

Class 9 Mini Project- Breast Cancer Cell Analysis

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2/13/2022

1. Exploratory data analysis

Preparing the data

```
#Saving the input data file into my Project Directory
fna.data <- "WisconsinCancer.csv"
#Input the data and store it under wisc.df
wisc.df <- read.csv(fna.data, row.names=1)
#Inspect data
View(wisc.df)
head(wisc.df)
```

```
##      diagnosis radius_mean texture_mean perimeter_mean area_mean
## 842302         M      17.99       10.38         122.80      1001.0
## 842517         M      20.57       17.77         132.90      1326.0
## 84300903        M      19.69       21.25         130.00      1203.0
## 84348301         M      11.42       20.38          77.58       386.1
## 84358402         M      20.29       14.34         135.10      1297.0
## 843786         M      12.45       15.70          82.57       477.1
##      smoothness_mean compactness_mean concavity_mean concave.points_mean
## 842302           0.11840           0.27760           0.3001           0.14710
## 842517           0.08474           0.07864           0.0869           0.07017
## 84300903          0.10960           0.15990           0.1974           0.12790
## 84348301          0.14250           0.28390           0.2414           0.10520
## 84358402          0.10030           0.13280           0.1980           0.10430
## 843786           0.12780           0.17000           0.1578           0.08089
##      symmetry_mean fractal_dimension_mean radius_se texture_se perimeter_se
## 842302           0.2419           0.07871           1.0950           0.9053           8.589
## 842517           0.1812           0.05667           0.5435           0.7339           3.398
## 84300903          0.2069           0.05999           0.7456           0.7869           4.585
## 84348301          0.2597           0.09744           0.4956           1.1560           3.445
## 84358402          0.1809           0.05883           0.7572           0.7813           5.438
## 843786           0.2087           0.07613           0.3345           0.8902           2.217
##      area_se smoothness_se compactness_se concavity_se concave.points_se
## 842302      153.40      0.006399      0.04904      0.05373      0.01587
## 842517       74.08      0.005225      0.01308      0.01860      0.01340
## 84300903      94.03      0.006150      0.04006      0.03832      0.02058
## 84348301      27.23      0.009110      0.07458      0.05661      0.01867
## 84358402      94.44      0.011490      0.02461      0.05688      0.01885
## 843786      27.19      0.007510      0.03345      0.03672      0.01137
```

```
##      symmetry_se fractal_dimension_se radius_worst texture_worst
## 842302      0.03003      0.006193      25.38      17.33
## 842517      0.01389      0.003532      24.99      23.41
## 84300903      0.02250      0.004571      23.57      25.53
## 84348301      0.05963      0.009208      14.91      26.50
## 84358402      0.01756      0.005115      22.54      16.67
## 843786      0.02165      0.005082      15.47      23.75
##      perimeter_worst area_worst smoothness_worst compactness_worst
## 842302      184.60      2019.0      0.1622      0.6656
## 842517      158.80      1956.0      0.1238      0.1866
## 84300903      152.50      1709.0      0.1444      0.4245
## 84348301      98.87      567.7      0.2098      0.8663
## 84358402      152.20      1575.0      0.1374      0.2050
## 843786      103.40      741.6      0.1791      0.5249
##      concavity_worst concave.points_worst symmetry_worst
## 842302      0.7119      0.2654      0.4601
## 842517      0.2416      0.1860      0.2750
## 84300903      0.4504      0.2430      0.3613
## 84348301      0.6869      0.2575      0.6638
## 84358402      0.4000      0.1625      0.2364
## 843786      0.5355      0.1741      0.3985
##      fractal_dimension_worst
## 842302      0.11890
## 842517      0.08902
## 84300903      0.08758
## 84348301      0.17300
## 84358402      0.07678
## 843786      0.12440
```

The `wisc.df$diagnosis` gives us the actual “answer” to whether a sample is benign or malignant, so we will exclude it from our data analysis

```
#Create new data frame that omits diagnosis column
wisc.data <- wisc.df[,-1]
```

```
#Create vector w/ diagnoses
diagnosis <- factor(wisc.df$diagnosis)
diagnosis
```

```
##      [1] M M M M M M M M M M M M M M M M M M M M M B B B M M M M M M M M M M M
##      [38] B M M M M M M M M M B B B B B M M B M M B B B B M B M M B B B B M B M M
##      [75] B M B M M B B B M M B M M M B B B M B B M M B B B M M B B B B M B B B
##     [112] B B B B B B M M M B M M B B B M M B M B M M B M M B B M B B B M B B B M B
##     [149] B B B B B B B B M B B B B M M B M B B M M B B M M B B B B M B B M M M B M
##     [186] B M B B B M B B M M B M M M M B M M M B M B B M B M M M M B B M M B B
##     [223] B M B B B B B M M B B M B B M M B M B B B B M B B B B B M B M M M M M M
##     [260] M M M M M M M B B B B B B M B M B B M B B B M B M M B B B B B B B B B B
##     [297] B M B B M B M B B B B B B B B B B B B M B B B M B M B B B B M M M B B
##     [334] B B M B M B M B B B M B B B B B B B M M M B B B B B B B B B B M M B M M
##     [371] M B M M B B B B B M B B B B B M B B B M B B M M B B B B B M B B B B B B
##     [408] B M B B B B B M B B M B B B B B B B B B B B M B M M B M B B B B B M B B
##     [445] M B M B B M B M B B B B B B B B M M B B B B B B M B B B B B B B B M B
##     [482] B B B B B B M B M B B M B B B B B M M B M B M B B B B M B B M B M B M M
```

```
## [519] B B B M B B B B B B B B B B M B M M B B B B B B B B B B B B B
## [556] B B B B B B B M M M M M M B
## Levels: B M
```

Q1. How many observations are in this dataset? There are 590 observations in wisc.data

```
str(wisc.data)
```

```
## 'data.frame': 569 obs. of 30 variables:
## $ radius_mean : num 18 20.6 19.7 11.4 20.3 ...
## $ texture_mean : num 10.4 17.8 21.2 20.4 14.3 ...
## $ perimeter_mean : num 122.8 132.9 130 77.6 135.1 ...
## $ area_mean : num 1001 1326 1203 386 1297 ...
## $ smoothness_mean : num 0.1184 0.0847 0.1096 0.1425 0.1003 ...
## $ compactness_mean : num 0.2776 0.0786 0.1599 0.2839 0.1328 ...
## $ concavity_mean : num 0.3001 0.0869 0.1974 0.2414 0.198 ...
## $ concave.points_mean : num 0.1471 0.0702 0.1279 0.1052 0.1043 ...
## $ symmetry_mean : num 0.242 0.181 0.207 0.26 0.181 ...
## $ fractal_dimension_mean : num 0.0787 0.0567 0.06 0.0974 0.0588 ...
## $ radius_se : num 1.095 0.543 0.746 0.496 0.757 ...
## $ texture_se : num 0.905 0.734 0.787 1.156 0.781 ...
## $ perimeter_se : num 8.59 3.4 4.58 3.44 5.44 ...
## $ area_se : num 153.4 74.1 94 27.2 94.4 ...
## $ smoothness_se : num 0.0064 0.00522 0.00615 0.00911 0.01149 ...
## $ compactness_se : num 0.049 0.0131 0.0401 0.0746 0.0246 ...
## $ concavity_se : num 0.0537 0.0186 0.0383 0.0566 0.0569 ...
## $ concave.points_se : num 0.0159 0.0134 0.0206 0.0187 0.0188 ...
## $ symmetry_se : num 0.03 0.0139 0.0225 0.0596 0.0176 ...
## $ fractal_dimension_se : num 0.00619 0.00353 0.00457 0.00921 0.00511 ...
## $ radius_worst : num 25.4 25 23.6 14.9 22.5 ...
## $ texture_worst : num 17.3 23.4 25.5 26.5 16.7 ...
## $ perimeter_worst : num 184.6 158.8 152.5 98.9 152.2 ...
## $ area_worst : num 2019 1956 1709 568 1575 ...
## $ smoothness_worst : num 0.162 0.124 0.144 0.21 0.137 ...
## $ compactness_worst : num 0.666 0.187 0.424 0.866 0.205 ...
## $ concavity_worst : num 0.712 0.242 0.45 0.687 0.4 ...
## $ concave.points_worst : num 0.265 0.186 0.243 0.258 0.163 ...
## $ symmetry_worst : num 0.46 0.275 0.361 0.664 0.236 ...
## $ fractal_dimension_worst: num 0.1189 0.089 0.0876 0.173 0.0768 ...
```

Q2. How many of the observations have a malignant diagnosis? 212 of the observations have a malignant diagnosis.

```
table(diagnosis)
```

```
## diagnosis
## B M
## 357 212
```

Q3. How many variables/features in the data are suffixed with `_mean`? 10 variables in the data are suffixed with `_mean`.

```
#Not quite sure how to reason this out but google helped
meanvect <- names(wisc.data)[grep("_mean", names(wisc.data))]
length(meanvect)
```

```
## [1] 10
```

2. Principal Component Analysis

```
#Check column means and st dev
colMeans(wisc.data)
```

```
##          radius_mean      texture_mean      perimeter_mean
##      1.412729e+01      1.928965e+01      9.196903e+01
##          area_mean      smoothness_mean      compactness_mean
##      6.548891e+02      9.636028e-02      1.043410e-01
##      concavity_mean      concave.points_mean      symmetry_mean
##      8.879932e-02      4.891915e-02      1.811619e-01
## fractal_dimension_mean      radius_se      texture_se
##      6.279761e-02      4.051721e-01      1.216853e+00
##      perimeter_se      area_se      smoothness_se
##      2.866059e+00      4.033708e+01      7.040979e-03
##      compactness_se      concavity_se      concave.points_se
##      2.547814e-02      3.189372e-02      1.179614e-02
##      symmetry_se      fractal_dimension_se      radius_worst
##      2.054230e-02      3.794904e-03      1.626919e+01
##      texture_worst      perimeter_worst      area_worst
##      2.567722e+01      1.072612e+02      8.805831e+02
##      smoothness_worst      compactness_worst      concavity_worst
##      1.323686e-01      2.542650e-01      2.721885e-01
##      concave.points_worst      symmetry_worst      fractal_dimension_worst
##      1.146062e-01      2.900756e-01      8.394582e-02
```

```
apply(wisc.data, 2, sd)
```

```
##          radius_mean      texture_mean      perimeter_mean
##      3.524049e+00      4.301036e+00      2.429898e+01
##          area_mean      smoothness_mean      compactness_mean
##      3.519141e+02      1.406413e-02      5.281276e-02
##      concavity_mean      concave.points_mean      symmetry_mean
##      7.971981e-02      3.880284e-02      2.741428e-02
## fractal_dimension_mean      radius_se      texture_se
##      7.060363e-03      2.773127e-01      5.516484e-01
##      perimeter_se      area_se      smoothness_se
##      2.021855e+00      4.549101e+01      3.002518e-03
##      compactness_se      concavity_se      concave.points_se
##      1.790818e-02      3.018606e-02      6.170285e-03
##      symmetry_se      fractal_dimension_se      radius_worst
##      8.266372e-03      2.646071e-03      4.833242e+00
##      texture_worst      perimeter_worst      area_worst
```

```
##          6.146258e+00          3.360254e+01          5.693570e+02
##      smoothness_worst      compactness_worst      concavity_worst
##          2.283243e-02          1.573365e-01          2.086243e-01
##      concave.points_worst      symmetry_worst      fractal_dimension_worst
##          6.573234e-02          6.186747e-02          1.806127e-02
```

```
#Perform PCA on wisc.data, use t() for transpose of the data, use scaling
wisc.pr <- prcomp(wisc.data, scale=TRUE)
```

```
#Summary of the PCA results
summary(wisc.pr)
```

```
## Importance of components:
##          PC1      PC2      PC3      PC4      PC5      PC6      PC7
## Standard deviation  3.6444 2.3857 1.67867 1.40735 1.28403 1.09880 0.82172
## Proportion of Variance 0.4427 0.1897 0.09393 0.06602 0.05496 0.04025 0.02251
## Cumulative Proportion 0.4427 0.6324 0.72636 0.79239 0.84734 0.88759 0.91010
##          PC8      PC9      PC10     PC11     PC12     PC13     PC14
## Standard deviation  0.69037 0.6457 0.59219 0.5421 0.51104 0.49128 0.39624
## Proportion of Variance 0.01589 0.0139 0.01169 0.0098 0.00871 0.00805 0.00523
## Cumulative Proportion 0.92598 0.9399 0.95157 0.9614 0.97007 0.97812 0.98335
##          PC15     PC16     PC17     PC18     PC19     PC20     PC21
## Standard deviation  0.30681 0.28260 0.24372 0.22939 0.22244 0.17652 0.1731
## Proportion of Variance 0.00314 0.00266 0.00198 0.00175 0.00165 0.00104 0.0010
## Cumulative Proportion 0.98649 0.98915 0.99113 0.99288 0.99453 0.99557 0.9966
##          PC22     PC23     PC24     PC25     PC26     PC27     PC28
## Standard deviation  0.16565 0.15602 0.1344 0.12442 0.09043 0.08307 0.03987
## Proportion of Variance 0.00091 0.00081 0.0006 0.00052 0.00027 0.00023 0.00005
## Cumulative Proportion 0.99749 0.99830 0.9989 0.99942 0.99969 0.99992 0.99997
##          PC29     PC30
## Standard deviation  0.02736 0.01153
## Proportion of Variance 0.00002 0.00000
## Cumulative Proportion 1.00000 1.00000
```

Q4. From your results, what proportion of the original variance is captured by the first principal components (PC1)?

The proportion of the original variance captured by the first principal component (PC1) is 44.27%

Q5. How many principal components (PCs) are required to describe at least 70% of the original variance in the data?

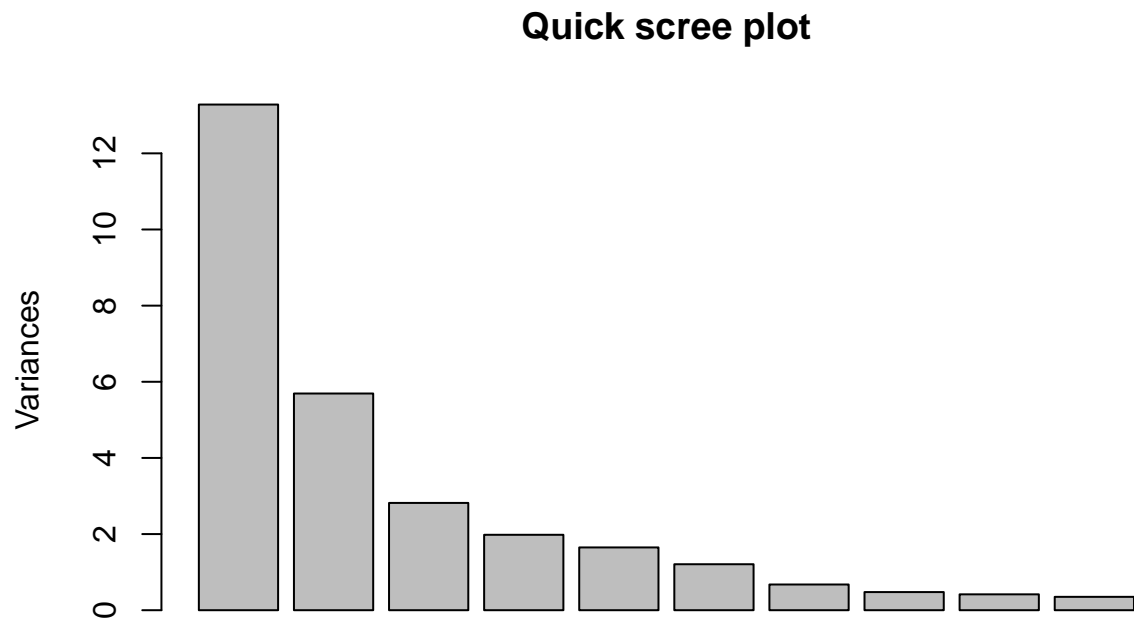
At least 3 PCs are required to describes at least 70% of the original variance in the data.

Q6. How many principal components (PCs) are required to describe at least 90% of the original variance in the data?

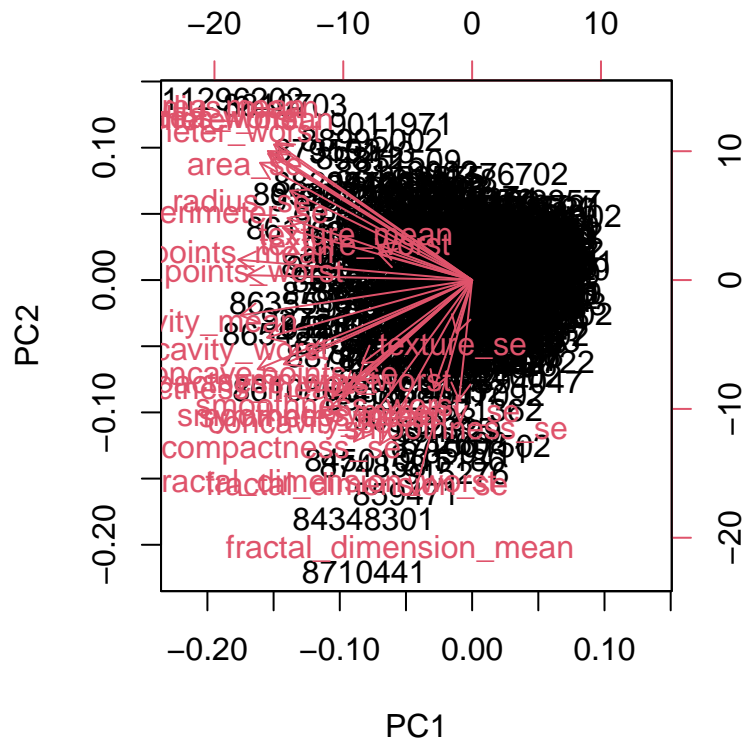
At least 7 PCs are required to describe at least 90% of the original variance in the data.

Interpreting PCA Results

```
#Scree plot  
plot(wisc.pr, main= "Quick scree plot")
```



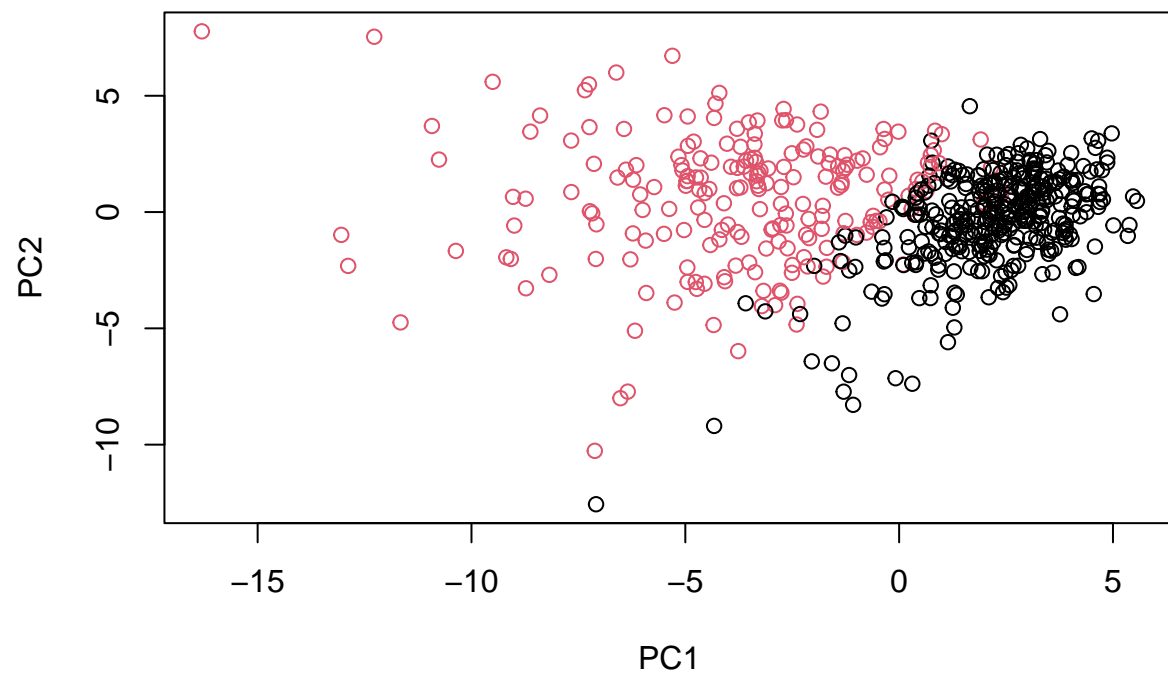
```
#Biplot  
biplot(wisc.pr)
```



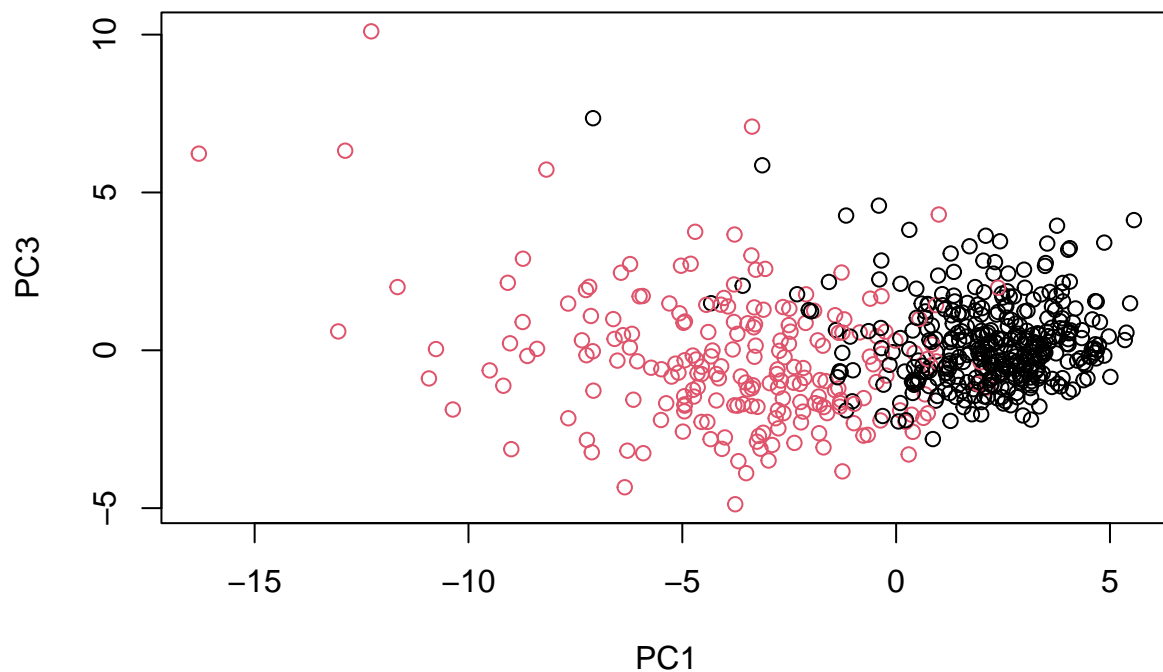
Q7. What stands out to you about this plot? Is it easy or difficult to understand? Why?

The biplot is basically impossible to interpret. It's unclear and you can't make any conclusions with assurance by looking at it.

```
#Scatterplot w/ PC1 & PC2
plot(wisc.pr$x[,1], wisc.pr$x[,2], col = diagnosis, xlab = "PC1", ylab = "PC2")
```



```
# Repeat for components 1 and 3  
plot(wisc.pr$x[,1], wisc.pr$x[,3], col = diagnosis,  
      xlab = "PC1", ylab = "PC3")
```

Q8. Generate a similar plot for principal components 1 and 3. What do you notice about these plots?

The plots are very similar however plotting PC1 vs PC3 shifts the plotted points downwards visually. Overall, there does not seem to be too much difference between the two plots.

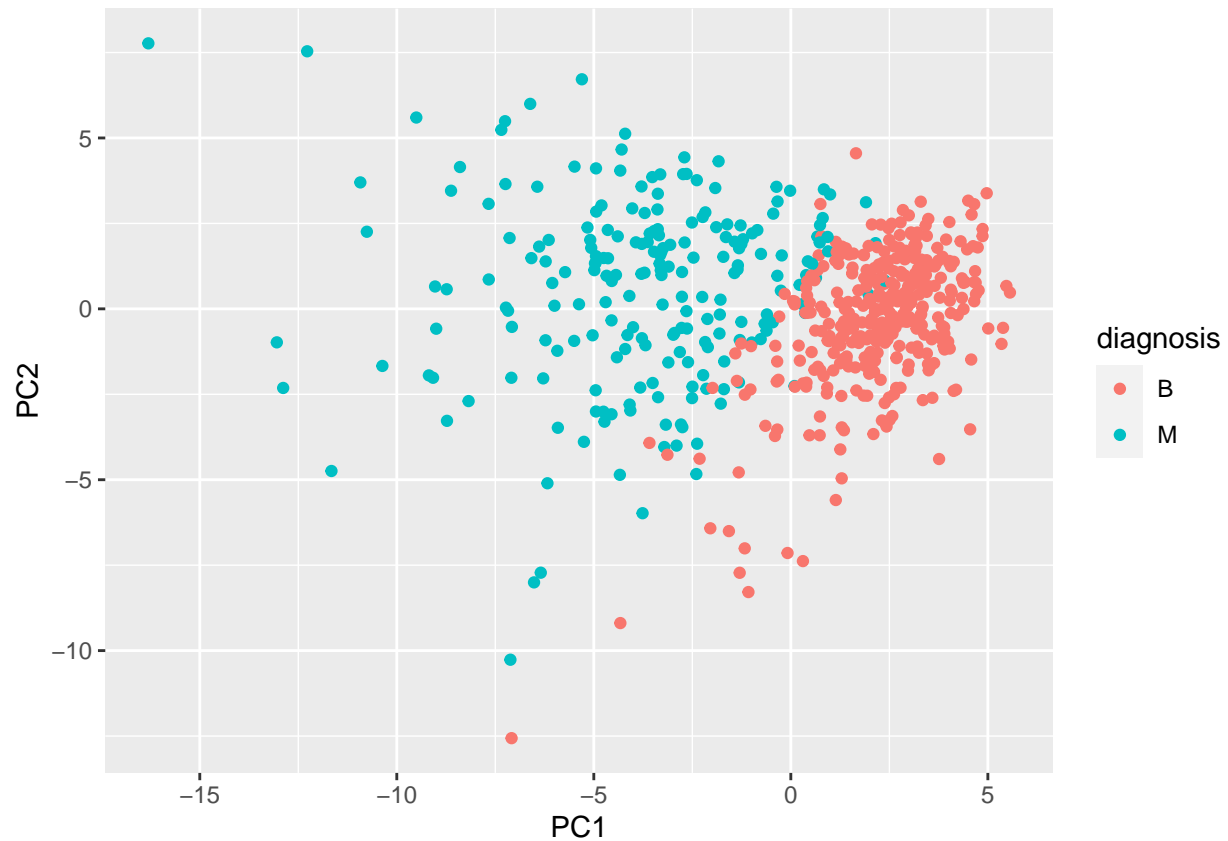
Plot with ggplot2

```
library(ggplot2)
```

ggplot uses data frames for input, and diagnosis vect must be converted to column

```
df <- as.data.frame(wisc.pr$x)
df$diagnosis <- diagnosis
```

```
#Scatter plot colored by dx
ggplot(df) + aes(PC1, PC2, col=diagnosis) + geom_point()
```



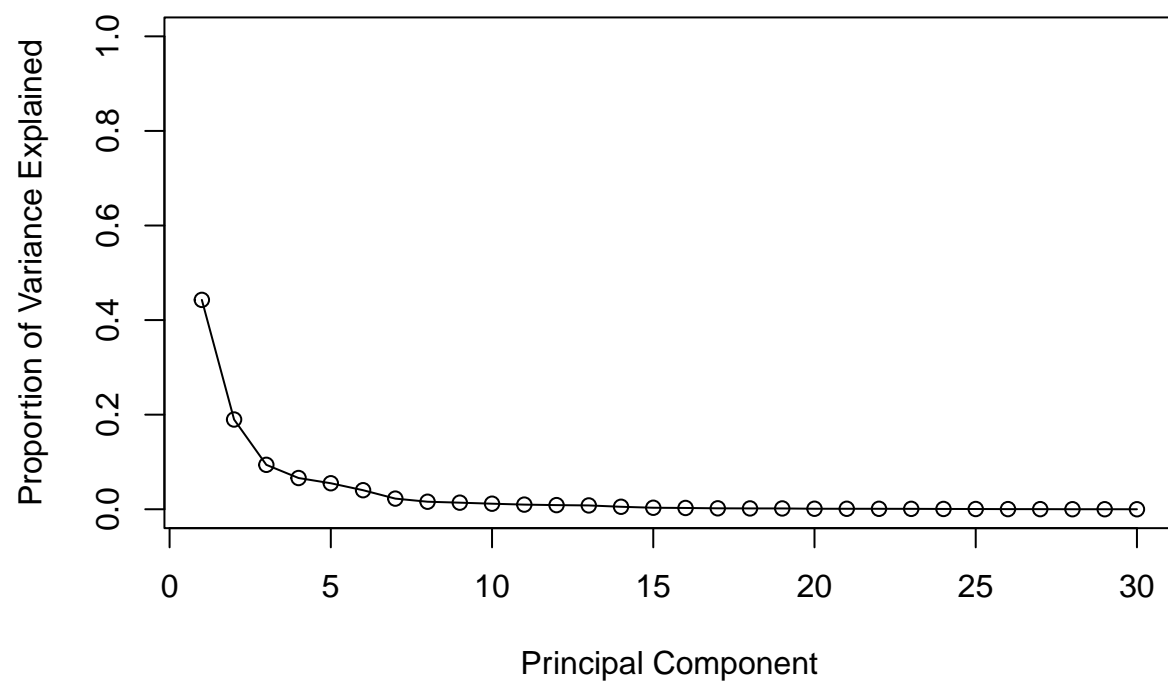
Variance explained

```
#The variance of e/ component
pr.var <- wisc.pr$sdev^2
head(pr.var)
```

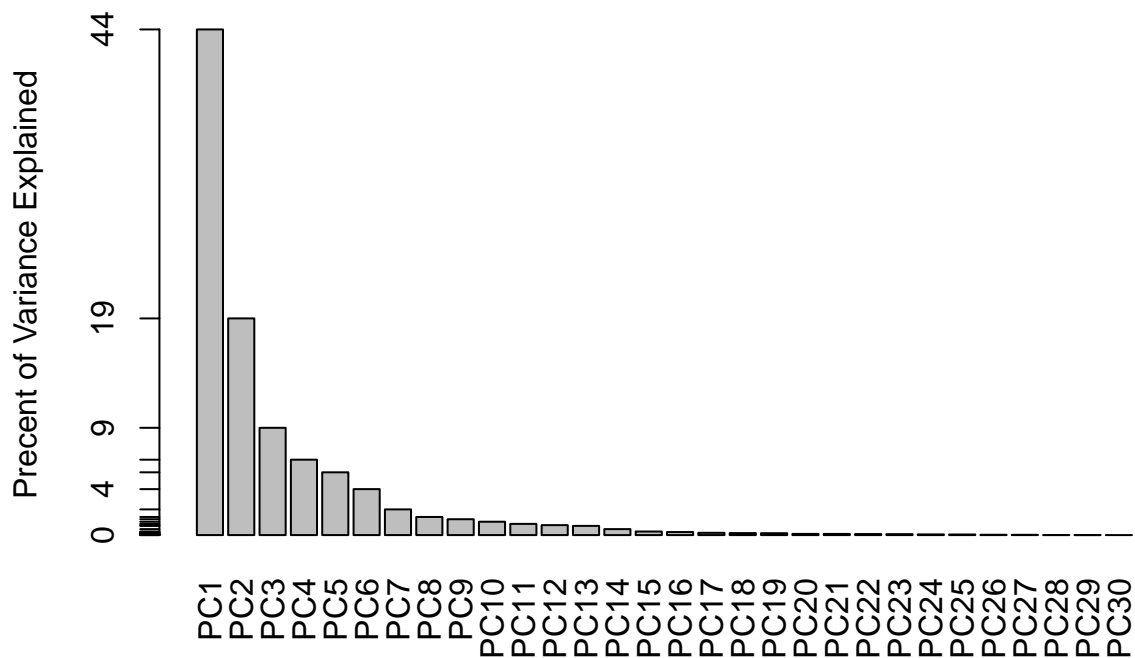
```
## [1] 13.281608  5.691355  2.817949  1.980640  1.648731  1.207357
```

```
#Variance explained by e/ PC
pve <- pr.var/sum(pr.var)
```

```
# Plot variance explained for each principal component
plot(pve, xlab = "Principal Component",
     ylab = "Proportion of Variance Explained",
     ylim = c(0, 1), type = "o")
```



```
# Alternative scree plot of the same data, note data driven y-axis
barplot(pve, ylab = "Precent of Variance Explained",
        names.arg=paste0("PC",1:length(pve)), las=2, axes = FALSE)
axis(2, at=pve, labels=round(pve,2)*100 )
```



Q9. For the first principal component, what is the component of the loading vector (i.e. `wisc.pr$rotation[,1]`) for the feature `concave.points_mean`?

It is -0.26085376

```
wisc.pr$rotation[,1]
```

```
##          radius_mean          texture_mean          perimeter_mean
##          -0.21890244          -0.10372458          -0.22753729
##          area_mean          smoothness_mean          compactness_mean
##          -0.22099499          -0.14258969          -0.23928535
##          concavity_mean          concave.points_mean          symmetry_mean
##          -0.25840048          -0.26085376          -0.13816696
## fractal_dimension_mean          radius_se          texture_se
##          -0.06436335          -0.20597878          -0.01742803
##          perimeter_se          area_se          smoothness_se
##          -0.21132592          -0.20286964          -0.01453145
##          compactness_se          concavity_se          concave.points_se
##          -0.17039345          -0.15358979          -0.18341740
##          symmetry_se          fractal_dimension_se          radius_worst
##          -0.04249842          -0.10256832          -0.22799663
##          texture_worst          perimeter_worst          area_worst
##          -0.10446933          -0.23663968          -0.22487053
##          smoothness_worst          compactness_worst          concavity_worst
##          -0.12795256          -0.21009588          -0.22876753
```

```
##      concave.points_worst      symmetry_worst fractal_dimension_worst
##      -0.25088597              -0.12290456      -0.13178394
```

Q10. What is the minimum number of principal components required to explain 80% of the variance of the data?

At least 5 principal components are required to explain 80% of the variance of the data.

Using alt programs for PCA (ade4 and factoextra)

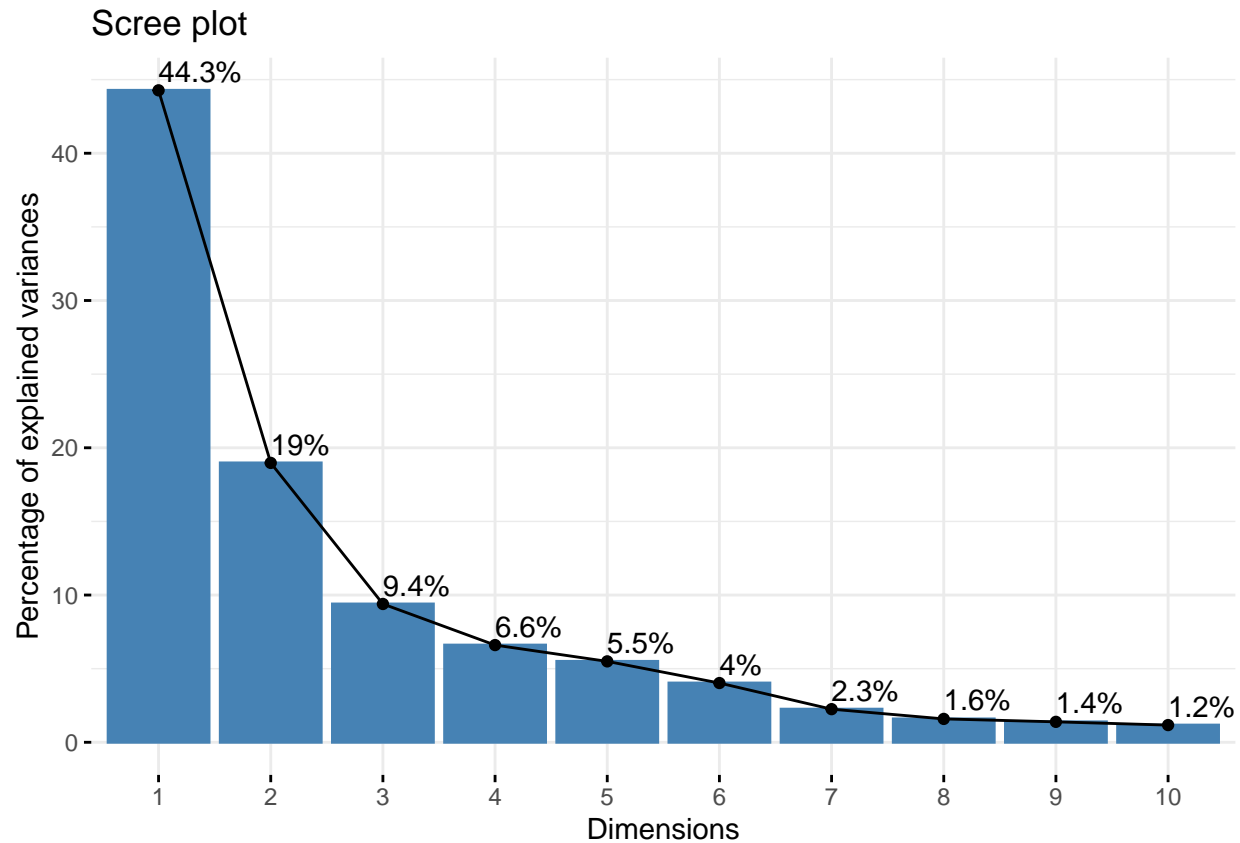
```
#Scree plot
library(ade4)
dudipca <- dudi.pca(df = wisc.data, center = TRUE, scale = TRUE, scannf = FALSE, nf = 30)
dudipca
```

```
## Duality diagramm
## class: pca dudi
## $call: dudi.pca(df = wisc.data, center = TRUE, scale = TRUE, scannf = FALSE,
##      nf = 30)
##
## $nf: 30 axis-components saved
## $rank: 30
## eigen values: 13.28 5.691 2.818 1.981 1.649 ...
##   vector length mode   content
## 1 $cw      30      numeric column weights
## 2 $lw     569      numeric row weights
## 3 $eig     30      numeric eigen values
##
##   data.frame nrow ncol content
## 1 $tab      569  30   modified array
## 2 $li      569  30   row coordinates
## 3 $l1      569  30   row normed scores
## 4 $co      30   30   column coordinates
## 5 $c1      30   30   column normed scores
## other elements: cent norm
```

```
library(factoextra)
```

```
## Welcome! Want to learn more? See two factoextra-related books at https://goo.gl/ve3WBa
```

```
#Scree plot
fviz_eig(dudipca, addlabels=TRUE)
```



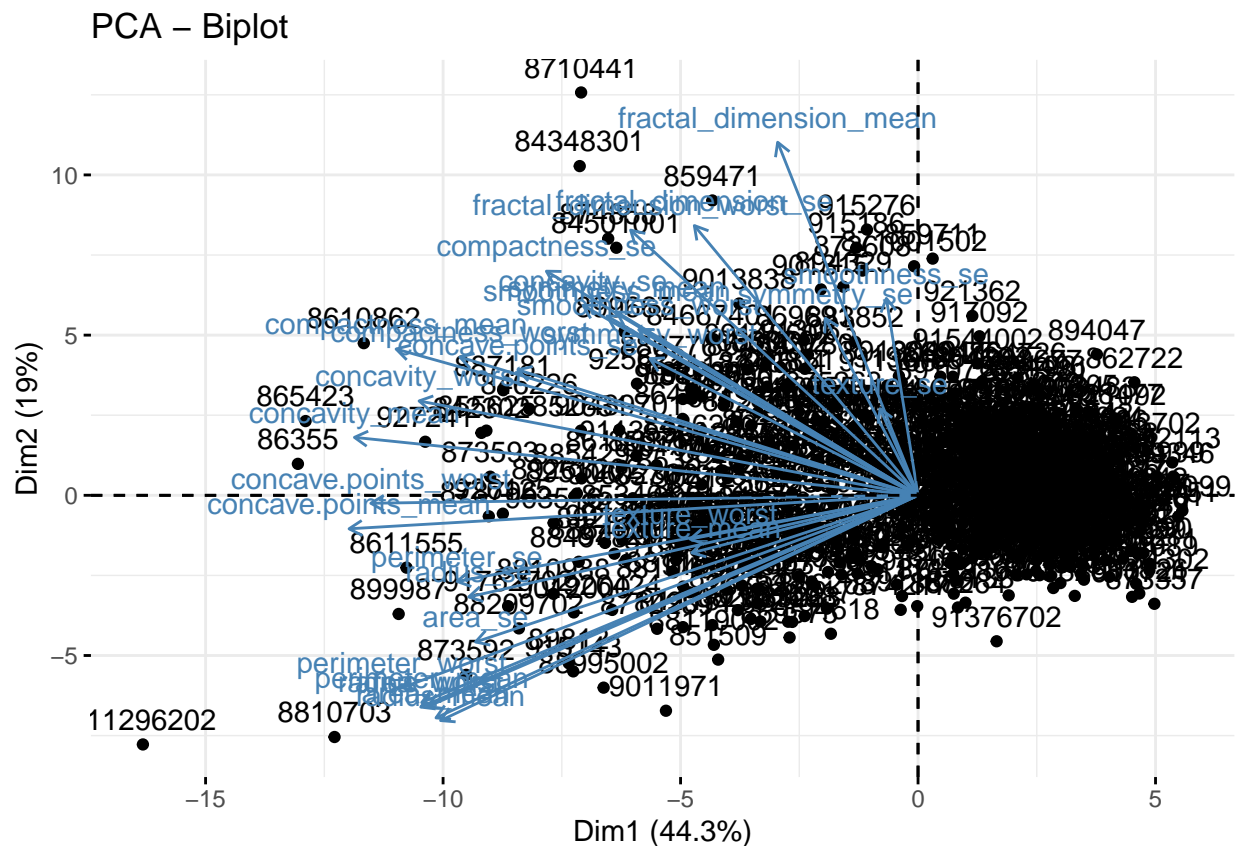
The above Scree plot shows PC1 containing 44.3% of variance, PC2 containing 19%, PC3 containing 9.4% and so on

```
get_eig(dudipca)
```

```
##          eigenvalue variance.percent cumulative.variance.percent
## Dim.1  1.328161e+01    4.427203e+01          44.27203
## Dim.2   5.691355e+00    1.897118e+01          63.24321
## Dim.3   2.817949e+00    9.393163e+00          72.63637
## Dim.4   1.980640e+00    6.602135e+00          79.23851
## Dim.5   1.648731e+00    5.495768e+00          84.73427
## Dim.6   1.207357e+00    4.024522e+00          88.75880
## Dim.7   6.752201e-01    2.250734e+00          91.00953
## Dim.8   4.766171e-01    1.588724e+00          92.59825
## Dim.9   4.168948e-01    1.389649e+00          93.98790
## Dim.10  3.506935e-01    1.168978e+00          95.15688
## Dim.11  2.939157e-01    9.797190e-01          96.13660
## Dim.12  2.611614e-01    8.705379e-01          97.00714
## Dim.13  2.413575e-01    8.045250e-01          97.81166
## Dim.14  1.570097e-01    5.233657e-01          98.33503
## Dim.15  9.413497e-02    3.137832e-01          98.64881
## Dim.16  7.986280e-02    2.662093e-01          98.91502
## Dim.17  5.939904e-02    1.979968e-01          99.11302
## Dim.18  5.261878e-02    1.753959e-01          99.28841
## Dim.19  4.947759e-02    1.649253e-01          99.45334
## Dim.20  3.115940e-02    1.038647e-01          99.55720
```

```
## Dim.21 2.997289e-02      9.990965e-02      99.65711
## Dim.22 2.743940e-02      9.146468e-02      99.74858
## Dim.23 2.434084e-02      8.113613e-02      99.82971
## Dim.24 1.805501e-02      6.018336e-02      99.88990
## Dim.25 1.548127e-02      5.160424e-02      99.94150
## Dim.26 8.177640e-03      2.725880e-02      99.96876
## Dim.27 6.900464e-03      2.300155e-02      99.99176
## Dim.28 1.589338e-03      5.297793e-03      99.99706
## Dim.29 7.488031e-04      2.496010e-03      99.99956
## Dim.30 1.330448e-04      4.434827e-04      100.00000
```

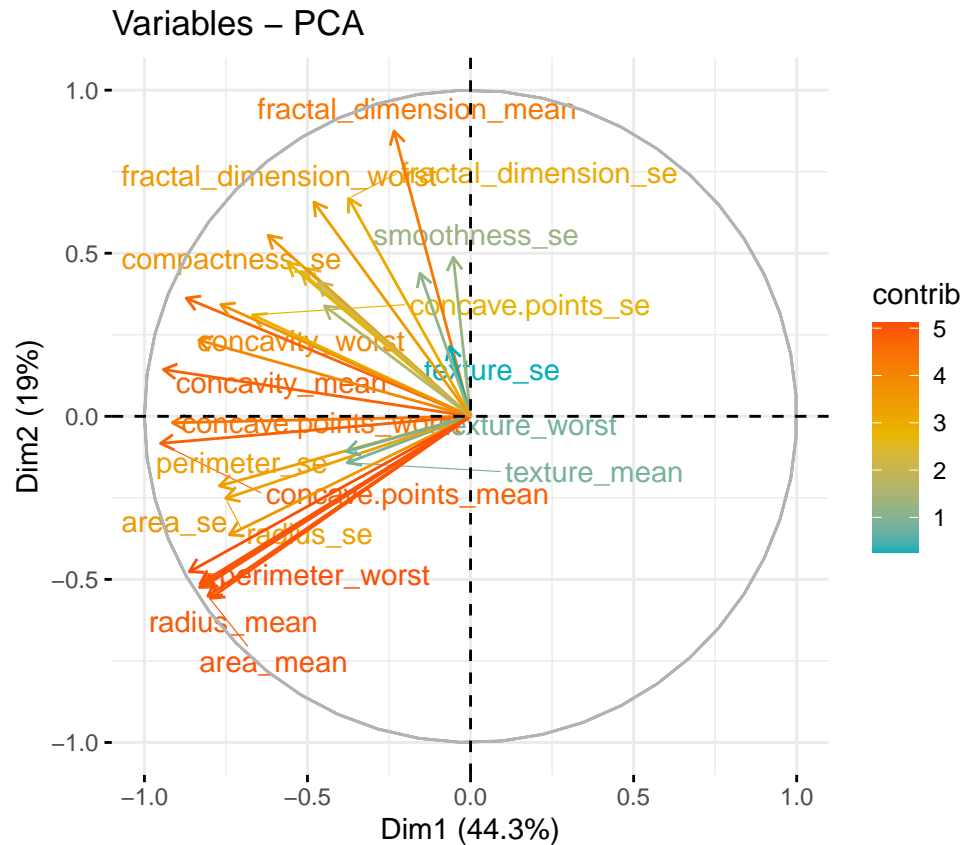
```
fviz_pca(dudipca)
```



The above biplot is extremely complicated

```
fviz_pca_var(dudipca, col.var="contrib",
             gradient.cols = c("#00AFBB", "#E7B800", "#FC4E07"),
             repel = TRUE # Avoid text overlapping
             )
```

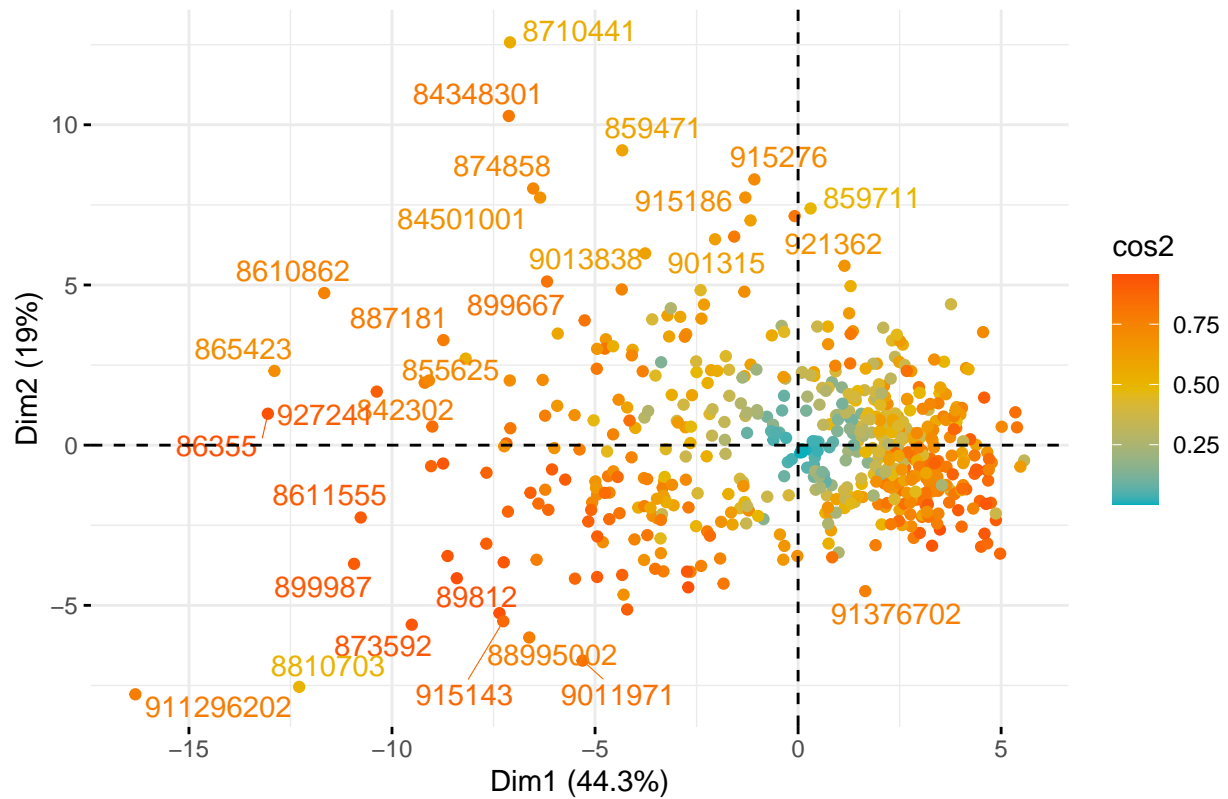
```
## Warning: ggrepel: 11 unlabeled data points (too many overlaps). Consider
## increasing max.overlaps
```



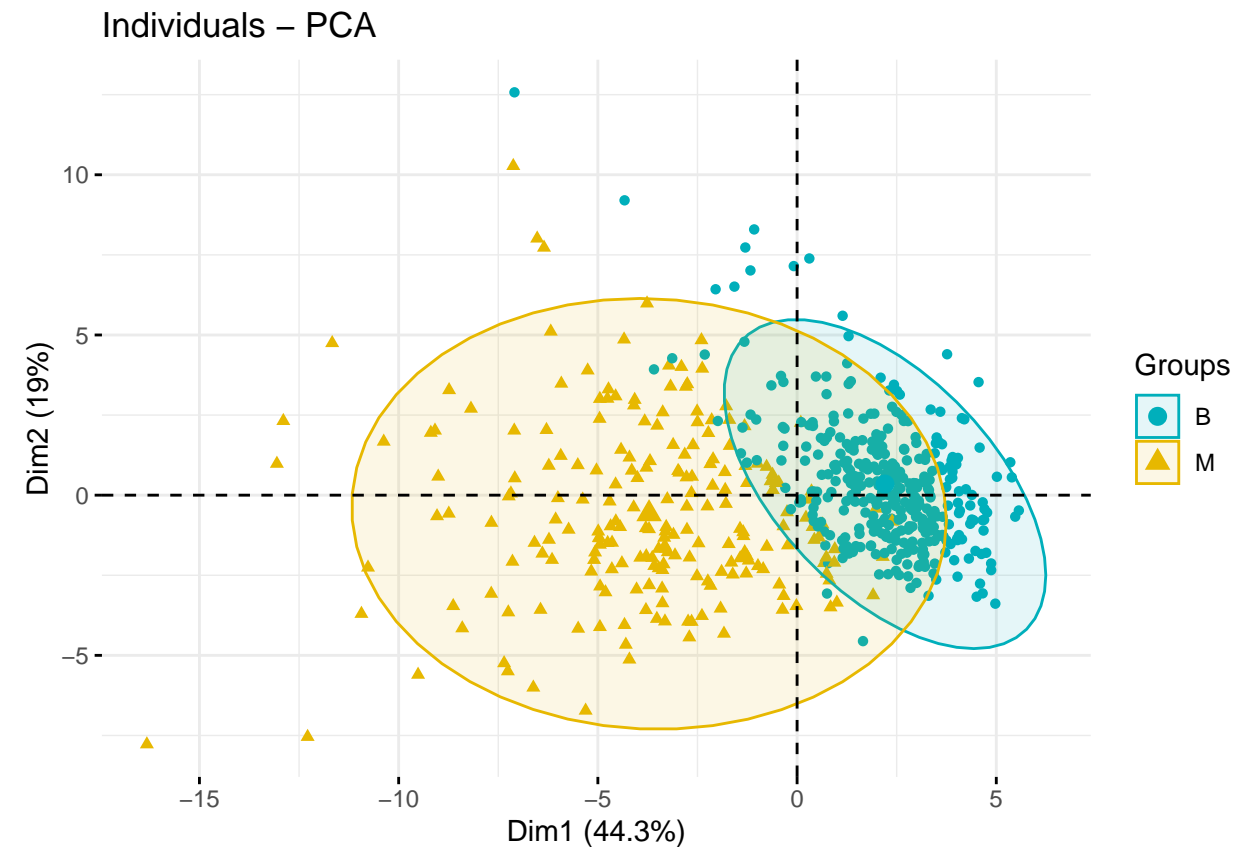
```
fviz_pca_ind(dudipca, col.ind = "cos2",
             gradient.cols = c("#00AFBB", "#E7B800", "#FC4E07"),
             repel = TRUE # Avoid text overlapping (slow if many points)
             )
```

```
## Warning: ggrepel: 540 unlabeled data points (too many overlaps). Consider
## increasing max.overlaps
```


Individuals – PCA



```
fviz_pca_ind(dudipca,
  label = "none", # hide individual labels
  habillage = wisc.df$diagnosis, # color by groups
  palette = c("#00AFBB", "#E7B800"),
  addEllipses = TRUE # Concentration ellipses
)
```



Hierarchical Clustering

```
#Scale the data
data.scaled <- scale(wisc.data)
```

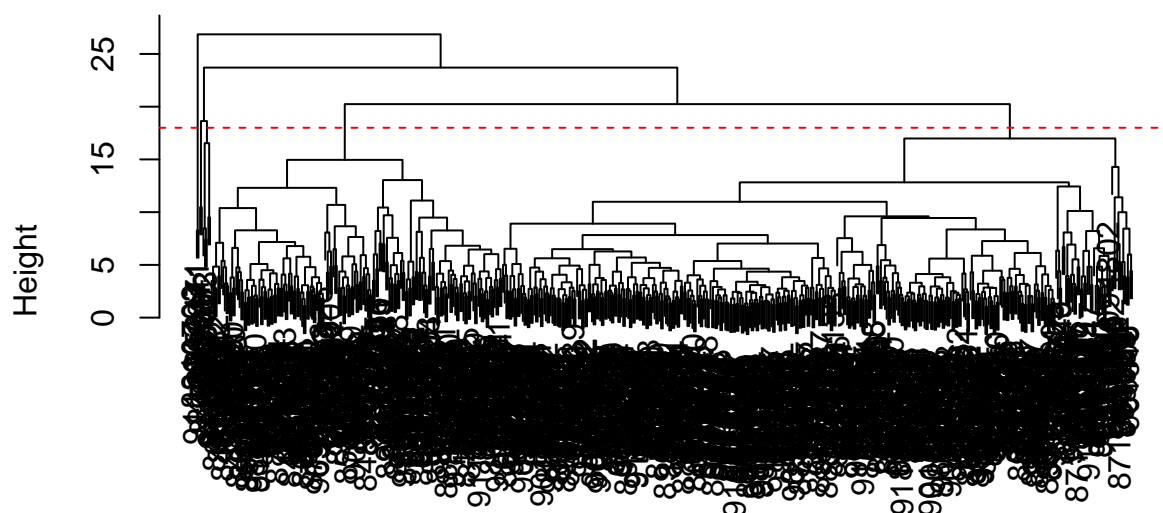
```
#Calc Euclidean dists b/w pairs of observations
data.dist <- dist(data.scaled)
```

```
wisc.hclust <- hclust(data.dist, method = "complete")
```

Q11. Using the `plot()` and `abline()` functions, what is the height at which the clustering model has 4 clusters? The height at which the clustering model has 4 clusters is approximately 18.

```
plot(wisc.hclust)
abline(h=18, col= "red", lty=2)
```

Cluster Dendrogram



```
data.dist
hclust(*, "complete")
```

```
#Cut the tree
wisc.hclust.clusters <- cutree(wisc.hclust, k=4)
```

```
#Compare our clusters to actual diagnoses
table(wisc.hclust.clusters, diagnosis)
```

```
##           diagnosis
## wisc.hclust.clusters  B  M
##           1  12 165
##           2   2   5
##           3 343  40
##           4   0   2
```

This showed that cluster 1 has mostly malignant cells, and cluster 3 has mostly benign cells.

```
wisc.hclust.clusters <- cutree(wisc.hclust, k=3)
table(wisc.hclust.clusters, diagnosis)
```

```
##           diagnosis
## wisc.hclust.clusters  B  M
##           1 355 205
##           2   2   5
##           3   0   2
```

```
wisc.hclust.clusters <- cutree(wisc.hclust, k=2)
table(wisc.hclust.clusters, diagnosis)
```

```
##              diagnosis
## wisc.hclust.clusters  B  M
##                   1 357 210
##                   2   0   2
```

```
wisc.hclust.clusters <- cutree(wisc.hclust, k=5)
table(wisc.hclust.clusters, diagnosis)
```

```
##              diagnosis
## wisc.hclust.clusters  B  M
##                   1  12 165
##                   2   0   5
##                   3 343  40
##                   4   2   0
##                   5   0   2
```

```
wisc.hclust.clusters <- cutree(wisc.hclust, k=6)
table(wisc.hclust.clusters, diagnosis)
```

```
##              diagnosis
## wisc.hclust.clusters  B  M
##                   1  12 165
##                   2   0   5
##                   3 331  39
##                   4   2   0
##                   5  12   1
##                   6   0   2
```

```
wisc.hclust.clusters <- cutree(wisc.hclust, k=10)
table(wisc.hclust.clusters, diagnosis)
```

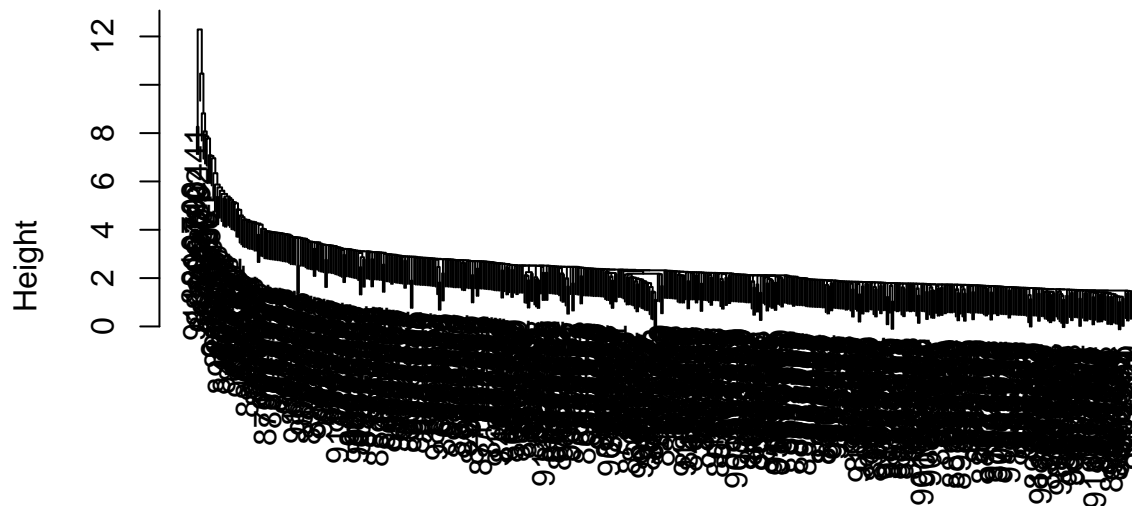
```
##              diagnosis
## wisc.hclust.clusters  B  M
##                   1  12  86
##                   2   0  59
##                   3   0   3
##                   4 331  39
##                   5   0  20
##                   6   2   0
##                   7  12   0
##                   8   0   2
##                   9   0   2
##                  10   0   1
```

Q12. Can you find a better cluster vs diagnoses match by cutting into a different number of clusters between 2 and 10?

No, 4 clusters seems to be the best match.

```
#Create plots using, single, complete, average, and ward.D2  
plot(hclust(data.dist, method = "single"))
```

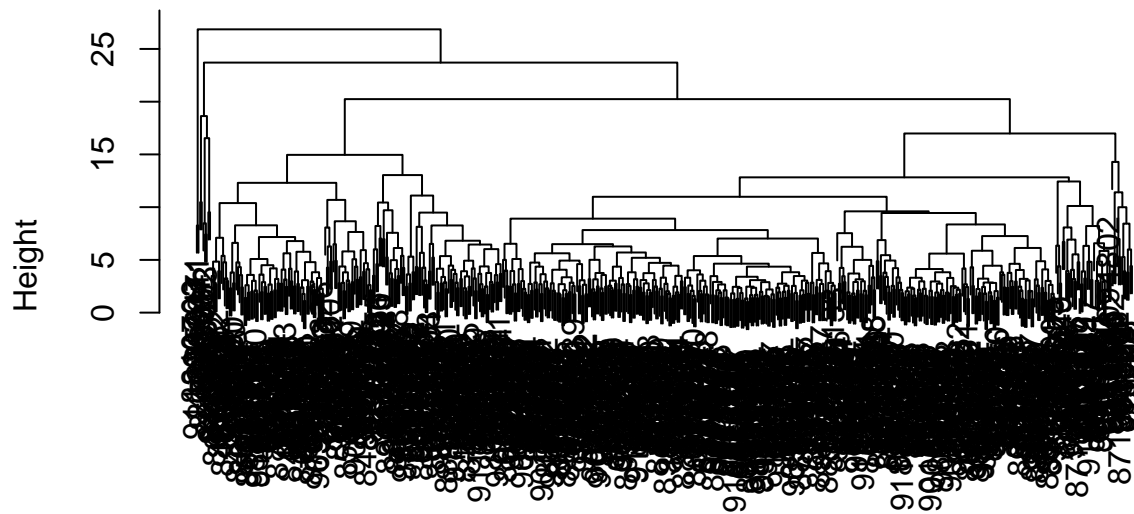
Cluster Dendrogram



data.dist
hclust (*, "single")

```
plot(hclust(data.dist, method = "complete"))
```

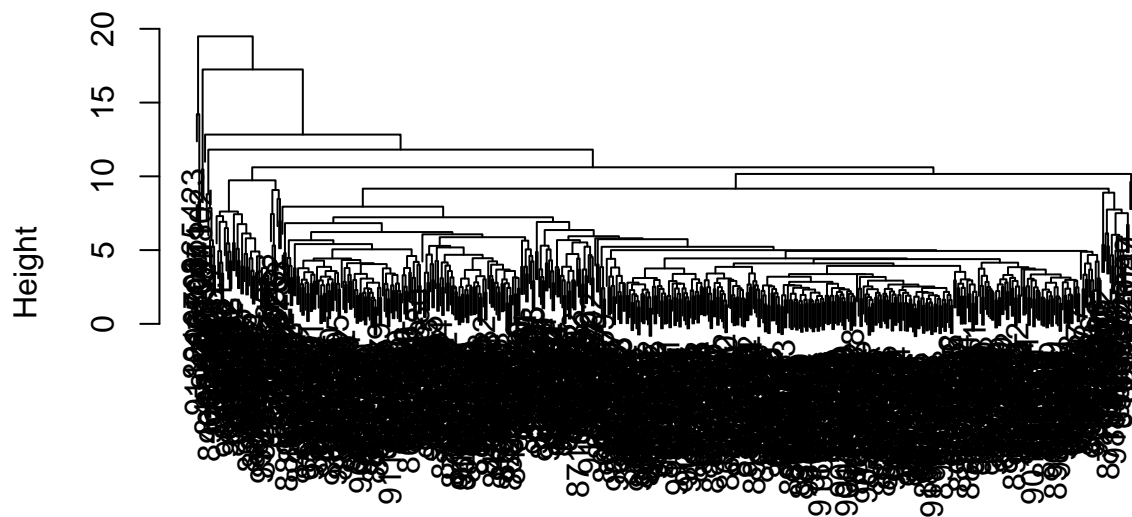
Cluster Dendrogram



data.dist
hclust (*, "complete")

```
plot(hclust(data.dist, method = "average"))
```

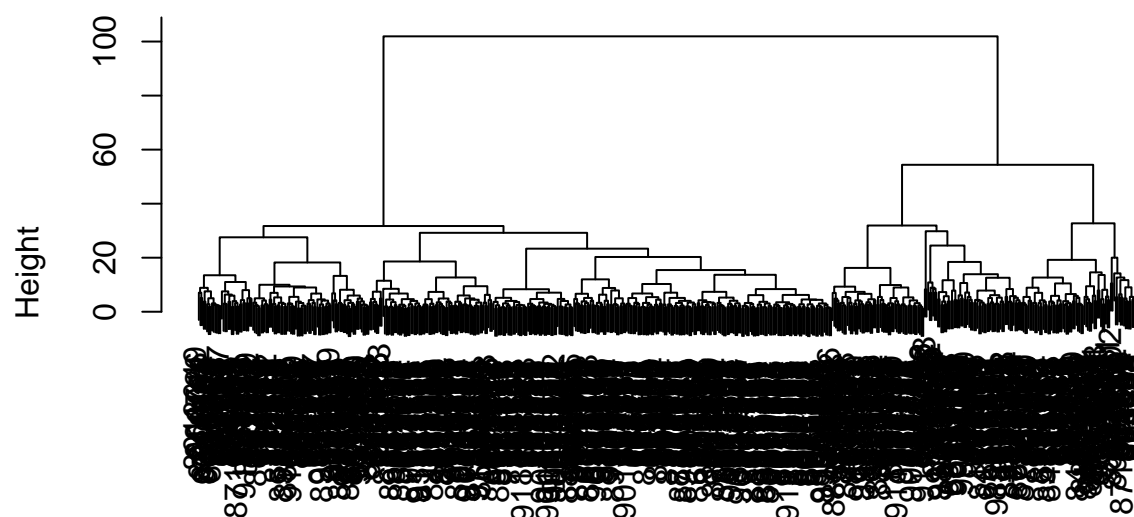
Cluster Dendrogram



data.dist
hclust (*, "average")

```
plot(hclust(data.dist, method = "ward.D2"))
```

Cluster Dendrogram



data.dist
hclust (*, "ward.D2")

Q13. Which method gives your favorite results for the same data.dist dataset? Explain your reasoning.

Using the ward.D2 method seems to give the “cleanest” looking dendrogram. The runner up is “complete.”

K Means Clustering

```
wisc.km <- kmeans(wisc.data, centers=2, nstart = 20)
```

```
table(wisc.km$cluster, diagnosis)
```

```
##      diagnosis
##      B      M
##  1     1   130
##  2   356    82
```

Q14. How well does k-means separate the two diagnoses? How does it compare to your hclust results?

The two diagnoses seem to have an obvious/significant split. It is overall similar to the hclust results when k=4.


```
table(wisc.km$cluster, cutree(wisc.hclust, k=4))
```

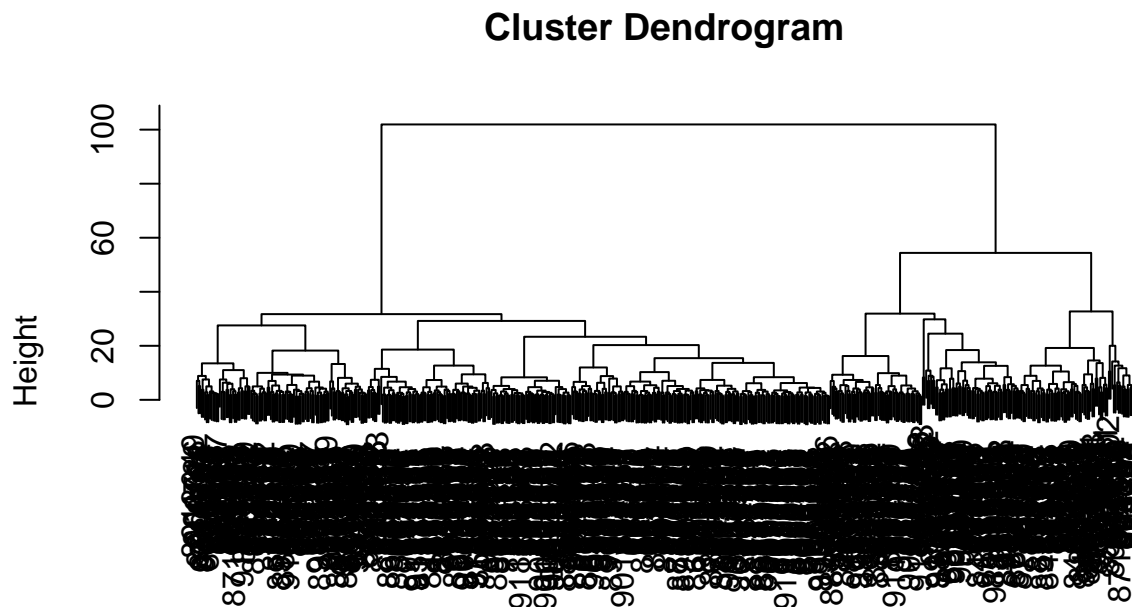
```
##
##      1    2    3    4
##  1 109    2   18    2
##  2   68    5  365    0
```

Combining Methods

Clustering on PCA Results

```
wisc.pr.hclust <- (hclust(data.dist, method = "ward.D2"))
```

```
plot(wisc.pr.hclust)
```



data.dist
hclust (*, "ward.D2")

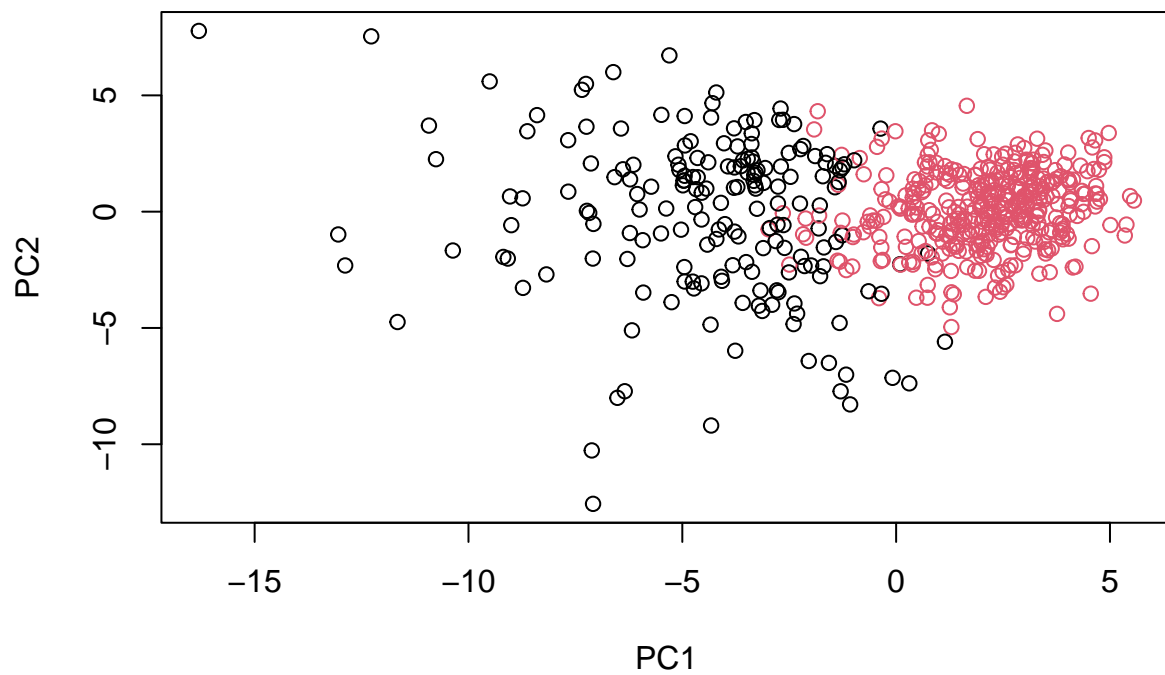
```
grps <- cutree(wisc.pr.hclust, k=2)
table(grps)
```

```
## grps
##  1    2
## 184 385
```

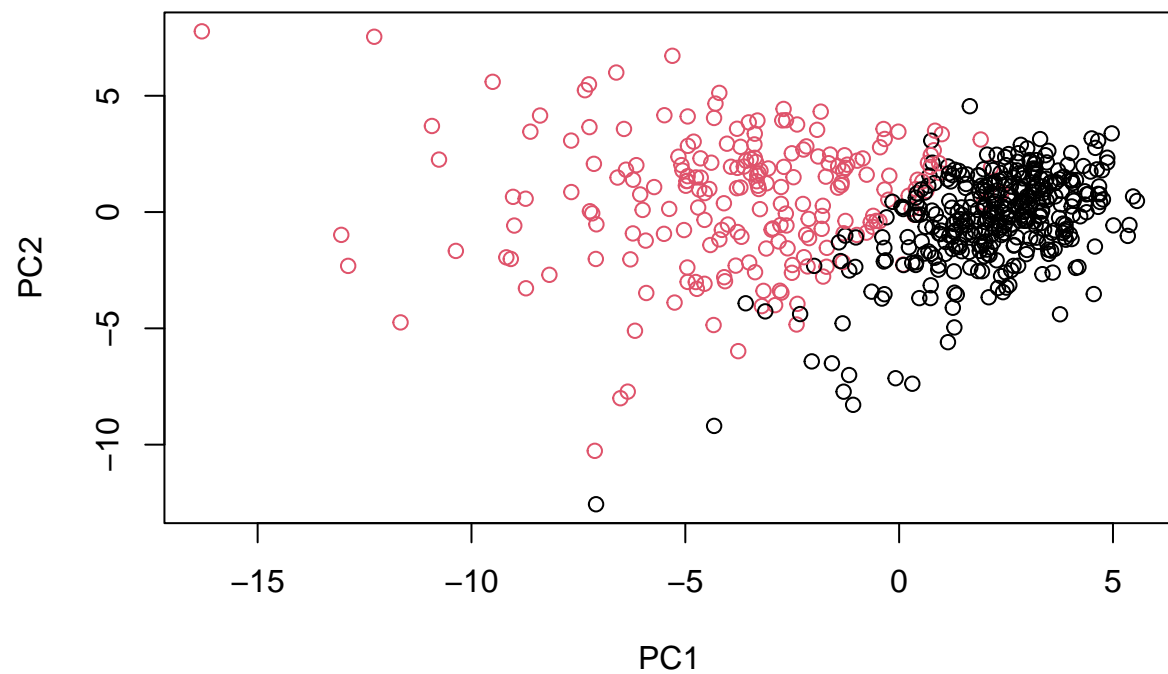
```
table(grps, diagnosis)
```

```
##      diagnosis  
## grps   B    M  
##    1  20 164  
##    2 337  48
```

```
plot(wisc.pr$x[,1:2], col=grps)
```



```
plot(wisc.pr$x[,1:2], col=diagnosis)
```



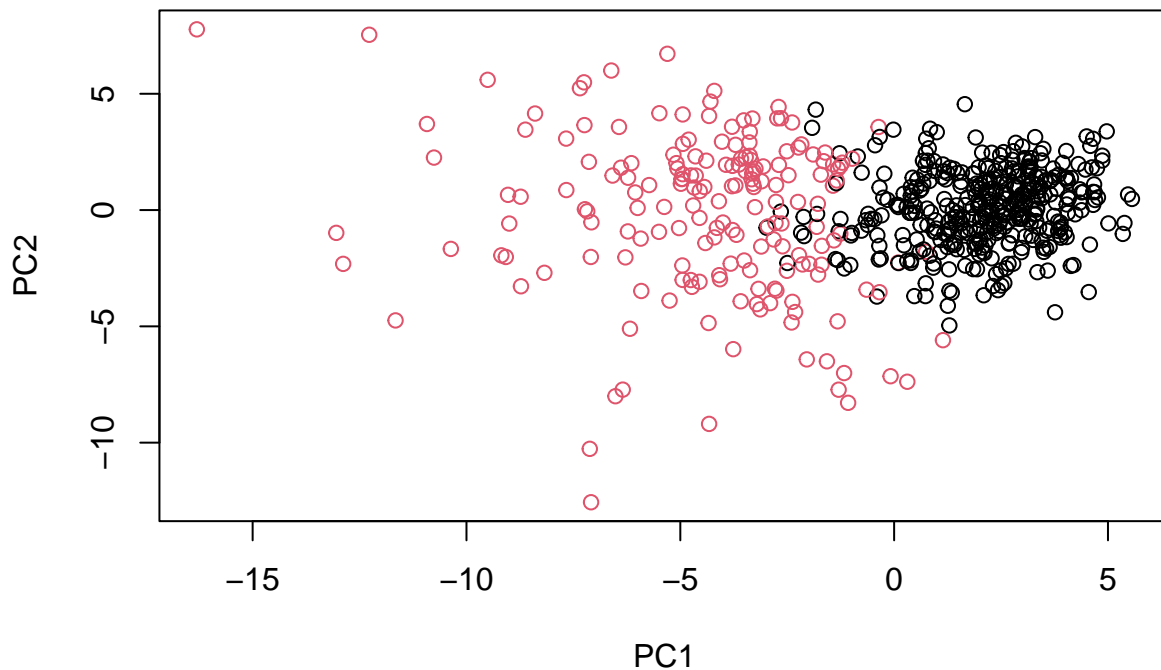
```
g <- as.factor(grps)
levels(g)
```

```
## [1] "1" "2"
```

```
g <- relevel(g,2)
levels(g)
```

```
## [1] "2" "1"
```

```
#plot w/ reordered factor
plot(wisc.pr$x[,1:2], col=g)
```



```
#clustering w/ 1st 7 PCs
wisc.pr.hclust.clusters <- cutree(wisc.pr.hclust, k=2)
```

```
table(wisc.pr.hclust.clusters, diagnosis)
```

```
##              diagnosis
## wisc.pr.hclust.clusters  B  M
##              1  20 164
##              2 337  48
```

Q15. How well does the newly created model with two clusters separate out the two diagnoses?

The two diagnoses seem to be separated significantly by using the new model with two clusters.

Q16. How well do the k-means and hierarchical clustering models you created in previous sections (i.e. before PCA) do in terms of separating the diagnoses? Again, use the `table()` function to compare the output of each model (`wisc.km$cluster` and `wisc.hclust.clusters`) with the vector containing the actual diagnoses.

The k-means and clustering models are roughly good at separating the diagnoses but they give ballpark results.

```
table(diagnosis)
```

```
## diagnosis
##      B      M
## 357 212
```

```
table(cutree(wisc.hclust, k=4), diagnosis)
```

```
##      diagnosis
##           B      M
##      1 12 165
##      2   2   5
##      3 343  40
##      4   0   2
```

```
table(wisc.km$cluster, diagnosis)
```

```
##      diagnosis
##           B      M
##      1   1 130
##      2 356  82
```

Sensitivity/Specificity

Sensitivity refers to a test's ability to correctly detect ill patients who do have the condition. In our example here the sensitivity is the total number of samples in the cluster identified as predominantly malignant (cancerous) divided by the total number of known malignant samples. In other words: $TP/(TP+FN)$.

Specificity relates to a test's ability to correctly reject healthy patients without a condition. In our example specificity is the proportion of benign (not cancerous) samples in the cluster identified as predominantly benign that are known to be benign. In other words: $TN/(TN+FP)$.

```
#hclust
```

```
#sensitivity  
165/212
```

```
## [1] 0.7783019
```

```
#Specificity  
343/357
```

```
## [1] 0.9607843
```

```
#k-means
```

```
#Sensitivity  
130/212
```

```
## [1] 0.6132075
```

```
#Specificity
356/357
```

```
## [1] 0.9971989
```

Q17. Which of your analysis procedures resulted in a clustering model with the best specificity?
How about sensitivity?

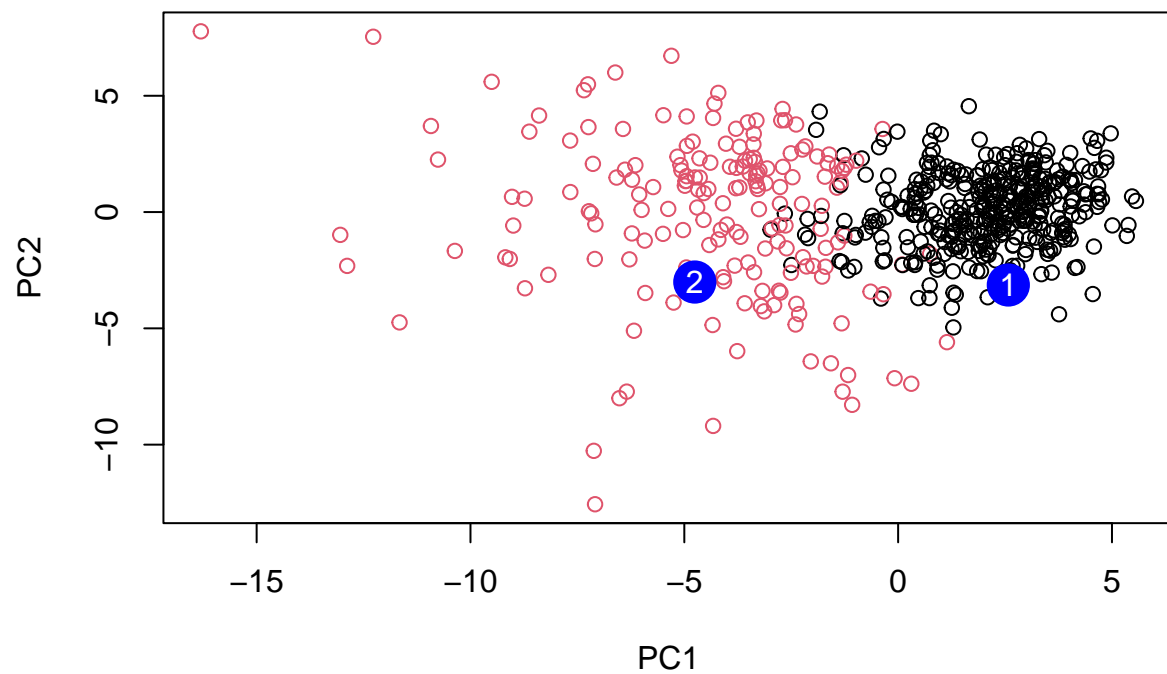
K-means gives the best specificity while hclustering gives the best sensitivity.

Prediction

```
#url <- "new_samples.csv"
url <- "https://tinyurl.com/new-samples-CSV"
new <- read.csv(url)
npc <- predict(wisc.pr, newdata=new)
npc
```

```
##          PC1          PC2          PC3          PC4          PC5          PC6          PC7
## [1,]  2.576616 -3.135913  1.3990492 -0.7631950  2.781648 -0.8150185 -0.3959098
## [2,] -4.754928 -3.009033 -0.1660946 -0.6052952 -1.140698 -1.2189945  0.8193031
##          PC8          PC9          PC10          PC11          PC12          PC13          PC14
## [1,] -0.2307350 0.1029569 -0.9272861 0.3411457  0.375921 0.1610764 1.187882
## [2,] -0.3307423 0.5281896 -0.4855301 0.7173233 -1.185917 0.5893856 0.303029
##          PC15          PC16          PC17          PC18          PC19          PC20
## [1,] 0.3216974 -0.1743616 -0.07875393 -0.11207028 -0.08802955 -0.2495216
## [2,] 0.1299153  0.1448061 -0.40509706  0.06565549  0.25591230 -0.4289500
##          PC21          PC22          PC23          PC24          PC25          PC26
## [1,] 0.1228233 0.09358453 0.08347651  0.1223396  0.02124121 0.078884581
## [2,] -0.1224776 0.01732146 0.06316631 -0.2338618 -0.20755948 -0.009833238
##          PC27          PC28          PC29          PC30
## [1,] 0.220199544 -0.02946023 -0.015620933  0.005269029
## [2,] -0.001134152 0.09638361  0.002795349 -0.019015820
```

```
plot(wisc.pr$x[,1:2], col=g)
points(npc[,1], npc[,2], col="blue", pch=16, cex=3)
text(npc[,1], npc[,2], c(1,2), col="white")
```



Q18. Which of these new patients should we prioritize for follow up based on your results?

Patient 2 (located in the red/malignant cluster) should be prioritized for follow-up.