# STAT 471: Midterm Review Practice Problems (Solutions).

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| N  | ote: this document is adapted from last year's midterm.                        |   |  |  |  |

## Instructions

- This exam is open-book / open-notes / open-internet. However, it is individual work. Communication among students is prohibited.
- Please complete your homework in R Markdown, using this document as a starting point. Show your R code using code chunks and add your text answers using **bold text**.
- When you are ready to submit, please compile your R Markdown file into a PDF. Then, submit this PDF through Canvas.
- While base R programming is acceptable, I strongly encourage you to use the tidyverse to complete your assignment. However, points will not be deducted if you use base R programming.
- In addition, please make sure to have clear labels and sensible titles on your plots.
- Make sure that readmission\_clean.csv and plot\_glmnet.R are in your working directory before beginning the exam.

## Introduction: Predicting readmission for diabetes patients

## Background and goals

Diabetes is a chronic medical condition affecting millions of Americans, but if managed well, patients can lead relatively normal lives. However, if improperly managed, diabetes can lead to patients being continuously admitted and readmitted to hospitals. Hospital readmissions represent a failure of the health system to provide adequate support to the patient and are extremely costly to the system. The goals are therefore to

- 1. identify important factors associated with readmission, and
- 2. predict whether a given patient will be readmitted.

#### Readmission data

In this exam, we will investigate a dataset originally from the Center for Clinical and Translational Research at Virginia Commonwealth University, covering diabetes patients across 130 U.S. hospitals from 1999 to 2008. Three former STAT 471 students Spencer Luster, Matthew Lesser and Mridul Ganesh brought this data set into the class through their final project. In this exam, we will use a cleaned subset of this data.

First, let's load a few libraries:

```
library(kableExtra)  # for printing tables
library(cowplot)  # for side by side plots
library(glmnetUtils)  # to run ridge and lasso
library(lubridate)  # for dealing with dates
library(maps)  # for creating maps
source("../../functions/plot_glmnet.R") # for lasso/ridge trace plots
library(tidyverse)  # for everything else
```

Let's load the data:

```
readmission = read_csv("readmission_clean.csv")
readmission
```

```
## # A tibble: 99,492 x 26
##
      race
                      gender age_group num_outpatient num_inpatient num_emergency
##
      <chr>
                      <chr>
                              <chr>
                                                  <dbl>
                                                                <dbl>
                                                                               <dbl>
##
    1 Caucasian
                      Female 80+
                                                      0
                                                                    0
                                                                                   0
##
   2 Caucasian
                      Female 80+
                                                      0
                                                                    0
                                                                                   0
                                                      0
                                                                                   0
##
   3 Caucasian
                      Male
                              20-59
                                                                    0
##
    4 AfricanAmerican Female 20-59
                                                      0
                                                                    0
                                                                                   0
##
   5 Caucasian
                      Male
                              20-59
                                                      Λ
                                                                    0
                                                                                   Λ
   6 AfricanAmerican Male
##
                              60 - 79
                                                      0
                                                                    0
                                                                                   0
                                                      Λ
##
    7 Caucasian
                      Female 20-59
                                                                    0
                                                                                   1
    8 Caucasian
                      Male
                              80+
                                                      0
                                                                    0
                                                                                   0
##
  9 Caucasian
                      Male
                              60 - 79
                                                      0
                                                                    0
                                                                                   0
##
## 10 AfricanAmerican Female 60-79
                                                      0
                                                                    0
                                                                                   0
## # ... with 99,482 more rows, and 20 more variables: num_medications <dbl>,
## #
       num_diagnoses <dbl>, adm_source <chr>, adm_type <chr>,
       time_in_hospital <dbl>, num_lab_procedures <dbl>, num_procedures <dbl>,
## #
## #
       discharge <chr>, max_glu_serum <chr>, A1Cresult <chr>, med_changed <chr>,
## #
       med_prescribed <chr>, insulin <chr>, metformin <chr>, glimepiride <chr>,
## #
       glipizide <chr>, glyburide <chr>, pioglitazone <chr>, rosiglitazone <chr>,
## #
       readmitted <dbl>
```

Each row corresponds to a hospital admission of a patient. There are 26 total variables, described below:

Demographic variables

race: patient's racegender: patient's genderage\_group: patient's age group

## Medical history

- num\_outpatient: number of outpatient visits by the patient in the year prior to the current admission
- $\bullet\,$  num\_inpatient: number of inpatient visits by the patient in the year prior to the current admission
- num\_emergency: number of emergency visits by the patient in the year prior to the current admission

- num medications: number of total medications the patient has taken
- num\_diagnoses: number of total diagnoses the patient has

#### Hospital admission details

- adm\_source: who referred the patient to the hospital
- adm\_type: type of admission
- time in hospital: length of stay in the hospital (in days)
- num\_lab\_procedures: number of lab procedures performed
- num procedures: number of non-lab procedures performed
- discharge: where the patient was discharged

#### Clinical results

- max\_glu\_serum: results of glucose serum test
- A1Cresult: results of A1c test

#### Medication details

- med\_changed: whether any medication was changed
- med prescribed: whether any medication was prescribed
- insulin: type of change (if any) to insulin medication
- metformin: type of change (if any) to insulin medication
- glimepiride: type of change (if any) to glimepiride medication
- glipizide: type of change (if any) to glipizide medication
- glyburide: type of change (if any) to glyburide medication
- pioglitazone: type of change (if any) to pioglitazone medication
- rosiglitazone: type of change (if any) to rosiglitazone medication

## $Readmission\ indicator$

• readmitted: whether the patient was readmitted to the hospital within 30 days of discharge

## Train/test split

Let's subsample 10000 observations for training and subsample a non-overlapping 2000 observations for testing:

```
set.seed(1)
n_total = nrow(readmission)
n_train = 10000
n_test = 2000
train_samples = sample(1:n_total, n_train)
test_samples = sample((1:n_total)[-train_samples], n_test)
readmission_train = readmission %>% filter(row_number() %in% train_samples)
readmission_test = readmission %>% filter(row_number() %in% test_samples)
```

## 1 Data wrangling and exploratory data analysis

First, let's do some exploratory data analysis on our training data readmission\_train.

1. For people in the age of 0–19, what are the frequencies of adm\_source? Present the result in a nice table.

Table 1: Frequencies of admission source for patiens in the age of 0-19

| adm_source                | frequency |
|---------------------------|-----------|
| Emergency Room            | 55        |
| Other                     | 5         |
| Physician Referral        | 23        |
| Transfer from Home Health | 1         |

Table 2: Case numbers for each gender-age combination

| Gender | Age     | Num outpatient | Num inpatient | Num emergency | Grand total |
|--------|---------|----------------|---------------|---------------|-------------|
| Female | 0-19    | 6              | 22            | 9             | 37          |
| Female | 20 - 59 | 604            | 1325          | 670           | 2599        |
| Female | 60-79   | 1059           | 1585          | 451           | 3095        |
| Female | 80 +    | 452            | 836           | 210           | 1498        |
| Male   | 0-19    | 0              | 30            | 1             | 31          |
| Male   | 20-59   | 417            | 976           | 395           | 1788        |
| Male   | 60-79   | 949            | 1370          | 293           | 2612        |
| Male   | 80+     | 331            | 399           | 95            | 825         |

```
caption = "Frequencies of admission source for patiens in the age of 0-19") %>%
kable_styling(position = "center")
```

2. For each unique (gender, age\_group) combination, calculate the sum of num\_outpatient, num\_inpatient, num\_emergency, then calculate the total sum of those three variables. Tabulate the result in a fancy table.

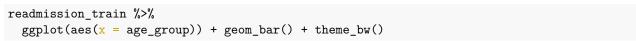
- ## `summarise()` has grouped output by 'gender'. You can override using the `.groups` argument.
  - 3. What fraction of the patients in the training data were readmitted?

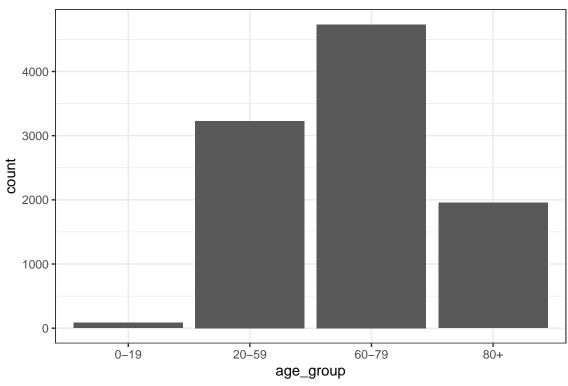
```
readmission_train %>% summarise(frac_readmitted = mean(readmitted))
```

```
## # A tibble: 1 x 1
## frac_readmitted
## <dbl>
## 1 0.117
```

We see that 0.117 of the patients in the training data were readmitted.

4. Produce a bar plot to display the breakdown of the patients by age group. What is the most prevalent age group in the training data?



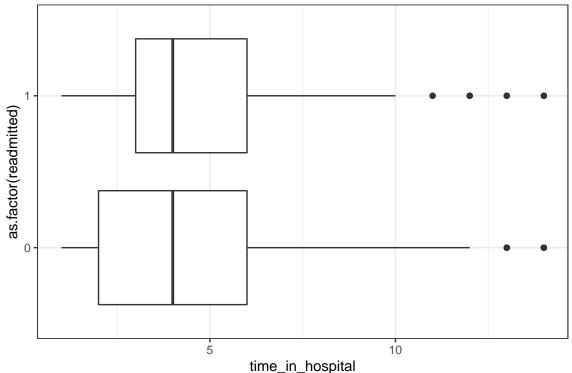


## The most prevalent age group is 60-79.

5. Produce a plot to show the relationship between time\_in\_hospital and readmitted. Using summarise, compute the median time in hospital separately for patients that were not readmitted and for those that were. Do these suggest that readmission rates vary based on time in hospital, and if so, what is the direction of the relationship?

(Hint: It may be useful to convert readmitted to a factor using as.factor(readmitted).)

```
readmission_train %>% ggplot(aes(x = time_in_hospital, y = as.factor(readmitted))) + geom_boxplot() + theme_bw()
```



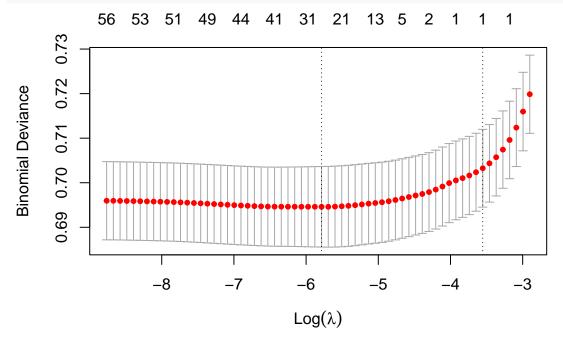
The median number of days in the hospital was 4 days for both groups of patients. This fact, along with the plot above, suggests that time in hospital is not associated with readmission.

## 2 Prediction via regularized logistic regression

i. Fit a 10-fold cross-validated lasso logistic regression to the training data.

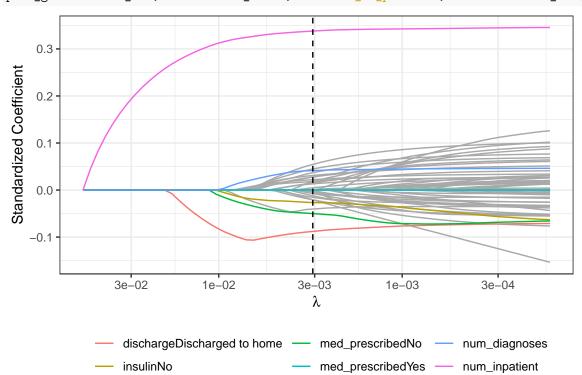
ii. Produce the CV plot, and give interpretations for this plot.

## plot(lasso\_fit)

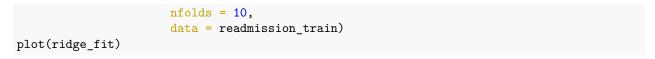


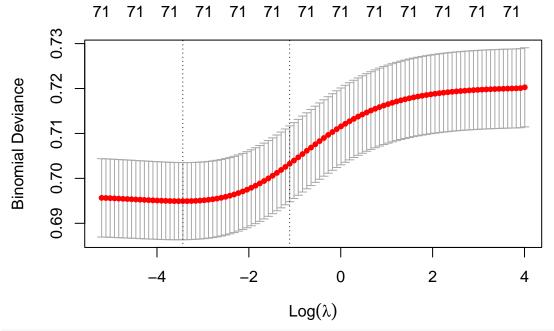
iii. Produce the trace plot based on lambda.min. Give interpretations.

plot\_glmnet(lasso\_fit, readmission\_train, features\_to\_plot = 6, lambda = lasso\_fit\$lambda.min)



iv. Repeat steps i-iii, this time using ridge regression





plot\_glmnet(ridge\_fit, readmission\_train, features\_to\_plot = 6, lambda = ridge\_fit\$lambda.min)

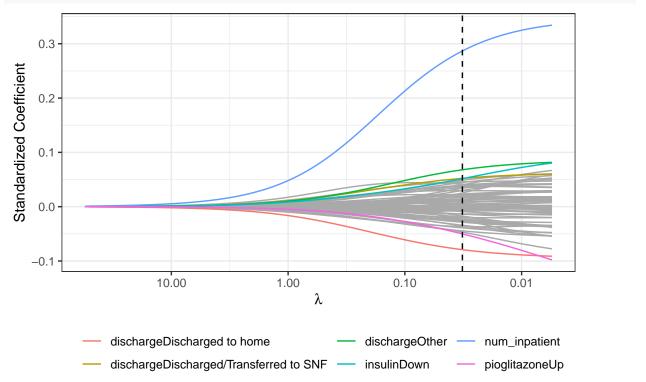


Table 3: Misclassification error of Lasso regression with different thresholds

| Threshold | Lasso error | Lasso FPR | Lasso FNR |
|-----------|-------------|-----------|-----------|
| 0.1       | 0.49        | 0.51      | 0.30      |
| 0.3       | 0.11        | 0.02      | 0.98      |
| 0.5       | 0.10        | 0.00      | 0.99      |
| 0.7       | 0.10        | 0.00      | 1.00      |

## 3 Model evaluation with different probability thresholds

i. Let us consider the following prediction rule. For a predicted  $\hat{y}$ , if it exceeds some threshold, we let the final prediction be one; otherwise, we let the final prediction be zero. For a grid of threshold = 0.1, 0.3, 0.5, 0.7, compute the mis-classification error, false positive rate (FP / N), and false negative rate (FN / P) for lasso regression using lambda.min. Which threshold is the best? Tabulate the results in a nice table.

```
get_metrics = function(yhat, y){
  err = mean(yhat != y)
  positive = sum(y)
  negative = length(y) - positive
  fpr = sum((yhat == 1) & (y == 0)) / negative
  fnr = sum((yhat == 0) & (y == 1)) / positive
  return(c(err, fpr, fnr))
}
groundtruth = readmission_test %>% select(readmitted) %>% pull()
lasso_predictions = predict(lasso_fit, newdata = readmission_test,
                            s = 'lambda.min', type='response') %>%
  as.numeric()
lasso_err = c(0, 0, 0, 0)
lasso_fpr = c(0, 0, 0, 0)
lasso_fnr = c(0, 0, 0, 0)
thresholds = c(0.1, 0.3, 0.5, 0.7)
for(i in 1:4) {
  lasso_predictions_binary = as.integer(lasso_predictions >= thresholds[i])
  curr_metrics = get_metrics(lasso_predictions_binary, groundtruth)
  lasso err[i] = curr metrics[1]
 lasso_fpr[i] = curr_metrics[2]
  lasso_fnr[i] = curr_metrics[3]
tibble(threshold = thresholds, error = lasso_err, fpr = lasso_fpr, fnr = lasso_fnr) %>%
   kable(format = "latex", row.names = NA,
        booktabs = TRUE, digits = 2,
        col.names = c("Threshold", "Lasso error", "Lasso FPR", "Lasso FNR"),
        caption = "Misclassification error of Lasso regression with different thresholds") %>%
   kable_styling(position = "center")
```

ii. Do the same thing for ridge. For the same level of threshold, which model is better?

Table 4: Misclassification error of Ridge regression with different thresholds

| Threshold | Ridge error | Ridge FPR | Ridge FNR |
|-----------|-------------|-----------|-----------|
| 0.1       | 0.50        | 0.53      | 0.30      |
| 0.3       | 0.11        | 0.01      | 0.97      |
| 0.5       | 0.10        | 0.00      | 1.00      |
| 0.7       | 0.10        | 0.00      | 1.00      |

```
as.numeric()
ridge_err = c(0, 0, 0, 0)
ridge_fpr = c(0, 0, 0, 0)
ridge_fnr = c(0, 0, 0, 0)
thresholds = c(0.1, 0.3, 0.5, 0.7)
for(i in 1:4) {
  ridge_predictions_binary = as.integer(ridge_predictions >= thresholds[i])
  curr_metrics = get_metrics(ridge_predictions_binary, groundtruth)
  ridge_err[i] = curr_metrics[1]
  ridge_fpr[i] = curr_metrics[2]
  ridge_fnr[i] = curr_metrics[3]
tibble(threshold = thresholds, error = ridge_err, fpr = ridge_fpr, fnr = ridge_fnr) %>%
    kable(format = "latex", row.names = NA,
       booktabs = TRUE, digits = 2,
        col.names = c("Threshold", "Ridge error", "Ridge FPR", "Ridge FNR"),
        caption = "Misclassification error of Ridge regression with different thresholds") %>%
    kable_styling(position = "center")
```