Computing Basics for Bioinformatics in R

Yaoyu Wang





Overview

- What is R
- Basic IO and data type
- Data type manipulation
- Set operation





- Programs for statistical analysis
 - SAS, SPSS, STATA, ...
- Programming Language
 - Assembly, FORTRAN, COBOL->B, C, S -> C++, JAVA
 - S = statistical programming language
 developed by Bell Labs in the late 70s
- Evoluation of S
 - S-Plus
 - R: Free & open source implementation of S
 - We can use R for Free!!.





- slow / fast
- Interpreted language : R, BASIC
- Compiled language : C, FORTRAN
- easy
- Great help manual
- Many web resource..(https://stat.ethz.ch/mailman/listinfo/r-help)
- Extensible
- Cross-platform: can run anywhere
- lots of libraries
- Can execture C, Fortran through Dynamic Link Library
- CRAN & Bioconductor





Basic of R

- R
 - Basic programming
 - Using packages
- stats
- Good reference
- http://www.r-project.org/



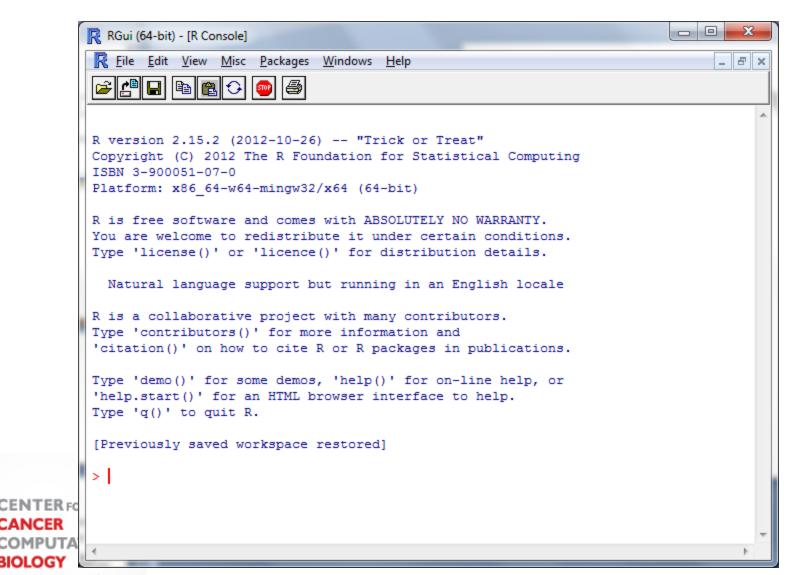


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Starting R





- Scalar
- Vector
- Matrix
- List
- Dataframe
- Factor





Data type: Scalar

- A single variable of numeric, character, and logical type
 - e.g. a=10 a variable a with value 10

Data type: Vector

- Statistical data = a set of (random) variables
- It can uniformly contain numeric, character, and logical values

e.g.
$$a=c(1,2,3,4)$$
 or $a=c("a", "b", "c","d")$





Data type: Matrix

 A set of vectors with the same length and same scalar data type

```
Example: A 2x2 matrix
```

```
> mdat =matrix(c(52,2,77, 11), nrow = 2, ncol=2, byrow=TRUE)
```

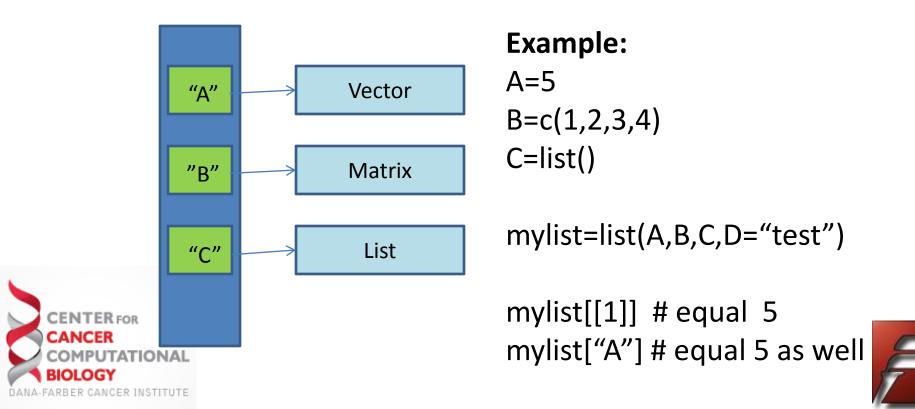
> mdat		[,1]	[,2]
	[1,]	52	2
	[2.]	77	11



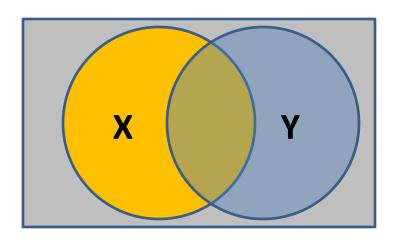


Data type: List

An object that contains a LIST of other objects



Set operation of vectors



- > xy=union(x,y)
- > xy_inter = intersect(x,y)
- > x_diff_y= setdiff(x,y)
- > x_in_y = is.element(x,y)
- $> x_{in}y = x[x \%in\% y]$





R Packages and Bioconductor

- There are many contributed packages that can be used to extend R basic function
- These libraries are created and maintained by the authors.
- BioConductor is an open source and open development software project for the analysis and comprehension of genomic data.
- http://www.bioconductor.org
- Download > Software > Installation Instructions

source("http://bioconductor.org/biocLite.R")
biocLite()



Basic Visualization for R

- Boxplot
- Scatterplots
- Histogram
- Heatmap

source("http://bioconductor.org/biocLite.R")
biocLite("gplots")

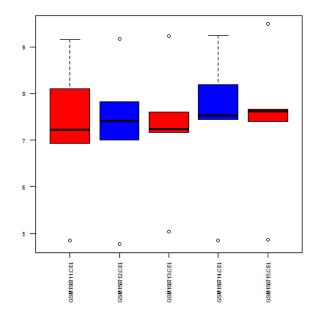
(want to see what's in gplots? Do > ls("package:gplots")





Generate Boxplot

```
boxplot(join.dat)
boxplot(join.dat, las=2)
boxplot(join.dat, las=2, cex.axis=0.8)
boxplot(join.dat, las=2, cex.axis=0.7, col=c("red, "blue""))
```

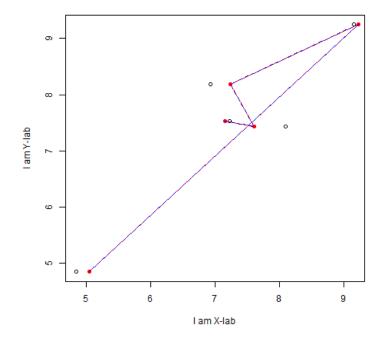






Scatter plot with connecting lines

```
plot(join.dat[,1], join.dat[,4], xlab="I am X-lab", ylab="I am Y-lab")
points(join.dat[,3], join.dat[,4])
points(join.dat[,3], join.dat[,4], col="red", pch=16)
lines(join.dat[,3], join.dat[,4], col="red", pch=16)
lines(join.dat[,3], join.dat[,4], col="blue", pch=16, lty="dashed")
```

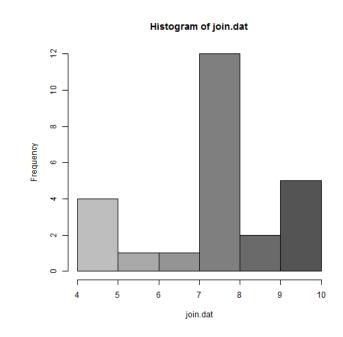






A histogram of all the probes intensity

join.dat=as.matrix(join.dat)
hist(join.dat)
hist(join.dat, col=colorpanel(10,"grey", "black"))

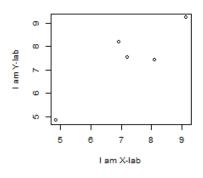


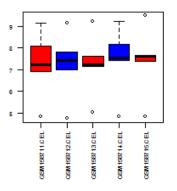




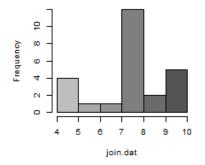
Multiple Plots Together

```
par(mfrow=c(2,2))
plot(join.dat[,1], join.dat[,4], xlab="I am X-lab", ylab="I am Y-lab")
boxplot(join.dat, las=2, cex.axis=0.7, col=c("red", "blue"))
hist(join.dat, col=colorpanel(10,"grey", "black"))
```





Histogram of join.dat





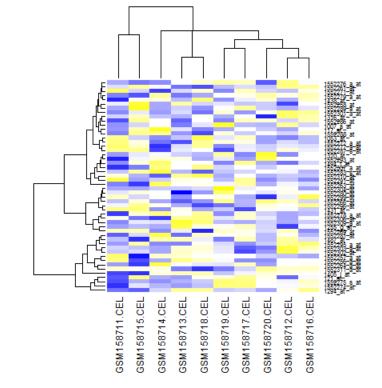


Heatmap

```
heatmap(join.dat)
heatmap(join.dat, col=colorpanel(100, "blue", "white", "yellow"))
heatmap(join.dat, col=colorpanel(100, "blue", "white", "yellow"),
margin=c(10,10))
```

heatmap(as.matrix(in.data[1:50,]), col=colorpanel(100, "blue", "white", "yellow"),

margin=c(10,10)



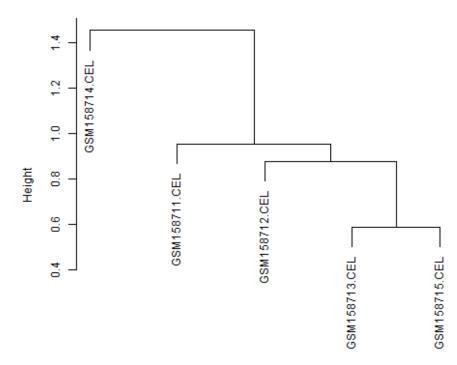




Hierachical Cluster Tree

dist(join.dat)
t(join.dat)
tree=dist(t(join.dat))
hclust(tree)

Cluster Dendrogram





tree hclust (*, "complete")



Exercise

Let's generate boxplots and figures for the expression patterns of all the **cell cycle** genes located on **chr17**

The probe/Gene name file is at:

/apps/ComputingBasics/cellcycle.chr17.txt





Just to get you start

```
# set target file and read in gexp.file="C:\\Users\\Yaoyu\\Desktop\\ComputingBasics\\example.rma.cll.txt" gexp=read.table(gexp.file, header=TRUE, sep="\t")
```

set probe file and read in probe.file="C:\\Users\\Yaoyu\\Desktop\\ComputingBasics\\cellcycle.chr17.txt" probe=read.table(probe.file, header=TRUE, sep="\t")





```
# Extract Target Probes from the gene expression matrix
cgexp=gexp[rownames(gexp) %in% probe[,1],]
png(file="boxplot.png")
boxplot(cgexp, las=2, cex.axis=0.7, col=c("red", "blue"))
dev.off()
cgexp=as.matrix(cgexp) # we can change the data type
png(file="hist.png")
hist.info=hist(cgexp, col=colorpanel(10,"grey", "black"))
dev.off()
#Heatmap
png("heatmap.png")
heatmap(cgexp, col=colorpanel(100, "blue", "white", "yellow"), margin=c(10,10))
dev.off()
```

