EEEB UN3005/GR5005Homework - Week 05 - Due 05 Mar 2019

Xun Zhao, xz2827

Homework Instructions: Complete this assignment by writing code in the code chunks provided. If required, provide written explanations below the relevant code chunks. Replace "USE YOUR NAME HERE" with your name in the document header. When complete, knit this document within RStudio to generate a pdf. Please review the resulting pdf to ensure that all content relevant for grading (i.e., code, code output, and written explanations) appears in the document. Rename your pdf document according to the following format: hw_week_05_firstname_lastname.pdf. Upload this final homework document to CourseWorks by 5 pm on the due date.

Problem 1 (3 points)

Think back to the research scenario described in last week's homework assignment: you're studying a bacterial pathogen of small mammals, and your pilot sampling efforts have found 9 infected animals out of 20 animals sampled. Much like last week, use grid approximation to construct the posterior for the probability of infection parameter (p). Use 1,000 points in your grid approximation, and assume a flat prior.

Now, let's take our posterior inference a bit further. Generate 10,000 samples from this posterior, assigning them to samples1. What is the mean value of samples1? What is the 90% HPDI of samples1?

```
grid = seq(0, 1, length.out = 1000)
prior = rep(1, 1000)
likelihood = dbinom(9, 20, prob = grid)
unstd.post = likelihood * prior
post = unstd.post / sum(unstd.post)
samples1 = sample(grid, 10000, replace = TRUE, prob = post)
print(mean(samples1))
```

```
## [1] 0.45434
print(HPDI(samples1, prob = 0.9))
```

```
## |0.9 0.9|
## 0.2822823 0.6206206
```

Problem 2 (2 points)

Now imagine that by reading the primary scientific literature, you find that prevalence of this particular bacterial pathogen in similar small mammal populations has never been reported above 0.5. Use grid approximation to construct a posterior distribution for the probability of infection parameter, this time with a prior that assumes that the probability of infection should be < 0.5. Note, a prior that is a constant below p = 0.5 and 0 above p = 0.5 is a mathematical representation of this assumption.

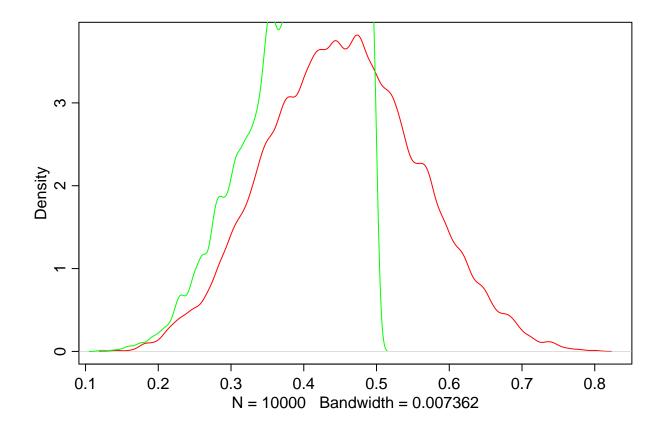
Generate 10,000 samples from this new posterior distribution (call them samples2). What is the mean value of samples2? What is the 90% HPDI of samples2?

Problem 3 (2 points)

Using the dens() function in the rethinking package, plot samples1 and samples2 together for visual comparison. The add = TRUE argument to your second dens() call will allow you to plot the samples overlaid. Using different colors for the two different vectors of posterior samples may also help distinguish them. It's just another argument in your dens() call (i.e., col = "red").

However you choose to depict them, using this visual comparison as an aid, how does the posterior represented by samples2 differ from samples1? What difference does the change in prior make?

```
dens(samples1, col = 'red')
dens(samples2, add = TRUE, col = 'green')
```



Problem 4 (2 points)

With further literature research, you discover that the true probability of infection is also extremely unlikely to be < 0.2. Modify your grid approximation to include a prior that represents the assumption that 0.2 . Generate 10,000 new posterior samples for <math>p using this prior (call these samples samples3).

What is the mean value of samples3? What is the 90% HPDI of samples3?

Does samples3 differ strongly from samples2? Why or why not? As in the previous problem, visual comparison of the posterior distributions may help significantly in answering these questions.

Problem 5 (1 points)

Assume that through further study you establish that the true probability of infection within the population is 0.3. Given this was the true probability of infection value, what was the probability of you initially observing 9 infected individuals out of 20 in your pilot study?