
DAY 2. A beginners guide to solving biological problems in R

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Course material:
<http://logic.sysbiol.cam.ac.uk/teaching/Rcourse/>

Original slides by Ian Roberts and Robert Stojnić

Day 1 review

1. Introduction to R and its environment
 - Running R from the command line
 - Rstudio
 - Creating and calculating with vectors
 - Loading packages
2. Data structures
 - Data frames, factors, matrices, lists
3. Data analysis example
 - reading from and writing to files
4. Programming techniques
 - manipulating data frames, looping, if statements
5. Statistics
 - distributions, tests, linear models

Day 2 Schedule

1. Writing scripts
 - Building scripts for routine analysis
 - Script automation using R batch mode
2. Extending R with customized functions
 - Learn how to write your own R functions
3. Advanced data analysis and basic R graphics
 - Integrate data from different sources and analyze
4. More on R graphics
 - Produce publication quality charts

Writing custom scripts & running R batch mode analysis

1

The R scripting language

Scripting

- A script is a series of instructions that when executed sequentially automates a task
 - A script is a good solution to a repetitive problem
- The art of good script writing is
 - understanding exactly what you want to do
 - expressing the steps as concisely as possible
 - making use of error checking
 - including descriptive comments
- R is a powerful scripting language, and embodies aspects found in most standard programming environments
 - procedural statements
 - loops
 - functions
 - conditional branching
- Scripts may be written in any standard text editor, e.g. notepad, gedit, kate
 - RGui (Mac and Windows) has a built-in text editor

An example script

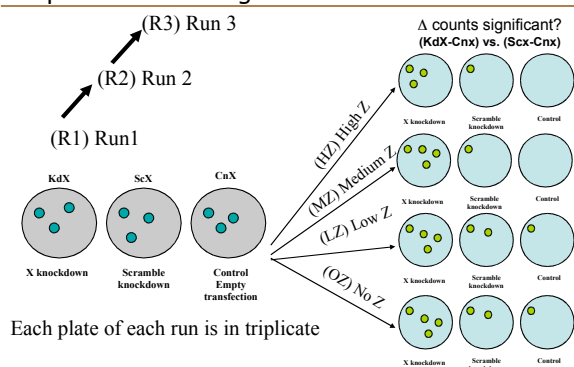
Scripting

- Colony forming assays provide a measure of cellular proliferation. They are used as read outs for various biological systems
 - A well controlled study may involve multiple samples, treatments and controls (probably replicated).
 - This produces a lot of 'count' data, ideally suited to routine script processing
- Encapsulating the analysis into an R script requires a clear understanding of the problem and data structure

CFA experimental design Scripting

- Expression of gene X may prevent cells from proliferating in high concentrations of compound Z. The theory is tested by knocking down gene X and growing cells in varying concentrations of compound Z.
 - Three repeat runs (same cell line)
 - Gene X knockdown --> KdX
 - Scramble gene X knockdown control --> ScX
 - Control (transfect empty vector) --> CnX
 - 4 concentrations of compound Z
 - High (HZ), Medium (MZ), Low (LZ), None (OZ)
 - The experiment is replicated over 3 successive weeks
 - Run1 (R1), Run2 (R2) and Run3 (R3)
 - 108 counts in total

Colony forming assay experimental design



Preparing the calculation(s) Scripting

- We need to make barplots of counts for the KDX-CNX and SCX-CNX for each concentration of Z.
- We will group the repeat runs & replicates, and take an average.
- A Wilcoxon Rank Sum test will tell us whether there is a significant level of protection for KDX in concentrations of Z
- We'll add in some data quality checks
 - Boxplots of repeat runs
 - Variance within replicates

We can copy & paste lines of code into a blank text document, try them out and keep the ones that work!

Importing data

Scripting

	R1						R2						R3					
Plate	KDX		SCX		CNX		KDX		SCX		CNX		KDX		SCX		CNX	
Replicate	1	2	3	1	2	3	1	2	3	1	2	3	1	2	3	1	2	3
OZ																		
LZ																		
MZ																		
HZ																		

Fine for humans, bad for R

Run	Plate	Treatment	Count
Run1	Plate1	Treatment1	Count1
Run1	Plate1	Treatment2	Count2
Run1	Plate1	Treatment3	Count3
Run1	Plate2	Treatment1	Count4
Run1	Plate2	Treatment2	Count5
Run1	Plate2	Treatment3	Count6
Run1	Plate3	Treatment1	Count7
Run1	Plate3	Treatment2	Count8
Run1	Plate3	Treatment3	Count9
Run1	Plate4	Treatment1	Count10
Run1	Plate4	Treatment2	Count11
Run1	Plate4	Treatment3	Count12
Run1	Plate5	Treatment1	Count13
Run1	Plate5	Treatment2	Count14
Run1	Plate5	Treatment3	Count15
Run1	Plate6	Treatment1	Count16
Run1	Plate6	Treatment2	Count17
Run1	Plate6	Treatment3	Count18
Run1	Plate7	Treatment1	Count19
Run1	Plate7	Treatment2	Count20
Run1	Plate7	Treatment3	Count21
Run1	Plate8	Treatment1	Count22
Run1	Plate8	Treatment2	Count23
Run1	Plate8	Treatment3	Count24
Run1	Plate9	Treatment1	Count25
Run1	Plate9	Treatment2	Count26
Run1	Plate9	Treatment3	Count27
Run1	Plate10	Treatment1	Count28
Run1	Plate10	Treatment2	Count29
Run1	Plate10	Treatment3	Count30
Run1	Plate11	Treatment1	Count31
Run1	Plate11	Treatment2	Count32
Run1	Plate11	Treatment3	Count33
Run1	Plate12	Treatment1	Count34
Run1	Plate12	Treatment2	Count35
Run1	Plate12	Treatment3	Count36
Run2	Plate1	Treatment1	Count37
Run2	Plate1	Treatment2	Count38
Run2	Plate1	Treatment3	Count39
Run2	Plate2	Treatment1	Count40
Run2	Plate2	Treatment2	Count41
Run2	Plate2	Treatment3	Count42
Run2	Plate3	Treatment1	Count43
Run2	Plate3	Treatment2	Count44
Run2	Plate3	Treatment3	Count45
Run2	Plate4	Treatment1	Count46
Run2	Plate4	Treatment2	Count47
Run2	Plate4	Treatment3	Count48
Run2	Plate5	Treatment1	Count49
Run2	Plate5	Treatment2	Count50
Run2	Plate5	Treatment3	Count51
Run2	Plate6	Treatment1	Count52
Run2	Plate6	Treatment2	Count53
Run2	Plate6	Treatment3	Count54
Run2	Plate7	Treatment1	Count55
Run2	Plate7	Treatment2	Count56
Run2	Plate7	Treatment3	Count57
Run2	Plate8	Treatment1	Count58
Run2	Plate8	Treatment2	Count59
Run2	Plate8	Treatment3	Count60
Run2	Plate9	Treatment1	Count61
Run2	Plate9	Treatment2	Count62
Run2	Plate9	Treatment3	Count63
Run2	Plate10	Treatment1	Count64
Run2	Plate10	Treatment2	Count65
Run2	Plate10	Treatment3	Count66
Run2	Plate11	Treatment1	Count67
Run2	Plate11	Treatment2	Count68
Run2	Plate11	Treatment3	Count69
Run2	Plate12	Treatment1	Count70
Run2	Plate12	Treatment2	Count71
Run2	Plate12	Treatment3	Count72
Run3	Plate1	Treatment1	Count73
Run3	Plate1	Treatment2	Count74
Run3	Plate1	Treatment3	Count75
Run3	Plate2	Treatment1	Count76
Run3	Plate2	Treatment2	Count77
Run3	Plate2	Treatment3	Count78
Run3	Plate3	Treatment1	Count79
Run3	Plate3	Treatment2	Count80
Run3	Plate3	Treatment3	Count81
Run3	Plate4	Treatment1	Count82
Run3	Plate4	Treatment2	Count83
Run3	Plate4	Treatment3	Count84
Run3	Plate5	Treatment1	Count85
Run3	Plate5	Treatment2	Count86
Run3	Plate5	Treatment3	Count87
Run3	Plate6	Treatment1	Count88
Run3	Plate6	Treatment2	Count89
Run3	Plate6	Treatment3	Count90
Run3	Plate7	Treatment1	Count91
Run3	Plate7	Treatment2	Count92
Run3	Plate7	Treatment3	Count93
Run3	Plate8	Treatment1	Count94
Run3	Plate8	Treatment2	Count95
Run3	Plate8	Treatment3	Count96
Run3	Plate9	Treatment1	Count97
Run3	Plate9	Treatment2	Count98
Run3	Plate9	Treatment3	Count99
Run3	Plate10	Treatment1	Count100
Run3	Plate10	Treatment2	Count101
Run3	Plate10	Treatment3	Count102
Run3	Plate11	Treatment1	Count103
Run3	Plate11	Treatment2	Count104
Run3	Plate11	Treatment3	Count105
Run3	Plate12	Treatment1	Count106
Run3	Plate12	Treatment2	Count107
Run3	Plate12	Treatment3	Count108

We will create a data frame of factors comprising Run, Plate, Treatment, Replicate
The response variable is counts.

We have 3 spreadsheets of data
Run1counts.csv
Run2counts.csv
Run3counts.csv

We will need to write a procedure that reads in the data
Note that our script will require a consistent data format

Factors

Response

Prepare for raw data

Script walkthrough 1

- Open a blank text document, and prepare to write this script
 - The data is contained in three files:
 - 11_CFA_Run1Counts.csv
 - 11_CFA_Run2Counts.csv
 - 11_CFA_Run3Counts.csv
 - Load in the data and concatenate it into a single data frame

```
# load in the data from the three runs into three separate data frames t1,
# t2, t3
t1 = read.csv("11_CFA_Run1Counts.csv")
t2 = read.csv("11_CFA_Run2Counts.csv")
t3 = read.csv("11_CFA_Run3Counts.csv")

# concatenate the three data frames into a single data frame
colony = rbind(t1, t2, t3)

# (or use one of the loops from yesterday...)
```

Example code:
11_CFAcountData.R

Import raw data

Script walkthrough 2

- Data is by default read in as factors, i.e. all input strings are enumerated and stored as numbers
- The three separate data frame have no indication of which number they came from. We will add a column indicating this:

```
# add the missing Run column - factors are stored as numbers !
runNum <- factor( rep( 1:3, each=36 ), labels=c("Run1","Run2","Run3") )
colony <- cbind( "Run" = runNum, colony )

# reorder factor levels in their natural order (instead of alphabetical)
colony$Treatment <- factor(colony$Treatment, c("OZ", "LZ", "MZ", "HZ"))
colony$Plate <- factor(colony$Plate, c("KDX","SCX","CNX"))

# show the full table
colony
```

The tapply function a brief digression

- Assume we have the following data for heights of 5 males and females:

```
data <- data.frame(gender=c("Male", "Male", "Female",  
                             "Female", "Female"), height=c(6, 6.1, 5.8, 6, 5.95))
```

gender	height
1 Male	6.00
2 Male	6.10
3 Female	5.80
4 Female	6.00
5 Female	5.95
- By calling `mean()` on the height column we can get the average of all 5 people, but how do we get average separately for males and females?
- `tapply()` lets us do exactly this
 - It applies a function to grouped data:
- `tapply(data$height, data$gender, mean)`
 data groups function

Undertake data analysis Script walkthrough 3

- We need the means of the triplicate counts for each Run
 - Broken down by plate type (KDX,SCX,CNX) and Z treatment concentration (OZ,LZ,MZ,HZ)

```
### Part 2. Investigating data ###  
tapply(colony$Count, list(colony$Run, colony$Plate,  
colony$Treatment), mean)
```

We can plot a graph of this. It gives us the variation in counts per run

```
par(mfrow=c(4,2,2,2))  
boxplot(Count~Run*Plate*Treatment, las=2, cex=0.2,  
data=colony)
```

Better still, lets plot a grouped bar chart of mean counts per plate type per Z treatment

```
barplot(tapply(colony$Count, list(colony$Plate,  
colony$Treatment), mean), beside=T)
```

```
## OZ  
Run1 98.33333 129.6667 108.3333  
Run2 180.33333 206.0000 188.6667  
Run3 282.33333 288.6667 265.6667  
  
## LZ  
Run1 75.0000 53.0000 21.66667  
Run2 136.3333 103.6667 32.00000  
Run3 157.0000 180.6667 46.66667  
  
## MZ  
Run1 29.66667 6.333333 0.3333333  
Run2 47.00000 11.66667 2.0000000  
Run3 73.00000 17.33333 3.666667  
  
## HZ  
Run1 6.333333 1.333333 0.3333333  
Run2 12.00000 0.333333 0.0000000  
Run3 18.66667 0.333333 0.0000000
```

Summarize & save the analysis Script walkthrough 4

- we need a reshaped, background corrected, table of results on which to perform our tests
- for clarity where possible use dollar (\$) notation (work only with data frames)

```
### Part 3. Summarizing data ###  
result <- tapply(colony$Count, list(colony$Treatment, colony$Plate), mean)  
result <- data.frame(result) # result of tapply is matrix, convert to dataframe  
result
```

```
# calculate kdx and scx values after background correction  
kdx = result$KDX - result$CNX  
scx = result$SCX - result$CNX
```

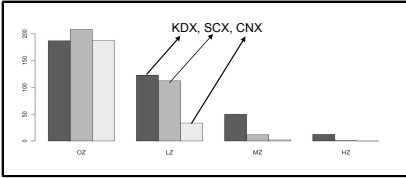
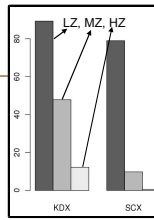
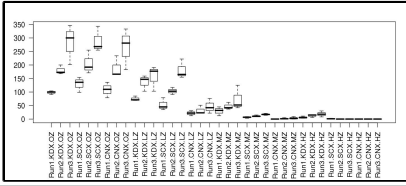
```
result <- cbind(kdx, scx)  
# remove the OZ entry  
result <- result[-1,]  
barplot(result, beside=T)
```

-ve subscripts
mean 'delete'

```
wilcox.test(result[,1], result[,2], paired=T)  
cor.test(result[,1], result[,2], paired=T)  
write.csv(result, "CPAresults.csv")
```

We can plot the results as a barchart, and undertake an appropriate two sample classical test

We find that the difference in means is not significant (would expect observation to occur 1:4 times), and that the scramble and knockdown counts have a 90% correlation



Bar chart shows run average counts of various plate types in different Z treatments. KDX is the only plate type that had growth in high Z.

```
> wilcox.test(result[,1],result[,2],paired=TRUE)

Wilcoxon signed rank test

data: result[, 1] and result[, 2]
V = 6, p-value = 0.25
alternative hypothesis: true location shift is not equal to 0
```

```
> cor.test(result[,1],result[,2],paired=TRUE)

Pearson's product-moment correlation

data: result[, 1] and result[, 2]
t = 2.5584, df = 1, p-value = 0.2372
alternative hypothesis: true correlation is not equal to 0
sample estimates:
cor
0.9313792
```

Would expect to see trend 1 in 4 times. There is a 93% correlation between the knockdown of gene X and scramble control and cell counts response when grown in compound Z.

Script steps review

Script walkthrough 5

- Excel formatted data needs to be exported as comma separated values text (or tab!)
- Get the data into R
 - `read.csv()` ... to assign the data to an object
- Produce exploratory plots
 - `boxplot()`
 - `barplot()`
- Undertake statistical tests
 - `cor.test()`
 - Spearman's rank correlation test
 - `wilcox.test()`
 - Wilcoxon test with two sets of paired data ... Mann-Whitney U test
- Write out the results
 - `write.csv()`
 - exports data as comma separated list
 - `save.image()`
 - could also save the R environment after analysis (we didn't do this)

Exercise

Colony forming assay script

- Enter the text of the count data script, and save the file.
 - To run the count data script in R, type
 - `source("filename")` # the script is available as 11_countData.R
- Each step of the script is executed, and the results displayed.
- We need to export the graphical output to a file, and the R objects also need to be saved.
 - Modify the script as follows:
Section 3, line directly after **apply** command insert:

```
jpeg(file="fig1.jpg",width=1600,height=800,res=150)
par(oma=c(4,2,2,2))
-- <boxplot commands in middle> ...
dev.off()
jpeg(file="fig2.jpg",width=675,height=900,res=150)
-- <barplot commands in middle> ...
dev.off()
```


Section 4, line directly after **result** command insert:

```
jpeg(file="fig3.jpg",width=1600,height=800,res=150)
-- <barplot commands in middle> ...
dev.off()
```

Batch processing R scripts

Scripting

- Scripts can be run without ever launching R, using R CMD batch mode.

quit R and type the following in a linux terminal

```
R CMD BATCH --no-restore 11_CFAcountData.R
```

or if you write all of graphical output to files:

```
Rscript 11_CFAcountData.R (works only with recent R versions)
```

User functions

2

Introducing ... User functions

- All R commands are functions.
- Functions perform calculations, possibly involving several arguments, then return a value to the calling statement.
- The calculation maybe any process, might or might not have return value
 - It need not be arithmetic
- User functions extend the capabilities of R by adapting or creating new tasks that are tailored to your specific requirements.
- User functions are a special kind of object

Defining a new function

- Parts of function definition: name, arguments, procedural steps, return value

```
sqXplusX <- function(x){  
  x^2 + x  
}
```
- **sqXplusX** is the function name
- **x** is the single argument to this function and it exists only within the function
- everything between brackets { } are procedural steps
- the **last** calculated value is the function return value
- after defining the function, we can use it:

```
> sqXplusX(10)  
[1] 110
```

Named and default arguments

- Example of function with more than one named argument:

```
powXplusX <- function(x, power=2){  
  x^power + x  
}
```
- Now we have two arguments. The second argument has a default value of 2.
- Arguments without default value are required, those with default values are optional.

```
> powXplusX(10)  
[1] 110  
> powXplusX(10, 3)  
[1] 1010
```

arguments matched based on **position**

```
> powXplusX(x=10, power=3)  
[1] 1010
```

arguments matched based on **name**

Assignments with arguments

User functions

```
sqXplusX <- function(x){  
  x^2 + x  
}
```

You can use a blank document in gedit, nedit or other text editor to hold these commands for you, then copy / paste the instructions into R

- Now try this ...

```
a <- matrix(1:100, ncol=10, byrow=T) # make some dummy data  
b <- sqXplusX(a) # transform a by sqXplusX, assign result to b  
b # to view the result
```

- sqXplusX user function is now an R object, check its arguments and list it in the current workspace

```
> args(sqXplusX)
```

```
> ls()
```

```
> sqXplusX
```

Don't add brackets to see the definition of sqXplusX

Assigned or anonymous ...

User functions

- Functions may be assigned a name, or anonymously created within an operation
 - Anonymous functions are really useful in `apply()` style procedures

```
apply(object, margin, function)
```

- E.g. I have a 10 x 10 matrix and want to square each item, and add the item to itself

```
a<-matrix(1:100, ncol=10, byrow=T)
```

```
a # to view new object
```

```
apply(a, c(1,2), function(x) x^2+x)
```

x is transiently assigned each item of a, and this is passed as an argument to the anonymous function

1 means by rows, 2 means by columns [1st or 2nd margin]
c(1,2) means do both rows and columns

Functions occupy their own space

User functions

- Objects created in functions are not available to the general environment unless returned.
 - they are said to be out of scope
 - Scope relates to the accessibility of an object.
- A function can only return one object.
- Custom functions disappear when R sessions end, unless the function object is saved in an Rdata file or sourced from a script.
 - A really useful function could be added to your .Rprofile file, and would always be ready for you at launch
- You could also make a package
 - Beyond the scope of the beginners course!!!!

Script / function tips

User functions

- If your script repeats the same style command more than twice, you should consider writing a function
- Writing functions makes your code more easily understandable because they encapsulate a procedure into a well-defined boundary with consistent input/output
- Functions should not be longer than one-to-two screens of code, keep functions clean and simple
- Look at other functions to get ideas for how to write your own ...
 - Display function code by entering the function's name without brackets.

File commands for extending scripts & user functions

Generic file commands

```
dir(...,pattern="txt")
```

Retrieve working directory file listing filtered by pattern. Note pattern is a regular expression, not a shell wildcard

```
glob2rx("*.txt")
```

Changes wildcards to regular expressions!

```
unlink(...)
```

Remove (permanently) a file from system

```
system(...)
```

Execute a shell command from within R

Result can not be coerced to an object, only available to linux R

```
> glob2rx("*.txt")  
[1] "^.*\\.txt$"
```

Text manipulation for extending scripts & user functions

- Text manipulation and file name mangling ... that's a technical term

```
grep(pattern, object)
```

- If pattern is not found, grep returns a 0 length object.
 - Test for null with `is.null()`

```
sub(pattern, replacement, object)
```

```
gsub(pattern, replacement, object)
```

- Sub replaces first occurrence only, gsub does them all.

```
cat("...", file=...)
```

- Outputs text to a file, or prints it on screen if file=""
 - cat requires "\n" to be given for new lines ... try ...

```
cat("Hello World!") ; cat("Hello World!",sep="\n") ; cat("Hello World!",sep="\n",file="world.txt")
```

- cat is extremely useful for writing scripts or generating reports on-the-fly

Error reporting for extending scripts & user functions

- Your code should report errors if inconsistency is detected.

```
stop(...)
```

- Stops execution of a function and reports a custom error message

```
is.family(...)
```

- Functions that can be used to test for a variety of conditions place them inside **if** structures to check that all is well

```
if( !is.numeric(x) ){ stop ("Non numeric value entered. Cannot continue.") }
```

If the object x is non numeric (e.g. Text has been entered when numbers were required), then stop execution and report message

The **is.family**

is.array	is.language	is.primitive
is.atomic	is.list	is.R
is.baseNamespace	is.loaded	is.R6
is.call	is.logical	is.real
is.character	is.matrix	is.recursive
is.class	is.mts	is.relistable
is.classDef	is.no	is.restart
is.classUnion	is.na	is.s4
is.complex	is.no.data.frame	is.startOfClass
is.data.frame	is.na.default	is.startOfMethod
is.duplicate	is.na.factor	is.startOfValue
is.environment	is.name	is.string
is.element	is.namespace	is.stepfun
is.empty.model	is.no.POSIXt	is.symbol
is.expression	is.null	is.symmetric
is.factor	is.numeric	is.symmetric.matrix
is.finite	is.numeric.Date	is.table
is.function	is.numeric.POSIXt	is.true
is.generic	is.numeric.version	is.ts
is.graphicsSymbol	is.object	is.unsorted
is.group	is.opened	is.unsorted
is.incomplete	is.infinite	is.unsorted
is.infinite	is.integer	is.unsorted
is.integer	is.integer.version	is.unsorted
is.pairlist	is.pairlist	is.unsorted

Temperature conversion exercise

User functions

- Centigrade to Fahrenheit conversion is given by

- F = 9/5 C + 32

- Write a function that converts between temperatures.

- The function will need two named arguments
 - temperature (**t**) is numeric
 - units (**unit**) is character
 - They will need default values, e.g. t=0, unit="c"

- The function should report an appropriate error if inappropriate values are given

```
if( !is.numeric(t) ) { .... }  
if( !unit %in% c("c","f") ) {...}
```

- The function should print out the temperature in F if given in C, and vice versa

Functions with named arguments are defined with the following syntax

```
myFunc<-function(arg=defaultValue,...)
```

- Why not add a third scale?

K=C+273.15

Example code:
12_convTemp.R

Building the solution

- It is difficult to write large chunks of code, instead start with something that works and build upon it

- E.g. to solve the temperature conversion exercise:

- start with the function powXplusX (from some slides ago)
- modify the argument names
- delete the old code, for now just print out the input arguments
- save the function file, load it into R and try it out
- now add the two lines for input checking from the previous slide
- try it out, see if passing a character for temperature gives the expected error
- now try to convert C into F and print out the result
- when that works, add the conversion from F to C

- If you get stuck, call us to help you !

Temperature conversion script

`convTemp<-function(t=0,unit="c"){ # convTemp is defined as a new user function requiring two arguments, t and unit, the default values are 0 and "c", respectively.`

```
  if( !is.numeric(t) ){
    stop("Non numeric temperature entered") # Exception error if character given for
    temperature
  }
  if(! (unit %in% c("c","f","k"))){
    stop("Unrecognized temperature unit. \n Enter either (c)entigrade, (f)ahrenheit
    or (k)elvin") # Exception error if unrecognized units entered
  }
  # Conversion for centigrade
  if(unit=="c"){
    fTemp <- 9/5 * t + 32
    kTemp <- t + 273.15
    output <- paste(t,"C is: \n",fTemp,"F \n",kTemp,"K \n")
    cat(output)
  }
  # Conversion for Fahrenheit
  if(unit=="f"){
    cTemp <- 5/9 * (t-32)
    kTemp <- cTemp + 273.15
    output <- paste(t,"F is: \n",cTemp,"C \n",kTemp,"K \n")
    cat(output)
  }
  # Conversion for Kelvin
  if(unit=="k"){
    cTemp <- t - 273.15
    fTemp <- 9/5 * cTemp + 32
    output <- paste(t,"K is: \n",cTemp,"C \n",fTemp,"F \n")
    cat(output)
  }
}
```

Units must be entered
in quotes, as it's a
character object

Example code:
12_convTemp.R

Advanced data processing

3

Combining data from multiple sources *Gene clustering example*

- R has powerful functions to combine heterogeneous data into a single data set
- Gene clustering example data:
 - five sets of differentially expressed genes from various experimental conditions
 - file with names of experimentally verified genes
- Gene clustering exercise:
 - 1.combine this dataset into a single table and cluster to see which conditions are similar
 - 2.repeat the clustering but only on a subset of experimentally verified genes

Combining gene tables

- input files have two columns: gene names and fold change
- we want to combine all five tables into a single table, with 0 for missing values

LPR2	3.5795	Psa	3.8529	Iola	3.0121	Iola	3.3019	brat	5.2812
fs(1yh	3.1376	vmd	3.6457	CG31368	2.8063	CG6919	2.9965	ct	4.828
CG6954	2.7482	ct	3.2011	Kr-h1	2.7262	CG31368	2.817	CG31163	4.3345
Psa	2.7012	fs(1yh	3.1489	svp	2.7005	CG5149	2.7675	LPR2	3.6882
zfh2	2.6247	bld	3.1229	miub	2.6475	Kr-h1	2.7647	vmd	3.6666
Fur1	2.4413	zfh2	2.6421	CG5149	2.5248	TER94	2.6286	zfh2	3.5314
ct	2.3804	RhoB7B	2.6022	run	2.4759	ina	2.5748	pros	3.4307
S	2.3674	pros	2.5679	ina	2.4302	CG11153	2.4795	Psa	3.3968
rux	2.3574	CG1124	2.5475	CG6954	2.4235	run	2.3631	fs(1yh	3.3869
RhoB7B	2.26	S	2.5404	CG11153	2.3845	CG14888	2.0938	CG31241	2.9973
CG14889	2.1735	pc	2.5111	Awd	2.2295	S	-2.0243	HmgZ	2.9226
pc	2.1421	Fur1	2.43	CG6919	2.1324	rux	-2.0668	Fur1	2.7469
pros	2.0882	PHDP	2.304	CG14888	2.087	pc	-2.3437	RhoB7B	2.7189
Kr-h1	-2.0447	CG31241	2.2802	Psa	-2.0276	corto	-2.5596	pc	2.6543
CG5149	-2.1521	rux	2.2232	rux	-2.093	fs(1yh	-2.6211	Toll-7	2.6161
ina	-2.2102	CG14889	2.1752	fs(1yh	-2.141	brat	-2.9904	rux	2.5975
CG14888	-2.4346	CG31163	2.1606	CG1124	-2.155	CG14889	2.3054	CG14889	2.3054
CG31368	-2.4793	HmgZ	2.0795	Fur1	-2.1588	ct	-3.3404	S	2.2304
Trim9	-2.616	svp	-2.0404	S	-2.2539	zfh2	-4.4947	CG1124	2.0216
Awd	-3.0395	TER94	-2.1807	corto	-2.2618	CG6954	-4.7244	Kr-h1	-2.1439
		cgf413	-2.4404	pc	-2.3017			ina	-2.1793
		brat	-2.7256	CG14889	-2.4393			CG5149	-2.1892
		CG31368	-2.7293	zfh2	-2.5884			run	-2.2194
		miub	-2.9055	HmgZ	-3.6328			Trim9	-2.251
		Awd	-3.1413	bld	-3.7627			cgf413	-2.3821
		Iola	-3.6982	brat	-3.7716			bld	-3.0293
								CG6919	-3.3719

Gene clustering

Script walkthrough 1

- To make the big table we first need to find out all the genes present in at least one of the files
- Make sure not to use factors in read.delim()

```
# start with an empty collection of genes
genes <- c()
for( fileNum in 1:5 ){
  # load in files 13_DiffGenes1.tsv ...
  t <- read.delim(paste("13_DiffGenes", fileNum, ".tsv", sep=""),
    as.is=TRUE, header=FALSE)

  # label the input columns to help code readability
  names(t) <- c("gene", "expression")
  genes <- union(genes, t$gene)
}

# for tidiness order our genes by name
genes <- sort(genes)

genes # show all genes
```

when loading in character data use **as.is=T** to prevent it being converted to factors!

union() is a set operation, combines two vectors by eliminating duplicates. There are also **intersect()** and **setdiff()**

Example code:
13_geneClustering.R

Gene clustering

Script walkthrough 2

- Using the complete list of genes, we can create the big table and fill in the values:

```
# make the destination table [rows = unique genes, cols = file numbers]
values <- matrix(0, nrow=length(genes), ncol=5)
rownames(values) <- genes # name the rows with the complete gene names

for(fileNum in 1:5){
  # read in the file again
  t <- read.delim(paste("13_DiffGenes", fileNum, ".tsv", sep=""),
    as.is=T, header=F)
  names(t) <- c("gene", "expression")

  # match the names of the genes to the rows in our big table
  index <- match(t$gene, rownames(values))
  # copy the expression levels
  values[index, fileNum] <- t$expression
}

# we use index to pick the rows in such way that they match the gene order in the input file
```

match() returns the index of first argument in the second, i.e. index of input file genes in the big table

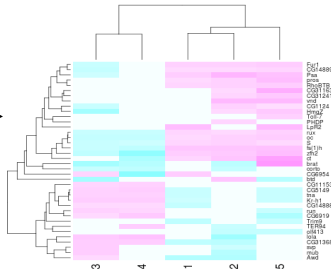
Gene clustering

Script walkthrough 3

- Now we can do hierarchical clustering:

```
heatmap(values, scale="none", col = cm.colors(256))
```

Values from the matrix are colour-coded. Rows and columns are re-arranged according to similarity



Gene clustering

Script walkthrough 4

- In a second part of our analysis, we want to produce the same heatmap but only based on a list of experimentally verified genes
- The problem is data is not formatted in the most convenient way:

genes	citation
oc,run,RhoBTB,CG5149,CG11153,S,Fur1	Segal et al, Development 2001
tna,Kr-h1,rux	Krejci et al, Development 2002

Gene clustering

Script walkthrough 5

- We load in this table, and only extract the gene names, then we use them to select a subset of **values** matrix

```
# load in the tab-delimited file with genes and citations
t.exp <- read.delim("13 ExperimentalGenes.tsv", as.is=T)
# split all gene names by "," and then flatten it out into a single vector
experim.genes <- unlist( strsplit(t.exp$genes, ",") )
```

`unlist()` flattens out a nested list into a single vector

`strsplit()` splits a vector of strings by a custom split character (","), the results is a list of split values for each element of input vector

```
# redo the heatmap by using just the genes in the experimentally verified set
is.experimental <- rownames(values) %in% experim.genes
heatmap(values[ is.experimental, ], scale="none", col = cm.colors(256))
```

Gene clustering review

- We load in the five tables twice - first to collect gene names, then to load expression values
- Based on expression table (**values**) we construct a clustered heatmap first on the whole set of genes, then on a selected subset
- Go through the code, try it out it and understand it
- Try answering the following questions:
 - what is **rownames(values)** ?
 - why is **rownames(values)[index]** and **t\$gene** giving the same output?
 - what is a difference between **rownames(values) %in% experim.genes** and **experim.genes %in% rownames(values)**

Example code:
13_geneClustering.R

Graphics

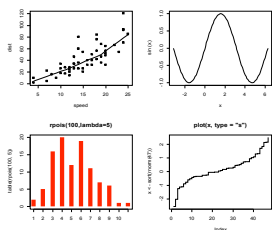
4

Starting out with R graphics

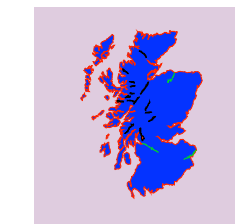
Graphics

- R provides several mechanisms for producing graphical output
 - Functionality depends on the level at which the user seeks interaction with R
 - graphics systems, packages, devices & engines
- High level graphics
 - Functions compute an appropriate chart based up on the information provided. Optional arguments may tailor the chart as required
 - Interaction is at traditional graphics system level. The user isn't required to know much about anything
- Low level graphics
 - The user interacts with the drawing device to build up a picture of the chart piece by piece.
 - This fine granular control is only required if you seek to do something exceptional
- R graphics produces plots using a painter's model
 - Elements of the graph are added to the canvas one layer at a time, and the picture built up in levels. Lower levels are obscured by higher levels, allowing for blending, masking and overlaying of objects.

High level vs. Low level plotting Graphics



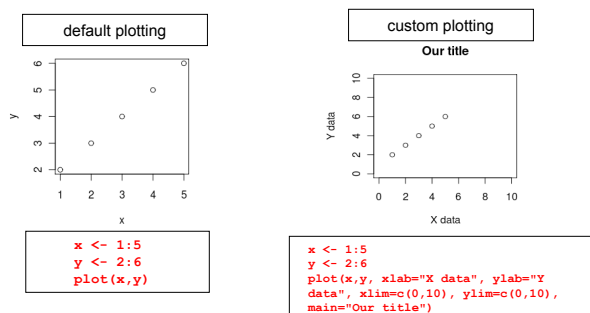
High level plotting
example (plot)



Low level plotting
(Scotland by blighty package)

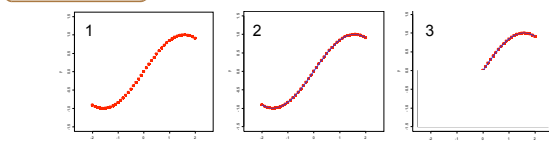
Essential plotting - plot()

- `plot()` is the main function for plotting, it takes x,y values to plot and also lots of graphical parameters (see **?par** for all of them)



R graphics uses a painter's model

```
x <- seq(-2, 2, 0.1)
y <- sin(x)
```



```
plot(y~x, ylim=c(-1.5,1.5),
xlim=c(-2.5,2.5),
col="red", pch=16, cex=1.4)
```

```
lines(y~x, ylim=c(-1.5,1.5),
xlim=c(-2.5,2.5), col="blue",
lty=1,lwd=2)
```

```
rect(-2.5,0,2.5,-1.5,
col="white", border="white")
```

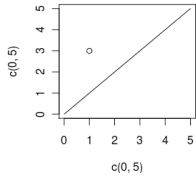
xlim, ylim = axis limits
col = line colour
pch = plotting character [**example (points)**]
cex = character expansion [scaling factor]

lty = line type
lwd = line width
rect = rectangle

Example code:
14_painterModel.R

Plotting x,y data - plot(), points(), lines()

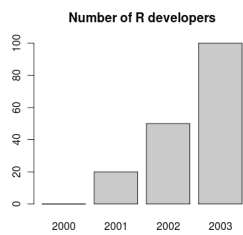
- **plot()** is used to start a new plot, accepts x,y data, but also data from some objects (like linear regression). Use the parameter **type** to draw points, lines, etc (see **?plot**)
- **points()** is used to add points to an existing plot
- **lines()** is used to add lines to an existing plot



```
plot(c(0, 5), c(0, 5), type="l") # draw as line from (0,0) to (5,5)
points(1, 3) # add a point at 1,3
```

Making bar plots - barplot()

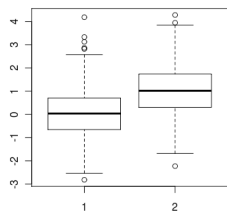
- visualizing a vector of data can be done with bar plots, using function **barplot()**



```
data <- c("2000"=0, "2001"=20, "2002"=50, "2003"=100)
barplot(data, main="Number of R developers")
```

Making box plots - boxplot()

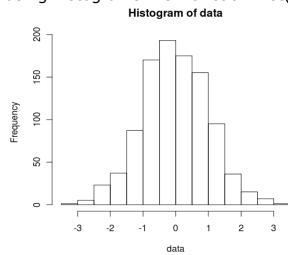
- when a spread of data needs to be visualised, we can use boxplots with function **boxplot()**



```
data1 <- rnorm(1000, mean=0)
data2 <- rnorm(1000, mean=1)
boxplot(data1, data2)
```

Making histograms - hist()

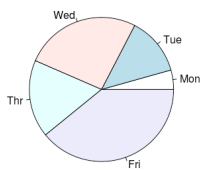
- when we need to look at the distribution of data, we can visualize it using histograms with function `hist()`



```
data <- rnorm(1000)
hist(data)
```

Pie charts - pie()

- to visualise percentages or parts of a whole we can use pie charts with function `pie()`



```
data <- c("Mon"=1, "Tue"=3, "Wed"=6, "Thr"=4, "Fri"=9)
pie(data)
```

Typical plotting workflow

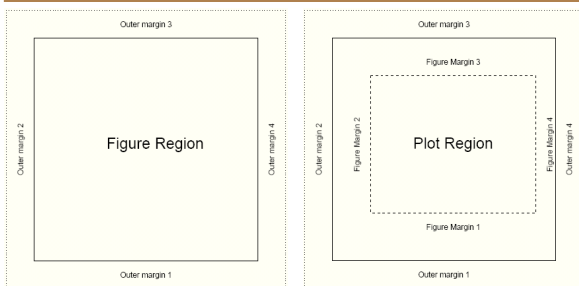
- Set the plot layout and style - `par()`
 - Set the number of plots you want per page
 - Set the outer margins of the figure region
 - The distance between the edge of the page and the figure region, or between adjacent plots if there are multiple figures per page
 - Set the inner margins of the plot
 - The distance between the plot axes and the labels & titles
 - Set the styles for the plot
 - Colours, fonts, line styles and weights
- Draw the plot - `plot(x,y, ...)`

Setting graphics layout and style - par()

- par()** Top level graphics function
- parameter specifies various page settings. These are inherited by subordinate functions, if no other styles are set.
 - Specific colours and styles may be set globally with **par**, but changed ad hoc in plotting commands
 - The global setting will remain unchanged, and reused in future plotting calls.
 - par** sets the size of page and figure margins
 - Margin spacing is in 'lines'
 - par** is responsible for controlling the number of figures that are plotted on a page
 - par** may set global colouring of axes, text, background, foreground, line styles (solid/dashed), if figures should be boxed or open etc. etc.

type **par()** to get a list of top down settings which may be set globally

Page settings with **par** Graphics



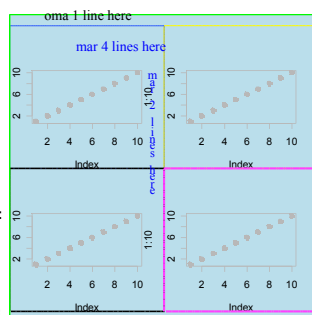
```
par(mfrow=c(1,1))  
one figure on page  
par(oma=c(2,2,2,2))  
equal outer margins
```

```
par(mar=c(5,4,4,2))  
Sets space for x & y labels, a main title, and a thin margin on the right
```

Numbering: bottom, left, top, right

Page layout plot exercise Graphics

- ```
par(mfrow=c(2,2))
• 2 x 2 figures per page
par(oma=c(1,0,1,0))
• 1 line spacing top and bottom
par(mar=c(4,2,4,2))
• 4 lines at bottom & top
• 2 lines left & right
par(bg="lightblue",fg="darkgrey")
• light blue background
• dark grey spots
par(pch=16,cex=1.4)
• Large circles for spots
• Execute 4 times with different colors:
plot(1:10)
box("figure",lty=3,col="blue")
• Draw a blue dashed line around plot
box("outer",lty=1,lwd=3,col="green")
• Draw a green solid line around figure
```



See how the figure margins overlap  
Using painter's model

15\_parExample.R

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## Plotting characters for plot() size and orientation

**pch=** ...

Sets one of the 26 standard plotting character used.

Can also use characters, such as "."

**cex=** ...

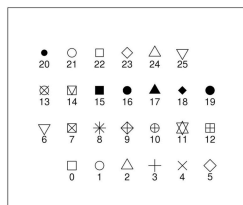
Character expansion. Sets the scaling factor of the printing character

**las=** ...

Axes label style. 1 normal, 2 rotated 90°

4 styles (0-3)

26 standard plotting characters



## Plotting characters exercise Graphics

16\_plottingChars.R

```
xCounter<-1
yCounter<-1
plotChar<-0
```

X-Y coordinates,  
Plotting character index counter

```
plot(NULL, xlim=c(0,8),
ylim=c(0,5), xaxt="n",
yaxt="n", ylab="", xlab=""
, main="26 standard plotting
characters")
```

Sets up an empty plotting area.  
Axis scale limits, xlim, ylim  
Don't draw axis ticks, xaxt, yaxt="n"  
Don't annotate axis, xlab, ylab=""  
Set a main title, main

```
while (plotChar < 26){
 if(xCounter < 7){
 xCounter <- xCounter+1
 } else {
 xCounter <- 1
 yCounter <- yCounter+1
 }
}
```

We want to print the characters in a  
7 x 4 grid. The if statement sets up  
the character plotting coordinates  
such that each time x = 7, make it 1  
again and increment the y axis by 1 at  
the same time

```
points(xCounter,yCounter,pch=plotChar,
cex=2)
text(xCounter,(yCounter-0.3),plotChar)
plotChar <- plotChar+1
}
```

While loop counts up to 25  
(0 to 25 = 26 iterations)  
And cycles through each pch  
available

## Annotating the plot

- plot accepts main title, subtitle, X label, Y label as standard arguments

```
plot(x, y, main="...", sub="...", xlab="...", ylab="...")
```

```
mtext(text="...", side=...)
```

- allows text to be written directly into the margin of a plot

```
text(x,y,labels="...")
```

- allows text to be written in the plot at x,y

```
legend(x,y, legend=...)
```

- produces a legend for the plot

## Appreciating drawing coordinates

- How do we know where to place items within the plot region when building up our customized graphs?
- Most of the time we can specify X,Y coordinates.
  - R calculates sensible pixel coordinates of plots from the data we provide. We don't need to worry about pixels, centimetre distances etc.
- `locator(...)`
  - Returns x,y coordinates from a mouse click within a plot
  - good for working out where to place legend items
- `identify(...)`
  - provides an id tag for the closest plotted point to a mouse click
  - useful if you want to label points on a chart
- `xy.coords(...)`
  - translates x,y coordinates into pixel coordinates
- Margin spacing is in lines
  - The exact distance is a factor of font family, style and size
  - Text may appear bunched or squashed if sufficient distance is not left between the axes and the caption

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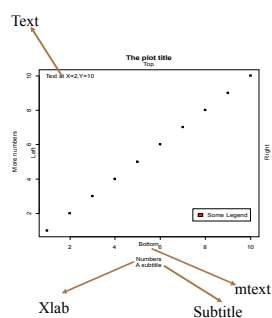
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## Building up a plot Graphics



### R code

```
par(mfrow=c(1,1))
par(bg="white",fg="black",cex=1)
par(oma=c(1,1,1,1))
par(mar=c(5,4,4,2)+0.1)

plot(1:10,main="The plot title",
sub="A subtitle", xlab="Numbers",
ylab="More numbers")

mtext(c("Bottom", "Left", "Top",
"Right"), c(1,2,3,4), line=.5)

text(2,10,"Text as you want to mouse click!")
legend(locator(1),"Some
Legend",fill="red")
```

align text left, right & centre with  
adj=(i,j) i.e centre is adj=(0.5,0.5), left  
is adj=(1,0) and right is adj=(0,1)

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## Plots with custom axes Graphics

- R `plot` doesn't support multiple Y axis by default
    - You have to make additional axes yourself!
  - Adding custom axis
- ```
axis(side=, at=, labels=, ...)
```
- If you want to specify custom axes, make sure you turn off the automatic axes in the plot / points call
- ```
plot(..., axes=F)
```

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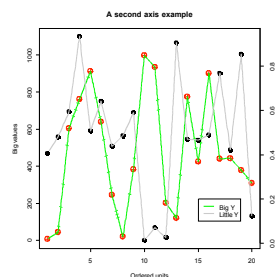
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## Adding a second Y axis

### Graphics



#### The trick

1. plot first Y series
2. use `par(new=T)` to overlay a second figure region
3. plot second series without axes
4. `axis(side=4, ...)` to add second Y axis
5. `mtext(side=4, ...)` to label second Y

## Example: The second Y series

### Graphics

```
x1<-1:20
y1<-sample(1000,20)
y2<-runif(20)
y2axis<-seq(0,1,.2)
```

Demo data

```
par(mar=c(4,4,4,4))
```

Set up equivalent figure margins

```
plot(x1,y1,type="p",pch=10,cex=2,col="red",
 main="A second axis example",
 ylab="Big values",ylim=c(0,1100),
 xlab="Ordered units")
points(x1,y1,type="l",lty=3,lwd=2,col="green")
```

Plot and label first Y series

Connect dots with a line

```
par(new=T)
```

Overlay a second plot region

```
plot(x1,y2,type="p",pch=20,cex=2,col="black",axes=FALSE,bty="n",xlab="",ylab="")
points(x1,y2,type="l",lty=3,lwd=2,col="grey")
```

Plot second Y series, but suppress labels

```
axis(side=4,at=pretty(y2axis))
mtext("Little values",side=4,line=2.5)
```

Anotate second Y axis

```
legend(15,0.2,c("Big Y","Little Y"),lty=1,lwd=2,col=c("green","grey"))
```

Add legend, note X,Y is on second Y axis scale

## Use of colour in R

### Graphics

- Colour is usually expressed as a hexadecimal code of Red, Green, and Blue counterparts
  - No good for humans.
- R supports numerous colour palettes which are available through several "colour" functions.
  - `colours()` # get inbuilt names of known colours
    - RGB primaries may take on a decimal intensity value of 0 to 255
      - 255 is #FF in hexadecimal
        - White is #FF FF FF
        - Black is #00 00 00
  - `rgb()` # converts red green blue intensities to colour
    - Strangely, likes decimalized intensities (ie. 0 is black, 1 is white)

```
> rgb(1,1,1)
[1] "FFFFFF"
```

```
> par(mfrow=c(2,2))
> plot(1:10,col="#FF00FF")
> plot(1:10,col=rgb(1,0,1))
> plot(1:10,col="magenta")
```

## Colour Ramps & Palettes Graphics

• Heatmaps use colour depth to convey data values. Cold colours are typically low values, and light colours are high state values. This is a colour ramp.

• R supports numerous graded colour charts. Specify *n*, to set the number of gradations required in the palette

```
rainbow(n)
heat.colors(n)
terrain.colors(n)
topo.colors(n)
cm.colors(n)
```

Rainbow Colours



Heat Colours



Terrain Colours



Topological Colours



Cyan Magenta Colours



19\_colourCharts.R

You can specify a user defined palette of indexed colours:

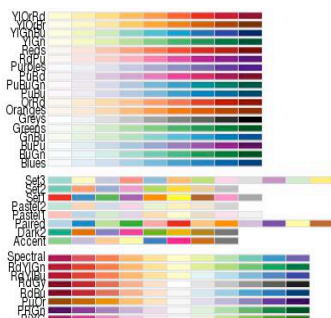
```
palette(rainbow(7)) # creates 7 indexed colours (1:7) based on
rainbow palette R O Y G B I V !!!
```

## Colour packages: RColorBrewer Graphics

- This add on package provides a series of well defined colour palettes. The colours in these palettes are selected to permit maximum visual discrimination
- Access the RColorBrewer library functions ...

```
library("RColorBrewer")
• Check out the available palettes
display.brewer.all(n=NULL, type="all", select=NULL, exact.n=TRUE)
• Define your own palette based on one of RColorBrewers'
myCol<-brewer.pal(n,...) # n=number of colours, "..." is the palette name
```

## RColorBrewer named palettes Graphics



## Saving plots to files

- Unless specified, R plots all graphics to the screen
- To send plots to a file, you need to set up an appropriate graphics device ...

```
postscript(file="a_name.ps", ...)
pdf(file="_.pdf", ...)
jpeg(file="_.jpg", ...)
png(file="_.png", ...)
```

- Each graphics device will have a specific set of arguments that dictate characteristics of the outputted file
  - `height=`, `width=`, `horizontal=`, `res=`, `paper=`
    - Top tip: jpeg, A4 @ 300 dpi, portrait, size in pixels
    - `jpg(file="my_Figure.jpg", height=3510, width=2490, res=300)`
    - Postscript & pdf work in inches by default, A4 = 8.3" x 11.7"

- Graphics devices need closing when printing is finished

```
dev.off()
```

for example:

```
png("tenPoints.png", width=300, height=300)
plot(1:10)
dev.off()
```

## Thoughts when plotting to a file Graphics

- Its very tempting to send all graphical output to a pdf file. Caution!
  - For high resolution publication quality images you need postscript. Set up postscript file capture with the following function

```
postscript("a_file.ps", paper="a4")
```

- postscript images can be converted to JPEG using ghostscript (free to download) for low resolution lab book photos and talks
- PDF images will grow too large for acrobat to render if plots contain many data points (e.g. Affymetrix MA plots)
- Automatically send multiple page outputs to separate image files using `...file="somename%02d.jpg"`
- Don't forget to close graphics devices (i.e. the file) by using
  - `dev.off()`

## Plotting exercise Graphics

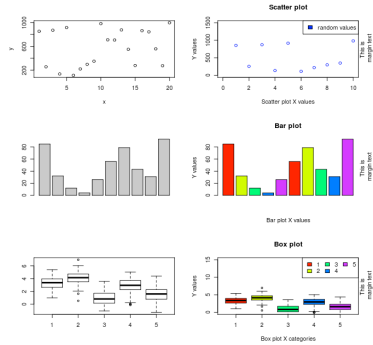
- Exercise:
  - Make a full A4 page figure comprising of 6 plots: 2 each of XY plot (`plot()`), barchart (`barplot()`) and box plots (`boxplot()`)
  - The two version of each plots should consist of: the default plot and a customised plot (change for instance colours, range, captions...)
  - Output the completed 6-panel figure to: screen, jpeg, postscript and pdf file
- Suggested route to solution:
  1. Generate some plotting data appropriate for each type of plot
  2. Write the code to produce the six plots, once plotting the data by using default plotting, one with some customisations you want
  3. To output the plot to screen, jpeg, postscript and pdf you will need to redo the plot multiple times - create a function to do a plotting and call it by redirecting graphical output to screen, jpeg file, poscript file and pdf file

20\_PanelPlotScript.R



## 6 Panel plots exercise

### Graphics



## References

- Official documentation on:
  - <http://cran.r-project.org/manuals.html>
- A good repository of R recipes:
  - Quick-R: <http://www.statmethods.net/>
- Don't forget that many packages come with tutorials ([vignettes](#))
- Website of this course:
  - <http://logic.sysbiol.cam.ac.uk/teaching/Rcourse/>
- R forums (stackoverflow & official):
  - <http://stackoverflow.com/questions/tagged/r>
  - <http://news.gmane.org/gmane.comp.lang.r.general>
- Plenty of textbooks to choose from, comprehensive list + reviews:
  - <http://www.r-project.org/doc/bib/R-books.html>

Thanks for your attention!

**END OF COURSE**