

Machine Learning pt. 1

Kelly_F

10/22/2021

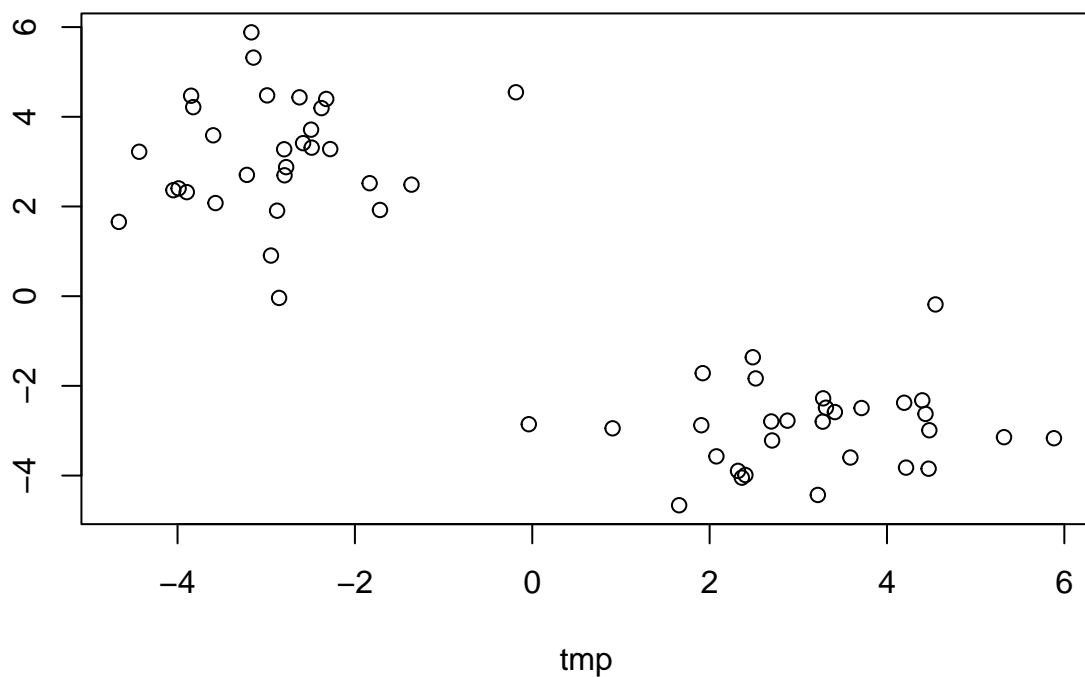
Clustering Methods

Kmeans clustering

k-means clustering: R function is “kmeans()”.

Test data used below to learn use of kmeans() function.

```
# Create Test Data  
tmp <- c(rnorm(30, 3), rnorm(30, -3))  
data <- cbind(tmp, rev(tmp)) #two columns, with second column the reverse of tmp vector  
  
# Plot data, which was constructed to have two clear clusters of data.  
plot(data)
```



Note: “clustering vector”(output) Tells you which cluster each element of the data belongs to.

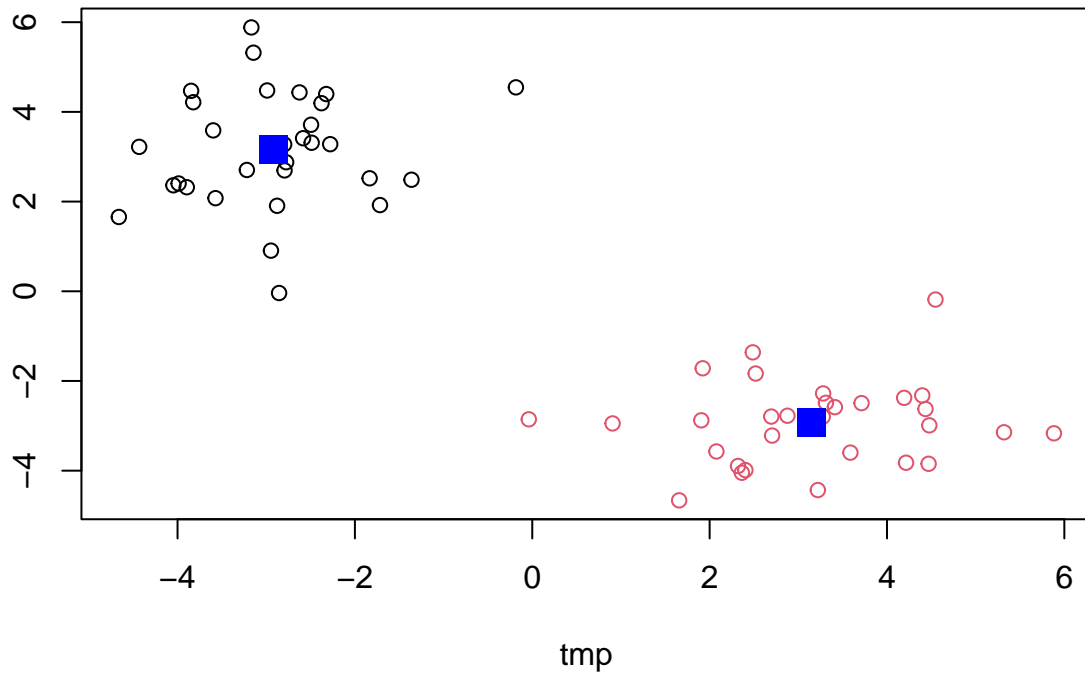
30 points in each cluster.

Q. What “compoennet” of your results object details cluster center?

2

Q. Plot x colored by the kmeans cluster assignemnt and add cluster centers as blue points.

```
plot(data, col=km_data$cluster)
points(km_data$centers, col="blue", pch=15, cex=2)
```



Hierarchical Clustering

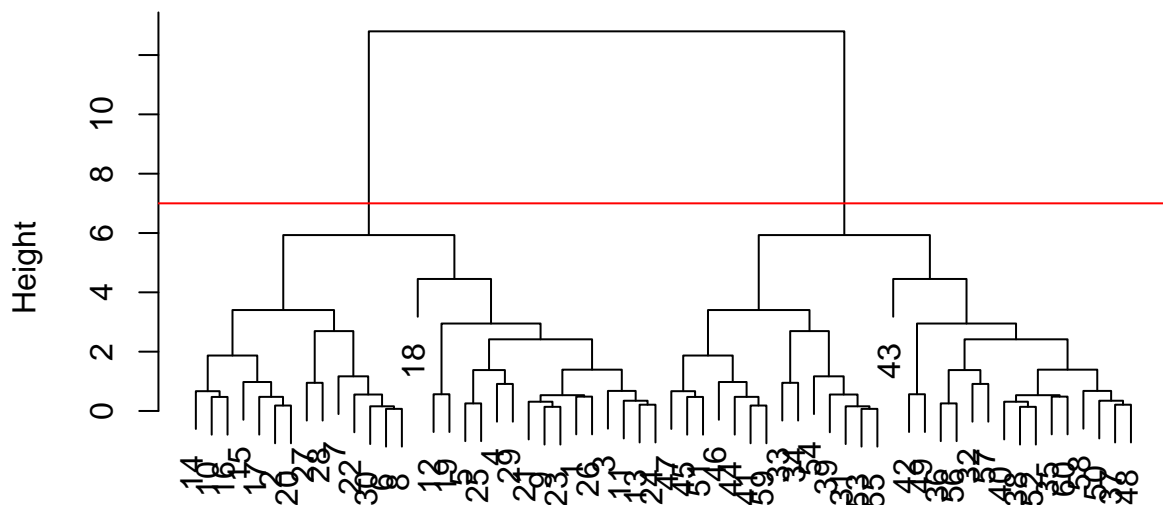
We will use the 'hclust()' function on the same data as kmeans example to see how this method works.

```
# hclust needs a distance matrix as an input
hc <- hclust(dist(data))
hc
```

```
##
## Call:
## hclust(d = dist(data))
##
## Cluster method   : complete
## Distance        : euclidean
## Number of objects: 60
```

```
# Plot Dendrogram to investigate between-cluster differences.
# Further distance on the dendrogram = more dissimilar
plot(hc)
abline(h=7, col="red")
```

Cluster Dendrogram



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

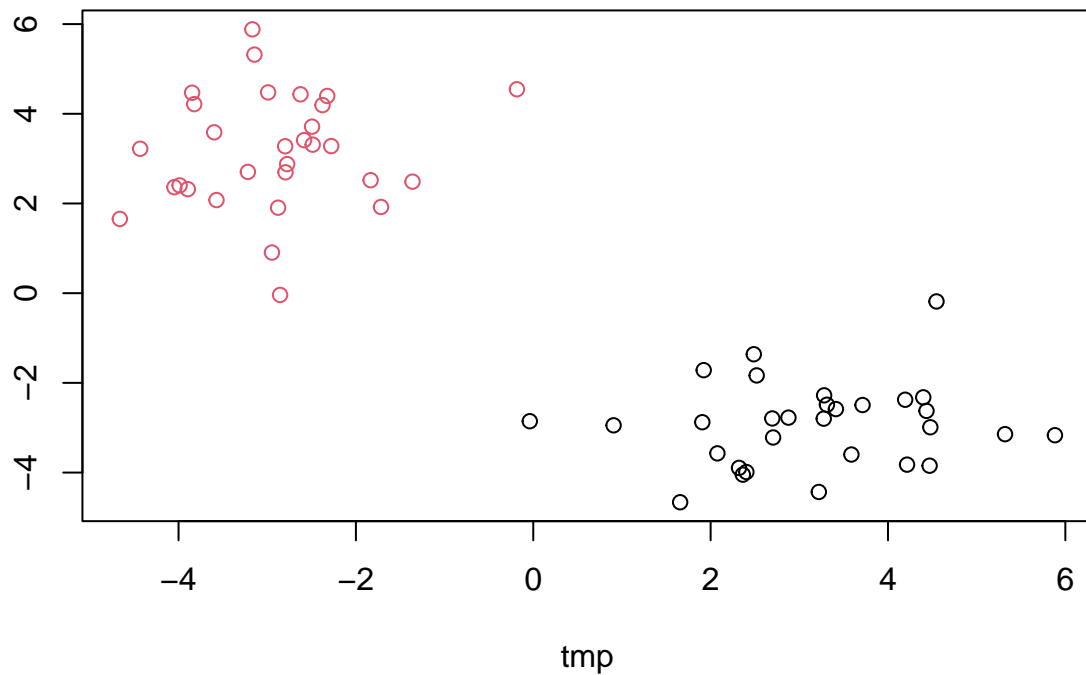
To find our membership vector we need to “cut” the tree into its respective branches (clusters). For this we will use the ‘cutree()’ function and tell it the height to cut at.

```
# Cut at height=7
cutree(hc, h=7)
```

[illegible]

```
# Alternatively, we can instead tell 'cutree()' how many clusters we want.
grps <- cutree(hc, k=2)
```

```
# Plot data and color by hclust grouping
plot(data, col=grps)
```



Kmeans recap

- clusters data, but must tell it how many centers you want.
- functions needs euclidean distances.

Hclust recap

- doesn't take raw data. Must give it a distance matrix.
- doesn't require euclidean distances as input.

Principal Component Analysis w/ UK Food Data

Analysis goal: is the diet composition across the 4 countries of interest different?

```
# Import data
url <- "https://tinyurl.com/UK-foods"
uk_food <- read.csv(url, row.names = 1)

# Determine the number of rows and columns in the dataframe
dim(uk_food)
```

```
## [1] 17  4
```

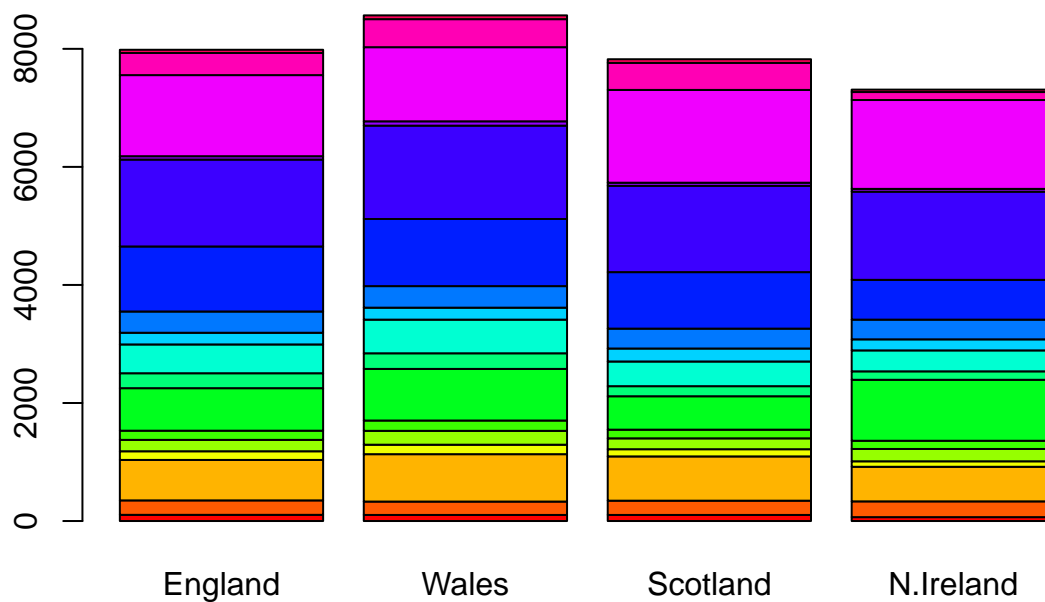
Q1. How many rows and columns are in your new data frame named x? What R functions could you use to answer this questions?

Initially, 17 rows, 5 columns. You can use 'dim()' function to determine this. Note, there are only 4 countries, so when .csv is read in, you must tell it that rownames are in the first column, row.names=1.

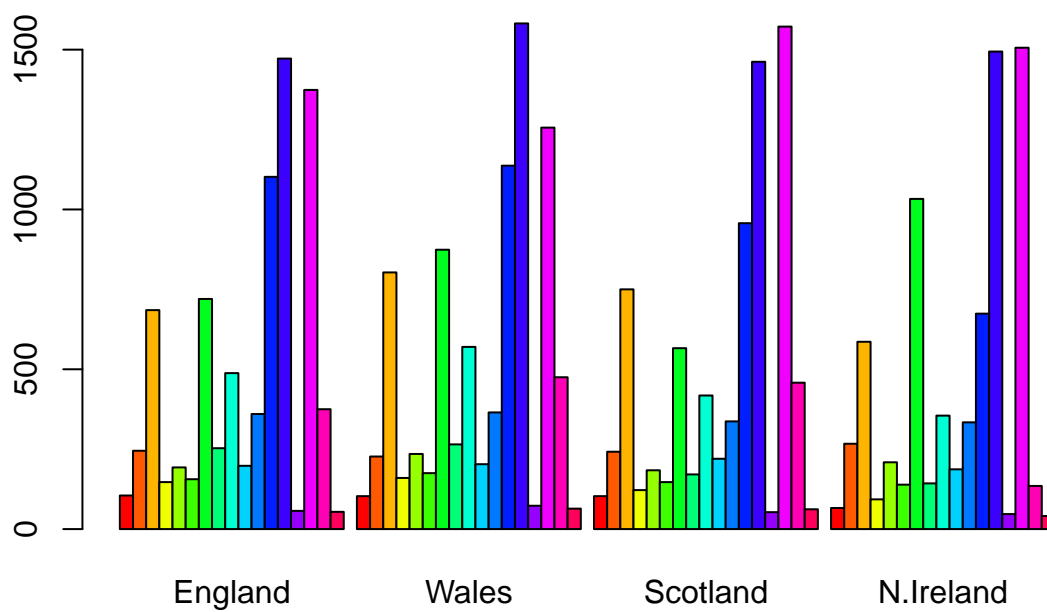
Q2. Which approach to solving the 'row-names problem' mentioned above do you prefer and why? Is one approach more robust than another under certain circumstances?

The row.names option when reading in the .csv, because it doesnt overwrite or obstruct the file you are importing.

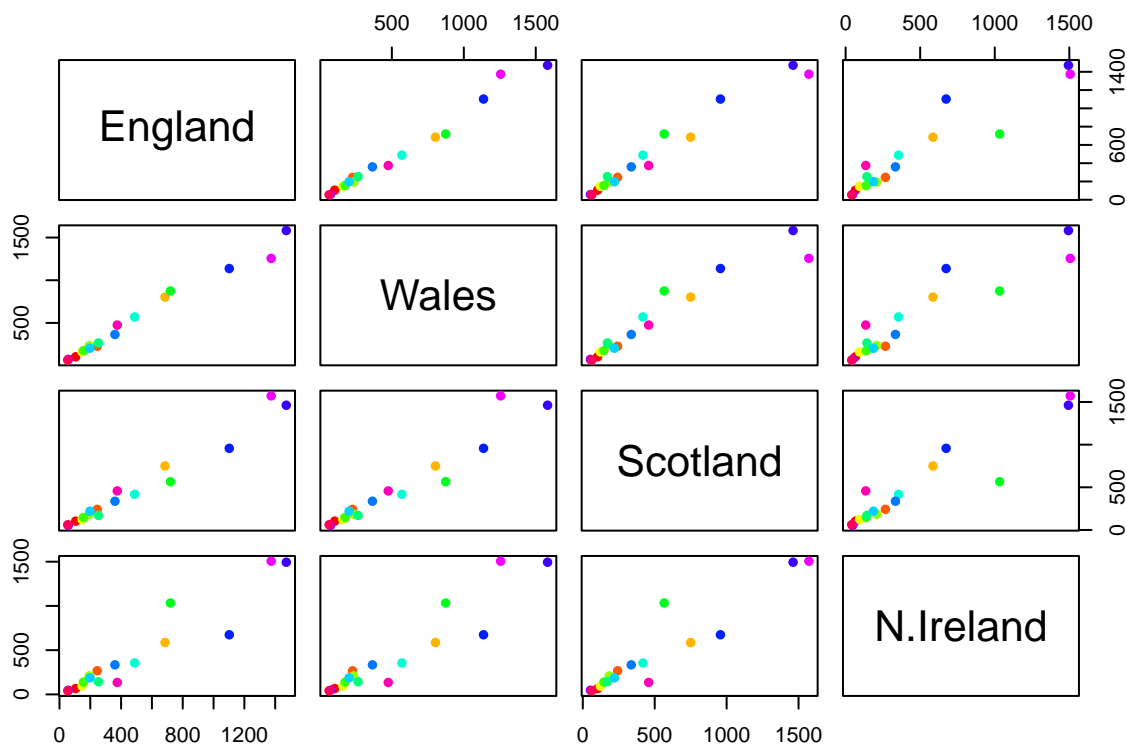
```
# Start to visualize data  
barplot(as.matrix(uk_food), col=rainbow(17))
```



```
barplot(as.matrix(uk_food), beside=T, col=rainbow(nrow(uk_food))) # plot each food category separately
```



```
mycols <- rainbow(nrow(uk_food)) #generate colors # of food categories
# Plot all possible pairwise correlation plots
pairs(uk_food, col=mycols, pch=16) #alternatively can set col=rainbow(17). Accomplishes the same thing
```



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

From the limited analysis we've completed, it seems that N. Ireland has lower diversity in what they eat compared to other countries. I.e., there are a few predominant food groups w/ high frequency, and the remaining food groups have lower frequency.

PCA to the rescue!

Here we will use the base R function for PCA, which is called 'prcomp()'.

```
# Run PCA
```

```
#As we noted in the lecture portion of class, prcomp() expects the observations to be rows and the vari
```

```
pca <- prcomp(t(uk_food)) # transpose dataframe for PCA analysis
summary(pca)
```

```
## Importance of components:
```

```
##
## Standard deviation      324.1502 212.7478 73.87622 4.189e-14
## Proportion of Variance  0.6744  0.2905  0.03503 0.000e+00
## Cumulative Proportion   0.6744  0.9650  1.00000 1.000e+00
```



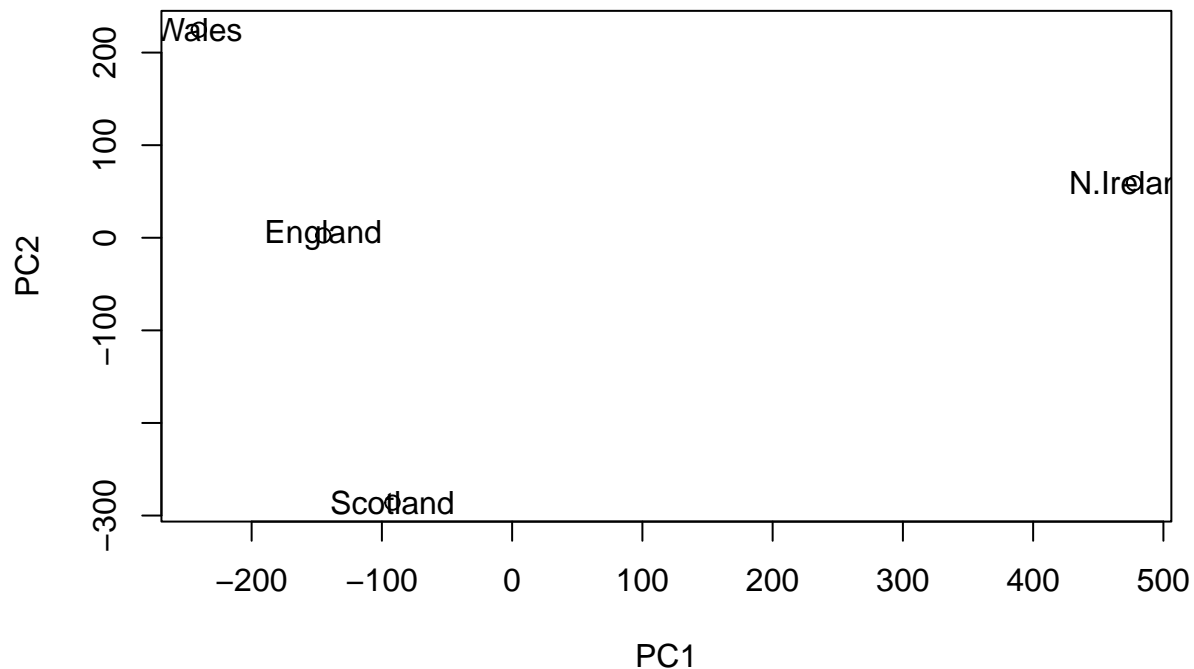
```
attributes(pca)
```

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

```
pca$x
```

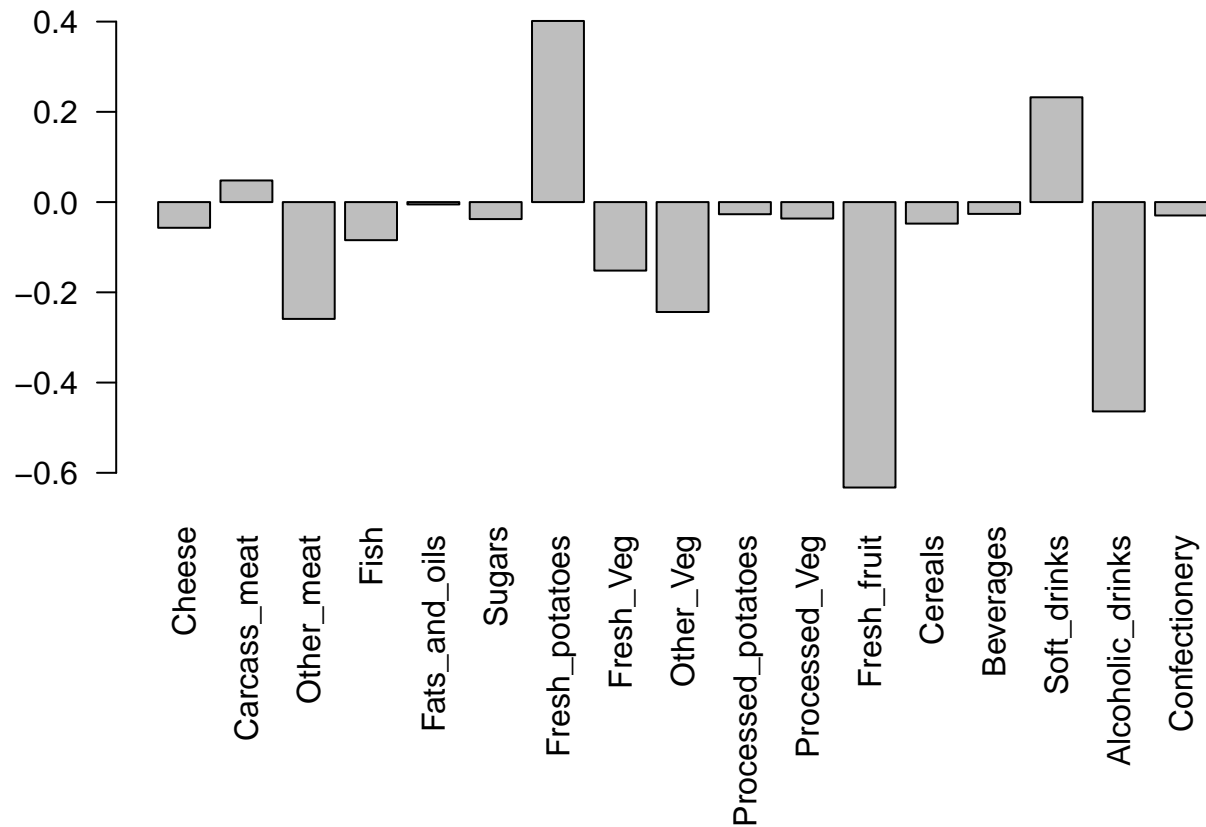
```
##           PC1           PC2           PC3           PC4
## England  -144.99315    2.532999 -105.768945  2.842865e-14
## Wales    -240.52915   224.646925   56.475555  7.804382e-13
## Scotland  -91.86934  -286.081786   44.415495 -9.614462e-13
## N.Ireland  477.39164    58.901862    4.877895  1.448078e-13
```

```
# Plot PCA
plot(pca$x[, 1:2])#plot PCA1 & 2 for each country
text(pca$x[,1], pca$x[,2], colnames(uk_food))
```



We can also examine the PCA “loadings” and investigate how much each individual food group contributes to each PC

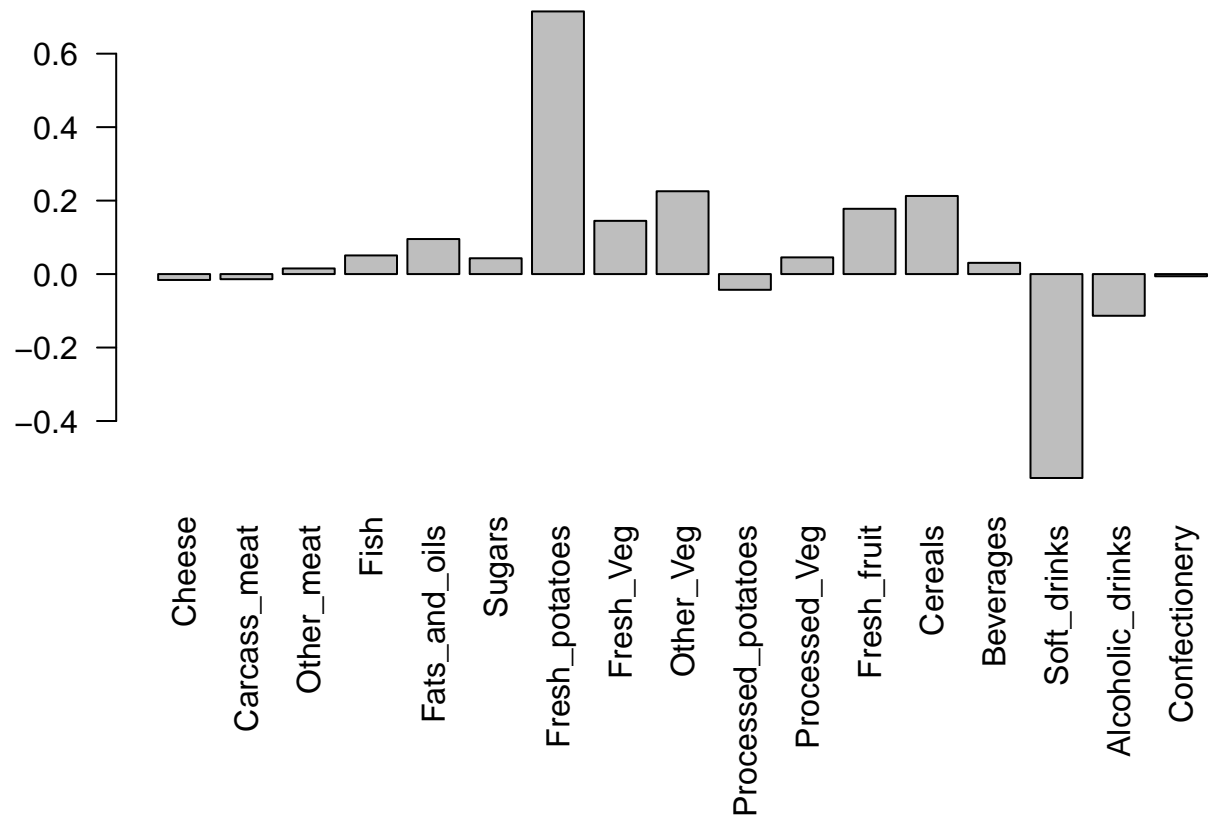
```
#Lets focus on PC1 as it accounts for > 90% of variance
par(mar=c(10, 3, 0.35, 0))
barplot(pca$rotation[, 1], las=2)
```



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

PC2 mainly tells us about differences between Wales, England, and Scotland. Wales has a high feature count for processed potatoes VS Scotland which has a high feature count for soft drinks.

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



PCA of RNA-seq data

```
url2 <- "https://tinyurl.com/expression-CSV"
rna.data <- read.csv(url2, row.names=1)
head(rna.data)
```

```
##      wt1 wt2 wt3 wt4 wt5 ko1 ko2 ko3 ko4 ko5
## gene1 439 458 408 429 420 90 88 86 90 93
## 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?

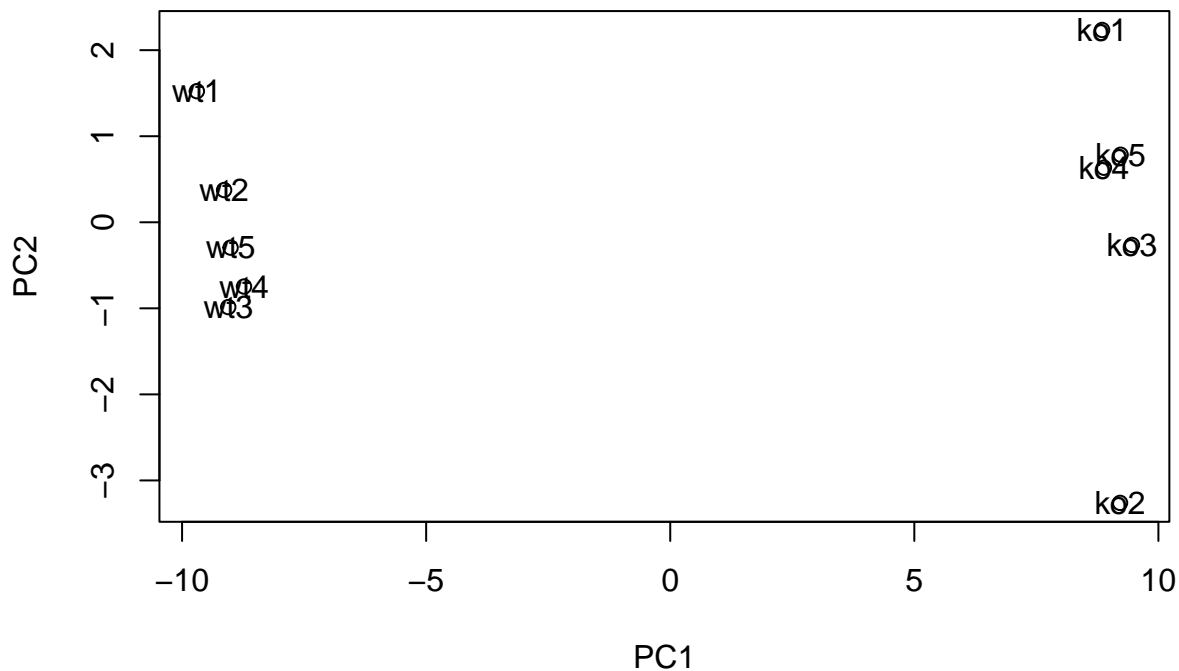
100 genes and 10 samples

```
dim(rna.data)
```

```
## [1] 100 10
```

```
# Generate PCA
pca_rna <- prcomp(t(rna.data), scale=TRUE) #transpose data before running PCA analysis

# Simple un polished plot of pc1 and pc2
plot(pca_rna$x[,1], pca_rna$x[,2], xlab="PC1", ylab="PC2")
text(pca_rna$x[, 1:2], labels= colnames(rna.data))
```



```
#Investigate what proportion of variance PC1 explains
summary(pca_rna)
```

```
## Importance of components:
##              PC1      PC2      PC3      PC4      PC5      PC6      PC7
## Standard deviation  9.6237  1.5198  1.05787  1.05203  0.88062  0.82545  0.80111
## Proportion of Variance 0.9262  0.0231  0.01119  0.01107  0.00775  0.00681  0.00642
## Cumulative Proportion 0.9262  0.9493  0.96045  0.97152  0.97928  0.98609  0.99251
##              PC8      PC9      PC10
## Standard deviation  0.62065  0.60342  3.348e-15
## Proportion of Variance 0.00385  0.00364  0.000e+00
## Cumulative Proportion 0.99636  1.00000  1.000e+00
```