

**Homework 10**

**Problem.** We consider a data set extracted from the 2017-2018 US National Health and Nutrition Examination Survey consisting of 230 participants aged between 20-25 years. For each participant, data were collected about body measures to estimate the prevalence of overweight and obesity. Data is stored in bodydata.csv

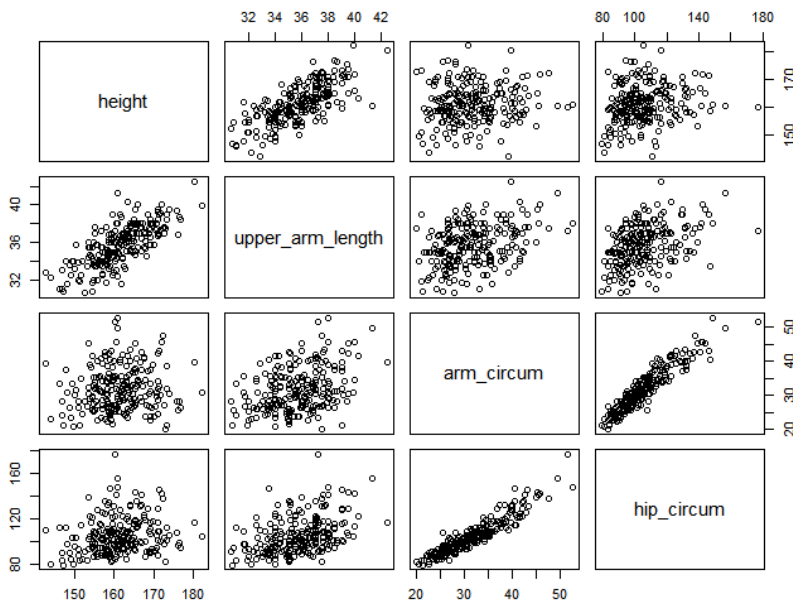
- Draw a scatterplot of the data in pairs using the R command pairs as in the notes.
- Compute the correlation of the data matrix. Note: some data row have missing data. You will need to use this version of the command: `cor(bodydata,use = "complete.obs")`
- Plot the correlation matrix using numbers as well as circles to display the size of correlation coefficients.
- Apply hierarchical clustering as in the lectures on the correlation plot using 2 clusters.
- Compute the p-values on the correlation matrix.
- Analyze the results: which variables are strongly correlated (correlation coefficient > 0.7) to each other? Which variables are not statistically correlated (use alpha = 0.05)?

**SOLUTION**

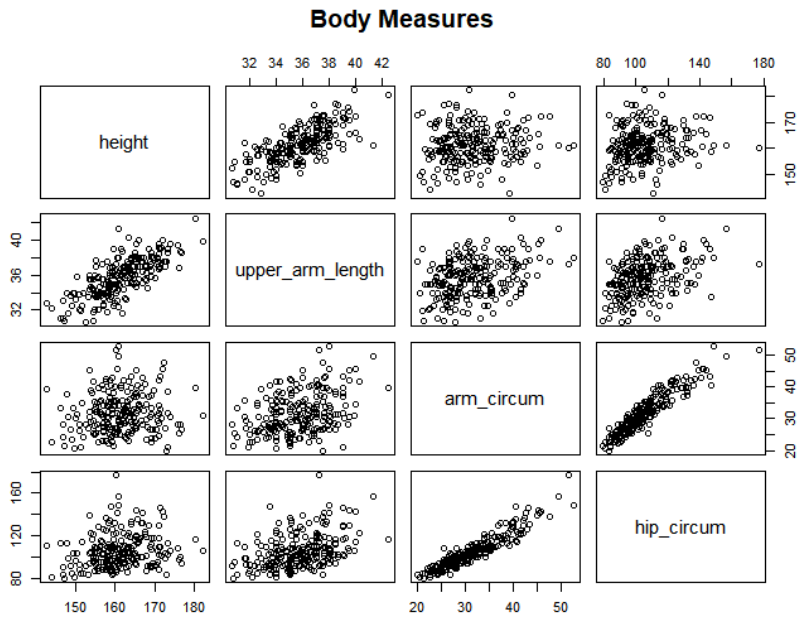
```
> bodydata <- read.csv("C:/Users/dlabate/Desktop/Teaching/ma4310/bodydata.csv")
> head(bodydata)
  height upper_arm_length arm_circum hip_circum
1  158.4             36.0      26.5      101.1
2  164.7             38.1      30.5       97.4
3  156.9             34.0      28.5      101.7
4  158.1             35.0      22.2       88.7
5  158.2             35.0      32.0      100.3
6  162.0             34.4      32.7       99.3
> dim(bodydata)
[1] 230  4
```

```
a)
> pairs(bodydata[c("height", "hip_circum")])

> pairs(bodydata[c("height", "upper_arm_length", "arm_circum", "hip_circum")])
```



```
>plot(bodydata, main = "Body Measures")
```



b)

```
> cor(bodydata,use = "complete.obs")
```

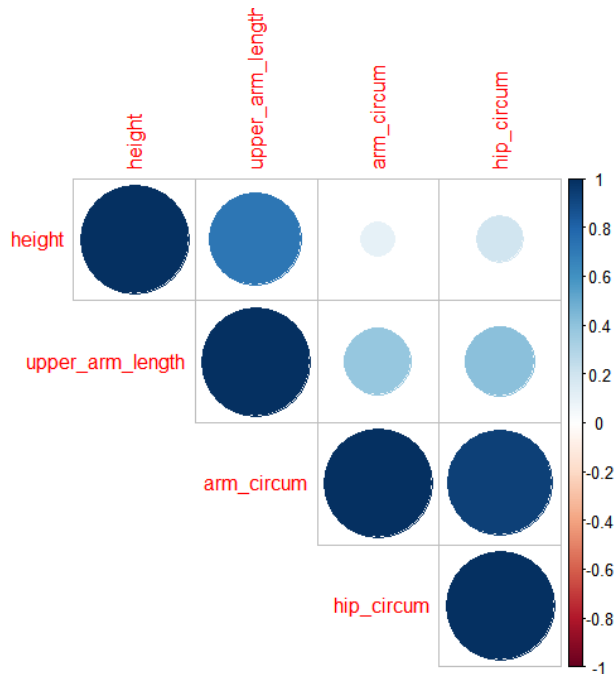
	height	upper_arm_length	arm_circum	hip_circum
height	1.0000000	0.7259228	0.1061935	0.1942494
upper_arm_length	0.7259228	1.0000000	0.3843140	0.4187889
arm_circum	0.1061935	0.3843140	1.0000000	0.9332575
hip_circum	0.1942494	0.4187889	0.9332575	1.0000000

c)

```
> corrplot(cor(bodydata,use = "complete.obs"),method = "number",type = "upper")
```

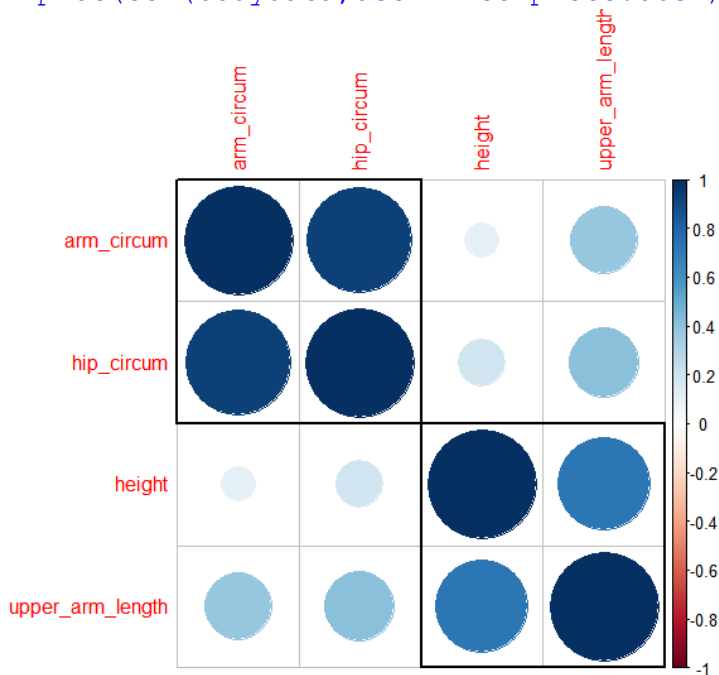


```
> corrplot(cor(bodydata,use = "complete.obs"),method = "circle",type = "upper")
```



d)

```
> corrplot(cor(bodydata,use = "complete.obs"), order = "hclust", addrect = 2)
```



e)

```
> X <- as.matrix(bodydata)
> res <- rcorr(X)
> round(res$P, 3)
```

	height	upper_arm_length	arm_circum	hip_circum
height	NA	0	0.176	0.004
upper_arm_length	0.000	NA	0.000	0.000
arm_circum	0.176	0	NA	0.000
hip_circum	0.004	0	0.000	NA

f)

**CONCLUSION:**

- The variables `arm_circum` and `hip_circum` are strongly correlated; so are the variables `height` and `upper_arm_length`.
- The variables `height` and `arm_circum` are not statistically correlated.

**Problem 2.** Load the Iris dataset in R

```
library(datasets)
data(iris)
```

It is a data frame with 150 samples (rows) and 5 variables (columns) named `Sepal.Length`, `Sepal.Width`, `Petal.Length`, `Petal.Width`, and `Species`.

- Generate a statistical summary of the data
- Apply PCA and LDA analysis to the data and generate a plot to represent the features with respect to 2 dimensions. Make sure to label the axes appropriately and display the different species using different colors or symbols.
- Concisely discuss the performance of the two methods.

**SOLUTION**

```
> library(datasets)
> data(iris)
```

a)

```
> summary(iris)
```

Sepal.Length	Sepal.Width	Petal.Length	Petal.Width	Species
Min. :4.300	Min. :2.000	Min. :1.000	Min. :0.100	setosa :50
1st Qu.:5.100	1st Qu.:2.800	1st Qu.:1.600	1st Qu.:0.300	versicolor:50
Median :5.800	Median :3.000	Median :4.350	Median :1.300	virginica :50
Mean :5.843	Mean :3.057	Mean :3.758	Mean :1.199	
3rd Qu.:6.400	3rd Qu.:3.300	3rd Qu.:5.100	3rd Qu.:1.800	
Max. :7.900	Max. :4.400	Max. :6.900	Max. :2.500	

b) We first apply PCA and display the PCA coordinates

```
> irisPCA=prcomp(iris[,1:4],scale=TRUE)
> summary(irisPCA)
```

Importance of components:

	PC1	PC2	PC3	PC4
Standard deviation	1.7084	0.9560	0.38309	0.14393
Proportion of Variance	0.7296	0.2285	0.03669	0.00518
Cumulative Proportion	0.7296	0.9581	0.99482	1.00000

Next we apply LDA and display the LDA coordinates

```
> library(MASS)
> irisLDA <- lda(Species ~ Sepal.Length + Sepal.Width + Petal.Length + Petal.Width
,iris, prior=c(1,1,1)/3)
```

(alternatively)

```
> irisLDA <- lda(Species~.,data=iris)
> irisLDA
```

Call:

```
lda(Species ~ ., data = iris)
```

Prior probabilities of groups:  
setosa versicolor virginica  
0.3333333 0.3333333 0.3333333

Group means:

	Sepal.Length	Sepal.Width	Petal.Length	Petal.Width
setosa	5.006	3.428	1.462	0.246
versicolor	5.936	2.770	4.260	1.326
virginica	6.588	2.974	5.552	2.026

Coefficients of linear discriminants:

	LD1	LD2
Sepal.Length	0.8293776	0.02410215
Sepal.Width	1.5344731	2.16452123
Petal.Length	-2.2012117	-0.93192121
Petal.Width	-2.8104603	2.83918785

Proportion of trace:

LD1	LD2
0.9912	0.0088

Next we display the first 2 principal components of the PCA and LDA methods

We start with the first two principal components PC1 and PC2 (figure next page)

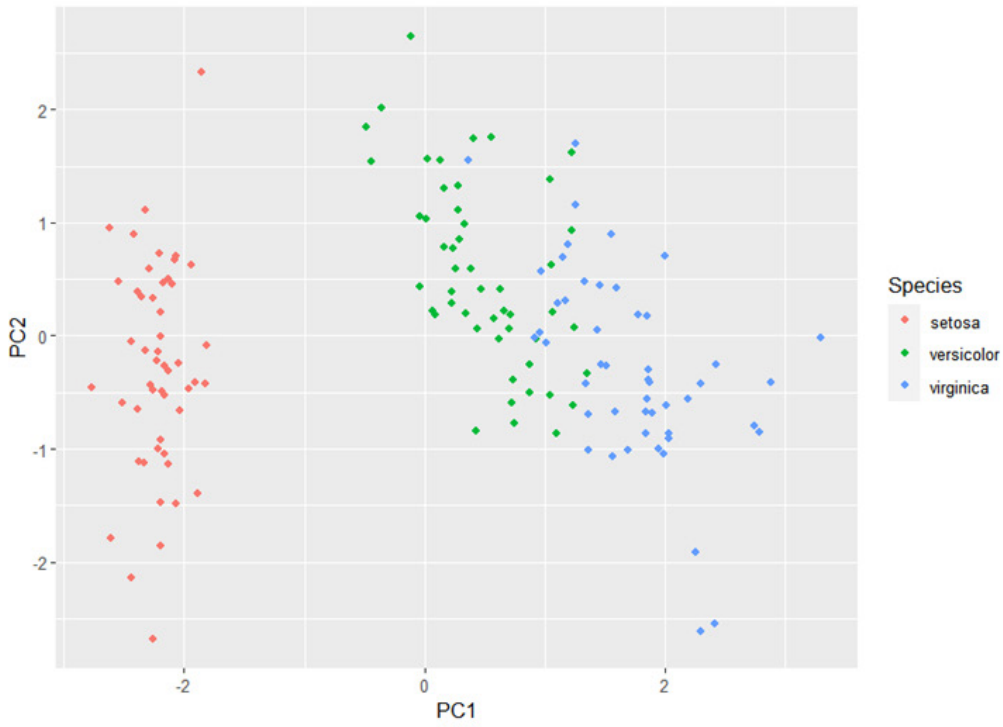
```
> irisPCA_frame <-data.frame(irisPCA$x,Species=iris$Species)
> ggplot(irisPCA_frame,aes(x=PC1,y=PC2,color=Species))+geom_point()+labs(title="PCA of iris data",x="PC1",y="PC2")
```

Next we plot the first two LDA components LD1 and LD2. For convenience we plot the PCA and LDA figures side by side in the next page

```
> irisLDAmode<-predict(irisLDA)
> irisLDA_frame <-data.frame(irisLDAmode$x,Species=iris$Species)
> ggplot(irisLDA_frame,aes(x=LD1,y=LD2,color=Species))+geom_point()+labs(title="LDA of iris data",x="LD1",y="LD2")
```

- d) The plots show that LDA separates the 3 species almost exactly using the single coordinate LD1. By contrast, The versicolor and virgica species has some overlap in the PCA projection. Thus, the LDA approach results in a more effective separation of the 3 species.**

PCA of iris data



LDA of iris data

