Probit regression for ordinal data: Categorical features

Setting:

• For each observations, we have 3 associated variables

X1: Nomial variable with 3 possible outcomes $\{1,2,3\}$ which we will generate from Mult(1, [0.2,0.3,0.5]) X2: Nomial variable with 3 possible outcomes $\{1,2,3\}$ which we will generate from Mult(1, [0.3,0.5,0.2]) Y: Ordinal variable with 5 levels, $\{1,2,3,4,5\}$ generated using the following process

```
\epsilon_i \sim N(0,1) \ z_i = \beta^T X_i + \epsilon \text{ where } X = [X1 == 1, X1 == 2, X2 == 1, X2 == 2] \text{ i.e. we drop category } (1,1) as the baseline g(z_i) = Y_i.
```

Here $\beta = [-3, 2, 2, -4]$ and function g will be the binning function which will bin data into 5 different bins corresponding to 5 possible ordinal outcome.

- We will try to model Y conditioned on other variables (X1, and X2) using probit regression with latent variable Z with rank likelihood on parameter Z.
- We have two unknown parameters β and z_i which will be sampled from the full conditional posterior distribution using blocked gibbs sampling.

Summary on modelling process

```
\epsilon_i \sim N(0,1) z_i = \beta^T X_i + \epsilon z_i \in R(Y) where R(Y) = \{z_i : z_i > z_j \text{ if } Y_i > Y_j \text{ and } z_i < z_j \text{ if } Y_i < Y_j \}
```

```
# Data generating process
set.seed(0)
n = 300
beta = c(-3, 2, 2, -4)
# noise term
epsilon = rnorm(n, mean = 0, sd = 1)
X1 = t(rmultinom(n, size = 1, prob = c(0.2, 0.3, 0.5)))
X2 = t(rmultinom(n, size = 1, prob = c(0.3, 0.5, 0.2)))
# X
X = cbind(X1[,2:3], X2[,2:3])
colnames(X) <- c('X1_cat2', 'X1_cat3', 'X2_cat2', 'X2_cat3')</pre>
# Z
Z = X%*\%beta + epsilon
# Cut-off points and Y
g = quantile(Z, probs = c(0.2, 0.4, 0.6, 0.8))
Y = rep(NA, n)
Y[Z \leq g[1]] = 1
Y[Z = g[1] \& Z < g[2]] = 2
Y[Z = g[2] \& Z < g[3]] = 3
Y[Z = g[3] \& Z < g[4]] = 4
Y[Z = g[4]] = 5
```

Prior specifications:

```
\beta \sim multiN(0, n(X^TX)^{-1})
```

Blocked Gibbs Sampling here consists of two major steps

1. Sample new β from its full conditional

$$\beta \mid z, X, Y, z \in R(Y) \sim multiN(\frac{n}{n+1}(X^TX)^{-1}X^Tz, \frac{n}{n+1}(X^TX)^{-1})$$

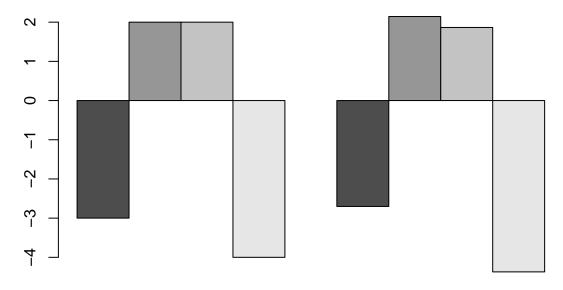
2. for each i, using inverse cdf method, sample new z_i from its full conditional which is a truncated normal distribution:

```
z_i \mid \beta, X, Y, z_i \in R(Y) \sim N(\beta^T x_i, 1) * I\{z_i \in (a, b)\}
where a = \max(z_i \text{ for } Y_i < Y_i) \text{ b} = \min(z_i \text{ for } Y_i > Y_i)
```

Note that in the gibbs sampling process, we force the first threshold (g1) to be at the true value in the data generating process otherwise the parameters will be unidentifiable (we can definitely fine infinite combinations of weights parameter to order the outcome according to their target Y: scaling, shifting, etc.)

```
# Blocked Gibbs Sampling
set.seed(1)
# Prior Parameter
g1 = g[1] #Fix g1 the first cutoff
Z_i = g1 + Y - 1.5
variance_beta = (n/(n+1))*solve(t(X)%*%X)
mean_beta_hat = (n/(n+1))*solve(t(X)%*%X)%*%t(X)
# Initialize the sampling matrix
S = 30000
SAMPLED_Z = matrix(nrow=S,ncol=n)
BETA = matrix(nrow=S,ncol=4)
for (round in 1:S) {
  # Step 1: Sample Beta
  mean_beta = mean_beta_hat%*%Z_i
  beta_sampled = dae::rmvnorm(mean = mean_beta,
                               V = variance beta, method = 'choleski')
  BETA[round,] <- beta_sampled</pre>
  # Step 2: Sample Z using inverse cdf appraoch
  for (i in 1:n) {
    # Get the lower and upper bound (a, b) of truncated normal
    a = max(-Inf, Z_i[Y<Y[i]], na.rm = TRUE)</pre>
    b = min(Z_i[Y>Y[i]], Inf, na.rm = TRUE)
    # Force the lowest cutoff to be at q1
    if (Y[i] == 1) {
      b = g1
    else if (Y[i] == 2) {
      a = g1
    }
    # Sample using inverse cdf
    ez = t(beta_sampled)%*%X[i,]
    u = runif(1, pnorm(a - ez), pnorm(b-ez))
```

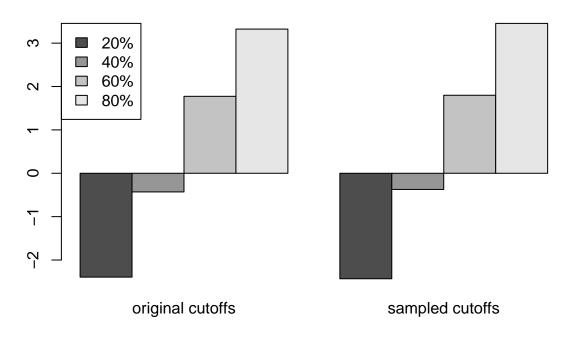
Blocked Gibbs Sampling Assessment: Beta



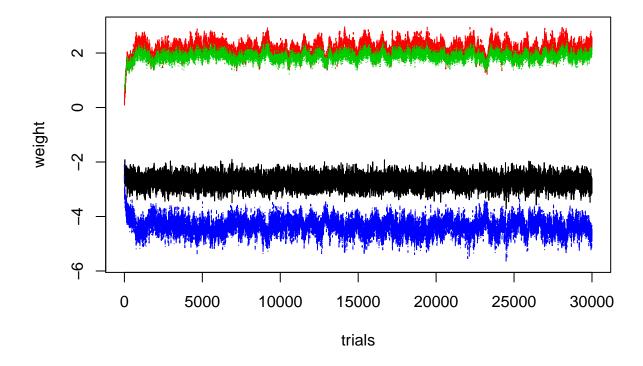
original beta

sampled beta

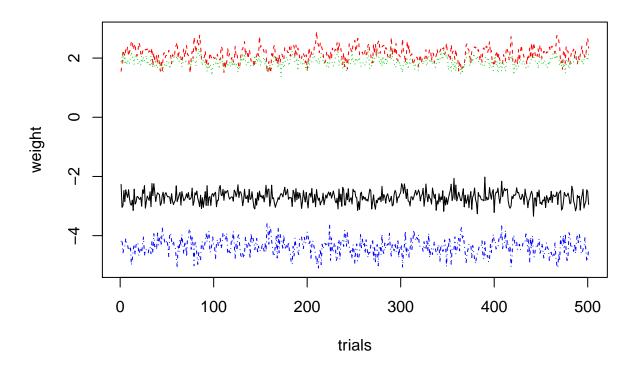
Blocked Gibbs Sampling Assessment: Cutoffs



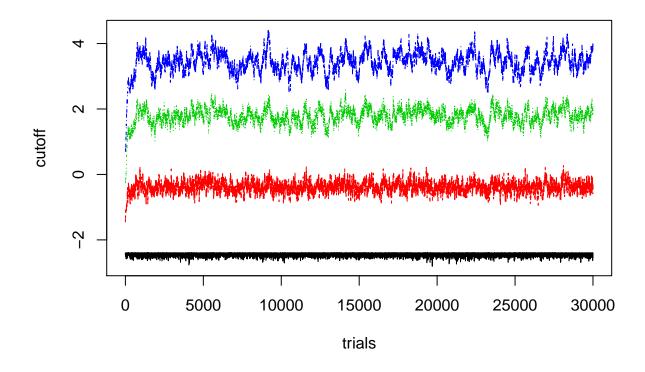
Checking stability of sampled Beta



Checking stability of sampled Beta: thining



Checking stability of sampled Cutoffs



Checking stability of sampled Cutoffs: thining

