Class 7: Machine Learning

Nhi To

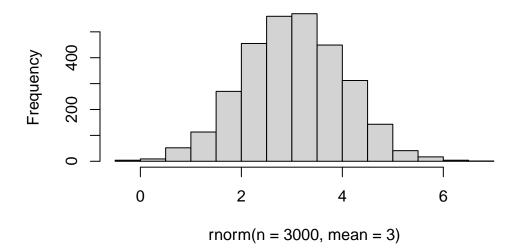
Today we wil explore unsupervised machine learning methods including clustering and dimensionality reduction methods.

Let's start by making up some data (where we know there are clear gorups) that we can use to test out different clustering methods.

We can use the 'rnorm()' function to help us here:

hist(rnorm(n=3000, mean=3))

Histogram of rnorm(n = 3000, mean = 3)



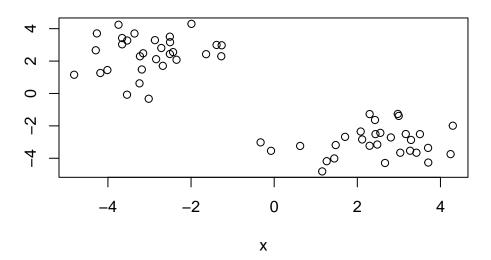
Make data with two "clusters'

```
x <- c( rnorm(30, mean=-3),
    rnorm(30, mean=+3) )

z <- cbind(x=x, rev(x))
head(z)</pre>
```

x [1,] -2.869380 3.291990 [2,] -2.510397 3.506521 [3,] -2.432219 2.553661 [4,] -1.262768 2.973889 [5,] -3.535593 3.270077 [6,] -2.839780 2.116586

plot(z)



How big is 'z'

nrow(z)

[1] 60

```
ncol(z)
```

[1] 2

K-means clustering

The main function in "base" R for K-means clustering is called 'kmeans()'

```
k <- kmeans(z, centers= 2)
k</pre>
```

K-means clustering with 2 clusters of sizes 30, 30

Cluster means:

```
x
1 2.434715 -2.994850
2 -2.994850 2.434715
```

Clustering vector:

Within cluster sum of squares by cluster:

```
[1] 62.58684 62.58684 (between_SS / total_SS = 87.6 %)
```

Available components:

```
[1] "cluster" "centers" "totss" "withinss" "tot.withinss" [6] "betweenss" "size" "iter" "ifault"
```

attributes(k)

```
$names
```

```
[1] "cluster" "centers" "totss" "withinss" "tot.withinss" [6] "betweenss" "size" "iter" "ifault"
```

\$class

[1] "kmeans"

Q. How many points lie in each cluster?

k\$size

[1] 30 30

Q. What component of our results tells us about the cluster membership (i.e. which pointlies in which cluster)?

k\$cluster

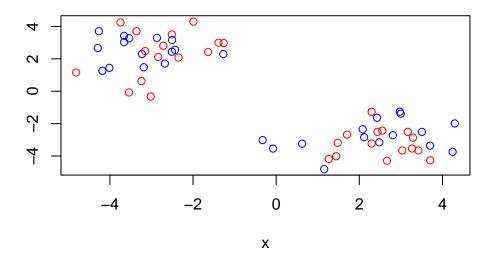
- - Q, Center of each cluster?

k\$centers

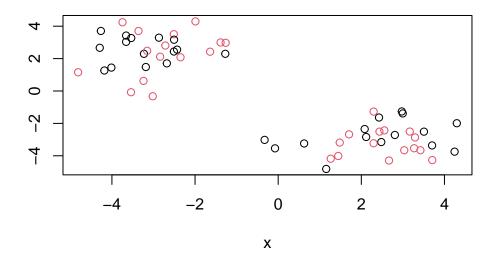
x 1 2.434715 -2.994850 2 -2.994850 2.434715

Q. Put this result info together and make a little "base R" plot of our clustering result. Also add the cluster center points to this plot.

```
plot (z, col= c("blue", "red"))
```

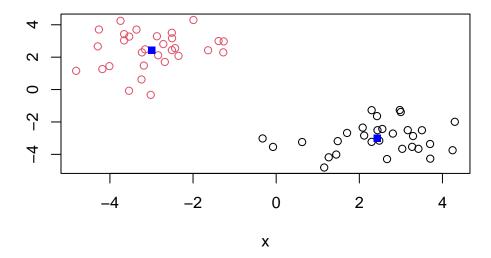


You can color by number



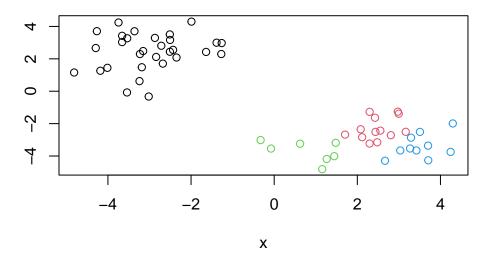
Plot colored by cluster membership:

```
plot(z, col= k$cluster)
points(k$centers, col="blue", pch=15)
```



Q. Run kmeans on our input 'z' and define 4 clusters making the same result visualiation plot as plot as one (plot of z colored by cluster membership)

```
k4 <- kmeans(z, centers=4)
plot(z, col= k4$cluster)</pre>
```



Hiearchial CLustering

The main function in base R for this called 'hclus()' it will take as input a distance matrix(key point is that you can't just give your "raw" data as input- you have to first calculate a distance martrix from your data).

```
d<- dist(z)
hc <- hclust(d)
hc</pre>
```

Call:

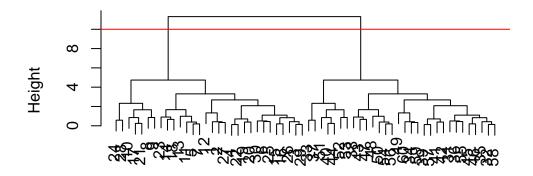
hclust(d = d)

Cluster method : complete
Distance : euclidean

Number of objects: 60

```
plot(hc)
abline(h=10, col="red")
```

Cluster Dendrogram

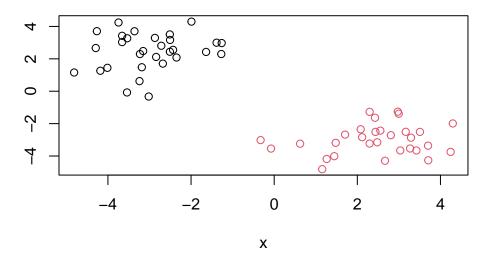


d hclust (*, "complete")

Once I inspect the "tree" I can "cut" the tree to yield my groupings or clusters. The function to do this is called 'cutree()'

```
grps<- cutree(hc, h=10)</pre>
```

plot(z, col=grps)



Hands on with Principal Component Analysis (PCA)

Let's examine some silly 17-dimensional data detailing food consumption in the UK (England, Scotland, Wales, and N. Ireland). Are these countries eating habits different or similr and if so how?

Data import

```
url <- "https://tinyurl.com/UK-foods"
x <- read.csv(url, row.names=1)
x</pre>
```

	England	Wales	${\tt Scotland}$	N.Ireland
Cheese	105	103	103	66
Carcass_meat	245	227	242	267
Other_meat	685	803	750	586
Fish	147	160	122	93
Fats_and_oils	193	235	184	209
Sugars	156	175	147	139
Fresh_potatoes	720	874	566	1033

Fresh_Veg	253	265	171	143
Other_Veg	488	570	418	355
Processed_potatoes	198	203	220	187
Processed_Veg	360	365	337	334
Fresh_fruit	1102	1137	957	674
Cereals	1472	1582	1462	1494
Beverages	57	73	53	47
Soft_drinks	1374	1256	1572	1506
Alcoholic_drinks	375	475	458	135
Confectionery	54	64	62	41

Q1. How many rows and columns are in your new data frame named x? What R functions could you use to answer this questions?

Answer: There are 17 rows and 4 columns

Q2. Which approach to solving the 'row-names problem' mentioned above do you prefer and why? Is one approach more robust than another under certain circumstances?

Answer:I approach "nrow()" and "ncol()" because it allows me to choose with one I would like to look at. If i use "dim()" I would be generating both.

nrow(x)

[1] 17

ncol(x)

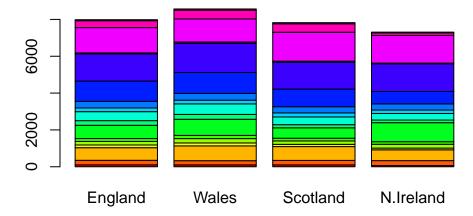
[1] 4

dim(x)

[1] 17 4

Q3. Changing what optional argument in the above barplot() function results in the following plot?

Answer:Changing the argument from barplot(as.matrix(x), beside=T, col=rainbow(nrow(x))) to barplot(as.matrix(x), beside=F, col=rainbow(nrow(x))) would result in a stacked barplot instead of a grouped barplot. The argument being changed was beside=T to beside=F.



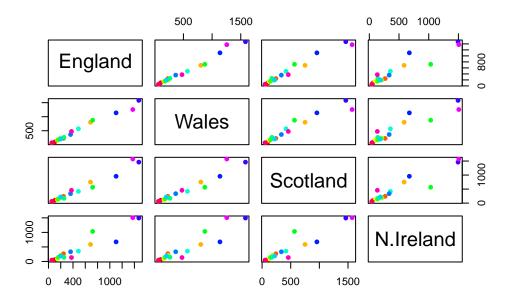
Q5: Generating all pairwise plots may help somewhat. Can you make sense of the following code and resulting figure? What does it mean if a given point lies on the diagonal for a given plot?

Ans: if the points are diagonal on a given plot, it means that the values of the two compared countries are similar. If the point does not align diagonally, it means that they are not similar.

Q6. What is the main differences between N. Ireland and the other countries of the UK in terms of this data-set?

Ans: The main differences between N. Ireland the other countries of the UK are the consumption of potatoes, fresh fruit, soft drinks, and alcoholic drinks. Ireland has consumed significantly more fresh potatoes than other countries of the UK. Ireland has consumed significantly less fresh fruit than the other countries of the UK. Ireland has consumed much more soft drinks than Wales. Ireland has consumed significantly less alcoholic dirnks than the other countries of the UK.

```
pairs(x, col=rainbow(nrow(x)), pch=16)
```



Looking at these types of "pairwise plots" can be helpful but it does not scale well and kind of sucks! There must be a better way...

PCA to the rescue!

The main function for PCA is base R is called 'prcomp()'. This function wants the transpose of our input data- i.e. the important foods in as columns and the countries as rows.

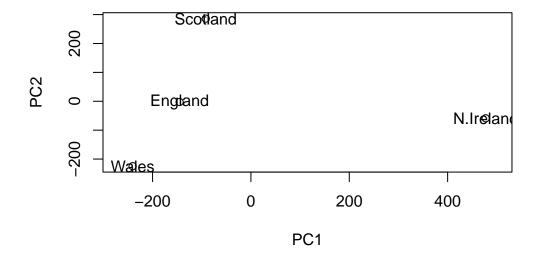
```
pca <- prcomp(t(x))
summary(pca)</pre>
```

Importance of components:

```
PC1 PC2 PC3 PC4
Standard deviation 324.1502 212.7478 73.87622 2.921e-14
Proportion of Variance 0.6744 0.2905 0.03503 0.000e+00
Cumulative Proportion 0.6744 0.9650 1.00000 1.000e+00
```

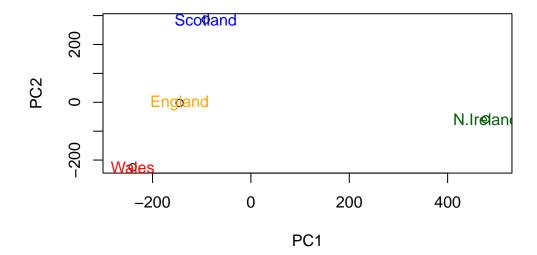
Q7. Complete the code below to generate a plot of PC1 vs PC2. The second line adds text labels over the data points.

```
# Plot PC1 vs PC2
plot(pca$x[,1], pca$x[,2], xlab="PC1", ylab="PC2", xlim=c(-270,500))
text(pca$x[,1], pca$x[,2], colnames(x))
```



Q8. Customize your plot so that the colors of the country names match the colors in our UK and Ireland map and table at start of this document.

```
plot(pca$x[,1], pca$x[,2], xlab="PC1", ylab="PC2", xlim=c(-270,500))
text(pca$x[,1], pca$x[,2], colnames(x), col=c("orange", "red", "blue", "darkgreen"))
```



Looking at these types of "pairwise plots" can be helpful but it does not scale well and kind of sucks! There must be a better way...

Let's see what is in our PCA result object 'pca'

attributes(pca)

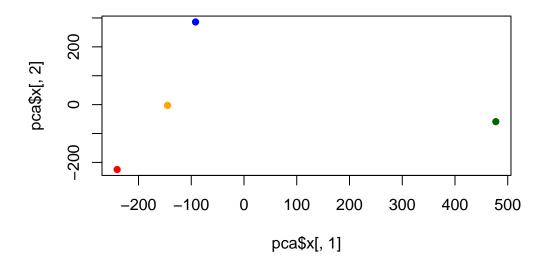
\$names [1] "sdev" "rotation" "center" "scale" "x" \$class [1] "prcomp"

The 'pca\$x' result object is where we will focus first as this details how the countries are related to each other in terms of our new "axis" (a.k.a "PCs", "eigenvectors", etc.)

head(pca\$x)

```
PC1
                             PC2
                                        PC3
                                                       PC4
England
          -144.99315
                       -2.532999 105.768945 -9.152022e-15
Wales
          -240.52915 -224.646925 -56.475555
                                             5.560040e-13
Scotland
           -91.86934
                      286.081786 -44.415495 -6.638419e-13
N.Ireland 477.39164
                      -58.901862
                                  -4.877895
                                             1.329771e-13
```

plot(pca\$x[,1], pca\$x[,2], pch=16, col=c("orange", "red", "blue", "darkgreen"))



We can look at the so-called PC "loadings" result object to see how the original foods contribute to our new PCs (i.e. how the original variables contribute to our new better PC variables)

pca\$rotation[,1]

Cheese	Carcass_meat	Other_meat	Fish
-0.056955380	0.047927628	-0.258916658	-0.084414983
Fats_and_oils	Sugars	Fresh_potatoes	Fresh_Veg
-0.005193623	-0.037620983	0.401402060	-0.151849942
Other_Veg	Processed_potatoes	Processed_Veg	Fresh_fruit
-0.243593729	-0.026886233	-0.036488269	-0.632640898
Cereals	Beverages	${\tt Soft_drinks}$	Alcoholic_drinks
-0.047702858	-0.026187756	0.232244140	-0.463968168
Confectionery			
-0.029650201			

Calculating how much variation in the original data each PC accounts for

```
v <- round( pca$sdev^2/sum(pca$sdev^2) * 100 )
v</pre>
```

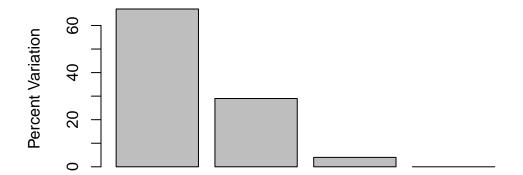
[1] 67 29 4 0

```
z <- summary(pca)
z$importance</pre>
```

```
PC1 PC2 PC3 PC4
Standard deviation 324.15019 212.74780 73.87622 2.921348e-14
Proportion of Variance 0.67444 0.29052 0.03503 0.000000e+00
Cumulative Proportion 0.67444 0.96497 1.00000 1.000000e+00
```

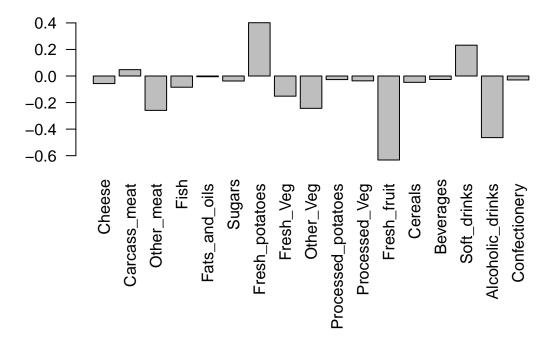
Plot v

```
barplot(v, xlab="Principal Component", ylab="Percent Variation")
```



Principal Component

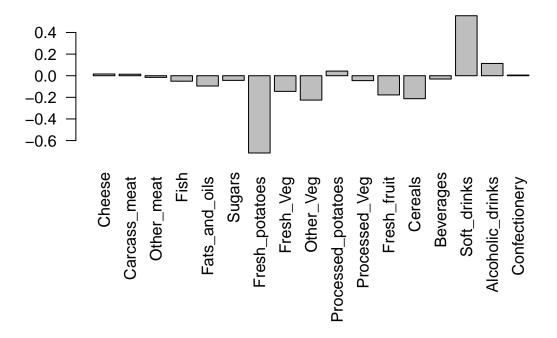
```
## Lets focus on PC1 as it accounts for > 90% of variance
par(mar=c(10, 3, 0.35, 0))
barplot( pca$rotation[,1], las=2 )
```



Q9: Generate a similar 'loadings plot' for PC2. What two food groups feature prominantely and what does PC2 maniply tell us about?

Answer: Fresh potatoes and soft drinks are the two food groups that are featured prominately in the 'loadings plot' for PC2. PC2 details the second-largest variance in the dataset, which means it captures variations not included in PC1.

```
## Lets focus on PC2
par(mar=c(10, 3, 0.35, 0))
barplot( pca$rotation[,2], las=2 )
```



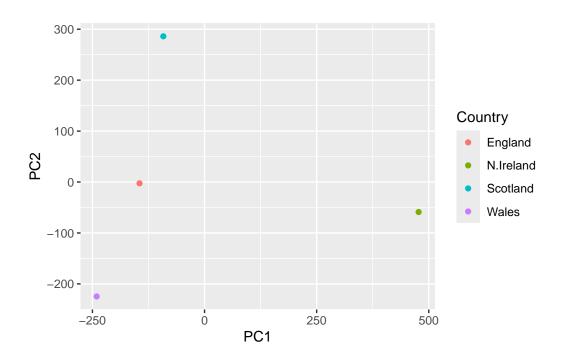
Using ggplot for the figures

```
library(ggplot2)

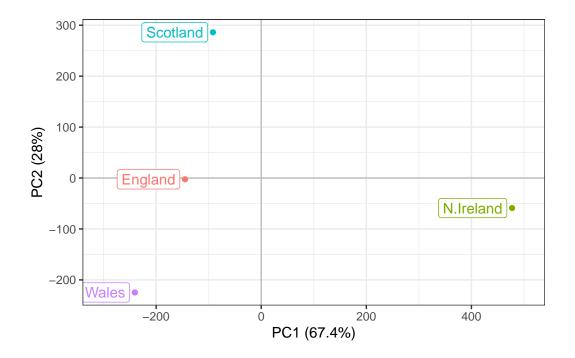
df <- as.data.frame(pca$x)

df_lab <- tibble::rownames_to_column(df, "Country")

# Our first basic plot
ggplot(df_lab) +
   aes(PC1, PC2, col=Country) +
   geom_point()</pre>
```

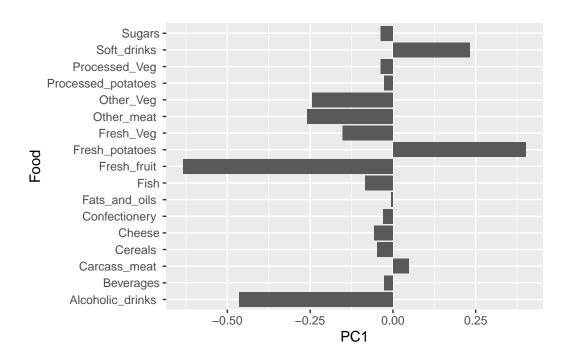


```
# a much nicer plot
ggplot(df_lab) +
  aes(PC1, PC2, col=Country, label=Country) +
  geom_hline(yintercept = 0, col="gray") +
  geom_vline(xintercept = 0, col="gray") +
  geom_point(show.legend = FALSE) +
  geom_label(hjust=1, nudge_x = -10, show.legend = FALSE) +
  expand_limits(x = c(-300,500)) +
  xlab("PC1 (67.4%)") +
  ylab("PC2 (28%)") +
  theme_bw()
```



```
ld <- as.data.frame(pca$rotation)
ld_lab <- tibble::rownames_to_column(ld, "Food")

ggplot(ld_lab) +
  aes(PC1, Food) +
  geom_col()</pre>
```



```
## PCA of RNA-seq data
url2 <- "https://tinyurl.com/expression-CSV"
rna.data <- read.csv(url2, row.names=1)
head(rna.data)</pre>
```

```
wt3
       wt1 wt2
                     wt4 wt5 ko1 ko2 ko3 ko4 ko5
       439 458
                408
                     429 420
                              90
                                  88
                                       86
                                           90
gene1
gene2
       219 200
                204
                     210 187 427 423 434 433 426
gene3 1006 989 1030 1017 973 252 237 238 226 210
gene4
       783 792
                829
                     856 760 849 856 835 885 894
gene5
       181 249
                204
                     244 225 277 305 272 270 279
gene6
       460 502
                491
                     491 493 612 594 577 618 638
```

Q10: How many genes and samples are in this data set?

Answer: There are 100 rows, which means there is 100 genes. There are 10 columns, which means there are 10 samples.

nrow(rna.data)

[1] 100

ncol(rna.data)

[1] 10

dim(rna.data)

[1] 100 10