

Module 12: Linear Regression, the g-prior, and model selection

Rebecca C. Steorts

Agenda

- ▶ Review of the Multivariate setup
- ▶ The g-prior
- ▶ Application to diabetes (Hoff, 9.2)
- ▶ Bayesian model selection (g-prior)
- ▶ Bayesian model averaging

Notation

- ▶ $X_{n \times p}$: regression features or covariates (design matrix)
- ▶ $\mathbf{x}_{p \times 1}$: i th row vector of the regression covariates
- ▶ $\mathbf{y}_{n \times 1}$: response variable (vector)
- ▶ $\beta_{p \times 1}$: vector of regression coefficients

Goal: Estimation of $p(y \mid X_{n \times p})$.

Dimensions: $y_i - \beta^T \mathbf{x}_i = (1 \times 1) - (1 \times p)(p \times 1) = (1 \times 1)$.

Multivariate Setup

Let's assume that we have data points (x_i, y_i) available for all $i = 1, \dots, n$.

- ▶ \mathbf{y} is the response variable

$$\mathbf{y} = \begin{pmatrix} y_1 \\ y_2 \\ \vdots \\ y_n \end{pmatrix}_{n \times 1}$$

- ▶ \mathbf{x}_i is the i th row of the design matrix $\mathbf{X}_{n \times p}$.

Consider the regression coefficients

$$\boldsymbol{\beta} = \begin{pmatrix} \beta_1 \\ \beta_2 \\ \vdots \\ \beta_p \end{pmatrix}_{p \times 1}$$

Multivariate Setup

$$y \mid X, \beta, \sigma^2 \sim MVN(X\beta, \sigma^2 I)$$

$$\beta \sim MVN(\beta_0, \Sigma_0)$$

Recall the posterior can be shown to be

$$\beta \mid \mathbf{y}, \mathbf{X} \sim MVN(\beta_n, \Sigma_n)$$

where

$$\beta_n = E[\beta \mid \mathbf{y}, \mathbf{X}, \sigma^2] = (\Sigma_o^{-1} + (X^T X)^{-1} / \sigma^2)^{-1} (\Sigma_o^{-1} \beta_0 + X^T \mathbf{y} / \sigma^2)$$

$$\Sigma_n = \text{Var}[\beta \mid \mathbf{y}, \mathbf{X}, \sigma^2] = (\Sigma_o^{-1} + (X^T X)^{-1} / \sigma^2)^{-1}$$

How do we specify β_0 and Σ_0 ?

The g-prior

To do the *least amount of calculus*, we can put a *g-prior* on β

$$\beta \mid \mathbf{X}, \mathbf{z} \sim \text{MVN}(0, g \sigma^2 (\mathbf{X}^T \mathbf{X})^{-1}).$$

$$\beta_n = E[\beta \mid \mathbf{y}, \mathbf{X}, \sigma^2] = \frac{g}{g+1} (\Sigma_o^{-1} + (X^T X)^{-1} / \sigma^2)^{-1} = \frac{g}{g+1} \hat{\beta}_{ols}$$

$$\Sigma_n = \text{Var}[\beta \mid \mathbf{y}, \mathbf{X}, \sigma^2] = \frac{g}{g+1} (X^T X)^{-1} / \sigma^2 = \frac{g}{g+1} \text{Var}[\hat{\beta}_{ols}]$$

- ▶ g shrinks the coefficients and can prevent overfitting to the data
- ▶ if $g = n$, then as n increases, inference approximates that using $\hat{\beta}_{ols}$

Variance component σ^2

What about a prior on $1/\sigma^2 = \lambda$

$$y \mid X, \beta, \sigma^2 \sim MVN(X\beta, \sigma^2 I) \quad (1)$$

$$\lambda \sim \text{Gamma}(\nu_0/2, \nu_0\lambda^{-1}/2) \quad (2)$$

Then the posterior can be shown to be

$$p(\lambda \mid y, X) \sim \text{Gamma}([\nu_0 + n]/2, [\nu_0\lambda^{-1} + SSR_g]/2)$$

where SSR_g is somewhat complicated (see Hoff for details, p. 158).

Variance component σ^2

The joint distribution can be written as

$$p(\lambda^{-1}, \beta \mid y, X) = p(\lambda^{-1} \mid y, X) \times p(\beta \mid y, X, \lambda^{-1})$$

Goal: simulate $(\lambda^{-1}, \beta) \sim p(\lambda^{-1}, \beta \mid y, X)$

Starting value (β_0, λ_0)

1. Simulate

$$\lambda^{-1} \sim p(\lambda^{-1} \mid y, X)$$

Gives us $(\lambda_1^{-1}, \beta_0)$

2. Use this updated value of λ_1^{-1} to simulate

$$\beta \sim p(\beta \mid y, X, \lambda^{-1})$$

Gives us $(\lambda_1^{-1}, \beta_1)$

Run the sampler for S iterations.

Application to diabetes (Exercise 9.2, part a)

As described in Exercise 7.6, suppose we have data on health-related variables of a population of 532 women.

In this exercise we will be modeling the conditional distribution of glucose level (glu) as a linear combination of the other variables, excluding the variable diabetes.

Model specification

$$y \mid X, \beta, \sigma^2 \sim MVN(X\beta, \lambda^{-1}I) \quad (3)$$

$$\beta \mid \lambda \sim MVN(0, g(X^T X)^{-1}) \quad (4)$$

$$\lambda^{-1} \sim \text{Gamma}(\nu_0/2, \nu_0 \lambda^{-1}/2) \quad (5)$$

We will use the g-prior, where

1. $g = n$
2. $\nu_0 = 1$
3. $\sigma_0^2 = \lambda^{-1} = \hat{\sigma}_{ols}^2 = 8.54$

Regression model on the g-prior

Fit a regression model using the g-prior with $g = n$, $\nu_0 = 2$ and $\sigma_0^2 = 1$. Obtain posterior confidence intervals for all of the parameters.

Regression model on the g-prior

Section 9.2.2 (Hoff) shows that under the g prior, $p(\sigma^2 \mid \mathbf{y}, \mathbf{X})$ and $p(\boldsymbol{\beta} \mid \mathbf{y}, \mathbf{X}, \sigma^2)$ are inverse gamma and multivariate normal distributions respectively.

Regression model on the g-prior

Therefore samples from the joint posterior $p(\sigma^2, \beta \mid \mathbf{y}, \mathbf{X}, \sigma^2)$ can be made with a Monte Carlo approximation.

We first center and scale all the variables so that there is no need to include an intercept in the model.

Regression model on the g-prior

```
library(knitr)
rm(list=ls())
azd_data = read.table("azdiabetes.dat", header = TRUE)
head(azd_data)
```

##	npreg	glu	bp	skin	bmi	ped	age	diabetes
## 1	5	86	68	28	30.2	0.364	24	No
## 2	7	195	70	33	25.1	0.163	55	Yes
## 3	5	77	82	41	35.8	0.156	35	No
## 4	0	165	76	43	47.9	0.259	26	No
## 5	0	107	60	25	26.4	0.133	23	No
## 6	5	97	76	27	35.6	0.378	52	Yes

Regression model on the g-prior

```
y = azd_data$glu  
# remove glu and diabetes  
X = as.matrix(azd_data[,c(-2,-8)])  
head(X)
```

```
##      npreg bp skin  bmi   ped age  
## [1,]      5 68   28 30.2 0.364  24  
## [2,]      7 70   33 25.1 0.163  55  
## [3,]      5 82   41 35.8 0.156  35  
## [4,]      0 76   43 47.9 0.259  26  
## [5,]      0 60   25 26.4 0.133  23  
## [6,]      5 76   27 35.6 0.378  52
```

Standardization

```
# standardize data to have mean 0 and variance 1  
ys = scale(y)  
Xs = scale(X)  
n = dim(Xs)[1]  
p = dim(Xs)[2]
```


Hyper-parameters

```
g = n  
nu0 = 2  
s20 = 1
```

Intermediate Matrices

```
# intermediate matrices
```

```
Hg = (g/(g+1)) * Xs %*% solve(t(Xs) %*% Xs) %*% t(Xs)
```

```
SSRg = t(ys) %*% ( diag(1,nrow=n) - Hg ) %*% ys
```

Monte carlo

```
# number of posterior samples
S = 1000

# generate posteriors
s2 = 1/rgamma(S, (nu0+n)/2, (nu0*s20 + SSRg)/2)
Vb = g * solve(t(Xs) %*% Xs)/(g+1)
Eb = Vb %*% t(Xs) %*% ys
E = matrix(rnorm(S*p, 0, sqrt(s2)),S,p)
beta_s = t( t(E %*% chol(Vb)) + c(Eb))

# transform coefficients to the original scale
sd_X = apply(X,2,sd)
Beta_a = sweep(beta_s,2,sd_X,FUN = "/")
```

The 95% posterior confidence intervals

```
# 95% credible interval  
Beta_CIa = apply(Beta_a, 2, quantile, c(0.025, 0.975))  
kable(data.frame(Beta_CIa))
```

	npreg	bp	skin	bmi	ped	age
2.5%	-0.0515254	-0.0005462	-0.0033335	0.0059186	0.1047847	0.014419
97.5%	0.0091583	0.0133459	0.0159199	0.0352423	0.5617721	0.035035

Model selection

- ▶ Often we have a large number of covariates.
- ▶ Using all of them induces poor statistical performance.
- ▶ How can we reduce the covariates and have good inference and prediction?
- ▶ Common method: Backwards and stepwise regression (slow).

Model selection

Suppose that we believe some of the regression coefficients are 0.

Come up with a prior distribution that reflects the probability of this occurring.

Consider

$$y_i = z_1 b_1 x_{i,1} + \dots z_p b_p x_{i,p},$$

where b_p is a real number and z_j indicate which regression coefficients are nonzero.

Note: $\beta_j = b_j \times z_j$.

Bayesian model selection

Bayesian model selection works by obtaining a posterior distribution for \mathbf{z} .

Assume a prior $p(\mathbf{z})$.

Then

$$p(\mathbf{z} \mid \mathbf{Y}, \mathbf{X}) = \frac{p(\mathbf{z})p(\mathbf{Y} \mid \mathbf{X}, \mathbf{z})}{\sum_{\mathbf{z}} p(\mathbf{z})p(\mathbf{Y} \mid \mathbf{X}, \mathbf{z})}$$

Bayesian model selection

Suppose we want to compare two models z_a and z_b . Consider

$$\text{odds}(z_a, z_b \mid \mathbf{Y}, \mathbf{X}) = \frac{p(z_a \mid \mathbf{Y}, \mathbf{X})}{p(z_b \mid \mathbf{Y}, \mathbf{X})} = \frac{p(z_a)}{p(z_b)} \times \frac{p(\mathbf{Y} \mid \mathbf{X}, z_a)}{p(\mathbf{Y} \mid \mathbf{X}, z_b)}$$

This is posterior odds = prior odds \times “Bayes factor”

“Bayes factor”: how much the data favor model z_a over model z_b

To obtain a posterior distribution over models, we must compute $p(\mathbf{Y} \mid \mathbf{X}, z)$ for *each* model under consideration.

Bayesian model selection

We must compute

$$p(\mathbf{Y} \mid \mathbf{X}, \mathbf{z}) = \int \int p(\mathbf{Y}, \beta, \sigma^2, \mid \mathbf{X}, \mathbf{z}) \quad (6)$$

$$\int \int p(\mathbf{Y} \mid \mathbf{X}, \mathbf{z}) p(\beta \mid \mathbf{X}, \mathbf{z}) p(\sigma^2). \quad (7)$$

To do the *least amount of calculus*, we can put a *g-prior* on β

$$\beta \mid \mathbf{X}, \mathbf{z} \sim \text{MVN}(0, g \sigma^2 (\mathbf{X}^T \mathbf{X})^{-1}).$$

Back to the g-prior

Given the g-prior

$$\beta \mid \mathbf{X}, \mathbf{z} \sim \text{MVN}(0, g \sigma^2 (\mathbf{X}^T \mathbf{X})^{-1}),$$

$p(\mathbf{Y} \mid \mathbf{X}, \mathbf{z})$ can be worked out in closed form (details p. 165).

Go through the details on your own.

Back to the g-prior

This results in being able to compute

$$\frac{p(\mathbf{Y} \mid \mathbf{X}, \mathbf{z}_a)}{p(\mathbf{Y} \mid \mathbf{X}, \mathbf{z}_b)} = (1 + n)^{(p_{z_b} - p_{z_a})/2} \times \left(\frac{s_{z_a}^2}{s_{z_b}^2} \right)^{1/2} \quad (8)$$

$$\times \left(\frac{s_{z_b}^2 + SSR_g^{z_b}}{s_{z_b}^2 + SSR_g^{z_a}} \right)^{(n+1)/2} \quad (9)$$

We have a ratio of the marginal probabilities, giving us a balance between model complexity and model fit.

Suppose p_{z_b} is large compared to p_{z_a} .

This causes a penalization of model z_b

Note that a large value of $SSR_g^{z_b}$ compared to $SSR_g^{z_a}$ will penalize model z_a .

Bayesian Model Averaging

Suppose that we have an estimate of β from which we can make predictions.

We may also want a list of relatively high probability models. We can use a Markov chain to search through the space of models for values of z with high posterior probability.

Bayesian model averaging

Suppose p is large. Then 2^p models to consider.

Instead let's use a Gibbs sampler to search through the space of models for values where \mathbf{z} has a high posterior probability.

Generate a new value of \mathbf{z} via

$$p(z_j \mid \mathbf{Y}, \mathbf{X}, \mathbf{z}_{-j}).$$

The full conditional that $z_j = 1$ can be written as $o_j / (o_j + 1)$.

$$o_j = \frac{p(z_j = 1 \mid \mathbf{Y}, \mathbf{X}, \mathbf{z}_{-j})}{p(z_j = 0 \mid \mathbf{Y}, \mathbf{X}, \mathbf{z}_{-j})} \quad (10)$$

$$= \frac{p(z_j = 1)p(\mathbf{Y} \mid \mathbf{X}, \mathbf{z}_{-j}, z_j = 1)}{p(z_j = 0)p(\mathbf{Y} \mid \mathbf{X}, \mathbf{z}_{-j}, z_j = 0)} \quad (11)$$

Bayesian model averaging

Note: we may also want to obtain posterior samples of β and σ^2 .

Using the conditional distributions from Section 9.2, we can sample from these directly.

The Gibbs sampling scheme requires using Section 9.2 and 9.3 (covered in lab).

Bayesian model averaging

$$\begin{array}{ccccc} \mathbf{z}^{(s)} & \longrightarrow & \sigma^{2(s)} & \longrightarrow & \boldsymbol{\beta}^{(s)} \\ \downarrow & & & & \\ \mathbf{z}^{(s+1)} & \longrightarrow & \sigma^{2(s+1)} & \longrightarrow & \boldsymbol{\beta}^{(s+1)} \end{array}$$

Figure 1: Start with $\mathbf{z}^{(s)}$. Then in random order update \mathbf{z}_j from its full conditional.

Bayesian model averaging

Generate

$$\{\mathbf{z}^{(s+1)}, \sigma^{2(s+1)}, \boldsymbol{\beta}^{(s+1)}\} :$$

1. Set $\mathbf{z} = \mathbf{z}^{(s)}$
2. For $j \in \{1, \dots, p\}$ in random order, replace z_j with a sample from

$$p(z_j \mid \mathbf{z}_{-j}, \mathbf{Y}, \mathbf{X})$$

3. Set $\mathbf{z}^{(s+1)} = \mathbf{z}$
4. Sample $\sigma^{2(s+1)} \sim p(\sigma^2 \mid \mathbf{z}^{(s+1)}, \mathbf{Y}, \mathbf{X})$
5. Sample $\boldsymbol{\beta}^{(s+1)} \sim p(\boldsymbol{\beta} \mid \mathbf{z}^{(s+1)}, \sigma^{2(s+1)}, \mathbf{Y}, \mathbf{X})$

Back to diabetes data (Exercise 9.2, b)

Let's perform Bayesian model averaging (as described in Section 9.3)

Obtain $P(\beta_j \neq 0 | y)$ as well as posterior confidence intervals for all of the parameters. Compare our results to that in part (a.)

Back to diabetes data (Exercise 9.2, b)

The following function `lpy.X` calculates the log of $p(\mathbf{y} \mid \mathbf{X})$, which we will use in implementing the Gibbs sampler for Bayesian model averaging.

```
## a function to compute the marginal probability
lpy.X <- function(y, X, g=length(y), nu0=1, s20=try(summary(lm(y~ -1+X)
  n = dim(X)[1]
  p = dim(X)[2]
  if (p==0) { Hg = 0; s20 = mean(y^2)}
  if (p>0){ Hg = (g/(g+1)) * X %>% solve(t(X) %>% X) %>% t(X) }
  SSRg = t(y) %>% ( diag(1, nrow=n) - Hg ) %>% y -
    .5*( n*log(pi) + p*log(1+g) + (nu0+n)*log(nu0*s20 + SSRg) - nu0*log
    lgamma( (nu0+n)/2 ) - lgamma(nu0/2)
}
```

Back to diabetes data (Exercise 9.2, b)

Let \mathbf{z} be the random binary vector of variable indicators. Generating samples of $p(\mathbf{z}, \sigma^2, \beta)$ from the joint posterior distribution is achieved with the following steps:

1. For $j \in \{1, \dots, p\}$ in random order, draw z_j from $p(z_j \mid \mathbf{z}_{-j}, \mathbf{y}, \mathbf{X})$.
2. Sample $\sigma^2 \sim p(\sigma^2 \mid \mathbf{z}, \mathbf{y}, \mathbf{X})$.
3. Sample $\beta \sim p(\beta \mid \mathbf{z}, \sigma^2, \mathbf{y}, \mathbf{X})$.

Gibbs sampler

```
## MCMC setup
g = n
nu0 = 1 # unit information prior
z = rep(1, p)
lpy.c = lpy.X(ys, Xs[,z==1,drop=FALSE])
S = 10
Z = matrix(NA, S, p)
# Sigma2 = numeric(S)
B = matrix(0, S, p)

## Gibbs sampler
for(s in 1:S){
  # if(s %% 100 ==0) {print(s)}
  # sample z
  for (j in sample(1:p)){
    zp = z
    zp[j] = 1 - zp[j]
    lpy.p = lpy.X(ys,Xs[, zp==1, drop=FALSE])
    r = (lpy.p - lpy.c) * (-1)^(zp[j]==0)
    zp[j] = rbinom(1, 1, 1/(1+exp(-r)))
    if(z[j] == zp[j]) {lpy.c = lpy.p}
  }
}
```

Results

The posterior probability $\Pr(\beta_j \neq 0 \mid \mathbf{y})$ is listed below for each predictor. Clearly all predictors are highly relevant to the response.

```
pprob_Z = apply(Z,2,mean)
pprob_Z = data.frame(matrix(pprob_Z,nr=1,nc=p))
names(pprob_Z) = names(azd_data[c(-2,-8)])
row.names(pprob_Z) = 'posterior including probability'

kable(pprob_Z)
```

	npreg	bp	skin	bmi	ped	age
posterior including probability	1	1	1	1	1	1

Results

The 95% posterior confidence intervals for all the parameters from Bayesian model averaging are listed below. The results are similar to those in part (a) because all the predictors are included in each iteration of Gibbs sampler all the time.

```
# transform coefficients to the original scale
#sd_X = apply(X,2,sd)
Beta_b = sweep(B,2,sd_X,FUN = "/")

# 95% credible interval
Beta_CIb = apply(Beta_b, 2, quantile, c(0.025, 0.975))

kable(data.frame(Beta_CIb), col.names = names(azd_data[c(-2,-8)]))
```

	npreg	bp	skin	bmi	ped	ag
2.5%	-0.0377503	0.0004040	-0.0004484	0.0111701	0.1530748	0.016852
97.5%	0.0046465	0.0116124	0.0150486	0.0277676	0.5196038	0.028107