

# Constructing a risk model for cervical cancer

Tom Ertman

8/24/2020

## Introduction

This dataset consists of 858 samples of 36 health features collected from female patients in regards to possible risk factors for cervical cancer. The object of this paper is to construct a risk model that includes factors from various diagnostic tests so that a subject's risk may be ranked from 0, indicating low or no risk, to 4, indicating a high risk of cervical cancer. Note that this dataset does not indicate whether a subject has cervical cancer.

## A complete feature list:

## [1] "Age"	"Number.of.sexual.partners"
## [3] "First.sexual.intercourse"	"Num.of.pregnancies"
## [5] "Smokes"	"Smokes..years."
## [7] "Smokes..packs.year."	"Hormonal.Contraceptives"
## [9] "Hormonal.Contraceptives..years."	"IUD"
## [11] "IUD..years."	"STDs"
## [13] "STDs..number."	"STDs.condylomatosis"
## [15] "STDs.cervical.condylomatosis"	"STDs.vaginal.condylomatosis"
## [17] "STDs.vulvo.perineal.condylomatosis"	"STDs.syphilis"
## [19] "STDs.pelvic.inflammatory.disease"	"STDs.genital.herpes"
## [21] "STDs.molluscum.contagiosum"	"STDs.AIDS"
## [23] "STDs.HIV"	"STDs.Hepatitis.B"
## [25] "STDs.HPV"	"STDs..Number.of.diagnosis"
## [27] "STDs..Time.since.first.diagnosis"	"STDs..Time.since.last.diagnosis"
## [29] "Dx.Cancer"	"Dx.CIN"
## [31] "Dx.HPV"	"Dx"
## [33] "Hinselmann"	"Schiller"
## [35] "Citology"	"Biopsy"

The features - Schiller - Hinselmann - Citology - Biopsy

Are all medical tests designed to detect cancerous cells on the cervix.

- Dx.CIN indicated a diagnoses of Cervical intraepithelial neoplasia
- Dx.HPV indicates a diagnoses of Human Papilloma Virus
- Dx.Cancer indicates a previous diagnoses of cancer
- Dx is unknown and dropped from the study

## Analysis:

There are a number of challenges with this dataset, namely the unbalanced nature of the positive results in the diagnostic tests which make constructing an accurate model difficult.

```
table(dfile$Schiller)
```

```
##
##    0    1
## 784   74
```

```
table(dfile$Hinselmann)
```

```
##
##    0    1
## 823   35
```

```
table(dfile$Citology)
```

```
##
##    0    1
## 814   44
```

```
table(dfile$Biopsy)
```

```
##
##    0    1
## 803   55
```

There are columns with an significant amount of missing data as illustrated here

```
z<-sapply(dfile,function(x){
  sum(is.na(x))
})
```

```
#features missing more than half of data
names(z[which(unnamed(z)>400)])
```

```
## [1] "STDs..Time.since.first.diagnosis" "STDs..Time.since.last.diagnosis"
```

We will deal with these issues by

- 1) for columns missing less than 25% of data, we will use imputation methods to assign values to missing features. For continuous data, we will substitute the median value for that column, for factors, the mode.
- 2) we will discard features with a large amount of missing data >25%

For data modeling, we will chose decision tree and randomforest algorithms using cross validation and feature tuning.

We begin by dropping our Dx feature, then dropping our two columns that have a very high amount of NA

```
#drop dx
dfile<-dfile[,-32]
#drop the two highest NA features
dfile<-dfile[,-28]
dfile<-dfile[,-27]
```

Now we'll impute values on our dataset with the impute function to assign values to missing data

```
df2<-imputeMissings::impute(dfile)
#names(df2[nearZeroVar(dfile,freqCut = 99/1)])
#df2<-df2[,-nearZeroVar(dfile)]
```

Our dataset is summarized here

```
summary(df2)
```

```
##      Age      Number.of.sexual.partners First.sexual.intercourse
##  Min.   :13.00   Min.    : 1.000           Min.    :10
##  1st Qu.:20.00   1st Qu.: 2.000           1st Qu.:15
##  Median :25.00   Median : 2.000           Median :17
##  Mean   :26.82   Mean    : 2.512           Mean    :17
##  3rd Qu.:32.00   3rd Qu.: 3.000           3rd Qu.:18
##  Max.    :84.00   Max.    :28.000           Max.    :32
##  Num.of.pregnancies  Smokes      Smokes..years.  Smokes..packs.year.
##  Min.    : 0.000      Min.    :0.00000  Min.    : 0.000  Min.    : 0.0000
##  1st Qu.: 1.000      1st Qu.:0.00000  1st Qu.: 0.000  1st Qu.: 0.0000
##  Median : 2.000      Median :0.00000  Median : 0.000  Median : 0.0000
##  Mean    : 2.258      Mean    :0.1434  Mean    : 1.201  Mean    : 0.4463
##  3rd Qu.: 3.000      3rd Qu.:0.00000  3rd Qu.: 0.000  3rd Qu.: 0.0000
##  Max.    :11.000      Max.    :1.0000  Max.    :37.000  Max.    :37.0000
##  Hormonal.Contraceptives  Hormonal.Contraceptives..years.  IUD
##  Min.    :0.0000      Min.    : 0.000           Min.    :0.00000
##  1st Qu.:0.0000      1st Qu.: 0.000           1st Qu.:0.00000
##  Median :1.0000      Median : 0.500           Median :0.00000
##  Mean    :0.6865      Mean    : 2.035           Mean    :0.09674
##  3rd Qu.:1.0000      3rd Qu.: 2.000           3rd Qu.:0.00000
##  Max.    :1.0000      Max.    :30.000           Max.    :1.00000
##  IUD..years.      STDs      STDs..number.  STDs.condylomatosis
##  Min.    : 0.0000  Min.    :0.00000  Min.    : 0.000  Min.    :0.00000
##  1st Qu.: 0.0000  1st Qu.:0.00000  1st Qu.: 0.000  1st Qu.:0.00000
##  Median : 0.0000  Median :0.00000  Median : 0.000  Median :0.00000
##  Mean    : 0.4446  Mean    :0.09207  Mean    : 0.155  Mean    :0.05128
##  3rd Qu.: 0.0000  3rd Qu.:0.00000  3rd Qu.: 0.000  3rd Qu.:0.00000
##  Max.    :19.0000  Max.    :1.00000  Max.    : 4.000  Max.    :1.00000
##  STDs.cervical.condylomatosis  STDs.vaginal.condylomatosis
##  Min.    :0           Min.    :0.000000
##  1st Qu.:0           1st Qu.:0.000000
##  Median :0           Median :0.000000
##  Mean    :0           Mean    :0.004662
##  3rd Qu.:0           3rd Qu.:0.000000
##  Max.    :0           Max.    :1.000000
##  STDs.vulvo.perineal.condylomatosis  STDs.syphilis
```

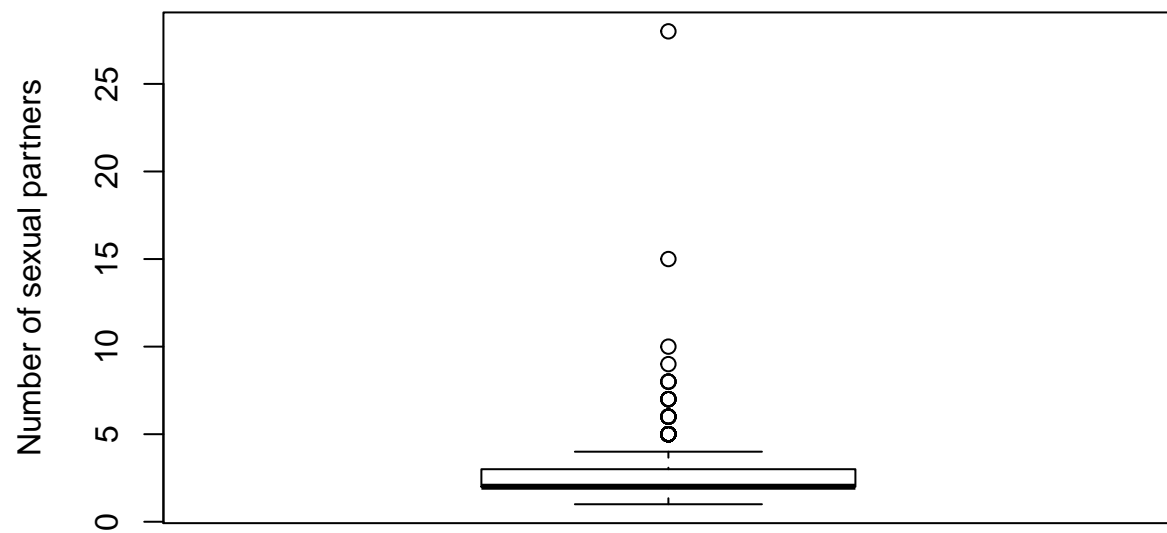
```

## Min.      :0.00000      Min.      :0.00000
## 1st Qu.:0.00000      1st Qu.:0.00000
## Median :0.00000      Median :0.00000
## Mean    :0.05012      Mean    :0.02098
## 3rd Qu.:0.00000      3rd Qu.:0.00000
## Max.    :1.00000      Max.    :1.00000
## STDs.pelvic.inflammatory.disease STDs.genital.herp
## Min.      :0.000000      Min.      :0.000000
## 1st Qu.:0.000000      1st Qu.:0.000000
## Median :0.000000      Median :0.000000
## Mean    :0.001166      Mean    :0.001166
## 3rd Qu.:0.000000      3rd Qu.:0.000000
## Max.    :1.000000      Max.    :1.000000
## STDs.molluscum.contagiosum STDs.AIDS STDs.HIV STDs.Hepatitis.B
## Min.      :0.000000      Min.      :0      Min.      :0.00000      Min.      :0.000000
## 1st Qu.:0.000000      1st Qu.:0      1st Qu.:0.00000      1st Qu.:0.000000
## Median :0.000000      Median :0      Median :0.00000      Median :0.000000
## Mean    :0.001166      Mean    :0      Mean    :0.02098      Mean    :0.001166
## 3rd Qu.:0.000000      3rd Qu.:0      3rd Qu.:0.00000      3rd Qu.:0.000000
## Max.    :1.000000      Max.    :0      Max.    :1.00000      Max.    :1.000000
## STDs.HPV STDs..Number.of.diagnosis Dx.Cancer
## Min.      :0.000000      Min.      :0.00000      Min.      :0.00000
## 1st Qu.:0.000000      1st Qu.:0.00000      1st Qu.:0.00000
## Median :0.000000      Median :0.00000      Median :0.00000
## Mean    :0.002331      Mean    :0.08741      Mean    :0.02098
## 3rd Qu.:0.000000      3rd Qu.:0.00000      3rd Qu.:0.00000
## Max.    :1.000000      Max.    :3.00000      Max.    :1.00000
## Dx.CIN Dx.HPV Hinselmann Schiller
## Min.      :0.00000      Min.      :0.00000      Min.      :0.00000      Min.      :0.00000
## 1st Qu.:0.00000      1st Qu.:0.00000      1st Qu.:0.00000      1st Qu.:0.00000
## Median :0.00000      Median :0.00000      Median :0.00000      Median :0.00000
## Mean    :0.01049      Mean    :0.02098      Mean    :0.04079      Mean    :0.08625
## 3rd Qu.:0.00000      3rd Qu.:0.00000      3rd Qu.:0.00000      3rd Qu.:0.00000
## Max.    :1.00000      Max.    :1.00000      Max.    :1.00000      Max.    :1.00000
## Citology Biopsy
## Min.      :0.00000      Min.      :0.0000
## 1st Qu.:0.00000      1st Qu.:0.0000
## Median :0.00000      Median :0.0000
## Mean    :0.05128      Mean    :0.0641
## 3rd Qu.:0.00000      3rd Qu.:0.0000
## Max.    :1.00000      Max.    :1.0000

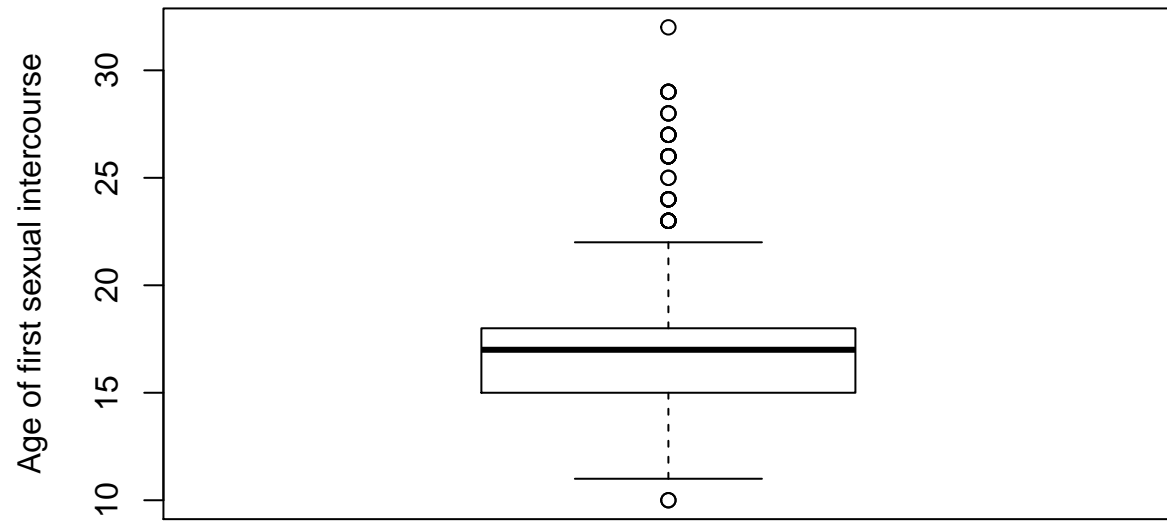
```

Here we examine possible outliers to our dataset

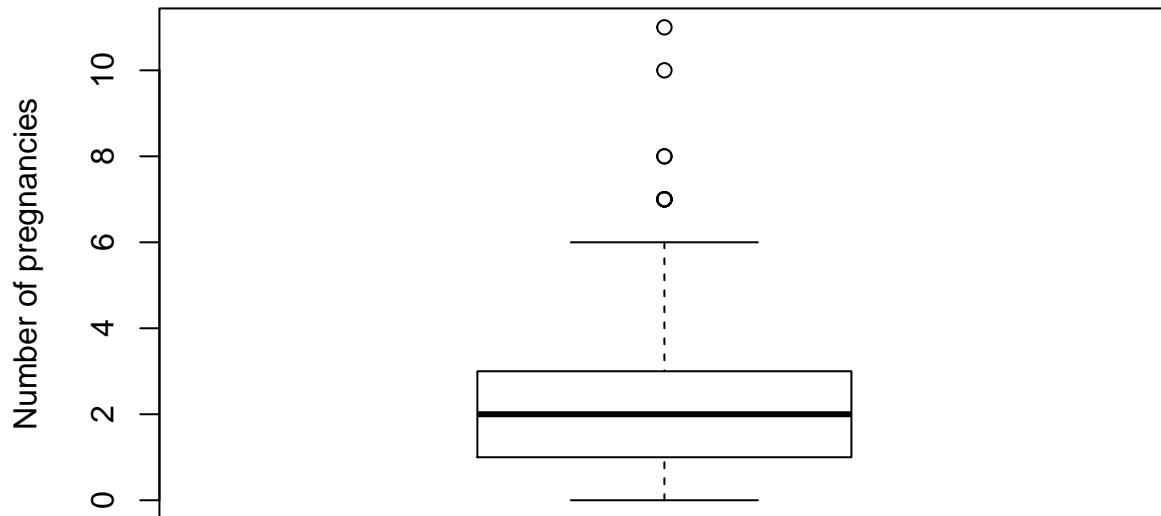
```
boxplot(df2$Number.of.sexual.partners,ylab="Number of sexual partners")
```



```
boxplot(df2$First.sexual.intercourse,ylab="Age of first sexual intercourse")
```



```
boxplot(df2$Num.of.pregnancies,ylab="Number of pregnancies")
```



Upon inspecting the data, we find an outlier to remove hence

```
out_sp<-outliers::outlier(df2$Number.of.sexual.partners)
df2[which(df2$Number.of.sexual.partners==out_sp),]
```

```
##      Age Number.of.sexual.partners First.sexual.intercourse Num.of.pregnancies
## 468   16                        28                        10                  1
##      Smokes Smokes..years. Smokes..packs.year. Hormonal.Contraceptives
## 468       1                5                5                0
##      Hormonal.Contraceptives..years. IUD IUD..years. STDs STDs..number.
## 468                                0  0                0  0                0
##      STDs.condylomatosis STDs.cervical.condylomatosis
## 468                      0                          0
##      STDs.vaginal.condylomatosis STDs.vulvo.perineal.condylomatosis
## 468                             0                          0
##      STDs.syphilis STDs.pelvic.inflammatory.disease STDs.genital.herpes
## 468                0                          0                0
##      STDs.molluscum.contagiosum STDs.AIDS STDs.HIV STDs.Hepatitis.B STDs.HPV
## 468                            0          0          0                0      0
##      STDs..Number.of.diagnosis Dx.Cancer Dx.CIN Dx.HPV Hinselmann Schiller
## 468                            0          0          0          0          0
##      Citology Biopsy
## 468          0      0
```

```
df2<-df2[!df2$Number.of.sexual.partners==out_sp,]
```

Finally, we create our aggregate risk factor:

```
data_set<-df2%>%mutate(risk_level=Hinselmann+Schiller+Citology+Biopsy)
data_set$risk_level<-factor(data_set$risk_level)
```

Now we separate into train and test sets

```
#draw a sample from our completed dataset
set.seed(1999)
index<-createDataPartition(data_set$risk_level,p=0.8)

#separate into train and test
training_set<-data_set[index$Resample1,]
test_set<-data_set[-index$Resample1,]
```

For our random forest model, we use 10 cross validations with a grid search for optimal parameters.

```
contrl=trainControl(method="cv",number=10,search="grid")

#execute the model on our training set
rf_model<-train(risk_level~.,data=training_set,method="rf",trControl=contrl)

#produce the confusion matrix for our results
confusionMatrix(predict(rf_model),training_set$risk_level)
```

```
## Confusion Matrix and Statistics
```

```
##
```

```
##           Reference
```

```
## Prediction    0    1    2    3    4
```

```
##           0 604    0    0    0    0
```

```
##           1   0   33    0    0    0
```

```
##           2   0   0   18    0    0
```

```
##           3   0   0   0   27    0
```

```
##           4   0   0   0   0    5
```

```
##
```

```
## Overall Statistics
```

```
##
```

```
##           Accuracy : 1
```

```
##           95% CI : (0.9946, 1)
```

```
## No Information Rate : 0.8792
```

```
## P-Value [Acc > NIR] : < 2.2e-16
```

```
##
```

```
##           Kappa : 1
```

```
##
```

```
## McNemar's Test P-Value : NA
```

```
##
```

```
## Statistics by Class:
```

```
##
```

```
##           Class: 0 Class: 1 Class: 2 Class: 3 Class: 4
```

```
## Sensitivity      1.0000  1.00000  1.0000  1.0000 1.000000
```

```
## Specificity      1.0000  1.00000  1.0000  1.0000 1.000000
```

```
## Pos Pred Value   1.0000  1.00000  1.0000  1.0000 1.000000
```

```
## Neg Pred Value   1.0000  1.00000  1.0000  1.0000 1.000000
```

```
## Prevalence       0.8792  0.04803  0.0262  0.0393 0.007278
```



```
## Detection Rate      0.8792  0.04803  0.0262  0.0393  0.007278
## Detection Prevalence 0.8792  0.04803  0.0262  0.0393  0.007278
## Balanced Accuracy   1.0000  1.00000  1.0000  1.0000  1.000000
```

Now we run our model on our test test:

```
final_rf<-predict(rf_model,newdata=test_set)
final_cf<-confusionMatrix(final_rf,test_set$risk_level)
final_cf
```

```
## Confusion Matrix and Statistics
##
##              Reference
## Prediction    0    1    2    3    4
##              0 151    0    0    0    0
##              1    0    8    0    0    0
##              2    0    0    4    0    0
##              3    0    0    0    6    0
##              4    0    0    0    0    1
##
## Overall Statistics
##
##              Accuracy : 1
##              95% CI : (0.9785, 1)
##      No Information Rate : 0.8882
##      P-Value [Acc > NIR] : 1.777e-09
##
##              Kappa : 1
##
##      McNemar's Test P-Value : NA
##
## Statistics by Class:
##
##              Class: 0 Class: 1 Class: 2 Class: 3 Class: 4
## Sensitivity      1.0000  1.00000  1.00000  1.00000  1.000000
## Specificity      1.0000  1.00000  1.00000  1.00000  1.000000
## Pos Pred Value   1.0000  1.00000  1.00000  1.00000  1.000000
## Neg Pred Value    1.0000  1.00000  1.00000  1.00000  1.000000
## Prevalence       0.8882  0.04706  0.02353  0.03529  0.005882
## Detection Rate    0.8882  0.04706  0.02353  0.03529  0.005882
## Detection Prevalence 0.8882  0.04706  0.02353  0.03529  0.005882
## Balanced Accuracy 1.0000  1.00000  1.00000  1.00000  1.000000
```

For decision tree model we set up a parameter tuning grid that varies the split, complexity parameter, max depth

```
split<-seq(1,20,2)
cp=seq(.001,.02,.002)
mdepth=seq(20,30,5)

parameters=as.matrix(expand.grid(msplit=split,pval=cp,mxdepth=mdepth))
```

We construct a loop that manually applies each parameter and records the accuracy

```
rpart_test<-function(msplit,p,mxdepth){

  contrl=rpart.control(minsplit=msplit,cp=p,maxdepth=mxdepth)
  dtree=rpart(data=training_set,risk_level~.,control=contrl)
  confusionMatrix(predict(dtree,type="class"),training_set$risk_level)$overall["Accuracy"]

}

acc<-matrix()
for ( i in seq(1,nrow(parameters))){

  acc[i]<-rpart_test(parameters[i,1],parameters[i,2],parameters[i,3])

}
```

Now we apply the optimal parameters to our model

```
index<-first(which(acc==max(acc)))
contrl<-rpart.control(minsplit=parameters[index,1],cp=parameters[index,2],maxdepth=parameters[index,3])

dtree<-rpart(data=training_set,risk_level~.,control=contrl)
confusionMatrix(predict(dtree,type="class"),training_set$risk_level)
```

## Confusion Matrix and Statistics

##

##           Reference

## Prediction   0   1   2   3   4

##           0 604   0   0   0   0

##           1   0 33   0   0   0

##           2   0   0 18   0   0

##           3   0   0   0 27   0

##           4   0   0   0   0 5

##

## Overall Statistics

##

##                   Accuracy : 1

##                   95% CI : (0.9946, 1)

##       No Information Rate : 0.8792

##       P-Value [Acc > NIR] : < 2.2e-16

##

##                   Kappa : 1

##

##   McNemar's Test P-Value : NA

##

## Statistics by Class:

##

##                           Class: 0 Class: 1 Class: 2 Class: 3 Class: 4

## Sensitivity               1.0000 1.00000 1.0000 1.0000 1.000000

## Specificity               1.0000 1.00000 1.0000 1.0000 1.000000

## Pos Pred Value           1.0000 1.00000 1.0000 1.0000 1.000000

```
## Neg Pred Value      1.0000  1.00000  1.0000  1.0000 1.000000
## Prevalence          0.8792  0.04803  0.0262  0.0393 0.007278
## Detection Rate      0.8792  0.04803  0.0262  0.0393 0.007278
## Detection Prevalence 0.8792  0.04803  0.0262  0.0393 0.007278
## Balanced Accuracy    1.0000  1.00000  1.0000  1.0000 1.000000
```

Using the optimized hyperparameters for our model we get

```
confusionMatrix(predict(dtree,newdata =test_set,type='class'),test_set$risk_level)
```

```
## Confusion Matrix and Statistics
##
##           Reference
## Prediction  0  1  2  3  4
##           0 151  0  0  0  0
##           1  0  8  0  0  0
##           2  0  0  4  0  0
##           3  0  0  0  6  0
##           4  0  0  0  0  1
##
## Overall Statistics
##
##           Accuracy : 1
##           95% CI : (0.9785, 1)
##           No Information Rate : 0.8882
##           P-Value [Acc > NIR] : 1.777e-09
##
##           Kappa : 1
##
##           McNemar's Test P-Value : NA
##
## Statistics by Class:
##
##           Class: 0 Class: 1 Class: 2 Class: 3 Class: 4
## Sensitivity      1.0000  1.00000  1.00000  1.00000 1.000000
## Specificity      1.0000  1.00000  1.00000  1.00000 1.000000
## Pos Pred Value   1.0000  1.00000  1.00000  1.00000 1.000000
## Neg Pred Value   1.0000  1.00000  1.00000  1.00000 1.000000
## Prevalence       0.8882  0.04706  0.02353  0.03529 0.005882
## Detection Rate   0.8882  0.04706  0.02353  0.03529 0.005882
## Detection Prevalence 0.8882  0.04706  0.02353  0.03529 0.005882
## Balanced Accuracy 1.0000  1.00000  1.00000  1.00000 1.000000
```

## Summary

Two models were run on our data and were both accurate in predicting aggregate risk levels associated with cervical cancer. However, we note that the variables used to construct each respective model differ in importance

```
rforest<-head(arrange(varImp(rf_model)$importance,desc(Overall)),10)
dctree<-head(arrange(varImp(dtree),desc(Overall)),10)
```

For our Random Forest model

```
rforest
```

```
##                                Overall
## Schiller                      100.000000
## Citology                      63.2970844
## Biopsy                       34.9116756
## Hinselmann                   16.9386855
## Age                          3.1996203
## Number.of.sexual.partners    1.6711225
## Hormonal.Contraceptives..years. 1.2961201
## First.sexual.intercourse     1.1688094
## Num.of.pregnancies          1.1453710
## IUD..years.                 0.7835994
```

and our Decision tree

```
dctree
```

```
##                                Overall
## Citology                      105.151529
## Biopsy                       84.341873
## Schiller                     66.952077
## Hinselmann                   51.930232
## Age                          8.145089
## Number.of.sexual.partners    6.690218
## IUD..years.                 6.675480
## Dx.Cancer                   3.821026
## STDs.genital.herpis         3.813718
## First.sexual.intercourse     3.318503
```

Using the specificity and sensitivi data for each test in combination with other tests we should be able to caculate how many subjects develop cancer and if all four diagnostic exams are necessary to make a diagnoses. With image samples from diagnostics test we should be able to identify cancerous cells via machine learning and possibly eliminate the need for painful tests such as a biopsy.