# Modeling

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#### **Imports**

```
library(dplyr)
library(ROCR)
```

#### Load Data

```
train <- readRDS(file = "../data/PIMA_train.Rds")
test <- readRDS(file = "../data/PIMA_test.Rds")</pre>
```

#### Baseline

```
tally(~ hasDiabetes, data = train)

## hasDiabetes
## 0 1
## 196 98
```

Any model we choose has to have an accuracy higher than 98/(98+196) = 33%. This is the baseline accuracy score.

### Logistic Regression Classifier

#### Training

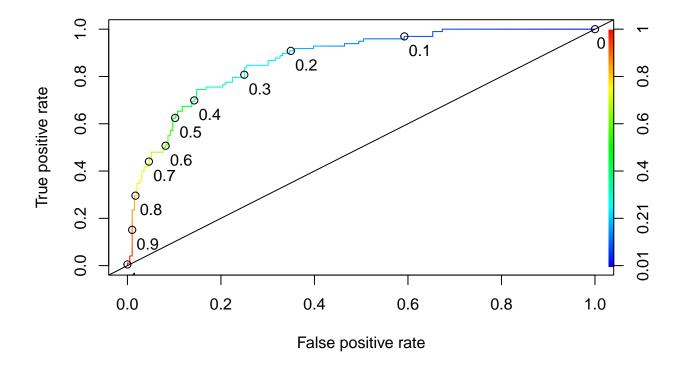
```
clf <- glm(hasDiabetes ~ ., family = binomial(link='logit'), data = train)
summary(clf)

##
## Call:
## glm(formula = hasDiabetes ~ ., family = binomial(link = "logit"),
## data = train)</pre>
```

```
##
## Deviance Residuals:
      Min
                    Median
##
                1Q
                                          Max
## -2.8732 -0.6344 -0.3526 0.5918
                                       2.2488
##
## Coefficients:
                             Estimate Std. Error z value Pr(>|z|)
                           -1.090e+01 1.516e+00 -7.190 6.48e-13 ***
## (Intercept)
## pregnancies
                            1.310e-01 6.738e-02
                                                  1.944 0.05192 .
## glucoseConcentration
                            4.080e-02 6.740e-03
                                                  6.054 1.41e-09 ***
## bloodPressure
                            2.284e-03 1.367e-02
                                                  0.167 0.86732
## skinThickness
                           -3.725e-06 2.118e-02
                                                  0.000 0.99986
## insulin
                           -1.764e-03 1.544e-03 -1.142 0.25325
                                                 2.731 0.00631 **
## bmi
                            9.483e-02 3.472e-02
## diabetesPedigreeFunction 1.472e+00 4.967e-01
                                                  2.963 0.00305 **
                            2.227e-02 2.099e-02
                                                 1.061 0.28881
## age
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
## (Dispersion parameter for binomial family taken to be 1)
##
##
      Null deviance: 374.27 on 293 degrees of freedom
## Residual deviance: 251.84 on 285 degrees of freedom
## AIC: 269.84
##
## Number of Fisher Scoring iterations: 5
saveRDS(clf, file = "../models/LogisticRegressionClassifier Full.Rds")
```

#### Model Evaluation on Train Set

```
predict <- predict(clf, type = 'response')</pre>
with(train,
     table(hasDiabetes, predict > 0.3))
##
## hasDiabetes FALSE TRUE
##
              0
                   147
                          49
##
              1
                    19
                          79
We can see in the confusion matrix that our accuracy is (79 + 147)/(147 + 49 + 19 + 79) = 0.7687. This is
with a cutoff of 0.3.
ROCRpred <- with(train,</pre>
                   prediction(predict, hasDiabetes))
ROCRperf <- performance(ROCRpred, 'tpr','fpr')</pre>
plot(ROCRperf, colorize = \frac{TRUE}{TRUE}, print.cutoffs.at = seq(0,1,0.1), text.adj = c(-0.2,1.7)); abline(0,1)
```



In the ROC curve we can see that our model is good (the curve is away from the diagonal). Since we care about not predicting a negative result for someone that is actually positive for diabetes (false negative rate), we want to have a larger true positive rate (1 - TPR = FNR). This means that we choose a cutoff near the blue part of the curve, the lower the cutoff the more cautious our model and the less accurate.

#### Model Evaluation on Test Set

#only after making decision of cutoff and dealing with missing values....