

# Lab1

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## Question 1. Test of outliers

a

```
dataset <- read.table("T1-9.dat")

mean <- colMeans(dataset[, 2:8])
Sx <- cov(dataset[, 2:8])
D2M <- mahalanobis(dataset[, 2:8], mean, Sx)
```

To answer this question properly we need to calculate again, as we did on the previous lab assignment, the Mahalanobis vector, which can be easily done.

	ARG	AUS	AUT	BEL	BER	BRA
6.6500027	3.9356705	4.3680848	2.4623238	7.6305669	2.1885513	
	CAN	CHI	CHN	COL	COK	CRC
3.0916940	4.1542602	3.0167240	5.3994790	19.8340006	4.4647286	
	CZE	DEN	DOM	FIN	FRA	GER
10.9014563	1.3174764	4.7284159	2.7556601	5.0862790	3.4467449	
	GBR	GRE	GUA	HUN	INA	IND
3.5572682	9.5403223	3.6846053	1.7086674	4.7128021	4.2129432	
	IRL	ISR	ITA	JPN	KEN	KORS
5.4848378	2.5421901	1.4553927	3.2695339	7.6254662	8.0349184	
	KORN	LUX	MAS	MRI	MEX	MYA
26.1671415	11.1088463	8.2371487	6.6649850	14.2309322	2.0429608	
	NED	NZL	NOR	PNG	PHI	POL
5.9522081	1.3505689	6.8880631	30.5072477	9.0658836	1.3311122	
	POR	ROM	RUS	SAM	SIN	ESP
2.3695737	6.3491309	3.7506462	35.0140631	8.0163948	1.4160474	
	SWE	SUI	TPE	THA	TUR	USA
0.7846063	3.2910927	10.1839962	7.8152191	8.0453684	9.1556967	

Once this have been done, we can proceed to run the chi-square test on this Mahalanobis data, which returns this data:

```
chisq.test(D2M)
```

Chi-squared test for given probabilities

data: D2M

X-squared = 364.07, df = 53, p-value < 2.2e-16

After that, and thanks to the outliers package (<https://cran.rproject.org/web/packages/outliers/index.html>) the outliers detection become considerably easier, proceeding as follows:

```
setNames(scores(D2M, type="chisq", prob = 0.999), dataset[, 1])
```

ARG	AUS	AUT	BEL	BER	BRA	CAN	CHI	CHN	COL	COK	CRC
FALSE	FALSE	FALSE	FALSE	FALSE	FALSE	FALSE	FALSE	FALSE	FALSE	FALSE	FALSE
CZE	DEN	DOM	FIN	FRA	GER	GBR	GRE	GUA	HUN	INA	IND
FALSE	FALSE	FALSE	FALSE	FALSE	FALSE	FALSE	FALSE	FALSE	FALSE	FALSE	FALSE
IRL	ISR	ITA	JPN	KEN	KORS	KORN	LUX	MAS	MRI	MEX	MYA
FALSE	FALSE	FALSE	FALSE	FALSE	FALSE	FALSE	FALSE	FALSE	FALSE	FALSE	FALSE
NED	NZL	NOR	PNG	PHI	POL	POR	ROM	RUS	SAM	SIN	ESP
FALSE	FALSE	FALSE	TRUE	FALSE	FALSE	FALSE	FALSE	FALSE	TRUE	FALSE	FALSE

```
SWE  SUI  TPE  THA  TUR  USA
FALSE FALSE FALSE FALSE FALSE FALSE
```

This output on the console shows, as a result of the `setNames` function, the country's acronym over a TRUE/FALSE indicator, which highlights the observations whose Mahalanobis distance lies beyond the indicated percentile.

In this first case, the function works with the given significance of  $\alpha=0.001$ . Analyzing the table above, it can be noticed how only two countries are being identified as outliers. Those are: Papua New Guinea (PNG, 30.5072477) and Samoa (SAM, 35.0140631), which are the two most extreme cases amongst all the calculated distances.

The study could be finished here but seems effective to go a bit deeper and try to modify this quantile through some modifications in order to find a better differentiation between outliers and average observations. More specifically, the Bonferroni correction enables the calibration of this parameter. Supposing that, for our dataset, each country would be compared once with each other country on the list, the number of total combinations comes to 1431. As a consequence, the chosen correction method results in a new  $\alpha$  value of  $1 - 0.761 = 0.239$ . Running again the `scores` function with this new parameter returns the following:

```
setNames(scores(D2M, type="chisq", prob = 0.761), dataset[, 1])
```

```
ARG  AUS  AUT  BEL  BER  BRA  CAN  CHI  CHN  COL  COK  CRC
FALSE FALSE FALSE FALSE FALSE FALSE FALSE FALSE FALSE FALSE TRUE FALSE
CZE  DEN  DOM  FIN  FRA  GER  GBR  GRE  GUA  HUN  INA  IND
FALSE FALSE FALSE FALSE FALSE FALSE FALSE FALSE FALSE FALSE FALSE FALSE
IRL  ISR  ITA  JPN  KEN  KORS  KORN  LUX  MAS  MRI  MEX  MYA
FALSE FALSE FALSE FALSE FALSE FALSE TRUE FALSE FALSE FALSE FALSE FALSE
NED  NZL  NOR  PNG  PHI  POL  POR  ROM  RUS  SAM  SIN  ESP
FALSE FALSE FALSE TRUE FALSE FALSE FALSE FALSE FALSE TRUE FALSE FALSE
SWE  SUI  TPE  THA  TUR  USA
FALSE FALSE FALSE FALSE FALSE FALSE
```

With this new limit, two new countries are identified as outliers: Cook Islands (COK, 19.8340006) and North Korea (KORN, 26.1671415). Those two values are far from being standard, therefore can be concluded that the correction is accurate. Beyond this conclusion,  $\alpha$  could be even more reduced, depending on where we want to place the “outlier threshold”. As a last example, setting  $\alpha$  to 0.35 makes also Mexico (MEX, 14.2309322) an outlier:

```
setNames(scores(D2M, type="chisq", prob = 0.65), dataset[, 1])
```

```
ARG  AUS  AUT  BEL  BER  BRA  CAN  CHI  CHN  COL  COK  CRC
FALSE FALSE FALSE FALSE FALSE FALSE FALSE FALSE FALSE FALSE TRUE FALSE
CZE  DEN  DOM  FIN  FRA  GER  GBR  GRE  GUA  HUN  INA  IND
FALSE FALSE FALSE FALSE FALSE FALSE FALSE FALSE FALSE FALSE FALSE FALSE
IRL  ISR  ITA  JPN  KEN  KORS  KORN  LUX  MAS  MRI  MEX  MYA
FALSE FALSE FALSE FALSE FALSE FALSE TRUE FALSE FALSE FALSE TRUE FALSE
NED  NZL  NOR  PNG  PHI  POL  POR  ROM  RUS  SAM  SIN  ESP
FALSE FALSE FALSE TRUE FALSE FALSE FALSE FALSE FALSE TRUE FALSE FALSE
SWE  SUI  TPE  THA  TUR  USA
FALSE FALSE FALSE FALSE FALSE FALSE
```

## b

The reason for this is that Euclidean distance simply computes the ordinary straight line between two points and the Mahalanobis distance takes the covariate into account, which leads to elliptic decision boundaries in 2D, therefore narrower than the Euclidean circular, which helps to properly identify the outliers, as done in the previous section

## Question 2. Test, confidence region and confidence intervals for a mean vector

Look at the bird data in file T5-12.dat and solve Exercise 5:20 of Johnson, Wichern. Do not use any extra R package or built-in test but code all required matrix calculations. You MAYNOT use loops!

a

```
bird_data <- read.table("T5-12.dat")
mu <- c(190, 275) #given mu values
x_bar <- colMeans(bird_data)

calcReqVals = function(data){
  n <- nrow(data)
  p <- ncol(data)
  conf <- 0.05
  S <- cov(data)
  eig <- eigen(S)
  x_ <- colMeans(data)

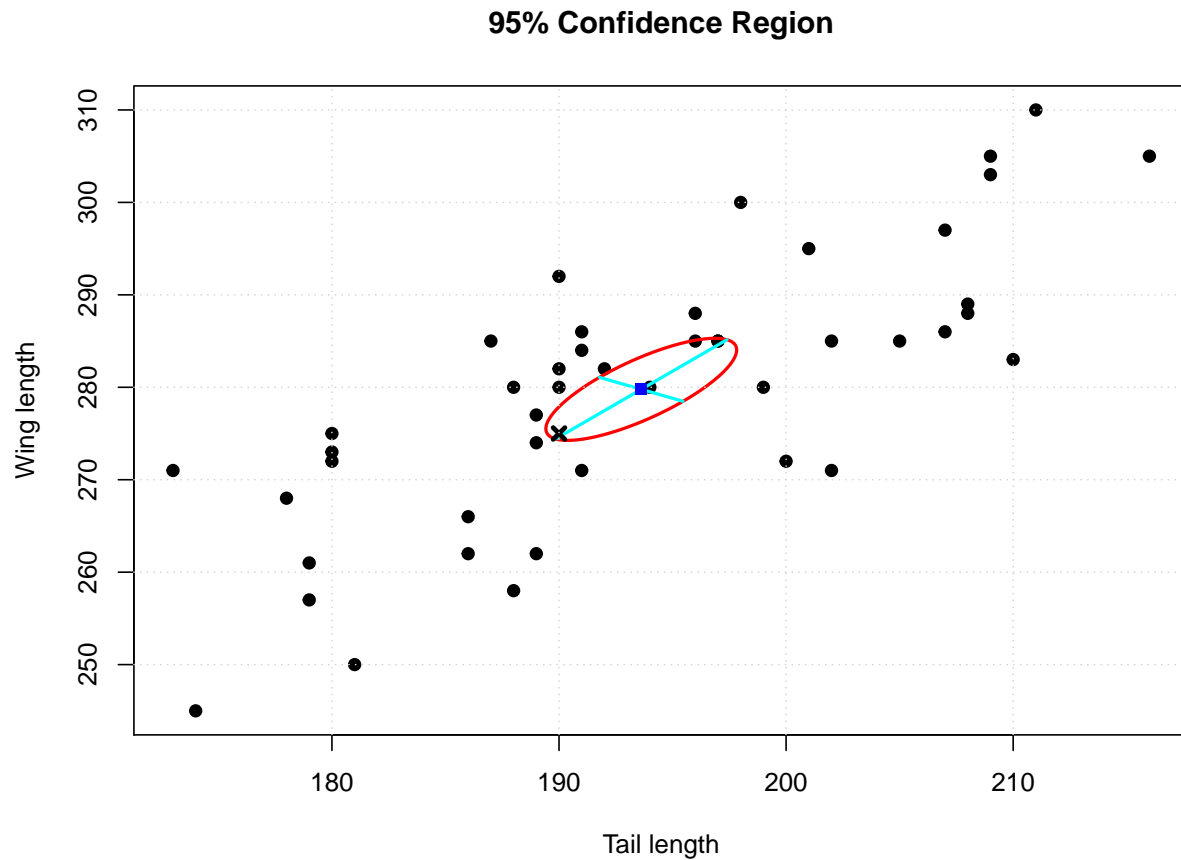
  quantile <- qf(1 - conf, df1=p, df2=n - p)
  scale <- sqrt(eig$values * p * (n - 1) * quantile / (n * (n - p)))
  scaled <- eig$vector %% diag(scale) # scale eigenvectors to length = square-root
  xMat <- rbind(x_[1] + scaled[1, ], x_[1] - scaled[1, ])
  yMat <- rbind(x_[2] + scaled[2, ], x_[2] - scaled[2, ])

  angles <- seq(0, 2 * pi, length.out=200)
  ellBase <- cbind(scale[1]*cos(angles), scale[2]*sin(angles)) # making a circle base...
  ellax <- eig$vector %% t(ellBase)

  return(list("ellax"=ellax, "xMat"=xMat, "yMat"=yMat))
}

out = calcReqVals(bird_data)

#Plotting the confidence region
plot(bird_data, pch=19, xlab="Tail length",
      ylab="Wing length", main="95% Confidence Region")
lines((out$ellax + x_bar)[1, ], (out$ellax + x_bar)[2, ], asp=1, type="l", lwd=2, col="red")
matlines(out$xMat, out$yMat, lty=1, lwd=2, col="cyan") #
points(mu[1], mu[2], pch=4, col="black", lwd=3)
grid()
points(x_bar[1], x_bar[2], type="p", col="blue", pch=15)
```



Since the known mean of the male data lies in the confidence interval for the mean of the female data, we fail to reject the hypothesis that the male and the female have the same mean.

We took most of the above code with understanding from stackoverflow: <https://stats.stackexchange.com/questions/9898/how-to-plot-an-ellipse-from-eigenvalues-and-eigenvectors-in-r>

We also asked for few help from Maria with respect to code and theory.

b

```
tsq95_intervals <- function(data) {
  conf = 0.05
  n <- nrow(data)
  p <- ncol(data)
  x_bar <- colMeans(data)
  S <- cov(data)
  offset <- sqrt(p * (n - 1) * qf(1 - conf, df1=p, df2=n - p) / (n - p) * diag(S) / n)
  rbind(x_bar - offset, x_bar + offset)
}

bon95_intervals <- function(data) {
  conf = 0.05
  n <- nrow(data)
```

```

p <- ncol(data)
x_bar <- colMeans(data)
S <- cov(data)
offset <- sqrt(diag(S) / n) * qt(1 - conf / (2 * p), df=n - 1)
rbind(x_bar - offset, x_bar + offset)
}

cat("\n95% T-square interval\n")

```

95% T-square interval

```
tsq95_intervals(bird_data)
```

```

      V1      V2
[1,] 189.4217 274.2564
[2,] 197.8227 285.2992

```

```
cat("\n95% Bonferroni interval\n")
```

95% Bonferroni interval

```
bon95_intervals(bird_data)
```

```

      V1      V2
[1,] 189.8216 274.7819
[2,] 197.4229 284.7736

```

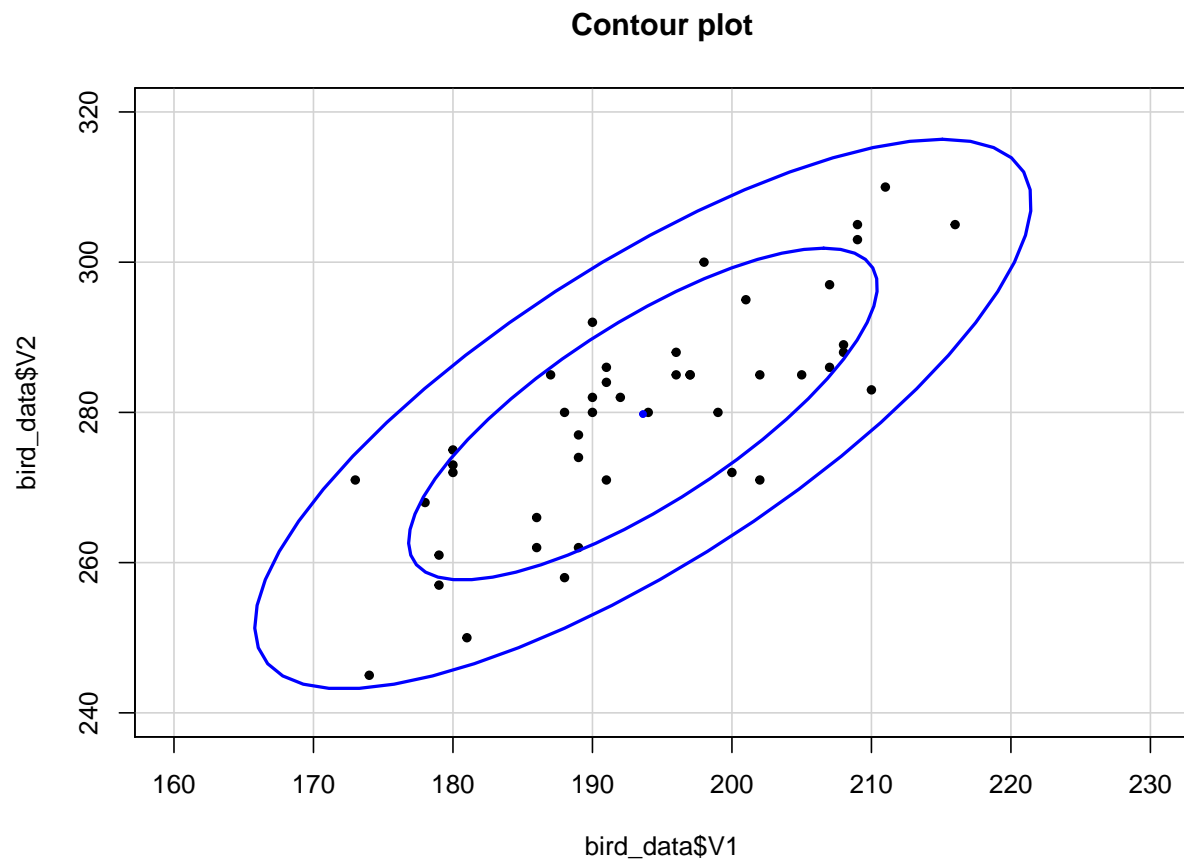
The only advantage we could find for T-squared intervals over Bonferroni interval is that, T-squared intervals account for the correlation between variates, while Bonferroni does not. Bonferroni is useful when we have to find the CI for the individual component.

**c**

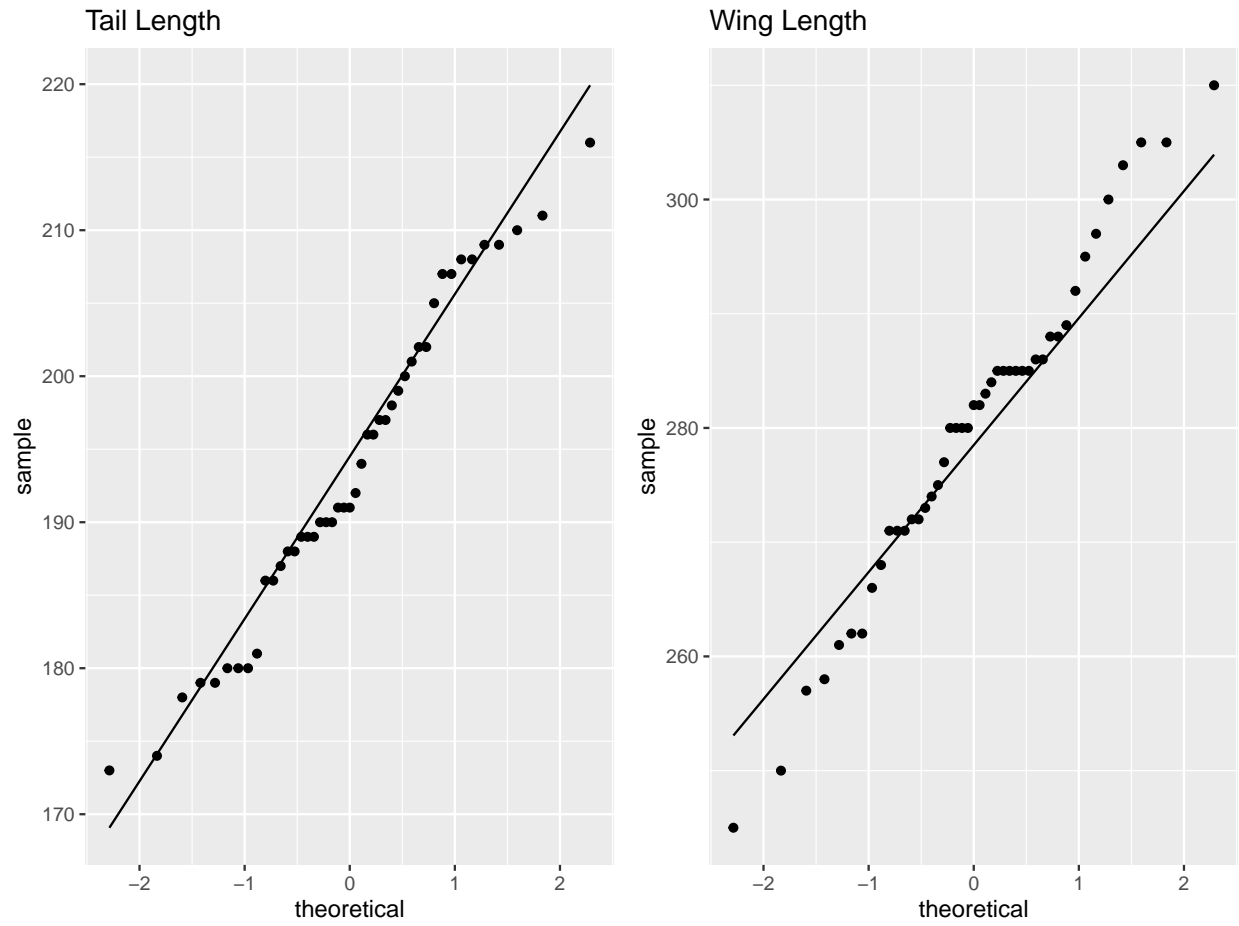
```

dataEllipse(x=bird_data$V1, y=bird_data$V2, pch=20, levels=c(0.68, 0.95),
            xlim=c(160, 230), ylim=c(240, 320), center.cex=0.5, main="Contour plot")

```



```
grid.arrange(ggplot(data = bird_data, aes(sample = V1)) +  
  stat_qq() + stat_qq_line() + ggtitle("Tail Length"),  
ggplot(data = bird_data, aes(sample = V2)) +  
  stat_qq() + stat_qq_line() +  
  ggtitle("Wing Length"),ncol=2)
```



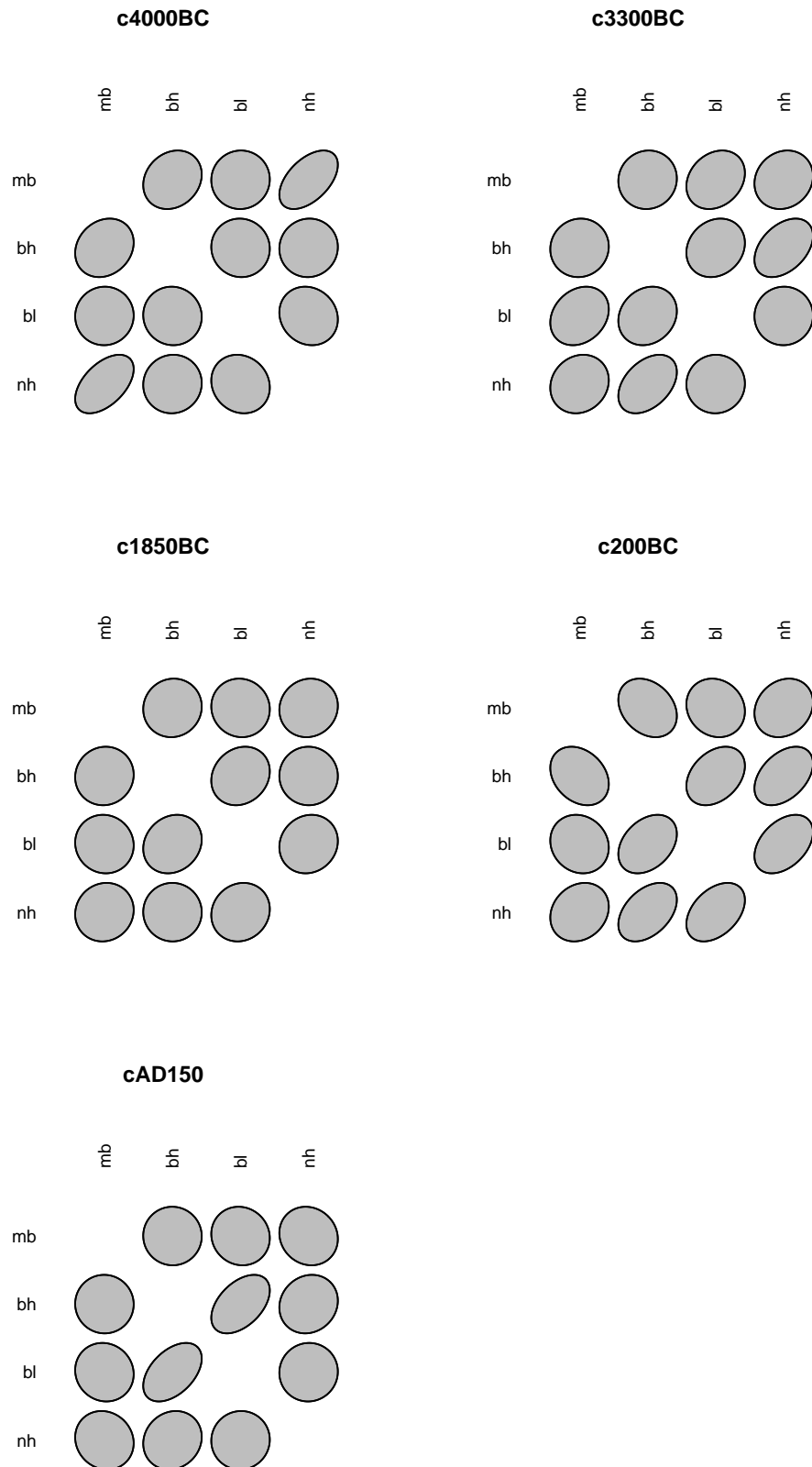
Since the data points go along the line in the qqplots, it shows that the data is normally distributed. A bivariate normal distribution would be a viable population model.



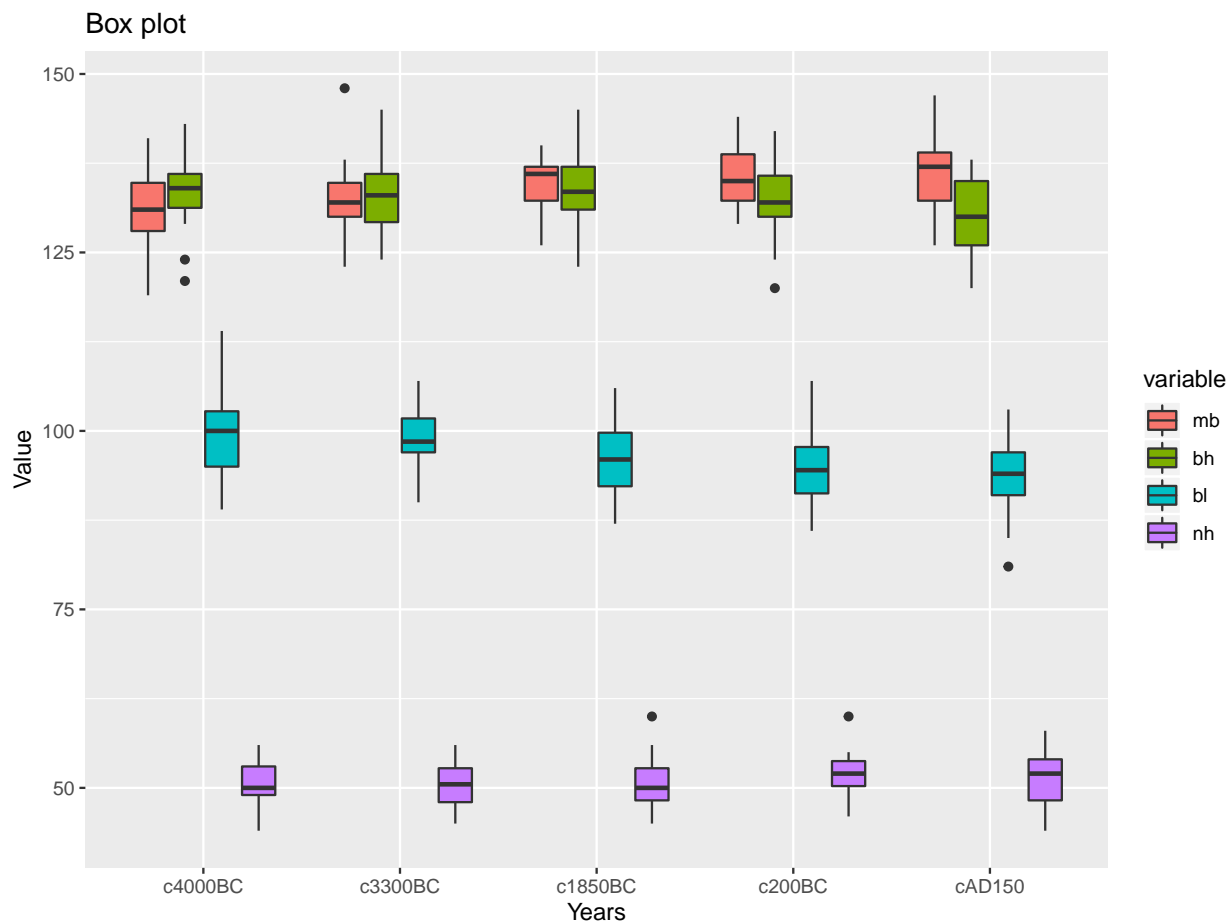


### Question 3. Comparison of mean vectors (one-way MANOVA)

a



From above correlation plot over 5 time period the common thing which we noticed is the positive correlation between basibregmatic height and basialiveolar length. Moreover, the correlation become more stronger between them over the period. In AD 150 the correlation between nh and mb was negative which was unprecedented.



From this the only thing which is informative is that the mean bl length decreased over the years while rest of the mean measurement did not follow any pattern. Also the distance between mean bh and mb increased towards the end of AD150. As maximal breadth of the skull increased, the height of the skull increased as well. Were brains were increasing in size ?

**b**

Table 1: Mean of covariates in different groups

Group	mb	bh	bl	nh
c4000BC	131.3667	133.6000	99.16667	50.53333
c3300BC	132.3667	132.7000	99.06667	50.23333
c1850BC	134.4667	133.8000	96.03333	50.56667
c200BC	135.5000	132.3000	94.53333	51.96667
cAD150	136.1667	130.3333	93.50000	51.36667

```
manova_fit <- manova(cbind(mb, bh, bl, nh) ~ sk$epoch, sk)
summary(manova_fit, test="Pillai")
```

```

      Df  Pillai approx F num Df den Df      Pr(>F)
sk$epoch    4 0.35331    3.512    16    580 4.675e-06 ***
Residuals 145
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

Pillai's trace is used as a test statistic in MANOVA. It is a positive valued statistic ranging from 0 to 1. Increasing values means that effects are contributing more to the model; you should reject the null hypothesis for large values.

Pillai's trace is considered to be the most powerful and robust statistic for general use, especially for departures from assumptions – Wikipedia

Here the p value is less than 0.05 so we reject the null hypothesis which is ,that the vector means are same across group.

**c**

```

t2_interval <- function(data, conf) {
  n <- nrow(data)
  p <- ncol(data)
  x_ <- colMeans(data)
  S <- cov(data)
  offset <- sqrt(p * (n - 1) * qf(1 - conf, df1=p, df2=n - p) / (n - p) * diag(S) / n)
  rbind(x_ - offset, x_ + offset)
}

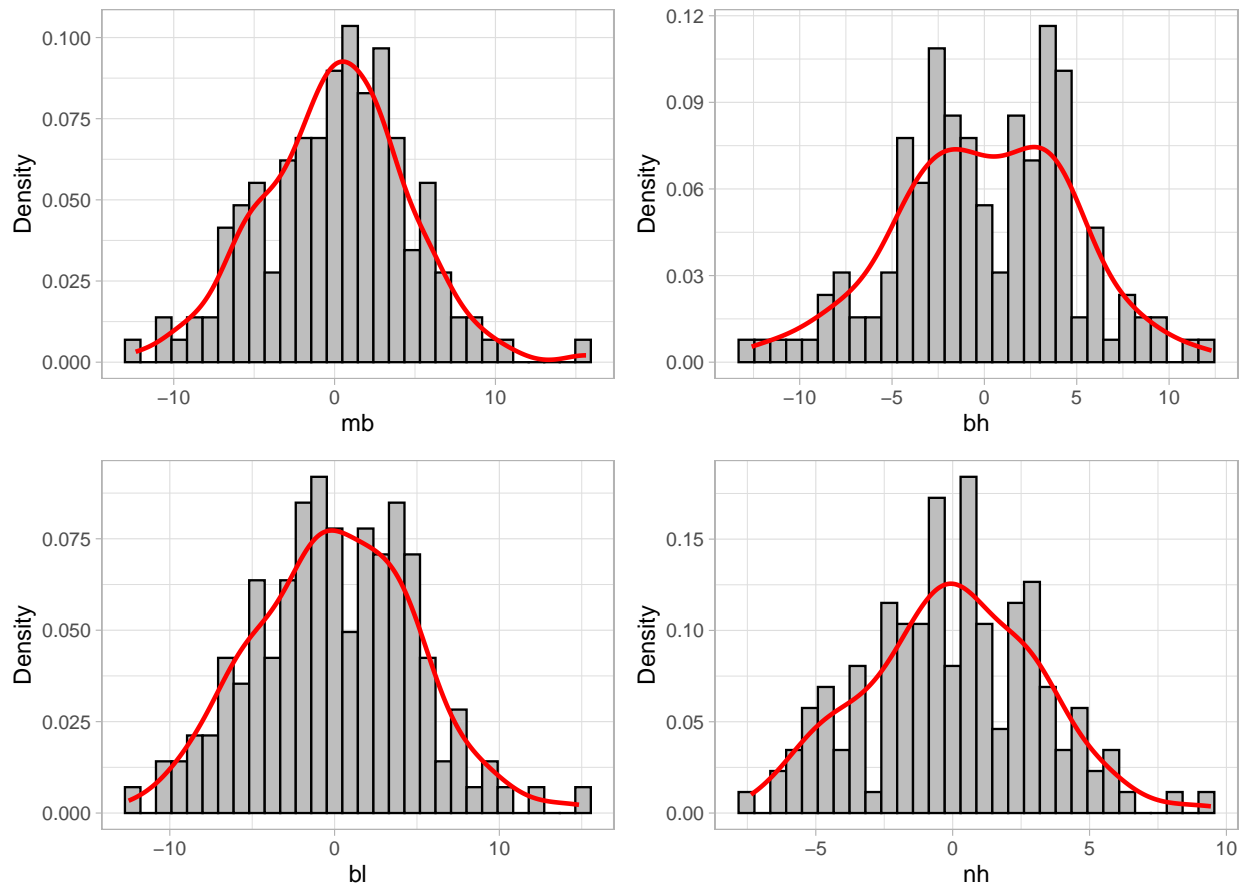
conf_interval <- t2_interval(sk[,2:5],0.05)

```

Table 2: 95% CI

mb	bh	bl	nh
132.7147	131.2755	95.076	50.10776
135.2320	133.8178	97.844	51.75890

Density plot and histogram



The variables are nearly normally distributed.

## Appendix

```
library(outliers)
library(heplots)
library(kableExtra)
library(gridExtra)
library(ggplot2)
library(reshape2)
library(ellipse)
library(car)
knitr::opts_chunk$set(
  message = FALSE,
  warning = FALSE,
  comment = NA,
  fig.width=8,
  fig.height=6
)
dataset <- read.table("T1-9.dat")

mean <- colMeans(dataset[, 2:8])
Sx <- cov(dataset[, 2:8])
D2M <- mahalanobis(dataset[, 2:8], mean, Sx)
setNames(D2M, dataset[, 1])
chisq.test(D2M)
setNames(scores(D2M, type="chisq", prob = 0.999), dataset[, 1])
setNames(scores(D2M, type="chisq", prob = 0.761), dataset[, 1])

setNames(scores(D2M, type="chisq", prob = 0.65), dataset[, 1])
bird_data <- read.table("T5-12.dat")
mu <- c(190, 275) #given mu values
x_bar <- colMeans(bird_data)

calcReqVals = function(data){
  n <- nrow(data)
  p <- ncol(data)
  conf <- 0.05
  S <- cov(data)
  eig <- eigen(S)
  x_ <- colMeans(data)

  quantile <- qf(1 - conf, df1=p, df2=n - p)
  scale <- sqrt(eig$values * p * (n - 1) * quantile / (n * (n - p)))
  scaled <- eig$vector %% diag(scale) # scale eigenvectors to length = square-root
  xMat <- rbind(x_[1] + scaled[1, ], x_[1] - scaled[1, ])
  yMat <- rbind(x_[2] + scaled[2, ], x_[2] - scaled[2, ])

  angles <- seq(0, 2 * pi, length.out=200)
  ellBase <- cbind(scale[1]*cos(angles), scale[2]*sin(angles)) # making a circle base...
  ellax <- eig$vector %% t(ellBase)

  return(list("ellax"=ellax, "xMat"=xMat, "yMat"=yMat))
}
```

```

out = calcReqVals(bird_data)

#Plotting the confidence region
plot(bird_data,pch=19, xlab="Tail length",
      ylab="Wing length", main="95% Confidence Region")
lines((out$ellax + x_bar)[1, ], (out$ellax + x_bar)[2, ], asp=1, type="l", lwd=2, col="red")
matlines(out$xMat, out$yMat, lty=1, lwd=2, col="cyan") #
points(mu[1], mu[2], pch=4, col="black", lwd=3)
grid()
points(x_bar[1],x_bar[2], type="p", col="blue", pch=15)
tsq95_intervals <- function(data) {
  conf = 0.05
  n <- nrow(data)
  p <- ncol(data)
  x_bar <- colMeans(data)
  S <- cov(data)
  offset <- sqrt(p * (n - 1) * qf(1 - conf, df1=p, df2=n - p) / (n - p) * diag(S) / n)
  rbind(x_bar - offset, x_bar + offset)
}

bon95_intervals <- function(data) {
  conf = 0.05
  n <- nrow(data)
  p <- ncol(data)
  x_bar <- colMeans(data)
  S <- cov(data)
  offset <- sqrt(diag(S) / n) * qt(1 - conf / (2 * p), df=n - 1)
  rbind(x_bar - offset, x_bar + offset)
}

cat("\n95% T-square interval\n")
tsq95_intervals(bird_data)

cat("\n95% Bonferroni interval\n")
bon95_intervals(bird_data)

dataEllipse(x=bird_data$V1, y=bird_data$V2, pch=20, levels=c(0.68, 0.95),
            xlim=c(160, 230), ylim=c(240, 320), center.cex=0.5, main="Contour plot")

grid.arrange(ggplot(data = bird_data, aes(sample = V1)) +
              stat_qq() + stat_qq_line() + ggtitle("Tail Length"),
              ggplot(data = bird_data, aes(sample = V2)) +
              stat_qq() + stat_qq_line() +
              ggtitle("Wing Length"),ncol=2)

sk <- Skulls
#aggregate(sk["c40,2:5"],by=list(sk$epoch), FUN =sd)

corr_mat <- cor(sk[sk$epoch=="c4000BC",2:5])
corr_mat2 <- cor(sk[sk$epoch=="c3300BC",2:5])
corr_mat3 <- cor(sk[sk$epoch=="c1850BC",2:5])
corr_mat4 <- cor(sk[sk$epoch=="c200BC",2:5])

```

```

corr_mat5 <- cor(sk[sk$epoch=="cAD150",2:5])

par(mfrow=c(3,2))

plotcorr(corr_mat,outline=TRUE,diag=FALSE,main="c4000BC")
plotcorr(corr_mat2, diag=FALSE,main="c3300BC")
plotcorr(corr_mat3, diag=FALSE,main ="c1850BC")
plotcorr(corr_mat4, diag=FALSE,main ="c200BC")
plotcorr(corr_mat5, diag=FALSE,main ="cAD150")

tall_sk <- reshape2::melt(sk, id="epoch")

ggplot(tall_sk) +
  geom_boxplot(aes(x=factor(epoch), y=value, fill=variable)) +
  ggtitle("Box plot") + xlab("Years") + ylab("Value")
mean_group <- aggregate(sk[,2:5],by=list(sk$epoch), FUN =mean)
colnames(mean_group)[1] <- "Group"

kable(mean_group,caption = "Mean of covariates in different groups") %>% kable_styling(latex_options = "none")

manova_fit <- manova(cbind(mb, bh, bl, nh) ~ sk$epoch, sk)
summary(manova_fit,test="Pillai")
t2_interval <- function(data, conf) {
  n <- nrow(data)
  p <- ncol(data)
  x_ <- colMeans(data)
  S <- cov(data)
  offset <- sqrt(p * (n - 1) * qf(1 - conf, df1=p, df2=n - p) / (n - p) * diag(S) / n)
  rbind(x_ - offset, x_ + offset)
}

conf_interval <- t2_interval(sk[,2:5],0.05)
kable(conf_interval,caption = "95\\% CI") %>%
  kable_styling(latex_options = "hold")
residual <- manova_fit$res %>% as.data.frame()

plot_hist_dens <- function (col_name){
  ggplot(residual,aes_string(col_name) ) +
    geom_histogram(aes(y=..density..),colour="black",fill="gray",bin=10) +
    geom_line(stat="density",colour="red",alpha=1,size=1) + theme_light() +
    xlab(col_name) + ylab("Density")
}

p1 <- plot_hist_dens("mb")
p2 <- plot_hist_dens("bh")
p3 <- plot_hist_dens("bl")
p4 <- plot_hist_dens("nh")

grid.arrange(p1, p2, p3, p4, ncol=2,nrow=2,top = "Density plot and histogram")

```