MCMC Short Course - Examples

George Casella

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Introduction

- 1. For each of the following, plot the function in R. Use various values for μ, θ, σ and τ .
 - (a) The censored data density arising from observing

$$Z = X \wedge Y = \min(X, Y),$$

where

$$X \sim \mathcal{N}(\theta, \sigma^2)$$
 and $Y \sim \mathcal{N}(\mu, \tau^2)$,

which is given by

$$\begin{bmatrix} 1 - \Phi\left(\frac{z-\theta}{\sigma}\right) \end{bmatrix} \times \tau^{-1}\varphi\left(\frac{z-\mu}{\tau}\right) \\ + \begin{bmatrix} 1 - \Phi\left(\frac{z-\mu}{\tau}\right) \end{bmatrix} \sigma^{-1}\varphi\left(\frac{z-\theta}{\sigma}\right)$$

where φ and Φ are the density and cdf of the normal $\mathcal{N}(0, 1)$ distribution.

(b) The mixture of two normal distributions,

$$p\mathcal{N}(\theta,\sigma^2) + (1-p)\mathcal{N}(\mu,\tau^2)$$
,

with density

$$p\sigma^{-1}\varphi\left(\frac{x-\theta}{\sigma}\right) + (1-p)\tau^{-1}\varphi\left(\frac{x-\mu}{\tau}\right)]$$

(c) The likelihood based on observing $(X_1, X_2, X_3) = (0, 5, 9)$ from the Student's t density proportional to

$$\sigma^{-1} \left(1 + \frac{(x-\theta)^2}{p\sigma^2} \right)^{-(p+1)/2} ,$$

with p = 1 and $\sigma = 1$ (the standard Cauchy).

• Plot.txt

Random Variable Generation

- 2. Check the R uniform random number generator:
 - (a) Generate 1,000 uniform random variables and make a histogram

(b) Generate uniform random variables (X_1, \ldots, X_n) and plot the pairs (X_i, X_{i+1}) to check for autocorrelation.

• Uniform.txt

3. (a) Generate a binomial (n, p) random variable with n = 25 and p = .2. Make a histogram and compare it to the binomial mass function, and to the R binomial generator.

• binomial.txt

(b) Generate 5,000 *logarithmicseries* random variables with mass function

$$P(X = x) = \frac{-(1-p)^x}{x \log p}, \quad x = 1, 2, \dots \quad 0$$

Make a histogram and plot the mass function.

• logarithmic.txt

- 4. In each case generate the random variables and compare to the density function
 - (a) Normal random variables using a Cauchy candidate in Accept/Reject
 - (b) Gamma(4.3, 6.2) random variables using a Gamma(4, 7).
 - (c) Truncated normal Standard normal truncated to $(2,\infty)$

• RandomVariables.txt

Integration

5. For the Bayes estimator

$$\delta(x) = \frac{\int_{-\infty}^{\infty} \frac{\theta}{1+\theta^2} e^{-(x-\theta)^2/2} d\theta}{\int_{-\infty}^{\infty} \frac{1}{1+\theta^2} e^{-(x-\theta)^2/2} d\theta}$$

- (a) Plot the integrand and use MCI to calculate the integral.
- (b) Monitor the convergence with the standard error of the estimate. Obtain three digits of accuracy with probability .95.

• BayesEstimator.txt

6. For a standard normal random variable Z, calculate P(Z > 2.5) using

- (a) Monte Carlo sums based on indicator functions
- (b) Importance sampling based on a candidate exponential $f(x) = e^{-(x-2.5)}$, x > 2.5.

Note that P(Z > 2.5) = .0062.

• TailProb.txt

Optimization

7. Use the R functions "optim" and "optimize" to find the maximum of

$$f(x) = \left[\cos(50x) + \sin(20x)\right]^2$$
.

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Compare to the results of a stochastic exploration.

• Optimize.txt

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- 8. For each of the likelihood functions in Exercise 1, find the maximum with "optimize" and stochastic exploration.
- 9. The follow are genotype data on blood type

Genotype	Probability	Observed	Probability	Frequency
AA	p_A^2	A	$p_A^2 + 2p_A p_O$	$n_A = 186$
AO	$2p_A p_O$			
BB	p_B^2	В	$p_B^2 + 2p_B p_O$	$n_B = 38$
BO	$2p_Bp_O$			
AB	$2p_A p_B$	AB	$p_A p_B$	$n_{AB} = 13$
00	p_O^2	0	p_O^2	$n_O = 284$

Because of dominance, we can only observe the genotype in the third column, with probabilities given by the fourth column. The interest is in estimating the allele frequencies p_A, p_B , and p_O (which sum to 1).

(a) Under a multinomial model, verify that the observed data likelihood is proportional to

$$(p_A^2 + 2p_A p_O)^{n_A} (p_B^2 + 2p_B p_O)^{n_B} (p_A p_B)^{n_{AB}} (p_O^2)^{n_O}$$

(b) With missing data Z_A and Z_B , verify the complete data likelihood

$$(p_A^2)^{Z_A} (2p_A p_O)^{n_A - Z_A} (p_B^2)^{Z_B} (2p_B p_O)^{n_B - Z_B} (p_A p_B)^{n_{AB}} (p_O^2)^{n_C}$$

(c) Verify that the missing data distribution is

$$Z_A \sim \text{binomial}\left(n_A, \frac{p_A^2}{p_A^2 + 2p_A p_O}\right) \text{ and } Z_B \sim \text{binomial}\left(n_B, \frac{p_B^2}{p_B^2 + 2p_B p_O}\right),$$

and write an EM algorithm to estimate p_A, p_B , and p_O

• EMBlood.txt

Metropolis-Hastings

- 10. Calculate the mean of a Gamma(4.3, 6.2) random variables using
 - (a) Accept-Reject with a Gamma(4,7) candidate.
 - (b) Metropolis-Hastings with a Gamma(4,7) candidate.
 - (c) Metropolis-Hastings with a Gamma(5,6) candidate.

In each case monitor the convergence.

• GammaAR.txt

11. The Institute for Child Health Policy (ICHP) at the University of Florida studies the effects of health policy decisions on children's health. A small portion of one of their studies follows.

The overall health of a child (metq) is rated on a 1-3 scale, with 3 being the worst. Each child is in an HMO (variable np, 1=nonprofit, -1=for profit). The dependent variable of interest (y_{ij}) is the use of an emergency room (erodds, 1=used emergency room, 0=did not). The question of interest is whether the status of the HMO affects the emergency room choice.

(a) An appropriate model is logistic regression

$$logit(p_{ij}) = a + bx_i + cz_{ij}, \quad i = 1, ..., k, \quad j = 1, ..., n_i.$$

where x_i is the HMO type, z_{ij} is the health status of the child, and p_{ij} is the probability of using an emergency room. Verify that the likelihood function is

$$\prod_{i=1}^{k} \prod_{j=1}^{n_i} \left(\frac{\exp(a + bx_i + cz_{ij})}{1 + \exp(a + bx_i + cz_{ij})} \right)^{y_{ij}} \left(\frac{1}{1 + \exp(a + bx_i + cz_{ij})} \right)^{1-y_{ij}}$$

(Here we are only distinguishing between for-profit and non-profit, so k = 2.)

- (b) Run a standard GLM on these data (at LogisticData.txt) and get the estimated mean and variance of a, b, and c.
- (c) Use normal candidate densities with mean and variance at the estimates in a Metropolis-Hastings algorithm that samples from the likelihood. Get histograms of the parameter values.

• MHLogistic.txt

Gibbs Sampling

12. Referring to Exercise 9, estimate p_A, p_B and p_O using a Gibbs sampler. Make a histogram of the samples.

• GibbsBlood.txt

13. A subset of the clinical mastitis data is

 $\begin{array}{l} 0, 0, 1, 1, 2, 2, 2, 2, 2, 2, 2, 4, 4, 4, 5, 5, 5, 5, 5, 5, 5, 6, \\ 6, 8, 8, 8, 9, 9, 9, 10, 10, 12, 12, 13, 13, 13, 13, 18, \\ 18, 19, 19, 19, 19, 20, 20, 22, 22, 22, 23, 25 \end{array}$

For each herd we model the mean number of occurrences as a Poisson mean λ_i , with a hierarchy to account for overdispersion. A model is

$$X_i \sim \text{Poisson}(\lambda_i),$$

$$\lambda_i \sim \text{Gamma}(\alpha, \beta_i),$$

$$\beta_i \sim \text{Gamma}(a, b),$$

where α , a, and b are specified. The posterior density of λ_i , $\pi(\lambda_i | \mathbf{x}, \alpha)$, can now be obtained from the Gibbs sampler

$$\lambda_i \sim \pi(\lambda_i | \mathbf{x}, \alpha, \beta_i) = \text{Gamma}(x_i + \alpha, 1 + \beta_i),$$

$$\beta_i \sim \pi(\beta_i | \mathbf{x}, \alpha, a, b, \lambda_i) = \text{Gamma}(\alpha + a, \lambda_i + b).$$

- (a) For $\alpha = .1, a = b = 1$ run the Gibbs sampler. Make histograms and monitor the convergence of λ_5 , λ_{15} , and β_{15}
- (b) Investigate the sensitivity of your answer to the specification of α, a and b.

• Mastitis.txt