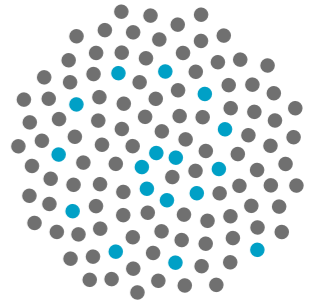


On Parameter Estimation for Markov Processes

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The problem

Often the most appropriate model for a stochastic system is that of a **discrete-state Markov process**.

In general, we can only observe the state of the process at successive, not necessarily equally-spaced, time points.

How can we estimate the parameters of our model from such observational data?

Epidemiological modelling example

The **stochastic SIS (susceptible-infective-susceptible) logistic** model is appropriate for modelling the spread of infections

- that do not confer any long lasting immunity, and
- where individuals become susceptible again after infection.

It is a continuous-time Markov chain ($m(t)$, $t \geq 0$) taking values in $S = \{0, 1, \dots, N\}$ with non-zero transition rates

$$q(m, m+1) = \lambda \frac{m}{N} (N-m), \quad m = 1, \dots, N-1,$$
$$q(m, m-1) = \mu m, \quad m = 1, \dots, N,$$

where λ , the per-interaction rate of infection, and μ , the per-individual rate of recovery, are both positive.

- Commonly we only know the number of people who are infected at successive, and not necessarily equally-spaced, time points.
- How do we estimate λ and μ , and the basic reproduction ratio $R_0 = \lambda/\mu$, from such observational data?

General approach

- Calculate the exact likelihood of observing the given data.
- Use a numerical search algorithm to compute the maximum likelihood estimators.
- The exact likelihood cannot usually be evaluated explicitly, so it must be computed numerically.

This combination provides a useful tool for fitting continuous-time Markov chains to real systems, but is computationally infeasible if the parameter space or the maximum population size is large.

Approximation approach

One way to achieve an approximate likelihood is to use diffusion approximations. If our model is **density-dependent**, that is the transition rates take the form

$$q_N(m, m+l) = Nf\left(\frac{m}{N}, l\right), \quad l \neq 0,$$

for a suitable function f , then we may derive a deterministic approximation and a Gaussian diffusion approximation for the density process $m(t)/N$, as N becomes large.

When the deterministic approximation has an **asymptotically stable fixed point**, we can accurately model the fluctuations of the density process about this fixed point by an **Ornstein-Uhlenbeck (OU) process**. The likelihood of an OU process is simply a Gaussian distribution.

Thus, we may approximate the exact likelihood by a Gaussian distribution, resulting in a substantial decrease in computational complexity.

Results

Comparison of the **General Approach** to the **Approximation Approach** for the stochastic SIS logistic model with $N = 50$ (number of individuals), $\lambda = 0.8$ (per-interaction rate of infection) and $\mu = 0.4$ (per-individual rate of recovery) and using one set of $n = 40$ observations:

| $n = 40$ | $\hat{\lambda}$ | $\hat{\mu}$ |
|------------------------|-----------------|-------------|
| General Approach | 1.14739 | 0.566294 |
| Approximation Approach | 1.10432 | 0.552027 |

Note that the maximum population size is small in the above comparison. When N is increased to 2000, where the General Approach is infeasible, and the estimates produced by the Approximation Approach improve:

| $n = 40$ | $\hat{\lambda}$ | $\hat{\mu}$ |
|------------------------|-----------------|-------------|
| Approximation Approach | 0.905564 | 0.447709 |

Our new method provides reasonably accurate estimates of parameters, in particular, when the maximum population size is large. This is precisely the situation in which the General Approach becomes infeasible, and thus the methods presented complement each other.

Reference

[1] Ross, J.V., Taimre, T. and Pollett, P.K. *On parameter estimation in population models*. Theoretical Population Biology (to appear).