



BGGN 213

More on R functions  
and packages

Lecture 7

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# Recap From Last Time:

- 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.

---

→ **Why**:

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

...

# Recap From Last Time:

- 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)  
  answer <- (x - rng[1]) / (rng[2] - rng[1])  
  
  if(plot) {  
    plot(answer, ...)  
  }  
  
  return(answer)  
}
```

**Name:** We can run the function by typing its name followed by brackets.

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rescale <- function(x, na.rm=TRUE, plot=FALSE, ...) {  
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    plot(answer, ...)  
  }  
  
  return(answer)  
}
```

**Name**: We can run 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)  
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  }  
  
  return(answer)  
}
```

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

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  answer <- (x - rng[1]) / (rng[2] - rng[1])  
  
  if(plot) {  
    plot(answer, ...)  
  }  
  
  return(answer)  
}
```

**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.

# Recap From Last Time:

- ▶ **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!

...

# Recap From Last Time:

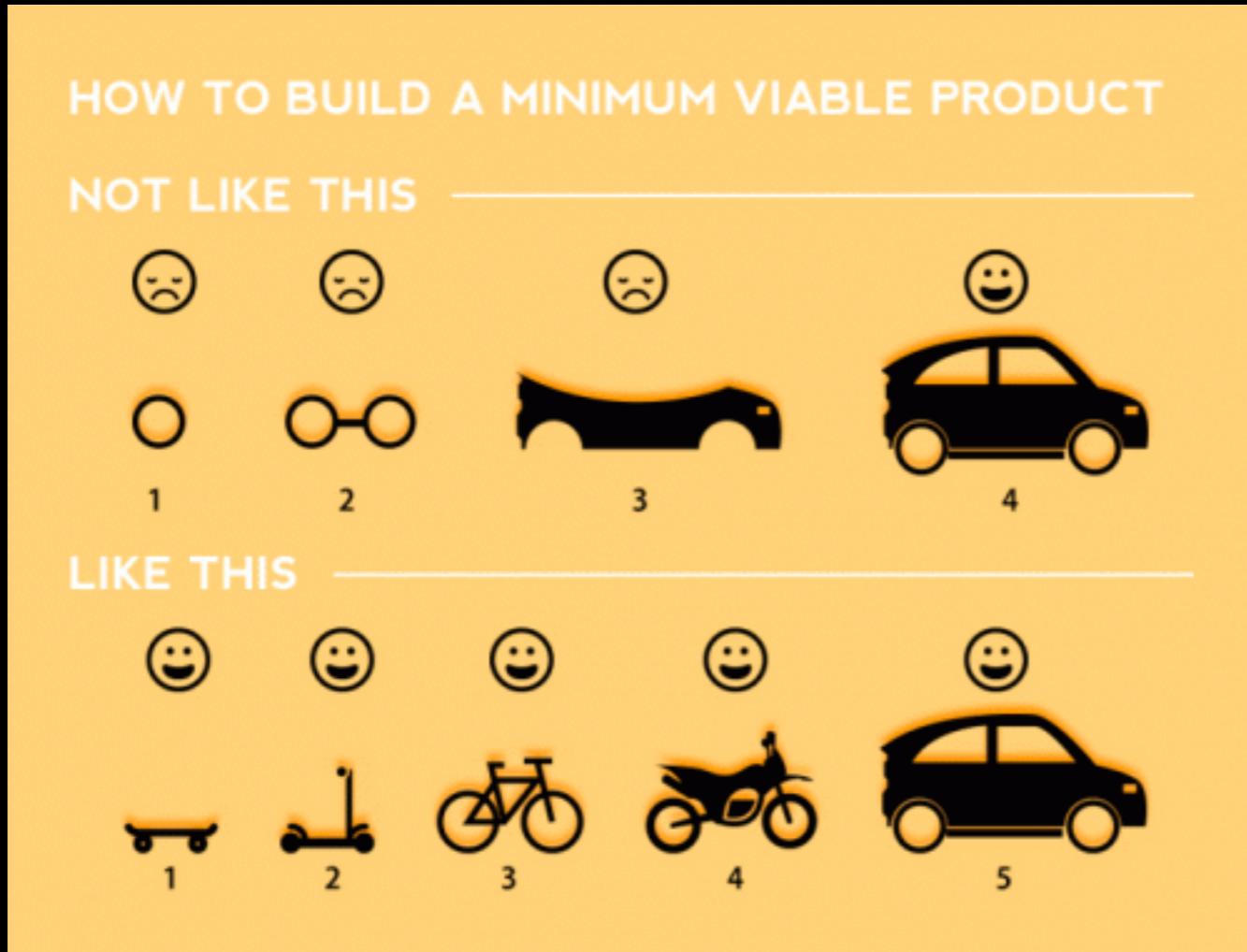
- **How:** Follow a step-by-step procedure to go from working code snippet to refined and tested function.

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
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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]

[ [MPA link](#) ]

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

# What is a function

```
1 name.of.function <- function(arg1, arg2) {  
  statements  
  return(something)  
}
```

- 1 **Name** (can be *almost* anything you want )
- 2 **Arguments** (i.e. input to your function)
- 3 **Body** (where the work gets done)

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)  
  
  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)  
  
  answer <- (x - rng[1]) / (rng[2] - rng[1])  
  
  if(plot) {  
    plot(answer, ...)  
  }  
  return(answer)  
}
```

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

```
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)  
  
  answer <- (x - rng[1]) / (rng[2] - rng[1])  
  
  if(plot) {  
    plot(answer, ...)  
  }  
  return(answer)  
}
```

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

# 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)
```

- 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?  
}
```

# 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() and sum() functions
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) )
}
```

# 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)
```

# 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) )  
}
```

# 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)  
  na.which <- which(na.in.both)  
  
  message("Found ", na.number, " NA's at position(s):",  
         paste(na.which, collapse=", "))  
  
  return( list(number=na.number, which=na.which) )  
}
```

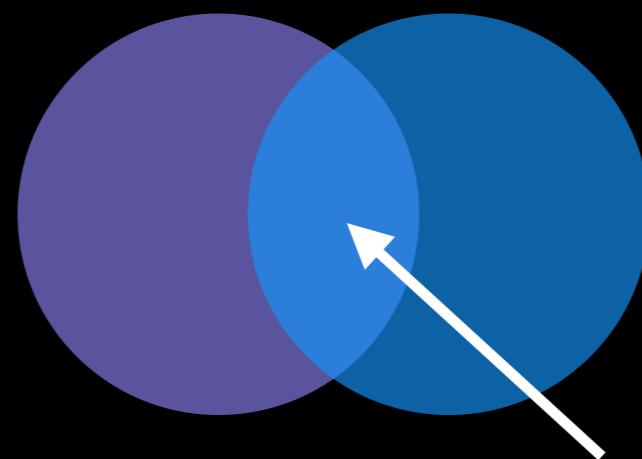
# Re-cap: Steps for function writing

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.

Break!

# One last example

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



intersect

Follow along!

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

Follow along!

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

# Simplify 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)

x <- df1$IDs
y <- df2$IDs

# Search for existing functionality to get us started...
??intersect

intersect(x, y)
[1] "gene2" "gene3"
```

Follow along!

```
# Close but not useful for returning indices yet.  
intersect(x, y)  
[1] "gene2" "gene3"
```

```
# Back to the documentation to find something more useful  
??intersect
```

Follow along!

```
# Close but not useful for returning indices yet.  
intersect(x, y)  
[1] "gene2" "gene3"  
  
# Back to the documentation to find something more useful  
?"%in%"  
  
# This looks like a more useful starting point - indices!  
x %in% y  
[1] FALSE TRUE TRUE
```

Follow along!

```
# Close but not useful for returning indices yet.  
intersect(x, y)  
[1] "gene2" "gene3"  
  
# Back to the documentation to find something more useful  
?"%in%"  
  
# 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 results to yield intersect
```

Follow along!

```
# Putting together  
cbind( x[ x %in% y ], y[ y %in% x ] )  
[,1] [,2]  
[1,] "gene2" "gene2"  
[2,] "gene3" "gene3"
```

```
# Make it into a first function
```

Follow along!

```
# Putting together
```

```
cbind( x[ x %in% y ], y[ y %in% x ] )  
[,1] [,2]  
[1,] "gene2" "gene2"  
[2,] "gene3" "gene3"
```

```
# Make it into a first function
```

```
gene_intersect <- function(x, y) {  
  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"
```

Follow along!

```
# Previous function for vector input
gene_intersect <- function(x, y) {
  cbind( x[ x %in% y ], y[ y %in% x ] )
}

# Lets change to take input data frames
gene_intersect2 <- function(df1, df2) {
  cbind( df1[ df1$IDs %in% df2$IDs, ],
         df2[ df2$IDs %in% df1$IDs, "exp"] )
}

# Correct but yucky format for 2nd colnames
gene_intersect2(df1, df2)
  IDs exp df2[df2$IDs %in% df1$IDs, "exp"]
2 gene2   1                               -2
3 gene3   1                               1
```

Follow along!

```
# Our input $IDs column name may change so lets add flexibility  
# By allowing 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 our current function.
```

Follow along!

```
# Looks complicated - simplify for human consumption!

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
```

```
# Looks much better

gene_intersect4 <- function(df1, df2, gene.colname="IDs") {

  df1.name <- df1[,gene.colname]
  df2.name <- df2[,gene.colname]

  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"),
                   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 message:
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/bggn213\\_W19/lectures/#6](https://bioboot.github.io/bggn213_W19/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.

# 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 ~13,250 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



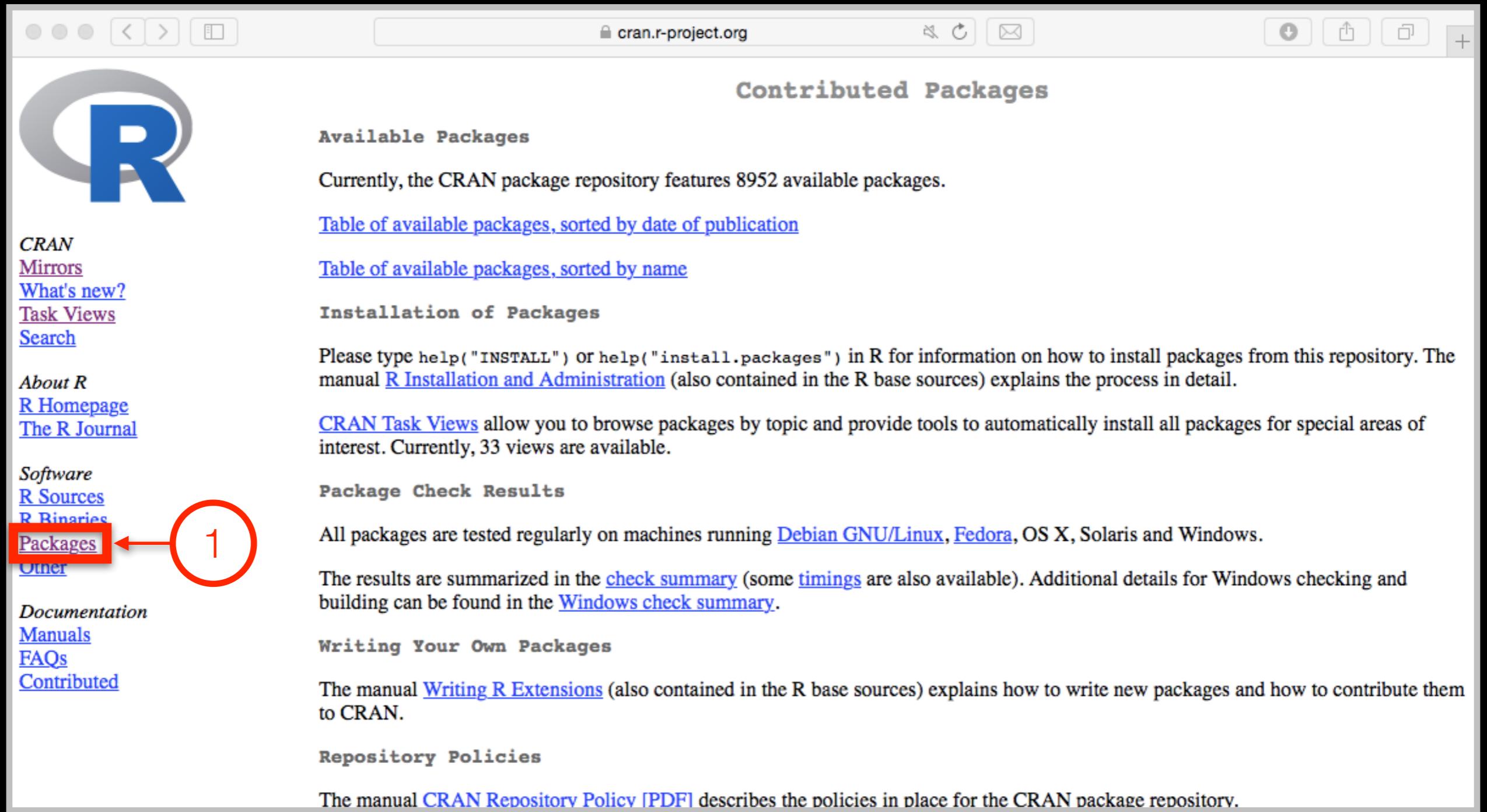
R packages can be of variable quality and often there are multiple packages with overlapping functionality.

**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>



The screenshot shows the CRAN website's 'Contributed Packages' page. The left sidebar has a large R logo and links for CRAN, Mirrors, What's new?, Task Views, Search, About R, R Homepage, and The R Journal. Below these are Software links: R Sources, R Binaries, Packages (which is highlighted with a red box and circled with a red arrow containing the number '1'), and Other. Documentation links include Manuals, FAQs, and Contributed. The main content area has sections for Available Packages, Installation of Packages, Package Check Results, Writing Your Own Packages, and Repository Policies. Each section contains descriptive text and links to manuals or summaries.

**Contributed Packages**

**Available Packages**

Currently, the CRAN package repository features 8952 available packages.

[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.

**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 [check summary](#) (some [timings](#) are also available). Additional details for Windows checking and building can be found in the [Windows check summary](#).

**Writing Your Own Packages**

The manual [Writing R Extensions](#) (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 3.6 has 1,473  
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

“Accessible, curated metagenomic data through ExperimentHub”

Pasolli *et al.*

**Nature Methods** 2017, 14:1023–1024.

# Installing a bioconductor package

```
> source("https://bioconductor.org/biocLite.R")  
> biocLite()  
> biocLite("GenomicFeatures")
```

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

# Installing a bioconductor package

```
> source("https://bioconductor.org/biocLite.R")  
> biocLite()  
> biocLite("GenomicFeatures")
```

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, rgl, rentrez, igraph,**  
**blogdown, shiny, msa, flexdashboard, 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 its 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

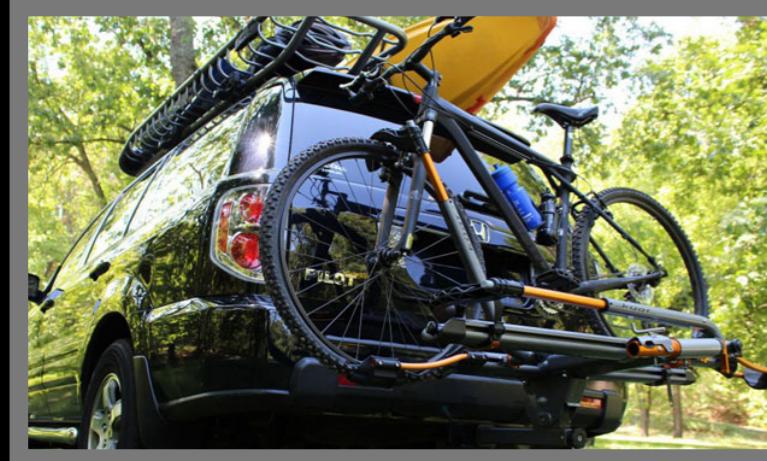
# Key Idea:



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!

# 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/> >

[Muddy Point Assessment Link]