

Running Head: HOSPITAL READMISSION ANALYSIS

# Hospital Readmission Prevalence: An analytics approach to reduce hospital readmissions

Anubha Adwani, Ankita Sharma, Ali Algarni, Imani Bansal

University of North Carolina Charlotte

## Table of Contents

Abstract .....	3
1. Introduction.....	4
2. Data Gathering and Analysis.....	4
2.1 Data Features.....	4
2.2 Data Pre-processing.....	5
2.3 Correlation Analysis .....	5
2.4 Regression .....	7
2.5 Chi Square Analysis .....	9
2.6 Principal Component Analysis .....	11
2.7 Pattern Mining.....	13
2.8 Feature Selection.....	14
2.8.1 Random Forest.....	14
2.8.2 Boruta Package.....	15
2.9 Classification .....	15
2.9.1 Decision Tree.....	16
2.9.2 Random Forest.....	17
2.9.3 Neural Network.....	18
2.9.4 K Nearest Neighbour.....	19
2.9.5 Evaluation of classification models.....	19
2.10 Clustering.....	20
2.10.1 PAM.....	20
2.10.2 Hierarchal Clustering.....	22
3. Findings and Analysis.....	23
4. References.....	24
5. Appendices.....	25

**Abstract**

Hospital readmission is the most important contributor towards total medical care expenditure and is an emerging indicator towards quality of care. Research studies show that nearly 15%-20% of the people discharged from the hospital are readmitted within 30 days from the date of discharge. Some of these readmissions are voluntarily, but some are preventable. The process of admitting a person, readmitting is equally tough for both the patient and the hospital. Reducing hospital readmissions is a win for both the patient and the hospital. Diabetes like any other chronic disease is associated with increased risk of hospital readmissions

Risk factor includes previous hospitalizations, age, gender, previous diagnoses and other socio economic barriers.

*Keywords:* diabetes, readmission, hospital, diagnoses.

## 1. Introduction

A significant proportion of medical costs are contributed by small percentage of chronic diseases like diabetes, heart problems, knee problems, etc. These costs are large due to the repeated readmission of the patient for the same problem. Prevention of unplanned or emergency readmissions is therefore increasingly gaining attention. The purpose of this report is to analyze the predictors in the unplanned readmissions and to describe the role of diagnoses of diabetes and glycemic control.

This report is about the analysis of a large clinical database, to examine the pattern of diabetes care in patients with diabetes that were admitted to hospitals and to reduce their readmissions. We examined the use of HbA1c results, the previous diagnoses of the patients, their hospital admission history.

## 2. Data Gathering and Analysis Techniques

### 2.1 Data features

The dataset given is a clinical dataset that contains historical records of patients pertaining to admission and readmission due to their diabetes. The dataset contains valuable but heterogeneous and incomplete data in terms of missing values, inconsistent records and high dimensionality. The data contains records that includes hospital admission information (inpatient, outpatient or emergency), demographic patient information (age, gender, race, weight), diagnoses documented by ICD-9 codes, laboratory data, pharmacy data, medical caregiver data. The database has 10k observations for 52 variables. The dataset has categorical, numerical, logical data types.

Binary Data	Readmitted
Numerical Data	rowID, num_lab_procedures, num_medications, num_outpatients, num_emergency, num_inpatient, time_in_hospital
Categorical Data	Race, gender, age, weight, medical_speciality, diag_1, diag_2, diag_3, A1C result, max_glu_serum, insulin, change, troglitazone, examide, glyburide,metformin, glipizide,repaglinide.

Table 1: Data Attributes and type

# HOSPITAL READMISSION ANALYSIS

```
> data<-read.csv("10kDiabetes.csv")
> str(data)
'data.frame': 10003 obs. of 52 variables:
 $ rowID      : int  1 2 3 4 5 6 7 8 9 10 ...
 $ race       : Factor w/ 7 levels "","?","AfricanAmerican",...: 5 5 5 3 3 5 5 5 5 5 ...
 $ gender     : Factor w/ 3 levels "","Female","Male": 2 2 2 2 3 2 2 3 3 ...
 $ age        : Factor w/ 11 levels "","[0-10)","[10-20)",...: 7 4 10 7 7 9 8 7 7 8 ...
 $ weight     : Factor w/ 9 levels "","?","[0-25)",...: 2 8 2 2 2 2 2 2 2 ...
 $ admission_type_id : Factor w/ 7 levels "","Elective",...: 2 7 5 3 3 2 2 3 1 2 ...
 $ discharge_disposition_id : Factor w/ 22 levels "","Admitted as an inpatient to this hospital",...: 3 3 11 3 3 3 18 3 3 3 ...
 $ admission_source_id : Factor w/ 11 levels "","Clinic Referral",...: 8 8 1 11 4 8 8 4 1 8 ...
 $ time_in_hospital : num  1 2 7 4 5 4 6 2 3 5 ...
 $ payer_code  : Factor w/ 17 levels "","?", "BC", "CH",...: 6 16 9 16 2 2 9 2 2 2 ...
 $ medical_specialty : Factor w/ 54 levels "","?", "Anesthesiology-Pediatric",...: 48 2 8 2 38 4 15 2 8 45 ...
 $ num_lab_procedures : num  35 8 12 33 31 29 46 49 54 47 ...
 $ num_procedures : num  4 5 0 1 0 0 1 1 0 2 ...
 $ num_medications : num  21 5 21 5 13 10 20 17 10 12 ...
 $ number_outpatient : num  0 0 0 0 0 0 0 2 0 0 ...
 $ number_emergency : num  0 0 0 0 0 0 0 1 0 0 ...
 $ number_inpatient : num  0 0 1 0 0 0 0 1 1 0 ...
 $ diag_1       : Factor w/ 459 levels "","?", "11", "110",...: 351 325 222 329 118 185 191 263 185 196 ...
 $ diag_2       : Factor w/ 431 levels "","?", "11", "110",...: 306 285 173 157 45 172 129 240 170 387 ...
 $ diag_3       : Factor w/ 462 levels "","?", "110", "112",...: 318 91 83 43 103 168 262 196 305 402 ...
 $ number_diagnoses : int  9 6 9 3 7 8 8 9 9 5 ...
 $ max_glu_serum : Factor w/ 5 levels "",">200",">300",...: 4 4 2 4 4 4 4 4 4 4 ...
 $ A1Cresult    : Factor w/ 5 levels "",">7",">8","None",...: 4 4 4 4 4 4 4 4 5 4 ...
 $ metformin    : Factor w/ 5 levels "","Down","No",...: 3 3 3 3 4 4 3 4 3 3 ...
 $ repaglinide  : Factor w/ 5 levels "","Down","No",...: 3 3 3 3 3 3 3 3 3 3 ...
 $ nateglinide  : Factor w/ 5 levels "","Down","No",...: 3 3 3 3 3 3 3 3 3 3 ...
 $ chlorpropamide : Factor w/ 4 levels "","No","Steady",...: 2 2 2 2 2 2 2 2 2 2 ...
 $ glimepiride  : Factor w/ 5 levels "","Down","No",...: 3 3 3 3 3 3 3 3 3 3 ...
 $ acetohexamide : Factor w/ 2 levels "","No": 2 2 2 2 2 2 2 2 2 2 ...
 $ glipizide    : Factor w/ 5 levels "","Down","No",...: 3 3 3 3 4 3 3 3 3 4 ...
 $ glyburide    : Factor w/ 5 levels "","Down","No",...: 3 3 3 3 3 3 3 3 3 3 ...
 $ tolbutamide  : Factor w/ 3 levels "","No","Steady": 2 2 2 2 2 2 2 2 2 2 ...
 $ pioglitazone : Factor w/ 5 levels "","Down","No",...: 3 3 3 3 3 3 3 3 3 3 ...
 $ rosiglitazone : Factor w/ 5 levels "","Down","No",...: 3 3 3 3 3 3 3 3 3 3 ...
 $ acarbose     : Factor w/ 4 levels "","No","Steady",...: 2 2 2 2 2 2 2 2 2 2 ...
 $ miglitol     : Factor w/ 5 levels "","Down","No",...: 3 3 3 3 3 3 3 3 3 3 ...
 $ troglitazone : Factor w/ 2 levels "","No": 2 2 2 2 2 2 2 2 2 2 ...
 $ tolazamide   : Factor w/ 3 levels "","No","Steady": 2 2 2 2 2 2 2 2 2 2 ...
 $ examide      : Factor w/ 2 levels "","No": 2 2 2 2 2 2 2 2 2 2 ...
```

Picture 1: Structure of the initial dataset

## 2.2 Data Pre-processing

The dataset contains many incomplete, redundant, noisy information. The first step towards analysis is to have quality data. Data quality includes accuracy, completeness, consistency and interpretability. In this step, we examine the structure of the data and its quality. We found that there were several features in the data that could not be treated well as they had high percentage of missing data. These features were weight (97% missing data), payer code (40%) and medical specialty (47%). The weight feature was removed due to its highest percentage of missing data. Medical specialty was maintained as it is relevant to the hospital care-giver information. The missing values in the data were handled by kNN imputation. Imputation is a class of procedures that aims to fill the missing values with the estimated ones based on the information available. The kNN imputation replaces missing data with the corresponding value from the nearest neighbor.

## 2.3 Correlation Analysis

Correlation was performed to measure the variability of variables. It is a way of measuring the extent to which two variables are related. Correlation coefficient represented the linear dependencies of two variables or sets of data. The value of correlation coefficient ranges from -1 to +1 with a value of  $\pm 0.1$  shows small effect,  $\pm 0.3$  shows medium effect and  $\pm 0.5$  shows large effect. Also, the p-values shows us the significance level of the correlation test performed. Value of p less than 0.05 shows the high significance of the test.

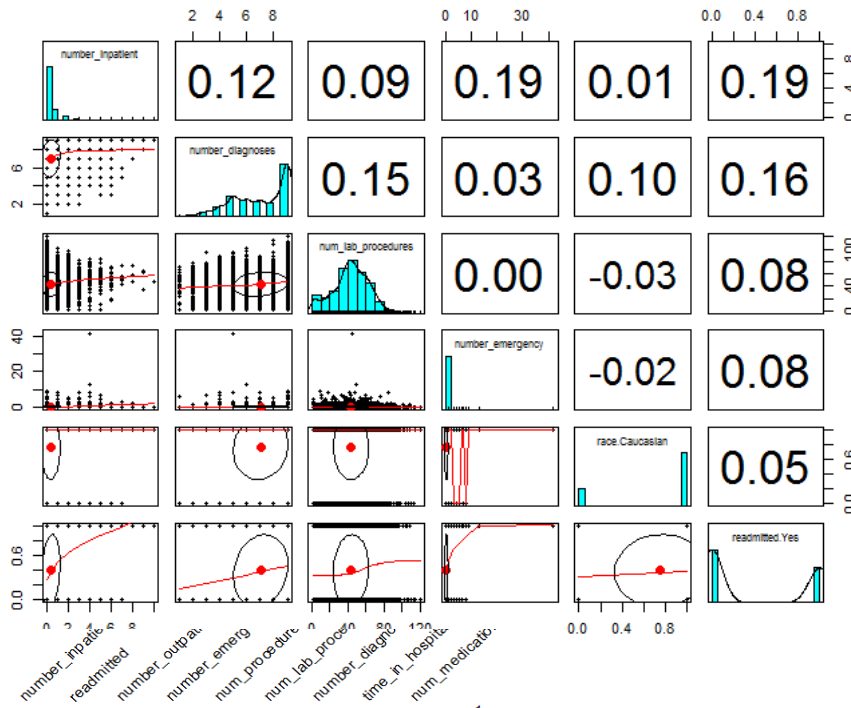
We have used two main functions `cor()` and `rcorr()` from Hmisc package to compute basic correlation coefficients. Pearson method is used for the correlation functions.

We have eliminated few of the variables that are not important for our analysis like payer code, index row, and description of the diagnoses. `cor()` and `rcorr()` were performed on total 42 variables and found out the most significant ones to list down here.

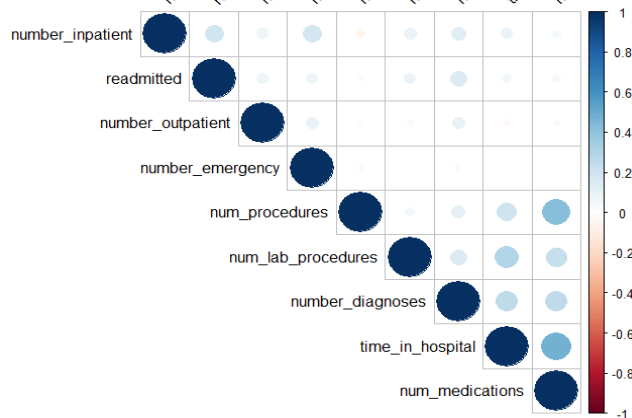
## HOSPITAL READMISSION ANALYSIS

Variables	Readmitted(r values)	Readmitted(p values)
race	0.04	0
gender	-0.01	0.21
age	0.08	0
admission_type_id	0	0.81
discharge_disposition_id	0	0.81
admission_source_id	-0.09	0
time_in_hospital	0.05	0
num_lab_procedures	0.08	0
num_procedures	-0.03	0.01
num_medications	0.04	0
number_outpatient	0.07	0
number_emergency	0.08	0
number_inpatient	0.19	0
number_diagnoses	0.16	0
max_glu_serum	0	0.64
A1C result	0	0.64
metformin	-0.02	0.05
repaglinide	0.02	0.02
nateglinide	0.01	0.24
chlorpropamide	0.01	0.46
glimepiride	0.02	0.09
glipizide	0.01	0.16

## HOSPITAL READMISSION ANALYSIS



This plot interprets that “number\_inpatient” and “num\_diagnoses” are positively correlated with “readmitted”.



The value of  $r$  is most significant for number\_inpatient and number\_diagnoses which is 0.19 and 0.16. Also, the p-value is  $<0.05$  which means these two variables are significantly correlated with the readmission and are important in predicting whether a patient will be readmitted or not.

*Corrplot: Using Method as Circle 1*

### 2.4 Regression

The next step in our hospital readmission analysis was regression which we have performed using logistic regression technique. Logistic regression is a method to predict an outcome variable that is categorical from predictor variables that are continuous and/or categorical.

In our analysis, the outcome variable is readmitted. Two models have been used to predict the value of outcome variable-first model consisting number\_inpatient as predictor variable and second model with number\_inpatient and number\_diagnoses as predictor variables. These two variables have been chosen as predictors because they have been found to be strongly correlated with the readmitted variable because of the correlation analysis.

Below is the summary of the models:

## HOSPITAL READMISSION ANALYSIS

```
> summary(regModel.1)

Call:
glm(formula = readmitted ~ number_inpatient, family = binomial(),
    data = reg_data)

Deviance Residuals:
    Min       1Q   Median       3Q      Max
-2.4368  -0.9293  -0.9293   1.4478   1.4478

Coefficients:
            Estimate Std. Error z value Pr(>|z|)
(Intercept)  -0.61627    0.02322  -26.54  <2e-16 ***
number_inpatient  0.50466    0.02790   18.09  <2e-16 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for binomial family taken to be 1)

    Null deviance: 13431  on 9999  degrees of freedom
Residual deviance: 13044  on 9998  degrees of freedom
AIC: 13048

Number of Fisher Scoring iterations: 4
```

```
> summary(regModel.2)

Call:
glm(formula = readmitted ~ number_inpatient + number_diagnoses,
    family = binomial(), data = reg_data)

Deviance Residuals:
    Min       1Q   Median       3Q      Max
-2.4372  -0.9974  -0.8249   1.3194   1.8350

Coefficients:
            Estimate Std. Error z value Pr(>|z|)
(Intercept)  -1.62204    0.08031  -20.20  <2e-16 ***
number_inpatient  0.46359    0.02788   16.63  <2e-16 ***
number_diagnoses  0.14380    0.01083   13.28  <2e-16 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for binomial family taken to be 1)

    Null deviance: 13431  on 9999  degrees of freedom
Residual deviance: 12862  on 9997  degrees of freedom
AIC: 12868

Number of Fisher Scoring iterations: 4
```

The odds ratio was found out for better understanding of the two models-

```
> exp(regModel.1$coefficients)
(Intercept) number_inpatient
0.5399532    1.6564174

> exp(regModel.2$coefficients)
(Intercept) number_inpatient number_diagnoses
0.1974953    1.5897764    1.1546509
```

Odds for number\_inpatient in 1st model is 1.65 which is greater than 1 that means as the predictor increases, the odds of the outcome increasing also increases. For the second model, it is 1.59 for number\_inpatient and 1.55 for number\_diagnoses which means if the values of these two variables will increase the odds of getting readmitted for patient will also increase.

```
> modelChi <- regModel.1$null.deviance - regModel.1$deviance
> modelChi
[1] 387.3111
> chidf <- regModel.1$df.null - regModel.1$df.residual
> chidf
[1] 1
> chisq.prob <- 1 - pchisq(modelChi, chidf)
> chisq.prob
[1] 0
> modelChi2 <- regModel.2$null.deviance - regModel.2$deviance
> modelChi2
[1] 569.679
> chidf2 <- regModel.2$df.null - regModel.2$df.residual
> chidf2
[1] 2
> chisq.prob2 <- 1 - pchisq(modelChi2, chidf2)
```

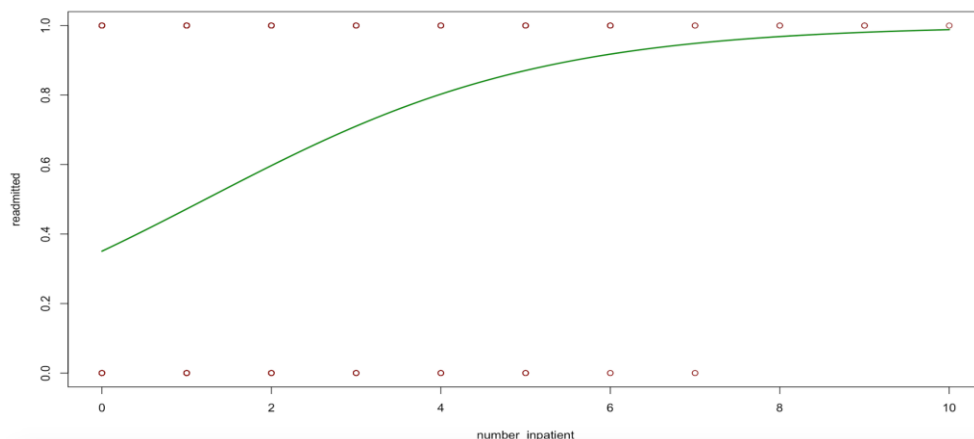


```
> chisq.prob2
```

```
[1] 0
```

The deviance for the 1st model is more as compared to the second one thus it can be concluded that the first model is better fit for predicting the outcome.

The relationship between number\_inpatient and readmitted can be visualized using the graph-



## 2.5 Chi Square Analysis

In case of categorical variables, we study their frequencies instead of the means as we do in numeric variables. This is because the mean of categorical variables does not convey any meaning, since they contain arbitrary values. To understand the relation between categorical variables, we implemented Pearson's chi square test of independence. This test is used to determine whether there is a significant association between two categorical variables or not. This test compares the observed frequencies in certain categories to the expected frequencies of the same category.

$$x^2 = (\text{observed} - \text{expected})^2 / \text{expected}$$

### Race and Readmission

We first took "race" and "readmitted" as a categorical feature and ran pearson's chi square test for them.

```
> count(diab_data1, 'race')
  race freq
1 AfricanAmerican 2090
2 Asian          55
3 Caucasian      7555
4 Hispanic       182
5 Other          121
> names(counts1) <- c("Race", "Readmitted", "Freq")
> counts1
```

Step 1: Frequency of each race of patients in the dataset.

```
race being
1 AfricanAmerican FALSE 1361
2 AfricanAmerican TRUE  729
3 Asian          FALSE  36
4 Asian          TRUE   19
5 Caucasian      FALSE 4442
6 Caucasian      TRUE 3111
7 Hispanic      FALSE 116
8 Hispanic      TRUE  65
9 Other         FALSE  80
10 Other        TRUE   41
```

Step 2: Frequency of each readmitted or not:

## HOSPITAL READMISSION ANALYSIS

## Step 3: Conduct the chi square test

```
> CrossTable(race_readmitted, chisq = TRUE, expected = TRUE, sresid = TRUE, format = "SPSS")
```

Cell Contents						
	Count	Expected Values	Chi-square Contribution	Row Percent	Column Percent	Total Percent
Total Observations in Table: 10000						
	african	caucasian	asian	hispanic	other	Row Total
TRUE	725 827.099 12.603 18.285% 34.756% 7.250% -3.550	3115 2996.351 4.698 78.562% 41.220% 31.150% 2.168	19 21.808 0.361 0.479% 34.545% 0.190% -0.601	65 71.766 0.638 1.639% 35.912% 0.650% -0.799	41 47.977 1.014 1.034% 33.884% 0.410% -1.007	3965 39.650%
FALSE	1361 1258.901 8.280 22.552% 65.244% 13.610% 2.878	4442 4560.650 3.087 73.604% 58.780% 44.420% -1.757	36 33.193 0.237 0.597% 65.455% 0.360% 0.487	116 109.234 0.419 1.922% 64.088% 1.160% 0.647	80 73.023 0.667 1.326% 66.116% 0.800% 0.816	6035 60.350%
Column Total	2086 20.860%	7557 75.570%	55 0.550%	181 1.810%	121 1.210%	10000

## Statistics for All Table Factors

## Pearson's Chi-squared test

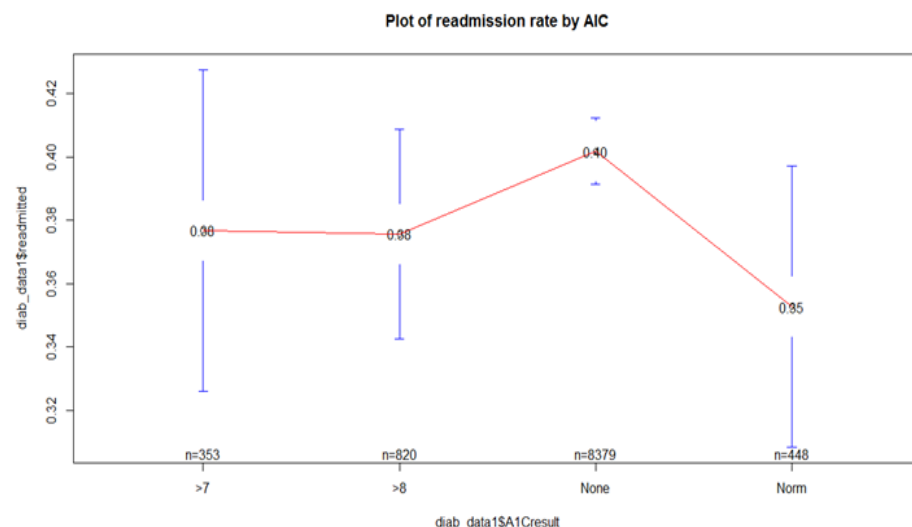
Chi^2 = 32.00584      d.f. = 4      p = 1.90785e-06

Minimum expected frequency: 21.8075

## Step 4: Result Interpretation:

The chi square test is highly significant with  $\chi^2(4) = 32.005$  and  $p < 0.05$ . This indicated that readmission rates vary along the race of the patient. Looking at the standard residuals, these results are non-significant for the Caucasian race, which show fairly even distribution for both readmitted (41%) and not admitted (59%). Within all other races, the standard residual varies much around 1.96. Like in the case of African race, expected readmission is higher than the observed frequency of readmitted patients.

Similarly, we found that “medical\_specialty” and “readmitted” are dependent with  $\chi^2(4) = 128.16$  and  $p < 0.05$ . It means that hospital care giver’s specialty played an important role in readmitting the patient. On the other hand, we found that “A1Cresult” and “readmitted” are independent with  $\chi^2(3) = 6.61$  and  $p > 0.05$ .



The graph shows that readmission rate is higher for patients for whom earlier A1C results were not considered or were not measured.

## 2.6 PCA

It is a multivariate technique that analyzes the data, in which observations are described by several inter correlated quantitative dependent variables. It is used to emphasize variation and bring out strong patterns in a dataset. The goal of doing PCA in our project was to extract the most important information from the data, compress the size of data by keeping only this important information and analyse the structure of observations and the variables.

Since this can be applied only to numeric variables, the attributes included in this were num\_lab\_procedures, num\_procedures, num\_medications, number\_outpatient, number\_emergency, number\_inpatient, time\_in\_hospital, number\_diagnoses and the response variable is readmitted.

### Step1: Correlation matrix of the data

```
> round(cor_intdata,2)
```

	time_in_hospital	num_lab_procedures	num_procedures	num_medications	number_outpatient	number_emergency	number_inpatient	number_diagnoses	readmitted
time_in_hospital	1.00	0.29	0.20	0.48	-0.03	-0.01	0.08	0.26	0.05
num_lab_procedures	0.29	1.00	0.05	0.24	-0.03	0.00	0.09	0.15	0.08
num_procedures	0.20	0.05	1.00	0.42	-0.02	-0.03	-0.06	0.10	-0.03
num_medications	0.48	0.24	0.42	1.00	0.03	0.01	0.04	0.25	0.04
number_outpatient	-0.03	-0.03	-0.02	0.03	1.00	0.08	0.08	0.10	0.07
number_emergency	-0.01	0.00	-0.03	0.01	0.08	1.00	0.19	0.03	0.08
number_inpatient	0.08	0.09	-0.06	0.04	0.08	0.19	1.00	0.12	0.19
number_diagnoses	0.26	0.15	0.10	0.25	0.10	0.03	0.12	1.00	0.16
readmitted	0.05	0.08	-0.03	0.04	0.07	0.08	0.19	0.16	1.00

```
> cortest.bartlett(diab_intdata, diag = TRUE)
R was not square, finding R from data
$chisq
[1] 8116.464

$ p.value
[1] 0

$df
[1] 36
```

Step2: Cortest Bartlett's Test: Along with correlation matrix, we run this test using cortest.barlette() function from 'psych' package. We ran this test on the raw data.

The test results imply that the results are significant with  $\chi^2(36) = 8116.46$  (having significance value  $<0.05$ ). This tells us that R-matrix is not an identity matrix. Therefore we proceed further with the PCA.

### Step 3: PCA with 9 factors

We create the first principal component model by taking all the numeric variables in the data and then deciding which components to keep.

```
pc1<- principal(diab_intdata, nfactors = 9)
```

## HOSPITAL READMISSION ANALYSIS

	RC8	RC2	RC6	RC1	RC5	RC3	RC4	RC7	RC9
SS loadings	1.01	1.01	1.00	1.00	1.00	1.00	1.00	1.00	0.97
Proportion Var	0.11	0.11	0.11	0.11	0.11	0.11	0.11	0.11	0.11
Cumulative Var	0.11	0.22	0.34	0.45	0.56	0.67	0.78	0.89	1.00
Proportion Explained	0.11	0.11	0.11	0.11	0.11	0.11	0.11	0.11	0.11
Cumulative Proportion	0.11	0.22	0.34	0.45	0.56	0.67	0.78	0.89	1.00

Mean item complexity = 1.1

Test of the hypothesis that 9 components are sufficient.

The thing to look here is the eigenvalues. The eigenvalues associated with each factor represents the variance explained by that linear component.

The result also displays the proportion of variance explained by each of the variable. Like component 8 explains 1.01 units of variance out of possible 9, to proportion this is

$$1.01/9 = 0.11.$$

These variables associated with the model are called values. We can find them by:

```
> pcl$values
```

```
[1] 2.0737141 1.4146662 1.0315128 0.9589719 0.8759846 0.7834234 0.7591759 0.6525837 0.4499673
```

According to, Kaiser's criterion, we retain only those components that have eigenvalue vectors greater than 1. In this case, we only retain 3 components.

#### Step 4: PCA with 3 factors

Now we do PCA again, only this time we have 3 factors instead of 9. The output shows the result of the second model.

```
> pc2<-principal(diab_intdata, nfactors=3)
> pc2
Principal Components Analysis
Call: principal(r = diab_intdata, nfactors = 3)
Standardized loadings (pattern matrix) based upon correlation matrix
```

	RC1	RC2	RC3	h2	u2	com
time_in_hospital	0.69	0.19	-0.27	0.58	0.42	1.5
num_lab_procedures	0.35	0.38	-0.51	0.52	0.48	2.7
num_procedures	0.68	-0.33	0.20	0.61	0.39	1.6
num_medications	0.84	0.00	0.00	0.70	0.30	1.0
number_outpatient	0.09	0.15	0.72	0.54	0.46	1.1
number_emergency	-0.02	0.38	0.45	0.35	0.65	1.9
number_inpatient	0.00	0.68	0.12	0.48	0.52	1.1
number_diagnoses	0.45	0.38	0.06	0.35	0.65	2.0
readmitted	0.02	0.61	0.03	0.38	0.62	1.0

```

SS loadings          RC1  RC2  RC3
1.97 1.45 1.10
Proportion Var      0.22 0.16 0.12
Cumulative Var      0.22 0.38 0.50
Proportion Explained 0.44 0.32 0.24
Cumulative Proportion 0.44 0.76 1.00

Mean item complexity = 1.5
Test of the hypothesis that 3 components are sufficient.

The root mean square of the residuals (RMSR) is 0.13
with the empirical chi square 11532.57 with prob < 0

Fit based upon off diagonal values = 0.38
```

```
> print.psych(pc2, cut = 0.3, sort = TRUE)
Principal Components Analysis
Call: principal(r = diab_intdata, nfactors = 3)
Standardized loadings (pattern matrix) based upon correlation matrix
```

	item	RC1	RC2	RC3	h2	u2	com
num_medications	4	0.84			0.70	0.30	1.0
time_in_hospital	1	0.69			0.58	0.42	1.5
num_procedures	3	0.68	-0.33		0.61	0.39	1.6
number_diagnoses	8	0.45	0.38		0.35	0.65	2.0
number_inpatient	7		0.68		0.48	0.52	1.1
readmitted	9		0.61		0.38	0.62	1.0
number_outpatient	5			0.72	0.54	0.46	1.1
num_lab_procedures	2	0.35	0.38	-0.51	0.52	0.48	2.7
number_emergency	6		0.38	0.45	0.35	0.65	1.9

```

SS loadings          RC1  RC2  RC3
1.97 1.45 1.10
Proportion Var      0.22 0.16 0.12
Cumulative Var      0.22 0.38 0.50
Proportion Explained 0.44 0.32 0.24
Cumulative Proportion 0.44 0.76 1.00

Mean item complexity = 1.5
Test of the hypothesis that 3 components are sufficient.

The root mean square of the residuals (RMSR) is 0.13
with the empirical chi square 11532.57 with prob < 0

Fit based upon off diagonal values = 0.38
```

We can interpret these results using print.psych() function as follows:

From these results we can interpret that num\_medication have a higher loading on component 1 i.e num\_diagnoses. So is num\_procedures. Component 1 explains 44% of the total variance. We can interpret that if a person is

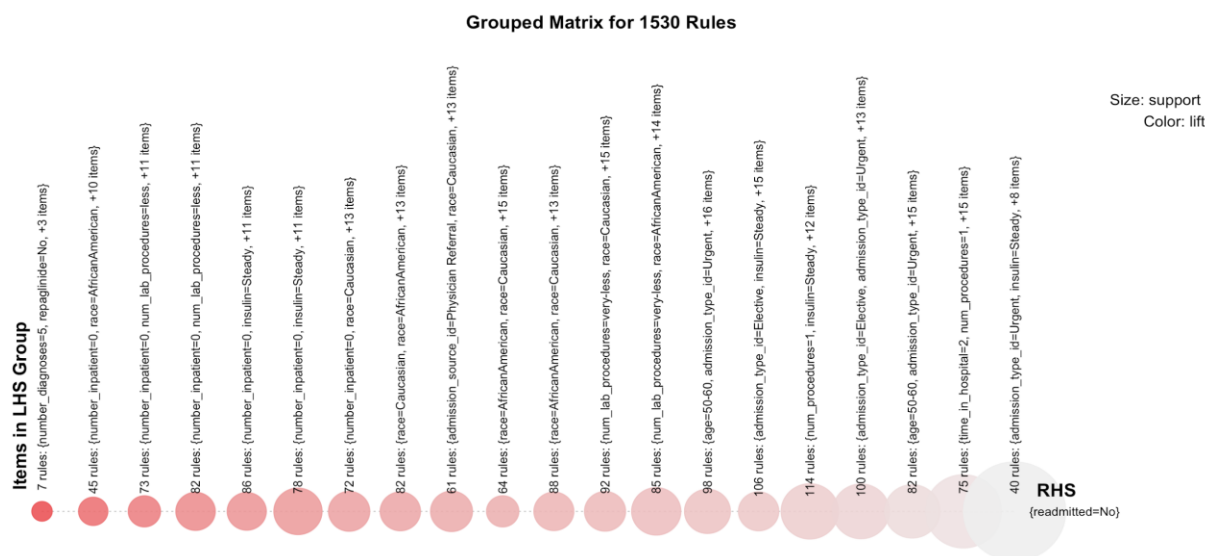
diagnosed with multiple symptoms, he has more number of procedures and so the number of medications increases

## 2.7 Pattern Mining

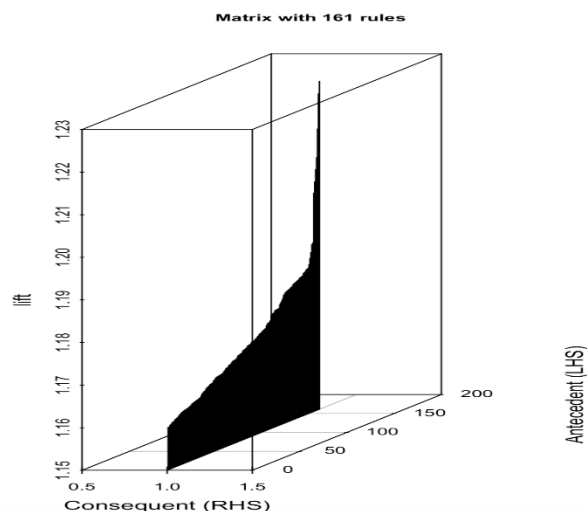
Pattern mining was done using apriori algorithm which is an influential algorithm for mining frequent item sets for Boolean association rules. This algorithm uses a bottom up approach where frequent subsets are extended one at a time.

Apriori was performed on 20 important variables that were extracted using feature selection method. Few of the variables that were numerical were first converted into factorial form such as num\_lab\_procedure and num\_medications and were given a label per values like 0-25 for very less, 25-50 for moderate and so on. Apriori was applied on transformed data by taking support value of 0.1, confidence value of 0.6, minimum length of rules as 2 and readmitted as RHS. Rules were sorted and a subset matrix was formed to eliminate the redundant rules which was done by providing the values in the lower subset triangle to be zero. Redundant rules were eliminated and pruned rules were identified from the entire set of rules. There were total 1530 pruned rules identified.

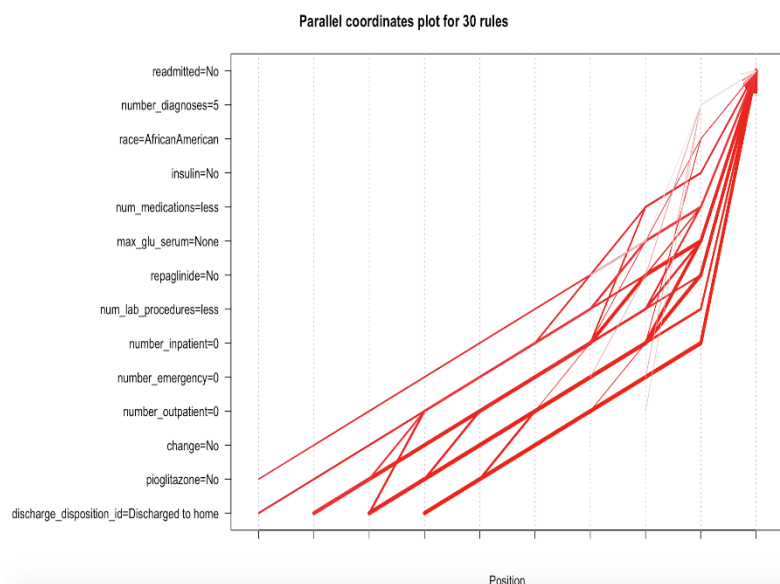
Below is the plot to visualize how these rules were categorized per lift and support values.



After the pruned rules, have been identified, a subset of rules was identified with confidence value greater than 0.7. This subset of rules contained total 161 rules which are important to predict frequent pattern that can lead to readmission values to be true or false.



3-D matrix graph to represent these rules with 'lift' on the y-axis and readmitted=No on the x-axis. Readmitted value is 'no' for almost all the rules because most of the rows have readmitted as false.



Parallel coordinate graph to visualize and understand the relationships.

### Interpretation-

From the parallel coordinate graph of the 30 highest lift valued rules it can be interpreted that patients with less number of outpatient, less number of inpatient, less number of medications, no insulin taken, less number of emergency, who have not taken pioglitazone and repaglinide medicines, and whose race is african american are less likely to be readmitted.

## 2.8 Feature Selection

To interpret the data in a more meaningful form, it is necessary to reduce the number of variables and remove attributes with less contribution in our data. Feature selection is important to subset important relevant attributes to build model and predict accurately. It helps to deal with curse of dimensionality, shortens the training time, improves results and avoids over fitting. We did feature selection using Random Forest and Boruta Package.

### 2.8.1 Random Forest

A popular automatic method for feature selection provided by the caret R package is called Recursive Feature Elimination or RFE. The algorithm is configured to explore all the possible subsets of the attribute. All the attributes are selected in this algorithm.

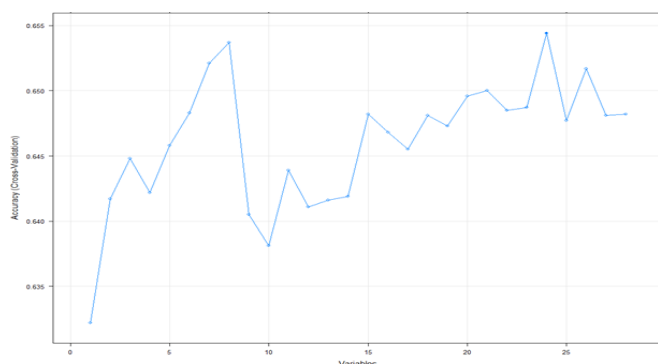
The output of rfe () is:

```
> print(results)
Recursive feature selection
Outer resampling method: Cross-validated (10 fold)
Resampling performance over subset size:
  Variables Accuracy  Kappa AccuracySD KappaSD Selected
1 0.6322 0.1779 0.015071 0.03073
2 0.6417 0.1931 0.014120 0.02946
3 0.6448 0.1879 0.014383 0.03199
4 0.6422 0.1897 0.010605 0.02250
5 0.6458 0.2002 0.012145 0.02902
6 0.6483 0.2027 0.014120 0.03328
7 0.6521 0.2092 0.013355 0.02927
8 0.6537 0.2140 0.012917 0.02867
9 0.6405 0.2072 0.012398 0.02901
10 0.6381 0.2067 0.006293 0.01650
11 0.6439 0.2183 0.010557 0.02737
12 0.6411 0.2115 0.014184 0.03514
13 0.6416 0.2140 0.016712 0.03898
14 0.6419 0.2155 0.015317 0.03658
15 0.6482 0.2291 0.009352 0.02339
16 0.6468 0.2320 0.008683 0.02070
17 0.6455 0.2276 0.013967 0.03348
18 0.6481 0.2321 0.015594 0.03782
19 0.6473 0.2309 0.013567 0.03382
20 0.6496 0.2340 0.016382 0.03891
21 0.6500 0.2329 0.016204 0.04029
22 0.6485 0.2294 0.012598 0.03102
23 0.6487 0.2283 0.016350 0.03954
24 0.6544 0.2400 0.017465 0.04151
25 0.6477 0.2291 0.018247 0.04404
26 0.6517 0.2354 0.015803 0.03894
27 0.6481 0.2273 0.016907 0.04204
28 0.6482 0.2275 0.013899 0.03436

The top 5 variables (out of 24):
  number_inpatient, discharge_disposition_id, number_diagnoses, number_outpatient, number_emergency
```

The output gives us top 5 important variables, number\_inpatient, discharge\_disposition\_id, number\_diagnoses, number\_outpatient and number\_emergency.





From this graph, we can interpret that 24 variables gives the most comparable result.

### 2.8.1 Boruta Package

The method performs top down search for relevant features by comparing original attributes importance with the importance achievable at random, progressively eliminating irrelevant features.

The output of running boruta() are:

```
> final_feature
Boruta performed 99 iterations in 11.06368 mins.
Tentatives roughfixed over the last 99 iterations.
22 attributes confirmed important: admission_source_id,
admission_type_id, age, change, diag_1 and 17 more;
10 attributes confirmed unimportant: AlCresult, diabetesMed, diag_2,
gender, glimepiride and 5 more;
> getSelectedAttributes(final_feature, withTentative = F)
[1] "race" "age" "admission_type_id" "discharge_disposition_id"
[5] "admission_source_id" "payer_code" "medical_specialty" "diag_1"
[9] "diag_3" "max_glu_serum" "repaglinide" "pioglitazone"
[13] "insulin" "change" "time_in_hospital" "num_lab_procedures"
[17] "num_procedures" "num_medications" "number_outpatient" "number_emergency"
[21] "number_inpatient" "number_diagnoses"
```

### 2.9 Classification:

We built different classification models and compared between them. We have tried to find the most accurate model based on evaluation parameters such as precision, recall, sensitivity, specificity and F1 measure. Also, we planned to finds any similarities between patterns extracted from each model. We used the following algorithms:

1. Decision tree
2. Random Forest
3. Neural Network
4. KNN

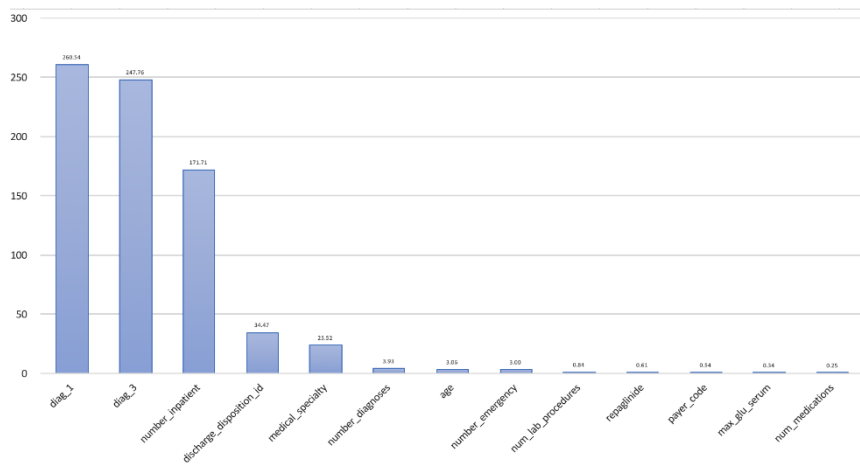
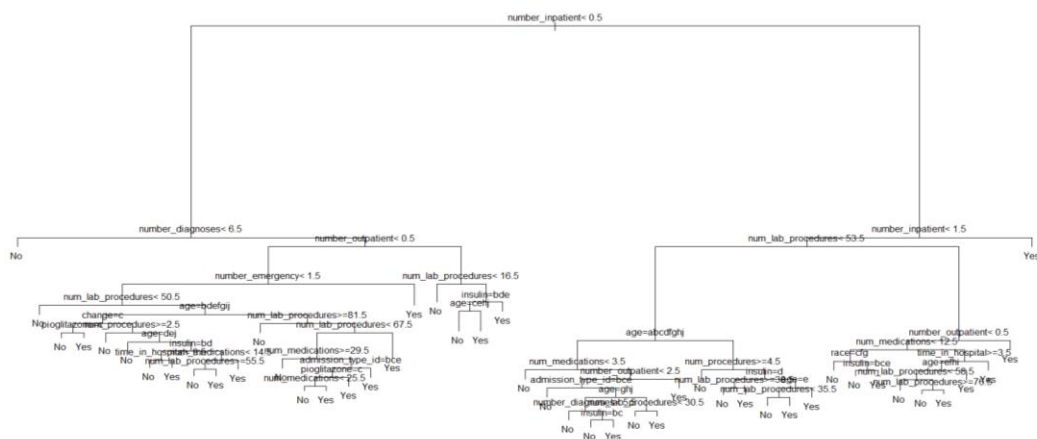
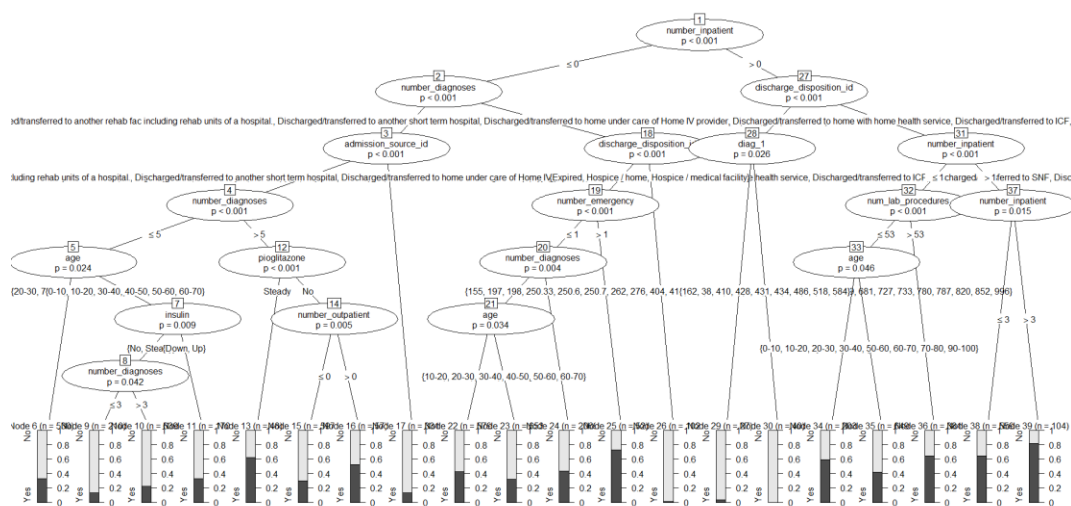
Data splitting used was 80% of data for training and 20% for testing. We also performed cross validation through KNN model, since it has built in function (traincontrol()) to do cross validation through building the model. For In the next section, we are going to explain each model in details.

### 2.9.1 Decision Tree:

Decision tree is one of the most common and simple classification algorithm. Decision tree uses specific attributes to split data and finally reach the final decision. So, it would use variables that have significant contribution to identify the label. We used three decision tree models in R:

- Tree function under tree package
- Ctree function under party package
- Rpart function under rpart package.

The accuracy and other parameters are same, but trees built were different. We have the following trees : 1- ctree model 2- rpart model



In addition, rpart model shows the importance of values used in the model. Here is a histogram that shows importance values of the attributes through building



## HOSPITAL READMISSION ANALYSIS

the tree.

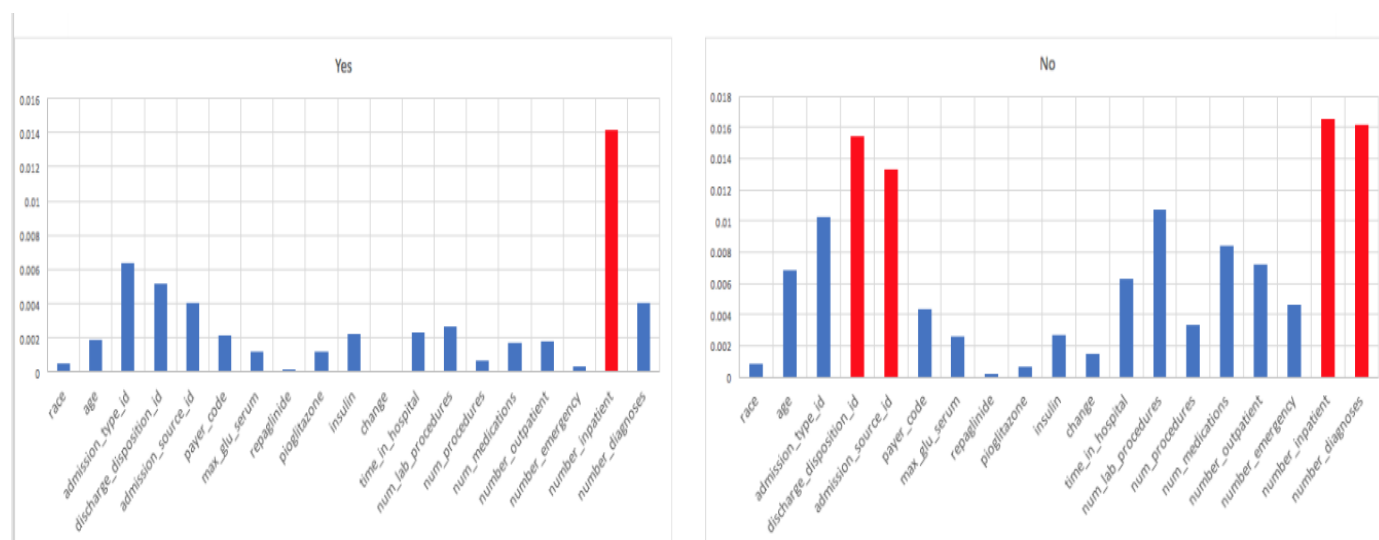
### 2.9.2 Random Forest:

In Random Forest, we used `randomForest` package and function build our model. We used 500 tree to be generated. We have the following result:

	No	Yes	MeanDecreaseAccuracy
race	0.0008203805381	0.0005078582110	0.00069820764057
age	0.0068196778544	-0.0019257768206	0.00336235185778
admission_type_id	0.0102117881901	-0.0063485931506	0.00366845320194
discharge_disposition_id	0.0153781618928	0.0051884862878	0.01134882100985
admission_source_id	0.0133083390921	-0.0040128602586	0.00647470050932
payer_code	0.0043160073784	-0.0021145496837	0.00177885965335
max_glu_serum	0.0025817475288	-0.0011858819010	0.00109318399397
repaglinide	0.0001943066917	-0.0001779874790	0.00004806117356
pioglitazone	0.0006898507979	0.0011626045612	0.00087434291725
insulin	0.0026939209814	-0.0022544108360	0.00073987735566
change	0.0014721113603	0.0000819060250	0.00091834062731
time_in_hospital	0.0062859387413	-0.0023394775733	0.00287620262069
num_lab_procedures	0.0106932091754	-0.0026821286106	0.00539172479252
num_procedures	0.0032938436840	-0.0006713071712	0.00172408692717
num_medications	0.0083757316157	-0.0017177005099	0.00439029492539
number_outpatient	0.0071667950047	0.0017591782052	0.00502700826698
number_emergency	0.0046643480445	-0.0002928679893	0.00270681849900
number_inpatient	0.0165101460698	0.0141871040026	0.01559349774107
number_diagnoses	0.0161997399412	0.0040099184158	0.01138715049516

We found some attributes that became significant comparing to others for both “yes” and “no” situation. For example, `number_inpatient` represented high contribution to ‘yes’ classification by 0.014 while rest of variable were between 0 and 0.006.

The following graph presented the contribution of each variable in both ‘yes’ and ‘no’ classification.



In addition, we used *inTrees* packages to extract and prune rules from *randomForest* model. Here is some rules extracted from our model.

len	fre	error	rules	predict
1	0.02 6	0.027	discharge_disposition_id %in% <ul style="list-style-type: none"> <li>Admitted as an inpatient to this hospital',</li> <li>'Discharged/transferred to a nursing facility certified under Medicaid but not certified under Medicare'</li> <li>'Discharged/transferred/referred another institution for outpatient services',</li> <li>'Discharged/transferred/referred to this institution for outpatient services',</li> <li>'Expired',</li> <li>Hospice / medical facility'</li> </ul>	No
1	0.40 5	0.387	age = 20-30, 50-60, 60-70	No
3	0.00 1	0.429	age = 60-70 admission_type_id = Emergency discharge_disposition_id %in% <ul style="list-style-type: none"> <li>Admitted as an inpatient to this hospital</li> <li>Discharged/transferred to a nursing facility certified under Medicaid but not certified under Medicare</li> <li>Discharged/transferred/referred to this institution for outpatient services</li> <li>Hospice / home</li> <li>Hospice / medical facility</li> </ul>	Yes

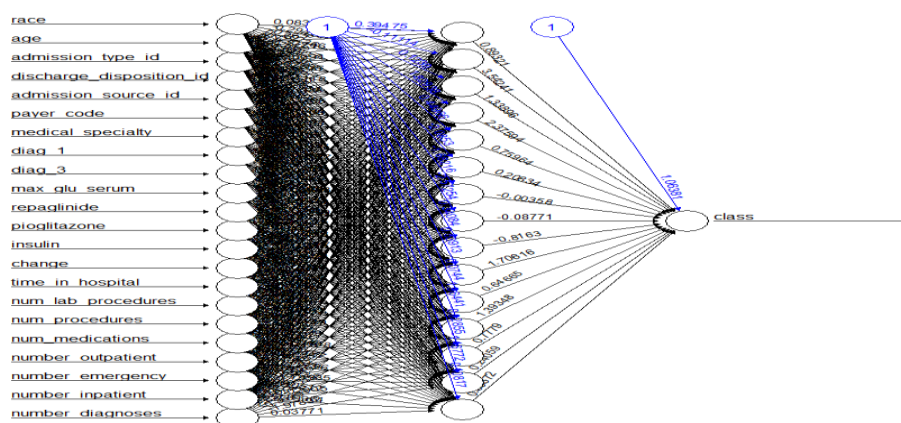
### 2.9.3 Neural Network

We used two models here just to compare between them. The first model uses multi-layer perceptron (MLP) algorithm in train function. Second model uses nnet function, and this model requires data to be scaled, centered and normalized. Here is the result of both model

nnet::nnet	<div> <p>Confusion Matrix and Statistics</p> <pre> Reference Prediction  1    2 1  1171    20 2   794    15  Accuracy : 0.593 95% CI : (0.5710959, 0.61 No Information Rate : 0.9825 P-Value [Acc &gt; NIR] : 1 </pre> </div>	caret::train	<div> <p>Confusion Matrix and Statistics</p> <pre> Reference Prediction No Yes No  1191    0 Yes  809    0  Accuracy : 0.5955 95% CI : (0.5736, 0.6171) No Information Rate : 1 P-Value [Acc &gt; NIR] : 1 </pre> </div>
------------	---	--------------	--

They have same accuracy, but MLP model faced some problem in calculating precision, recall and F1 since there is zero values for true negative(TN) and false negative(FN). We used *Neuralnet* package to do NN model and visualize it. Here is the plot diagram:

## HOSPITAL READMISSION ANALYSIS



### 2.9.4 KNN

Under caret package, we used train () function with knn method. We also used cross validation in training our model. We used auto tuning to identify the best k value that has the highest accuracy. Here is the result of our model

```
> pat_knn
k-Nearest Neighbors

8000 samples
 22 predictor
 2 classes: 'No', 'Yes'

No pre-processing
Resampling: Cross-Validated (10 fold)
Summary of sample sizes: 7199, 7201, 7199, 7200, 7201, 7199, ...
Resampling results across tuning parameters:
```

k	Accuracy	Kappa
5	0.5851236750	0.09088818120
7	0.5921228994	0.09713599938
9	0.5946224367	0.09564066758
11	0.6001204115	0.10154393027
13	0.6009972889	0.10151416487
15	0.6058721320	0.10809779363
17	0.6083747947	0.10986679431
19	0.6116265172	0.11452334964
21	0.6122476107	0.11327335808
23	0.6133719869	0.11324041265
25	0.6119982365	0.10787254509
27	0.6117457373	0.10506040089
29	0.6118698002	0.10403982495
31	0.6154952711	0.11011163538
33	0.6137447998	0.10593940019
35	0.6162448010	0.11006149104
37	0.6148718309	0.10661819938
39	0.6163719850	0.10853762168
41	0.6171257342	0.10898750447
43	0.6178737035	0.10930327988

Accuracy was used to select the optimal model using the largest value.  
The final value used for the model was k = 43.

The highest accuracy of our classification model achieved by using k= 43. We tuned our model by selecting length -20, so model tested 20 k values starting 5 and ended by 43. If we extended the length of tuning model, we may get better results than k = 43.

### 2.9.5 Evaluation our classification models:

Here is the table that comparing different classification algorithms:

Model	Accuracy	TP	TN	FP	FN	precision	recall	F-Meas s
Decision tree	0.638	213	1036	596	128	0.89	0.64	0.75
Random Forest	0.632	339	925	470	266	0.78	0.66	0.75
KNN	0.60	189	1016	175	620	0.62	.85	0.71
NN (train)	0.595	0	1191	809	0	NA	NA	NA
NN (nnet)	0.593	15	1171	794	20	0.98	0.60	0.74

Table shows that decision tree achieved the highest accuracy among other model while lowest

accuracy went for NN models. NN is significant in achieving true negative (TN) among other models, but very low in true positive (TP). This is obvious in both precision and recall values. F-measure showed that both decision tree and random forest higher than others. However, we still have kind of bias here because decision tree didn't include all attributes (variables with levels less than 53), while other model included all data fields.

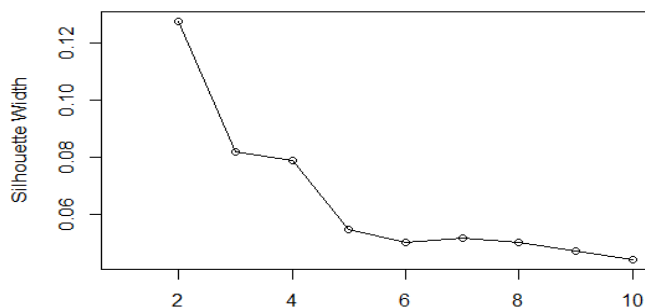
By seeing the results of models, we found some variables that become very important to assign accurate label. Number\_inpatient, for example, showed high importance in decision tree, used as the root of tree in both trees we built, and showed importance in classification using random forest model. Number\_emergancnt, discharge\_disposition\_id, diag\_1, diag\_3 represented importance in classification.

## 2.10 Clustering

It is an explanatory data analysis technique used for identifying groups in the data set. Each group contains similar profile per the set criteria. The clusters have high intra similarity while low inter similarity. We implemented clustering to cluster patients based on their previous diagnoses. Hospital readmission is not only based on present symptoms but also on the medical history of the patient.

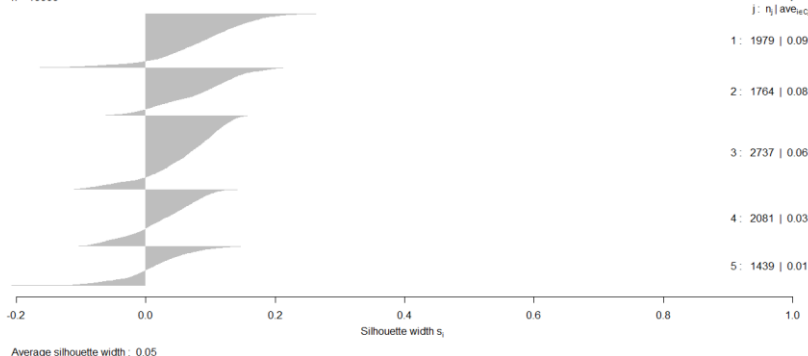
### 2.10.1 PAM

Since we had mixed data type, we implemented PAM (Partitioning Around Medoids) for clustering. This algorithm is based on  $k$  representative medoids among the dataset observations. The `pam()` function takes 2 inputs, dataset and the number of clusters( $k$ ). To find the optimal number of clusters we tried `silhouette()` function. This function is used to interpret and validate the consistency of the clusters.



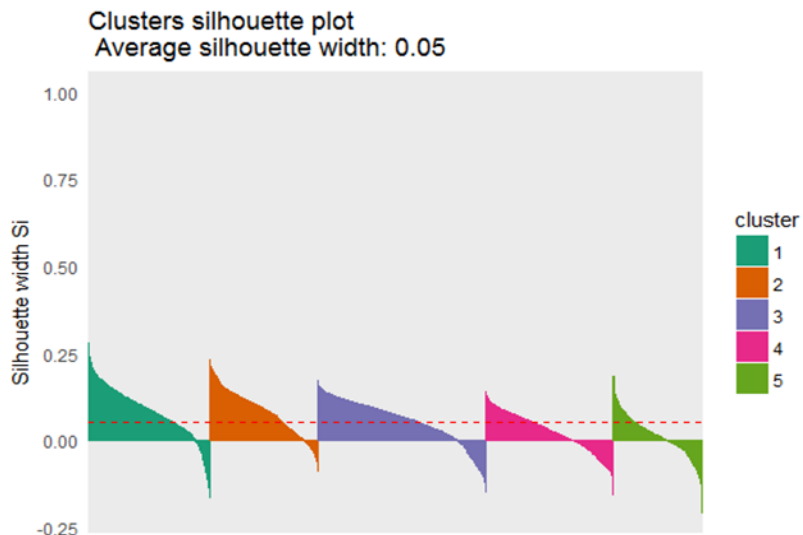
From this silhouette graph, we can see that the line tends to straighten out at 5. The silhouette plot shows the cluster width. Larger the width, better the clusters. So we chose  $k=5$  for our `pam()` function.

Silhouette plot of (x = res\$clustering, dist = gower\_dist)  
n = 10000



This plot shows 5 clusters, with total number of data items in each and the cluster width. Like cluster 1 has 1979 items with the width of 0.09.

## HOSPITAL READMISSION ANALYSIS



This graph is more colored visualization of the silhouette plot using “factoextra” package. The lines below zero denotes that some items are wrongly clustered.

We can find the members of each cluster using member() function:

```
> member.c
[1] 1 1 2 3 3 1 1 4 4 1 5 4 1 3 5 3 4 5 4 2 3 1 3 1 4 5 3 4 1 2 3 1 1 5 4 4 5 5 4 4 5
[60] 4 1 3 1 1 1 3 1 3 4 4 5 3 4 1 3 3 5 1 3 4 1 5 3 5 1 4 5 3 1 1 2 4 4 4 2 4 5 3 1 4
[119] 2 3 4 5 1 3 5 5 4 4 5 3 1 1 4 4 4 1 4 3 4 5 4 4 4 4 3 1 1 4 1 1 3 1 4 4 3 4 1 5 1
[178] 1 3 1 5 4 3 3 4 3 5 5 5 4 4 5 1 5 5 2 5 3 1 4 5 5 4 3 3 4 5 1 4 5 3 1 5 4 4 4 4 5
[237] 1 5 3 3 3 3 3 5 3 4 4 5 3 3 4 5 5 1 1 4 1 1 3 4 3 3 5 2 4 4 1 4 5 1 3 4 3 4 1 2 4
[296] 1 1 1 3 1 4 4 4 4 5 5 4 1 1 3 3 5 2 5 4 1 1 5 4 4 4 5 5 3 1 4 4 4 5 2 1 4 1 4 3 1
[355] 4 4 4 4 1 1 4 1 4 4 3 5 4 3 5 4 2 4 1 4 5 1 1 4 3 4 3 4 1 3 1 3 1 4 1 5 5 2 3 1 5
[414] 3 5 3 5 3 4 4 1 5 4 3 5 5 1 4 5 3 4 5 4 1 3 3 1 3 1 4 4 5 4 3 4 4 4 5 4 5 1 4 3 4
[473] 3 3 1 3 3 4 5 4 5 5 5 5 5 5 1 1 4 5 4 4 1 5 1 1 1 4 3 4 1 5 4 5 1 5 1 3 3 3 3 3 3
[532] 4 4 1 3 5 4 5 4 5 1 4 3 4 4 5 5 1 3 1 5 5 4 3 3 4 1 4 5 5 5 3 4 5 4 1 1 4 4 5 3 1
[591] 1 5 4 4 4 4 5 3 5 1 5 4 3 4 4 5 5 4 1 1 2 4 1 4 5 4 1 4 4 3 3 4 5 1 3 3 2 4 5 5 5
```

It shows that data item 1 belongs to cluster 1, item 8 belongs to cluster 4 and so on.

### Cluster 1:

```
> printout.table(summary)
[[1]]
      race      age      admission_type_id
AfricanAmerican: 436 60-70 :864 Elective :1531
Asian          : 15 70-80 :539 Emergency : 136
Caucasian      :2160 50-60 :508 Newborn   :  0
Hispanic       : 42 80-90 :289 Not Available: 226
Other          : 36 40-50 :272 Not Mapped  : 17
              : 30-40 :116 Urgent       : 759
              (other):101

      discharge_disposition_id
Discharged to home :1933
Discharged/transferred to home with home health service :301
Discharged/transferred to SNF :192
Discharged/transferred to another rehab fac including rehab units of a hospital.: 54
Discharged/transferred to another type of inpatient care institution : 53
Discharged/transferred to another short term hospital : 43
(Other) :113

      admission_source_id payer_code medical_specialty
Physician Referral :2206 MC :1740 Cardiology :791
Transfer from a hospital :193 BC :233 Family/GeneralPractice :279
Emergency Room :146 HM :202 Surgery-General :264
Clinic Referral : 80 SP :140 InternalMedicine :185
Transfer from another health care facility: 28 UN :124 Orthopedics-Reconstructive:145
HMO Referral : 16 MD : 91 Orthopedics :144
(Other) : 20 (Other):159 (Other) :881

      diag_1      diag_3      max_glu_serum      repaglinide      pioglitazone      insulin      change      time_in_hospital
414 : 513 250 : 596 >200: 35 Down : 3 Down : 0 Down :164 Ch: 913 Min. : 1.00
715 : 173 401 : 326 >300: 17 No :2663 No :2495 No :1723 No:1776 1st Qu.: 2.00
410 : 114 414 : 125 None:2545 Steady: 20 Steady: 190 Steady: 628 Median : 3.00
996 : 83 272 : 83 Norm: 92 Up : 3 Up : 4 Up :174 Mean : 3.86
427 : 79 427 : 73
722 : 72 428 : 59 Max. :14.00
(Other):1655 (Other):1427

      num_lab_procedures num_procedures num_medications number_outpatient number_emergency number_inpatient
Min. : 1.00 Min. :0.000 Min. : 1.00 Min. : 0.0000 Min. :0.00000 Min. :0.0000
1st Qu.: 20.00 1st Qu.:1.000 1st Qu.:11.00 1st Qu.: 0.0000 1st Qu.:0.00000 1st Qu.:0.0000
Median : 35.00 Median :2.000 Median :15.00 Median : 0.0000 Median :0.00000 Median :0.0000
Mean : 33.88 Mean :2.364 Mean :17.11 Mean : 0.3061 Mean :0.05727 Mean :0.2283
3rd Qu.: 46.00 3rd Qu.:3.000 3rd Qu.:21.00 3rd Qu.: 0.0000 3rd Qu.:0.00000 3rd Qu.:0.0000
Max. :114.00 Max. :6.000 Max. :81.00 Max. :36.0000 Max. :5.00000 Max. :7.0000
```



## HOSPITAL READMISSION ANALYSIS

## Cluster 2:

```
[[2]]
      race      age      admission_type_id
AfricanAmerican: 697 60-70 :904 Elective : 238
Asian : 14 70-80 :573 Emergency :2045
Caucasian :2324 30-60 :356 Newborn : 0
Hispanic : 55 80-90 :532 Not Available: 204
Other : 37 40-50 :303 Not Mapped : 14
      30-40 :127 Urgent : 626
      (Other):132

Discharged to home
Discharged/transferred to home with home health service
Discharged/transferred to SNF
Discharged/transferred to another rehab fac including rehab units of a hospital
Discharged/transferred to another short term hospital
Discharged/transferred to another type of inpatient care institution
(Other)

discharge_disposition_id
:1895
:457
:448
: 68
: 62
: 46
:151

admission_source_id payer_code medical_specialty
Emergency Room :2134 MC :2173 InternalMedicine :1650
Physician Referral : 670 HM : 207 Family/GeneralPractice: 467
Transfer from another health care facility : 115 BC : 188 Cardiology : 243
Transfer from a hospital : 104 SP : 141 Emergency/Trauma : 243
Clinic Referral : 58 MD : 114 Surgery-General : 88
Transfer from a Skilled Nursing Facility (SNF): 28 UN : 109 Nephrology : 59
(Other) : 18 (Other): 195 (Other) : 377

diag_1 diag_3 max_glu_serum repaglinide pioglitazone insulin change time_in_hospital
428 : 359 401 : 402 >200: 79 Down : 2 Down : 10 Down : 561 Ch:2504 Min. : 1.000
410 : 134 250 : 244 >300: 52 No :3061 No :2798 No : 462 No: 623 1st Qu.: 3.000
486 : 123 276 : 185 None:2913 Steady: 57 Steady: 300 Steady:1623 Mean : 5.000
414 : 110 427 : 144 Norm: 83 Up : 7 Up : 19 Up : 481 Mean : 5.321
682 : 91 428 : 93 3rd Qu.: 7.000
434 : 87 414 : 83 Max. :14.000
(Other):2223 (Other):1976

num_lab_procedures num_procedures num_medications number_outpatient number_emergency number_inpatient
Min. : 1.00 Min. :0.0000 Min. : 1.00 Min. : 0.0000 Min. : 0.0000 Min. : 0.0000
1st Qu.: 37.00 1st Qu.:0.0000 1st Qu.: 8.00 1st Qu.: 0.0000 1st Qu.: 0.0000 1st Qu.: 0.0000
Median : 48.00 Median :1.0000 Median :16.00 Median : 0.0000 Median : 0.0000 Median : 0.0000
Mean : 48.04 Mean :1.193 Mean :17.54 Mean : 0.3256 Mean : 0.1666 Mean : 0.5692
3rd Qu.: 61.50 3rd Qu.:2.000 3rd Qu.:22.00 3rd Qu.: 0.0000 3rd Qu.: 0.0000 3rd Qu.: 1.0000
Max. :113.00 Max. : 6.000 Max. :75.00 Max. :27.0000 Max. :42.0000 Max. :10.0000
```

## Cluster 3:

```
[[3]]
      race      age      admission_type_id
AfricanAmerican: 952 70-80 :1483 Elective : 106
Asian : 26 80-90 : 756 Emergency :3237
Caucasian :3074 30-60 : 658 Newborn : 1
Hispanic : 84 60-70 : 419 Not Available: 225
Other : 48 40-50 : 403 Not Mapped : 7
      30-40 : 174 Urgent : 608
      (Other): 291

Discharged to home
Discharged/transferred to SNF
Discharged/transferred to home with home health service
Expired
Discharged/transferred to another short term hospital
Not Mapped
(Other)

discharge_disposition_id
:2627
: 569
:445
:130
: 81
: 62
:270

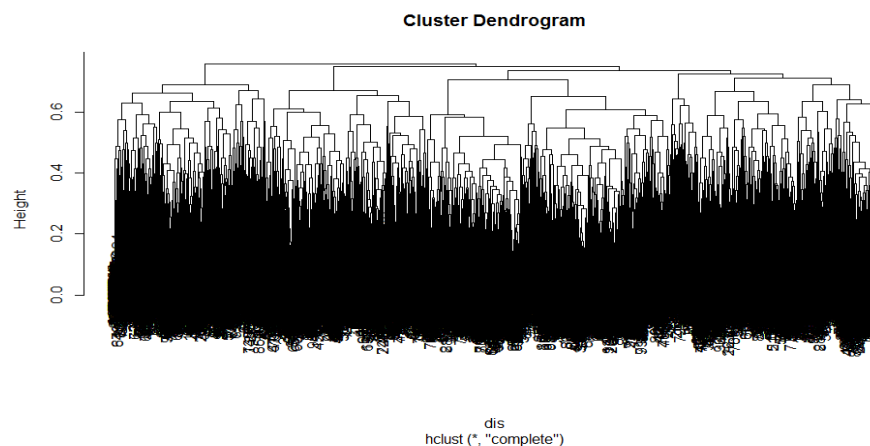
admission_source_id payer_code medical_specialty
Emergency Room :3259 MC :3006 InternalMedicine :2448
Physician Referral : 456 HM : 285 Family/GeneralPractice: 641
Transfer from another health care facility : 208 BC : 227 Emergency/Trauma : 313
Transfer from a hospital : 109 SP : 181 Cardiology : 267
Transfer from a Skilled Nursing Facility (SNF): 82 UN : 139 Surgery-General : 82
Clinic Referral : 44 MD : 118 Nephrology : 74
(Other) : 26 (Other):228 (Other): 359

diag_1 diag_3 max_glu_serum repaglinide pioglitazone insulin change time_in_hospital
786 : 346 250 : 487 >200: 83 Down : 0 Down : 3 Down : 217 Ch: 859 Min. : 1.000
428 : 218 401 : 314 >300: 57 No :4146 No :4001 No :2974 No:3325 1st Qu.: 2.000
486 : 168 428 : 264 None:3878 Steady: 35 Steady: 174 Steady: 806 Median : 3.000
410 : 155 276 : 232 Norm: 166 Up : 3 Up : 6 Up : 187 Mean : 4.141
427 : 123 427 : 176 3rd Qu.: 5.000
434 : 123 414 : 157 Max. :14.000
(Other):3051 (Other):2554

num_lab_procedures num_procedures num_medications number_outpatient number_emergency number_inpatient
Min. : 1.00 Min. :0.0000 Min. : 1.00 Min. : 0.0000 Min. : 0.0000 Min. : 0.0000
1st Qu.: 36.00 1st Qu.:0.0000 1st Qu.: 8.00 1st Qu.: 0.0000 1st Qu.: 0.0000 1st Qu.: 0.0000
Median : 46.00 Median :0.0000 Median :12.00 Median : 0.0000 Median : 0.0000 Median : 0.0000
Mean : 45.29 Mean :0.9336 Mean :13.09 Mean : 0.2333 Mean : 0.1135 Mean : 0.3535
3rd Qu.: 58.00 3rd Qu.:1.0000 3rd Qu.:17.00 3rd Qu.: 0.0000 3rd Qu.: 0.0000 3rd Qu.: 0.0000
Max. :120.00 Max. : 6.0000 Max. :60.00 Max. :21.0000 Max. : 9.0000 Max. : 9.0000
```

From here we found that cluster 1 has the patients who had previous history of circulatory and connective tissue problems, were readmitted on physician's referral and having max\_glu\_serum not measured earlier.. Cluster 2 represents patients who have cardiac problems in the past and were admitted at emergency and having stable insulin. Cluster 3 represents patients having respiratory problems with the age of 70-80 and max\_glu\_serum>200.

## 2.10.2 Hierarchical Clustering:



We also tried to implement hclust() from cluster package to plot dendrograms.

hclust() function was

implemented using complete linkage and average linkage. Complete linkage method uses maximum distance between the data clusters. While average linkage method uses the mean of the distance between the clusters. Members.c represents members of the clusters using complete linkage method, while members.a represents average linkage method.

```
> table(member.c, member.a)
```

member.c	member.a				
	1	2	3	4	5
1	2329	3	1	0	3
2	362	0	0	0	0
3	2115	1	1	0	0
4	3203	0	0	0	0
5	1970	7	3	2	0

This table shows that 2329 items belong to cluster 1, while 7 items were wrongly predicted for cluster 1.

### 3. Findings and Analysis

We have some findings through our analysis. We have focused on significant variables through our results in classification, which are: number\_inpatient, number\_diagnoses, ICD 9 diag\_1, ICD 9 diag\_3, and Discharge disposition\_id. By checking these important variables, we found some patterns. We found that if number\_inpatient > 7.5, label will be 'Yes'. We also find that if number\_emergency more than 8.5, the class will be 'yes'. However, we found many similarities between many variables by filtering both 'Yes' and 'No' observation.

We also examine ICD 9 codes frequency for both 'Yes' and 'No' observations. We found that code 428 is most frequent in Yes observation, while 414 for No observation. 428 is about heart attack and there are researches confirmed the relationship between diabetes and heart attack risk [5]. In our data, 428 has 8% of readmitted cases (342 out of 3965) and 4% of not readmitted cases (294 out of 6035). Diabetes, like other chronic medical condition, is associated with increased risk of hospital readmission. Efforts to reduce readmission should be multifactorial and encompass both general and specific diabetic measures. We found certain factors played important role in readmission analysis.

**4. References:**

- <https://www.analyticsvidhya.com/blog/2016/03/select-important-variables-boruta-package/>
- <http://machinelearningmastery.com/feature-selection-with-the-caret-r-package/>
- <http://www.sthda.com/english/wiki/partitioning-cluster-analysis-quick-start-guide-unsupervised-machine-learning>
- <http://www.sthda.com/english/wiki/cluster-analysis-in-r-unsupervised-machine-learning>
- <https://cran.r-project.org/web/packages/dendextend/vignettes/introduction.html>
- [http://www.joslin.org/info/diabetes and heart disease an intimate connection.html](http://www.joslin.org/info/diabetes_and_heart_disease_an_intimate_connection.html)



## 5. Appendices:

### R Code:

```
setwd("/Users/Imani/Desktop/UNCC/KDD/Final Project")
#converting all empty and ? to NA
diab_data<- read.csv("10kDiabetes.csv", header = TRUE, na.strings = c("", "?", "NA"))
head(diab_data)
str(diab_data)
```

#### #Part 1

```
#removing weight, since they are too sparse to process
diab_data[["weight"]]<-NULL
```

#### #Part 2

#### #CORR RCORR

```
diab_cordata <- subset(diab_data, select =
c("time_in_hospital", "num_lab_procedures", "num_procedures", "num_medications", "number_outpatient", "number_emergency", "number_inpatient", "number_diagnoses", "readmitted"))
cor_data<-
cor(diab_data$time_in_hospital, diab_data$num_lab_procedures, use="complete.obs", method="pearson")
cor_data
```

```
cor_cordata<-cor(diab_cordata, use="complete.obs", method="pearson")
cor_cordata
```

#### #rcorr

```
install.packages("Hmisc", dependencies = TRUE)
library(Hmisc)
tnlp<-rcorr(diab_data$time_in_hospital, diab_data$num_lab_procedures, type = "pearson")
tnlp
```

```
install.packages("plyr", dependencies = TRUE)
library('plyr')
count(diab_data, 'race')
count(diab_data, 'gender')
```

#### #chi-square analysis

```
counts1 <- ddply(diab_data,.(diab_data$race, diab_data$readmitted), nrow)
counts1
```

## HOSPITAL READMISSION ANALYSIS

```

names(counts1)<-c("Race","Readmitted","Freq")

african<-c(725,1361)
caucasian<-c(3115,4442)
asian<-c(19,36)
hispanic<-c(65,116)
other<-c(41,80)

race_readmitted <- cbind(african,caucasian,asian,hispanic,other)
row.names(race_readmitted) = c("TRUE","FALSE")

install.packages("gmodels")
library(gmodels)
CrossTable(race_readmitted, chisq = TRUE, expected = TRUE, sresid = TRUE, format =
"SPSS")

install.packages("mlogit")
library(mlogit)

model1 <- glm(diab_data$readmitted ~
diab_data$number_inpatient+diab_data$number_outpatient+diab_data$num_lab_procedures,data
a = diab_data, family = binomial())
model2 <- glm(diab_data$readmitted ~
diab_data$number_inpatient+diab_data$number_outpatient+diab_data$number_diagnoses,data
= diab_data, family = binomial())
summary(model1)
summary(model2)

anova(model1,model2)

#FEATURE SELECTION
library(mlbench)

# feature selection using random forest
control2 <- rfeControl(functions=rfFuncs, method="cv", number=10)
# run the RFE algorithm
results <- rfe(patient[,1:32], patient[,33], sizes=c(1:32), rfeControl=control2)
# summarize the results
print(results)
# list the chosen features

```

## HOSPITAL READMISSION ANALYSIS

```

predictors(results)
# plot the results
plot(results)

```

### # feature selection using Boruta

```

install.packages("Boruta")
library(Boruta)
patient_boruta <- Boruta(readmitted~., data = patient, doTrace = 2)
final_feature<- TentativeRoughFix(patient_boruta)
getSelectedAttributes(final_feature, withTentative = F) # shows selected attributes

```

### #CLASSIFICATION

```
#####Training data #####
```

```

pat_ind <- sample(1:nrow(patient),0.8*nrow(patient))
train <- new_pat[pat_ind,]
test <- new_pat[-pat_ind,]

```

```
#####Decision Tree #####
```

```

# decision tree using party package
ctree <- party::ctree(class~.,train)
plot(tree)

```

```
#decision tree using rpart package
```

```

tree_rp <- rpart::rpart(class~.,train, method = "class", minsplit = 1, minbucket = 1, cp = 0.001)
#decision tree using tree package:
tree <- tree::tree(class~.,train)

```

```
#####Random Forest #####
```

```

rf <- randomForest(class~., data = train[,c(1:6,10:23)], importance=TRUE, ntree=500)
library(inTrees)
# extract rules from rf model
rf_rules <- extractRules(RF2List(pat_rf),new_pat[,c(1:6,10:22)])

```

## HOSPITAL READMISSION ANALYSIS

```
##### KNN #####
```

```
control <- trainControl(method="repeatedcv", number=10, repeats=3) # 10 fold cross validation
control_knn <- trainControl(method="cv", number=10) # 10 fold cross validation
clus<-makeCluster(spec=8,type="PSOCK")
registerDoParallel(clus)
pat_knn <- train(class ~ ., data = train, method = "knn", trControl=control_knn,tuneLength = 20)
stopCluster(clus)
```

```
##### Neural Network #####
```

```
# using caret package
clus<-makeCluster(spec=8,type="PSOCK")
registerDoParallel(clus)
pat_nn <- train(class ~ ., data = train, method="mlp", metric='Accuracy', tuneGrid=expand.grid(.size=1:15))
stopCluster(clus)
```

```
#another NN models
```

```
# scale
```

```
preprocessParams <- preProcess(tn[,1:22], method=c("scale"))
tn2 <- predict(preprocessParams, tn[,1:22])
tn2$class <- new_pat$readmitted
```

```
# normalize
```

```
preprocessParams2 <- preProcess(tn[,1:22], method=c("range"))
tn3 <- predict(preprocessParams2, tn[,1:22])
tn3$class <- tn$readmitted
```

```
tn_train <- tn3[pat_ind,]
```

```
tn_test <- tn3[-pat_ind,]
```

```
#build another NN package
```

```
library(neuralnet)
```

```
n <- names(tn_train)
```

```
f <- as.formula(paste('class ~', paste(n[!n %in% "class"], collapse = " + ")))
```

```
nnnn <- neuralnet(f,data=tn_train,hidden=15,linear.output=FALSE)
```

## HOSPITAL READMISSION ANALYSIS

```
xnnx <- compute(nnnn,tn_test[1:22])
xnnx
plot(nnnn)
```

```
# another NN package
library(nnet)
ideal <- class.ind(tn$readmitted)
ANN = nnet(tn[pat_ind,1:22], ideal[pat_ind,], size=10)
nnp <- predict(ANN, tn[-pat_ind,1:22], type="class")
caret::confusionMatrix(tn_test$class,nnp)
```

```
##### valdiating models #####
```

```
#valdiate tree
val_tree <- predict(tree, test)
table(val == test[,23])
```

```
val_rp <- predict(tree_rp, test, type= "class")
table(val_rp == test[,23])
```

```
val_ctree <- predict(ctree, test)
table(val_ctree == test[,23])
```

```
val_knn2 <- predict(pat_knn2, test)
```

```
# validate knn
val_knn <- predict(pat_knn, test)
table(val_knn == test[,23])
```

```
val_knn2 <- predict(pat_knn2, test)
table(val_knn2 == test[,23])
```

```
# validate random forest
val_rf <- predict(pat_rf, test)
table(val_rf == test[,23])
```

```
# validate NN
```

## HOSPITAL READMISSION ANALYSIS

```
val_nn <- predict(pat_nn, test)
val_nnnn <- predict(nnnn, test2)
table(val_nn == test[,23])
```

#CLUSTERING

```
patient_data<- read.csv("patient.csv")
str(patient_data)
patient_data[["X"]]<-NULL
patient_data[["payer_code"]]<-NULL
library(cluster)
#scatter plot
plot(time_in_hospital~ diag_1, patient_data)
?daisy
dis<- daisy(patient_data, metric = "gower")
print(dis, digits = 3)
head(dis)

#complete linkage
hc.c<-hclust(dis)
plot(hc.c,hang = -1, cex = 0.6)

#average linkage
hc.a<- hclust(dis, method = "average")

#cluster membership
member.c<-cutree(hc.c, 5)
member.c

member.a<- cutree(hc.a,5)
table(member.c, member.a)

#cluster means
aggregate(patient_data, list(member.c), mean)

#silhoutte plot
library(cluster)
windows()
plot(silhouette(cutree(hc.c,5), dis))
```

## HOSPITAL READMISSION ANALYSIS

```
table(member.a, patient_data$readmitted)
```

```
#dendextend
```

```
library(dendextend)
```

```
dend<-as.dendrogram(hc.c)
```

```
d1=color_branches(dend,k=5)
```

```
plot(d1)
```

```
plot(hc.c, type = "phylogram", show.tip.label = TRUE,
```

```
  edge.color = "red", edge.width = 1, edge.lty = 1,
```

```
  tip.color = "blue")
```

```
install.packages("devtools")
```

```
library(devtools)
```

```
devtools::install_github("kassambara/factoextra")
```

```
library(factoextra)
```

```
#clustering
```

```
clust_data<- patient_data
```

```
head(patient_num_data)
```

```
medians<- apply(clust_data,2,median)
```

```
mads<- apply(clust_data,2,mad)
```

```
clust_data = scale(clust_data,center=medians,scale=mads)
```

```
patient.dist<-dist(clust_data)
```

```
#hclust uses complete linkage
```

```
patient.clust<-hclust(patient.dist)
```

```
plot(patient.clust,labels=patient_data$readmitted,main='Default from hclust')
```

```
#dendextend package
```

```
install.packages("dendextend")
```

```
install.packages("colorspace")
```

```
library(dendextend)
```

```
library(colorspace)
```

```
k <- 4
```

```
cols <- rainbow_hcl(k)
```

## HOSPITAL READMISSION ANALYSIS

```

dend <- as.dendrogram(patient.clust)
dend <- color_branches(dend, k = k)
plot(dend)
labels_dend <- labels(dend)
groups <- cutree(dend, k=4, order_clusters_as_data = FALSE)
dends <- list()
for(i in 1:k) {
  labels_to_keep <- labels_dend[i != groups]
  dends[[i]] <- prune(dend, labels_to_keep)
}
par(mfrow = c(2,2))
for(i in 1:k) {
  plot(dends[[i]],
       main = paste0("Tree number ", i))
}
groups.12 = cutree(patient.clust,12)
table(groups.12)

#Try clustering
library(dplyr)
library(cluster)
library(ggplot2)

?daisy

gower_dist<-daisy(patient_data, metric = "gower")
summary(gower_dist)
gower_mat <- as.matrix(gower_dist)
#find most similar data pairs
patient_data[which(gower_mat == min(gower_mat[gower_mat != min(gower_mat)]),arr.ind =
TRUE)[1, ], ]

sil_width <- c(NA)
for(i in 2:10){

  pam_fit <- pam(gower_dist,
                diss = TRUE,
                k = i)

  sil_width[i] <- pam_fit$silinfo$avg.width

```



```
}
```

```
#Silhouette analysis measures how well an observation is clustered and  
#it estimates the average distance between clusters. The silhouette plot displays a  
#measure of how close each point in one cluster is to points in the neighboring clusters.  
#plot sil width, higher the better
```

```
plot(1:10, sil_width,  
     xlab = "Number of clusters",  
     ylab = "Silhouette Width")  
lines(1:10, sil_width)
```