HW4

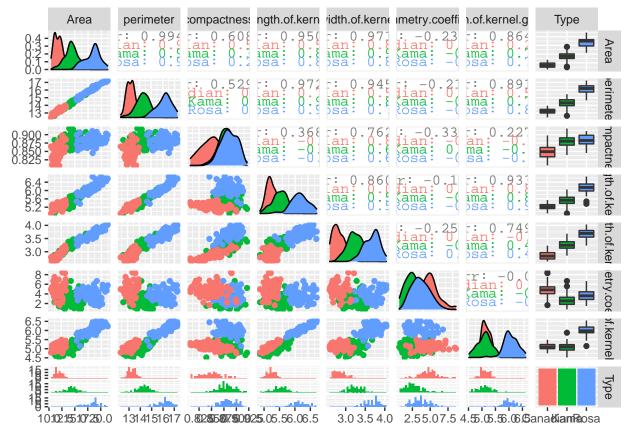
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```
##Problem 1
library(readr)
seeds <- read_csv("/Users/davidschultheiss/Downloads/seeds.csv")</pre>
## Parsed with column specification:
## cols(
##
     Area = col_double(),
##
     perimeter = col_double(),
##
     compactness = col_double(),
##
     length.of.kernel = col_double(),
##
     width.of.kernel = col_double(),
##
     asymmetry.coefficient = col_double(),
##
     length.of.kernel.groove = col_double(),
##
     Type = col_double()
## )
seeds$Type[which(seeds$Type==1)]="Kama"
seeds$Type[which(seeds$Type==2)]="Rosa"
seeds$Type[which(seeds$Type==3)]="Canadian"
seeds$Type = factor(seeds$Type)
  a)
library(GGally)
## Warning: package 'GGally' was built under R version 3.6.2
## Loading required package: ggplot2
## Warning: package 'ggplot2' was built under R version 3.6.2
## Registered S3 method overwritten by 'GGally':
    method from
##
##
     +.gg
          ggplot2
```

ggpairs(data= seeds, mapping= aes(color= seeds\$Type))

```
## 'stat_bin()' using 'bins = 30'. Pick better value with 'binwidth'.
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## 'stat_bin()' using 'bins = 30'. Pick better value with 'binwidth'.
```



Length of Kernel Groove, Compactness, and Asymmetry Coefficient. I chose these because all have weak correlations (>.34), and clustering for different Types.

b)

library(tidyverse)

```
## v tibble 2.1.3 v dplyr 0.8.3
## v tidyr 1.0.0 v stringr 1.4.0
## v purrr 0.3.3 v forcats 0.4.0
```

```
## -- Conflicts -----
                                            ----- tidyverse_conflicts() --
## x dplyr::filter() masks stats::filter()
## x dplyr::lag()
                    masks stats::lag()
set.seed(1)
testsample = sample(1:nrow(seeds), .2*nrow(seeds))
test = seeds[testsample, ]
training = seeds[-testsample, ]
test = select(test, Type, length.of.kernel.groove, compactness,
              asymmetry.coefficient)
training = select(training, Type, length.of.kernel.groove, compactness,
              asymmetry.coefficient)
  c) KNN -
library(class)
knn.fit = knn(training[,-1], test[,-1], training$Type, k=1)
miss = length(which(knn.fit != test$Type))
(rate = (miss / length(knn.fit)) * 100)
## [1] 30.95238
knn.fit = knn(training[,-1], test[,-1], training$Type, k=5)
miss = length(which(knn.fit != test$Type))
(rate = (miss / length(knn.fit)) * 100)
## [1] 28.57143
knn.fit = knn(training[,-1], test[,-1], training$Type, k=10)
miss = length(which(knn.fit != test$Type))
(rate = (miss / length(knn.fit)) * 100)
## [1] 23.80952
LDA -
library(MASS)
##
## Attaching package: 'MASS'
## The following object is masked from 'package:dplyr':
##
##
       select
```

```
ldafit = lda(data = training, Type ~ length.of.kernel.groove + compactness +
      asymmetry.coefficient)
lda.pred = predict(ldafit, test)
table(test$Type, lda.pred$class)
##
##
              Canadian Kama Rosa
##
     Canadian
                    11
##
                      4
     Kama
                           9
                                1
     Rosa
                      0
                               14
mean(test$Type != lda.pred$class)
## [1] 0.1904762
QDA -
qdafit = qda(data = training, Type ~ length.of.kernel.groove + compactness +
               asymmetry.coefficient)
qda.pred = predict(qdafit, test)
table(test$Type, qda.pred$class)
##
##
              Canadian Kama Rosa
##
     Canadian
                    12
##
     Kama
                     5
                           8
                                1
##
     Rosa
                           0
                               14
mean(test$Type != qda.pred$class)
## [1] 0.1904762
  d) The discriminant analysis models both have lower misclassification rates than KNN, and perform better
    with our data.
\#\#Problem 2
tumor <- read_csv("/Users/davidschultheiss/Downloads/tumor.csv")</pre>
## Parsed with column specification:
## cols(
##
     Diagnosis = col_character(),
    Radius = col_double(),
##
##
    Texture = col_double();
    Perimeter = col_double(),
##
##
     Area = col_double(),
     Smoothness = col_double(),
##
```

```
##
     Compactness = col_double(),
##
     Concavity = col_double(),
     'Concave Points' = col double(),
##
     Symmetry = col_double(),
##
     'Fractal Dimension' = col_double()
##
## )
  a)
is.factor(tumor$Diagnosis)
## [1] FALSE
tumor$Diagnosis = factor(tumor$Diagnosis)
ggpairs(data= tumor, mapping= aes(color= Diagnosis))
## 'stat_bin()' using 'bins = 30'. Pick better value with 'binwidth'.
## 'stat_bin()' using 'bins = 30'. Pick better value with 'binwidth'.
## 'stat_bin()' using 'bins = 30'. Pick better value with 'binwidth'.
## 'stat_bin()' using 'bins = 30'. Pick better value with 'binwidth'.
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## 'stat_bin()' using 'bins = 30'. Pick better value with 'binwidth'.
## 'stat_bin()' using 'bins = 30'. Pick better value with 'binwidth'.
     Diagnosi: Radius
                     Texture
                            Perimete
                                     Area
                                           noothne impactne Concavity scave Po Symmetrical Dimer
                                                                                          D:
m
     023850123850 10152025 1020304040802160 500050005000500250 0.0.2.30.0.0.0.2.30.005005200500520050050050050
```

Benign is red, and Malignant is teal.

Obviously variables like perimeter, area, and radius are highly correlated. Compactness and Concavity are also related.

No Variables look particularly reliable for predicting Diagnosis. Variables that could be useful are radius, texture, perimeter, area, compactness, concavity, and concave points.

I will be using three variables. Texture, area, and concavity. Area and concavity do have a .67 correlation, but most of our significant variables are highly related.

I think that KNN being non-parametric and having a decent amount of observations to train the data will make it the most reliable method.

b)

```
set.seed(1)
testsample = sample(1:nrow(tumor), .2*nrow(tumor))
test = tumor[testsample, ]
training = tumor[-testsample, ]
```

c)

```
##
## Call:
## glm(formula = Diagnosis ~ Texture + Area + Concavity, family = binomial,
##
       data = training)
##
## Deviance Residuals:
##
      Min
                 1Q
                      Median
                                   3Q
                                           Max
  -3.1928
           -0.2628 -0.0941
                               0.0384
##
                                        3.0324
##
## Coefficients:
##
                 Estimate Std. Error z value Pr(>|z|)
## (Intercept) -14.004711
                           1.652111 -8.477 < 2e-16 ***
## Texture
                 0.214226
                            0.051518
                                       4.158 3.21e-05 ***
## Area
                 0.011207
                            0.001525
                                       7.349 2.00e-13 ***
## Concavity
                26.062158
                            3.799056
                                       6.860 6.88e-12 ***
## ---
## Signif. codes: 0 '*** 0.001 '** 0.01 '* 0.05 '.' 0.1 ' 1
##
## (Dispersion parameter for binomial family taken to be 1)
##
##
       Null deviance: 610.92 on 455 degrees of freedom
## Residual deviance: 172.77 on 452 degrees of freedom
## AIC: 180.77
##
## Number of Fisher Scoring iterations: 7
```

All of our variables are significant.

d)

```
glm.probs = predict(glm.fit, test, type= 'response')
glm.pred = rep(0, nrow(test))
glm.pred[glm.probs > 0.5] = 1
table(test$Diagnosis, glm.pred)
##
              glm.pred
##
                0 1
##
     Benign
               77 3
     Malignant 4 29
##
Misclassification rate is 7/113 or 6.19%
  e) False Negative Rate = 4/81 or 4.94\%. False Positive Rate = 3/32 or 9.37\%
  f)
glm.pred = rep(0, nrow(test))
glm.pred[glm.probs > 0.3] = 1
table(test$Diagnosis, glm.pred)
              glm.pred
##
##
                0 1
               74 6
##
     Benign
     Malignant 1 32
Misclassification rate holds at 6.19%. FNR falls to 1.33%. FPR rises to 15.79%.
  g)
test = dplyr::select(test, Diagnosis, Texture, Area, Concavity)
training = dplyr::select(training, Diagnosis, Texture, Area, Concavity)
knn.fit = knn(training[,-1], test[,-1], training$Diagnosis, k=10)
miss = length(which(knn.fit != test$Diagnosis))
(rate = (miss / length(knn.fit)) * 100)
## [1] 8.849558
Misclassification for KNN is 8.85%
ldafit = lda(data = training, Diagnosis ~ Texture + Area + Concavity)
lda.pred = predict(ldafit, test)
table(test$Diagnosis, lda.pred$class)
##
##
               Benign Malignant
##
                   80
     Benign
                              27
     Malignant
                     6
##
```

```
mean(test$Diagnosis != lda.pred$class)
```

[1] 0.05309735

Misclassification rate for LDA is 5.3%. LDA is the lowest overall.

h) LDA is the best for this researcher. It has better predictive capability with a lower misclassification rate than logistic regression. KNN is not able to examine individual variables.