

Heart Disease (Logistic and K-NN)

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Logistic regression & K-Nearest Neighbors.

Performing classification analysis on a data combining 5 popular heart disease datasets already available independently but not combined before. In this dataset, 5 heart datasets are combined over 11 common features which makes it the largest heart disease dataset available so far for research purposes.

The five datasets used for its curation are: Cleveland, Hungarian, Switzerland, Long Beach VA, and Statlog (Heart) Data Set.

```
#rm(List = ls())
setwd("C:/Users/micah/OneDrive/Documents/R/Heart disease")
data1 <- read.csv("~/R/Heart
disease/heart_statlog_cleveland_hungary_final.csv")
```

```
head(data1)
```

```
##   age sex chest.pain.type resting.bp.s cholesterol fasting.blood.sugar
## 1  40  1             2         140          289                0
## 2  49  0             3         160          180                0
## 3  37  1             2         130          283                0
## 4  48  0             4         138          214                0
## 5  54  1             3         150          195                0
## 6  39  1             3         120          339                0
##   resting.ecg max.heart.rate exercise.angina oldpeak ST.slope target
## 1           0          172              0      0.0        1         0
## 2           0          156              0      1.0        2         1
## 3           1           98              0      0.0        1         0
```

```
## 4      0      108      1      1.5      2      1
## 5      0      122      0      0.0      1      0
## 6      0      170      0      0.0      1      0
```

```
dim(data1)
```

```
## [1] 1190  12
```

```
sum(is.na(data1))
```

```
## [1] 0
```

The data contains 1190 observations with 12 variables. It should also be noted that the data has no missing values.

Investigate correlation among predictors.

```
correlation_matrix <- cor(data1)
```

```
library(GGally)
```

```
## Warning: package 'GGally' was built under R version 4.3.1
```

```
## Loading required package: ggplot2
```

```
## Warning: package 'ggplot2' was built under R version 4.3.1
```

```
## Registered S3 method overwritten by 'GGally':
```

```
##   method from
```

```
##   +.gg    ggplot2
```

```
library(ggplot2)
```

```
ggcorr(data1, label = TRUE, label_alpha = .7)
```



Based off the correlation matrix plot, note the high correlation between ST.slope and old peak, exercise angina, max heart rate. Also note the high correlation between maximum heart rate and chest pain type.

Data types

Change the data types of categorical variables such as sex from numerical to factor variables and label categories.

```
table(data1$sex)

##
##  0  1
## 281 909

class(data1$sex)

## [1] "integer"

data1$sex <- factor(data1$sex,
                    levels=c(0,1),
                    labels = c("Female","Male"))

table(data1$sex)

##
## Female  Male
##   281    909
```

```

#table(data1$chest.pain.type)
data1$chest.pain <- factor(data1$chest.pain.type,
                           levels=c(1,2,3,4),
                           labels=c("typical angina","atypical angina",
                                     "non-anginal pain","asymptomatic"))

table(data1$chest.pain)

##
##   typical angina  atypical angina non-anginal pain    asymptomatic
##              66              216              283              625

#table(data1$fasting.blood.sugar)
data1$fasting.sugar <- factor(data1$fasting.blood.sugar,
                              levels=c(0,1),
                              labels=c("False", "True"))

table(data1$fasting.sugar)

##
## False  True
##   936   254

data1$resting.ecg <- factor(data1$resting.ecg,
                            levels=c(0,1,2),
                            labels=c("Normal","ST-T abnormality",
                                      "Left ventricular hypertrophy"))

table(data1$resting.ecg)

##
##              Normal              ST-T abnormality
##              684              181
## Left ventricular hypertrophy
##              325

data1$exercise.angina <- factor(data1$exercise.angina,
                                levels=c(0,1),
                                labels=c("No", "Yes"))

table(data1$exercise.angina)

##
## No Yes
## 729 461

data1$ST.slope[data1$ST.slope == 0] <- 1
data1$ST.slope <- factor(data1$ST.slope,
                        levels=c(1,2,3),
                        labels=c("upsloping","flat","downsloping"))

table(data1$ST.slope)

##
##   upsloping      flat downsloping
##       527       582       81

```

```
data1$target <- factor(data1$target,
                        levels=c(0,1),
                        labels=c("No disease","Heart disease"))
table(data1$target)

##
##      No disease Heart disease
##           561           629
```

Descriptive Statistics

#Continuous variables

```
attach(data1)

par(mfrow=c(3,2))
hist(age)
hist(resting.bp.s, main="Histogram of resting blood pressure")
hist(cholesterol)
hist(max.heart.rate, main="Histogram of maximum heart rate achieved")
hist(oldpeak)

#Calculate means, medians, standard deviation and IQR
library(dplyr)

## Warning: package 'dplyr' was built under R version 4.3.1

##
## Attaching package: 'dplyr'

## The following objects are masked from 'package:stats':
##
##      filter, lag

## The following objects are masked from 'package:base':
##
##      intersect, setdiff, setequal, union

descriptive <- data1 %>%
  select(age,resting.bp.s,cholesterol,max.heart.rate,oldpeak) %>%
  summarise_all(list(min,max,mean,sd,median,IQR))

print(descriptive)

##   age_fn1 resting.bp.s_fn1 cholesterol_fn1 max.heart.rate_fn1 oldpeak_fn1
## 1      28                0                0                60         -2.6
##   age_fn2 resting.bp.s_fn2 cholesterol_fn2 max.heart.rate_fn2 oldpeak_fn2
## 1      77             200             603             202          6.2
##   age_fn3 resting.bp.s_fn3 cholesterol_fn3 max.heart.rate_fn3 oldpeak_fn3
## 1 53.72017       132.1538       210.3639       139.7328    0.9227731
##   age_fn4 resting.bp.s_fn4 cholesterol_fn4 max.heart.rate_fn4 oldpeak_fn4
## 1 9.358203       18.36882       101.4205       25.51764    1.086337
```

```
## age_fn5 resting.bp.s_fn5 cholesterol_fn5 max.heart.rate_fn5 oldpeak_fn5
## 1      54              130             229             140.5         0.6
## age_fn6 resting.bp.s_fn6 cholesterol_fn6 max.heart.rate_fn6 oldpeak_fn6
## 1      13              20             81.75            39          1.6
```



```
#Factor variables
#Calculate proportions for multiple variables
combined_proportions <- data1 %>%
  tidyr::gather(key = "variable", value = "value", sex, chest.pain,
fasting.sugar,resting.ecg, exercise.angina,
                ST.slope,target) %>%
  group_by(variable, value) %>%
  summarize(count = n()) %>%
  group_by(variable) %>%
  mutate(proportion = count / sum(count))

## Warning: attributes are not identical across measure variables; they will
be
## dropped

## `summarise()` has grouped output by 'variable'. You can override using the
## `.groups` argument.

print(combined_proportions)

## # A tibble: 18 × 4
## # Groups:   variable [7]
##   variable      value count proportion
```

##	<chr>	<chr>	<int>	<dbl>
## 1	ST.slope	downsloping	81	0.0681
## 2	ST.slope	flat	582	0.489
## 3	ST.slope	upsloping	527	0.443
## 4	chest.pain	asymptomatic	625	0.525
## 5	chest.pain	atypical angina	216	0.182
## 6	chest.pain	non-anginal pain	283	0.238
## 7	chest.pain	typical angina	66	0.0555
## 8	exercise.angina	No	729	0.613
## 9	exercise.angina	Yes	461	0.387
## 10	fasting.sugar	False	936	0.787
## 11	fasting.sugar	True	254	0.213
## 12	resting.ecg	Left ventricular hypertrophy	325	0.273
## 13	resting.ecg	Normal	684	0.575
## 14	resting.ecg	ST-T abnormality	181	0.152
## 15	sex	Female	281	0.236
## 16	sex	Male	909	0.764
## 17	target	Heart disease	629	0.529
## 18	target	No disease	561	0.471

Logistic Regression

```
logit1 <-
glm(target~sex+chest.pain+fasting.sugar+resting.ecg+exercise.angina+ age+
resting.bp.s+cholesterol+max.heart.rate+oldpeak,data=data1,family = binomial)

summary(logit1)
```

```
##
## Call:
## glm(formula = target ~ sex + chest.pain + fasting.sugar + resting.ecg +
##     exercise.angina + age + resting.bp.s + cholesterol + max.heart.rate +
##     oldpeak, family = binomial, data = data1)
##
## Coefficients:
##
##               Estimate Std. Error z value
Pr(>|z|)
## (Intercept)      -1.5840687   1.1012453  -1.438
0.1503
## sexMale           1.3100306   0.2083787   6.287
3.24e-10
## chest.painatypical angina      -0.2692564   0.3805399  -0.708
0.4792
## chest.painnon-anginal pain      0.0126788   0.3396094   0.037
0.9702
## chest.painasymptomatic      1.6176821   0.3317116   4.877
1.08e-06
## fasting.sugarTrue      0.9337392   0.2141872   4.359
1.30e-05
## resting.ecgST-T abnormality     -0.1022413   0.2546271  -0.402
```

```

0.6880
## resting.ecgLeft ventricular hypertrophy 0.3008586 0.1969282 1.528
0.1266
## exercise.anginaYes 1.1486696 0.1873302 6.132
8.69e-10
## age 0.0171370 0.0101248 1.693
0.0905
## resting.bp.s 0.0053326 0.0047236 1.129
0.2589
## cholesterol -0.0029118 0.0009465 -3.076
0.0021
## max.heart.rate -0.0164854 0.0038229 -4.312
1.62e-05
## oldpeak 0.6419186 0.0894538 7.176
7.18e-13
##
## (Intercept)
## sexMale ***
## chest.painatypical angina
## chest.painnon-anginal pain
## chest.painasymptomatic ***
## fasting.sugarTrue ***
## resting.ecgST-T abnormality
## resting.ecgLeft ventricular hypertrophy
## exercise.anginaYes ***
## age .
## resting.bp.s
## cholesterol **
## max.heart.rate ***
## oldpeak ***
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
##
## (Dispersion parameter for binomial family taken to be 1)
##
## Null deviance: 1645.80 on 1189 degrees of freedom
## Residual deviance: 948.59 on 1176 degrees of freedom
## AIC: 976.59
##
## Number of Fisher Scoring iterations: 5

logit2 <- glm(target~sex+chest.pain+fasting.sugar+exercise.angina+age
              +cholesterol+max.heart.rate+oldpeak,data=data1,family =
binomial)
summary(logit2)

##
## Call:
## glm(formula = target ~ sex + chest.pain + fasting.sugar + exercise.angina
+

```



```
##      age + cholesterol + max.heart.rate + oldpeak, family = binomial,
##      data = data1)
##
## Coefficients:
##              Estimate Std. Error z value Pr(>|z|)
## (Intercept)    -1.2677854   0.9610527  -1.319   0.1871
## sexMale         1.2898138   0.2071517   6.226 4.77e-10 ***
## chest.painatypical angina -0.3557215   0.3779937  -0.941   0.3467
## chest.painnon-anginal pain -0.0476018   0.3381876  -0.141   0.8881
## chest.painasymptomatic    1.5529701   0.3295448   4.712 2.45e-06 ***
## fasting.sugarTrue    0.9424443   0.2127570   4.430 9.44e-06 ***
## exercise.anginaYes    1.1484854   0.1854946   6.191 5.96e-10 ***
## age              0.0221528   0.0097354   2.275   0.0229 *
## cholesterol       -0.0024700   0.0009143  -2.701   0.0069 **
## max.heart.rate     -0.0153385   0.0037406  -4.101 4.12e-05 ***
## oldpeak           0.6511051   0.0886588   7.344 2.07e-13 ***
## ---
## Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
##
## (Dispersion parameter for binomial family taken to be 1)
##
##      Null deviance: 1645.80  on 1189  degrees of freedom
## Residual deviance:  952.53  on 1179  degrees of freedom
## AIC: 974.53
##
## Number of Fisher Scoring iterations: 5
```

The `anova()` function can also be used to compare nested logistic regression models to determine if adding additional predictors significantly improves the model fit.

```
anova(logit2, logit1, test = "Chisq")

## Analysis of Deviance Table
##
## Model 1: target ~ sex + chest.pain + fasting.sugar + exercise.angina +
##      age + cholesterol + max.heart.rate + oldpeak
## Model 2: target ~ sex + chest.pain + fasting.sugar + resting.ecg +
##      exercise.angina +
##      age + resting.bp.s + cholesterol + max.heart.rate + oldpeak
##   Resid. Df Resid. Dev Df Deviance Pr(>Chi)
## 1      1179      952.53
## 2      1176      948.59  3    3.9419   0.2678
```

P-value > 0.05, therefore, the complex model does not significantly improve the fit compared to the simpler model.

```
# Get the coefficients of the model
coefficients <- summary(logit2)$coefficients

# Transform the coefficients to odds ratios
```

```
odds_ratios <- exp(coefficients[, "Estimate"])
odds_ratios

##                (Intercept)                sexMale
##                0.2814542                3.6321101
## chest.painatypical angina chest.painnon-anginal pain
##                0.7006677                0.9535134
## chest.painasymptomatic                fasting.sugarTrue
##                4.7254846                2.5662465
## exercise.anginaYes                age
##                3.1534132                1.0224000
## cholesterol                max.heart.rate
##                0.9975331                0.9847785
## oldpeak
##                1.9176589
```

Confidence intervals

```
confidence_intervals <- exp(confint(logit2))
```

Waiting for profiling to be done...

```
confidence_intervals
```

```
##                2.5 %    97.5 %
## (Intercept)    0.04230963 1.8391194
## sexMale        2.43336409 5.4864693
## chest.painatypical angina 0.33442881 1.4768618
## chest.painnon-anginal pain 0.49392466 1.8655904
## chest.painasymptomatic 2.49668065 9.1156647
## fasting.sugarTrue 1.69851791 3.9142978
## exercise.anginaYes 2.19576124 4.5468569
## age            1.00313139 1.0421973
## cholesterol    0.99572405 0.9993036
## max.heart.rate 0.97753338 0.9919863
## oldpeak        1.61682066 2.2895689
```

Split into train and test data

```
set.seed(2)
```

```
train_indices <- sample(seq_len(nrow(data1)), size = 0.7*nrow(data1))
```

```
train <- data1[train_indices,]
```

```
test <- data1[-train_indices,]
```

```
glm.fit <- glm(target~sex+chest.pain+fasting.sugar+exercise.angina+age
               +cholesterol+max.heart.rate+oldpeak, data=train, family =
binomial)
```

```
glm.prob =predict(glm.fit, test, type="response")
```

#Compute the predictions using test data.

```
glm.pred=rep("No", 357)
```

#Probability above 0.5 is predicted as Up

```

glm.pred[glm.prob>.50]="Yes"
table(glm.pred,test$target)

##
## glm.pred No disease Heart disease
##      No      139      25
##      Yes      37     156

#correctly predicting heart disease / no heart disease (83%)
(139+156)/357

## [1] 0.8263305

#prediction error (17%)
1 - (139+156)/357

## [1] 0.1736695

```

Using the trained model, the probability of predicting correctly (heart disease / no heart disease) is 0.83. Therefore, the prediction error is approximately 17%.

K-Nearest Neighbors

```

library(class)
train.x <-
cbind(train$sex,train$chest.pain,train$fasting.sugar,train$exercise.angina,

train$age,train$cholesterol,train$max.heart.rate,train$oldpeak)
test.x <-
cbind(test$sex,test$chest.pain,test$fasting.sugar,test$exercise.angina,
      test$age,test$cholesterol,test$max.heart.rate,test$oldpeak)
train.heart <- train$target

#Prediction accuracy with k = 1
set.seed(1)
knn.pred<-knn(train.x,test.x,train.heart,k=1)
table(knn.pred,test$target)

##
## knn.pred      No disease Heart disease
## No disease      130      35
## Heart disease      46     146

mean(knn.pred == test$target)

## [1] 0.7731092

```

77% highest K-NN prediction accuracy.

Therefore, the Logistic Model (83%) predicts better than the non-parametric K-Nearest Neighbor model (77%).