

 Covered the When, Why, What and How of writing your own R functions.

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 - When: When you find yourself doing the same thing 3 or more times with repetitive code consider writing a function.

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 - When: When you find yourself doing the same thing 3 or more times with repetitive code consider writing a function.

<u>₩hy</u>:

- 1. Makes the purpose of the code more clear
- 2. Reduces mistakes from copy/paste
- 3. Makes updating your code easer
- 4. Reduces code duplication and facilitates re-use.

. . .

- Covered the When, Why, What and How of writing your own R functions.
 - What: A function is defined with:
 - 1. A user selected name,
 - 2. A comma separated set of input arguments, and
 - 3. Regular R code for the function body including an optional output return value e.g.

```
fname <- function(arg1, arg2) { paste(arg1,arg2) }

Name Input arguments Function body
```

Every function in R has the same parts

```
rescale <- function(x, na.rm=TRUE, plot=FALSE, ...) {
   rng <-range(x, na.rm=na.rm)</pre>
   answer <- (x - rng[1]) / (rng[2] - rng[1])
   if(plot) {
     plot(answer, ...)
  return(answer)
```

```
rescale <- function(x, na.rm=TRUE, plot=FALSE, ...) {
   rng <-range(x, na.rm=na.rm)</pre>
   answer <- (x - rng[1]) / (rng[2] - rng[1])
   if(plot) {
     plot(answer, ...)
   return(answer)
```

Arguments: We can supply values for these variables that appear in the function body.

```
rescale <- function(x, na.rm=TRUE, plot=FALSE,
   rng <-range(x, na.rm=na.rm)</pre>
   answer <- (x - rng[1]) / (rng[2] - rng[1])
   if(plot) {
     plot(answer, ...)
   return(answer)
```

Arguments: We can supply values for these variables that appear in the function body.

```
rescale <- function(x, na.rm=TRUE, plot=FALSE, ...) {
    rng <-range(x, na.rm=na.rm)
    answer <- (x - rng[1]) / (rng[2] - rng[1])
    if(plot) {
       plot(answer, ...)
    }
    recturn(answer)
}</pre>
```

Body: R will run this code whenever we call the function by typing its name followed by brackets.

Arguments: We can supply values for these variables that appear in the function body.

```
rescale <- function(x, na.rm=TRUE, plot=FALSE, ...) {
    rng <-range(x, na.rm=na.rm)
    answer <- (x - rng[1]) / (rng[2] - rng[1])
    if(plot) {
       plot(answer, ...)
    }
    resturn(answer)
}</pre>
```

Return value: The function will return this value

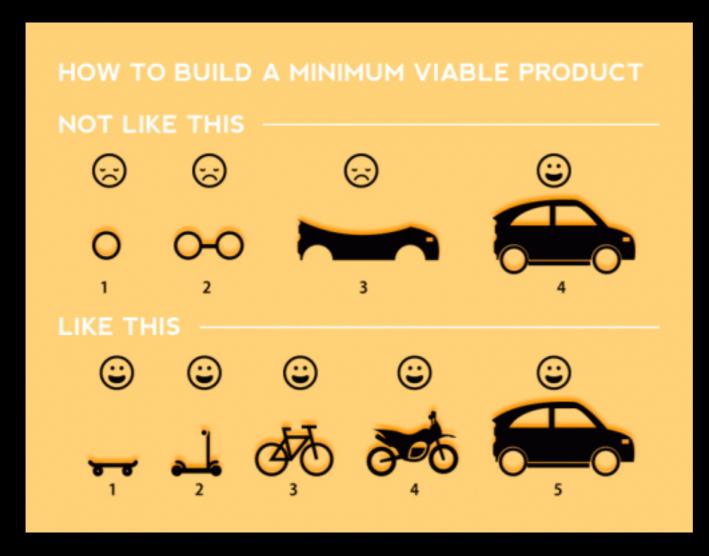
Body: R will run this code whenever we call the function by typing its name followed by brackets.

- How: Follow a step-by-step procedure to go from working code snippet to refined and tested function.
 - 1. Start with a simple problem and write a working snippet of code.
 - 2. Rewrite for clarity and to reduce duplication
 - 3. Then, and only then, turn into an initial function
 - 4. Test on small well defined input
 - 5. Report on potential problem by failing early and loudly!

- How: Follow a step-by-step procedure to go from working code snippet to refined and tested function.
 - 1. Start with a simple problem and write a working snippet of code.
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Recap...

1. Start with a simple problem and write a working snippet of code.



Build that skateboard before you build the car.

A limited but functional thing is very useful and keeps the spirits high.

[Image credit: Spotify development team]

Back by popular demand More examples of how to write your own functions!

Revisit our first example function from last day... source("http://tinyurl.com/rescale-R")

```
rescale <- function(x, na.rm=TRUE, plot=FALSE, ...) {
   rng <-range(x, na.rm=na.rm)</pre>
   answer <- (x - rng[1]) / (rng[2] - rng[1])
   if(plot) {
     plot(answer, ...)
   return(answer)
```

```
# Test fail
rescale( c(1,10,"string") )
```

The functions warning() and stop()

- The functions warning() and stop() are used inside functions to handle and report on unexpected situations
- They both print a user defined message (which you supply as a character input argument to the warning() and stop() functions).
- However, warning() will keep on going with running the function body code whereas stop() will terminate the action of the function.
- A common idiom is to use stop("some message") to report on unexpected input type or other problem early in a function,
 - i.e. "Fail early and loudly!"

```
rescale2 <- function(x, na.rm=TRUE, plot=FALSE, ...) {
   if( !is.numeric(x) ) {
      stop("Input x should be numeric", call.=FALSE)
   rng <-range(x, na.rm=na.rm)</pre>
   answer <- (x - rng[1]) / (rng[2] - rng[1])
   if(plot) {
      plot(answer, ...)
   return (answer)
```

```
rescale2 <- function(x, na.rm=TRUE, plot=FALSE, ...) {
   if( !is.numeric(x) ) {
      stop("Input x should be numeric", call.=FALSE)
   rng <-range(x, na.rm=na.rm)</pre>
   answer \leftarrow (x - rng[1]) / (rng[2] - rng[1])
   if(plot) {
      plot(answer, ...)
   return (answer)
```

Suggested steps for writing your functions

- 1. Start with a simple problem and get a working snippet of code
- 2. Rewrite to use temporary variables (e.g. x, y, df, m etc.)
- 3. Rewrite for clarity and to reduce calculation duplication
- 4. Turn into an initial function with clear useful names
- 5. Test on small well defined input and (subsets of) real input
- 6. Report on potential problem by failing early and loudly!
- 7. Refine and polish

Side-Note: What makes a good function?

- Correct
- Understandable (remember that functions are for humans and computers)
- Correct + Understandable = Obviously correct
- Use sensible names throughout. What does this code do?

```
baz <- foo(df, v=0)
df2 < replace_missing(df, value=0)</pre>
```

 Good names make code understandable with minimal context. You should strive for self-explanatory names

More examples

 We want to write a function, called both_na(), that counts how many positions in two input vectors, x and y, both have a missing value

```
# Should we start like this?
both_na <- function(x, y) {
    # something goes here?
}</pre>
```

No! Always start with a simple definition of the problem

 We should start by solving a simple example problem first where we know the answer.

```
# Lets define an example x and y x <- c(1, 2, NA 3, NA) y <- c(NA, 3, NA 3, 4)
```

 Here the answer should be 1 as only the third position has NA in both inputs x and y.

Tip: Search for existing functionality to get us started...

Get a working snippet of code first that is close to what we want

```
# Lets define an example x and y x <- c(1, 2, NA, 3, NA) y <- c(NA, 3, NA, 3, 4)
```

```
# Use the is.na() function
is.na(x)
[1] FALSE FALSE TRUE FALSE TRUE
```

Get a working snippet of code first that is close to what we want

```
# Lets define an example x and y x <- c(1, 2, NA, 3, NA) y <- c(NA, 3, NA, 3, 4)
```

```
# Use the is.na() function
is.na(x)
[1] FALSE FALSE TRUE FALSE TRUE
```

Q. How many TRUE values are there?

Get a working snippet of code first that is close to what we want

```
# Lets define an example x and y x <- c(1, 2, NA, 3, NA) y <- c(NA, 3, NA, 3, 4)
```

```
# Use the is.na() function
is.na(x)
[1] FALSE FALSE TRUE FALSE TRUE

sum( is.na(x) )
[1] 2

# Putting together!
sum( is.na(x) & is.na(y) )
[1] 1
```

Then rewrite your snippet as a *first* function

```
# Lets define an example x and y
x <- c(1, 2, NA, 3, NA)
y <- c(NA, 3, NA, 3, 4)
```

```
# Our working snippet
sum( is.na(x) & is.na(y) )
```

```
# No further simplification necessary
both_na <- function(x, y) {
   sum( is.na(x) & is.na(y) )
}</pre>
```

Test on various inputs (a.k.a. eejit proofing)

 We have a function that works in at least one situation, but we should probably check it works in others.

```
x <- c(NA, NA, NA)
y1 <- c(1, NA, NA)
y2 <- c(1, NA, NA, NA)

both_na(x, y1)
[1] 2

# What will this return?
both_na(x, y2)</pre>
```

Report on potential problem by failing early and loudly!

• The generic warning with recycling behavior of the last example may not be what you want as it could be easily missed especially in scripts.

```
both_na2 <- function(x, y) {
  if(length(x) != length(y)) {
    stop("Input x and y should be the same length")
  }
  sum( is.na(x) & is.na(y) )
}</pre>
```

Refine and polish: Make our function more useful by returning more information

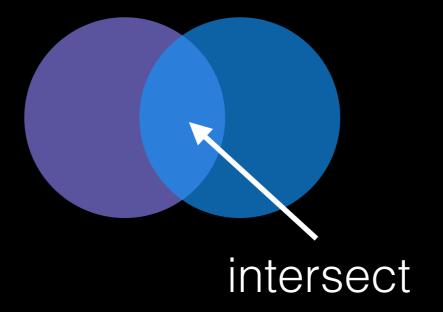
```
both na3 <- function(x, y) {
  if(length(x) != length(y)) {
    stop("Input x and y should be vectors of the same length")
  na.in.both <- (is.na(x) & is.na(y))
  na.number <- sum(na.in.both)</pre>
  na.which <- which(na.in.both)</pre>
 message("Found ", na.number, " NA's at position(s):",
          paste(na.which, collapse=", ") )
  return( list(number=na.number, which=na.which) )
```

Re-cap: Steps for function writing

- 1. Start with a simple problem and get a working snippet of code
- 2. Rewrite to use temporary variables
- 3. Rewrite for clarity and to reduce calculation duplication
- 4. Turn into an initial function
- 5. Test on small well defined input and (subsets of) real input
- 6. Report on potential problem by failing early and loudly!
- 7. Refine and polish,
- 8. Document and comment within the code on your reasoning.

One last example

Find common genes in two data sets and return their associated data (from each data set)



source("http://tinyurl.com/rescale-R")

source("http://tinyurl.com/rescale-R")

Alono.

df1

*	IDs ‡	ехр	‡
1	gene1		2
2	gene2		1
3	gene3		1

df2

*	IDs ‡	exp ‡
1	gene2	-2
2	gene4	NA
3	gene3	1
4	gene5	2

source("http://tinyurl.com/rescale-R")

```
# Start with a simple version of the problem
df1 <- data.frame(IDs=c("gene1", "gene2", "gene3"),
                    \exp=c(2,1,1),
                    stringsAsFactors=FALSE)
df2 <- data.frame(IDs=c("gene2", "gene4", "gene3", "gene5"),
                    \exp=c(-2, NA, 1, 2),
                    stringsAsFactors=FALSE)
# Simplify further to single vectors
x <- df1$IDs
y <- df2$IDs
# Now what do we do?
                                                        IDs
                                                             exp
                                      IDs
                                           exp
                                                      1 gene2
                                     1 gene1
                                                      2 gene4
                                                                NA
                                     2 gene2
                                                      3 gene3
                                     gene3
                                                       gene5
```

10h alo

```
# Start with a simple version of the problem
df1 <- data.frame(IDs=c("gene1", "gene2", "gene3"),
                   \exp=c(2,1,1),
                   stringsAsFactors=FALSE)
df2 <- data.frame(IDs=c("gene2", "gene4", "gene3", "gene5"),
                   \exp=c(-2, NA, 1, 2),
                   stringsAsFactors=FALSE)
# Simplify further to single vectors
                                                      IDs
                                                             IDs
x < - df1$IDs
                                                             gene2
                                                      gene1
y <- df2$IDs
                                                             gene4
                                                      gene2
                                                             gene3
                                                      gene3
# Now what do we do?
                                                             gene5
```

Tip: Search for existing functionality to get us started...

```
10h 3/0
```

```
# Simplify further to single vectors
x < - df1$IDs
y <- df2$IDs
# Search for existing functionality to get us started ...
??intersect
intersect(x, y)
#> [1] "gene2" "gene3"
                                                         IDs
                                                                IDs
                                                                gene2
                                                         gene1
                                                                gene4
                                                         gene2
                                                                gene3
                                                         gene3
                                                                gene5
```

Close but not useful for returning indices yet.

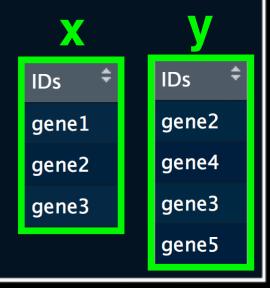
```
ON Blong
```

```
# Simplify further to single vectors
x < - df1$IDs
y <- df2$IDs
# Search for existing functionality to get us started ...
??intersect
intersect(x, y)
#> [1] "gene2" "gene3"
                                                        IDs
                                                               IDs
# Back to search results...
                                                               gene2
                                                        gene1
?intersect
                                                               gene4
                                                        gene2
                                                               gene3
                                                        gene3
                                                               gene5
```

Close but not useful for returning indices yet.

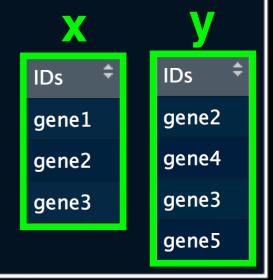
```
TONON Along!
```

```
# This looks like a more useful starting point - indices!
x %in% y
#> [1] FALSE TRUE TRUE
```



```
# This looks like a more useful starting point - indices!
x %in% y
#> [1] FALSE TRUE TRUE

x[x %in% y]
#> [1] "gene2" "gene3"
```



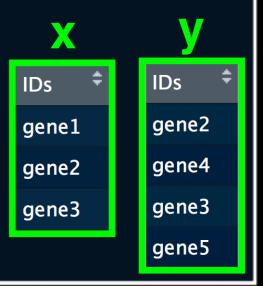
```
# This looks like a more useful starting point - indices!
x %in% y
#> [1] FALSE TRUE TRUE
x[x % in% y]
#> [1] "gene2" "gene3"
y[ y %in% x ]
#> [1] "gene2" "gene3"
# We can now cbind() these these results ...
cbind(x[x%in%y],y[y%in%x])
                                                         IDs 🕏
                                                   IDs
#> [,1] [,2]
                                                         gene2
                                                   gene1
#> [1,] "gene2" "gene2"
                                                         gene4
                                                   gene2
#> [2,] "gene3" "gene3"
                                                         gene3
                                                   gene3
                                                         gene5
```

```
# This looks like a more useful starting point - indices!
x %in% y
#> [1] FALSE TRUE TRUE

x[x %in% y]
#> [1] "gene2" "gene3"

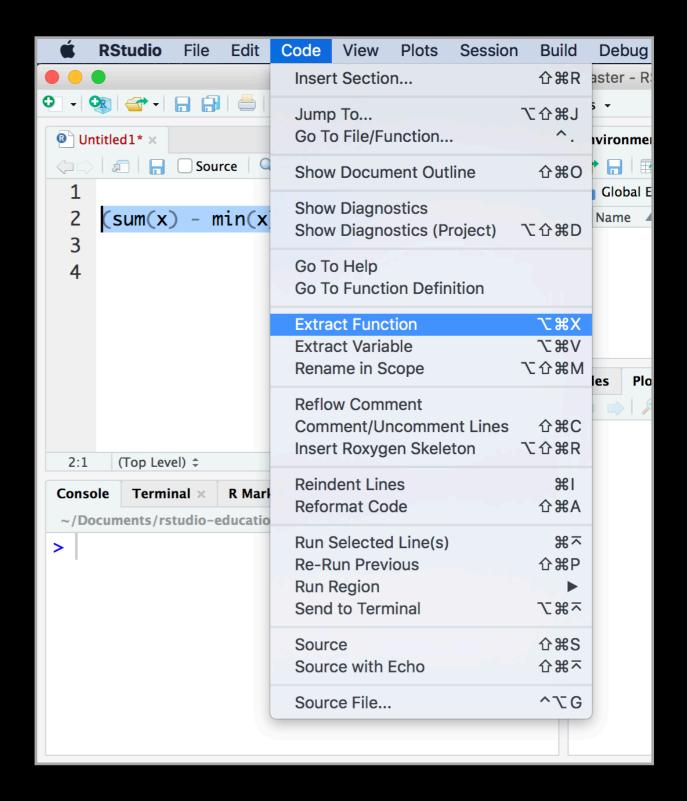
y[ y %in% x ]
#> [1] "gene2" "gene3"
```

Working Snippet!



RStudio Tip: Extract function

- You can use the Extract
 Function feature of the
 RStudio IDE to turn a piece of code into a function.
 - Code > Extract Function
- It is good practice to double check it has set your function input arguments correctly and of course further test, refine and polish...



gene3

gene5

gene3

```
# Our previous working snippet ...
cbind(x[x%in%y],y[y%in%x])
# Make this snippet into a first function
gene intersect <- function(x, y) {</pre>
  cbind(x[x%in%y],y[y%in%x])
# Looks good so far but we need to work with data frames
gene intersect(x, y)
#> [,1] [,2]
#> [1,] "gene2" "gene2"
                                                       IDs
                                                 IDs
#> [2,] "gene3" "gene3"
                                                       gene2
                                                 gene1
                                                       gene4
                                                 gene2
```

```
# Our previous working snippet ...
cbind(x[x%in%y],y[y%in%x])
# Make this snippet into a first function
gene intersect <- function(x, y) {</pre>
 cbind(x[x%in%y],y[y%in%x])
# Looks good so far but we need to work with data frames
gene intersect(x, y)
#> [,1] [,2]
#> [1,] "gene2" "gene2"
#> [2,] "gene3" "gene3"
```

*	IDs ‡	exp ‡
1	gene1	2
2	gene2	1
3	gene3	1

*	IDs ‡	exp ‡
1	gene2	-2
2	gene4	NA
3	gene3	1
4	gene5	2

1042/2

source("http://tinyurl.com/rescale-R")

df⁻

*	IDs ‡	exp ‡
1	gene1	2
2	gene2	1
3	gene3	1

df2

*	IDs ‡	exp ‡
1	gene2	-2
2	gene4	NA
3	gene3	1
4	gene5	2

N.B. Our input \$IDs column name may change:

So lets add flexibility by allowing the user to specify the gene containing column name

```
# Experiment first to make sure things are as we expect
gene.colname="IDs"
df1[,gene.colname]
#> [1] "gene1" "gene2" "gene3"
```

```
# Next step: Add df1[,gene.colname] etc to function.
gene intersect3 <- function(df1, df2, gene.colname="IDs") {
  cbind( df1[ df1[,gene.colname] %in%
             df2[,gene.colname], ],
        exp2=df2[ df2[,gene.colname] %in%
             df1[,gene.colname], "exp"] )
# Works but the function is not kind on the reader
gene intersect3(df1, df2)
#> IDs exp exp2
#> 2 gene2 1 -2
#> 3 gene3 1 1
```

```
# Improve by simplifying for for human consumption
gene intersect4 <- function(df1, df2, gene.colname="IDs")
  df1.name <- df1[,gene.colname]</pre>
  df2.name <- df2[,gene.colname]</pre>
  df1.inds <- df1.name %in% df2.name
  df2.inds <- df2.name %in% df1.name
  cbind( df1[ df1.inds, ],
         exp2=df2[ df2.inds, "exp"] )
```

```
# Improve by simplifying for for human consumption
gene intersect4 <- function(df1, df2, gene.colname="IDs")
  df1.name <- df1[,gene.colname]</pre>
  df2.name <- df2[,gene.colname]</pre>
  df1.inds <- df1.name %in% df2.name
  df2.inds <- df2.name %in% df1.name
  cbind( df1[ df1.inds, ],
         exp2=df2[ df2.inds, "exp"] )
# Getting closer!
gene intersect4(df1, df2)
#> IDs exp exp2
#> 2 gene2 1 -2
#> 3 gene3 1 1
```

```
# Test, break, fix, text again
df1 <- data.frame(IDs=c("gene1", "gene2", "gene3"),</pre>
                  \exp=c(2,1,1),
                  stringsAsFactors=FALSE)
df3 <- data.frame(IDs=c("gene2", "gene2", "gene5", "gene5"),
                  \exp=c(-2, NA, 1, 2),
                  stringsAsFactors=FALSE)
# Works but could do with more spit and polish!
gene intersect4(df1, df3)
#> IDs exp exp2
#> 1 gene2 1 -2
#> 2 gene2 1 NA
#> Warning in data.frame(..., check.names = FALSE): row
names were found from a short variable and have been
discarded
```

```
# Additional features we could add
# - Catch and stop when user inputs weird things
# - Use different colnames for matching in df1 and df2,
# - Match based on the content of multiple columns,
# - Optionally return rows not in df1 or not in df2 with NAs
# - Optionally sort results by matching column
# - etc...

merge(df1, df2, by="IDs")
#> IDs exp.x exp.y
#> 1 gene2 1 -2
```

#> 2 gene3 1 1

For more details refer to sections 2-5 in last days handout!

https://bioboot.github.io/bimm143_F19/lectures/#6

Remember Section 1B (question 6) is your last days homework (see also scoring rubric).

The **Sections 2** to **5** are there for your benefit.

Your turn...

 Write a function grade() to determine an overall grade from a vector of student homework assignment scores dropping the lowest single alignment score.

```
# student 1
c(100, 100, 100, 100, 100, 100, 90)
# student 2
c(100, NA, 90, 90, 90, 90, 97, 80)
```

Your turn...

• Write a function grade() to determine an overall grade from a vector of student homework assignment scores dropping the lowest single alignment score.

```
# student 1
c(100, 100, 100, 100, 100, 100, 100, 90)

# student 2
c(100, NA, 90, 90, 90, 90, 97, 80)

# now grade all students in an example class
url <- "https://tinyurl.com/gradeinput"</pre>
```

PHONION

CRAN & Bioconductor

Major repositories for **R packages** that extend R functionality

CRAN: Comprehensive R Archive Network

- CRAN is a network of mirrored servers around the world that administer and distribute R itself, R documentation and R packages (basically add on functionality!)
- There are currently ~15,133 packages on CRAN in the areas of finance, bioinformatics, machine learning, high performance computing, multivariate statistics, natural language processing, etc. etc.

https://cran.r-project.org/

Side-note: R packages come in all shapes and sizes











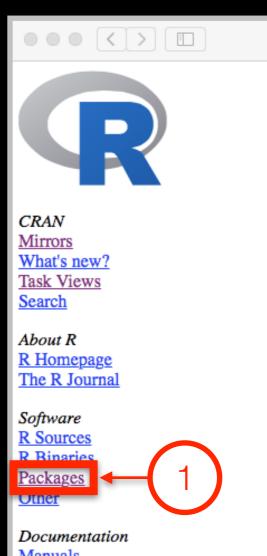
R packages can be of variable quality and often there are multiple packages with overlapping functionality.

Refer to relevant publications, package citations, update/maintenance history, documentation quality and your own tests!

The journal has sufficient experience with CRAN and Bioconductor resources to endorse their use by authors. We do not yet provide any endorsement for the suitability or usefulness of other solutions.

From: "Credit for Code". Nature Genetics (2014), 46:1

https://cran.r-project.org



Manuals
FAQs
Contributed

Contributed Packages

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Available Packages

Currently, the CRAN package repository features 8952 available packages.

a cran.r-project.org

Table of available packages, sorted by date of publication

Table of available packages, sorted by name

Installation of Packages

Please type help("INSTALL") or help("install.packages") in R for information on how to install packages from this repository. The manual R Installation and Administration (also contained in the R base sources) explains the process in detail.

<u>CRAN Task Views</u> allow you to browse packages by topic and provide tools to automatically install all packages for special areas of interest. Currently, 33 views are available.

Package Check Results

All packages are tested regularly on machines running Debian GNU/Linux, Fedora, OS X, Solaris and Windows.

The results are summarized in the <u>check summary</u> (some <u>timings</u> are also available). Additional details for Windows checking and building can be found in the <u>Windows check summary</u>.

Writing Your Own Packages

The manual <u>Writing R Extensions</u> (also contained in the R base sources) explains how to write new packages and how to contribute them to CRAN.

Repository Policies

The manual CRAN Repository Policy [PDF] describes the policies in place for the CRAN package repository.

Installing a package

RStudio > Tools > Install Packages

- install.packages("bio3d")
- > library("bio3d")

Bioconductor

R packages and utilities for working with high-throughput genomic data

http://bioconductor.org



More pragmatic:

Bioconductor is a software repository of R packages with some rules and guiding principles.

Version 3.9 had 1,741 software packages.

Bioconductor has emphasized Reproducible Research since its start, and has been an early adapter and driver of tools to do this.

"Bioconductor: open software development for computational biology and bioinformatics"

Gentleman et al

Genome Biology 2004, 5:R80

"Orchestrating high-throughput genomic analysis with Bioconductor"

Huber et al

Nature Methods 2015, 12:115-121

Installing a bioconductor package

First get the BiocManager package from CRAN

- > install.packages("BiocManager")
- > library(BiocManager)

Install bioconductor core packages with:

> install()

See: http://www.bioconductor.org/install/

Installing a bioconductor package

First get the BiocManager package from CRAN

- > install.packages("BiocManager")
- > library(BiocManager)

Install bioconductor core packages with:

> install()

Install specific packages, e.g. "AnnotationDbi", with:

> install("AnnotationDbi")

See: http://www.bioconductor.org/install/

Your Turn: Form a group of 3, pick a package to explore and install, Report back to the class.

ggplot2, bio3d, rmarkdown, dplyr, rgl, pheatmap, rentrez, reprex, igraph, blogdown, shiny, msa, flexdashborad, phyloseq, datapasta

Questions to answer:

- How does it extend R functionality? (i.e. What can you do with it that you could not do before?)
- How is it's documentation, vignettes, demos and web presence?
- Can you successfully follow a tutorial or vignette to get started quickly with the package?
- Can you find a GitHub or Bitbucket site for the the package with a regular heartbeat?

[Collaborative Google Doc Link] <- See Website

Your Turn: Form a group of 3, pick a package to explore and install, Report back to the class.

ggplot2, bio3d, rmarkdown, rgl, dplyr, rentrez, reprex blogdown, shiny, msa, flexdashborad, phyloseq,

Questions to answer:

- How does it extend R functionality? (i.e. What can you do with it that you could not do before?)
- How is it's documentation, vignettes, demos and web presence?
- Can you successfully follow a tutorial or vignette to get started quickly with the package?
- Can you find a GitHub or Bitbucket site for the the package with a regular heartbeat?

[Collaborative Google Doc Link] <- See Website

Key Idea:



Using existing base functions in R is like riding a bus - it is relatively straightforward you just need to know which bus to use and know where to get on and get off.



Being able to use **CRAN & bioconductor** packages and functions is like having access to UBER - they can take you more places but may only cover big cities.



Writing your **own functions** in R is like driving an SUV with kayak & bike on top - it takes more work, you need to know how to get there. Ultimately, however it will give you the flexibility to go completely new places!

Summary

- R is a powerful data programming language and environment for statistical computing, data analysis and graphics.
- Introduced R syntax and major R data structures
- Demonstrated using R for exploratory data analysis and graphics.
- Exposed you to the why, when, and how of writing your own R functions.
- Introduced CRAN and Bioconductor package repositories.

[Muddy Point Assessment Link]

Learning Resources

- DataCamp. Online tutorials using R in your browser.
 - < https://www.datacamp.com/ >
- R for Data Science. A new O'Reilly book that will teach you how to do data science with R, by Garrett Grolemund and Hadley Wickham.
 - < http://r4ds.had.co.nz/ >