

Module 11: Lesson 1 Lecture Notes

Callum Arnold

7/19/2021

Contents

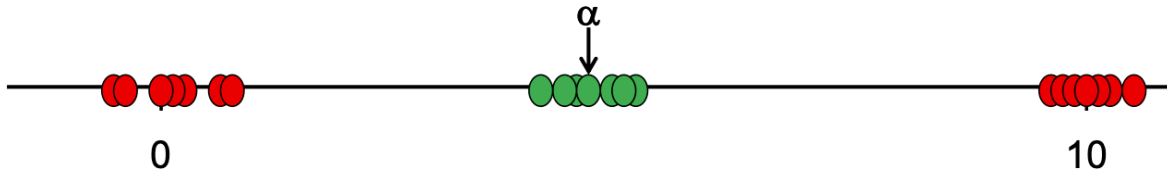
Simulation	1
Introducing simulation	1
Simulating Markov models	2
Simulating non-Markov models	4
Discussion	5
When infections and removals happen at the same time, do we consider to be Markov or non-Markov	5
In the non-Markov model example, the time to infection is exponential, so why is it not a Markov model	6

Simulation

Introducing simulation

- Suppose we have a stochastic epidemic model (e.g. SIR)
- Simulation = producing a realisation of the model (i.e. possible outcome)
 - Producing an outcome according to the correct distribution of all possible outcomes
- For the SIR model
 - producing a set of infection and removal times according to the correct distribution inherent in the model
- Why is it useful?
 - Helps us understand model behaviour
 - It is useful for testing our inference procedure (finding or estimating model parameters) e.g. validate our method against data from the simulation where we know the “truth”
- For example:
 - Perform N ($=1000$) simulations
 - Have model with parameter vector θ fixed at θ_T
 - For each simulation, estimate the model parameters to get $\theta_1, \dots, \theta_N$
 - The average of the model estimates should be close to the true value of θ_T
- Also useful for model checking
 - Let’s say we estimate model parameter α

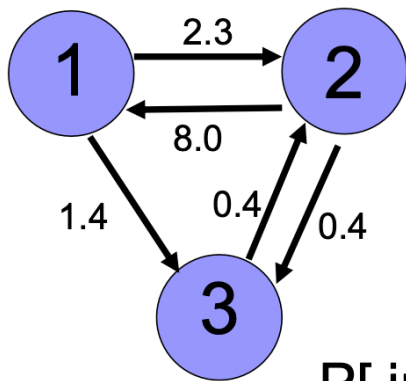
- For each value of α , we perform a large number of simulations and see whether the output of each is similar to the actual data
- Suppose we have a mixture model, e.g. looking at seroprevalence data where some people infected and others aren't, and x_k is the IgG value
 - $x_k \sim N(0, 1)$ with probability 0.5, and $x_k \sim N(10, 1)$, if we just fit a single parameter, we will be way off and we can check our predictions against the data to show that our model guess will never look like the data



- Simulations can be used for prediction
 - e.g. estimate epidemic model parameters up to time T and simulate forward using these estimates

Simulating Markov models

- Let the state space be denoted $S = \{1, 2, 3, \dots, n\}$
 - S is the set of states the MC can visit and each state can be multidimensional
- We care about the tendency of the chain to move from $i \rightarrow j$
 - $\Pr(X(t + dt) = j | X(t) = i) = q_{ij}dt + o(dt)$
- The chain stays in state i for time T_i
 - $T_i \sim \text{Exp}(\sum_{j \neq i} q_{ij})$
 - $P(T_i > t) = \exp(-\sum_{j \neq i} q_{ij}t)$
- When it leaves state i , the chain jumps to state j with probability $q_{ij} / \sum_{j \neq i} q_{ij}$
- The time spent in state i and the choice of where to jump to **are independent**, and they are also independent of the same quantities in other states and at other times



Here $q_{12} = 2.3$, $q_{21} = 8.0$ etc

If chain enters state 1, time spent there is

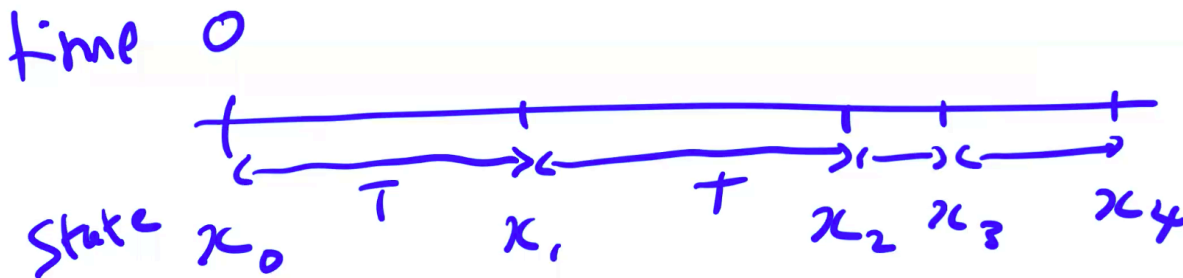
$$T_1 \sim \text{Exp}(2.3 + 1.4) = \text{Exp}(3.7)$$

$$P[\text{jump to state 2 next}] = \frac{2.3}{2.3 + 1.4}$$

Gillespie algorithm (next event simulation)

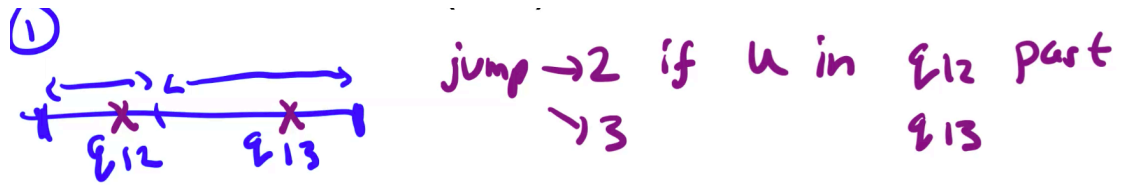
- All that is needed is to generate the time spent in a state, and the next state that is visited

Algorithm:



- Initialise ($t=0$, $X(0) = x_0$)
- for state i :
 - Calculate L (sum of the jump rates out of state i)
 - * $L = \sum_{j \neq i} q_{ij}$
 - Generate time spent in state i
 - * $T_i \sim \text{Exp}(L)$
 - Sample from a uniform distribution ($u = U[0, 1]$)
 - If $u < \frac{\sum_{j \leq 1, j \neq i} q_{ij}}{L}$
 - * $k = 1$
 - * Divide by L to normalize so it's between 0 and 1
 - Else if $u < \frac{\sum_{j \leq 2, j \neq i} q_{ij}}{L}$
 - * $k = 2$

- Continue until k determined



- Update current time

- Record t and k

* Time of next event = $t + T_i$

* State jumped to is k

- The algorithm outputs a sequence of times (t_k) and a corresponding sequence of states (x_k)

Example: general epidemic model (SIR)

- Due to the Poisson process infection mechanism and exponentially distributed infectious period
 - $\{[S(t), I(t)] : t \geq 0\}$
- If the chain is currently at (s, i) , then it can jump to:
 - $(s - 1, i + 1)$ (infection) at rate $\beta si/N$
 - $(s, i - 1)$ (removal) at rate γi
- Therefore, the time spent in (s, i)
 - $T_{(s,i)} \sim \text{Exp}([\beta si/N] + \gamma i)$
- Once the chain leaves (s, i)
 - Probability of infection $(s - 1, i + 1)$: $\frac{\beta s}{\beta s + N\gamma}$
 - Probability of recovery $(s, i - 1)$: $\frac{N\gamma}{\beta s + N\gamma}$
 - Calculate probabilities by dividing rate of interest by sum of the rates
- Apply the algorithm iteratively
- Can sometimes be useful to keep track of the type of each event e.g. infection or recovery

Simulating non-Markov models

- Same idea as before
 - Generating time until next event
 - But independence properties of Markov chain are lost so we need to explicitly generate the times of future events as the algorithm evolves
- In the Markov SIR model:
 - Infections occur according to a Poisson process of rate $\beta S_t I_t / N$
 - Each infective remains so for a period of time $T_I \sim \text{Exp}(\gamma)$
- In the non-Markov SIR model
 - A common generalisation is to let the infectious period distribution T_I be non-exponential e.g. constant, Gamma

- Infectious period T_I drawn from specified distribution with parameter vector θ
- Now two model parameters β and θ
- To simulate the epidemic
 - Generate removal time of each individual as they become infected
 - * The time of next removal is known, as is the identity of the removed individual
 - Generate possible time-to-next infection
 - * $T \sim \text{Exp}(\beta si/N)$
 - * If $T < \text{time of next removal}$, next event is an infection
 - Otherwise, next removal occurs
 - If removal occurs first, then i in $T \sim \text{Exp}(\beta si/N)$ changes, so we no longer have the correct distribution for time to next infection and it needs to be updated

Example: non-Markov SIR model

- Lets assume we have a fixed infectious period
 - $T_I = c$
- We need \vec{r} which contains the removal times of all the current infectives
- Initialize:
 - $S = N - 1$
 - $I = 1$
 - $t = 0$
 - $\vec{r} = (c)$
 - * We only have one infective, so \vec{r} only contains 1 removal time ($0 + c$)
- While $i > 0$:
 - $T \sim \text{Exp}(\beta SI/N)$
 - * Draw from the infectious period distribution
 - $R = \min(r)$
 - * Time of the next removal
 - if $t + T < R$ (current time + potential time to next infection < time of next removal)
 - * $S = S - 1, I = I + 1$
 - * Add new removal time $r \leftarrow (t + T + c)$
 - * Update current time $t = t + T$

Discussion

When infections and removals happen at the same time, do we consider to be Markov or non-Markov

Things are actually happening instantaneously, it's just that we make them discrete when we aggregate data.

In the non-Markov model example, the time to infection is exponential, so why is it not a Markov model

- The joint distribution of the number of susceptibles and infected at each time
 - Markov model is a markov chain
 - Not a markov chain in non-markov model as infectious period distribution is non-exponential
-