisms of disease spread.

- Only one variable is observed (P&I deaths per day) so construct a model containing only one variable.
- Think about how disease spreads and express your thoughts with mathematical notation.
- Derive a differential equation that models the process of disease transmission.
- Analyze the model and determine its strengths and weaknesses/limitations.

# Make (Biological) Assumptions Clear

- 1 *Assume* the disease is transmitted by contact between an infected individual and a susceptible individual.
- 2 *Assume* the latent period (delay between being infected and becoming infectious) is so short that it can be ignored (technically assume it is zero).
- 3 *Assume* all members of the population are identical and respond identically to the disease. In particular, all susceptible individuals are equally susceptible and all infected individuals are equally infectious.
- 4 *Assume* the population size is fixed during the epidemic, *i.e.*, ignore births, migration, and deaths from causes other than the disease, and count individuals who have died from the disease as part of the population.

# About Assumptions. . .

- Note that the first assumption on the [previous slide](#) is actually correct.
- The other assumptions are wrong, but are reasonable approximations.
- It is best to *start as simple as possible and add complexity later*, in order to:
  - obtain a model that actually succeeds in explaining [the data](#) with as few assumptions as possible;
  - check that inferences we draw from our model(s) are robust to the inclusion of more biological details/realism.

# What variables should we include in our model?

- Independent variable: time ( $t$ )
- Dependent variable: Many options, e.g.,
  - Incidence (number of new infections per unit time)
  - Prevalence (total current number of infected individuals)
  - Death rate (number of deaths per unit time)
  - Death toll (number of deaths so far)
- So, what would be best?
- Not deaths, because whether or not you die may be unrelated to how much you transmit.
- But deaths are what **we observe!** What to do?!?
- Make another assumption. . .

## Additional assumption(s)

- We actually want to know incidence or prevalence, but we **observe deaths**.
- Under what circumstances would daily deaths be a good estimate of incidence? (*i.e.*, What must we assume in addition to the **assumptions we have already made**.)
  - 5 **Assume** that the time from infection to death is exactly the same (a certain number of days) for every individual who dies.
  - 6 **Assume** that the probability of dying from the disease is the same for every individual who is infected.
- Then daily death counts are proportional to daily incidence a certain number of days in the past, *i.e.*, the “mortality curve” that **we observe** is a translated and scaled version of the “epidemic curve” (new cases per day).

## So... what variables should we include in our model?

- Independent variable: time ( $t$ )
- Dependent variable: one of:
  - Incidence (number of new infections per unit time)
  - Prevalence (total current number of infected individuals)
- Which one?
- Choose prevalence ( $I$ ) because anybody who is currently infectious can infect others, so it will probably be easier to formulate a transmission model based on prevalence. (Try not to lose sight of underlying biological mechanisms.)
- But our **mortality curve** is related to incidence, not prevalence!?! Argh. What to do?!?
- Let's work with prevalence and see how it works out. Maybe we'll be able to derive the incidence curve from a model based on prevalence.