

Challenges of Teaching Bayesian Statistics to Undergraduates

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A motivating example for undergraduates

- ▶ A major challenge for teaching undergraduates is finding motivating applications
- ▶ In this talk, I will describe a shiny app that displays a Bayesian analysis of disease outbreak data
- ▶ This cursory example does not include any distributions or computing
- ▶ The lecture is meant to pique interest and give a high-level overview of Bayesian methods

This is a good motivating example

- ▶ Disease outbreak modeling is obviously topical
- ▶ Fiddling with curves is fun!
- ▶ There are only two parameters and they are interpretable
- ▶ Data is sparse early in the outbreak, so priors are crucial
- ▶ Leads to a good discussion of the role of prior and ideas for setting priors

Background

- ▶ The lecture would begin with a description of the data and model
- ▶ The SIR model is a differential equation, but it could be presented as a discrete time model (or without equations)
- ▶ Markdown document with background material

Description of the SIR model for a disease outbreak

Description of the data

We will use the *Influenza in a boarding school in England, 1978* in the packages *outbreaks* for illustration. The data are described in the packages as:

These data comprise of a time series of influenza cases in a boarding school in England. The original data were available only in a figure with some additional data in the main text; hence, the exact numbers vary depending on the source. These data are from Chapter 9 of De Vries et al. (1996). The index case was infected by 1978-01-10, and had febrile illness from 1978-01-15 to 1978-01-18. 512 boys out of 763 became ill.

Example model fits are given below, and interactive model fitting can be performed at

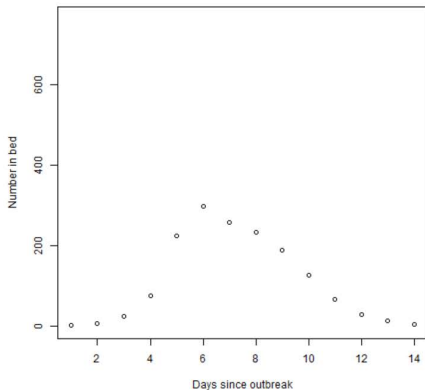
<https://shiny.stat.ncsu.edu/bjreich/SIR/>

```
library(outbreaks)
influenza_england_1978_school
```

```
##      date in_bed convalescent
## 1 1978-01-22      3           0
## 2 1978-01-23      8           0
## 3 1978-01-24     26           0
## 4 1978-01-25     76           0
## 5 1978-01-26    225           9
## 6 1978-01-27    298          17
## 7 1978-01-28    258         105
## 8 1978-01-29    233         162
## 9 1978-01-30    189         176
## 10 1978-01-31    128         166
## 11 1978-02-01     68         150
## 12 1978-02-02     29          85
## 13 1978-02-03     14          47
## 14 1978-02-04      4          20
```

```
Y <- influenza_england_1978_school[,2]
nt <- length(Y)      # Number of time sets
N <- 763              # Population size
I0 <- 1               # Initial number of infected

plot(Y,ylim=c(0,N),xlab="Days since outbreak",ylab="Number in bed")
```



Description of the SIR model

Description of the SIR model

The Susceptible-Infected-Recovered (SIR) model is one of the most basic models of disease spread. At time t , let S_t , I_t and R_t be the number of people in each condition, with $S_t + I_t + R_t = N$ for all t . The states evolve according to the differential equations

$$\frac{dS_t}{dt} = -\beta S_t I_t / N \quad \frac{dI_t}{dt} = \beta S_t I_t / N - \gamma I_t.$$

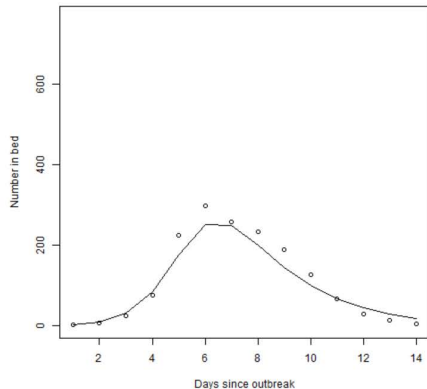
The parameter β controls the rate of new infections and the parameter γ controls the recovery rate. We will use a discrete approximation to these curves with hourly time steps, so $dt = 1/24$. Here are the curves for a few values of the parameters β and γ .

```
SIR <- function(I0,N,beta,gamma,nt,m){
  # Initial states
  i <- I0
  s <- N-I0
  dt <- 1/m
  I <- rep(0,nt)

  for(t in 1:nt){
    for(j in 1:m){
      s <- s - dt*beta*i*s/N
      i <- i + dt*beta*i*s/N - dt*gamma*i
    }
    I[t] <- i
  }
  return(I)}

# Reasonable fit
beta <- 1.7
gamma <- 0.5
I <- SIR(I0=I0,N=N,beta=beta,gamma=gamma,nt=nt,m=24)
plot(Y,ylim=c(0,N),xlab="Days since outbreak",ylab="Number in bed")
```

```
# Reasonable fit
beta <- 1.7
gamma <- 0.5
I <- SIR(I0=I0,N=N,beta=beta,gamma=gamma,nt=nt,m=24)
plot(Y,ylim=c(0,N),xlab="Days since outbreak",ylab="Number in bed")
lines(I)
```

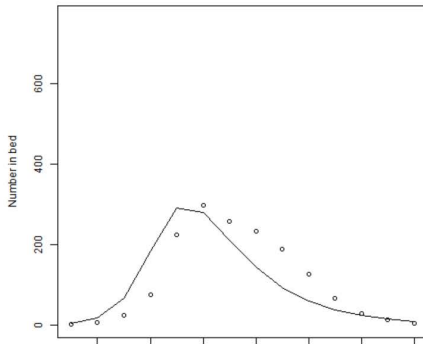


Larger infection rate

Larger infection rate

Increasing β gives a steeper curve because the disease spreads faster.

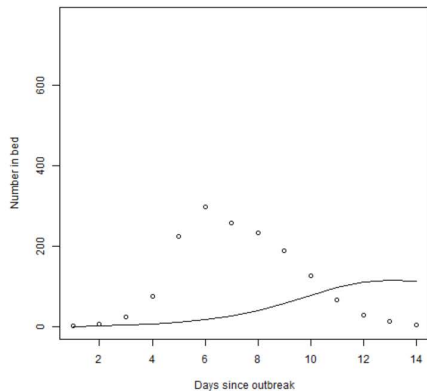
```
beta <- 2.0
gamma <- 0.5
I <- SIR(I0=I0,N=N,beta=beta,gamma=gamma,nt=nt,m=24)
plot(Y,ylim=c(0,N),xlab="Days since outbreak",ylab="Number in bed")
lines(I)
```



Smaller infection rate

Decreasing β flattens the curve.

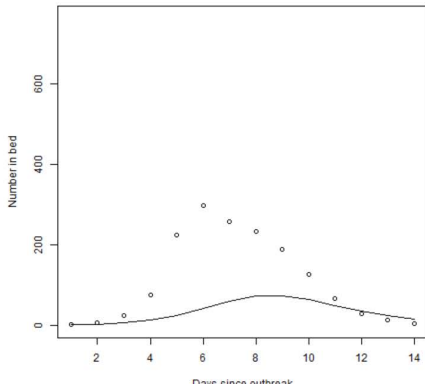
```
beta <- 1.0  
gamma <- 0.5  
I <- SIR(I0=I0,N=N,beta=beta,gamma=gamma,nt=nt,m=24)  
plot(Y,ylim=c(0,N),xlab="Days since outbreak",ylab="Number in bed")  
lines(I)
```



Larger recovery rate

Increasing γ gives fewer overall infections because people are infectious for a shorter amount of time.

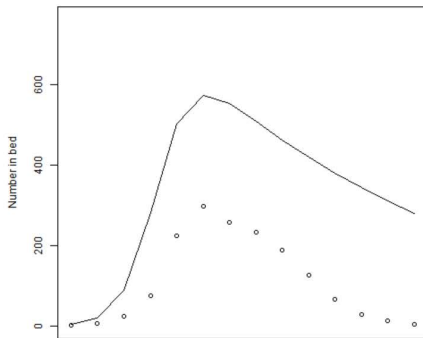
```
beta <- 1.7
gamma <- 1.0
I <- SIR(I0=I0,N=N,beta=beta,gamma=gamma,nt=nt,m=24)
plot(Y,ylim=c(0,N),xlab="Days since outbreak",ylab="Number in bed")
lines(I)
```



Smaller recovery rate

Decreasing γ gives more overall infections because people are infectious for a longer amount of time.

```
beta <- 1.7
gamma <- 0.1
I <- SIR(I0=I0,N=N,beta=beta,gamma=gamma,nt=nt,m=24)
plot(Y,ylim=c(0,N),xlab="Days since outbreak",ylab="Number in bed")
lines(I)
```



Definition of parameters

- ▶ The model *parameters* are β and γ
- ▶ Definition of parameters: fixed but unknown constants in the probability model
- ▶ If we knew the parameters we could:
 - ▶ Compare this outbreak to others
 - ▶ Forecast spread
 - ▶ Predict effects of interventions

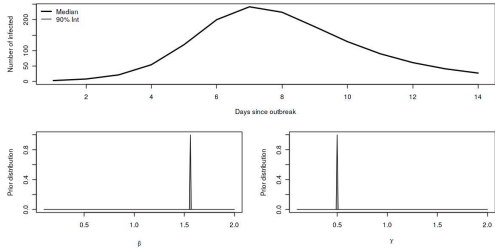
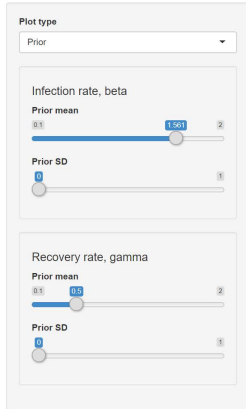
Discussion questions

- ▶ **Question 1:** At the onset of the outbreak, how would you estimate the parameters?
- ▶ **Question 2:** What are the consequences of using the wrong values?

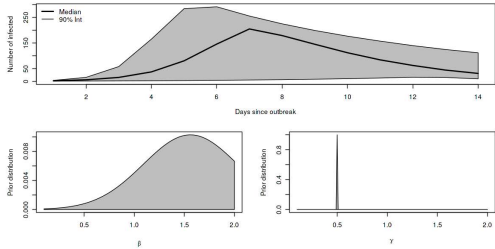
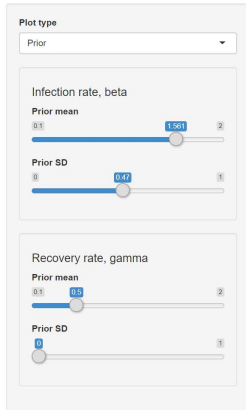
Definition of the prior distribution

- ▶ We rarely know the parameters at the onset of a study; this is why we collect data
- ▶ Rather than select a particular value, we place an uncertainty distribution on the parameters
- ▶ This is called the *prior distribution*
- ▶ You can explore priors and their effect on the curve [here](#)

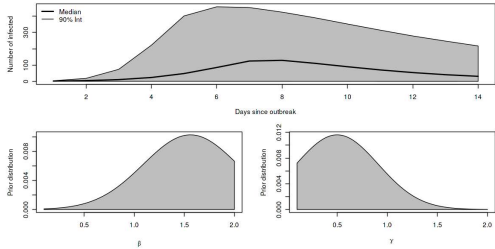
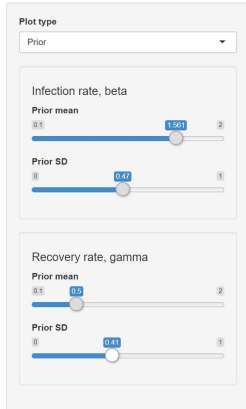
Bayesian analysis of a disease outbreak



Bayesian analysis of a disease outbreak



Bayesian analysis of a disease outbreak



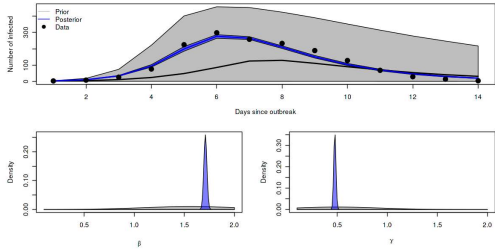
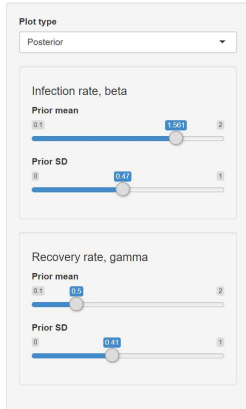
Discussion questions

- ▶ **Question 1:** At the onset of the outbreak, where can you get information about the parameters?
- ▶ **Question 2:** What would you do if there is no prior information?

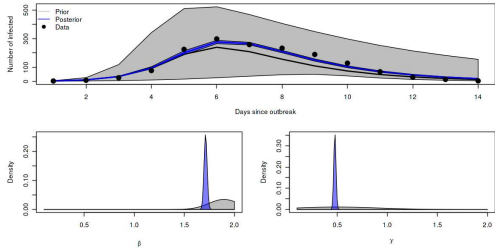
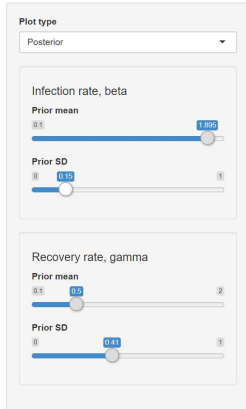
Definition of the posterior distribution

- ▶ After the outbreak we have two sources of information: the prior and the data
- ▶ *Bayesian learning* combines these two sources of information
- ▶ The uncertainty distribution of the parameters after observing the data is the *posterior distribution*
- ▶ You can explore Bayesian learning at [here](#)

Bayesian analysis of a disease outbreak



Bayesian analysis of a disease outbreak



Discussion questions

- ▶ **Question 1:** Does the SIR model seem to fit the data well?
What to do if it doesn't?

- ▶ **Question 2:** How much affect does changing the prior have on the posterior?

More information about the shiny app

- ▶ The material presented here can be found on the website for our book¹
- ▶ The website is <https://bayessm.org/> (“Link to Supplemental Materials” → “Bayesian modeling of a disease outbreak”)
- ▶ We also have many datasets, worked examples, GUIs, beamer slides and solutions to homework problems
- ▶ Thanks!

¹Reich and Ghosh (2019). Bayesian Statistical Methods. CRC Press.