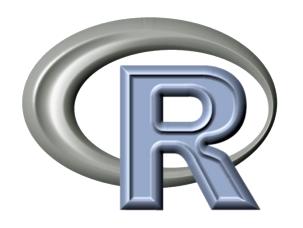
### An Introduction to R

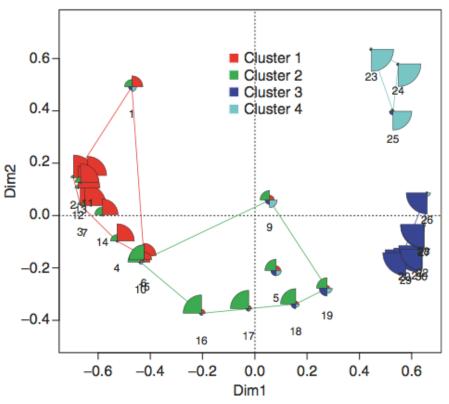
Chris Miller, PhD Washington University St Louis

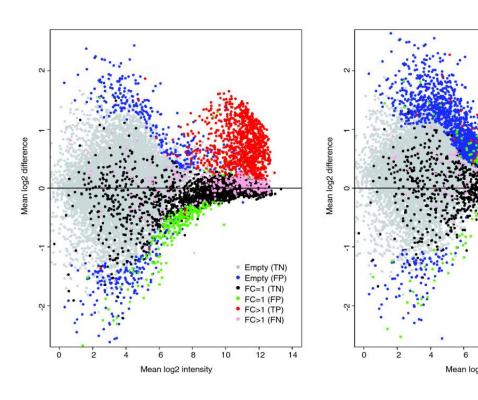


## What is R?

A free software environment for statistical computing and graphics

#### Ordination of fuzzy clusters (PCoA)





Empty (TN)
Empty (FP)
FC=1 (TN)
FC=1 (FP)
FC>1 (TP)
FC>1 (FN)

# Why is R useful?

- Data management and manipulation
- Well established system of packages and documentation
- Support for rich statistical simulation and modeling
- Active development and dedicated community
- Cutting-edge graphical data visualization
- Free!

## Things R is less good at

- BIG data
- There is a learning curve from many other languanges

- A common paradigm is to use other tools to massage your data into a bite size chunk, then import that into R for exploration/vizualization
  - e.g. Generate coverage in 10,000 bp bins from a bam file using mosdepth, run stats and make pretty plots of them with R

#### Where to learn more about R

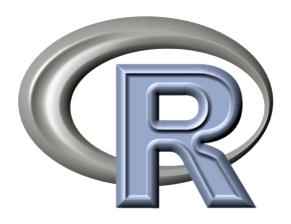
- The R Project Homepage: <a href="http://www.r-project.org">http://www.r-project.org</a>
- Quick R Homepage: <a href="http://www.statmethods.net">http://www.statmethods.net</a>
- Bioconductor: <a href="http://www.bioconductor.org">http://www.bioconductor.org</a>
- An Introduction to R (long!): <a href="http://cran.r-project.org/doc/manuals/R-intro.html">http://cran.r-project.org/doc/manuals/R-intro.html</a>
- Google there are tons of tutorials, guides, demos, packages and more

## R for Biologists

- Bioconductor (<a href="http://bioconductor.org">http://bioconductor.org</a>)
  - 2,140 packages (21-August, 2022):
    - Variant detection: coding changes, PolyPhen database
    - Annotation: pathway analysis, access GO, KEGG, NCBI and many others
    - High-throughput assays: flow cytometry, mass spec
    - Transcription factor binding detection
- Ecology (see: <a href="http://cran.r-project.org/web/views/Environmetrics.html">http://cran.r-project.org/web/views/Environmetrics.html</a>)
  - Ordination
  - Cluster Analysis
  - Ecological Theory
  - Population Dynamics
  - Spatial Data Analysis
- Phylogenetics and Evolution (see: <a href="http://cran.r-project.org/web/views/Phylogenetics.html">http://cran.r-project.org/web/views/Phylogenetics.html</a>)
  - Ancestral State Reconstruction
  - Phylogenetic Inference
  - Trait Evolution

# Obtaining R

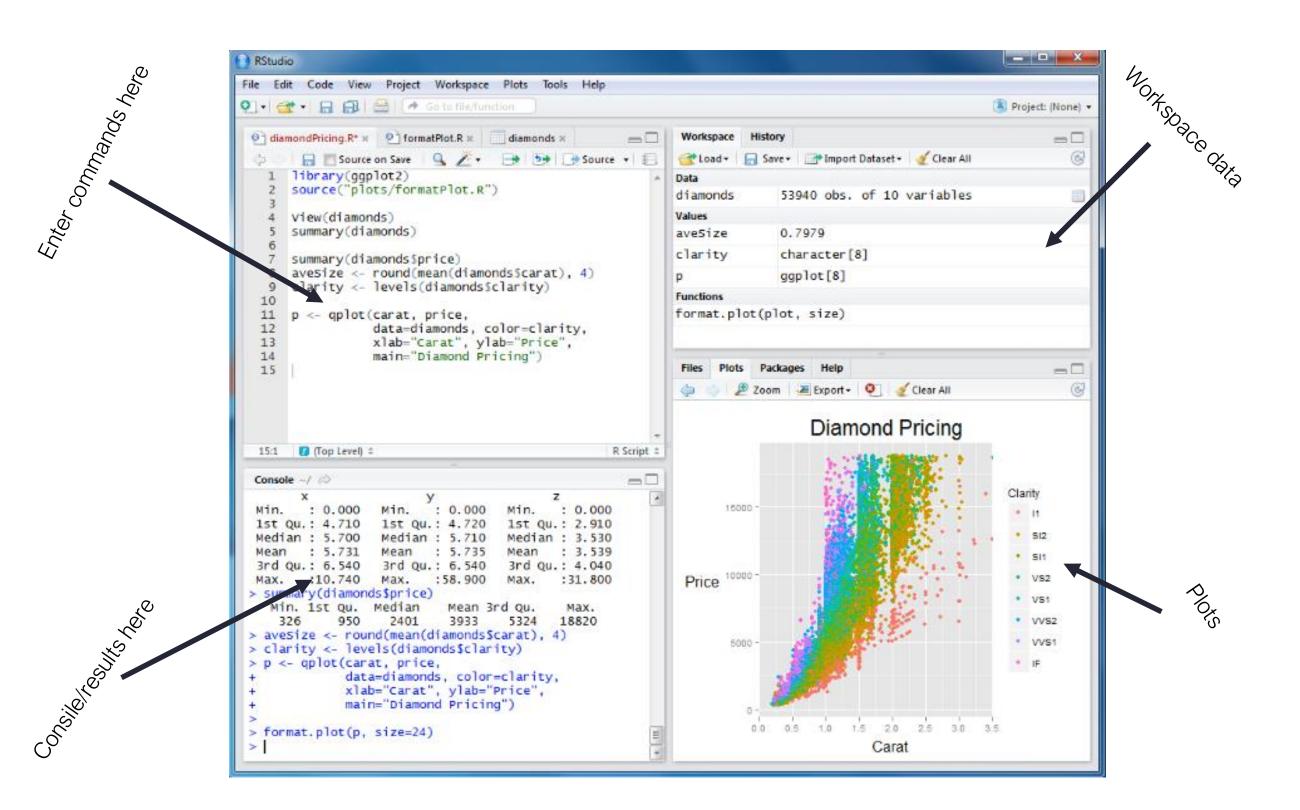
• Windows, Mac or Linux OS: <a href="https://www.r-project.org">https://www.r-project.org</a>



## Running R

- Install a R Integrated Development Environment (IDE)
  - RStudio: <a href="http://www.rstudio.com">http://www.rstudio.com</a>
  - Makes working with R much easier, particularly for a new R user
  - Run on Windows, Mac or Linux OS
- Or from the command line, type R

#### R Studio



## Basic R functionality

#### Calculator $(4+5^2)/3.14$ • +, -, /, \*, ^, log(), exp(), sqrt(), [1] 9.235669 abs(), cos(), sin(), tan(), ... Set Variables / y=13.4y=c(1,2,3,4,5)>y >y Vectors [1] 13.4 [1] 1 2 3 4 5 Sequences y=rep(2,10)[1] 2 2 2 2 2 2 2 2 2 2 2 [1] 2345678 y=2:8**Statistics** t.test(7:34, 5:29) t = 1.6348, df = 50.999, p-value = 0.1082

alternative hypothesis: true difference in means is not equal to 0

95 percent confidence interval:

-0.797982 7.797982 sample estimates: mean of x mean of y 20.5 17.0

# Manipulation I

n[c(1,3,5)]

n[n<50]

n[n>8 & n!=50]

[1] 50

[1] 3 12 50 103

[1] 3 7 12

[1] 3 12 103

[1] 3 7 12

[1] 12 103

## Manipulation II

n+1

[1] 4 8 13 51 104

sum(n)

[1] 175

mean(n)

[1] 35

var(n)

[1] 1796.5

min(n)

[1] 3

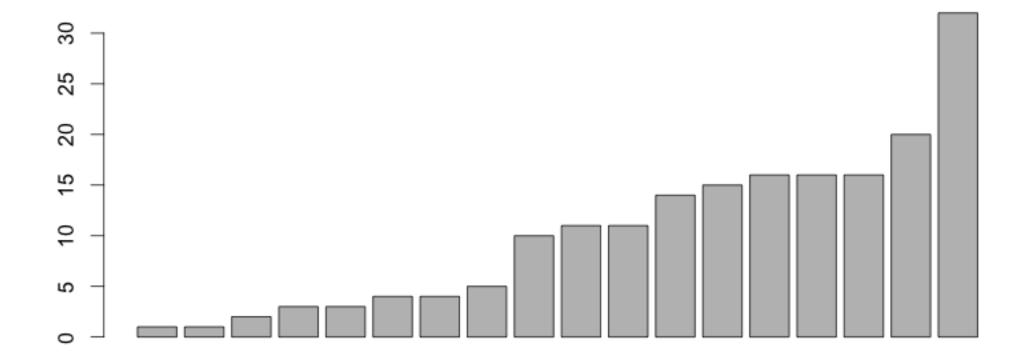
max(n)

[1] 103

## Basic Visualization I

y=c(1,1,2,3,3,4,4,5,10,11,11,14,15,16,16,16,20,32)

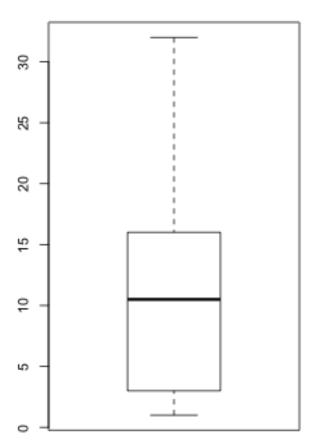
barplot(y)



## Basic Visualization II

y=c(1,1,2,3,3,4,4,5,10,11,11,14,15,16,16,16,20,32)

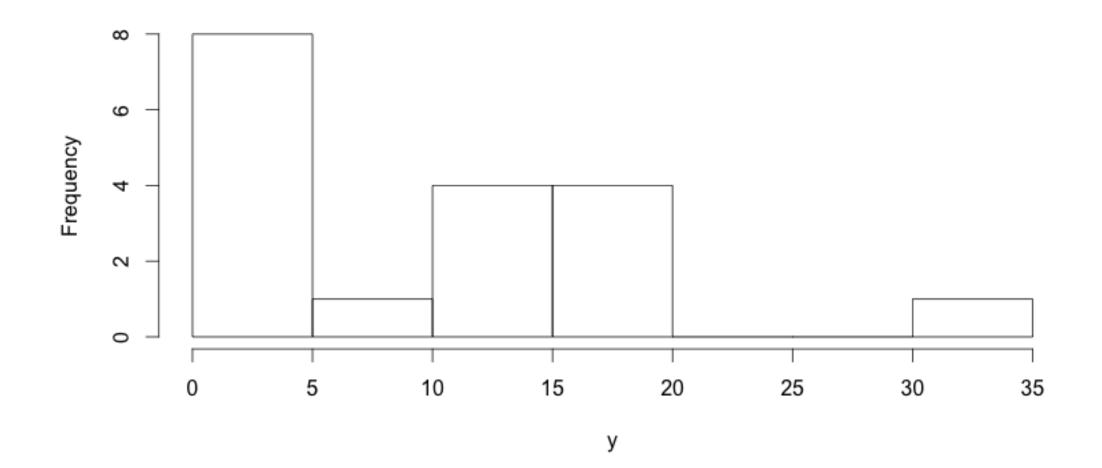
boxplot(y)



### **Basic Visualization III**

y=c(1,1,2,3,3,4,4,5,10,11,11,14,15,16,16,16,20,32)

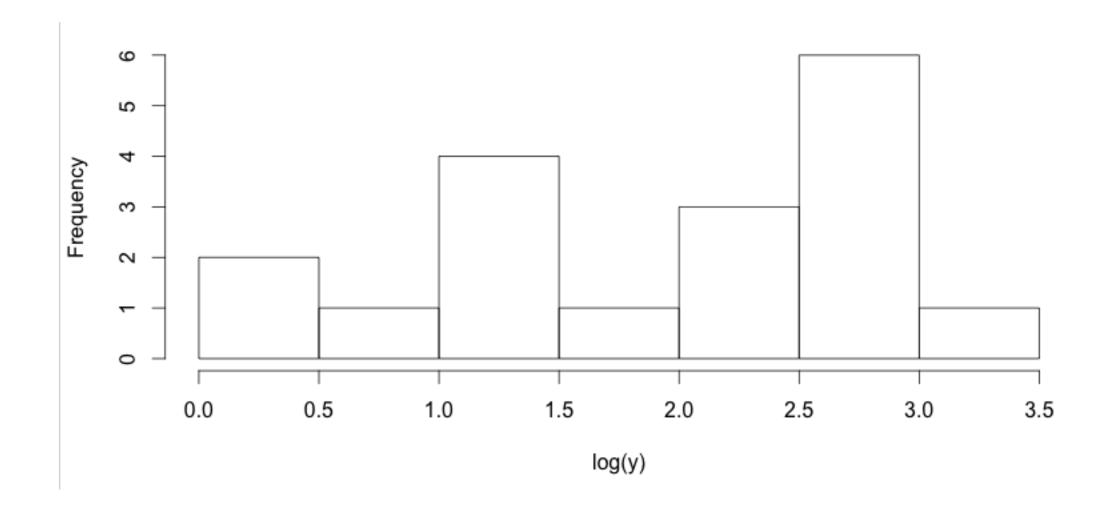
# hist(y)



### Basic Visualization III.i

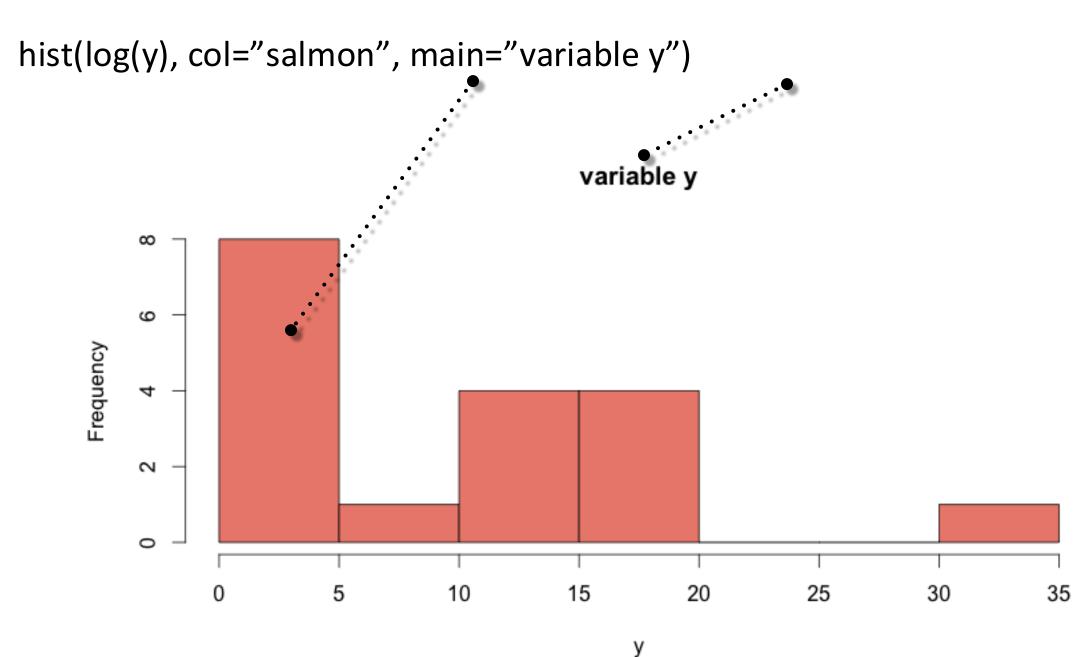
y=c(1,1,2,3,3,4,4,5,10,11,11,14,15,16,16,16,20,32)

hist(log(y))



## Basic Visualization III.ii

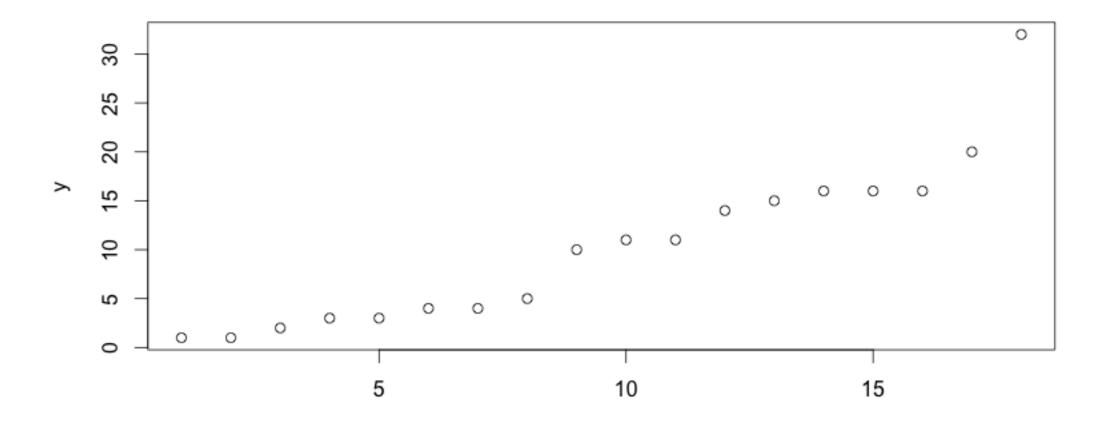
y=c(1,1,2,3,3,4,4,5,10,11,11,14,15,16,16,16,20,32)



### **Basic Visualization IV**

y=c(1,1,2,3,3,4,4,5,10,11,11,14,15,16,16,16,20,32)

plot(y)

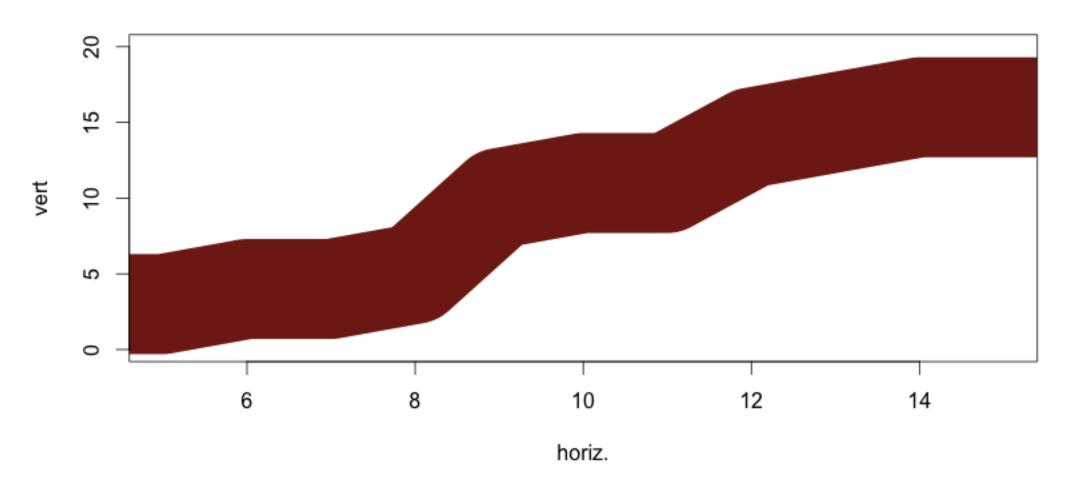


### Basic Visualization IV.ii

y=(1,1,2,3,3,4,4,5,10,11,11,14,15,16,16,16,20,32)

plot(y, type="l", col="dark red", lwd=100, main="y variable", ylim=c(0,20), xlim=c(5,15), ylab="vert", xlab="horiz.")

#### y variable



## Help

- Do you need to remember all of the variables?
- ? is your friend
- ?plot

plot {graphics}

#### Generic X-Y Plotting

#### Description

Generic function for plotting of R objects. For more details about the graphical parameter arguments, see par.

For simple scatter plots, <u>plot.default</u> will be used. However, there are plot methods for many R objects, including <u>functions</u>, <u>data.frames</u>, <u>density</u> objects, etc. Use methods (plot) and the documentation for these.

#### Usage

```
plot(x, y, ...)
```

R Documentation

type

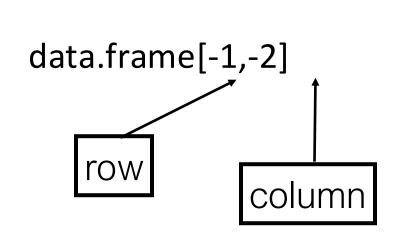
what type of plot should be drawn. Possible types are

- . "p" for points,
- "1" for lines,
- "b" for both,
- . "c" for the lines part alone of "b",
- "o" for both 'overplotted',
- "h" for 'histogram' like (or 'high-density') vertical lines,
- "s" for stair steps,
- . "S" for other steps, see 'Details' below,
- "n" for no plotting.

#### read.csv

- A data.frame is essentially a table
- columns can be mixed types
  - numeric, text strings
- rows must be same type

	clostridia	proteobacteria	bacteroides
01_healthy	22	54	245
02_healthy	26	65	265
03_healthy	34	66	262
01_sick	32	32	116
02_sick	12	24	101
03_sick	9	18	87



	clostridia	bacteroides
02_healthy	26	265
03_healthy	34	262
01_sick	32	116
02_sick	12	101
03_sick	9	87

## Data Frame Manipulations

#### https://is.gd/bacteriacsv

	clostridia	proteobacteria	bacteroides
01_healthy	22	54	245
02_healthy	26	65	265
03_healthy	34	66	262
01_sick	32	32	116
02_sick	12	24	101
03_sick	9	18	87

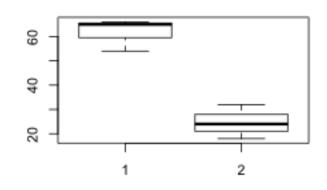
bac\$proteobacteria

[1] 54 65 66 32 24 18

t.test(bac\$proteobacteria[1:3], bac\$proteobacteria[4:6])

p-value = 0.002725

boxplot(bac\$proteobacteria[1:3],
 bac\$proteobacteria[4:6])



## Getting Help in R

#### ?write.table

#### Description

write.table prints its required argument x (after converting it to a data frame if it is not one nor a matrix) to a file or connection.

#### Usage

#### **Arguments**

the object to be written, preferably a matrix or data frame. If not, it is attempted to coerce x to a data frame.

file either a character string naming a file or a connection open for writing. "" indicates output to the console.

logical. Only relevant if file is a character string. If TRUE, the output is appended to the file. If FALSE, any existing file of the name is destroyed.

Exercise: try to write out our `bac` data frame to a tab-separated file with no quoting around the data

## Getting Help in R

- R syntax can be weird
- At first you don't even know what you don't know
- Google is your friend! (lots of StackOverflow answers, R blogs)
- Don't use LLM answers blindly, but do use them and learn from them
  - set up positive and negative controls

## Exercise

• https://github.com/genome/bfx-workshop/tree/master/lectures/week\_06