Class 7 Lab

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Hands on with Principal Component Analysis

Examine a 17-dimensional data detailing food consumption in England, Wales, Scotland, and Northern Ireland.

Read the provide input file

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

17 rows, 5 columns, using nrow() and ncol()

```
url<- "https://tinyurl.com/UK-foods"
x <- read.csv(url) ##To remove numbers from row, could add ', row.names=1' after url
nrow(x)</pre>
```

[1] 17

```
ncol(x)
```

[1] 5

head(x)

	Х	England	Wales	Scotland	N.Ireland
1	Cheese	105	103	103	66
2	Carcass_meat	245	227	242	267
3	${\tt Other_meat}$	685	803	750	586
4	Fish	147	160	122	93
5	Fats_and_oils	193	235	184	209
6	Sugars	156	175	147	139

```
#View(x)
```

Remove the first column so it starts with the value of each food and not a number

```
rownames(x) <- x[,1]
x <- x[,-1]
head(x)
```

	England	Wales	Scotland	${\tt N.Ireland}$
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

```
dim(x)
```

[1] 17 4

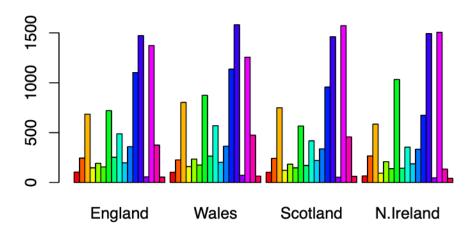
#DO NOT run again, will remove another column, can be fixed by rerunning code above so that

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 minus indexing technique can be tricky because it requires that you only run functional code once or else it will continue to remove values from the dataset. For this reason, I think the technique below might be more functional. $x \leftarrow read.csv(url, row.names=1)$

Make a fun rainbow plot of this data

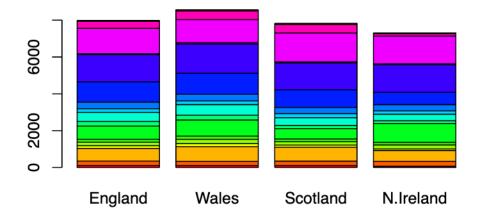
```
barplot(as.matrix(x), beside=T, col=rainbow(nrow(x)))
```



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

Setting 'beside' as equal to "F". The beside line of code changes the arrangement of the bars in the plot.

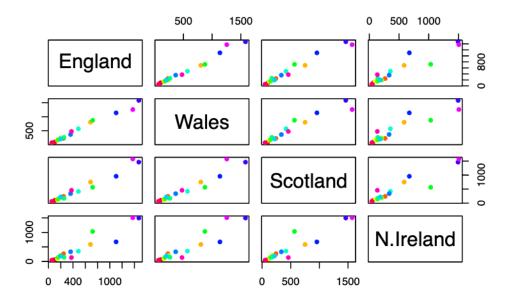
```
barplot(as.matrix(x), beside=F, col=rainbow(nrow(x)))
```



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?

This kind of display is only workable with small datasets and is called a scatterplot matrix. Each variable is listed in a line and plotted against each other. In the first row, England is the y axis. The x axis depends on the other intersecting country. So, row 1 is England as y axis and column 1 is England as x axis. If a point lies on a diagonal, it means that there is a perfect correlation between the two variables, ie they are both the same value.

```
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 (time consuming, laborious, error prone)!

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

N. Ireland has a noticeable difference in the bright green dot towards the center of the dataset, as it lies astray from the diagonal line. However, it is hard to deduce much specifically with so many variables and different plots.

PCA to the rescue

The main function for PCA in base R is called 'prcomp()'. This function wants the transpose of our input data - i.e. the important food categories 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

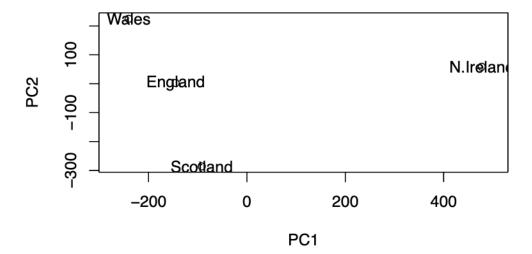
The 'pca\$x' result object is where we will focus first as this details how the countries are related to each other in terms of our new "axis" (ie "PCs", "eigenvectors", etc.).

head(pca\$x)

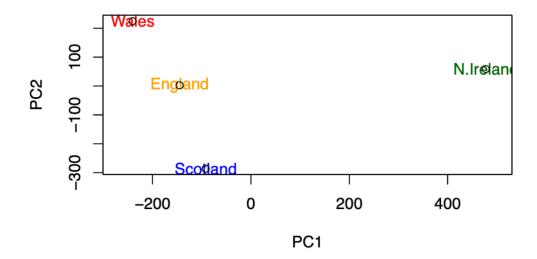
```
PC1
                              PC2
                                         PC3
                                                       PC4
England
          -144.99315
                       -2.532999 105.768945 -9.152022e-15
Wales
          -240.52915 -224.646925 -56.475555
                 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

Q7. Complete the code below to generate a plot of PC1 vs PC2. The second line adds text labels over the data points. plot(pcax[,1],pcax[,2], pch=16, col=c("orange", "red", "blue", "darkgreen"), xlab="PC1", ylab= "PC2", xlim=c(-270,500), text(pcax[,1],pcax[,2], colnames(x)))

```
pca$x [,2] <- -pca$x [,2]
#PCA1 is correct, PCA2 is opposite for some reason? Y axis needs to be mult by -1
plot(pca$x[, 1], pca$x[, 2], xlab="PC1", ylab="PC2", xlim=c(-270,500))
text(pca$x[, 1], pca$x[, 2], colnames(x))</pre>
```



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.



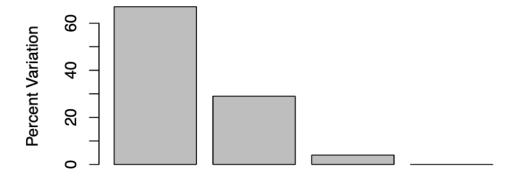
How much variation in the original data set does each PC account for (by proportion to 100)?

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

[1] 67 29 4 0

Summarize as a barplot

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



Principal Component

Variable Loadings

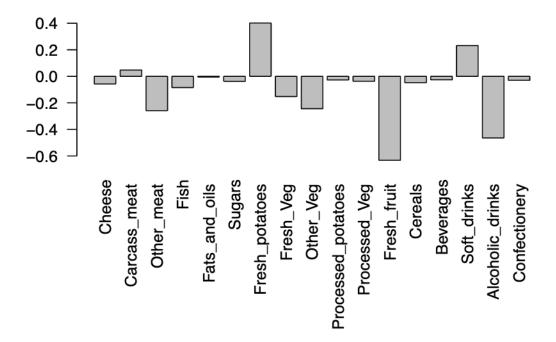
We can look at the so-caled PC loadings result to see how the original foods contribute to our new PCs (ie how the original variables contribute to our new, better variables).

pca\$rotation[,1]

Cheese	Carcass_meat	Other_meat	Fish
-0.056955380	0.047927628	-0.258916658	-0.084414983
${\tt Fats_and_oils}$	Sugars	Fresh_potatoes	${\tt Fresh_Veg}$
-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			

Positive values indicate more impact than others, negative values indicate less impact than others

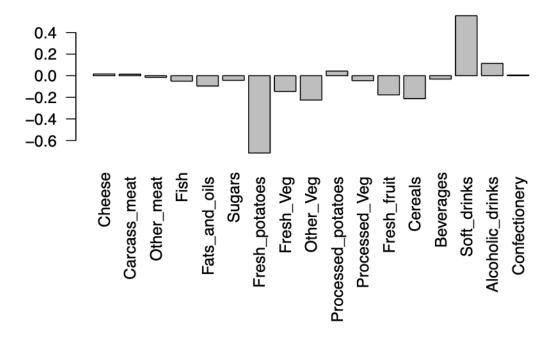
```
#Graphical focus on PC1, largest positive loading scores 'push' N. Ireland to the right posi
par(mar=c(10,3,0.35,0))
barplot(pca$rotation[,1], las=2)
```



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

Fresh potatoes and soft drink are the most dominant bars in this graph. Soft drinks are significantly higher than average and fresh potatoes are significantly lower.

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



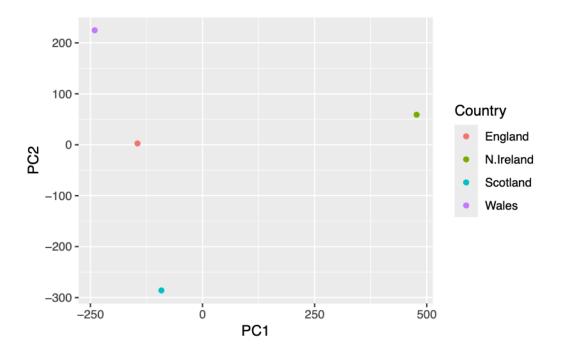
Using ggplot for figures

```
library(ggplot2)

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

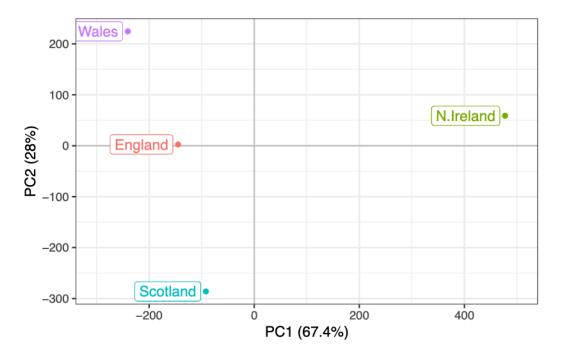
df_lab <- tibble::rownames_to_column(df, "Country")

ggplot(df_lab) +
  aes(PC1, PC2, col=Country) +
  geom_point()</pre>
```



Fancier but a pain to make graph

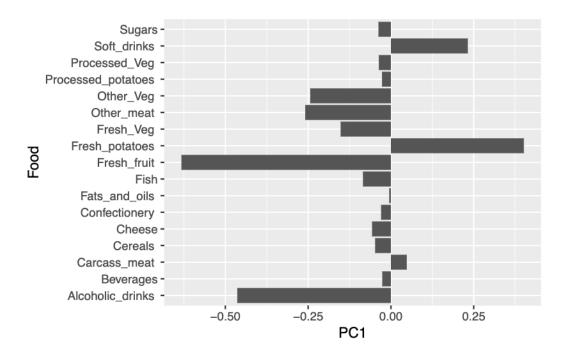
```
ggplot(df_lab) +
  aes(PC1, PC2, col=Country, label=Country) +
  geom_hline(yintercept = 0, col="gray") +
  geom_vline(xintercept = 0, col="gray") +
  geom_point(show.legend = FALSE) +
  geom_label(hjust=1, nudge_x = -10, show.legend = FALSE) +
  expand_limits(x = c(-300,500)) +
  xlab("PC1 (67.4%)")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



Plot of loadings and PC contributions

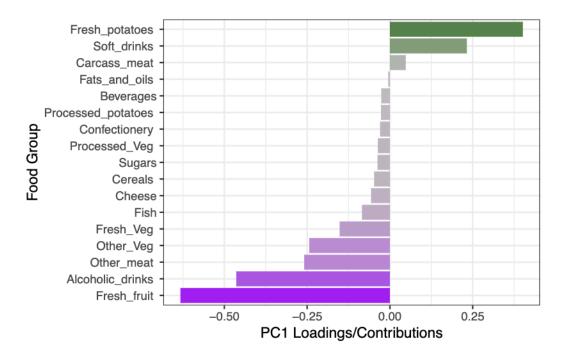
```
ld <- as.data.frame(pca$rotation)
ld_lab <- tibble::rownames_to_column(ld, "Food")

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



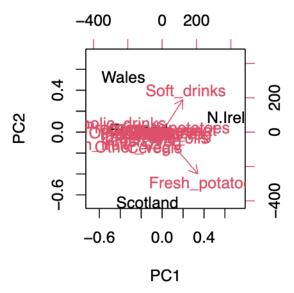
Prettier one

```
ggplot(ld_lab) +
  aes(PC1, reorder(Food, PC1), bg=PC1) +
  geom_col() +
  xlab("PC1 Loadings/Contributions") +
  ylab("Food Group") +
  scale_fill_gradient2(low="purple", mid="gray", high="darkgreen", guide=NULL) +
  theme_bw()
```



Biplots are a good option for small datasets. There is a central group of red aroows pointing to the red word labels for each variable.

biplot(pca)



PCA of RNA-seq data

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

```
    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?

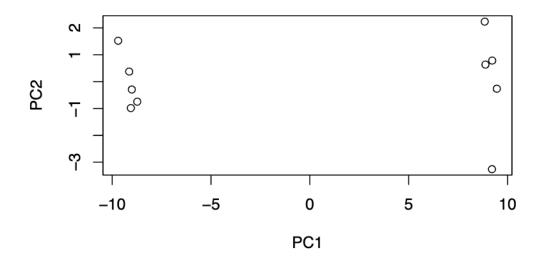
10 genes, 100 samples

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

[1] 100 10

Run a PCA and plot the results

```
pca <- prcomp(t(rna.data), scale=TRUE)
#what does the t indicate here?
plot(pca$x[,1], pca$x[,2], xlab="PC1", ylab="PC2")</pre>
```



summary(pca)

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 3.345e-15 Proportion of Variance 0.00385 0.00364 0.000e+00 Cumulative Proportion 0.99636 1.00000 1.000e+00

Notice 92% of variance is contained in PC1.

plot(pca, main="Quick scree plot")

Quick scree plot



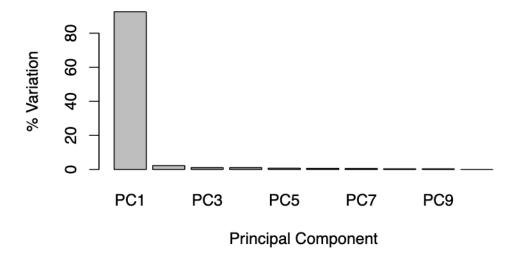
Let's make the scree plot on our own.

```
# Variance captured per PC
pca.var <- pca$sdev^2

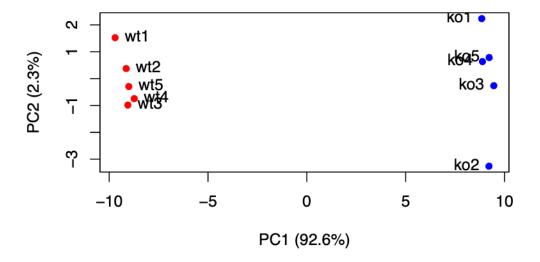
#percent variance
pca.var.per <- round(pca.var/sum(pca.var)*100, 1)
pca.var.per</pre>
```

```
[1] 92.6 2.3 1.1 1.1 0.8 0.7 0.6 0.4 0.4 0.0
```

My Scree Plot



More attractive and more useful, labeling specific points by name and with colors

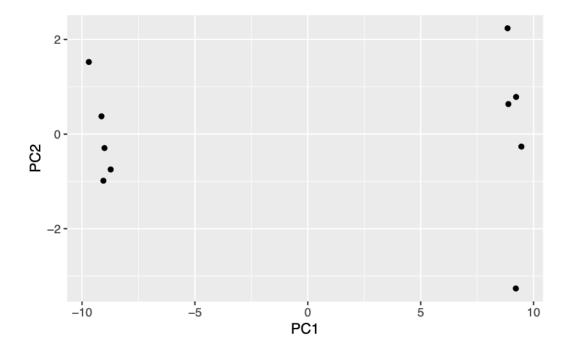


Let's graph it by ggplot

```
library(ggplot2)

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

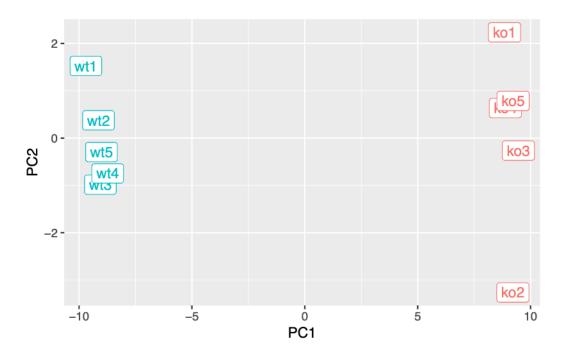
ggplot(df) +
  aes(PC1, PC2) +
  geom_point()</pre>
```



Add a condition specific color, sample label aesthetic for WT vs knockout

```
df$samples <- colnames(rna.data)
df$condition <- substr(colnames(rna.data),1,2)

p <- ggplot(df) +
          aes(PC1, PC2, label=samples, col=condition) +
          geom_label(show.legend = FALSE)
p</pre>
```



Add title, caption, x and y specificity, and a theme

PCA of RNASeq Data

PC1 clealy seperates wild-type from knock-out samples

