Final Project Report

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Introduction

Breast cancer is the most common cancer worldwide and the most common cancer diagnosed in the US (Mayo). Each year in the US, about 264,000 cases of breast cancer are diagnosed in women and about 2,400 in men (CDC).

Early diagnosis of the condition is crucial to improve the survival rate and relieve suffering in patients. Mammography is an effective X-ray imaging technology that detects breast cancer early. Classically, benign or malignant breast tumors are diagnosed by radiologists' interpretation of mammograms based on clinical parameters. However, diagnosing cancer is challenging even for the most skilled doctors. Since masses are heterogeneous, clinical parameters supply limited information on mammography mass. The symptoms are often shared with diseases and conditions that are unrelated to cancer, leading doctors to improperly diagnose the disease.

Cancerous lumps are often confused for blocked milk ducts, breast cysts, and other benign conditions. According to an expansive study conducted by Dartmouth College, the University of Vermont, and the Fred Hutchinson Cancer Research Center, and published in the March 2015 issue of the Journal of American Medical Association, approximately 13% of the diagnoses missed Stage 1 breast cancer. Meanwhile, 48% failed to detect atypia hyperplasia, a precursor to breast cancer. A significant number also over-diagnosed atypia hyperplasia.

There is, therefore, an urgent need to find new tools that can identify patients with breast cancer. Our study aims to build supervised machine-learning models to predict the diagnosis of breast cancer and understand the most important variables, to assist doctors and radiologists in accurately interpreting mammography imaging.

We built 3 models in total: Lasso penalized logistic regression, SVM, and Random Forest.

Data

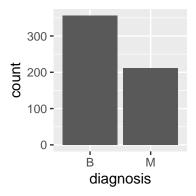
We obtain the Breast Cancer Wisconsin (Diagnosis) Data Set from Kaggle. The dataset contains diagnosis results and features of the cell nuclei computed from a digitized image of a fine needle aspirate (FNA) of a breast mass for 568 patients. The size of the nucleus is expressed by the features radius and area. The shape is expressed by the features smoothness, concavity, compactness, concave points, symmetry, and fractal dimension. The perimeter expresses both the size and shape of the nucleus. A higher value of shape features corresponds to a less regular contour and, therefore, to a higher probability of malignancy. For each of the features the mean value, worst value (mean of the three largest values), and standard error are computed for each image, resulting in 30 features of 568 images.

Data Processing

The original dataset contains a blank column '...33', so we dropped it. We also dropped the 'id' column, and rename several columns that contains blank space in their names.

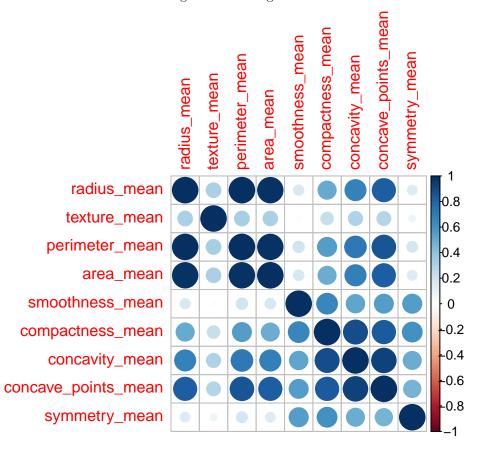
In order to fit SVM on the data, we encode the diagnosis variable into a factor variable with level 1 and -1: We partition the data into training and testing sets using a 70-30 percentage split(70% of the original data as the training set, and 30% as the testing set):

EDA



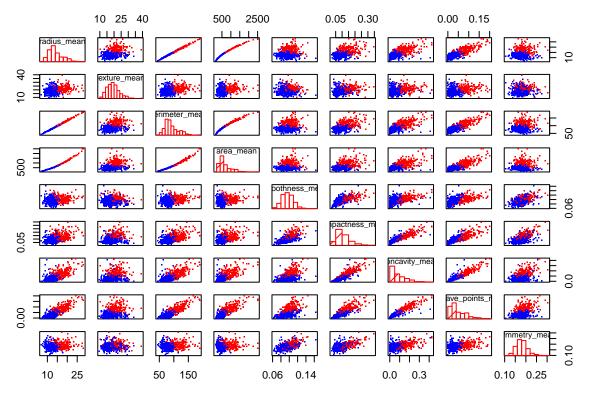
The bar plot shows that there is a larger number of benign than malignant cancer.

We divide the data into 3 categories according to their features.

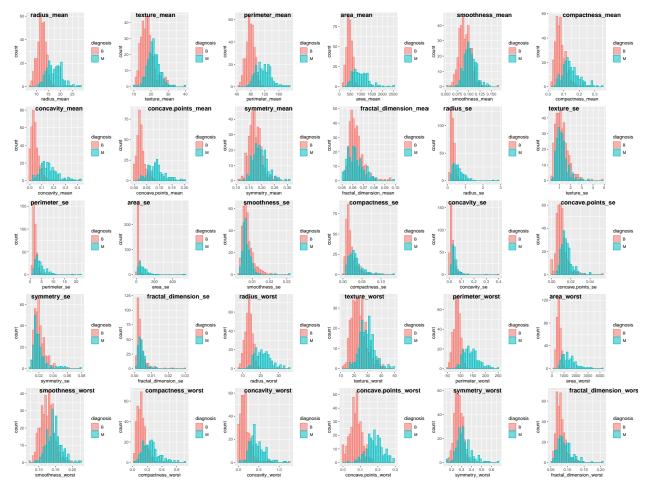


Major observations:

- Radius_mean, perimeter_mean, and area_mean are highly correlated.
- Compactness_mean, concavity_mean and concave_points_mean are highly correlated.



We observe from the scatterplot matrix that the two classifications seem to be generally separable, with distinct regions in the scatterplot visualization of the association with the predictors.



After deriving the histogram comparing distributions of predictors based on the two classifications, we would like to find features with little overlap between benign and malignant classes which will likely to be significant for diagnosis.

Methodology

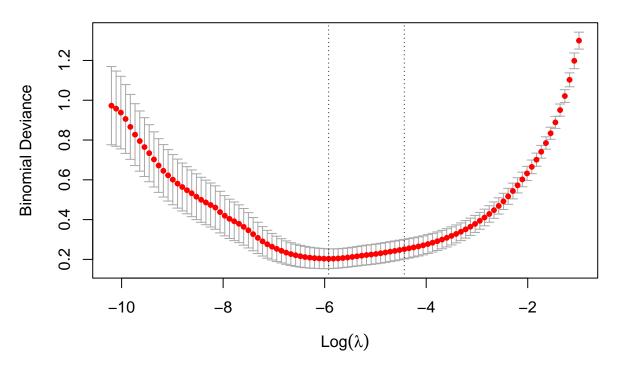
Model 1: LASSO-Penalized Logistic Regression

We decided to use a LASSO-penalized logistic regression model to perform variable selection by gauging insights into which predictors are the most contributive, since less significant variables are forced to be exactly zero, and the most significant variables are kept in the final model.

Model Selection

We included the interaction term symmetry_worst*concave_points_mean to explore the interaction between symmetery and the mean for number of concave portions of the contour.

29 25 26 24 22 18 17 15 10 9 7 6 5 5 4 3 2



[1] 0.002682998

##	32 x 1 sparse Matrix of class	"dgCMatrix"				
##		s0				
##	(Intercept) -28.47476478					
##	radius_mean .					
##	texture_mean	0.05803386				
##	perimeter_mean					
##	area_mean .					
##	smoothness_mean .					
##	compactness_mean .					
##	concavity_mean .					
##	concave_points_mean 18.2161782					
##	symmetry_mean	•				
	fractal_dimension_mean	•				
##	radius_se	4.79328840				
##	texture_se	-0.54137861				
##	perimeter_se					
##	area_se	0.04492504				
##	smoothness_se	91.67837368				
##	t compactness_se -43.3963					
##	concavity_se .					
##	concave_points_se .					
	symmetry_se	•				
	fractal_dimension_se	-82.24760167				
##	radius_worst	0.60106634				
	texture_worst	0.22789172				
##	perimeter_worst	•				
	area_worst	•				
##	smoothness_worst	17.86872342				
##	compactness_worst	•				

```
## concavity_worst 4.40330813
## concave_points_worst 20.93831516
## symmetry_worst 6.54485491
## fractal_dimension_worst .
## concave_points_mean:symmetry_worst 28.55102927
```

Prediction

Using the logistic regression model, besides classification we also want to understand uncertainty - more specifically, predictive probabilities that a tumor is benigh or malignant given the values of the predictors:

textur <u>eomæ</u>	<u>erapdiintestmenesesse</u> oot	hmusspastraetsa <u>l</u>	selimlenstewtusesnworts	tness <u>ca</u> wityz <u>st</u> wore	stsyoimus probastilistidicted_class
10.38 0.147	1.095 0.905 153.40.006	0.049 0.006	25.38 17.33 0.162	0.712 0.265	0.460 1.000 M
$17.77 \ 0.070$	$0.543 0.734\ 74.080.005$	0.013 0.004	$24.99 \ 23.41 \ 0.124$	0.242 0.186	$0.275 \ 1.000 \ \mathrm{M}$
$21.82\ 0.094$	$0.3061.002\;\; 24.320.006$	0.035 0.004	$15.49 \ 30.73 \ 0.170$	0.539 0.206	$0.438 \ 0.995 \ \mathrm{M}$
$22.61\ 0.080$	$0.2121.169\ 19.210.006$	0.059 0.008	$15.03 \ 32.01 \ 0.165$	0.694 0.221	$0.360 \ 0.954 \ \mathrm{M}$
$20.13 \ 0.053$	$0.4731.240\ 45.400.006$	0.012 0.002	$19.07 \ 30.88 \ 0.146$	0.291 0.161	$0.303 \ 0.995 \ \mathrm{M}$
$20.68\ 0.103$	$0.5691.073\ 54.180.007$	0.025 0.004	$20.96 \ 31.48 \ 0.179$	0.478 0.207	$0.371 \ 1.000 \ \mathrm{M}$
$22.15\ 0.095$	$0.7581.017 \ 112.400.006$	0.019 0.002	$27.32 \ 30.88 \ 0.151$	0.537 0.239	0.277 1.000 M
$15.71 \ 0.031$	$0.185 0.748\ 14.670.004$	0.019 0.002	$14.50\ 20.49\ 0.131$	0.189 0.073	0.318 0.002 B
$20.25 \ 0.077$	0.8531.849 93.540.011	0.027 0.004	$21.31 \ 27.26 \ 0.134$	0.345 0.149	0.234 1.000 M
$18.70 \ 0.052$	$0.4821.030\ \ 41.000.006$	0.034 0.006	$16.82\ 28.12\ 0.164$	0.696 0.155	$0.476 \ 0.995 \ \mathrm{M}$

[1] 0.9883041

We achieved a prediction accuracy of 0.9883. To interpret the predictions, we see that a patient with tumor with texture_mean of 10.38, concave_points_mean of 0.147, radius_se of 1.095, texture_se of 0.905, area_se of 153.400, smoothness_se of 0.006, compactness_se of 0.049, fractal_dimension_se of 0.006, radius_worst of 25.380, texture_worst of 17.33, smoothness_worst of 0.162, concavity_worst of 0.712, concave_points_worst of 0.265, symmetry_worst of 0.460 is expected to have a 100% of being diagnosed as malignant tumor. Whereas... is expected to have a 87.4% of being diagnosed as malignant tumor.

Model 2: SVM

Linear Kernel SVM

We use the predictors selected by the LASSO penalized logistic regression as predictors for the support vector machine model:

If two predictors have high correlation, we only use one of them:

```
##
## Parameter tuning of 'svm':
##
## - sampling method: 10-fold cross validation
##
## - best parameters:
    cost
##
##
     0.1
##
## - best performance: 0.02275641
##
## - Detailed performance results:
##
                error dispersion
      cost
```

```
## 1 1e-03 0.09301282 0.03304092
## 2 1e-02 0.05038462 0.02042181
## 3 1e-01 0.02275641 0.02208879
## 4 1e+00 0.03275641 0.02065577
## 5 5e+00 0.04025641 0.02412695
## 6 1e+01 0.04025641 0.02412695
## 7 1e+02 0.04032051 0.02117315
##
## Call:
## best.tune(METHOD = svm, train.x = diagnosis_binary ~ texture_mean +
       concave_points_mean + radius_se + texture_se + area_se + smoothness_se +
##
       compactness_se + fractal_dimension_se + radius_worst + texture_worst +
##
       smoothness_worst + concavity_worst + concave_points_worst + symmetry_worst +
##
       concave_points_mean * symmetry_worst, data = cancer_train, ranges = list(cost = c(0.001,
##
       0.01, 0.1, 1, 5, 10, 100)), kernel = "linear")
##
##
## Parameters:
     SVM-Type: C-classification
##
   SVM-Kernel: linear
##
##
         cost: 0.1
## Number of Support Vectors: 55
##
   (28 27)
##
##
##
## Number of Classes: 2
##
## Levels:
## -1 1
##
          truth
## predict -1 1
       -1 98 5
##
##
        1 0 68
## [1] 0.02923977
The misclassification rate is 0.02923.
Radial Kernel SVM
##
## Parameter tuning of 'svm':
## - sampling method: 10-fold cross validation
## - best parameters:
##
   cost gamma
##
      1 0.5
## - best performance: 0.04794872
```

- Detailed performance results:

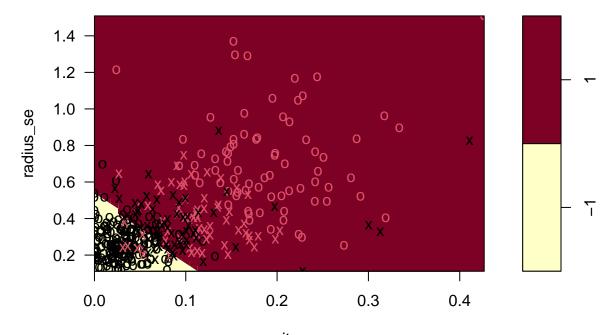
```
error dispersion
      cost gamma
## 1 1e-01
              0.5 0.35019231 0.06877558
## 2 1e+00
              0.5 0.04794872 0.03456232
              0.5 0.05044872 0.03147495
## 3 1e+01
## 4 1e+02
              0.5 0.05044872 0.03147495
## 5
    1e+03
            0.5 0.05044872 0.03147495
## 6 1e-01
             1.0 0.35019231 0.06877558
## 7 1e+00
              1.0 0.24192308 0.08101092
## 8 1e+01
              1.0 0.20929487 0.09689376
## 9 1e+02
              1.0 0.20929487 0.09689376
## 10 1e+03
              1.0 0.20929487 0.09689376
## 11 1e-01
              2.0 0.35019231 0.06877558
              2.0 0.34762821 0.06673515
## 12 1e+00
## 13 1e+01
              2.0 0.34012821 0.05872625
## 14 1e+02
              2.0 0.34012821 0.05872625
## 15 1e+03
              2.0 0.34012821 0.05872625
## 16 1e-01
              3.0 0.35019231 0.06877558
## 17 1e+00
              3.0 0.35019231 0.06877558
## 18 1e+01
              3.0 0.35019231 0.06877558
## 19 1e+02
              3.0 0.35019231 0.06877558
## 20 1e+03
             3.0 0.35019231 0.06877558
## 21 1e-01
              4.0 0.35019231 0.06877558
## 22 1e+00
              4.0 0.35019231 0.06877558
## 23 1e+01
              4.0 0.35019231 0.06877558
## 24 1e+02
              4.0 0.35019231 0.06877558
## 25 1e+03
              4.0 0.35019231 0.06877558
##
## Call:
  best.tune(METHOD = svm, train.x = diagnosis_binary ~ texture_mean +
       concave_points_mean + radius_se + texture_se + area_se + smoothness_se +
##
##
       compactness_se + fractal_dimension_se + radius_worst + texture_worst +
##
       smoothness_worst + concavity_worst + concave_points_worst + symmetry_worst +
       concave_points_mean * symmetry_worst, data = cancer_train, ranges = list(cost = c(0.1,
##
##
       1, 10, 100, 1000), gamma = c(0.5, 1, 2, 3, 4)), kernel = "radial")
##
##
## Parameters:
     SVM-Type: C-classification
##
##
   SVM-Kernel: radial
##
         cost: 1
##
## Number of Support Vectors:
##
##
   (130 160)
##
##
## Number of Classes: 2
##
## Levels:
  -1 1
##
          truth
## predict -1 1
##
       -1 94 2
```

1 4 71

[1] 0.03508772

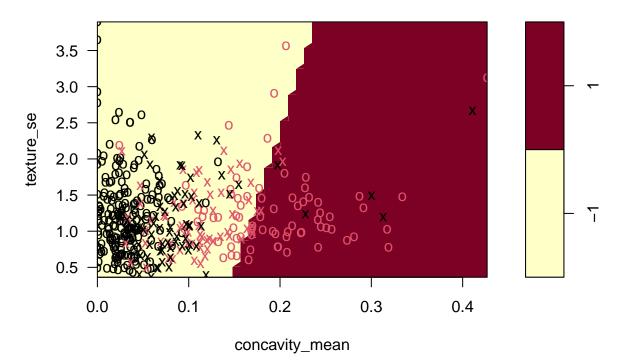
The misclassification rate is 0.03509, which is greater than that of the linear kernel which suggests that the two classes are likely to be linearly separable so that we can find a separating hyperplane using the linear kernel.

SVM classification plot



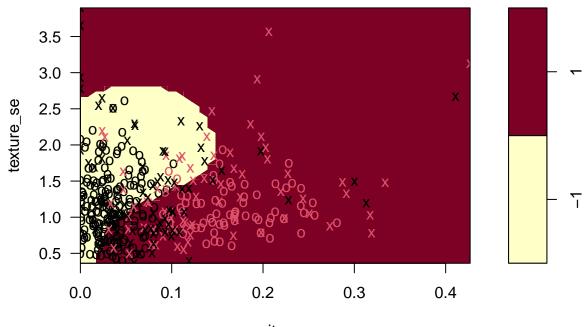
Linear

concavity_mean SVM classification plot

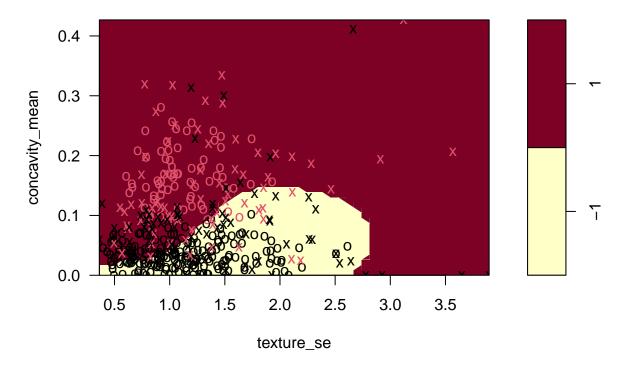


Radial

SVM classification plot



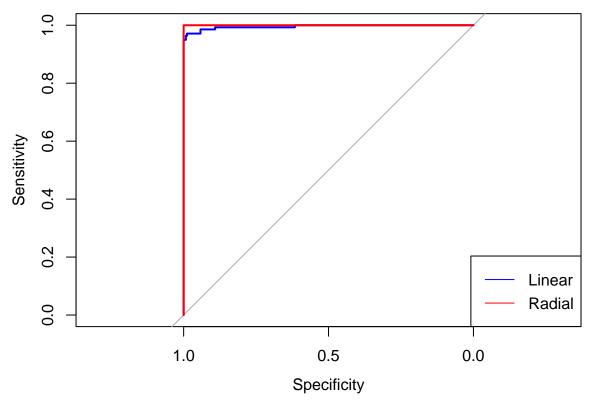
concavity_mean SVM classification plot

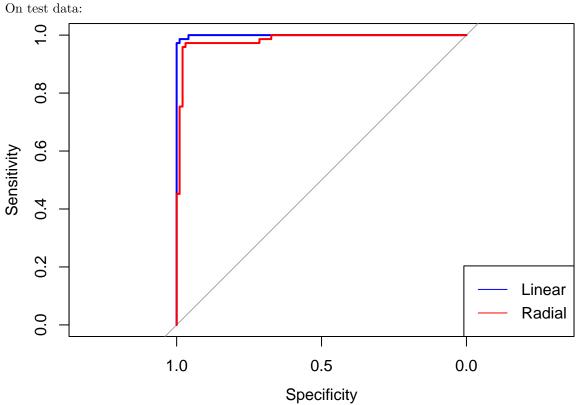


ROC (Linear SVM)

We visualize the ROC curves:

On train data:





Even though the radial kernel fits the training data more closely due to its higher complexity, the linear kernel performs better on the test data (since the data is likely to be linearly separable as explained above), we decided to select the model with the linear kernel.

Model 3: Random Forest

For Random Forest, we decided to use group 'Mean' and group 'Worst' separately. This is because, we want to understand the potential difference in prediction given the different level of severity of the patients' conditions. As we have mentioned in the introduction, 'Worst' measures mean of the three largest values. By building a model with only 'Mean' and 'Worst' variables, we think we can better understand the extreme cases.

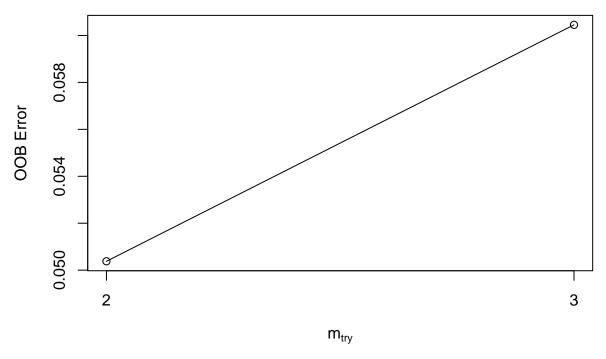
First, We select variables by running a preliminary random forest model with all the variables, to rank their importance.

##		MeanDecreaseGini
##	radius_mean	8.5616407
##	texture_mean	3.0559818
##	perimeter_mean	5.9008019
##	area_mean	9.8692050
##	smoothness_mean	1.1478325
##	compactness_mean	1.6773967
##	concavity_mean	10.1532690
##	concave_points_mean	18.7904551
##	symmetry_mean	0.6545792
##	<pre>fractal_dimension_mean</pre>	0.7849650
##	radius_se	2.4824456
##	texture_se	0.8773616
##	perimeter_se	2.8646560
##	area_se	6.6899822
##	smoothness_se	1.1900223
##	compactness_se	0.9471109
##	concavity_se	1.1550698
##	concave_points_se	1.0486457
##	symmetry_se	0.8049179
##	<pre>fractal_dimension_se</pre>	1.1728247
##	radius_worst	18.0287062
##	texture_worst	3.6374436
##	perimeter_worst	20.0049651
##	area_worst	17.7615481
##	smoothness_worst	2.9276817
##	compactness_worst	3.0046846
##	concavity_worst	6.6002452
##	concave_points_worst	25.2111650
##	symmetry_worst	1.8577452
##	${\tt fractal_dimension_worst}$	1.4519769

After testing different variables based on their important, and taking into accounts the collinearity issue we discussed in the EDA section, we decided to select the following variables as the predictors: concave_points_worst, area_worst, perimeter_worst, radius_worst, concave_points_mean, perimeter_mean, concavity_worst, area_se.

We chose mtry=2, because after tuning mtry, we found that mtry=2 has the lowest OOB error.

```
## mtry = 2 00B error = 5.04%
## Searching left ...
## Searching right ...
## mtry = 3 00B error = 6.05%
## -0.2 0.01
```



We chose the number of tree to be 500. The number of trees should be chosen carefully, since a high performance of the individual models might lead to overfitting when the number of trees is very high. However, taking 50 trees caused a lower accuracy than taking 500 trees. Therefore, this higher number of trees is chosen.

area_worst + perimeter_worst

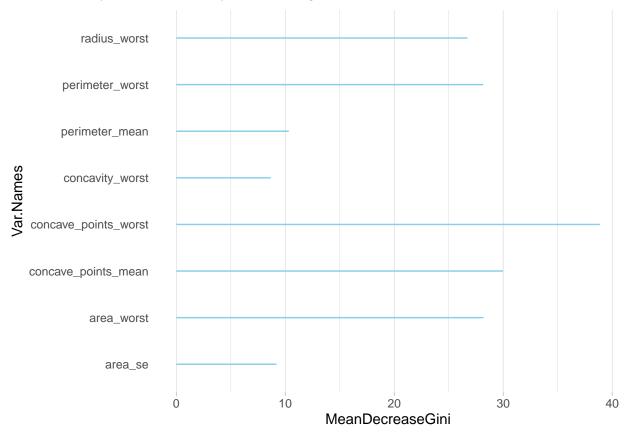
```
##
## Call:
##
    randomForest(formula = diagnosis_binary ~ concave_points_worst +
                  Type of random forest: classification
##
##
                        Number of trees: 500
## No. of variables tried at each split: 2
##
##
           OOB estimate of error rate: 5.04%
##
  Confusion matrix:
            1 class.error
               0.02713178
##
  -1 251
            7
       13 126
               0.09352518
  Confusion Matrix and Statistics
##
##
             Reference
##
  Prediction -1
                  1
##
           -1 96
                  5
               2 68
##
           1
##
##
                  Accuracy : 0.9591
##
                    95% CI: (0.9175, 0.9834)
       No Information Rate: 0.5731
##
##
       P-Value [Acc > NIR] : <2e-16
##
##
                     Kappa: 0.9159
##
```

Mcnemar's Test P-Value: 0.4497

##

```
##
##
               Sensitivity: 0.9796
               Specificity: 0.9315
##
            Pos Pred Value: 0.9505
##
##
            Neg Pred Value: 0.9714
                Prevalence: 0.5731
##
##
            Detection Rate: 0.5614
      Detection Prevalence: 0.5906
##
##
         Balanced Accuracy: 0.9555
##
##
          'Positive' Class : -1
##
```

The RF model yields a 96% accuracy for the testing set.



From the plot we can see that the most important predictors are: concave_points_worst, concave_points_mean, area_worst, perimeter_worst, and radius_worst. Interestingly, perimeter_worst has high gini coefficient, but perimeter_mean ranks second from the last.

Conclusion

Concavity is the severity of concave portions of the contour. A high concavity means that the boundary of the cell nucleus has indentations, and thus is rather rough than smooth. Concave points id the number of concave portions of the contour of the cell nucleus.