Support Vector Machines(SVMs) Tutorial

Sonali Narang 11/12/2019

Support Vector Machines(SVMs)

A Support Vector Machine (SVM) is a discriminative classifier formally defined by a separating hyperplane. Given labeled training data, the algorithm outputs an optimal hyperplane which categorizes new examples.

The Breast Cancer Dataset

699 Observations, 11 variables Predictor Variable: Class-benign or malignant

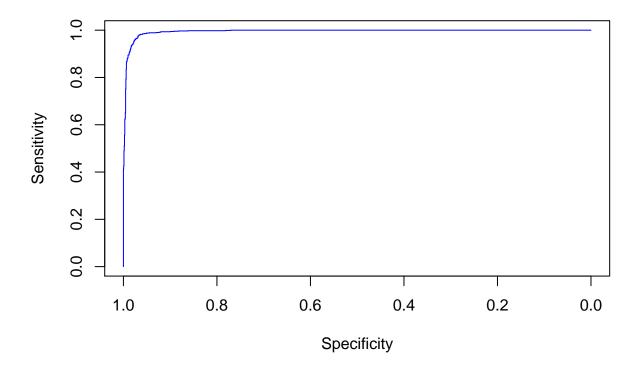
```
data(BreastCancer)
#bc = BreastCancer %>%
# mutate_if(is.character, as.numeric)
\#bc[is.na(bc)] = 0
BreastCancer_num = transform(BreastCancer, Id = as.numeric(Id),
                         Cl.thickness = as.numeric(Cl.thickness),
                         Cell.size = as.numeric(Cell.size),
                         Cell.shape = as.numeric(Cell.shape),
                         Marg.adhesion = as.numeric(Marg.adhesion),
                         Epith.c.size = as.numeric(Epith.c.size),
                         Bare.nuclei = as.numeric(Bare.nuclei),
                         Bl.cromatin = as.numeric(Bl.cromatin),
                         Normal.nucleoli = as.numeric(Normal.nucleoli),
                         Mitoses = as.numeric(Mitoses))
BreastCancer_num[is.na(BreastCancer_num)] = 0
train_size = floor(0.75 * nrow(BreastCancer_num))
train_pos <- sample(seq_len(nrow(BreastCancer_num)), size = train_size)</pre>
train classification <- BreastCancer num[train pos, ]</pre>
test_classification <- BreastCancer_num[-train_pos, ]</pre>
```

 $\#\#\mathrm{SVM}$

```
set.seed(1112)
control = trainControl(method = "repeatedcv", repeats = 5, classProbs = T, savePredictions = T)
svm = train(Class ~ Id + Cl.thickness + Cell.size + Cell.shape + Marg.adhesion + Epith.c.size + Bare.nu
svm
```

```
## Support Vector Machines with Linear Kernel
##
```

```
## 524 samples
## 10 predictor
    2 classes: 'benign', 'malignant'
##
## No pre-processing
## Resampling: Cross-Validated (10 fold, repeated 5 times)
## Summary of sample sizes: 472, 472, 472, 471, 471, ...
## Resampling results:
##
##
     Accuracy
                Kappa
    0.9667774 0.9259438
## Tuning parameter 'C' was held constant at a value of 1
##Receiver operating characteristic(ROC) curve
roc(predictor = svm$pred$malignant, response = svm$pred$obs)$auc
## Setting levels: control = benign, case = malignant
## Setting direction: controls < cases
## Area under the curve: 0.9946
plot(x = roc(predictor = svm$pred$malignant, response = svm$pred$obs)$specificities, y = roc(predictor =
## Setting levels: control = benign, case = malignant
## Setting direction: controls < cases
## Setting levels: control = benign, case = malignant
## Setting direction: controls < cases
```



Test Set

```
svm_test = predict(svm, newdata = test_classification)
confusionMatrix(svm_test, reference = test_classification$Class)
```

```
## Confusion Matrix and Statistics
##
##
              Reference
## Prediction benign malignant
##
     benign
                  111
                               5
     malignant
                             55
##
##
##
                  Accuracy: 0.9486
                    95% CI : (0.9046, 0.9762)
##
##
       No Information Rate: 0.6571
##
       P-Value [Acc > NIR] : <2e-16
##
                     Kappa : 0.8854
##
##
    Mcnemar's Test P-Value : 1
##
##
               Sensitivity: 0.9652
##
##
               Specificity: 0.9167
            Pos Pred Value: 0.9569
##
            Neg Pred Value: 0.9322
##
                Prevalence: 0.6571
##
```

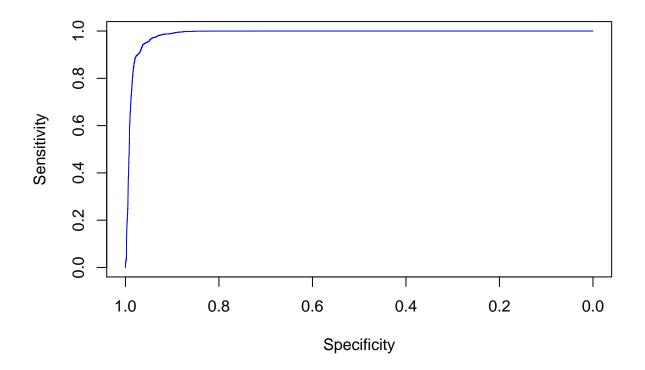
```
## Detection Rate : 0.6343
## Detection Prevalence : 0.6629
## Balanced Accuracy : 0.9409
##
## 'Positive' Class : benign
##
```

SVM with a radial kernel

```
set.seed(1112)
control = trainControl(method = "repeatedcv", repeats = 5, classProbs = T, savePredictions = T)
svm = train(Class ~ Id + Cl.thickness + Cell.size + Cell.shape + Marg.adhesion + Epith.c.size + Bare.nu
svm
## Support Vector Machines with Radial Basis Function Kernel
##
## 524 samples
## 10 predictor
   2 classes: 'benign', 'malignant'
## No pre-processing
## Resampling: Cross-Validated (10 fold, repeated 5 times)
## Summary of sample sizes: 472, 472, 472, 471, 471, ...
## Resampling results across tuning parameters:
##
##
    C
            Accuracy
                       Kappa
      0.25 0.9462256 0.8845771
##
##
      0.50 0.9481487 0.8886031
##
      1.00 0.9519513 0.8964471
##
      2.00 0.9511821 0.8947132
##
      4.00 0.9500355 0.8923302
##
      8.00 0.9508047 0.8939457
##
     16.00 0.9492662 0.8907169
     32.00 0.9500282 0.8923031
##
##
      64.00 0.9504128 0.8930790
     128.00 0.9508047 0.8939448
##
## Tuning parameter 'sigma' was held constant at a value of 0.7192712
## Accuracy was used to select the optimal model using the largest value.
## The final values used for the model were sigma = 0.7192712 and C = 1.
##Receiver operating characteristic(ROC) curve
roc(predictor = svm$pred$malignant, response = svm$pred$obs)$auc
## Setting levels: control = benign, case = malignant
## Setting direction: controls < cases
```

Area under the curve: 0.9873

```
plot(x = roc(predictor = svm$pred$malignant, response = svm$pred$obs)$specificities, y = roc(predictor = svm$pred$cobs)$specificities, y = ro
```



Test Set

```
svm_test = predict(svm, newdata = test_classification)
confusionMatrix(svm_test, reference = test_classification$Class)

## Confusion Matrix and Statistics
##

## Reference
## Prediction benign malignant
## benign 109 2
## malignant 6 58
```

```
##
##
                  Accuracy : 0.9543
                    95% CI : (0.9119, 0.9801)
##
##
       No Information Rate: 0.6571
       P-Value [Acc > NIR] : <2e-16
##
##
##
                     Kappa : 0.9001
##
##
    Mcnemar's Test P-Value : 0.2888
##
##
               Sensitivity: 0.9478
               Specificity: 0.9667
##
##
            Pos Pred Value: 0.9820
            Neg Pred Value: 0.9062
##
##
                Prevalence: 0.6571
##
            Detection Rate: 0.6229
##
      Detection Prevalence : 0.6343
         Balanced Accuracy: 0.9572
##
##
          'Positive' Class : benign
##
##
```

Homework

- 1. Choose an appropriate machine learning dataset and use SVM with two different kernels. Campare the results.
- 2. Attempt using SVM after using a feature selection method. Do the results improve? Explain.