

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

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# Agenda

# 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  $p(y \mid x)$ .

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  $i = 1, \dots, n$ .

- ▶  $y$  is the response variable

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

- ▶  $x_i$  is the  $i$ th row of the design matrix  $X_{n \times p}$ .

Consider the regression coefficients

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

## 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

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

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

How do we specify  $\beta_0$  and  $\Sigma_0$ ?

## The g-prior

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

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

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

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

- ▶  $g$  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 $\sigma^2$

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 $\sigma^2$

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  $(\beta_0, \lambda_0)$

1. Simulate

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

Gives us  $(\lambda_1^{-1}, \beta_0)$  2. Use this updated value of  $\lambda_1^{-1}$  to simulate

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

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

Run the sampler for  $S$  iterations.



## Back to the oxygen uptake example

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

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

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

We will use the g-prior, where

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

## 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 ( $\text{glu}$ ) as a linear combination of the other variables, excluding the variable diabetes.

## 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  
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

## 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
2.5%	-0.0515254	-0.0005462	-0.0033335	0.0059186	0.1047847
97.5%	0.0091583	0.0133459	0.0159199	0.0352423	0.5617721

# 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  $\beta$

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

## Back to the g-prior

Given the g-prior

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

$p(\mathbf{Y} \mid \mathbf{X}, \mathbf{z})$  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 are content with a estimate of  $\beta$  from which we can make predictions.

We may also want a list of relatively high probablitiy 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  $\beta$  and  $\sigma^2$ .

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  $z_j$  from its full conditional.



# Bayesian model averaging

Generate

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

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

## 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
  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
  SSRg = t(y) %*% ( diag(1, nrow=n) - Hg ) %*% y -
    .5*( n*log(pi) + p*log(1+g) + (nu0+n)*log(nu0*s20 + SSR
    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])
```

## 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
```

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