

# A pipeline to analyse time-course gene expression data immediate

**Abstract** The phenotypic diversity of cells is governed by a complex equilibrium between their genetic identity and their environmental interactions: Understanding the dynamics of gene expression is a fundamental question of biology. However, analysing time-course transcriptomic data raises unique challenging statistical and computational questions, requiring the development of novel methods and software. Using as case study time-course transcriptomics data from mice exposed to different strains of influenza, this workflows provides a step-by-step tutorial of the methodology used to analyse time-course data: (1) normalization of the micro-array dataset; (2) differential expression analysis using functional data analysis; (3) clustering fo time-course data; (4) interpreting clusters with GO term and KEGG pathway enrichment analysis.

## Keywords

time-course gene expression data, clustering, differential expression, workflow

#TODO list

- Make clear that it applies to RNASeq and how to call the different functions to RNASeq
- Check that clustering can deal with count data.

```
library(devtools)
install_github("NelleV/moanin")

library(moanin)
```

EAP:

- I got an error in installation script for packages:

```
epurdom-0:2019timecourse-rnaseq-pipeline epurdom$ Rscript scripts/install.R
Loading required package: devtools
Loading required package: usethis
Loading required package: ggfortify
Loading required package: ggplot2
Loading required package: kableExtra
Error in contrib.url(repos, "source") :
  trying to use CRAN without setting a mirror
Calls: install.packages -> contrib.url
In addition: Warning message:
In library(package, lib.loc = lib.loc, character.only = TRUE, logical.return = TRUE, :
  there is no package called <e2><80><98>kableExtra<e2><80><99>
Execution halted
```

## Introduction

Gene expression studies provide simultaneous quantification of the level of mRNA from all genes in a sample. High-throughput studies of gene expression have a long history, starting with microarray technologies in the 1990s through to single-cell technologies. While many expression studies are designed to compare the gene expression between distinct groups, there is also a long history of time-course expression studies. Such studies compare the gene expression across time by measuring mRNA levels from samples collected at different timepoints<sup>1</sup>. Such time-course studies can vary from measuring a few distinct time points, to sampling ten to twenty time points. These longer time series are particularly interested in investigating development over time. More recently, a new variety of time course studies have come from single-cell sequencing experiments [1, 2, 3] which can sequencing single cells which are at different stages of development; in this case, the time point is the stage of the cell in the process of development – a value that is not known but estimated from the data as its “pseudo-time.”

While there are many methods that have been proposed for discrete aspects of time course data, the entire workflow for analysis of such data remains difficult, particularly for long, developmental time series. Most methods proposed for time course data are concerned with detecting genes that are changing over time (differential expression analysis), examples being edge [4], functional component analysis based models [5], time-course permutation tests [6], and multiple testing strategies to combine single time point differential expression analysis [7]. However, with long time course datasets, particularly in developmental systems, a massive number of genes will show *some* change. For example, in a study of mice lung tissues infected with influenza that we consider in this workflow, over 50% of genes are shown to be changing over time. The task in these settings is often not to detect changes in genes, but to categorize them into biologically interpretable patterns.

We present here a workflow for such an analysis that consists of 4 main parts (Figure 1):

- Quality control and normalization;
- Identification of genes that are differentially expressed;
- Clustering of genes into distinct temporal patterns;
- Biological interpretation of the clusters.

This workflow represents an integration of both novel implementations of previously established methods and new methodologies for the settings of developmental time series. It relies on several standard packages for analysing gene expression data, some specific for time-course data, others broadly used by the community. We provide the various steps of the workflow as functions in a R package called moanin.

<sup>1</sup>Because the collection of the mRNA is often destructive, samples at different time points are generally from different biological samples; longitudinal studies, for example tracking the same subject over time, are certainly possible, but not directly considered here.

**Figure 1.** Workflow for analyzing time-course datasets.

## Analysis of the dynamical response of mouse lung tissue to influenza

This workflow is illustrated using data from a micro-array time-course experiment, exposing mice to three different strains of influenza, and collecting lung tissue during 14 time-points after infection (0, 3, 6, 9, 12, 18, 24, 30, 36, 48, 60 hours, then 3, 5, and 7 days later) [8]. The three strains of influenza used in the study are (1) a low pathogenicity seasonal H1N1 influenza virus (A/Kawasaki/UTK4/2009 [H1N1]), a mildly pathogenic virus from the 2009 pandemic season (A/California/04/2009 [H1N1]), and a highly pathogenic H5N1 avian influenza virus (A/Vietnam/1203/2004 [H5N1]). Mice were injected with  $10^5$  PFU of each virus. An additional 42 mice were injected with a lower dose of the Vietnam avian influenza virus ( $10^3$  PFU).

### EAP: We need to add comment about microarray versus sequencing data

By combining gene expression time-course data with virus growth data, the authors show that the inflammatory response of lung tissue is gated until a threshold of the virus concentration is exceeded in the lung. Once this threshold is exceeded, a strong inflammatory and cytokin production occurs. This results provides evidence that the pathology response is non-linearly regulated by virus concentration.

### Quality control and normalization

The first steps of analysis of gene expression data is always to do normalization and quality control checks of the data. In what follows, we show an example of this for the influenza data using generic methods; these steps are not specific to time course data, but could be done for any gene expression analysis.

First let's load the data. The package `moanin` contains the normalized data and metadata of [8].

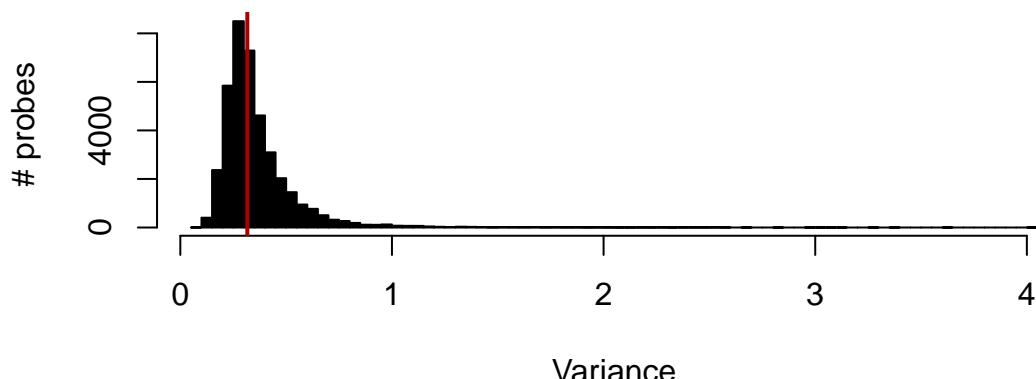
```
# Now load in the metadata
data(shoemaker2015)
meta = shoemaker2015$meta
data = shoemaker2015$data
```

### Exploratory analysis and quality control

Typically, two quality control and exploratory analysis steps are also performed before and after normalization: (1) low dimensionality embedding of the samples; (2) correlation plots between each samples. In both cases, we expect a strong biological signal, while replicate samples should be strongly clustered or correlated with one another.

Before performing any additional exploratory analysis, let us only keep highly variable genes: we keep for this step only the top 50% most variable genes.

```
variance_cutoff = 0.5
variance_per_genes = apply(data, 1, mad)
min_variance = quantile(variance_per_genes, c(variance_cutoff))
variance_filtered_data = data[variance_per_genes > min_variance,]
```

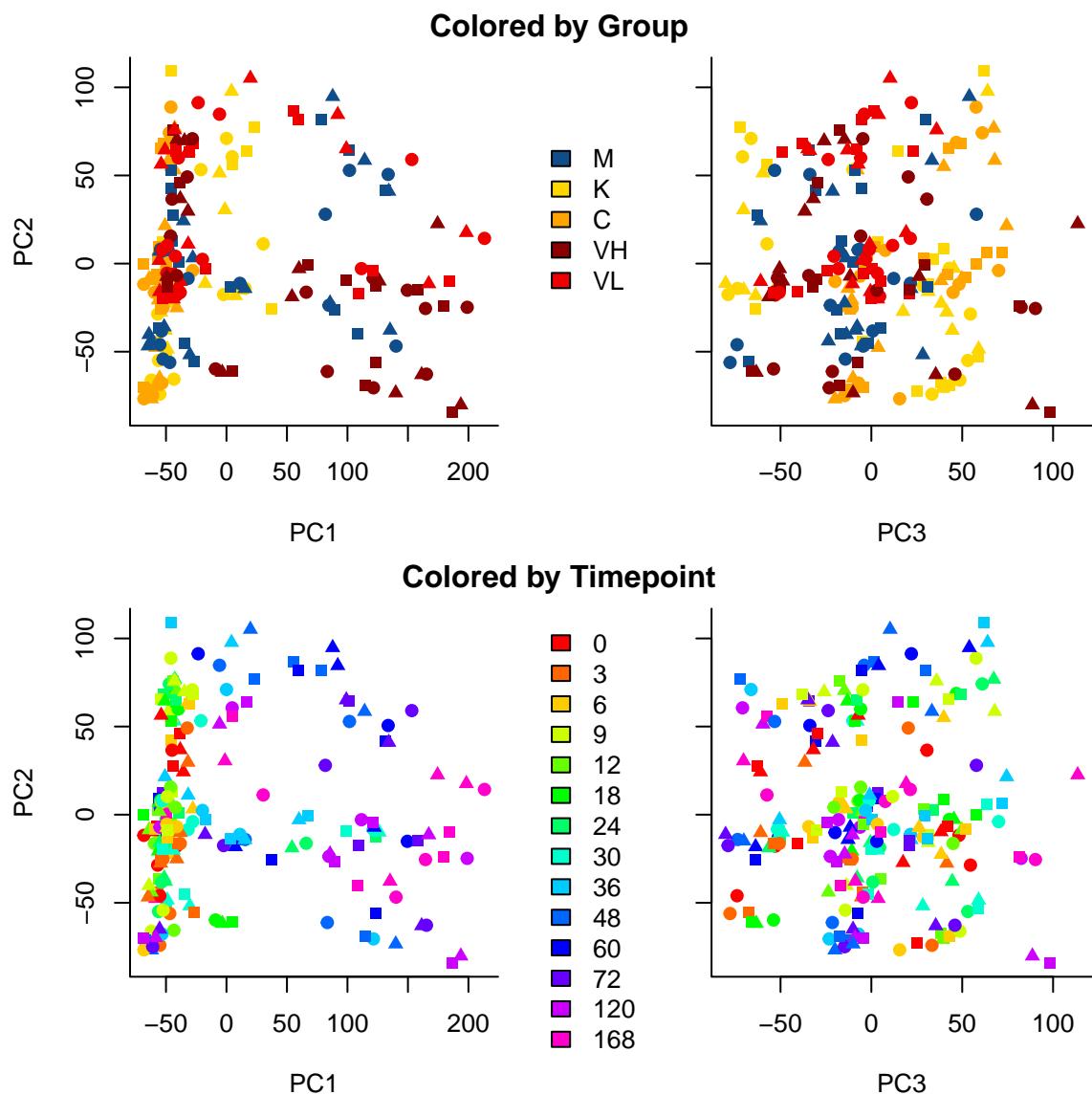


Let us first perform the PCA analysis. Here, we perform a PCA of rank 3 of the centered and scaled gene expression data.

```
pca_data = prcomp(t(variance_filtered_data), rank=3, center=TRUE, scale=TRUE)
percent_var = round(100 * attr(pca_data, "percentVar"))
```

We then plot the two first PC components, and color each sample by (1) its condition; (2) its sampling time. We use different plotting symbols for each replicate. We also plot the second and third components in the second column.

```
par(mfrow=c(2, 2))
par(mar=c(4.5, 4.5, 1.5, 1.5))
plot(
  pca_data$x[, c("PC1", "PC2")],
  col=ann_colors$Group[meta$Group],
  pch=ann_markers$Replicate[as.factor(meta$Replicate)], bty="l")
mtext(at=275, text="Colored by Group", line=0.5, font=2, adj=0.5)
legend(x=250, y=25, legend=names(ann_colors$Group),
       fill=ann_colors$Group, bty="n", xpd=NA, yjust=0.5)
par(mar=c(4.5, 5.5, 1.5, .1))
plot(
  pca_data$x[, c("PC3", "PC2")],
  col=ann_colors$Group[meta$Group],
  pch=ann_markers$Replicate[as.factor(meta$Replicate)],
  ylab="", bty="l", yaxt="n")
axis(2, labels=FALSE)
par(mar=c(4.5, 4.5, 1.5, 1.5))
plot(
  pca_data$x[, c("PC1", "PC2")],
  col=ann_colors$Timepoint[as.factor(meta$Timepoint)],
  pch=ann_markers$Replicate[as.factor(meta$Replicate)], bty="l")
mtext(at=275, text="Colored by Timepoint", line=0.5, font=2, adj=0.5)
legend(x=250, y=-15, legend=names(ann_colors$Timepoint),
       fill=ann_colors$Timepoint, bty="n", xpd=NA, yjust=0.5)
par(mar=c(4.5, 5.5, 1.5, .1))
plot(
  pca_data$x[, c("PC3", "PC2")],
  col=ann_colors$Timepoint[as.factor(meta$Timepoint)],
  pch=ann_markers$Replicate[as.factor(meta$Replicate)],
  ylab="", bty="l", yaxt="n")
axis(2, labels=FALSE)
```



Then, we plot the pearson correlation across each samples. We order the samples by their Group (treatment) and Timepoint (the time of sampling). Additionally, in this example, we order each treatment by strength of the pathogenicity of the treatment: Control, Kawasaki, California, low dose of Vietnam, then high dose of Vietnam.

```
# Reorder the conditions such that:
#   - Control is before any influenza treatment
#   - Each treatment is ordered from low to high pathogeny
meta$Group = factor(meta$Group, levels(meta$Group)[c(3, 2, 1, 5, 4)])

# Reorder genes on condition, time, and replicate
ord = order(
  meta$Group,
  meta$Timepoint,
  meta$Replicate)

variance_filtered_data = variance_filtered_data[, ord]
data_corr = cor(variance_filtered_data, method="pearson")
data_corr_meta = meta[ord, ]
data_corr_meta$Timepoint = as.factor(data_corr_meta$Timepoint)
# use aheatmap from the NMF package for a heatmap
aheatmap(
  data_corr,
  Colv=NA, Rowv=NA,
  annCol=data_corr_meta[, c("Group", "Timepoint")],
```

```
annRow=data_corr_meta[, c("Group", "Timepoint")],
annLegend=TRUE,
annColors=ann_colors,
main="Correlation plot")
```



We can already see interesting patterns emerging from the correlation plot. First, the cross-correlation amongst samples taking from the control mice is higher than the cross correlation amongst the rest of the treatments. Second, the influenza-infected mice mildly react until time point 36. Third, the less pathogenic the strain is, the closer the samples are to the control condition. Fourth, the Vietnam samples at time point 120 and 168 are the one that are the most different from control samples.

#### Differential expression analysis of time-course data

##### Approaches to DE analysis in time-course data

The next step in a gene expression analysis is typically to run a differential expression analysis, generally to find genes different between different conditions. For time-course data, there are two different approaches for determining differentially expressed genes,

- 1) Per-time point analysis, where we consider each time point a different condition and determine what genes are changing between time points, or between conditions at a single time-point.

- 2) Global analysis, where we consider the expression pattern globally over time, and consider what genes have either different patterns between conditions or a changing pattern (i.e. non-constant) over time. A common approach first step is to fit a spline model to each gene [4], and then use that spline model to test for different kinds of differential expression across time.

The per-time point analysis is using classical differential expression approaches, and is often the approach advocated when dealing with small time-course datasets, where there are only a few time points [9, 10, 11]. For long time-course datasets, however, a separate test for each time-point results in creating many different tests, for example one for every time point, the results of which are difficult to integrate. We find in practice that the global analysis simplifies analysis and of longer time courses data, with per-time point analysis reserved for particularly interesting comparisons of individual time-points.

We note that we could have time course data on either a single condition (to identify genes changing over time) or time course data on multiple conditions (such as the influenza dataset we are considering), which will alter slightly the types of questions we are interested in. In what follows, we will focus on the situation where we have multiple conditions. **EAP: Any reason one and not the other. Maybe should change this sentence**

Use with `moanin` provides functionality for performing both of these types of approaches, though our focus is on the global approach. In both situations, we first need to set up a object (a `splines_model` object) to hold the meta data, as well as information for fitting the spline model.

We start by creating the `splines_model` object using the `create_splines_model` function.

```
splines_model = create_splines_model(meta, degrees_of_freedom=6)
```

The `splines_model` object contains a number of metadata and options used throughout this analysis: the condition and timepoints of each samples, the formula object used or the basis or the degrees of freedom of the model. The metadata `data.frame` object should contain at least two columns: one named `Group`, containing the treatment effect, and a second one named `Timepoint` containing the timepoint information.

		Group	Replicate	Timepoint
GSM1557140	0	K	1	0
GSM1557141	1	K	2	0
GSM1557142	2	K	3	0
GSM1557143	3	K	1	12
GSM1557144	4	K	2	12
GSM1557145	5	K	3	12

If no formula is provided, it will default to the following: `formula = ~Group:ns(Timepoint, df=degrees_of_fredoom) + Group + 0.`

**EAP: General comment: you need a few more `print()` and `head()` of some objects to demonstrate what the output looks like**

### Weekly differential expression analysis

`moanin` provides a simple interface to perform a timepoint by timepoint differential expression analysis. This is traditionally done by the user defining the comparisons (called `contrasts` in linear models). Under the hood, simply calls `limma` [9] on the set of contrasts provided. By default, it expects RNA-Seq contact counts, and will estimate voom weights. We turn

Here, we show an example where we define our contrasts to be the difference between the control mouse (“M”) and the mouse infected with the high dose of the influenza strain A/Vietnam/1203/04 (H5N1) (“VL”) for each time point, but the function works with any form contrasts [9].

First, create the contrasts for all timepoints between the two groups of interest:

```
contrasts = create_timepoints_contrasts("M", "VL", splines_model)
```

This creates a vector of contrasts to be tested, one for each timepoint:

Contrasts
M.0-VL.0
M.3-VL.3
M.6-VL.6
M.9-VL.9
M.12-VL.12
M.18-VL.18
M.24-VL.24
M.30-VL.30
M.36-VL.36
M.48-VL.48
M.60-VL.60
M.72-VL.72
M.120-VL.120
M.168-VL.168

EAP: Not sure if kable is better than just printout?

Then run the differential expression analysis on all of those timepoints jointly using the function DE\_timepoints.

```
# Define contrast
weekly_de_analysis = DE_timepoints(
  data, splines_model, contrasts,
  use_voom_weights=FALSE)
```

EAP: Sort of strange to have argument splines\_model for function that doesn't use splines. Perhaps there is a generic name that could be used for the argument

This results in a table of the results, where each row corresponds to a gene and the columns correspond to the p-value (pval), log-fold change (lfc) and adjusted p-value (qval); the order of the genes in the table is the same as the input data. Here we show the results for the first two timepoints and the first ten genes:

	M.0-VL.0-pval	M.0-VL.0-qval	M.0-VL.0-lfc	M.3-VL.3-pval	M.3-VL.3-qval	M.3-VL.3-lfc
NM_009912	0.273	0.475	2.202	0.767	0.876	1.158
NM_008725	0.851	0.924	2.028	0.139	0.302	1.457
NM_007473	0.088	0.219	0.678	0.917	0.959	0.764
ENSMUST00000094955	0.008	0.037	1.186	0.935	0.969	-0.633
NM_001042489	0.419	0.621	1.212	0.761	0.872	0.599
NM_008159	0.123	0.278	1.102	0.117	0.268	0.737
NM_001013813	0.057	0.159	0.630	0.171	0.348	0.975
AK039774	0.015	0.059	0.637	0.062	0.171	0.971
NM_013782	0.875	0.938	0.594	0.000	0.001	1.180
NM_028622	0.767	0.876	1.648	0.946	0.974	-1.865

EAP: For the output of this, the column names are in format M.0-VL.0-pval, for example. This may be hard to parse algorithmically down the road because you are using the same character for the contrast (-) as the separator for the pval. So for example, if you wanted to split the contrast name from the pval or lfc part, it wouldn't be very easy. Maybe underscore \_ would be better separator?

We will repeat this, comparing each of the treatments to the control ("M") (code not printed here, as it is a replicate of the above, see accompanying Rmarkdown document).

Let's look at the distribution of genes found differentially expressed per week between control and each of the influenza strains.

EAP: Should timely\_barplot be a function of moanin?

NV: I'd rather not include these types of functions that are hard to test and clutter the namespace. In order to provide reasonable visualization, they also often need to be tweaked specifically to match the data (here, the ylim is set specifically to the data

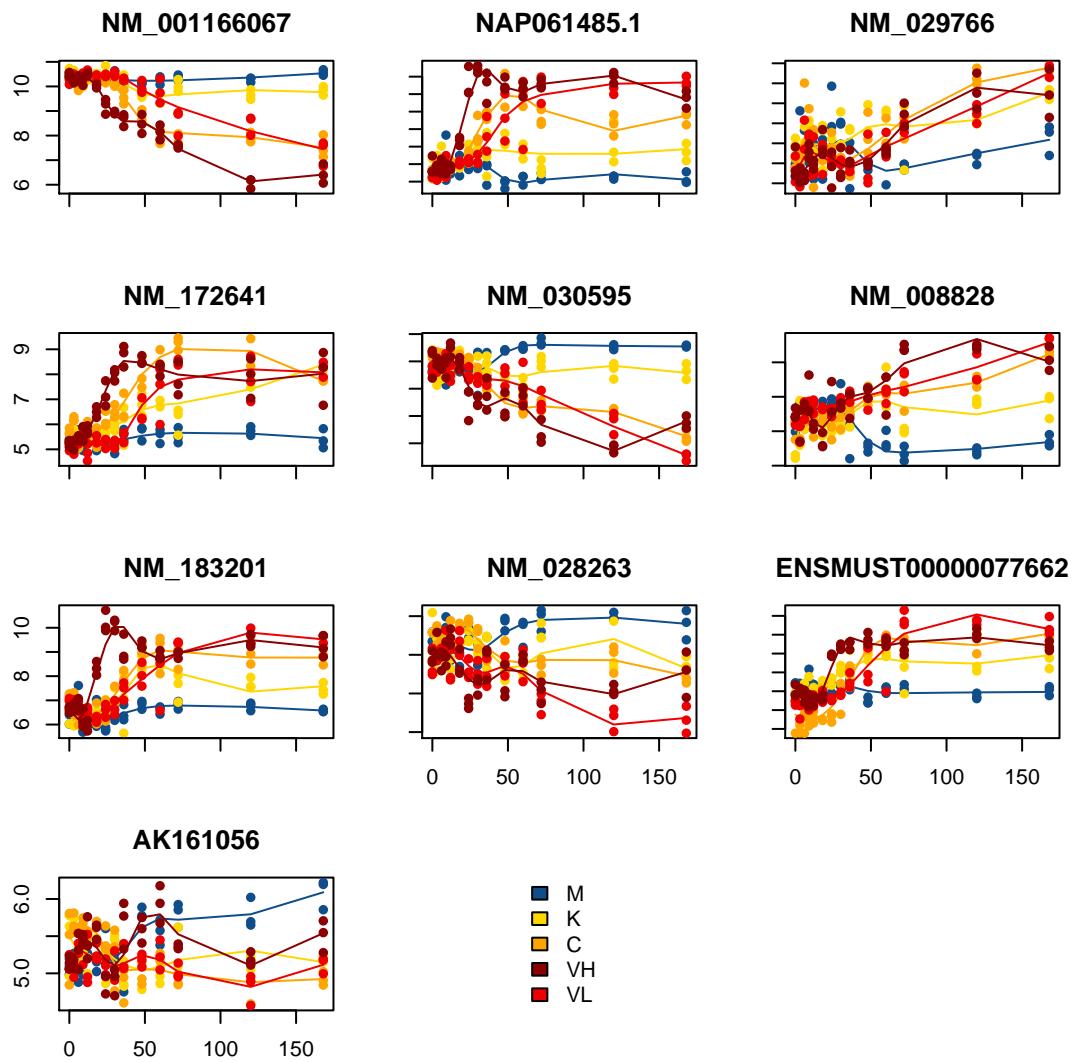


Such an analysis can demonstrate some general trends, with clearly more genes being differentially expressed at later time points, and the Vietnam high-dose showing perhaps an earlier onset than the low-dose.

However, the distribution of the number of genes found differentially expressed by considering each timepoint independently highlights the challenges of such approach. We can see that some timepoints have many less genes found significantly differentially expressed (e.g. timepoint 6H and 18H of the Kawasaki strain). While there may be biological differences at those time points for some genes, it seems unlikely that the large majority of genes differentially expressed at timepoint 3H stop being differentially expressed at 6H and then jump back to being differentially expressed at 9H. A more likely explanation is that there are some technical or biological artifact about the samples for 6H that is creating higher variation and thus less ability to detect significance.

Another difficulty with such an approach is making sense of the general temporal structure for any particular gene, as different genes will have different combinations of timepoints DE. For the comparison of the Kawasaki strain to the control, for example, there are `length(signifCombos)` genes found DE in some timepoints, and there are `length(tabCombos)` different combinations of timepoints for which they are DE. Some of these make sense, such as DE in timepoints 48H-168H (`tabCombos["48,60,72,120,168"]` genes), but many are very fragmentary. For example there are `tabCombos["48,60,120,168"]` genes which are DE in timepoints 48,60,120,168H, but *not* in the 72H. Many of these genes are likely to have not made the cutoff for significance in 72H, but don't show real differences in the overall trend between 48H-168H.

Here are plots of the first 10 such genes, where we can see that some might show some meaningful changes between 60 and 72H, but others clearly just have a single replicate that is different or increased variability at 72H.



As a summary, classic differential expression methods are appropriate for unordered treatments, but fail to make use of the temporal structure of the data.

#### Time-course differential expression analysis between two groups

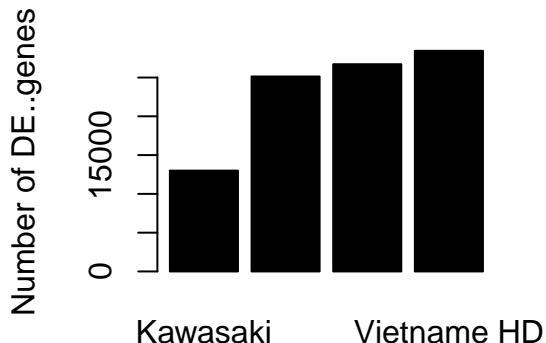
To leverage this temporal structure, Storey et al [4] proposed to model each gene in time-course micro-array with a splines function, and to use a log-ratio likelihood test to detect differentially expressed genes.

moanin extends this idea by providing functionality to compare time course data between different treatment conditions, using a similar mechanism of contrasts – only now the contrasts are differences between the estimated means.

```
# Differential expression analysis
timecourse_contrasts = c("M-K", "M-C", "M-VL", "M-VH")

# The function takes the data (data.frame or named matrix), the meta data
# (data.frame containing a timepoint and group column, the first corresponding
# <c2><a0> to the time-course information, the latter corresponding to the
# treatment).
pvalues = DE_timecourse(
  data, splines_model, timecourse_contrasts,
  use_voom_weights=FALSE)
pvalues = apply(pvalues, 2, p.adjust)
```

The number of genes found differentially expressed ranges from around 12000 to 29000 depending on the strain and dosage of influenza virus given to the mice. This corresponds to between 30% to 70% of the genes found differentially expressed in this time-course experiment.



The next step in a classical differential expression analysis is typically to assess the effect of the treatment by looking at the log fold change. Computing the log fold change on a time-course experiment is not trivial: one can be interested in the average log-fold change across time, or the cumulative log-fold change. Sometimes a gene can be over-expressed at the beginning of the time-course data, and then over-expressed at the end of the experiment. As a result, moanin provides a number of possible ways to compute the log fold change across the whole time-course.

First, `moanin` provides a simple interface to compute the log-fold change for each individual timepoints.

```
log_fold_change_timepoints = estimate_log_fold_change(
    data, splines_model, timecourse_contrasts, method="timely")
```

	M-K:1	M-K:7	M-K:11	M-K:14	M-K:2
NM_009912	-0.0061424	0.0224948	-0.0515947	-0.1143973	0.0614619
NM_008725	2.6547855	0.2734387	-0.6882925	0.4718807	-1.7463319
NM_007473	-0.1603623	0.1017655	0.5079573	0.0595773	0.4250185
ENSMUST00000094955	0.3505920	0.6463011	0.2564796	0.3599102	0.4177116
NM_001042489	0.0889750	0.2321381	-0.1891501	0.1786113	0.3320644
NM_008159	-0.3863770	0.0298395	-0.0862755	-0.3266351	-0.1893062

This matrix can then be used to visualize the log-fold change for each contrast per timepoint.

Sometimes, a single value per gene and per contrast is more useful. Here is a table of the possible ways to compute log-fold change values.

Name	Formula	Description
timely	$lfc(t)$	Function of time
sum	$\sum_t lfc(t)$	Sum of log fold change.
abs_sum	$\sum_t   lfc(t)  $	Always positive
max	$\max_t   lfc(t)  $	Always positive
min	$\min_t   lfc(t)  $	Always positive
epicon	\$\$	Captures overall strength of response and overall direction

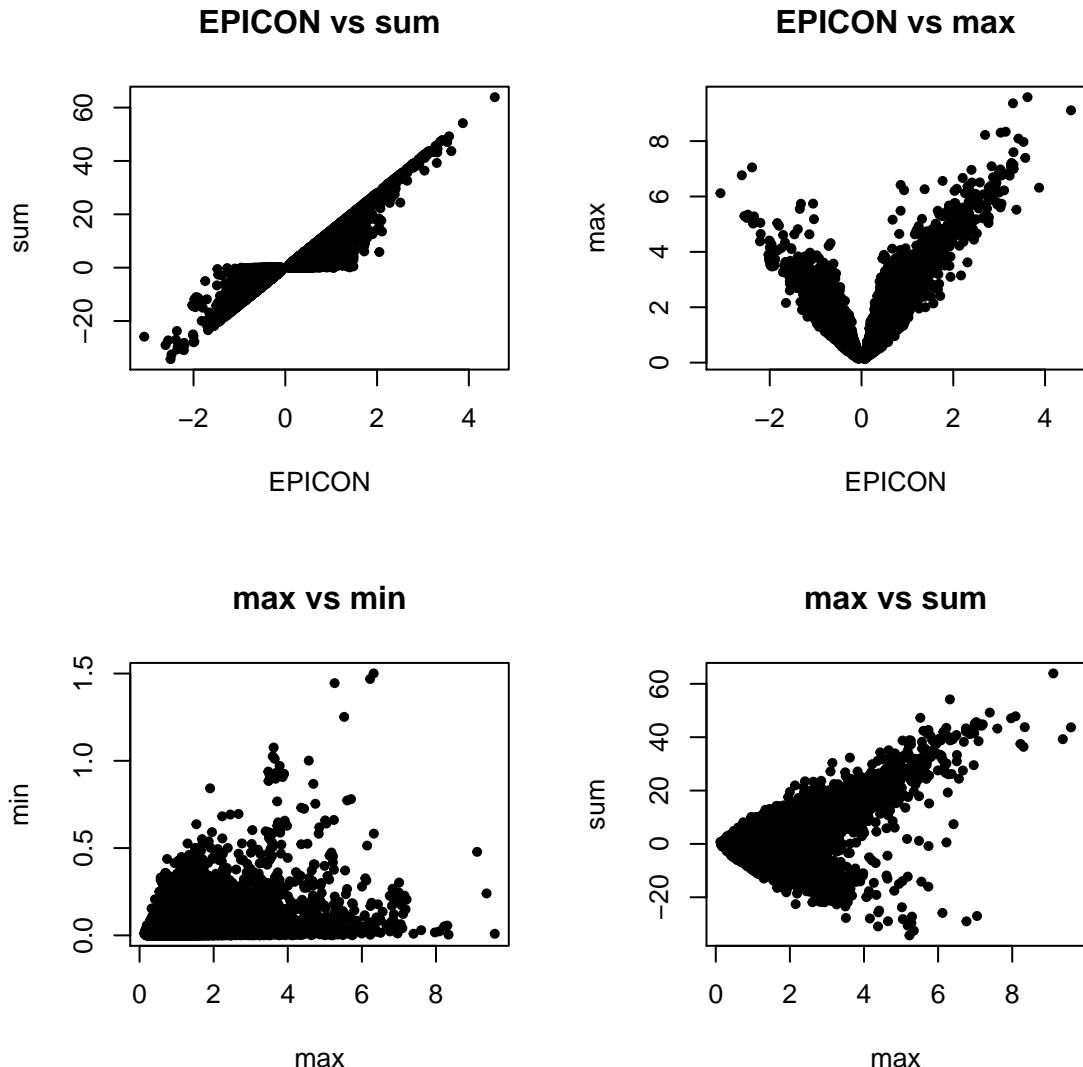
EAP: epicon is going to be weird for general use

```
log_fold_change_epicon = estimate_log_fold_change(
    data, splines_model, timecourse_contrasts, method="epicon")

log_fold_change_sum = estimate_log_fold_change(
    data, splines_model, timecourse_contrasts, method="sum")

log_fold_change_max = estimate_log_fold_change(
    data, splines_model, timecourse_contrasts, method="max")
```

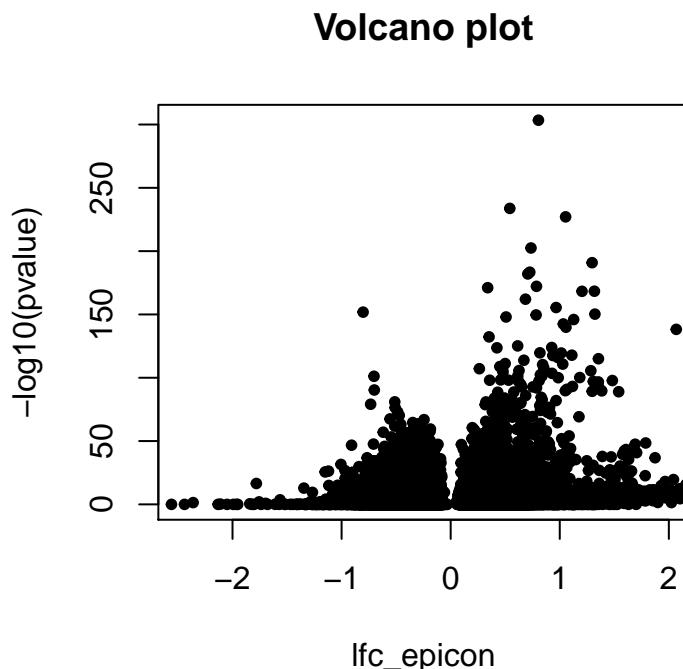
```
log_fold_change_min = estimate_log_fold_change(
    data, splines_model, timecourse_contrasts, method="min")
```



From the single measures of log-fold change and the p-value, we can now look at the volcano plot. Here is an example of a volcano plot for the comparison of the control to the Kawasaki strain, using the EPICON log fold change computation.

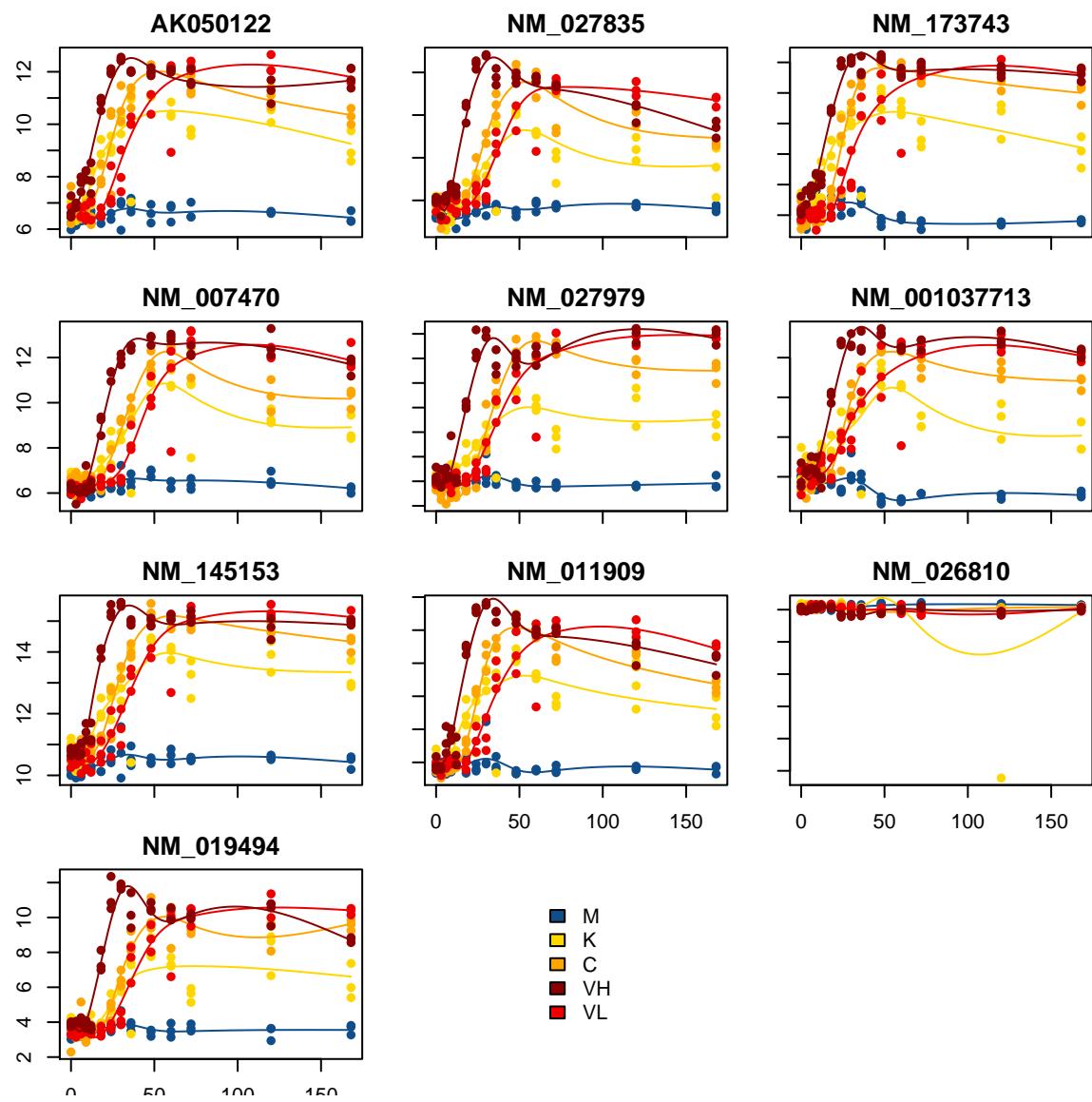
```
pvalue = qvalues[, "M-K"]
lfc_epicon = log_fold_change_epicon[, "M-K"]
names(lfc_epicon) = row.names(log_fold_change_epicon)

plot(lfc_epicon, -log10(pvalue), pch=20, main="Volcano plot",
      xlim=c(-2.5, 2))
```



As another sanity check, `moanin` provides a simple utility function to visualize gene time-course data. Here, we plot the 10 genes with the smallest p-values.

```
top_DE_genes_pval = names(sort(pvalue)[1:10])
plot_splines_data(data[top_DE_genes_pval, ], splines_model,
  colors=ann_colors$Group, smooth=TRUE,
  mar=c(1.5, 2.5, 2, 0.1))
```



And here we visualize the genes with the largest absolute EPICON log-fold change.

```
top_DE_genes_lfc = names(
  sort(abs(lfc_epicon),
  decreasing=TRUE)[1:10])
plot_splines_data(data[top_DE_genes_lfc, ], splines_model,
  colors=ann_colors$Group, smooth=TRUE,
  mar=c(1.5,2.5,2,0.1))
```



Thanks to those visualization, we can see that genes often follow similar patterns of expression, although on a different scale for each gene. We can leverage this observation to cluster the genes into groups of similar patterns of transcriptomic response.

### Clustering of time-course data

The very large number of genes found differentially expressed impair any interpretation one would attempt: with 70% of the genome found differentially expressed, all pathways are affected by the treatment. Hence the next step of the workflow to cluster gene expression according to their dynamical response to the treatment. Before clustering the genes, we first reduce the set of genes of interest to genes that are (1) significantly found differentially expressed; (2) “highly” differentially expressed. To do this, we first aggregate all p-values obtained during the time-course differential expresison step in a single p-value using Fisher’s method [12]. Then we select all the genes which have a Fisher adjusted p-value below 0.05 and a log fold change of at least two between at least one condition and one time-point. Reducing the set of genes on which to perform the clustering allows to estimate the centroids with more stability.

```
# Then rank by fisher's p-value and take max the number of genes of interest
# Filter out q-values for the pvalues table
fishers_pval = pvalues_fisher_method(pvalues)
fishers_qval = p.adjust(fishers_pval)

genes_to_keep = row.names(
  log_fold_change_max[
    (rowSums(log_fold_change_max > 2) > 0) &
```

```
(fishers_qval < 0.05), ])
# Keep the data corresponding to the genes of interest in another variable.
y = as.matrix(data[genes_to_keep, ])
```

After filtering, we are left with 3950 genes. We can then apply a clustering. As observed by looking at genes found differentially expressed, many genes share a similar gene expression pattern, but on different scale. We thus propose the following adaptation of k-means:

- **Splines estimation:** for each gene, fit the splines function with the basis of your choice.
- **Rescaling splines:** for each gene, rescale the estimated splines function such that the values are bounded between 0 and 1.
- **K-means:** apply k-means on the rescaled fitted splines to estimate the centroids.
- **Assign scores and labels to all genes:** then assign a score and a label to all gene based on a goodness-of-fit measure on the raw data. By default, the `splines_kmeans_score_and_label` function only labels the best 50% of genes.

The first three steps are performed jointly by the `splines_kmeans` function.

```
# First fit the kmeans clusters
kmeans_clusters = splines_kmeans(
  y, splines_model, n_clusters=8,
  random_seed=42,
  n_init=20)
```

We then use the `plot_splines_data` function to visualize the centroids obtained with the splines k-means model.

```
plot_splines_data(
  kmeans_clusters$centroids, splines_model,
  colors=ann_colors$Group,
  smooth=TRUE)
```

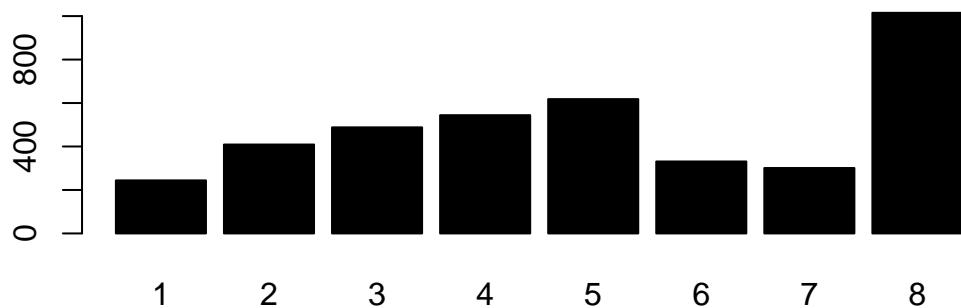
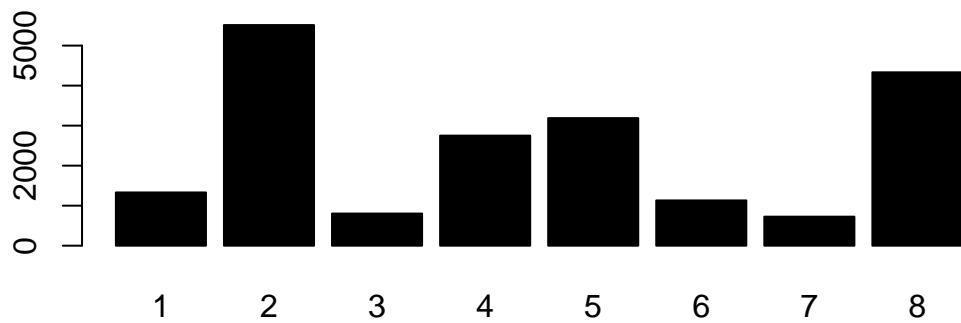


The scoring and labeling can be done via the `splines_kmeans_score_and_label` function.

```
#<c2><a0>Then assign scores and labels to all the data, using a goodness-of-fit
# scoring function.
scores_and_labels = splines_kmeans_score_and_label(
  data, kmeans_clusters)
labels = scores_and_labels$labels

# Let's keep only the list of genes that have a label.
labels = unlist(labels[!is.na(labels)])
```

Before performing the next steps, let us investigate in more detail the differences between the labels provided by the splines k-means model and the scoring and labeling step.

**splines k-means labeling****score and label**

The scoring and label step allows to label genes that were removed during the filtering step, yet are good matches to the centroids found during the clustering.

**Confusion matrix**

214	0	3	0	0	0	1	2	1
1	320	0	37	0	37	0	0	2
22	0	226	0	13	0	60	20	3
0	77	0	454	0	0	1	0	4
0	0	4	0	457	0	0	59	5
0	26	0	12	3	131	0	0	6
3	0	31	3	0	0	246	1	7
0	0	0	0	18	0	0	985	8
1	2	3	4	5	6	7	8	

**Looking at specific clusters in details.**

Now, let us look more in detail some specific clusters. Cluster 8 seems particularly interesting: it captures genes with strong differences between the different influenza treatments and the control, while the control remains relatively flat.

Heatmaps are useful to investigate the range of expression patterns for specific genes. Here, we are going to plot heatmaps of the normalized gene expression patterns and the rescaled gene expression patterns side by side.

First, select the genes of interest.

```
cluster_to_plot = 8
genes_to_plot = names(labels[labels == cluster_to_plot])

layout(matrix(c(1,2), nrow=1), widths=c(1.5, 2))

data_to_plot = data[genes_to_plot, ]
submeta = meta
ord = order(
  submeta$Group,
  submeta$Timepoint,
  submeta$Replicate)
submeta$Timepoint = as.factor(submeta$Timepoint)
```

```
data_to_plot = data_to_plot[, ord]
submeta = submeta[ord, ]

res = aheatmap(
  data_to_plot,
  Colv=NA,
  color="YlGnBu",
  annCol=submeta[, 
    c("Group", "Timepoint")],
  annLegend=FALSE,
  annColors=ann_colors,
  main=paste("Cluster", cluster_to_plot, "(raw)"),
  treeheight=0, legend=FALSE)

# Now use the results of the previous call to aheatmap to reorder the genes.
aheatmap(
  moanin:::rescale_values(data_to_plot)[res$rowInd, ],
  Colv=NA,
  Rowv=NA,
  annCol=submeta[, 
    c("Group", "Timepoint")],
  annLegend=TRUE,
  annColors=ann_colors,
  main=paste("Cluster", cluster_to_plot, "(rescaled)"),
  treeheight=0)
```



Those two heatmaps demonstrate that the clustering method successfully cluster genes that are on different scales, and yet share the same dynamical response to the treatments.

#### How to choose the number of clusters.

A common question that arises when performing clustering is how to choose the number of clusters. A choice for the number of clusters  $K$  depends on the goal. In this particular case, the end goal is not the clustering, but to facilitate interpretation of the differential expression analysis step. As a result, the number of clusters should not exceed the number of gene sets the user wants to interpret. This allows to set a maximum number of clusters. Let us assume here that this number is 20 clusters.

Once the maximum number of clusters is set, several strategies allow to identify the number of clusters:

- **Elbow method.** First introduced in 1953 by Thorndike [13], the elbow methods looks at the total with cluster sum of squares as a function of the number of clusters (WCSS). When adding clusters doesn't decrease the WCSS by a sufficient amount, one can consider stopping. This method thus provides visual aid to the user to choose the number of clusters, but often the "elbow" is hard to see on real data, where the number of clusters is not clearly defined.
- **Silhouette method.** Similarly to the Elbow method, the Silhouette method refers to a method of validation of consistency within clusters, and provides visual aid to choose the number of clusters.
- **Stability methods** Stability methods are more computationally intensive than any other method, as they rely on assessing the stability of the clustering for every  $k$  to a small randomization of the data. The user is then invited to choose the number of cluster based on a number of similarity measures.

First, let us run the clustering for all possible clusters of interest. We will, for each clustering experiment, conserve (1) with within cluster sum of squares; (2) the labels assigned to all genes.

```
all_possible_n_clusters = c(5, 9, 11, 15)
all_clustering = list()
wss_values = list()

i = 1
for(n_cluster in all_possible_n_clusters){
  clustering_results = splines_kmeans(
    y, splines_model,
    n_clusters=n_cluster, random_seed=42,
    n_init=10)
  wss_values[i] = sum(clustering_results$WCSS_per_cluster)
  all_clustering[[i]] = clustering_results$clusters
  i = i + 1
}
```

**Elbow method** The Elbow method to choose the number of clusters relies on visualization aid to choose the number of cluster. The method relies on plotting the within cluster sum of squares as a function of the number of clusters. At some point, the WCSS will stop dropping, giving an angle in the graph. The number of cluster is chosen at this "Elbow point."

```
plot(all_possible_n_clusters, wss_values,
      type="b", pch=19, frame=FALSE,
      xlab="Number of clusters K",
      ylab="Total within-clusters sum of squares")
```



**Average silhouette method** The silhouette value is a measure of how similar a data point is to its own cluster (cohesion) compared to other clusters (separation).

```
# function to compute average silhouette for k clusters
average_silhouette = function(labels, y) {
  silhouette_results = silhouette(unlist(labels[1]), dist(y))
  return(mean(silhouette_results[, 3]))
}

# extract the average silhouette
average_silhouette_values = list()
i = 1
for(i in 1:length(all_clustering)){
  clustering_results = all_clustering[i]
  average_silhouette_values[i] = average_silhouette(clustering_results, y)
  i = i + 1
}

plot(all_possible_n_clusters, average_silhouette_values,
  type="b", pch=19, frame=FALSE,
  xlab="Number of clusters K",
  ylab="Average Silhouettes")
```



**Looking at the stability of the clustering** On real data, the number of clusters is not only unknown but also ambiguous: it will depend on the desired clustering resolution of the user. Yet, in the case of biological data, stability and reproducibility of the results is necessary to ensure that the biological interpretation of the results hold when the data or the model is exposed to reasonable perturbations.

Methods that rely on the stability of the clustering results to choose  $k$  thus ensure that the biological interpretation of the clusters hold with perturbation to the data. In addition, simulation where the data is generated with a well defined  $k$  show that the clustering is more stable for the correct of number of the clusters.

Most methods method to find the number of clusters with stability measures only provide visual aids to guide the user. The first element often visualized is the consensus matrix: the consensus matrix is an  $n \times n$  matrix that stores the proportion of clustering in which two items are clustered together. A perfect consensus matrix ordered such as each elements that belong to the same cluster are adjacent to one another which show blocks along the diagonal close to 1.

To perform such analysis, the first step is run the clustering several times on a resampled dataset—either using bootstrap or subsampling.

Using the bootstrapping strategy:

```
n_genes = dim(y)[1]
indices = sample(1:dim(y)[1], n_genes, replace=TRUE)

bootstrapped_y = y[indices, ]
```

Using the subsampling strategy, keeping 80% of the genes:

```
subsample_proportion = 1
indices = sample(1:dim(y)[1], n_genes * subsample_proportion, replace=FALSE)
subsampled_y = y[indices, ]
```

Here we plot to stability matrix of the top 1000 genes for  $k = 5$  and  $k = 20$ .

```
stability_5 = read.table("results/stability_5.tsv", sep="\t")
consensus_matrix_stability_5 = consensus_matrix(stability_5,
                                                scale=FALSE)
aheatmap(consensus_matrix_stability_5[1:1000, 1:1000], Rowv=FALSE,
         Colv=FALSE,
         treeheight=0)
```



```
stability_20 = read.table("results/stability_20.tsv", sep="\t")
consensus_matrix_stability_20 = consensus_matrix(stability_20,
                                                 scale=FALSE)
aheatmap(consensus_matrix_stability_20[1:1000, 1:1000], Rowv=FALSE,
         Colv=FALSE,
         treeheight=0)
```



**The model explorer strategy** The model explorer algorithm [14] proposes to estimate the number of clusters exploiting the observation that if the number of clusters is correct, the clustering results are stable to bootstrapping. The distribution of similarities between bootstrapped results for each  $k$  can thus be compared for different values of  $k$  and guide the user in the choice of number of clusters.

The model explorer strategy works as follows. First, choose a similarity measure between two partitions or clusters  $S(C_1, C_2)$ . Examples are the normalized mutual information or Fowlkes-Mallows. Then perform  $n$  bootstrap experiments to estimate the cluster centroids, followed by a step of assigning a label to all data points. Finally, compute the pairwise similarity measure between all bootstrapped partition, and plot the cumulative density of the obtained scores.

```
plot_model_explorer(all_labels)
```



From this plot, we can deduce that  $k = 5$  is more stable than  $k = 3$  and  $k = 4$ , but not as stable as  $k = 2$ . The model explorer strategy, in addition to visualizing the diversity of the centroids, can thus help assessing an adequate number of clusters.

Now, replot the same model explorer, but only for the clustering experiments  $k = 6, k = 7, k = 8, k = 9$  and  $k = 10$  so that we can see more clearly the stability measures in that range.

```
clusters = c("C6", "C7", "C8", "C9", "C10")
selected_labels = all_labels[clusters]
plot_model_explorer(selected_labels)
```



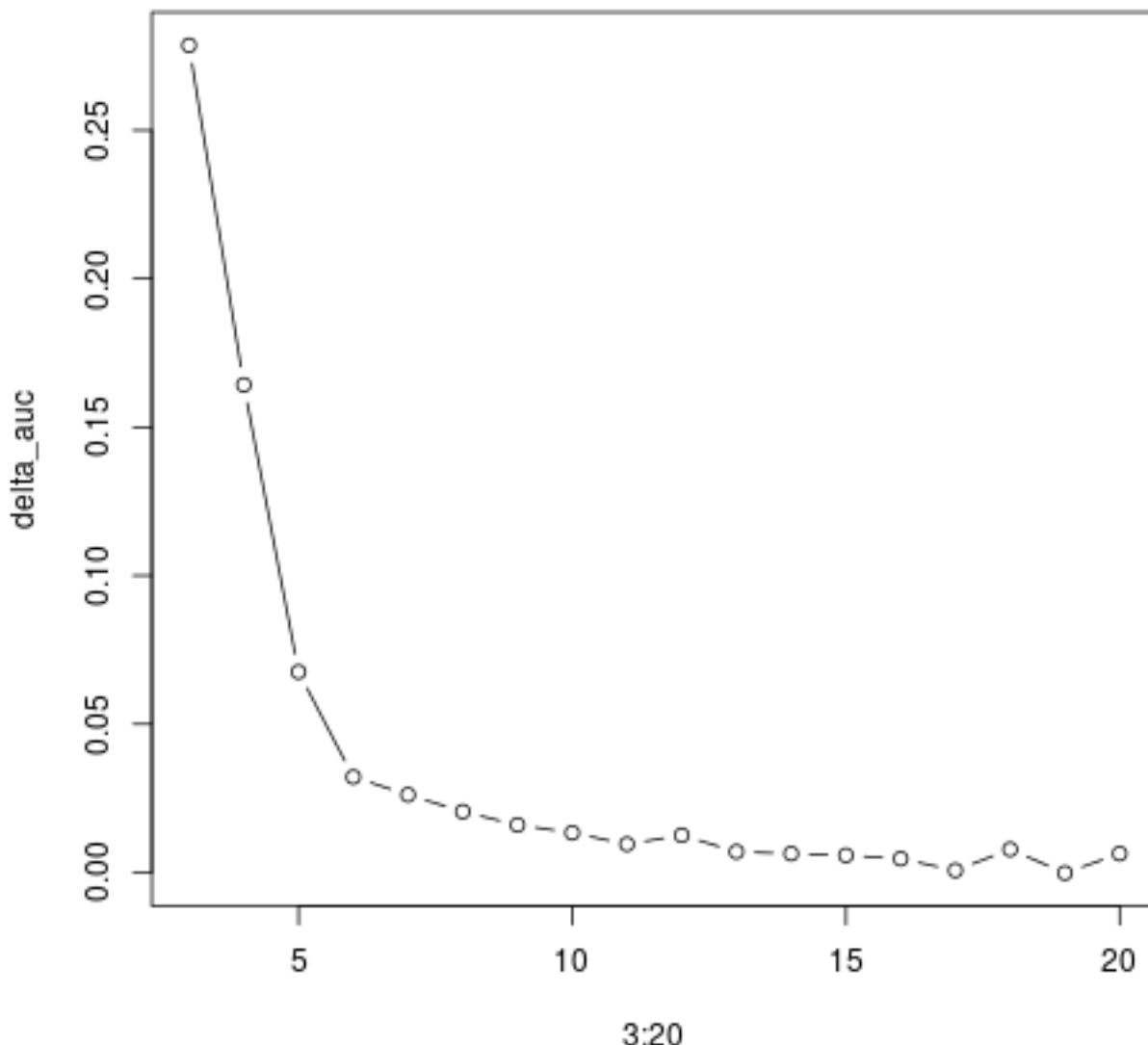
**Consensus clustering as a way to find  $k$**  The consensus clustering [15] relies on a similar idea but instead of looking at the cumulative density of similarity measures of bootstrapped clustering, the authors suggests plotting the cumulative density of elements of the consensus matrix.

```
plot_cdf_consensus(all_labels)
```



The stability of the clustering based on the consensus matrix can then be measured via a single number by looking at the area under the curve: the more stable the clustering, the closer to 0 or 1 will be the entries of consensus matrix. The consensus clustering strategy thus suggest at looking at either the AUC as a function of the number of the clusters or the “improvement” in the AUC as a function of the number of cluster.





The consensus clustering method suggest that the most stable is  $k = 2$ , which separates over-expressed genes from under-expressed genes. While it is indeed a very stable clustering, it does not capture the range of gene expression patterns present in the data. This shows the limitation of such method on real data, where the number of clusters is not clearly defined.

#### Downstream analysis of clusters.

Once good clusters are obtained, the next step is to leverage the clustering to ease interpretation. Classic enrichment analysis step can then be performed in the gene set defined in each cluster: KEGG pathway enrichment analysis, GO term enrichment analysis, motif enrichment analysis, etc.

First, let us clean up the gene obtained and only select the genes we are going to use in the enrichment analysis. One can either use the whole set of genes, only the set of differentially expressed genes in each cluster, or a subset of genes that fit well to a cluster (based on some criterion).

#### Finding enriched pathways using biomaRt and KEGGprofile

Let us first tackle the case of pathway enrichment analysis. We will leverage the packages `biomaRt` [16] and `KEGGprofile` [17] for this step. `KEGGprofile` is a package that easily allows to perform pathway enrichment analysis on a set of genes labeled with the ensembl annotation. We thus need to convert the gene

names into the appropriate format. This is where `biomaRt` comes in handy: it enables easy conversion from one gene annotation to another. Here, we will use it to convert the gene names from the Refseq annotation to the ensembl one.

```
ensembl = useMart("ensembl")
ensembl = useDataset("mmusculus_gene_ensembl", mart=ensembl)

cluster = 8
labels = unlist(labels)
gene_names = names(labels)
genes = gene_names[labels == cluster]

# convert gene names
genes = getBM(attributes=c("ensembl_gene_id", "entrezgene_id"),
              filters="refseq_mrna", values=genes,
              mart=ensembl)[["entrezgene_id"]]
genes = as.vector(unlist(genes))
pathways = find_enriched_pathway(
  genes, species="mmu",
  download_latest=TRUE)$stastic
```

	Pathway	Percentage	Adj. p-value
05169	Epstein-Barr virus infection	37	0
04060	Cytokine-cytokine receptor interaction	31	0
04668	TNF signaling pathway	45	0
05164	Influenza A	38	0
04621	NOD-like receptor signaling pathway	34	0
05162	Measles	37	0
04630	JAK-STAT signaling pathway	33	0
05160	Hepatitis C	34	0
04064	NF-kappa B signaling pathway	39	0
04620	Toll-like receptor signaling pathway	40	0

### Finding enriched GO terms

To find GO terms, we use `biomaRt` to find the mapping between GO terms and gene mapping. The GO enrichment library `topGO` [18] expects the GO term to gene mapping to be a list where each item is a mapping between a gene name and a GO term ID vector.

```
$NM_199153
[1] "GO:0016020" "GO:0016021" "GO:0007186" "GO:0004930" "GO:0007165"
[6] "GO:0050896" "GO:0050909"

$NM_201361
[1] "GO:0016020" "GO:0016021" "GO:0003674" "GO:0008150" "GO:0005794"
[6] "GO:0005829" "GO:0005737" "GO:0005856" "GO:0005874" "GO:0005739"
[11] "GO:0005819" "GO:0000922" "GO:0072686"
```

`biomaRt` queries results a matrix with two named columns of gene names and GO term ID.

```
genes = getBM(attributes=c("go_id", "refseq_mrna"),
              values=gene_names,
              filters="refseq_mrna",
              mart=ensembl)

# Create gene to GO id mapping
gene_id_go_mapping = create_go_term_mapping(genes)
```

Once the gene ID to GO mapping list is created, `moanin` provides a simple interface to `topGO` to fetch enriched GO terms. Here, we show an example of running a GOrterm enrichment on the “Biological process” ontology (BP).

```
assignments = labels == cluster

go_terms_enriched = find_enriched_go_terms(
    assignments,
    gene_id_go_mapping, ontology="BP")
```

	GO ID	Description	Annotated	Significant	Expected	P-value	adj. p-value
5	GO:0003002	regionalization	172	153	134.41	0.00026	0.2629
6	GO:0048562	embryonic organ morpho- genesis	152	135	118.78	0.00045	0.3323
7	GO:0010970	transport along mi- crotubule	82	75	64.08	0.00046	0.3323
8	GO:0006805	xenobiotic metabolic process	49	47	38.29	0.00059	0.3729
9	GO:0060173	limb develop- ment	92	84	71.89	0.00068	0.3820
10	GO:0019395	fatty acid oxidation	53	51	41.42	0.00134	0.6194
11	GO:0055114	oxidation- reduction process	536	461	418.84	0.00139	0.6194
12	GO:0060070	canonical Wnt signaling pathway	149	131	116.43	0.00147	0.6194
13	GO:0090596	sensory organ morpho- genesis	134	118	104.71	0.00219	0.8517
14	GO:0060411	cardiac septum morpho- genesis	33	32	25.79	0.00295	1.0000
15	GO:0010811	positive regulation of cell- substrate ad...	81	73	63.30	0.00384	1.0000
16	GO:0042738	exogenous drug catabolic process	22	22	17.19	0.00437	1.0000
17	GO:1901381	positive regulation of potassium ion tra...	22	22	17.19	0.00437	1.0000
18	GO:0007368	determination of left/right symmetry	66	60	51.57	0.00536	1.0000
19	GO:0060078	regulation of postsynaptic mem- brane pote...	66	60	51.57	0.00536	1.0000

## Session information

```

sessionInfo()

## R version 3.6.0 (2019-04-26)
## Platform: x86_64-pc-linux-gnu (64-bit)
## Running under: Ubuntu 19.04
##
## Matrix products: default
## BLAS:    /usr/lib/x86_64-linux-gnublas/libblas.so.3.8.0
## LAPACK:  /usr/lib/x86_64-linux-gnulapack/liblapack.so.3.8.0
##
## locale:
## [1] C
##
## attached base packages:
## [1] stats4     parallel   splines    stats      graphics   grDevices utils
## [8] datasets   methods    base
##
## other attached packages:
## [1] RColorBrewer_1.1-2 moanin_0.0.0      topGO_2.32.0
## [4] SparseM_1.77    GO.db_3.6.0       AnnotationDbi_1.42.1
## [7] IRanges_2.14.12 S4Vectors_0.18.3  graph_1.58.2
## [10] viridis_0.5.1   viridisLite_0.3.0 KEGGprofile_1.22.0
## [13] RCurl_1.95-4.12 bitops_1.0-6      NMF_0.21.0
## [16] Biobase_2.40.0  BiocGenerics_0.26.0 cluster_2.0.9
## [19] rngtools_1.3.1.1 pkgmaker_0.27    registry_0.5-1
## [22] ggplot2_3.1.1   biomaRt_2.36.1   BiocStyle_2.8.2
## [25] kableExtra_1.1.0 knitr_1.22      limma_3.36.5
## [28] usethis_1.5.0   devtools_2.0.2   rmarkdown_1.12
##
## loaded via a namespace (and not attached):
## [1] colorspace_1.4-1      rprojroot_1.3-2
## [3] XVector_0.20.0        fs_1.3.1
## [5] rstudioapi_0.10       remotes_2.0.4
## [7] bit64_0.9-7          ClusterR_1.1.9
## [9] KEGG.db_3.2.3         xml2_1.2.0
## [11] codetools_0.2-16      doParallel_1.0.14
## [13] pkgload_1.0.2         ade4_1.7-13
## [15] gridBase_0.4-7        png_0.1-7
## [17] FD_1.0-12            readr_1.3.1
## [19] compiler_3.6.0        httr_1.4.0
## [21] backports_1.1.4      Matrix_1.2-17
## [23] assertthat_0.2.1      lazyeval_0.2.2
## [25] cli_1.1.0             htmltools_0.3.6
## [27] prettyunits_1.0.2     tools_3.6.0
## [29] gmp_0.5-13.5          gtable_0.3.0
## [31] glue_1.3.1            reshape2_1.4.3
## [33] dplyr_0.8.0.1         BiocWorkflowTools_1.6.2
## [35] Rcpp_1.0.1             Biostrings_2.48.0
## [37] NMI_2.0                ape_5.3
## [39] nlme_3.1-139           iterators_1.0.10
## [41] xfun_0.6               stringr_1.4.0
## [43] ps_1.3.0               testthat_2.1.1
## [45] rvest_0.3.3            gtools_3.8.1
## [47] XML_3.98-1.19          zlibbioc_1.26.0
## [49] MASS_7.3-51.4           scales_1.0.0
## [51] hms_0.4.2              curl_3.3
## [53] yaml_2.2.0              memoise_1.1.0
## [55] gridExtra_2.3           TeachingDemos_2.10
## [57] stringi_1.4.3           RSQLite_2.1.1
## [59] desc_1.2.0               foreach_1.4.4
## [61] permute_0.9-5           pkgbuild_1.0.3
## [63] bibtex_0.4.2             geometry_0.4.1
## [65] rlang_0.4.0              pkgconfig_2.0.2

```

```

## [67] matrixStats_0.54.0      evaluate_0.13
## [69] lattice_0.20-38        purrr_0.3.2
## [71] bit_1.1-14            processx_3.3.1
## [73] tidyselect_0.2.5       plyr_1.8.4
## [75] magrittr_1.5           bookdown_0.9
## [77] R6_2.4.0               DBI_1.0.0
## [79] mgcv_1.8-28           pillar_1.3.1
## [81] withr_2.1.2            abind_1.4-5
## [83] KEGGREST_1.20.2        tibble_2.1.1
## [85] crayon_1.3.4           progress_1.2.0
## [87] grid_3.6.0              blob_1.1.1
## [89] callr_3.2.0            git2r_0.25.2
## [91] vegan_2.5-4             digest_0.6.18
## [93] webshot_0.5.1          xtable_1.8-4
## [95] munsell_0.5.0           magic_1.5-9
## [97] sessioninfo_1.1.1

```

## Conclusion

This workflow provides a tutorial for the analysis of time-course gene expression data in R, illustrated through the analysis of mice lung tissue exposed to different influenza strain. It covers four main steps; (1) quality control and normalization; (2) differential expression analysis; (3) clustering of time-course gene expression data; (4) downstream analysis of clusters.

## Software and data availability

The source code for this workflow can be found at <https://github.com/NelleV/2019timecourse-rnaseq-pipeline>. Data used in this workflow are available from NCBI GEO, accession GSE95601.

## Author contributions

NV and EP wrote the workflow.

## Competing interests

The authors declare that they have no competing interests.

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*I confirm that the funders had no role in study design, data collection and analysis, decision to publish, or preparation of the manuscript.*

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