## ISYE 6501 Homework 10

2023-10-24

### Question 14.1

The breast cancer data set breast-cancer-wisconsin.data.txt has missing values.

#### Part 1

Use the mean/mode imputation method to impute values for the missing data.

First, I loaded and cleaned the data set. As you can see from the summary, there are 16 missing data points in the Bare\_nuclei column.

```
#load data
cancer_data <- read.csv("~/Z. OMSA/Intro to Analytics Modeling/Week 10 Homework/breast-cancer-wisconsin
                         header=FALSE)
#add column names
colnames(cancer_data) <- c("Sample_code_number",</pre>
                            "Clump_thickness",
                            "Uniformity_of_cell_size",
                            "Uniformity_of_cell_shape",
                            "Marginal_adhesion",
                            "Single_epithelial_size",
                            "Bare_nuclei",
                            "Bland_chromatin",
                            "Normal_nucleoli",
                            "Mitoses",
                            "Class")
#update binary variables
cancer_data$Class[cancer_data$Class == 2] <- 0 #benign</pre>
cancer_data$Class[cancer_data$Class == 4] <- 1 #malignant</pre>
#change ? to NA
cancer_data[cancer_data == "?"] <- NA</pre>
#make Bare nuclei column numeric
cancer_data$Bare_nuclei <- as.numeric(cancer_data$Bare_nuclei)</pre>
#summarize data
summary(cancer_data)
```

```
1st Qu.: 870688
                        1st Qu.: 2.000
                                          1st Qu.: 1.000
##
##
    Median : 1171710
                        Median: 4.000
                                          Median : 1.000
##
    Mean
           : 1071704
                        Mean
                               : 4.418
                                          Mean
                                                 : 3.134
                        3rd Qu.: 6.000
                                          3rd Qu.: 5.000
##
    3rd Qu.: 1238298
##
    Max.
           :13454352
                        Max.
                               :10.000
                                          Max.
                                                 :10.000
##
##
    Uniformity_of_cell_shape Marginal_adhesion Single_epithelial_size
##
           : 1.000
                              Min.
                                     : 1.000
                                                 Min.
                                                       : 1.000
##
    1st Qu.: 1.000
                              1st Qu.: 1.000
                                                 1st Qu.: 2.000
##
    Median : 1.000
                              Median : 1.000
                                                 Median : 2.000
    Mean
           : 3.207
                              Mean
                                     : 2.807
                                                 Mean
                                                         : 3.216
    3rd Qu.: 5.000
                              3rd Qu.: 4.000
##
                                                 3rd Qu.: 4.000
           :10.000
##
    Max.
                              Max.
                                      :10.000
                                                         :10.000
                                                 Max.
##
##
     Bare_nuclei
                      Bland_chromatin Normal_nucleoli
                                                             Mitoses
##
    Min.
           : 1.000
                             : 1.000
                                       Min.
                                               : 1.000
                                                                 : 1.000
                      Min.
                                                          Min.
                      1st Qu.: 2.000
                                                          1st Qu.: 1.000
##
    1st Qu.: 1.000
                                        1st Qu.: 1.000
    Median : 1.000
                      Median : 3.000
                                       Median : 1.000
                                                          Median : 1.000
                                               : 2.867
##
    Mean
           : 3.545
                             : 3.438
                                                                 : 1.589
                      Mean
                                       Mean
                                                          Mean
##
    3rd Qu.: 6.000
                      3rd Qu.: 5.000
                                        3rd Qu.: 4.000
                                                          3rd Qu.: 1.000
##
    Max.
           :10.000
                      Max.
                             :10.000
                                       Max.
                                               :10.000
                                                          Max.
                                                                 :10.000
    NA's
           :16
##
##
        Class
##
   Min.
           :0.0000
##
   1st Qu.:0.0000
##
   Median :0.0000
##
   Mean
           :0.3448
##
    3rd Qu.:1.0000
##
           :1.0000
    Max.
##
```

As you can see in the code block below, the 16 missing data points represent less than 5% of the data set. Therefore, it is appropriate to use imputation methods to address missing data.

```
#total missing data points
missing_data <- sum(is.na(cancer_data))
missing_data</pre>
```

## [1] 16

```
#percentage of missing data points
pct_missing <- missing_data/nrow(cancer_data)
pct_missing</pre>
```

```
## [1] 0.02288984
```

Next, I created a histogram to visualize the spread of Bare\_nuceli data. I used this as a baseline to compare with my imputed data sets. The imputed data sets should have a very similar distribution to the baseline.

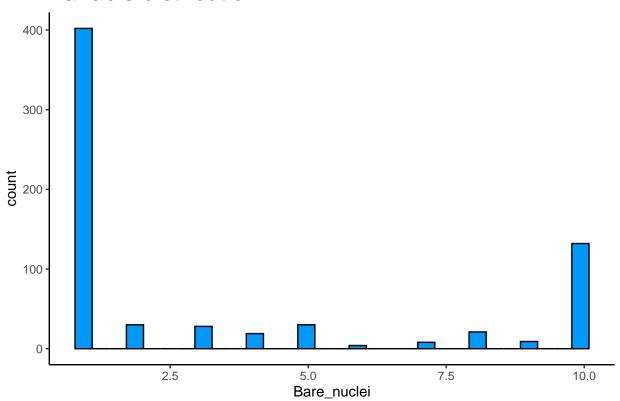
```
#visualize spread of data
ggplot(cancer_data, aes(Bare_nuclei)) +
  geom_histogram(color = "#000000", fill = "#0099F8") +
```

```
ggtitle("Variable distribution") +
theme_classic() +
theme(plot.title = element_text(size = 18))
```

## 'stat\_bin()' using 'bins = 30'. Pick better value with 'binwidth'.

## Warning: Removed 16 rows containing non-finite values ('stat\_bin()').

# Variable distribution



Then, I calculated the mean of the Bare\_nuclei column and used this value to replace the missing data.

```
#copy data set
cancer_data_mean <- cancer_data
#impute with mean
mean <- round(mean(cancer_data_mean$Bare_nuclei, na.rm=TRUE), 2)
print(paste0("Mean: ", mean))

## [1] "Mean: 3.54"

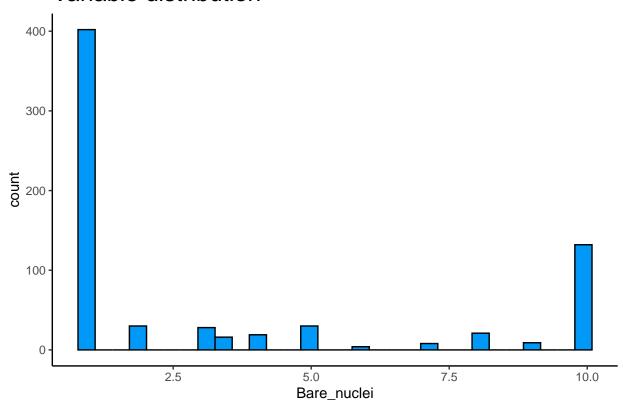
cancer_data_mean[is.na(cancer_data_mean)] <- mean</pre>
```

The code block below shows the new spread of data.

```
#visualize spread of data after imputation
ggplot(cancer_data_mean, aes(Bare_nuclei)) +
  geom_histogram(color = "#000000", fill = "#0099F8") +
  ggtitle("Variable distribution") +
  theme_classic() +
  theme(plot.title = element_text(size = 18))
```

## 'stat\_bin()' using 'bins = 30'. Pick better value with 'binwidth'.

# Variable distribution



I also used the mode to impute missing values.

```
#copy data set
cancer_data_mode <- cancer_data
#impute with mode
mode <- Mode(cancer_data_mode$Bare_nuclei, na.rm =TRUE)
print(paste0("Mode: ", mode))

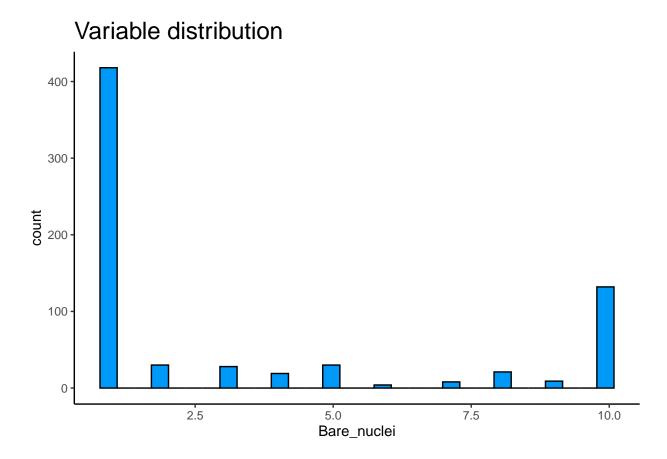
## [1] "Mode: 1"

cancer_data_mode[is.na(cancer_data_mode)] <- mode</pre>
```

The code block below shows the new spread of data.

```
#visualize spread of data after imputation
ggplot(cancer_data_mode, aes(Bare_nuclei)) +
  geom_histogram(color = "#000000", fill = "#0099F8") +
  ggtitle("Variable distribution") +
  theme_classic() +
  theme(plot.title = element_text(size = 18))
```

## 'stat\_bin()' using 'bins = 30'. Pick better value with 'binwidth'.



# Part 2 Use regression to impute values for the missing data.

I used the mice() function to automate the process of imputation with regression. I input my data set, a method for imputation ("norm.predict" for linear regression), and a value for "m" (the number of multiple imputations). The code block below shows the imputed values for the missing data over 1 cycle.

```
#copy data set
cancer_data_regression <- cancer_data
#impute with regression using mice()
mice_data <- mice(cancer_data_regression, method = "norm.predict", m=1)</pre>
```

```
## 1 1 Bare_nuclei
## 2 1 Bare_nuclei
## 3 1 Bare_nuclei
## 4 1 Bare_nuclei
## 5 1 Bare_nuclei
## 5 1 Bare_nuclei
##see imputed values over the five trials
mice_data[["imp"]]$Bare_nuclei
```

```
##
## 24 7.191237
## 41 3.419208
## 140 1.188951
## 146 1.579936
## 159 1.260453
## 165 1.428797
## 236 1.943842
## 250 1.562574
## 276 1.740990
## 293 6.432884
## 295 1.303253
## 298 1.186205
## 316 2.083334
## 322 1.469119
## 412 1.179806
## 618 1.059370
```

##

iter imp variable

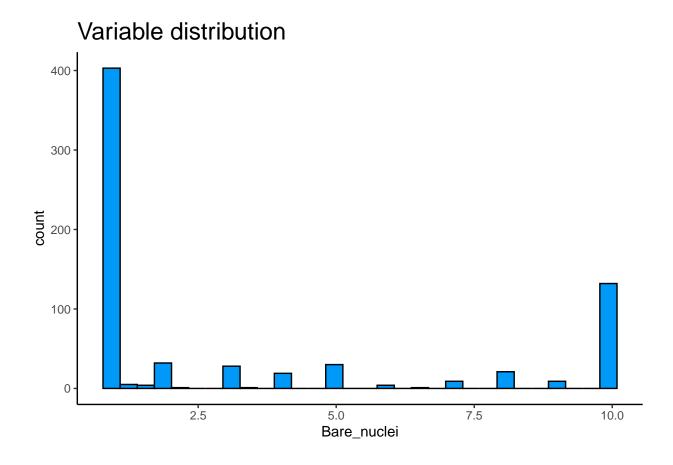
I used the complete() function to replace missing values in my original data set with the values imputed by the mice() function.

```
#update original data set
cancer_data_regression <- complete(mice_data, action = 1)</pre>
```

The code block below shows the new spread of data.

```
#visualize spread of data after imputation
ggplot(cancer_data_regression, aes(Bare_nuclei)) +
  geom_histogram(color = "#000000", fill = "#0099F8") +
  ggtitle("Variable distribution") +
  theme_classic() +
  theme(plot.title = element_text(size = 18))
```

## 'stat\_bin()' using 'bins = 30'. Pick better value with 'binwidth'.



Use regression with perturbation to impute values for the missing data.

I also used the mice() function to impute values using regression with perturbation. I used the "norm.nob" instead of the "norm.predict" method to impute with added variability.

```
#copy data set
cancer_data_regression_perturb <- cancer_data</pre>
#imputed values
mice_data <- mice(cancer_data_regression_perturb, method = "norm.nob", m=1)</pre>
##
##
    iter imp variable
##
         1 Bare_nuclei
##
         1 Bare_nuclei
##
            Bare_nuclei
     3
##
     4
            Bare_nuclei
     5
            Bare_nuclei
##
mice_data[["imp"]]$Bare_nuclei
```

## 1

Part 3

```
## 24
       7.4977412
## 41
       2.3875268
## 140 0.3901479
## 146 -1.2845248
## 159 -2.7607492
## 165 2.1056021
## 236 3.2466977
## 250 6.4299315
## 276
       4.1241822
## 293 8.2781692
## 295
       2.5457169
## 298 1.8990268
## 316 2.2256577
## 322 2.9330411
## 412 2.4196539
## 618 0.5741458
```

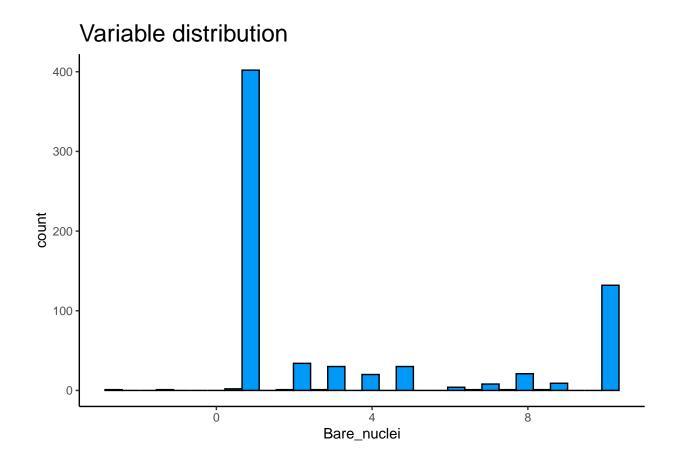
Then, I used the complete() function to merge the imputed values with my original data set.

```
#update original data set
cancer_data_regression_perturb <- complete(mice_data, action=1)</pre>
```

The code block below shows the new spread of data.

```
#visualize spread of data after imputation
ggplot(cancer_data_regression_perturb, aes(Bare_nuclei)) +
  geom_histogram(color = "#000000", fill = "#0099F8") +
  ggtitle("Variable distribution") +
  theme_classic() +
  theme(plot.title = element_text(size = 18))
```

## 'stat\_bin()' using 'bins = 30'. Pick better value with 'binwidth'.



#### Part 4 (Optional)

Compare the results and quality of classification models (e.g., SVM, KNN) build using the data sets from questions 1, 2, and 3.

As you can see in the code block below, knn model accuracy is the same for all 4 data sets. This is likely because very few values were changed in each data set. If more values were imputed, it is likely we would observe more variability.

```
#fit knn model
knn_model <- kknn(Class~., train_data, test_data, kernel="rectangular", scale=TRUE)

#make predictions and determine accuracy
predictions <- round(predict(knn_model))
accuracy <- sum(test_data$Class == predictions)/nrow(test_data)

#print result
print(paste0(data_set_descriptions[[ind]], "model accuracy: ", round(accuracy, 4)*100, "%"))
ind <- ind+1
}

## [1] "Imputation with Mean model accuracy: 96.79%"

## [1] "Imputation with Mode model accuracy: 96.79%"

## [1] "Imputation with Regression model accuracy: 96.79%"

## [1] "Imputation with Regression with Perturbation model accuracy: 96.79%"</pre>
```

# Question 15.1

Describe a situation or problem from your job, everyday life, current events, etc., for which optimization would be appropriate. What data would you need?

Google Maps uses optimization methods to provide users with efficient and real-time route recommendations. The technology considers traffic estimates, distance to location, speed limits, road closures, and driver preferences (e.g., avoid highways), among other factors to suggest a route that minimizes travel time.