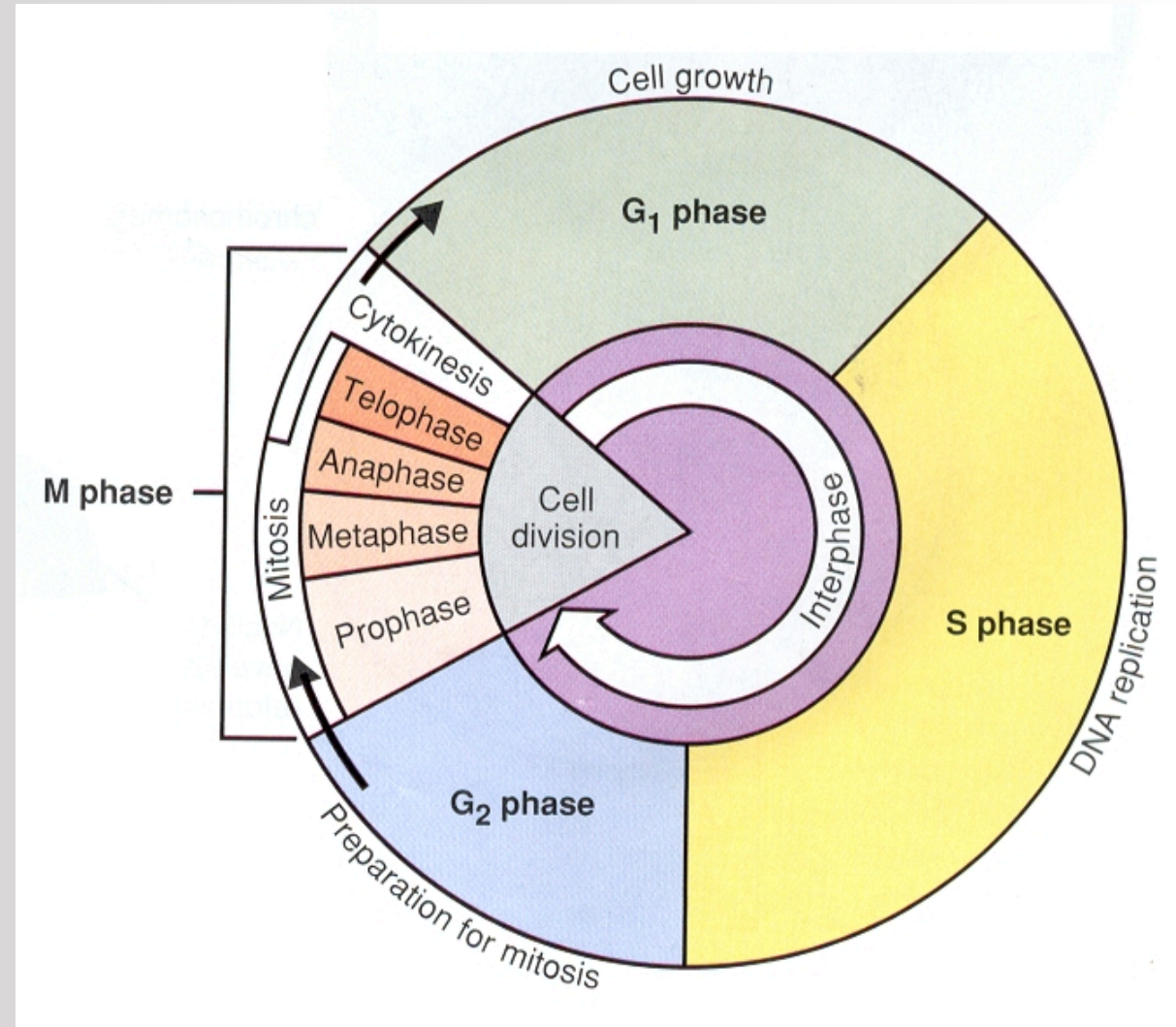


Analyze RNA-seq data and prepare
for quantitative PCR experiment

M2D4

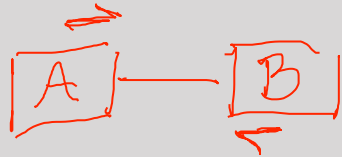
qPCR of p21 and GAPDH

- p21 (also CDKN1A)
 - Regulator of cell cycle progression at G1; arrest in G1 in response to stress
 - Tightly controlled by tumor suppressor p53
- GAPDH (glyceraldehyde 3-phosphate dehydrogenase)
 - Catalyzes glycolysis
 - housekeeping gene: constitutively and highly expressed in cells



Primer design guidelines

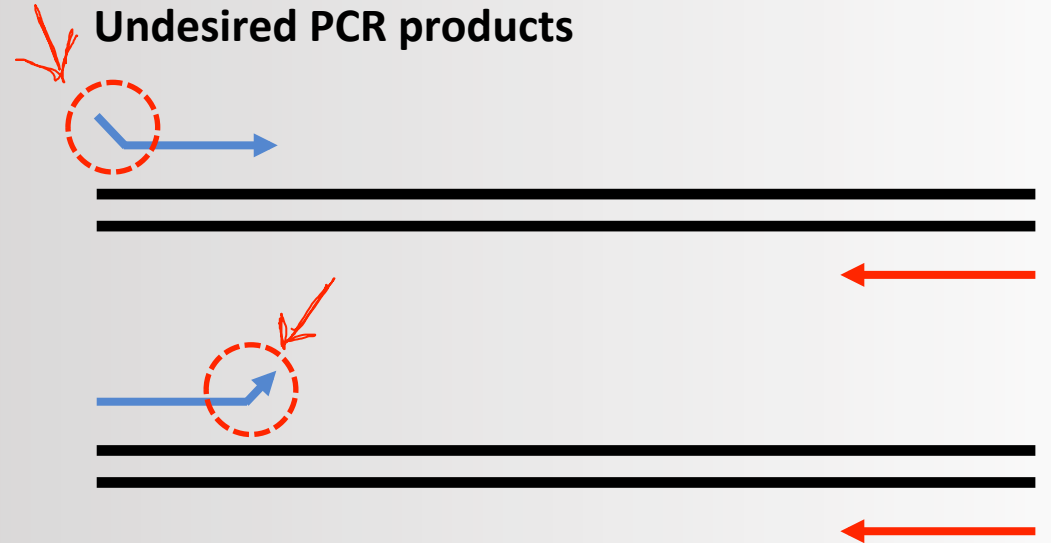
- Length \approx 100 bp
- GC-content = 50-60%
- 3' base preferentially a G or C
- $T_m \approx 60^\circ\text{C}$
- Must span exon-exon junction
- Does the location of mismatches matter?
- Does the length of alternative products matter?



Desired PCR product



Undesired PCR products



Our RNA-seq data



Seed cells



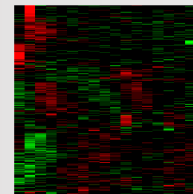
**Induce DNA damage
(etoposide)**

ACAUGGAUUA



TGTACCTAAT

**Purify RNA and
prepare cDNA**



RNA-seq

Our data structure –
preprocessed by Amanda

Rowranges
(Chromosomes,
number of exons,
IDs, etc.)

colData
(DLD-1 or BRCA2-/-,
+/- etoposide)

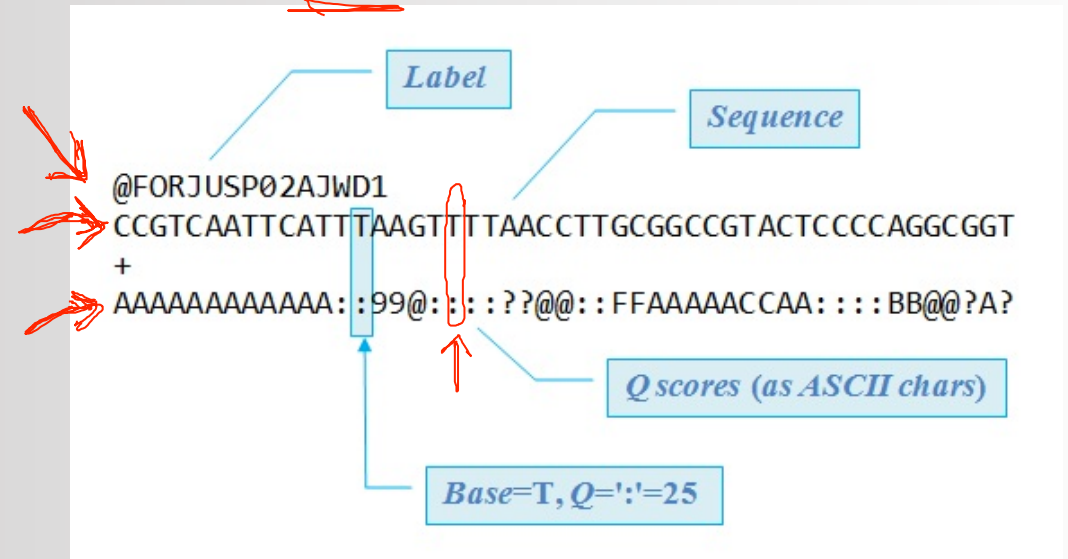
assay
(i.e. “counts”)

What does raw RNA-seq data look like?

The data:

- Forward (+ Reverse) read
- Quality score
- Barcode to identify sample
- Label to map sample to flow cell

Example .fastq file

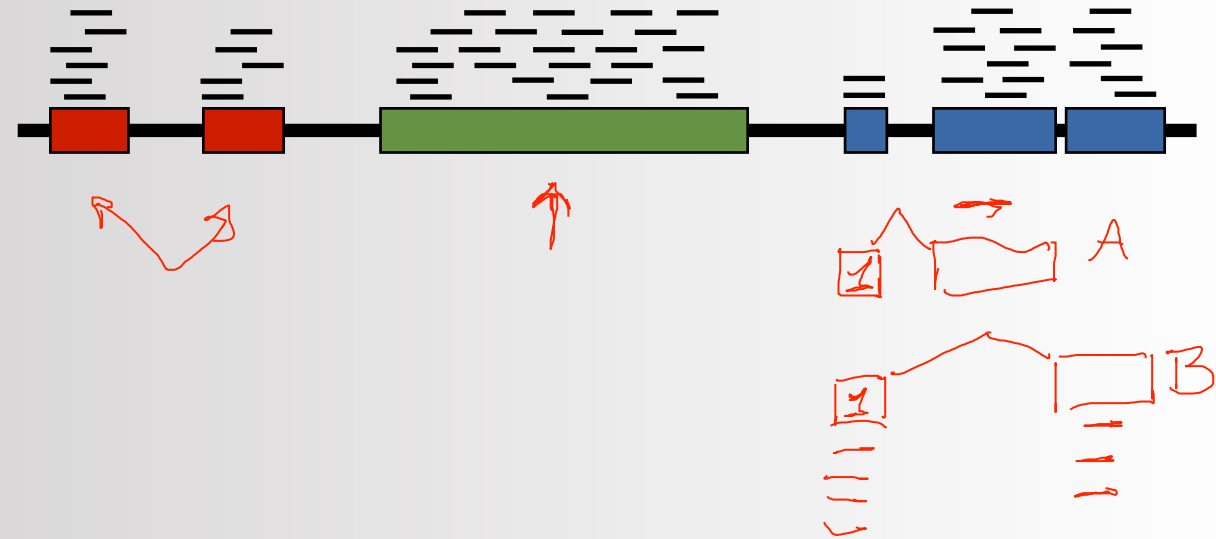


From RNA-seq data to transcriptomics

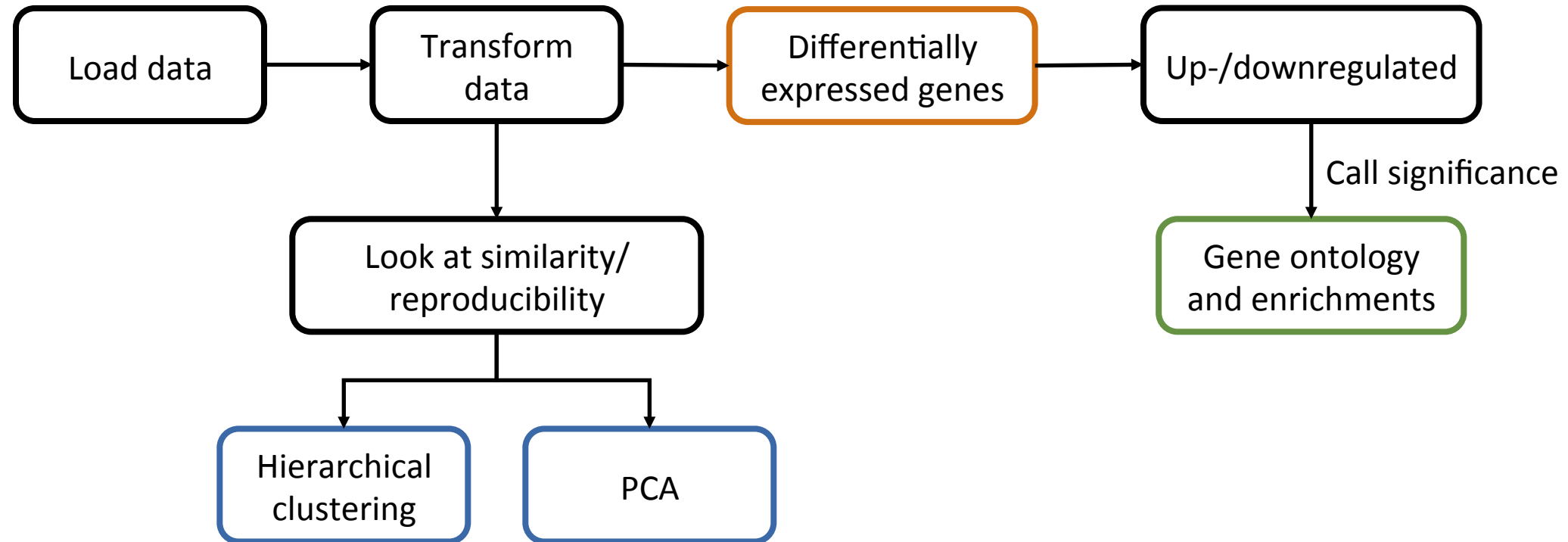
We can have from 8 million to 1.5 billion reads!

Processing the data:

- We count the number of times a gene is expressed
- We adjust for the total number of reads
- We adjust for the length of the gene

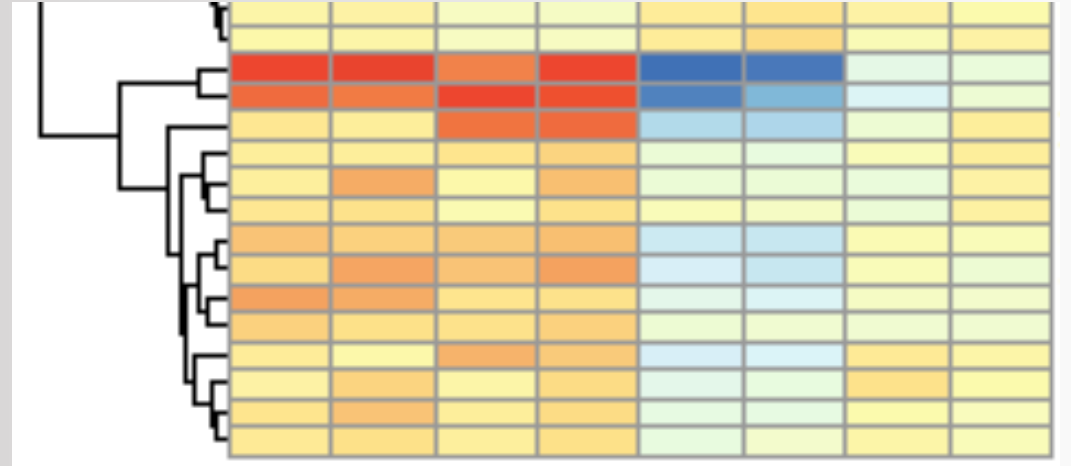


What you will do today



Are specific genes differentially expressed?

genes



gene X
gene Y
gene Z
.
.
.

DLD-1, etoposide
BRCA2-/-, etoposide
DLD-1, no drug
BRCA2-/-, no drug

samples

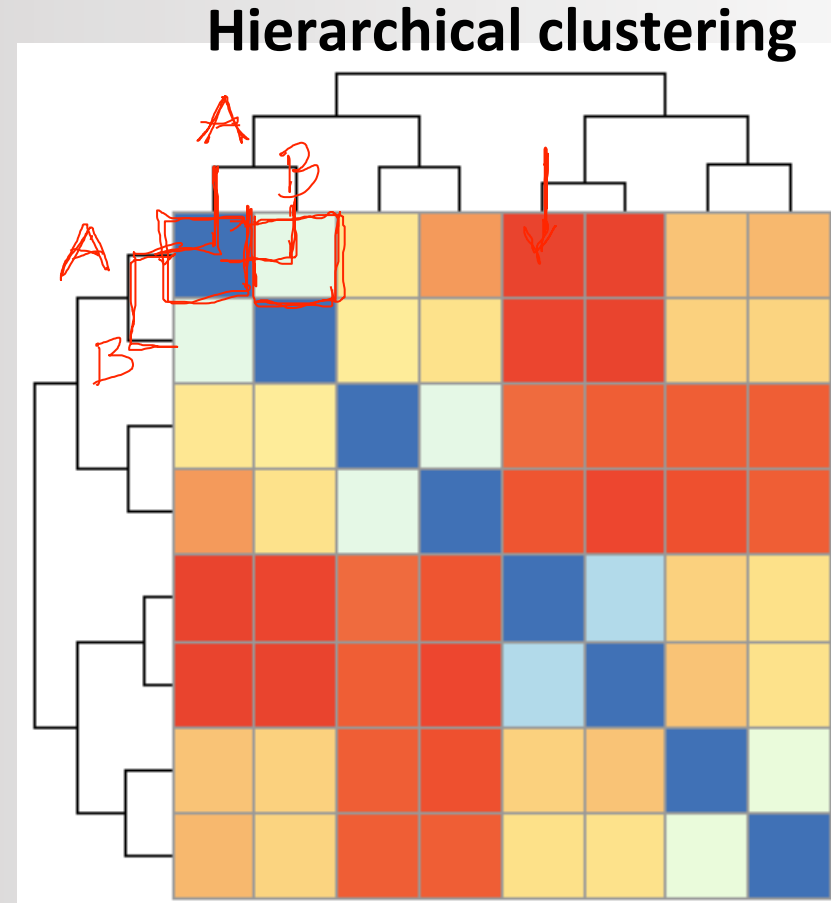
Which samples are most similar?

- Distance from one sample to another
- Symmetrical matrix

Handwritten scribble

$$\vec{x}_A = a_1, a_2, \dots, a_n$$

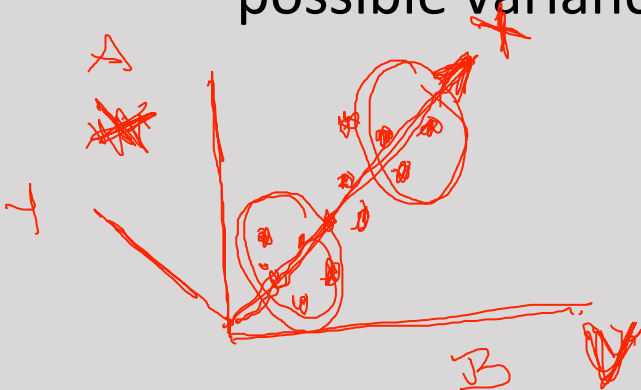
$$\vec{x}_B = b_1, b_2, \dots, b_n$$



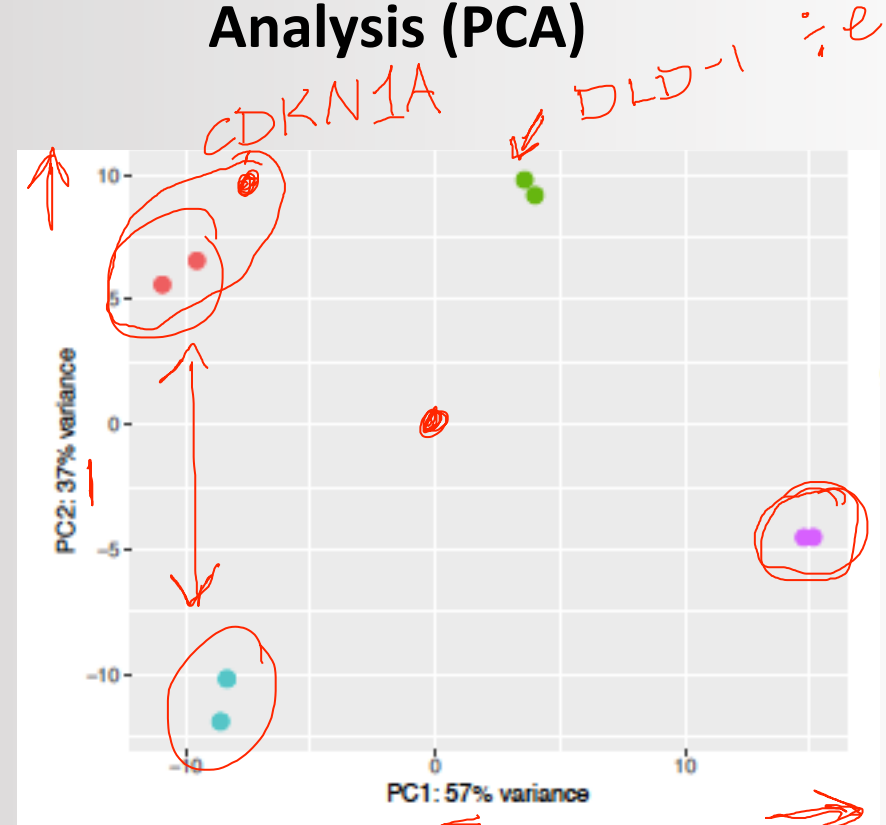
Samples:	DLD-1	BRCA2 -/-
Etoposide:	+	+
	-	-

Do samples change together?

- $\approx 20,000$ genes across 8 samples
 - Can we reduce the number of variables?
- PCA reduces dimensionality
 - Each component has the largest possible variance



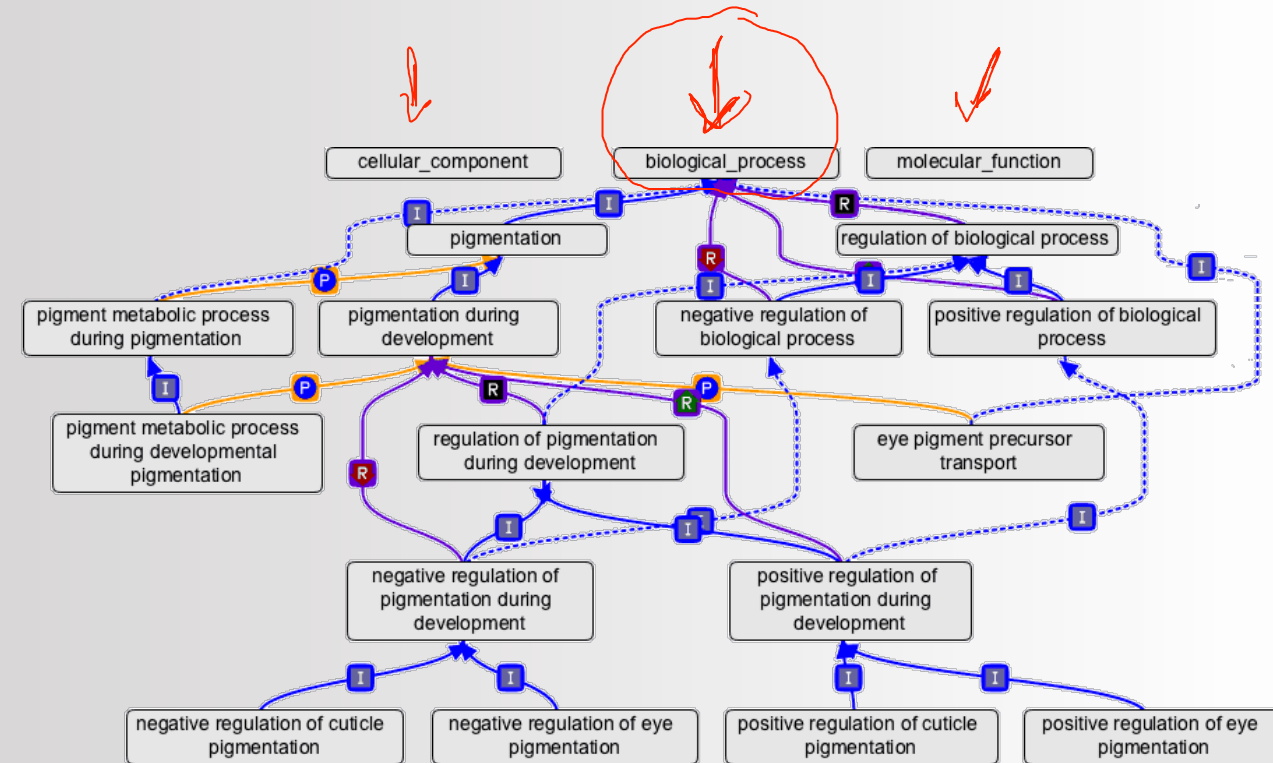
Principal Component Analysis (PCA)



Samples:	DLD-1	BRCA2 -/-
Etoposide:	+	+
	-	-

What is the function of genes?

- Manual annotation of $\approx 20,000$ genes is not feasible!
- GO (Gene Ontology) provides automatic annotations of biological function

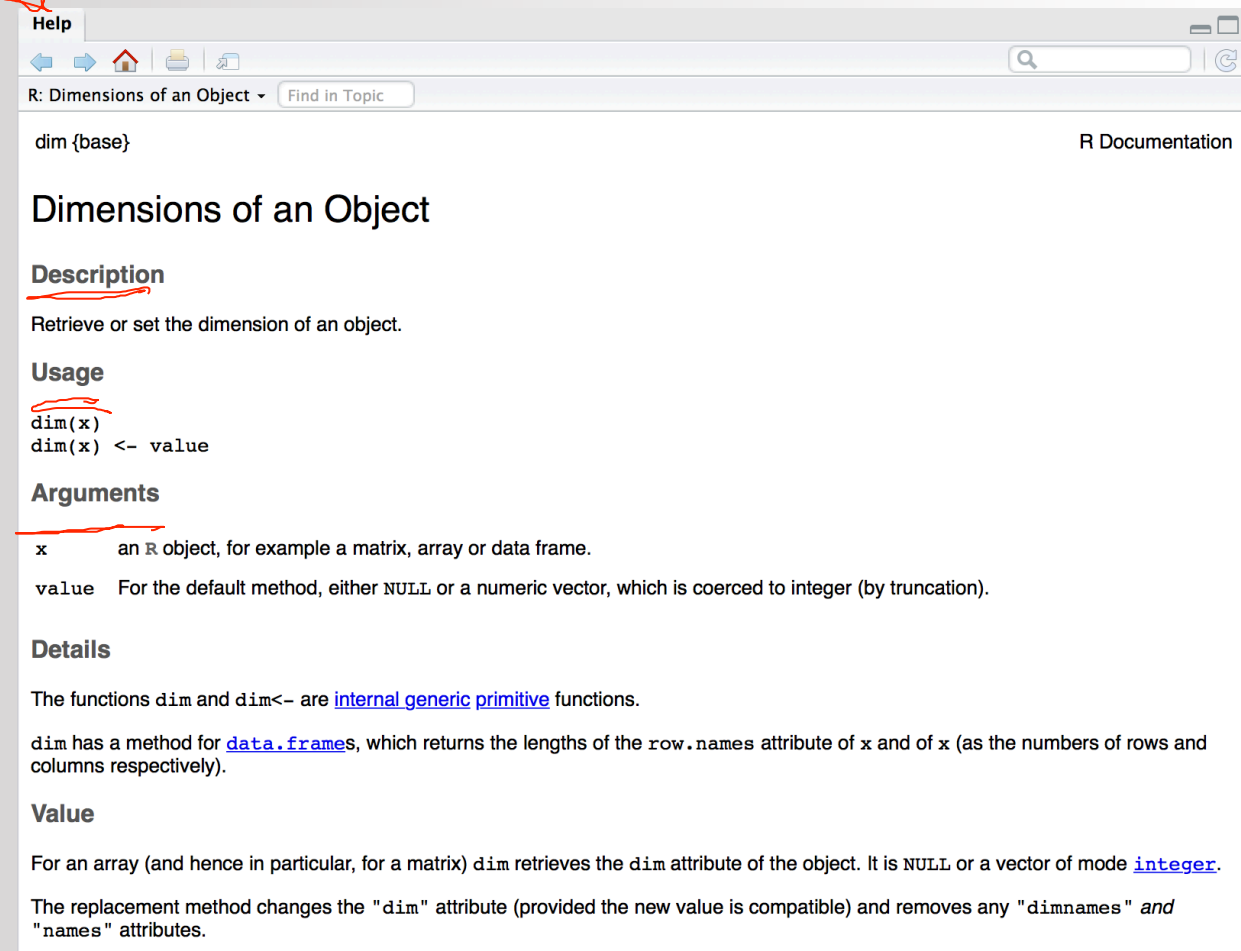


Getting help in R

Method 1
`?function`

Method 2
`help(function)`

Example: Type in `?dim` or `help(dim)`

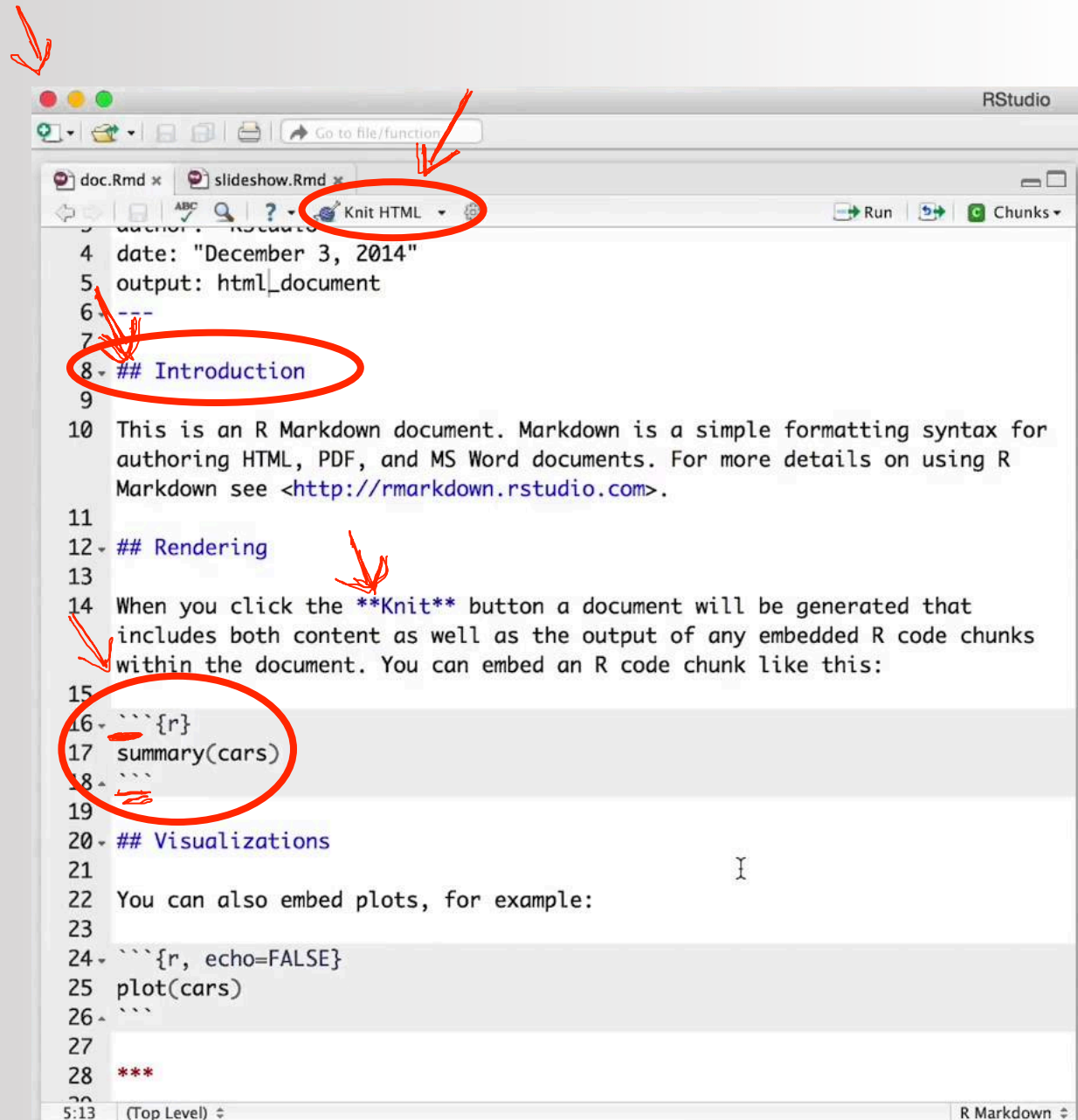


The screenshot shows the R help window for the `dim` function. The window title is "Help" and the browser address bar shows "R: Dimensions of an Object". The main content area displays the following information:

- dim {base}** (R Documentation)
- Dimensions of an Object**
- Description**: Retrieve or set the dimension of an object.
- Usage**:
`dim(x)`
`dim(x) <- value`
- Arguments**:
 - `x`: an R object, for example a matrix, array or data frame.
 - `value`: For the default method, either `NULL` or a numeric vector, which is coerced to integer (by truncation).
- Details**: The functions `dim` and `dim<-` are [internal generic primitive](#) functions. `dim` has a method for [data.frames](#), which returns the lengths of the `row.names` attribute of `x` and of `x` (as the numbers of rows and columns respectively).
- Value**: For an array (and hence in particular, for a matrix) `dim` retrieves the `dim` attribute of the object. It is `NULL` or a vector of mode [integer](#). The replacement method changes the "dim" attribute (provided the new value is compatible) and removes any "dimnames" and "names" attributes.

Writing code in R Markdown

- File → New File → R Markdown...
- Run selected line:
 - Cmd + Enter
- Run current chunk:
 - Cmd + shift + Enter
- Insert chunk:
 - Alt + Cmd + I



R Markdown Cheat Sheet

learn more at rmarkdown.rstudio.com

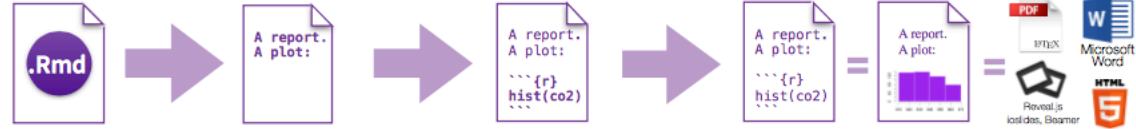
rmarkdown 0.2.50 Updated: 8/14



1. Workflow

R Markdown is a format for writing reproducible, dynamic reports with R. Use it to embed R code and results into slideshows, pdfs, html documents, Word files and more. To make a report:

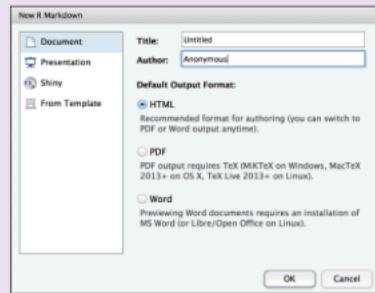
- i. **Open** - Open a file that uses the .Rmd extension.
- ii. **Write** - Write content with the easy to use R Markdown syntax
- iii. **Embed** - Embed R code that creates output to include in the report
- iv. **Render** - Replace R code with its output and transform the report into a slideshow, pdf, html or ms Word file.



2. Open File

Start by saving a text file with the extension .Rmd, or open an RStudio Rmd template

- In the menu bar, click **File ▶ New File ▶ R Markdown...**
- A window will open. Select the class of output you would like to make with your .Rmd file
- Select the specific type of output to make with the radio buttons (you can change this later)
- Click OK



3. Markdown

Next, write your report in plain text. Use markdown syntax to describe how to format text in the final report.

syntax

Plain text
End a line with two spaces to start a new paragraph.
italics and *italics*
bold and **bold**
superscript²
~~strikethrough~~
[\[link\]\(www.rstudio.com\)](http://www.rstudio.com)

Header 1
Header 2
Header 3
Header 4
Header 5
Header 6

endash: --
emdash: ---
ellipsis: ...
inline equation: $A = \pi r^2$
image:

horizontal rule (or slide break):

> block quote

* unordered list
* item 2
+ sub-item 1
+ sub-item 2

1. ordered list
2. item 2
+ sub-item 1
+ sub-item 2

Table Header	Second Header
Table Cell	Cell 2
Cell 3	Cell 4

becomes

Plain text
End a line with two spaces to start a new paragraph.
italics and *italics*
bold and **bold**
superscript²
~~strikethrough~~
[link](http://www.rstudio.com)

Header 1
Header 2
Header 3
Header 4
Header 5
Header 6

Header 4
Header 5
Header 6

endash: --
emdash: ---
ellipsis: ...
inline equation: $A = \pi r^2$



horizontal rule (or slide break):

block quote

- unordered list
- item 2
 - sub-item 1
 - sub-item 2

- ordered list
- item 2
 - sub-item 1
 - sub-item 2

Table Header	Second Header
Table Cell	Cell 2
Cell 3	Cell 4

4. Choose Output

Write a YAML header that explains what type of document to build from your R Markdown file.

YAML

A YAML header is a set of key: value pairs at the start of your file. Begin and end the header with a line of three dashes (---)

```
---  
title: "Untitled"  
author: "Anonymous"  
output: html_document  
---
```

This is the start of my report. The above is metadata saved in a YAML header.

The RStudio template writes the YAML header for you

The output value determines which type of file R will build from your .Rmd file (in Step 6)

output: html_document html file (web page)

output: pdf_document pdf document

output: word_document Microsoft Word .docx

output: beamer_presentation beamer slideshow (pdf)

output: ioslides_presentation ioslides slideshow (html)



Homework Due M2D5

- Peer review of methods

- before you leave today I will give you a copy of a classmate's methods homework to review

- review for clarity, completeness, and organization (see prompts on the wiki for full description)

- you must submit typed comments as a separate document using the "numbering method"

- turn your comments into the instructors M2D5, not to the peer you reviewed