

# Mixed effect probit regression

## Genotypic fungal resistance

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April 28, 2016

# Outline

- Probit regression
- Bayesian probit regression
  - Data augmentation
- Bayesian mixed effect probit regression
- Extensions
  - Ordinal categorical data
  - Nominal categorical data
  - Bayesian logistic regression

# Probit regression

Consider the model

$$Y_i \stackrel{ind}{\sim} \text{Ber}(\theta_i)$$

where, for the  $i$ th observation,

- $Y_i$  is binary indicating *success* and
- $\theta_i$  is the probability of success.

A probit regression model assumes

$$\theta_i = \Phi(X_i^\top \beta)$$

where

- $X_i$  are the explanatory variables for the  $i$ th observation,
- $\Phi$  is the standard normal cumulative distribution function, and
- $\beta$  is the vector of parameters to be estimated.

# Low birth weight

low	age	lwt	race	smoke	ptl	ht
Min. :0.0000	Min. :14.00	Min. : 80.0	1:96	Min. :0.0000	Min. :0.0000	Min. :0.00000
1st Qu.:0.0000	1st Qu.:19.00	1st Qu.:110.0	2:26	1st Qu.:0.0000	1st Qu.:0.0000	1st Qu.:0.00000
Median :0.0000	Median :23.00	Median :121.0	3:67	Median :0.0000	Median :0.0000	Median :0.00000
Mean :0.3122	Mean :23.24	Mean :129.8		Mean :0.3915	Mean :0.1958	Mean :0.06349
3rd Qu.:1.0000	3rd Qu.:26.00	3rd Qu.:140.0		3rd Qu.:1.0000	3rd Qu.:0.0000	3rd Qu.:0.00000
Max. :1.0000	Max. :45.00	Max. :250.0		Max. :1.0000	Max. :3.0000	Max. :1.00000
ui	ftv	bwt				
Min. :0.0000	Min. :0.0000	Min. : 709				
1st Qu.:0.0000	1st Qu.:0.0000	1st Qu.:2414				
Median :0.0000	Median :0.0000	Median :2977				
Mean :0.1481	Mean :0.7937	Mean :2945				
3rd Qu.:0.0000	3rd Qu.:1.0000	3rd Qu.:3487				
Max. :1.0000	Max. :6.0000	Max. :4990				

```
m = glm(low~., family=binomial(link=probit), data=birthwt[, -10]); summary(m)
```

Call:

```
glm(formula = low ~ ., family = binomial(link = probit), data = birthwt[,  
-10])
```

Deviance Residuals:

Min	1Q	Median	3Q	Max
-1.8848	-0.8271	-0.5217	0.9903	2.2445

Coefficients:

	Estimate	Std. Error	z value	Pr(> z )
(Intercept)	-1.31431	0.24893	-5.280	1.29e-07 ***
age	-0.09774	0.11482	-0.851	0.39466
lwt	-0.27281	0.12217	-2.233	0.02555 *
race2	0.74961	0.31431	2.385	0.01708 *
race3	0.52183	0.25557	2.042	0.04117 *
smoke	0.56910	0.23469	2.425	0.01531 *
ptl	0.31968	0.20835	1.534	0.12495
ht	1.11161	0.41664	2.668	0.00763 **
ui	0.46517	0.27930	1.665	0.09581 .
ftv	0.02832	0.10161	0.279	0.78050

---  
Signif. codes: 0 '\*\*\*' 0.001 '\*\*' 0.01 '\*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for binomial family taken to be 1)

Null deviance: 234.67 on 188 degrees of freedom  
Residual deviance: 201.03 on 179 degrees of freedom  
AIC: 221.03

Number of Fisher Scoring iterations: 5

# Bayesian probit regression

Consider the model

$$\begin{aligned} Y_i &\overset{ind}{\sim} \text{Ber}(\theta_i) \\ \theta_i &= \Phi(X_i^\top \beta) \end{aligned}$$

with prior

$$\beta \sim N(b, B)$$

The posterior distribution is

$$\begin{aligned} p(\beta|y) &\propto p(y|\beta)p(\beta) \\ &\propto \left[ \prod_{i=1}^n \Phi(X_i' \beta)^{y_i} [1 - \Phi(X_i' \beta)]^{1-y_i} \right] e^{-(\beta-b)^\top B^{-1}(\beta-b)/2} \end{aligned}$$

But neither  $p(\beta|y)$  nor  $p(\beta_p|y, \beta_{-p})$  are a known distribution.

# Data augmentation

An alternative construction of the model is

$$\begin{aligned} Y_i &= I(\zeta_i > 0) \\ \zeta_i &\stackrel{\text{ind}}{\sim} N(X_i' \beta, 1) \end{aligned}$$

Note that

$$\begin{aligned} \theta_i &= P(Y_i = 1) \\ &= P(\zeta_i > 0) \\ &= P(X_i' \beta + \epsilon > 0) \quad \epsilon \sim N(0, 1) \\ &= P(\epsilon > -X_i' \beta) \\ &= P(\epsilon < X_i' \beta) \quad \text{symmetry of standard normal} \\ &= \Phi(X_i' \beta) \end{aligned}$$

Thus, this is equivalent to the probit regression model.

## Posterior distribution

Now, the likelihood is

$$p(y|\zeta) \propto \prod_{i=1}^n [\mathbf{I}(\zeta_i > 0)\mathbf{I}(y_i = 1) + \mathbf{I}(\zeta_i \leq 0)\mathbf{I}(y_i = 0)]$$

and

$$\zeta_i \stackrel{\text{ind}}{\sim} N(X_i'\beta, 1) \quad \beta \sim N(b, B)$$

Therefore the *complete data likelihood* is

$$p(y, \zeta|\beta) \propto \prod_{i=1}^n N(\zeta_i|X_i'\beta, 1) [\mathbf{I}(\zeta_i > 0)\mathbf{I}(y_i = 1) + \mathbf{I}(\zeta_i \leq 0)\mathbf{I}(y_i = 0)]$$

Thus the posterior distribution is

$$p(\beta, \zeta|y) \propto p(y|\zeta, \beta)p(\zeta, \beta) = p(y|\zeta)p(\zeta|\beta)p(\beta) = p(y, \zeta|\beta)p(\beta)$$

and we will derive the full conditionals for  $p(\beta|\zeta, y)$  and  $p(\zeta|\beta, y)$ .



# Full conditional for $\beta$

The full conditional for  $\beta$  is

$$\begin{aligned} p(\beta | \dots) &\propto p(y|\zeta)p(\zeta|\beta)p(\beta) \\ &\propto p(\zeta|\beta)p(\beta) \\ &= [\prod_{i=1}^n N(\zeta_i|X_i'\beta, 1)] N(\beta|b, B) \\ &= N(\zeta|X\beta, I)N(\beta|b, B) \end{aligned}$$

and thus  $\beta | \dots \sim N(\hat{\beta}, \hat{\Sigma}_\beta)$  with

$$\begin{aligned} \hat{\Sigma}_\beta &= [B^{-1} + X^\top X]^{-1} \\ \hat{\beta} &= \hat{\Sigma}_\beta [B^{-1}b + X^\top \zeta] \end{aligned}$$

## Full conditional for $\zeta$

The full conditional for  $\zeta$  is

$$\begin{aligned} p(\zeta | \dots) &\propto p(y|\zeta)p(\zeta|\beta)p(\beta) \\ &\propto p(y|\zeta)p(\zeta|\beta) \\ &= \prod_{i=1}^n N(\zeta_i | X_i' \beta, 1) [I(\zeta_i > 0)I(y_i = 1) + I(\zeta_i \leq 0)I(y_i = 0)] \end{aligned}$$

Thus the  $\zeta_i$  are conditionally independent with distribution

$$p(\zeta_i | y_i, \beta) = \begin{cases} N(\zeta_i | X_i' \beta, 1) I(\zeta_i > 0) & \text{if } y_i = 1 \\ N(\zeta_i | X_i' \beta, 1) I(\zeta_i \leq 0) & \text{if } y_i = 0 \end{cases}$$

These can be drawn using the modified inverse cdf method.

```

mcmc = function(n_iter, y, X, beta0, Sigma_beta) {
  n = nrow(X)
  p = ncol(X)

  # Precalculate quantities
  y = (as.numeric(y)==1)
  n1 = sum( y)
  n0 = sum(!y)
  XX = t(X)%*%X
  Si = solve(Sigma_beta)
  Sib = Si%*%beta0

  # Saving structures
  beta_keep      = matrix(NA, n_iter, p)
  zeta_keep      = matrix(NA, n_iter, n)

  # Initial values
  m = glm(y~X-1, family=binomial(probit))
  beta = coef(m)
  zeta = rep(NA,n)

  for (i in 1:n_iter) {
    # Sample zeta
    Xb = X%*%beta
    cut = pnorm(0,Xb)
    zeta[ y] = qnorm(runif(n1, cut[ y], 1), Xb[ y], 1)
    zeta[!y] = qnorm(runif(n0, 0, cut[!y]), Xb[!y], 1)

    # Sample beta
    S_hat = solve(Si+XX)
    b_hat = S_hat %*% (Sib+t(X)%*%zeta)
    beta = mvrnorm(1, b_hat, S_hat)

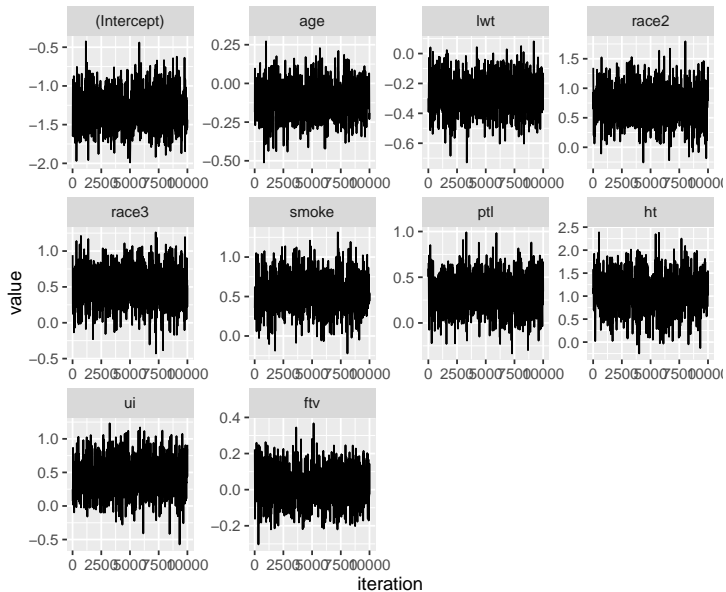
    # Record values
    beta_keep[i,] = beta
    zeta_keep[i,] = zeta
  }
}

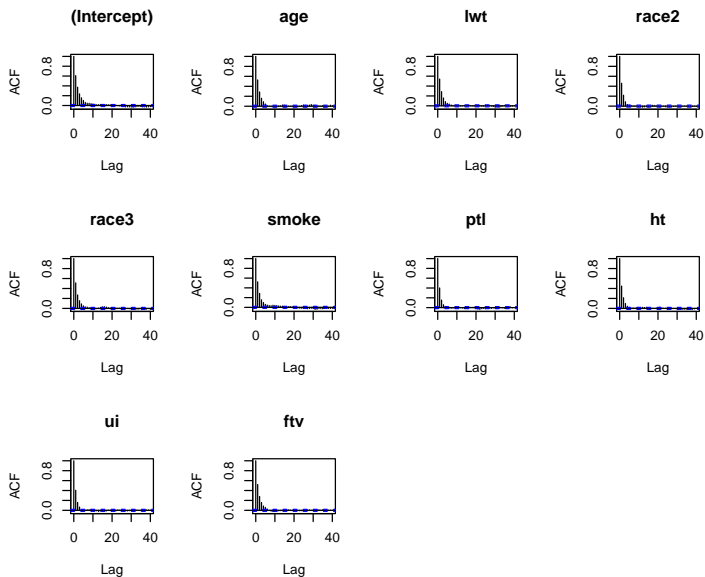
```

# Run the MCMC

```
X = model.matrix(m) # Constructs the design matrix
p = ncol(X)
n_iter = 10000
system.time(out <- mcmc(n_iter, birthwt$low, X, rep(0,p), 3*diag(p)))
```

```
      user  system elapsed
2.763    0.009    2.775
```





# Credible intervals

Source: local data frame [10 x 4]

	variable	ess	lb	ub
	(fctr)	(dbl)	(dbl)	(dbl)
1	(Intercept)	2958	-1.75	-0.81
2	age	2773	-0.33	0.12
3	lwt	2516	-0.52	-0.05
4	race2	4766	0.11	1.32
5	race3	3389	-0.01	0.98
6	smoke	3069	0.09	1.01
7	ptl	5416	-0.07	0.72
8	ht	3692	0.26	1.87
9	ui	4269	-0.08	0.98
10	ftv	2910	-0.18	0.22

# Probit regression with random effects

Consider the probit regression model

$$\begin{aligned} Y_i &= I(\zeta_i > 0) \\ \zeta &\sim N(\tilde{X}\tilde{\beta}, 1) \end{aligned}$$

where

$$\tilde{X} = [X \quad Zm] \quad \tilde{\beta} = (\beta, \alpha)^\top$$

where  $X$  is the design matrix for fixed effects and  $Zm$  is the design matrix for the random effects. A common assumption is that the random effects are  $\alpha \sim N(0, \sigma^2 I)$ . Thus the distribution on  $\tilde{\beta}$  is

$$\tilde{\beta} = \begin{pmatrix} \beta \\ \alpha \end{pmatrix} \sim N\left(\begin{bmatrix} b \\ 0 \end{bmatrix}, \begin{bmatrix} B & 0 \\ 0 & \sigma^2 I \end{bmatrix}\right)$$

where the precision is

$$\begin{bmatrix} B & 0 \\ 0 & \sigma^2 I \end{bmatrix}^{-1} = \begin{bmatrix} B^{-1} & 0 \\ 0 & \frac{1}{\sigma^2} I \end{bmatrix}$$



# Full posterior

The full posterior is

$$p(\zeta, \beta, \alpha, \sigma^2 | y) \propto p(y | \zeta) p(\zeta | \tilde{\beta}) p(\tilde{\beta} | \sigma^2) p(\sigma^2)$$

We have already derived the full conditionals

- $p(\tilde{\beta} | \dots)$
- $p(\zeta | \dots)$

but we need the full conditional for  $\sigma^2$  to implement a Gibbs sampler.

Full conditional for  $\sigma^2$ 

If we choose  $\sigma \sim \text{Unif}(0, 10)$  and there are  $U$  random effects, then

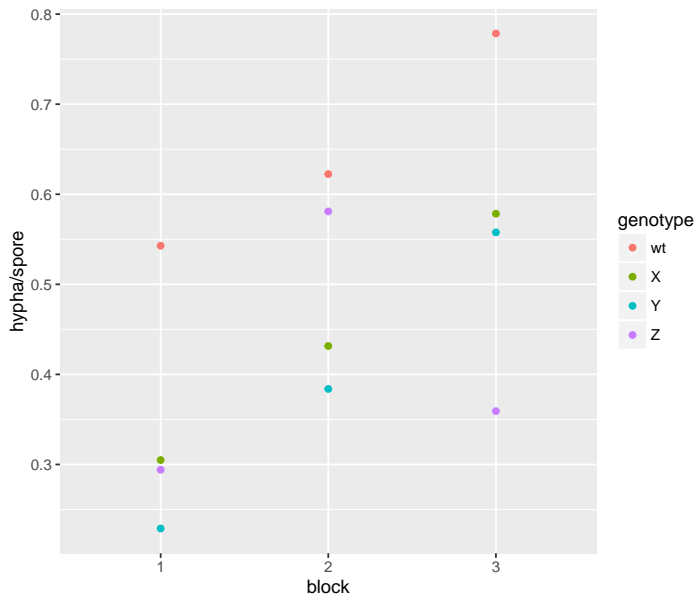
$$\begin{aligned}
 p(\sigma^2 | \dots) &\propto p(y|\zeta)p(\zeta|\tilde{\beta})p(\tilde{\beta}|\sigma^2)p(\sigma^2) \\
 &= p(\tilde{\beta}|\sigma^2)p(\sigma^2) \\
 &\propto p(\alpha|\sigma^2)p(\sigma^2) \\
 &\propto \prod_{i=1}^U N(\alpha_i|0, \sigma^2) \frac{1}{\sigma} \text{I}(0 < \sigma^2 < 100) \\
 &\propto (\sigma^2)^{-U/2} e^{-\frac{1}{2\sigma^2} \alpha' \alpha} (\sigma^2)^{-1/2} \text{I}(0 < \sigma^2 < 100) \\
 &= (\sigma^2)^{-\frac{U-1}{2}-1} e^{-\frac{\alpha' \alpha}{2\sigma^2}} \text{I}(0 < \sigma^2 < 100)
 \end{aligned}$$

Thus  $\sigma^2 \sim \text{IG}([U - 1]/2, \alpha' \alpha/2)$  truncated to be smaller than 100. This can be drawn using the modified inverse cdf method.

# Genotypic resistance to corn fungus

	X	genotype	block	spore	hypha	prop	pot
1	1	X	1	82	25	0.3048780	X1
6	6	X	2	95	41	0.4315789	X2
11	11	X	3	102	59	0.5784314	X3
16	16	Y	1	83	19	0.2289157	Y1
21	21	Y	2	99	38	0.3838384	Y2
26	26	Y	3	104	58	0.5576923	Y3
31	31	Z	1	102	30	0.2941176	Z1
36	36	Z	2	105	61	0.5809524	Z2
41	41	Z	3	103	37	0.3592233	Z3
46	46	wt	1	140	76	0.5428571	wt1
51	51	wt	2	143	89	0.6223776	wt2
56	56	wt	3	158	123	0.7784810	wt3

# Corn fungus data set



```
Generalized linear mixed model fit by maximum likelihood (Laplace Approximation) ['glmerMod']
Family: binomial (probit)
Formula: cbind(hypha, spore ~ hypha) ~ block + genotype + (1 | pot)
Data: d
Control: glmerControl(optimizer = "bobyqa")
```

AIC	BIC	logLik	deviance	df.resid
95.3	98.7	-40.6	81.3	5

Scaled residuals:

Min	1Q	Median	3Q	Max
-1.45760	-0.35765	0.05486	0.36506	1.32376

Random effects:

Groups	Name	Variance	Std.Dev.
pot	(Intercept)	0.01773	0.1331

Number of obs: 12, groups: pot, 12

Fixed effects:

	Estimate	Std. Error	z value	Pr(> z )
(Intercept)	0.05126	0.12429	0.412	0.680040
block2	0.42497	0.13027	3.262	0.001106 **
block3	0.60818	0.13006	4.676	2.92e-06 ***
genotypeX	-0.55654	0.14700	-3.786	0.000153 ***
genotypeY	-0.68630	0.14725	-4.661	3.15e-06 ***
genotypeZ	-0.62691	0.14500	-4.324	1.53e-05 ***

---

Signif. codes: 0 '\*\*\*' 0.001 '\*\*' 0.01 '\*' 0.05 '.' 0.1 ' ' 1

Correlation of Fixed Effects:

	(Intr)	block2	block3	gntypX	gntypY
block2		-0.530			
block3		-0.526	0.522		
genotypeX		-0.520	-0.019	-0.027	
genotypeY		-0.514	-0.027	-0.035	0.454
genotypeZ		-0.525	-0.012	-0.012	0.450

```

mcmc = function(n_iter, y, X, Zm, beta0, Sigma_beta) {
  require(Matrix)
  n = nrow(X)
  p = ncol(X)
  q = ncol(Zm)

  # Initial values
  m = glm(y~0+X, family=binomial(probit))
  beta = c(coef(m), rnorm(q))
  zeta = rep(NA, n)

  # Precalculate quantities
  y = (as.numeric(y)==1)
  n1 = sum( y)
  n0 = sum(!y)
  X = cbind(X, Zm)
  XX = t(X)%*%X
  Si = solve(Sigma_beta)
  Sib = Si%*%beta0
  a = (q-1)/2

  # Saving structures
  beta_keep = matrix(NA, n_iter, p)
  alpha_keep = matrix(NA, n_iter, q)
  sigma_keep = rep(NA, n_iter)

  for (i in 1:n_iter) {
    # Sample zeta
    Xb = X%*%beta
    cut = pnorm(0, as.numeric(Xb))
    zeta[ y] = qnorm(runif(n1, cut[ y], 1), Xb[ y], 1)
    zeta[!y] = qnorm(runif(n0, 0, cut[!y]), Xb[!y], 1)

    # Sample sigma
    alpha = beta[p+1:q]
    b = sum(alpha^2)/2
  }
}

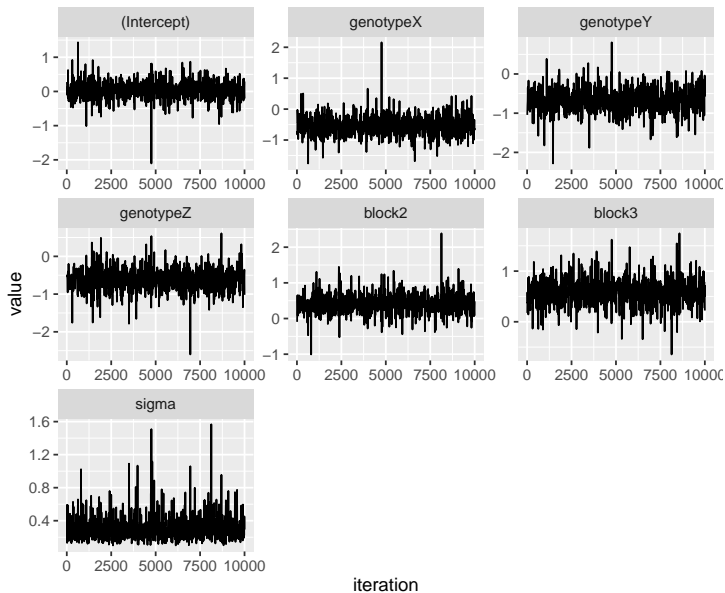
```

```
# Turn into binary data
dd = ddply(d, .(genotype,block,pot), function(x) {
  data.frame(y=c(rep(1,x$hypha),rep(0,x$spore-x$hypha)))
})

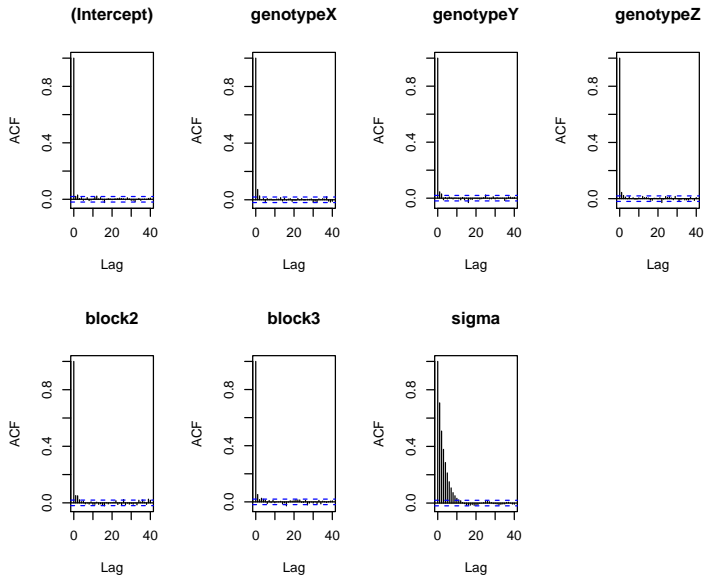
m = glmer(y~genotype+block+(1|pot), family=binomial(probit), dd)

X = model.matrix(m)
Z = as.matrix(getME(m,"Z"))
p = ncol(X)
n_iter = 10000
system.time(out <- mcmc(n_iter, dd$y, X, Z, rep(0,p), 10*diag(p)))

      user  system elapsed
95.382   0.011  95.514
```







# Credible intervals

Source: local data frame [7 x 4]

	variable (fctr)	ess (dbl)	lb (dbl)	ub (dbl)
1	(Intercept)	8747	-0.45	0.55
2	genotypeX	9417	-1.12	0.05
3	genotypeY	10051	-1.25	-0.09
4	genotypeZ	9572	-1.17	-0.03
5	block2	9012	-0.09	0.93
6	block3	8831	0.08	1.09
7	sigma	1688	0.13	0.68

## Contrasts to compare other genotypes

```
t(with(betas, data.frame("X-Y" = quantile(genotypeX-genotypeY, c(.025,.975)),
                        "Y-Z" = quantile(genotypeY-genotypeZ, c(.025,.975)),
                        "X-Z" = quantile(genotypeX-genotypeZ, c(.025,.975)), check.names=FALSE)))
```

	2.5%	97.5%
X-Y	-0.4448574	0.7118965
Y-Z	-0.6486938	0.5173782
X-Z	-0.5104321	0.6529829

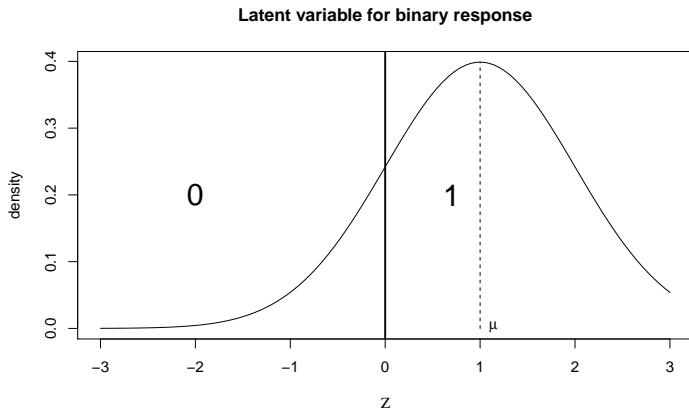
## $t$ priors

Suppose we want  $\beta_j \stackrel{\text{ind}}{\sim} t_{v_j}(b_j, B_j)$ . We can write this prior hierarchically via

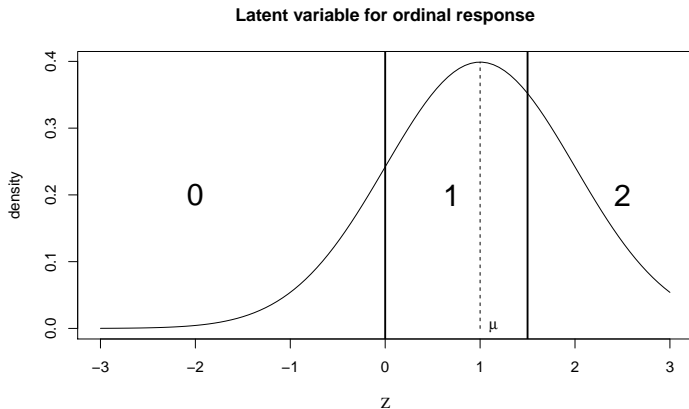
$$\beta_j | \tau_j^2 \stackrel{\text{ind}}{\sim} N(b_j, \tau_j^2), \quad \tau_j^2 \sim \text{Inv} - \chi^2(v_j, B_j).$$

Now the MCMC can proceed exactly as before, but with the additional full conditional for  $(\tau_1^2, \dots, \tau_J^2)$  which will be independent inverse  $\chi^2$  distributions.

# Binary response



# Ordinal response with 3 categories



## Unordered categorical response

Suppose  $Y_i$  is random variable with support  $1, \dots, K$  and

$$Pr(Y_i = k) = \theta_{ik}$$

where  $\theta_{ik}$  may depend on explanatory variables for both  $i$  and  $k$ . For example, an individual is shopping for fruit then perhaps the age of the individual and the price of the fruits will affect the shopper's choice.

We can model this using data augmentation by introducing a latent utility  $\zeta_{ik}$  for each shopper-fruit combination. Then the response is

$$Y_i = \operatorname{argmax}_k \zeta_{ik}$$

and there is great flexibility in how the  $\zeta_{ik}$  are modeled.

# Bayesian logistic regression

$$Y_i = I(\zeta_i > 0)$$

$$\zeta_i \stackrel{\text{ind}}{\sim} \text{Logistic}(X_i' \beta, 1)$$

```
Warning: data was combined!
N: 183, P: 10
Burn-in complete: 0.06 sec. for 500 iterations.
Expect approx. 0.12 sec. for 1000 samples.
Sampling complete: 0.13 sec. for 1000 iterations.
```

	X1	lb	ub
1 (Intercept)	-3.26	-1.43	
2	age	-0.57	0.24
3	lwt	-0.98	-0.14
4	race2	0.27	2.42
5	race3	0.05	1.82
6	smoke	0.24	1.84
7	ptl	-0.13	1.25
8	ht	0.68	3.46
9	ui	-0.19	1.75
10	ftv	-0.32	0.41