Complete the following computations on gene means of the Golub data set.
a) Compute the mean expression values for every gene among "ALL" patients.

## a) Rscript:

```
source("http://www.bioconductor.org/biocLite.R")
biocLite()
biocLite("multtest")
library(multtest)
data(golub)
gol.fac <- factor (golub.cl, levels=0:1, labels= c ("ALL","AML"))
meanALL<- apply(golub[,gol.fac=="ALL"],1,mean)
meanALL</pre>
```

b) Compute the mean expression values for every gene among "AML" patients.

# b). Rscript:

```
library(multtest)
data(golub)
gol.fac <- factor (golub.cl, levels=0:1, labels= c ("ALL","AML"))
meanALL<- apply(golub[,gol.fac=="ALL"],1,mean)
meanALL</pre>
```

c) Give the biological names of the three genes with the largest mean expression value among "ALL" patients.

# c)Rscript:

```
library(multtest); data(golub)
gol.fac <- factor(golub.cl,levels=0:1, labels= c("ALL","AML"))
mall <- apply(golub[,gol.fac=="ALL"], 1, mean)
maml <- apply(golub[,gol.fac=="AML"], 1, mean)
o <- order(abs(mall), decreasing=TRUE)
print(golub.gnames[o[1:3],2])
```

#### Answer:

- [1] "GB DEF = Chromosome 1q subtelomeric sequence D1S553"
- [2] "37 kD laminin receptor precursor/p40 ribosome associated protein gene"
- [3] "RPS14 gene (ribosomal protein S14) extracted from Human ribosomal protein S14 gene"

d) Give the biological names of the three genes with the largest mean expression value among "AML" patients.

```
d). Rscript:
library(multtest); data(golub)
gol.fac <- factor(golub.cl,levels=0:1, labels= c("ALL","AML"))
mall <- apply(golub[,gol.fac=="ALL"], 1, mean)
maml <- apply(golub[,gol.fac=="AML"], 1, mean)
o <- order(abs(maml), decreasing=TRUE)
print(golub.gnames[o[1:3],2])
```

#### Answer:

- [1] "GB DEF = mRNA fragment for elongation factor TU (N-terminus)"
- [2] "GB DEF = HLA-B null allele mRNA"
- [3] "Globin, Beta"

Complete the following computations using the Golub data set.

- a) Save the expression values of the first five genes (in the first five rows) for the AML patients in a csv file "AML5.csv."
- A) Rscript:

```
library(multtest); data(golub)
expval<-rbind(golub[1:5,28:38])
write.table (expval, file= "AML5.csv")</pre>
```

#### Answer:

Attached file AML5.csv

b) Save the expression values of the first five genes for the ALL patients in a plain text file "ALL5.txt."

### **Rscript:**

```
library(multtest); data(golub)
eall<-rbind(golub[1:5,1:27])
write.table(eall, file="ALL5.txt")</pre>
```

#### Answer:

Attached file ALL5.txt

c) Compute the standard deviation of the expression values on the first patient, of the 100th to 200th genes (total 101 genes). Rscript:

```
library(multtest); data(golub) stddev<-sd(golub[100:200,1]) stddev
```

#### Answer:

[1] 0.9174976

d) Compute the standard deviation of the expression values of every gene, across all patients. Find the number of genes with standard deviations greater than 1.

### **Rscript:**

```
library(multtest); data(golub)
stdd<-sd(golub[,1:38])
stdd

i<-1
ctr<-0
for (i in golub[i,1:38])
    {
    i<-i+1
    std<-sd(golub[i,1:38])
    if(!is.na(std > 1))
      {
       ctr<-ctr+1
    }
}
ctr</pre>
```

#### Answer:

Standard deviation of expression value of every gene - stdd [1] 0.9998404 number of genes with standard deviations greater than 1-ctr [1] 3

e) Do a scatter plot of the  $101_{st}$  gene expressions against the  $102_{nd}$  gene expressions, labeling the x-axis and the y-axis with the genes' biological names. Do this using xlab= and ylab= control options.

### **Rscript:**

plot(golub[101,1:38],golub[102,1:38],xlab=golub.gnames[101,2],ylab=golub.gnames[102,2])

### **Answer:**

Answer2e.png

Complete a-c using the ALL data set.

Load the ALL data from the ALL library, and use str and openVignette() for further orientation.

a) Use exprs(ALL[,ALL\$BT=="B1"] to extract the gene expressions from the patients in disease stage B1. Produce a histogram of these gene expressions.

# **Rscript:**

```
library(ALL)
data(ALL)
vignette()
hist(exprs(ALL[,ALL$BT=="B1"]),xlab="Gene
Expression",ylab="Frequency",main="Gene expression of patients in disease stage
B1")
```

#### Answer:

Answer3a.png

b) Compute the mean gene expressions over these B1 patients.

# **Rscript**:

```
library(ALL)
data(ALL)
vignette()
meangene<-apply(exprs(ALL[,ALL$BT=="B1"]),1,mean)
meangene</pre>
```

#### Answer:

In Rscript.

c) Give the gene identifiers of the three genes with the largest mean.

### **Rscript:**

library(ALL) data(ALL)

```
vignette()
B1<-exprs(ALL[,ALL$BT=="B1"])
meangene<-apply(B1,1,mean)
meangene
o <- order(abs(meangene), decreasing=TRUE)
print(B1[o[1:3],1])</pre>
```

#### Answer:

```
AFFX-hum_alu_at 31962_at 31957_r_at 13.72101 13.36514 13.05370
```

To complete a and b, work with the "trees" data set that comes with R.

a) Find the type of the trees data object.

# **Rscipt:**

class(trees)

### **Answer:**

"data.frame"

b) Produce a figure with two overlaid scatterplots: girth versus height and girth versus volume. Do the height plot with blue "+" symbols, and do the volume plot with red "o" symbols. You need to set the ylim= control option so that all points from the two plots can show up on the merged figure.

### **Rscript**:

```
plot(trees[,1],trees[,2],col="blue",pch="+", ylim=c(1,90),xlim=c(1,32),xlab="Girth",ylab="Height and Volume",main="Overlaid scatter plot") points(trees[,3],col="red") legend("bottomright",c("Height","Volume"),col=c("blue","red"),lty=c(1,1))
```

#### Answer:

Answer4b.png