

2022 SISG Bayesian Statistics for Genetics R Notes: Binomial Sampling

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Binomial Data

Introduction

In these notes, in the context of binomial sampling, we look at:

- Specifying a prior distribution
- Prediction.
- Testing.
- Logistic regression.

We analyze allele specific expression (ASE) data, and low birth weight data.

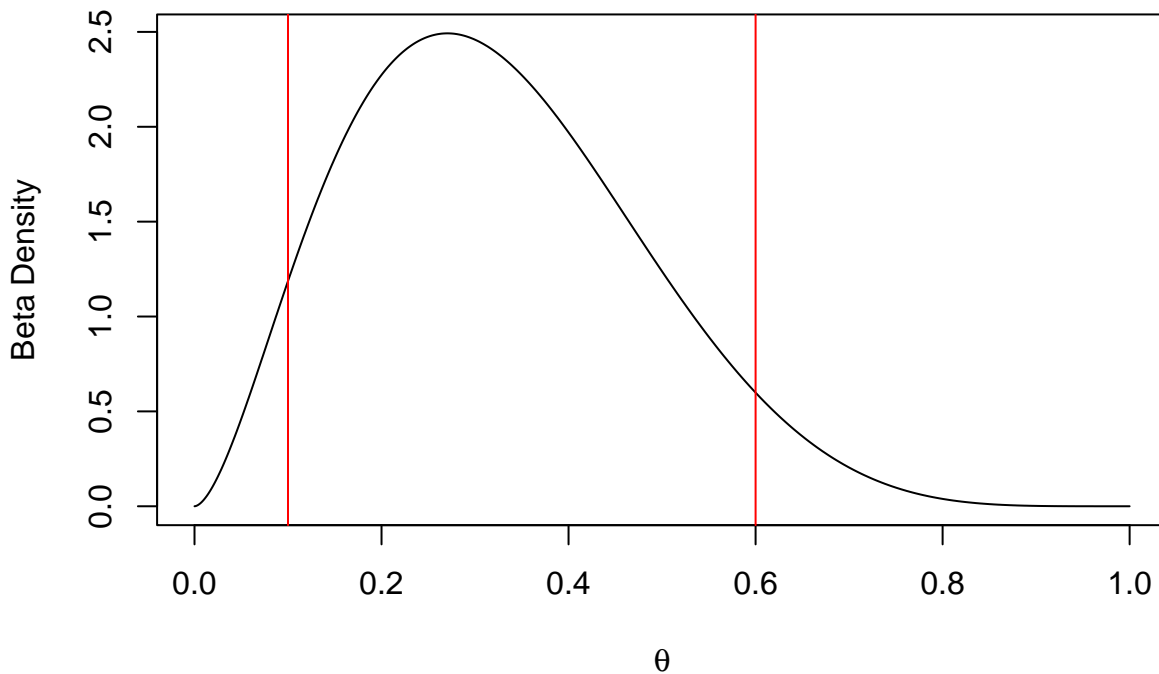
Specifying a prior distribution

The code below finds the beta distribution, i.e. the a and the b , with 5% and 95% points of 0.1 and 0.6.

```
# Function to find a and b
priorch <- function(x,q1,q2,p1,p2){
  (p1-pbeta(q1,x[1],x[2]))^2 + (p2-pbeta(q2,x[1],x[2]))^2 }
p1 <- 0.05
p2 <- 0.95
q1 <- 0.1
q2 <- 0.6
opt <- optim(par=c(1,1),fn=priorch,q1=q1,q2=q2,p1=p1,p2=p2,
             control=list(abstol=1e-8))
cat("a and b are ",opt$par,"\n")
## a and b are 2.730616 5.667462
```

The code below finds the beta distribution, i.e. the a and the b , with 5% and 95% points of 0.1 and 0.6.

```
probvals <- seq(0,1,.001)
plot(probvals,dbeta(probvals,shape1=opt$par[1],shape2=opt$par[2]),
     type="l", xlab=expression(theta),ylab="Beta Density")
abline(v=q1,col="red")
abline(v=q2,col="red")
```



Differences in Binomial Proportions

We consider an example in which we wish to compare allele frequencies between two populations.

Let θ_1 and θ_2 be the allele frequencies in the NE and US population from which the samples were drawn, respectively.

The allele frequencies were 10.69% and 13.21% with sample sizes of 650 and 265, in the NE and US samples, respectively.

We assume independent $\text{Beta}(1,1)$ priors on each of θ_1 and θ_2 .

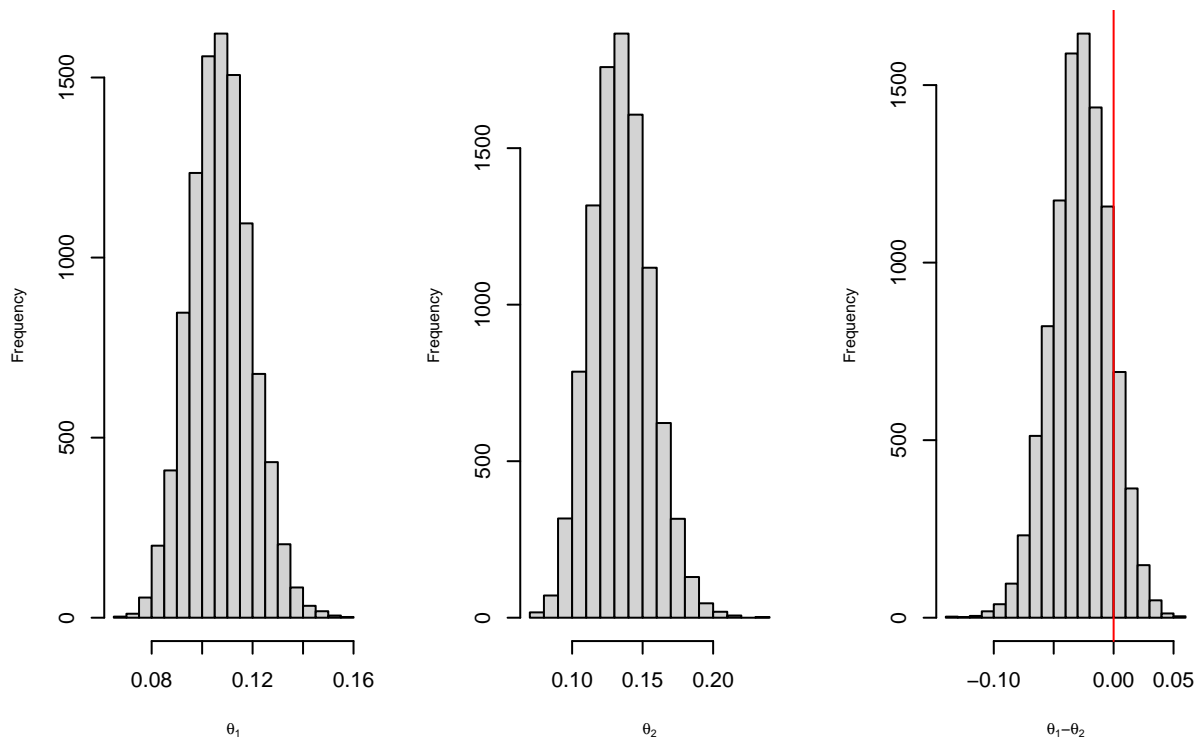
The y_1 and y_2 data (i.e. the numbers with the allele in the two populations) were reconstructed from figures in the original paper in which only the denominators and the frequencies were given, hence the `floor` function.

```
N1 <- 650
y1 <- floor(N1*.1069)
N2 <- 265
y2 <- floor(N2*.1321)
nsamp <- 10000
a <- b <- 1
post1 <- rbeta(nsamp,y1+a,N1-y1+b)
post2 <- rbeta(nsamp,y2+a,N2-y2+b)
```

The key step is in constructing a sample estimate of the difference in probabilities $\theta_1 - \theta_2$.

```
par(mfrow=c(1,3))
hist(post1,xlab=expression(theta[1]),main="",cex.lab=0.7)
hist(post2,xlab=expression(theta[2]),main="",cex.lab=0.7)
#
hist(post1-post2,xlab=expression(paste(theta[1],"-",
theta[2])),main="",cex.lab=0.7)
abline(v=0,col="red")
```

```
sum(post1-post2>0)/nsamp
## [1] 0.1269
```



ASE Data

Analysis of ASE data

```
download.file("http://faculty.washington.edu/kenrice/sisgbayes/ASEgene.txt",
             destfile = "ASEgene.txt")
ASEdat <- read.table("ASEgene.txt",header=TRUE)
head(ASEdat)
##      Y      N
## 1  62   107
## 2  33    59
## 3 658  1550
## 4  14    61
## 5  57   153
## 6 218   451
dim(ASEdat)
## [1] 4844    2
ngenes <- dim(ASEdat)[1]
pvals <- NULL
for (i in 1:ngenes){
  pvals[i] <- binom.test(ASEdat$Y[i],ASEdat$N[i],
                        p=0.5,alternative="two.sided")["p.value"]
}
```

```
# Function to evaluate Bayes factors for a binomial
# likelihood and beta prior, and a point null at p0
```

```

BFbinomial <- function(N,y,a,b,p0){
  logPrH0 <- lchoose(N,y) + y*log(p0) + (N-y)*log(1-p0)
  logPrH1 <- lchoose(N,y) + gamma(a+b) - lgamma(a) - lgamma(b) + lgamma(y+a) + lgamma(N-y+b) -lgam
  logBF <- logPrH0 - logPrH1
  list(logPrH0=logPrH0,logPrH1=logPrH1,logBF=logBF)
}

nsim <- 5000
a <- 1
b <- 1
p0 <- 0.5

postprob <- logBFR <- rep(0,ngenes)
pcutoff <- 0.05/length(pvals)
for (i in 1:ngenes){
  BFcall <- BFbinomial(ASEdat$N[i],ASEdat$Y[i],a,b,p0)
  logBFR[i] <- -BFcall$logBF
  postprob[i] <- pbeta(0.5,a+ASEdat$Y[i],b+ASEdat$N[i]-ASEdat$Y[i])
}
cat("log BFR > log(150) = ",sum(logBFR>log(150)),"\n")
## log BFR > log(150) = 197
cat("log BFR > log(20) = ",sum(logBFR>log(20)),"\n")
## log BFR > log(20) = 359
cat("p-values > ",pcutoff,sum(pvals<pcutoff),"\n")
## p-values > 1.032205e-05 111
cat("postprobs < 0.01 and > 0.99 ",sum(postprob<0.01),sum(postprob>0.99),"\n")
## postprobs < 0.01 and > 0.99 278 242

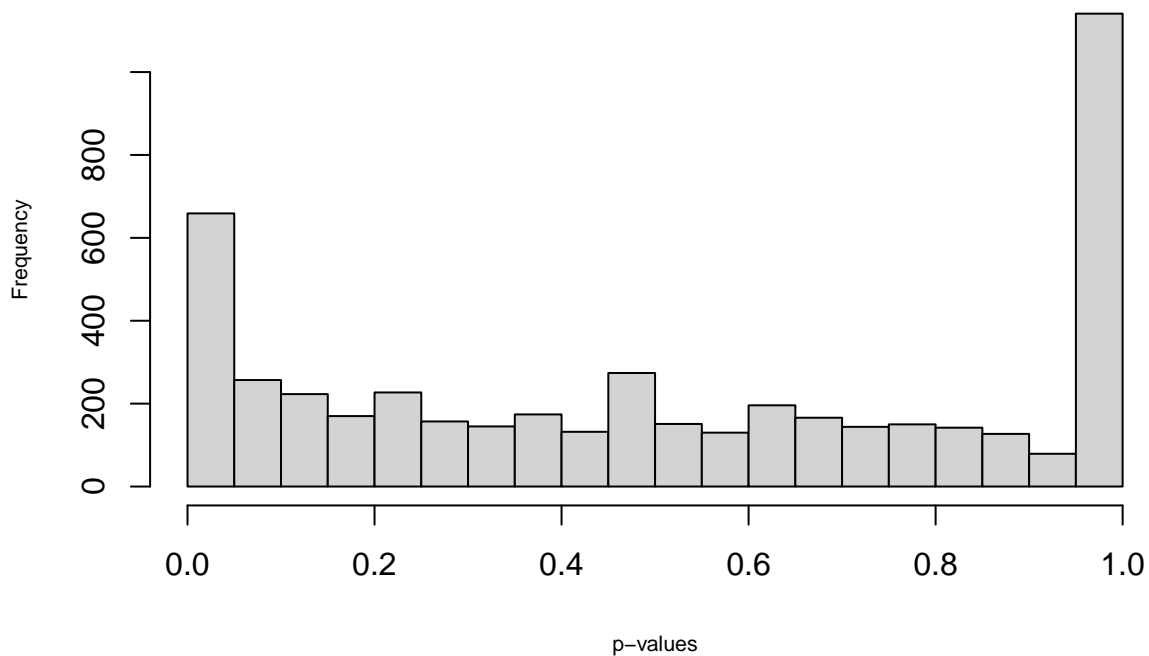
```

Histogram of p -values for ASE data

```

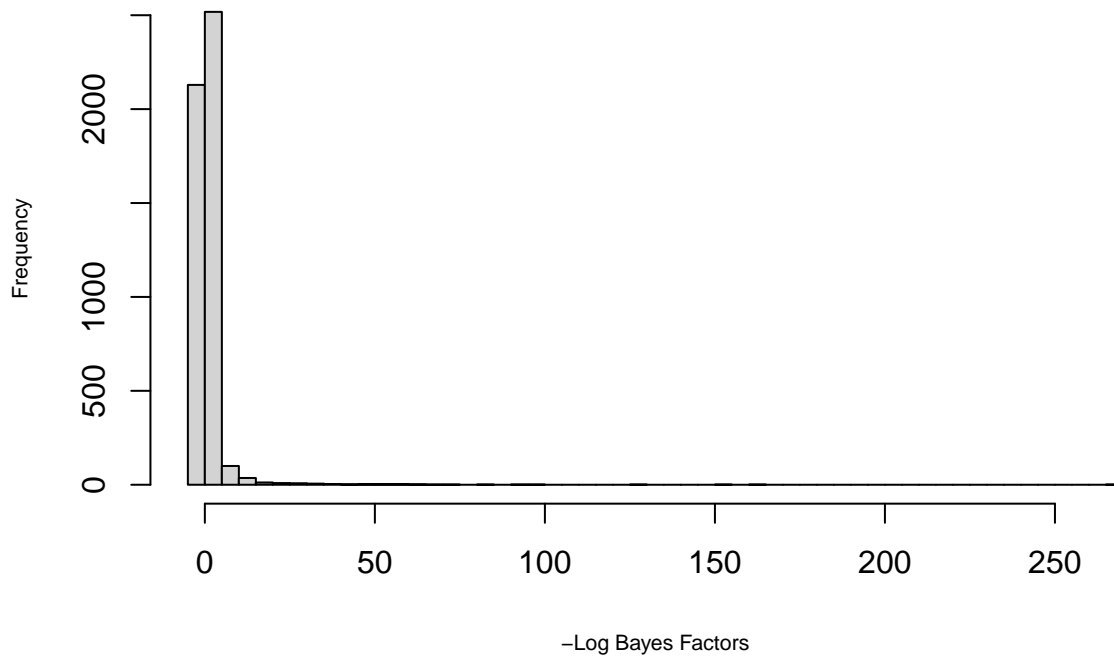
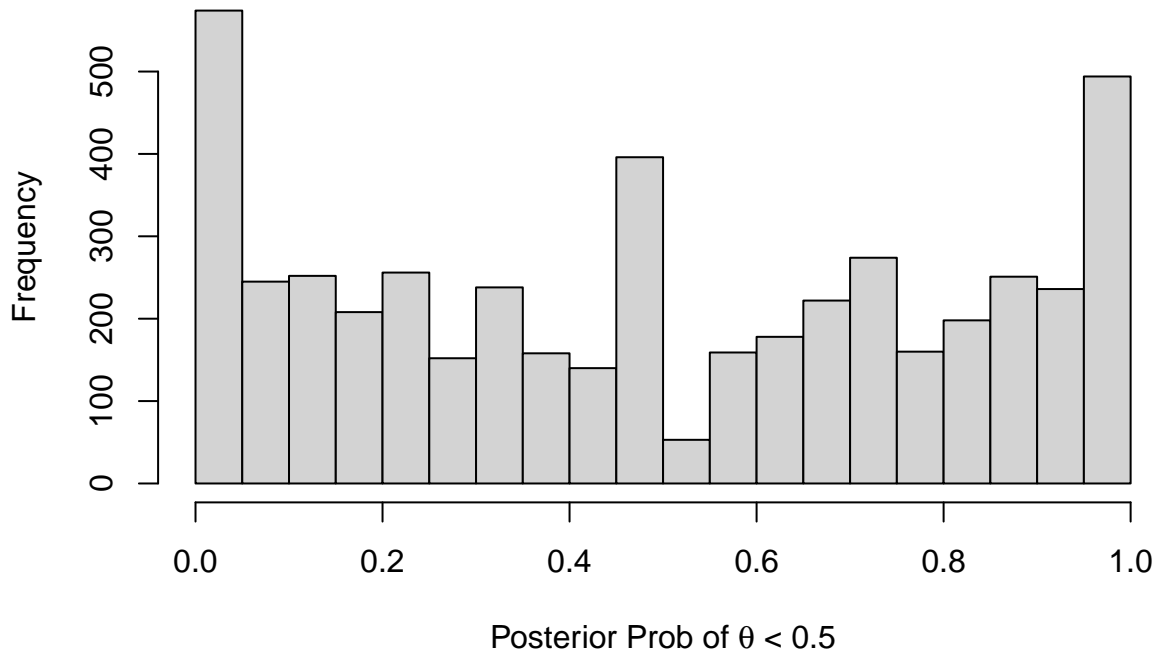
hist(pvals,xlab="p-values",main="",nclass=20,cex.lab=0.7,pch=16,cex=0.5)

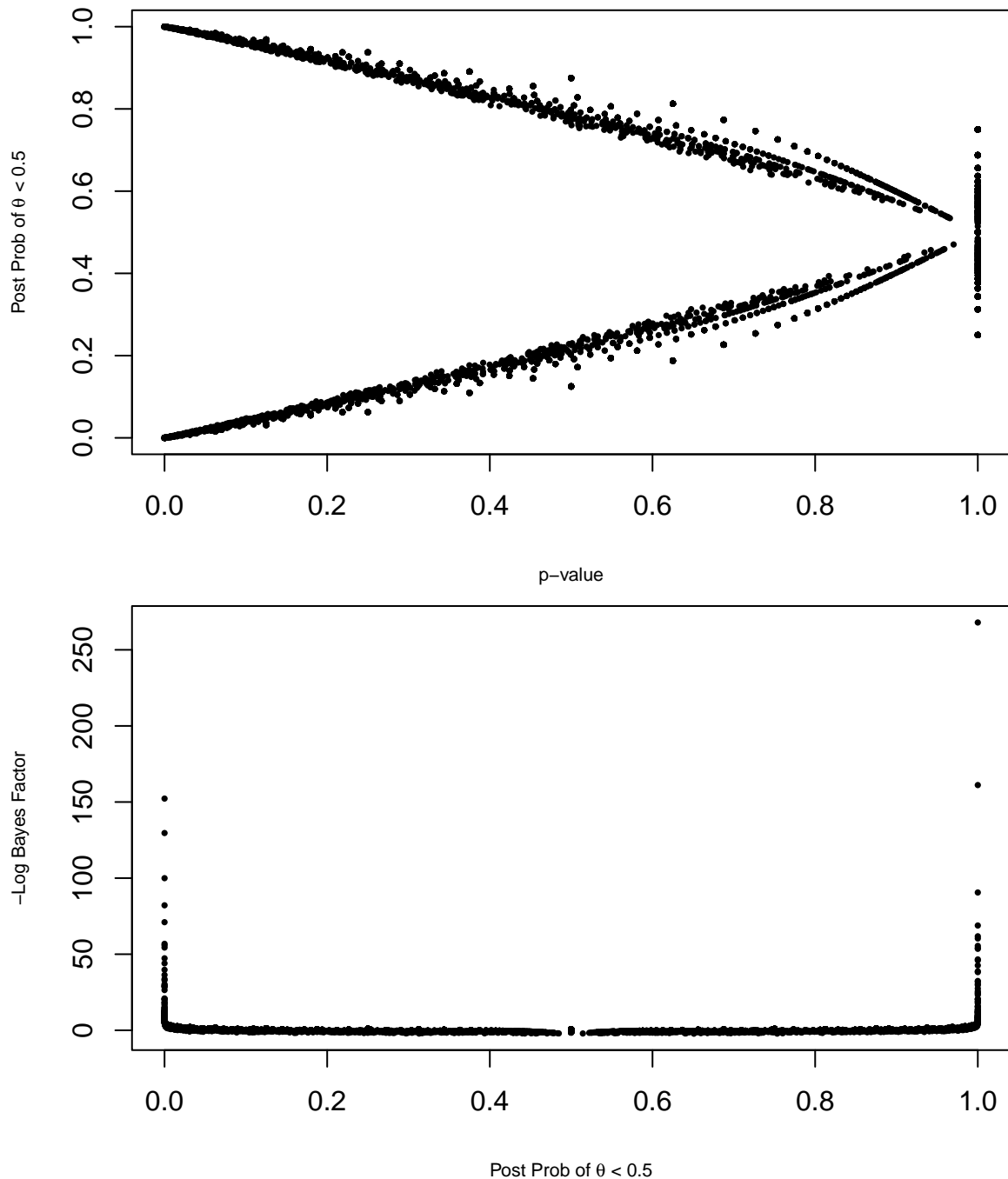
```



Histogram of posterior probabilities for ASE data

```
hist(postprob,nclass=20,xlab=expression(paste("Posterior Prob of ",theta," < 0.5")),main="",cex.label=0
```





Logistic Regression

Birth weight analysis

Hosmer and Lemeshow (2004) present data on 189 births to women participating in a larger study at Baystate Medical Center (Springfield, MA), with information on the following variables:

- LOW: Low birth weight; ($0 = \geq 2500\text{g}$; $1 = < 2500\text{g}$)
- AGE: Mother's age
- LWT: Mother's weight
- RACE: Listed race of mother; ($1 = \text{white}$; $2 = \text{black}$; $3 = \text{other}$)

- SMOKE: Smoking status during pregnancy; (0 = no; 1 = yes)
- HT: History of hypertension; (0 = no; 1 = yes)
- UI: Presence of uterine irritability; (0 = no; 1 = yes)
- FTV: Number of physician visits during the first trimester.

This data (`lowbwt`) can be found in the `brinla` package, which contains code and data from *Bayesian Regression with INLA* by Wang et al. (2018).

Birth weight analysis: load packages and data

```
# devtools::install_github("julianfaraway/brinla")
# remotes::install_github("julianfaraway/brinla")
library(brinla)
library(dplyr)
library(tidyr)
data(lowbwt)
knitr::opts_chunk$set(dev = 'pdf')
```

Frequentist logistic regression with glm

```
bwt.glm <- glm(LOW ~ AGE + LWT + RACE + SMOKE + HT + UI + FTV,
              data=lowbwt, family=binomial())
round(coef(summary(bwt.glm)), 3)
##               Estimate Std. Error z value Pr(>|z|)
## (Intercept)    0.455      1.185   0.384   0.701
## AGE           -0.021      0.036  -0.570   0.568
## LWT           -0.017      0.007  -2.409   0.016
## RACE2          1.290      0.528   2.445   0.015
## RACE3          0.919      0.436   2.106   0.035
## SMOKE1         1.042      0.395   2.634   0.008
## HT1            1.885      0.695   2.713   0.007
## UI1            0.904      0.449   2.015   0.044
## FTV            0.059      0.172   0.344   0.731
```

Logistic regression with INLA (default prior)

```
bwt.inla <- inla(LOW ~ AGE + LWT + RACE + SMOKE + HT + UI + FTV,
               data=lowbwt, family = "binomial", Ntrials = 1,
               control.compute = list(dic = TRUE, cpo = TRUE))
round(bwt.inla$summary.fixed[,1:5], 3)
##               mean      sd 0.025quant 0.5quant 0.975quant
## (Intercept)  0.567 1.186    -1.727    0.555    2.925
## AGE         -0.021 0.036    -0.092   -0.020    0.049
## LWT         -0.018 0.007    -0.032   -0.017   -0.005
## RACE2        1.340 0.528     0.316    1.336    2.386
## RACE3        0.946 0.436     0.104    0.941    1.816
## SMOKE1       1.075 0.395     0.315    1.070    1.867
## HT1          1.973 0.694     0.663    1.955    3.391
## UI1          0.933 0.449     0.052    0.933    1.813
## FTV          0.056 0.172    -0.289    0.059    0.386
```

Logistic regression example: comparison of estimates

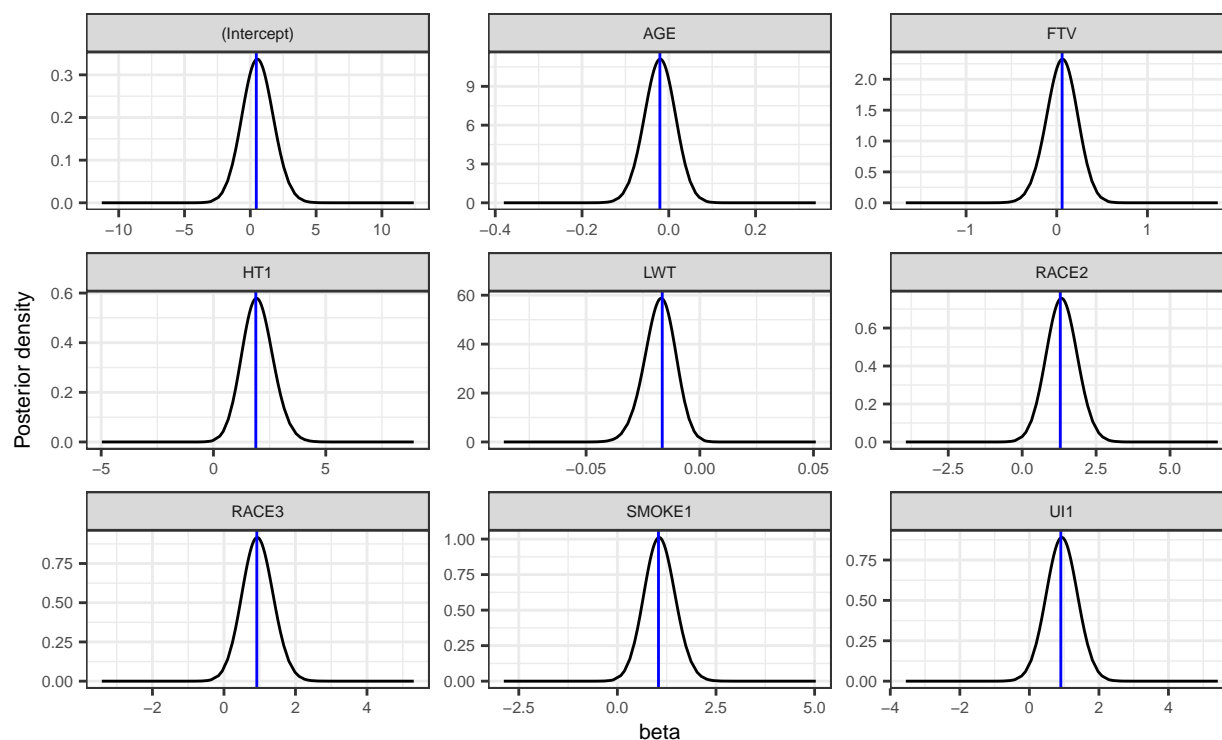
```
temp <- cbind(coef(summary(bwt.glm))[,1:2],bwt.inla$summary.fixed[,1:2])
temp
```

	Estimate	Std. Error	mean	sd
(Intercept)	0.45481245	1.185407237	0.56680173	1.185561299
AGE	-0.02050558	0.035952662	-0.02068758	0.035960525
LWT	-0.01652405	0.006859658	-0.01760162	0.006853204
RACE2	1.28975584	0.527614670	1.34028840	0.527672589
RACE3	0.91906824	0.436301131	0.94553399	0.436260790
SMOKE1	1.04159102	0.395478361	1.07502341	0.395394355
HT1	1.88506202	0.694814773	1.97347903	0.694086373
UI1	0.90414991	0.448602226	0.93278784	0.448574454
FTV	0.05911925	0.171995642	0.05591469	0.171967864

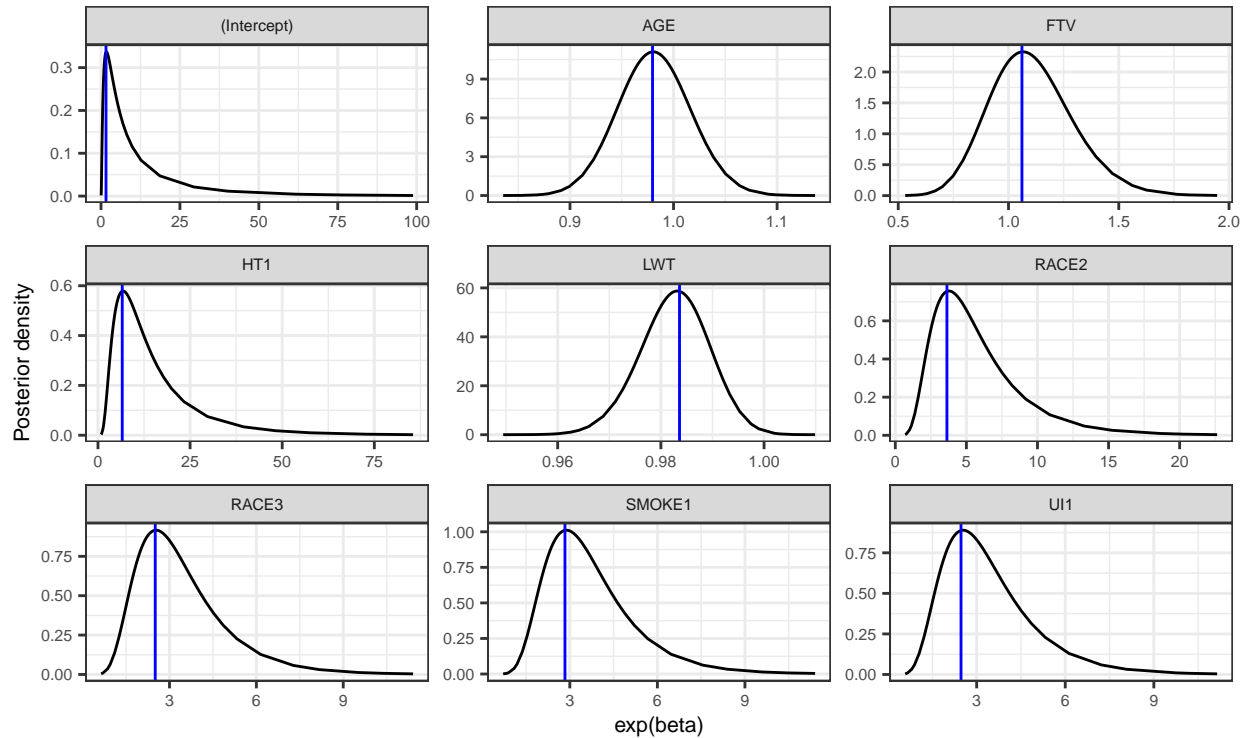
We calculate $\Pr(\beta_j > 0|y)$, $j = 0, \dots, 8$.

```
round(unlist(lapply(bwt.inla$marginals.fixed,
function(x) 1-inla.pmarginal(0, x))), 3)
```

	AGE	LWT	RACE2	RACE3	SMOKE1
(Intercept)	0.680	0.003	0.995	0.986	0.998
HT1		FTV			
	0.999	0.981	0.632		



Method \oplus INLA \oplus glm



Method \perp INLA \perp glm

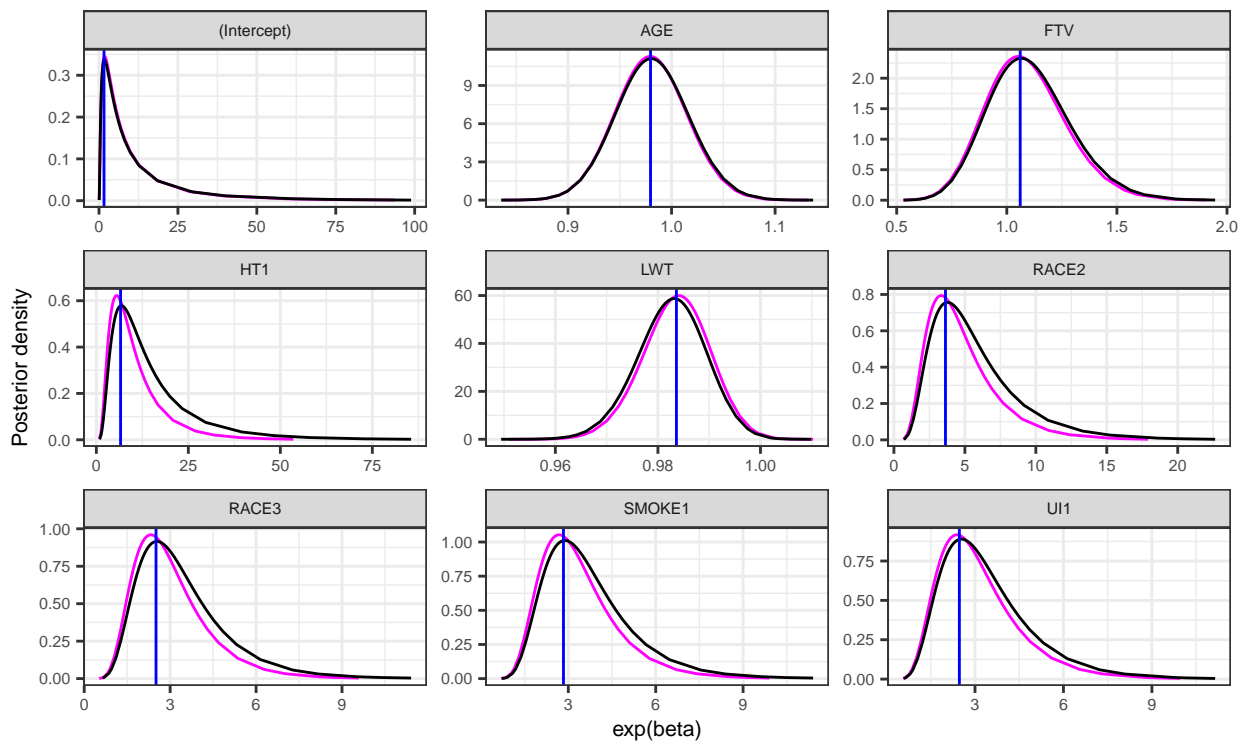
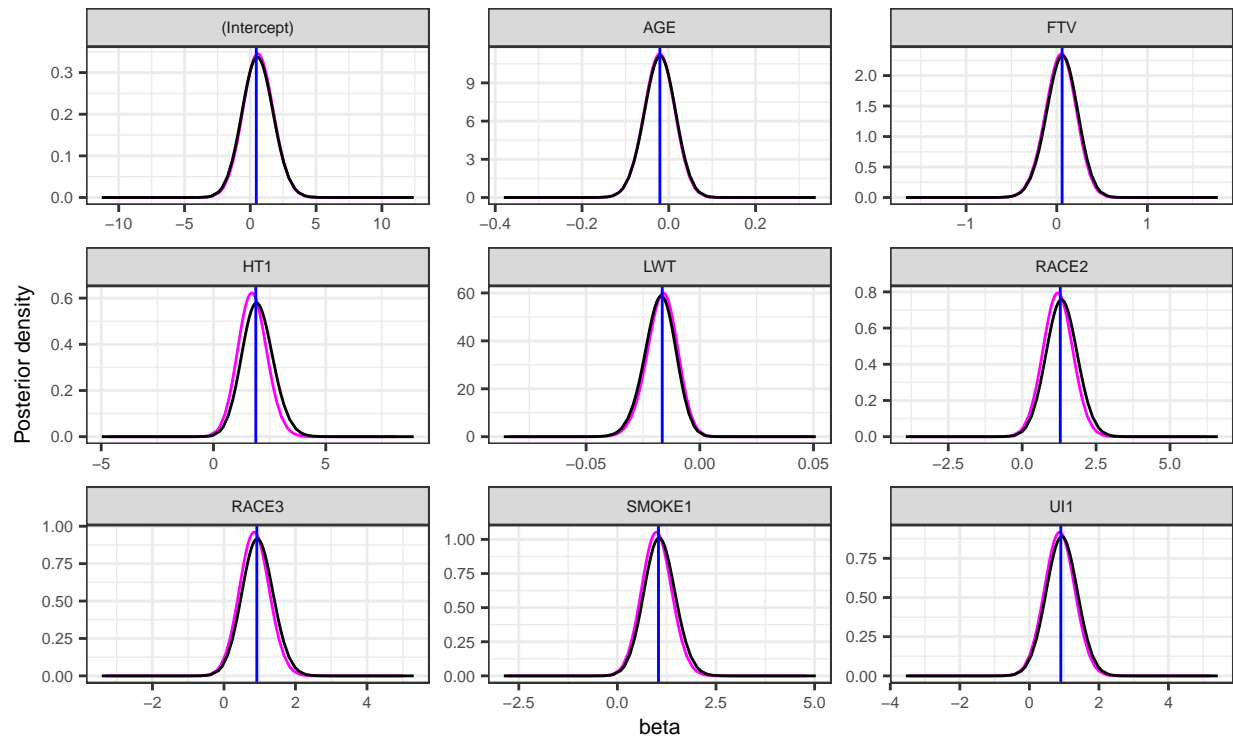
```
alt.prior.fixed <- list(mean.intercept = 0, prec.intercept = 1 / 10^2,
                        mean = 0, prec = 1 / 2^2)
bwt.inla.alt <- inla(LOW ~ AGE + LWT + RACE + SMOKE + HT + UI + FTV,
                    data=lowbwt, family = "binomial", Ntrials = 1,
                    control.compute = list(dic = TRUE, cpo = TRUE),
                    control.fixed = alt.prior.fixed)
round(bwt.inla.alt$summary.fixed, 3)
```

	mean	sd	0.025quant	0.5quant	0.975quant	mode	kld
## (Intercept)	0.607	1.157	-1.632	0.595	2.909	0.573	0
## AGE	-0.022	0.035	-0.092	-0.021	0.047	-0.021	0
## LWT	-0.017	0.007	-0.030	-0.016	-0.004	-0.016	0
## RACE2	1.210	0.503	0.229	1.208	2.201	1.204	0
## RACE3	0.854	0.416	0.047	0.850	1.679	0.844	0
## SMOKE1	0.999	0.379	0.267	0.995	1.756	0.987	0
## HT1	1.740	0.643	0.510	1.729	3.036	1.706	0
## UI1	0.869	0.435	0.013	0.869	1.721	0.870	0
## FTV	0.043	0.170	-0.298	0.046	0.369	0.051	0

We calculate $\Pr(\beta_j > 0|y)$, $j = 0, \dots, 8$.

```
round(unlist(lapply(bwt.inla.alt$marginals.fixed,
                  function(x) 1-inla.pmarginal(0, x))), 3)
```

	AGE	LWT	RACE2	RACE3	SMOKE1
## (Intercept)	0.696	0.270	0.004	0.992	0.981
## HT1	0.998	0.977	0.605		



Prediction

Predictions from a Binomial Distribution

We now consider prediction.

Assume $y|\theta \sim \text{binomial}(N, \theta)$ and $\theta \sim \text{beta}(a, b)$.

We suppose we wish to predict the number of successes Z from M trials.

The predictive distribution is

$$\Pr(z|y) = \binom{M}{z} \frac{\Gamma(N+a+b)}{\Gamma(y+a)\Gamma(N-y+b)} \frac{\Gamma(a+y+z)\Gamma(b+N-y+M-z)}{\Gamma(a+b+N+M)}$$

for $z = 0, \dots, M$.

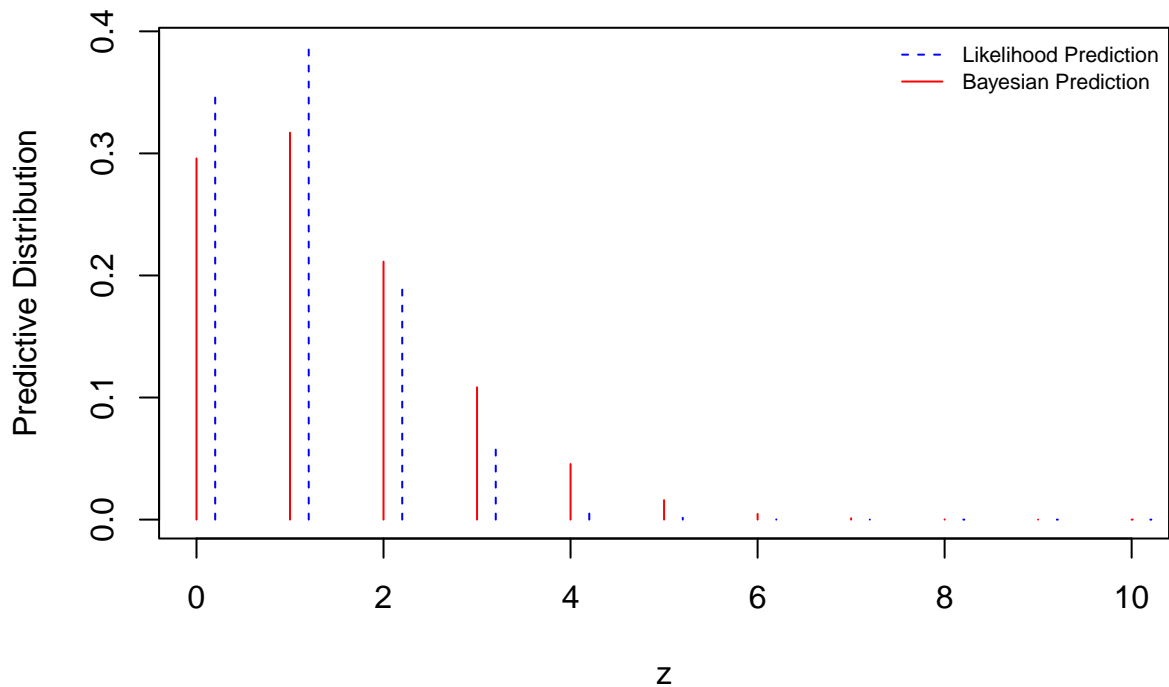
We demonstrate with a uniform prior and observing $y = 2$ successes from $N = 20$ trials, and suppose we wish to predict the number of successes we will see in 10 additional trials.

```
# User written function
binomialpred <- function(a,b,y,N,z,M){
  lchoose(M,z) + lgamma(a+b+N) - lgamma(a+y) - lgamma(b+N-y) +
    lgamma(a+y+z) + lgamma(b+N-y+M-z) - lgamma(a+b+N+M)
}
# Set up the prior and data
a <- b <- 1
y <- 2
N <- 20
M <- 10
```

Along with the Bayesian predictive distribution, we also include a simple approach in which we assume simply take a $\text{binomial}(M, y/N)$ distribution, i.e. assuming the probability is known to be the sample fraction.

```
binpred <- NULL
z <- seq(0,M)
sumcheck <- 0
for (i in 1:(M+1)){
  binpred[i] <- exp(binomialpred(a,b,y,N,z[i],M))
  sumcheck <- sumcheck + binpred[i]
}
likpred <- dbinom(z,M,prob=y/N)
cat("Sum of probs = ",sumcheck,"\n")
## Sum of probs = 1

plot(binpred~z,type="h",col="red",ylim=c(0,max(likpred,binpred)),
     ylab="Predictive Distribution")
points(z+.2,likpred,type="h",col="blue",lty=2)
legend("topright",legend=c("Likelihood Prediction",
    "Bayesian Prediction"),lty=2:1,col=c("blue","red"),bty="n")
```

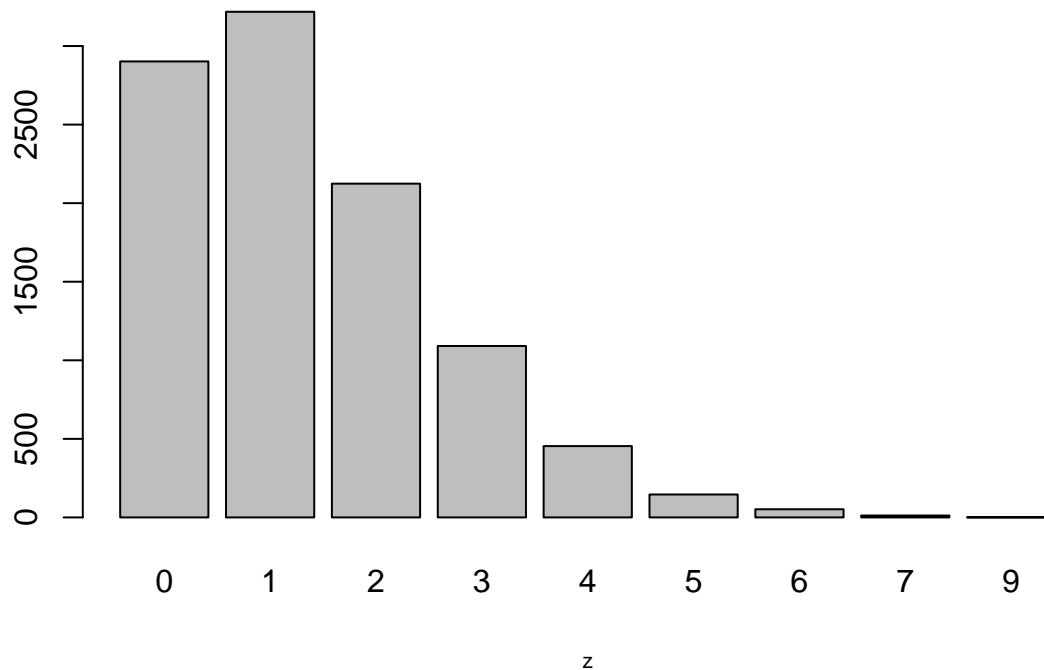


We now simulate directly via:

- Sampling from $\theta^{(s)} \sim p(\theta|y)$, $s = 1, \dots, S$.
- Sampling from $z^{(s)} \sim p(z|\theta)$, $s = 1, \dots, S$.

```
a <- b <- 1; y <- 2; N <- 20; M <- 10
nsim <- 10000
theta <- z <- NULL # This is inefficient but makes method clear
for (s in 1:nsim){
  theta[s] <- rbeta(1,a+y,b+N-y)
  z[s] <- rbinom(1,M,theta[s])
}

barplot(table(z),xlab="z",cex.lab=0.7)
```



Exercises

- Experiment with the priors $\text{Beta}(a, a)$ for the ASE example. In particular, for $a = 2$:
 - Obtain a histogram of the posterior probabilities $\Pr(\theta < 0.5|y)$, across genes.
 - Plot these posterior probabilities versus the versions under $a = 1$, and comment.
 - How sensitive are the (log) Bayes factors to the prior specification?
 - For how many genes would we reject $H_0 : \theta = 0.5$ if we use a rule of $1/\text{BF} > 150$?
- Redo the logistic regression birthweight example, using the default priors, but with the smoking variable only in the model. Compare with a frequentist smoking only example with the `glm` function.