

Advanced disease modelling - Practical

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Motivation

We have data on a flu epidemic in an English boarding school in 1978 source. Among the 763 boys in the school, 512 were infected and put in bed during their infection, which allows us to follow the number of infected boys over time.

```
time = c(3,4,5,6,7,8,9,10,11,12,13,14,15)
infected = c(31,82,216,299,269,242,190,125,81,52,25,22,7)
data=cbind(time, infected)
```

The objective of this exercise is to recover the parameters associated with this epidemic. We use for this an SIR-type model, which can be simulated with the following code:

```
library(deSolve)
```

```
## Warning: package 'deSolve' was built under R version 4.3.3
```

```
##SIR model
SIR<-function(t,x,parms){
  ##taille de chaque compartiment et de la population
  S = x[1]
  I = x[2]
  R = x[3]
  N = x[1]+x[2]+x[3]

  ##valeurs des parametres
  lambda = parms["lambda"]
  gamma = parms["gamma"]

  ##variations
  dS=-lambda*S*I/N
  dI=lambda*S*I/N-gamma*I
  dR=gamma*I
  res = c(dS,dI,dR)
  list(res)
}

simulate_SIR=function(parameters){

  #parameters
  parms = c(parameters["lambda"],parameters["gamma"])
  N=parameters["N"]

  #initial conditions
  init <- c(N-parameters["initI"],parameters["initI"],0)
```

```

#simulation
temps <- seq(0,15)
solveSIR <- lsoda(y =init, times=temps, func = SIR,
                  parms = parms)
solutionSIR=as.data.frame(solveSIR)
names(solutionSIR)=c("time","S","I","R")

#merge with data
sir_data=merge(data,solutionSIR)

return(sir_data)
}

theta_init =c("lambda"=1.7,"gamma"=0.44,"initI"=1, "N"=763)
simul=simulate_SIR(theta_init)
simul

```

##	time	infected	S	I	R
## 1	3	31	708.92573	39.81687	14.25740
## 2	4	82	604.66673	112.66174	45.67153
## 3	5	216	414.29233	228.36691	120.34076
## 4	6	299	226.78000	296.87794	239.34207
## 5	7	269	118.18036	276.76512	368.05452
## 6	8	242	67.95755	217.71490	477.32755
## 7	9	190	44.76844	158.47863	559.75293
## 8	10	125	33.23410	111.17714	618.58876
## 9	11	81	27.02160	76.52240	659.45599
## 10	12	52	23.45243	52.11566	687.43191
## 11	13	25	21.30197	35.27341	706.42461
## 12	14	22	19.96202	23.78333	719.25465
## 13	15	7	19.10746	15.99762	727.89492

The column “infected” contains the data. The columns S, I and R correspond to the number of individuals in the compartments S, I and R respectively. The column time indicates the time in days.

Question 1

Using the previous code, visualise the number of infected boys over time and compare the data with model simulation for various parameters

Question 2

We assume that the number of infected boys observed each day follows a Poisson distribution whose parameter is the number of infected simulated by the model.

Prior distributions are chosen as follows: $\text{Unif}[0,10]$ for the transmission rate λ , $\text{Unif}[0,1]$ pour le cure rate γ .

- With R, write a function that calculates the model’s likelihood for a given set of parameters. Write another function that calculates the model’s log-likelihood (logarithm of the likelihood).
- With R, write a function that calculates the model’s posterior distribution for a given set of parameters. Write another function that calculates the model’s log-posterior (logarithm of the posterior).

Help: These R functions might be helpful: `dunif`, `dpois`.

Question 3

Implement the Metropolis-Hastings algorithm in R.

We can use the following proposal distribution:

```
#SIMULATE PARAMETER SET FROM PROPOSAL
```

```
proposal = function(theta){  
  return(c("lambda"=rnorm(1,mean=theta[["lambda"]],sd=0.01),  
          "gamma"=rnorm(1,mean=theta[["gamma"]],sd=0.01),  
          "initI"=1,  
          "N"=763))  
}  
  
proposal_density = function(theta_before, theta_after){  
  return(dnorm(theta_after[["lambda"]],mean=theta_before[["lambda"]],sd=0.01)*  
         dnorm(theta_after[["gamma"]],mean=theta_before[["gamma"]],sd=0.01))  
}
```

Help: in order to evaluate the acceptance probability, you can use this function: runif

Question 4

Apply the algorithm to the data and comment the results. Can you check if the MCMC chain has converged?

Question 5

Compare the fitted model and the data. Try to include uncertainty in fitted parameters in your simulations.