# **Assignment 15**

#### **Anthony Cunningham**

```
knitr::opts_chunk$set(echo = TRUE, warning = FALSE, message = FALSE)
# Change working dir in RMarkdown cell
knitr::opts knit$set(root.dir =
'C:/Users/AC069015/kumc_applied_stats/data_824_data_viz_and_acquisition'
library(dplyr)
## Attaching package: 'dplyr'
  The following objects are masked from 'package:stats':
##
##
      filter, lag
  The following objects are masked from 'package:base':
##
##
##
      intersect, setdiff, setequal, union
library(tidyr)
library(ggplot2)
library(readr)
library(datasets)
library(readx1)
library(circlize)
## Warning: package 'circlize' was built under R version 4.2.1
## -----
```

```
library(FactoMineR)

## Warning: package 'FactoMineR' was built under R version 4.2.1

library(factoextra)

## Warning: package 'factoextra' was built under R version 4.2.1

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

library(pheatmap)

## Warning: package 'pheatmap' was built under R version 4.2.1

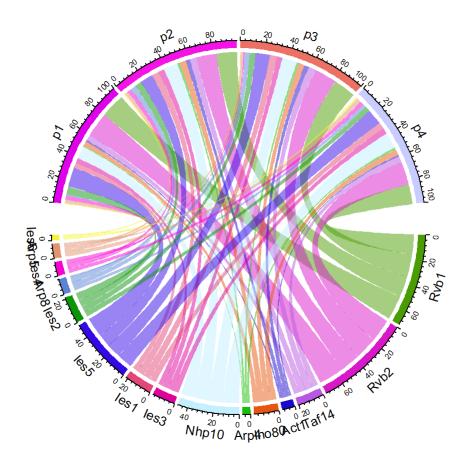
library(cluster)
library(factoextra)
```

#### Exercise 1

```
dta <- read_excel("datasets/Interaction_proteins.xls", skip=1)

dta_new <- dta %>% mutate(
   p1 = rowMeans(select(., contains("_P1")))*100,
   p2 = rowMeans(select(., contains("_P2")))*100,
   p3 = rowMeans(select(., contains("_P3")))*100,
   p4 = rowMeans(select(., contains("_P4")))*100
) %>% select(p1, p2, p3, p4)

rownames(dta_new) <-dta$Proteins
chordDiagram(as.matrix(dta_new))</pre>
```



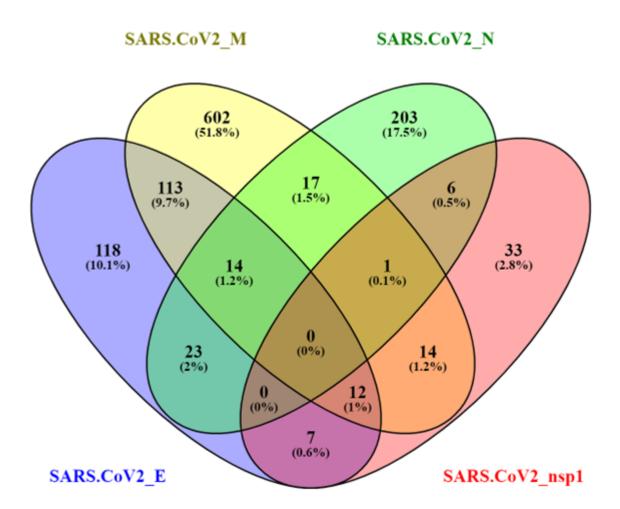
This circle diagram visually depicts the average value for 16 proteins (labeled at the bottom-half) across 4 separate replicates. Larger widths indicate higher averages for a particular protein. From the diagram, we notice that the proteins with the highest average values are Rvb1 and Rvb2, followed by les5 and Nhp10. On the other hand, proteins les3 and les6 consistently displayed the smalled average values.

## Exercise 2

**Examples of Missing Data Structures:** 

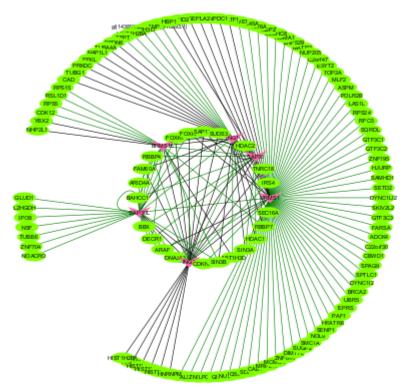
- MCAR: Blood pressure monitor malfunctions at random times, resulting in blood pressure measures not being recorded for some routine patient visits.
- MAR: An overworked nurse in the emergency department doesn't capture patient weight or blood pressure
  measurements during very busy periods in order to maximize patient throughput. While missingness of
  these measures aren't related to the values themselves, they are missing for non-random patients
  (associated with time and missingness of the other measures that are also not captured).
- MNAR: An outpatient clinic asks all patients about whether (and to what extent) they use tobacco,
  marijuana, alcohol and hard drugs. It's plausible that users of substances that are banned or illegal in the
  clinic's area are more likely to not answer this question than non-users (i.e. response is missing), for fear of
  disclosure to law enforcement agencies.

### Exercise 3



Venn Diagram

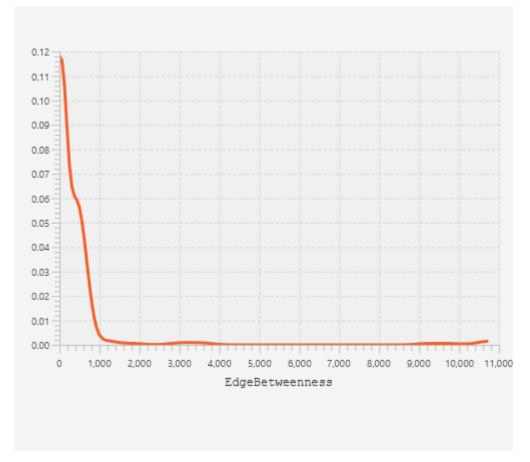
## Exercise 4



Network Graph - Circular Layout

#### **Network Properties:**

- Clustering Coefficient = 0.111 this indicates a large degree of separation between nodes, on average (i.e. a randomly-selected node is unlikely to have triangular connections)
- Degree Distribution: This network appears to be a scale-free network, with only a few nodes with many edges; most nodes only have a few edges:

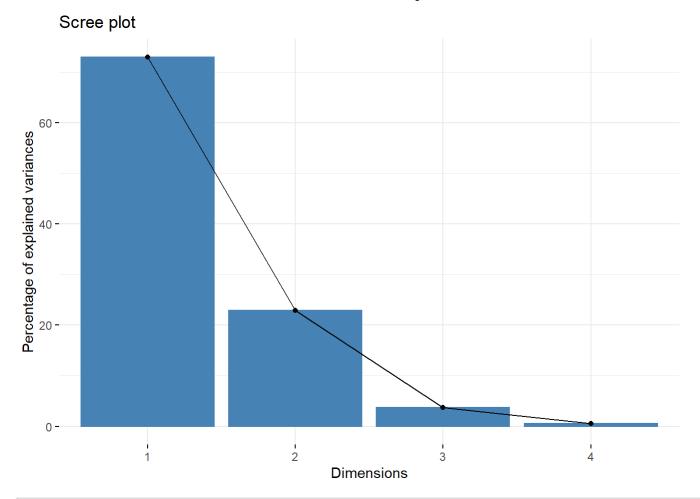


- Diameter = 4, which indicates that this network is not very extensive: the maximum number of nodes that the 2 nodes furthest away from each other is is 4.
- Avg. Path Length = 2.267, again, indicating a shallow network.

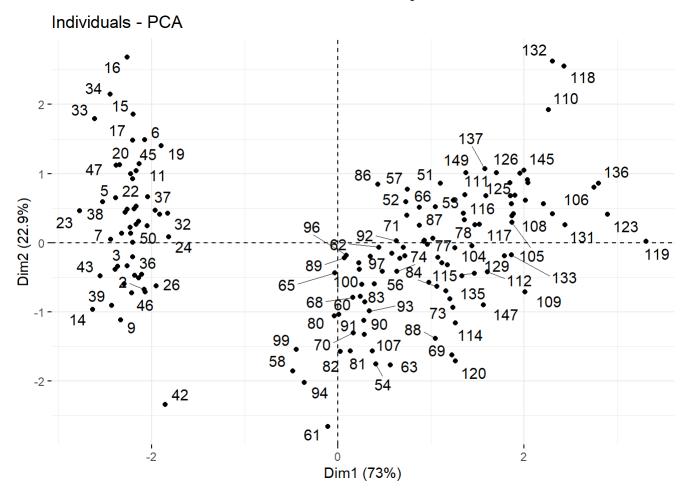
#### Exercise 5

• Use any data (iris is a good data) to visualize the relationship between variables using PCA, CA, or MCA.

```
data("iris")
res.PCA <- PCA(iris %>% select(-Species), graph = FALSE)
fviz_eig(res.PCA)
```

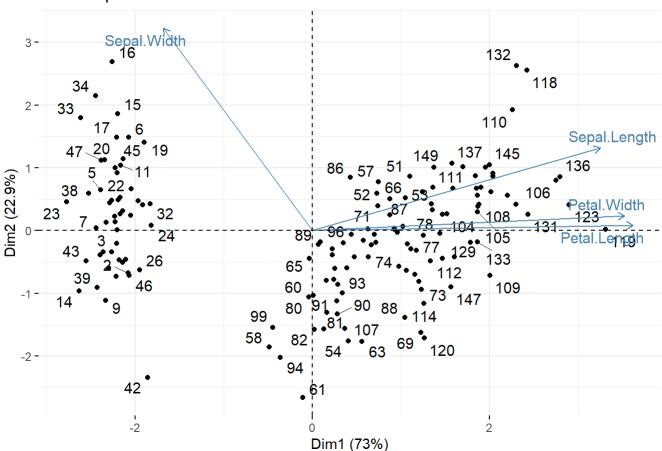


fviz\_pca\_ind(res.PCA, repel = TRUE)



fviz\_pca\_biplot(res.PCA, repel = TRUE)



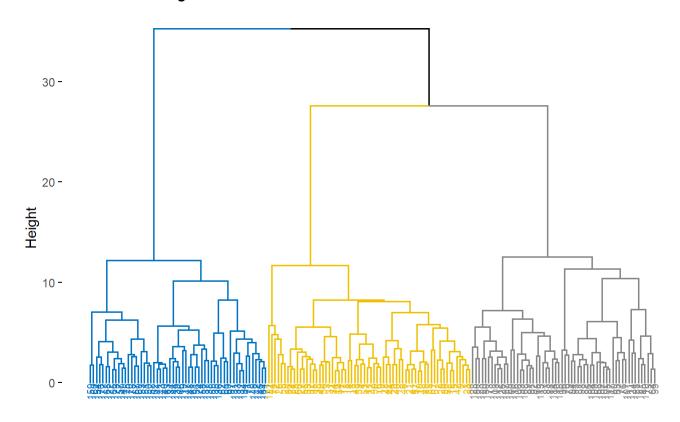


• Use any data to visualize variables and observations using heat map and hierarchical clustering method. You can use the same data that you used when doing clustering in Clustvis.

```
wine <- read_csv("datasets/wine-clustering.csv")
wine_scaled <- scale(wine)

res.hc <- hclust(dist(wine_scaled), method = "ward.D2")
fviz_dend(res.hc, cex=0.5, k=3, palette="jco")</pre>
```

#### Cluster Dendrogram



pheatmap(t(wine\_scaled), cuttree\_cols=3)

