

ADVANCED BAYESIAN MODELING

BINOMIAL HIERARCHICAL MODEL IN R/JAGS:
RAT TUMOR RESULTS

Recall:

- ▶ Rat tumor hierarchical model with (diffuse) independent exponential hyperpriors
- ▶ Run JAGS model in R using `rjags` package
- ▶ `x` has posterior samples of α and β .

Now to analyze results ...

```
> summary(x)
```

```
Iterations = 3501:13500
```

```
Thinning interval = 1
```

```
Number of chains = 1
```

```
Sample size per chain = 10000
```

1. Empirical mean and standard deviation for each variable,
plus standard error of the mean:

	Mean	SD	Naive SE	Time-series SE
alpha	3.427	1.360	0.01360	0.1453
beta	20.496	8.053	0.08053	0.8360

2. Quantiles for each variable:

	2.5%	25%	50%	75%	97.5%
alpha	1.595	2.447	3.156	4.09	6.871
beta	9.722	14.599	18.946	24.40	41.474

Information from summary:

$$E(\alpha \mid y) \approx 3.4$$

$$\sqrt{\text{var}(\alpha \mid y)} \approx 1.4$$

$$E(\beta \mid y) \approx 20$$

$$\sqrt{\text{var}(\beta \mid y)} \approx 8$$

Approx. 95% central posterior intervals:

$$\alpha : (1.6, 6.9)$$

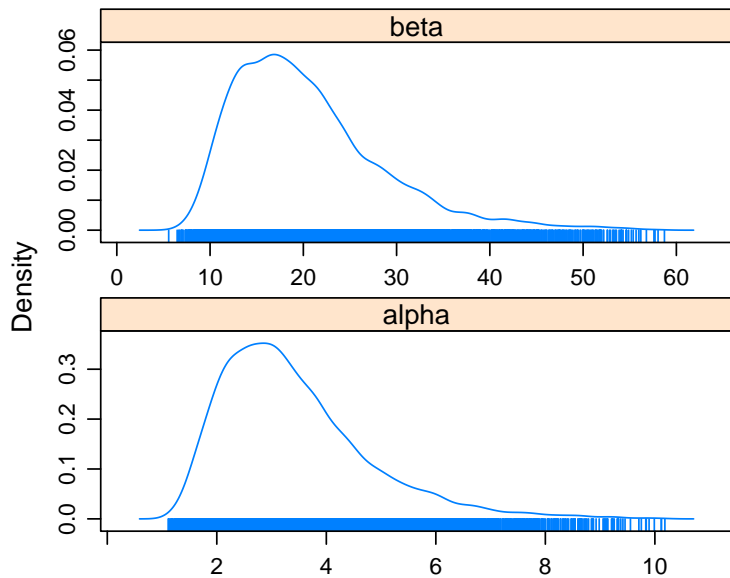
$$\beta : (9.7, 41)$$

Get estimated posterior densities of α and β :

```
> require(lattice)
```

Loading required package: lattice

```
> densityplot(x)
```

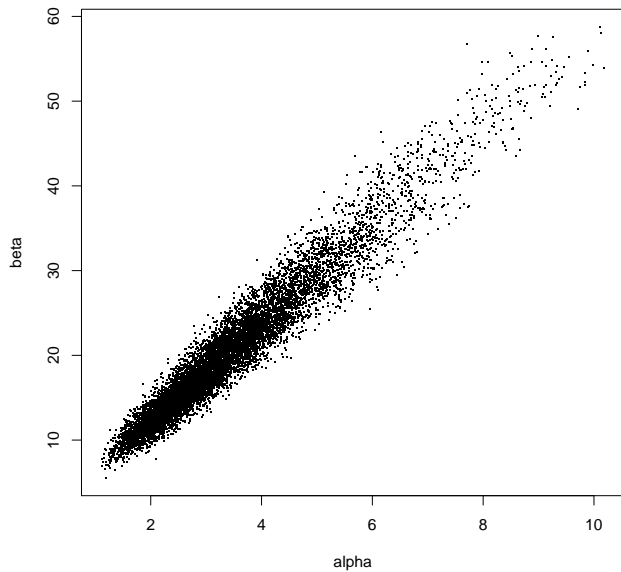


Examine joint posterior distribution of α and β :

```
> alpha <- as.matrix(x)[,"alpha"]
```

```
> beta <- as.matrix(x)[,"beta"]
```

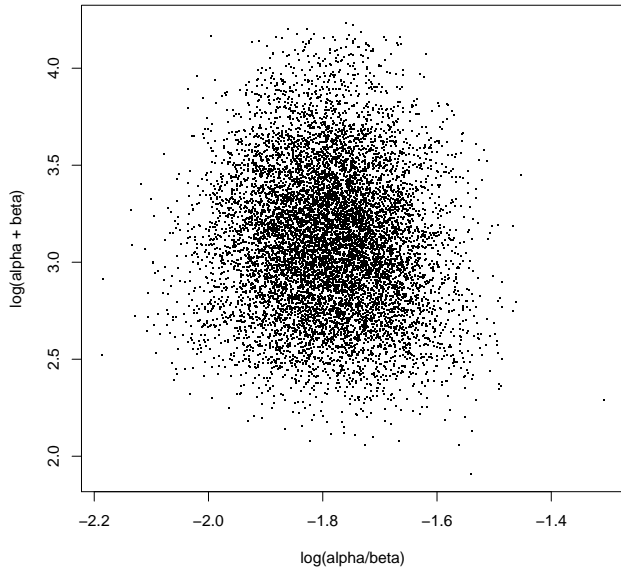
```
> plot(alpha, beta, pch=".", cex=2)
```

May be more meaningful to graph α/β and $\alpha + \beta$, and to use log scales:

```
> plot(log(alpha/beta), log(alpha+beta), pch=".", cex=2)
```

(See BDA3, Sec. 5.3.)



Were exponential hyperpriors diffuse enough?

Try different (less informative?) prior ...

Alternative Model

Recall flat hyperprior proposed in BDA3, Sec. 5.3:

$$p(\phi_1, \phi_2) \propto 1 \quad \phi_1 = \frac{\alpha}{\alpha + \beta} \in (0, 1) \quad \phi_2 = (\alpha + \beta)^{-1/2} \in (0, \infty)$$

Solving,

$$\alpha = \phi_1 / \phi_2^2 \quad \beta = (1 - \phi_1) / \phi_2^2$$

Approximate that improper hyperprior by a wide but proper one:

$$\phi_1 \sim \text{U}(0, 1) \quad \phi_2 \sim \text{U}(0, 1000) \quad \text{independent}$$

File rattumor2.bug:

```
model {  
  
  for (j in 1:length(y)) {  
    y[j] ~ dbin(theta[j], N[j])  
    theta[j] ~ dbeta(alpha, beta)  
  }  
  
  alpha <- phi1 / phi2^2  
  beta <- (1-phi1) / phi2^2  
  
  phi1 ~ dunif(0,1)  
  phi2 ~ dunif(0,1000)  
  
}
```

Try to run JAGS (in R using rjags) as before:

```
> m <- jags.model("rattumor2.bug", d)
```

```
Compiling model graph
```

```
  Resolving undeclared variables
```

```
  Allocating nodes
```

```
Graph information:
```

```
  Observed stochastic nodes: 71
```

```
  Unobserved stochastic nodes: 73
```

```
  Total graph size: 223
```

```
Initializing model
```

```
|+++++
```

```
| 26%
```

```
Error: Error in node phi1
```

```
Slicer stuck at value with infinite density
```

Origin of error is obscure, relating to misbehavior of a built-in “sampler” in JAGS.

Three options:

- ▶ Make hyperprior on ϕ_2 more informative:

```
phi2 ~ dunif(0,10)
```

- ▶ Truncate the beta distribution away from its problematic endpoints:

```
theta[j] ~ dbeta(alpha, beta) T(0.0001,0.9999)
```

- ▶ Turn off the sampler causing the problem.

We choose the third option. Turning off a sampler generally causes JAGS to fall back on another sampler that may not have the same problem.

In R:

```
> set.factory("bugs::BinomSlice","sampler",FALSE)
```

Now try again:

```
> m <- jags.model("rattumor2.bug", d)
```

```
...
```

```
> update(m, 2500)
```

```
|*****| 100%
```

```
> x <- coda.samples(m, c("alpha","beta"), n.iter=10000)
```

```
|*****| 100%
```

Plot posterior jointly distributed samples of α/β and $\alpha + \beta$, on log scales:

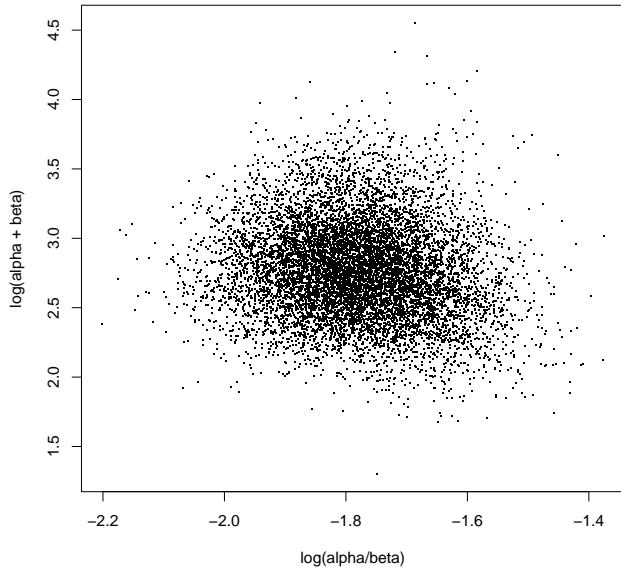
```
> alpha <- as.matrix(x)[,"alpha"]
```

```
> beta <- as.matrix(x)[,"beta"]
```

```
> plot(log(alpha/beta), log(alpha+beta), pch=".", cex=2)
```

Compare BDA3, Fig. 5.3.

(Also compare with previous model results – try yourself.)

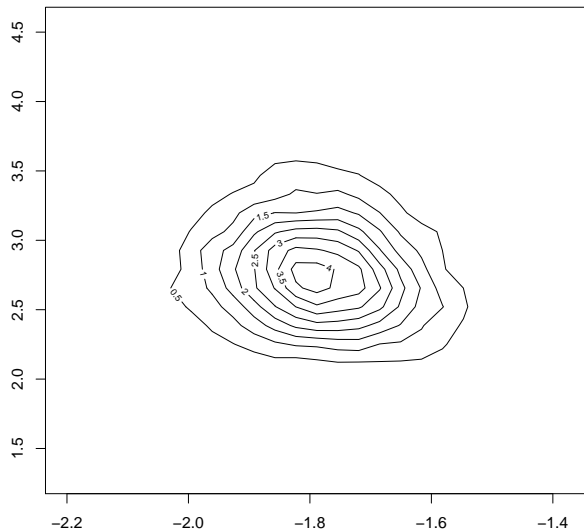


Try also a contour plot of estimated joint posterior density:

```
> library(MASS)
```

```
> contour(kde2d(log(alpha/beta), log(alpha+beta)))
```

Compare BDA3, Fig. 5.3.



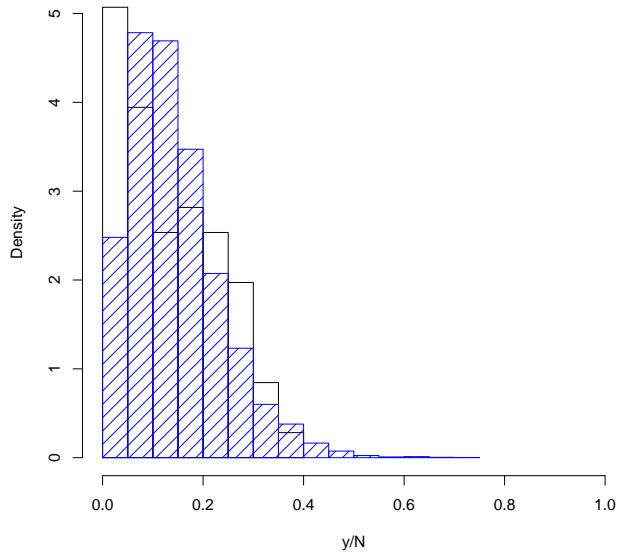
Now consider the posterior predictive distribution of $\tilde{\theta}$, the tumor probability for a “new” experiment, exchangeable with the others.

Can directly simulate $\tilde{\theta}$ using posterior α and β samples:

```
> thetatilde <- rbeta(10000, alpha, beta)
```

Then plot as histogram, on same plot with naive empirical histogram:

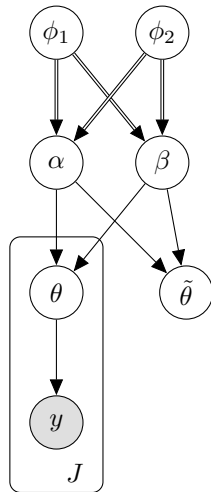
```
> with(d, hist(y/N, freq=FALSE, main="", xlim=c(0,1)))  
> hist(thetatilde, freq=FALSE, density = 10, col="blue", border="blue",  
+      add=TRUE)
```



We can add $\tilde{\theta}$ to the DAG.

Note that it is a single (scalar) node with no observed descendants. It is conditionally independent of θ , given its parents (α and β).

Since $\tilde{\theta}$ is just another node, we can alternatively simulate it using JAGS ...



```

model {

  for (j in 1:length(y)) {
    y[j] ~ dbin(theta[j], N[j])
    theta[j] ~ dbeta(alpha, beta)
  }

  thetatilde ~ dbeta(alpha, beta)

  alpha <- phi1 / phi2^2
  beta <- (1-phi1) / phi2^2

  phi1 ~ dunif(0,1)
  phi2 ~ dunif(0,1000)

}

```

Remark:

Since JAGS uses simulation, seeded differently each time, you should expect slightly different results in each run.

Setting an R seed does *not* set the JAGS seed.

To set the JAGS seed, see the JAGS manual.