

# R implementation

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## 1 Loaded functions:

```
#source("/media/Data/Dropbox/humanR/01funcs.R")
rm(list=ls())
#setwd("/media/Data/Dropbox/humanR/PD/")
#setwd("~/Dropbox/humanR/PD/")
###load("PD.Rdata", .GlobalEnv)
#lsos(pat="")
```

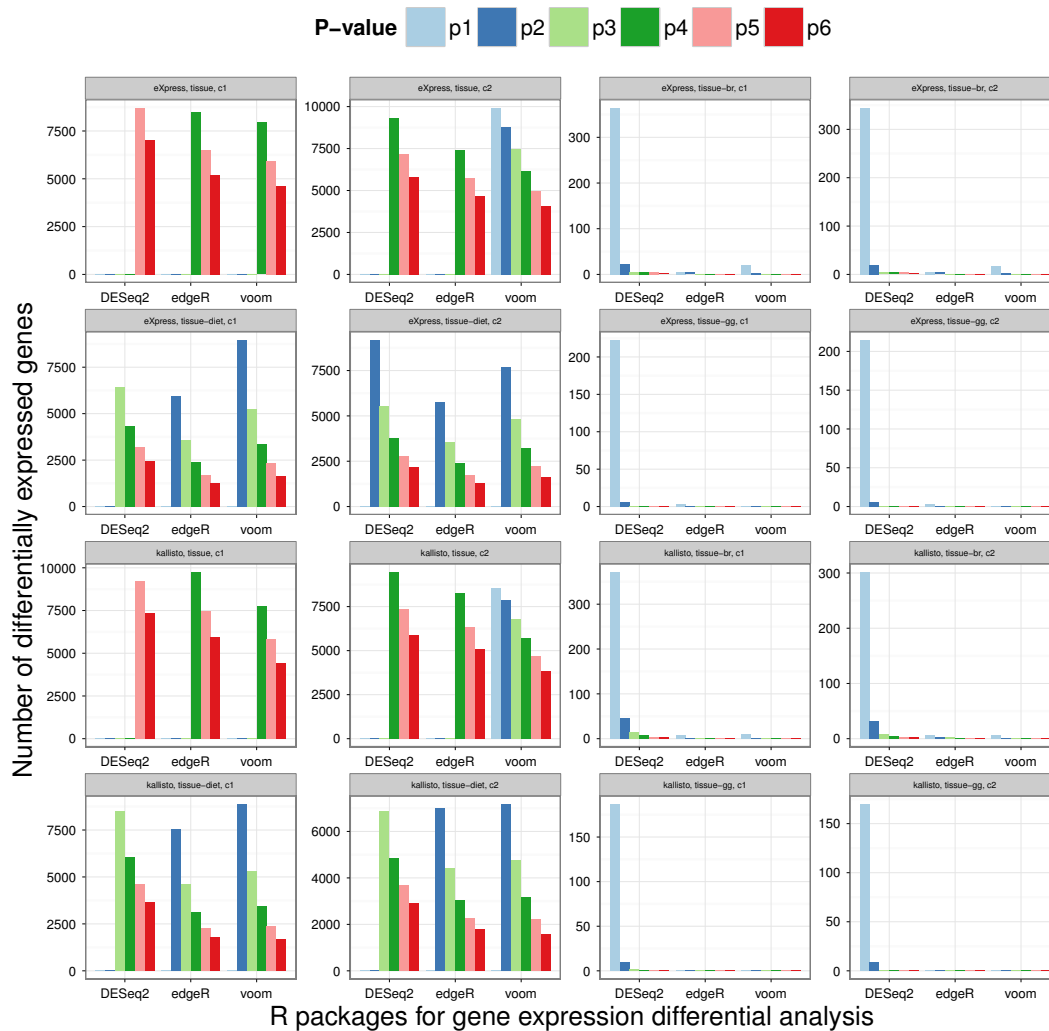
## 2 Load packages.

```
pkgs <- c('xlsx', 'caret', 'leaps', 'glmnet', 'lattice',
          'latticeExtra', 'dplyr', 'tidyr', 'grid')
lapply(pkgs, require, character.only = TRUE)
```

## 3 1 Differentially expressed genes

4 Differentially expressed genes are counted from mapping **both gills and ganglia** sequenced samples to  
5 reference transcriptome built from all samples.

```
read.table("./data/summary.raw.all.txt") %>%
  ggplot(aes(
    x = V1,
    y = V8,
    fill = V6)) +
  theme_bw() +
  geom_bar(stat = "identity",
    position = "dodge") +
  facet_wrap(~ V4 + V5 + V7,
    ncol = 4,
    scales = "free") +
  scale_fill_brewer(type = "qual", palette = "Paired",
    name = "P-value") +
  labs(x = "R packages for gene expression differential analysis",
    y = "Number of differentially expressed genes") +
  theme(legend.position = "top",
    axis.text.x = element_text(vjust = .5, size = 6),
    axis.text.y = element_text(vjust = .5, size = 6),
    strip.text = element_text(size = 4),
    axis.ticks = element_line(size = .2),
    axis.ticks.length = unit(.05, "cm"))
```



6

7 Differentially expressed genes are counted from mapping **gills** sequenced samples to reference tran-

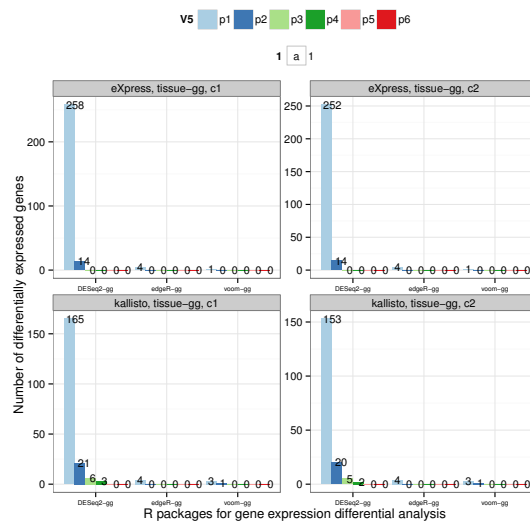
8 scriptome built from all samples.

```
read.table("./data/summary.gg.txt") %>%
```

```

ggplot(aes(
  x = V2,
  y = V8,
  fill = V5)) +
  geom_bar(stat = "identity",
    position = "dodge") +
  geom_text(aes(x = V2,
    y = V8,
    ymax = V8,
    label = V8,
    size = 1,
    hjust = 0),
    position = position_dodge(width = 1)) +
  facet_wrap(~ V3 + V4 + V6,
    ncol = 2,
    scales = "free") +
  scale_fill_brewer(type = "qual", palette = "Paired") +
  theme_bw() +
  theme(legend.position = "top",
    axis.text.x = element_text(vjust = .5,
      size = 6)) +
  labs(x = "R packages for gene expression differential analysis",
    y = "Number of differentially expressed genes")

```



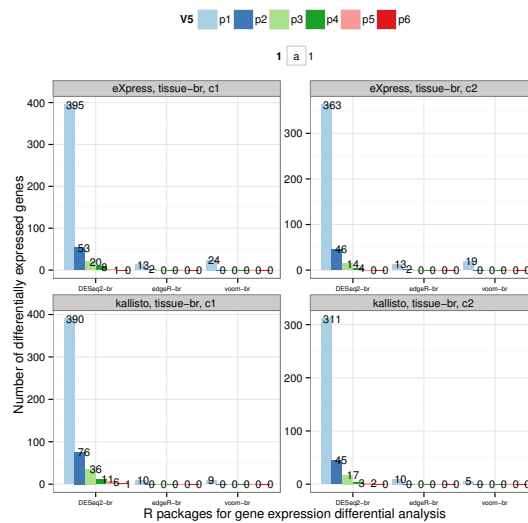
9  
10 Differentially expressed genes are counted from mapping **ganglia** sequenced samples to reference tran-  
11 scriptome built from all samples.

```
read.table("./data/summary.br.txt") %>%
```

```

ggplot(aes(
  x = V2,
  y = V8,
  fill = V5)) +
  geom_bar(stat = "identity",
    position = "dodge") +
  geom_text(aes(x = V2,
    y = V8,
    ymax = V8,
    label = V8,
    size = 1,
    hjust = 0),
    position = position_dodge(width = 1)) +
  facet_wrap(~ V3 + V4 + V6,
    ncol = 2,
    scales = "free") +
  scale_fill_brewer(type = "qual", palette = "Paired") +
  theme_bw() +
  theme(legend.position = "top",
    axis.text.x = element_text(vjust = .5,
      size = 6)) +
  labs(x = "R packages for gene expression differential analysis",
    y = "Number of differentially expressed genes")

```



**1.1 Increasing DEG by changing the trimming rates of raw reads**  
 Getting gene expression by mapping the original raw reads **without trimming** to the gills de novo transcriptome.

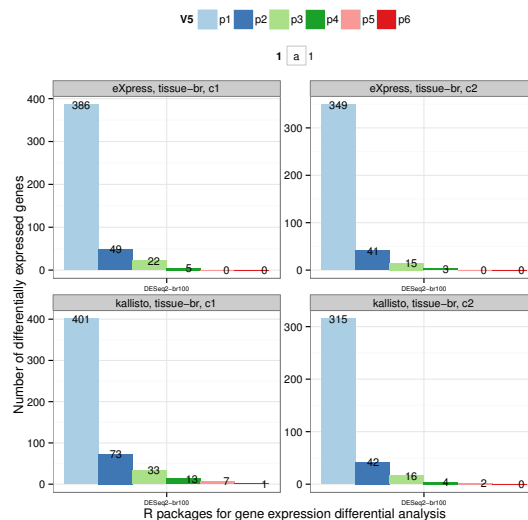
† De novo assembly was carried out with trimmed reads though

```
read.table("./data/summary.br_35078.txt") %>%
```

```

ggplot(aes(
  x = V2,
  y = V8,
  fill = V5)) +
  geom_bar(stat = "identity",
    position = "dodge") +
  geom_text(aes(x = V2,
    y = V8,
    ymax = V8,
    label = V8,
    size = 1,
    hjust = 0),
    position = position_dodge(width = 1)) +
  facet_wrap(~ V3 + V4 + V6,
    ncol = 2,
    scales = "free") +
  scale_fill_brewer(type = "qual", palette = "Paired") +
  theme_bw() +
  theme(legend.position = "top",
    axis.text.x = element_text(vjust = .5,
      size = 6)) +
  labs(x = "R packages for gene expression differential analysis",
    y = "Number of differentially expressed genes")

```



## 1.2 Increasing DEGs by changing the normalization strategy: Fast abundance quantification *kallisto*

The below graph shows the number of differentially expressed genes when raw reads were normalized separately for each biological sample.

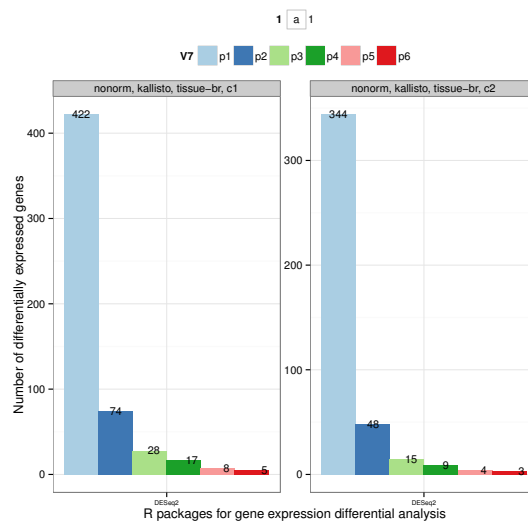
```
read.table("./data/summary.br.nonorm.txt") %>%
```

† All the analyses before were done on normalized reads by grouping all biological samples together

```

ggplot(aes(
  x = V1,
  y = V10,
  fill = V7)) +
  geom_bar(stat = "identity",
    position = "dodge") +
  geom_text(aes(x = V1,
    y = V10,
    ymax = V10,
    label = V10,
    size = 1,
    hjust = 0),
    position = position_dodge(width = 1)) +
  facet_wrap(~ V4 + V5 + V6 + V8,
    ncol = 2,
    scales = "free") +
  scale_fill_brewer(type = "qual", palette = 'Paired') +
  theme_bw() +
  theme(legend.position = "top",
    axis.text.x = element_text(vjust = .5,
      size = 6)) +
  labs(x = "R packages for gene expression differential analysis",
    y = "Number of differentially expressed genes")

```



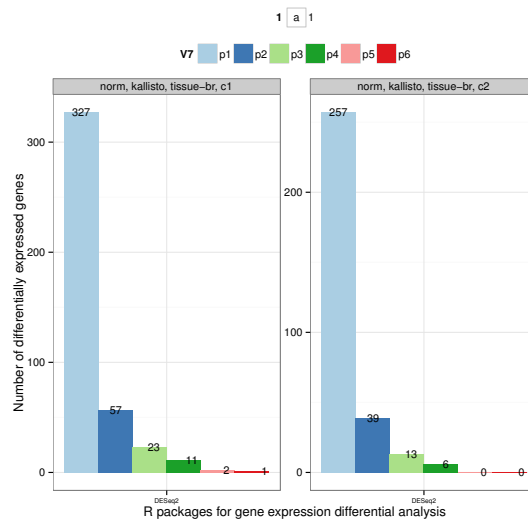
21 The below graph shows the number of differentially expressed genes when raw reads were **NOT** normal-  
 22 ized.  
 23

```
read.table("./data/summary.br.norm.txt") %>%
```

```

ggplot(aes(
  x = V1,
  y = V10,
  fill = V7)) +
  geom_bar(stat = "identity",
    position = "dodge") +
  geom_text(aes(x = V1,
    y = V10,
    ymax = V10,
    label = V10,
    size = 1,
    hjust = 0),
    position = position_dodge(width = 1)) +
  facet_wrap(~ V4 + V5 + V6 + V8,
    ncol = 2,
    scales = "free") +
  scale_fill_brewer(type = "qual", palette = "Paired") +
  theme_bw() +
  theme(legend.position = "top",
    axis.text.x = element_text(vjust = .5,
      size = 6)) +
  labs(x = "R packages for gene expression differential analysis",
    y = "Number of differentially expressed genes")

```



### 1.3 Increasing DEGs by changing the normalization strategy: abundance quantification with alignment *express*

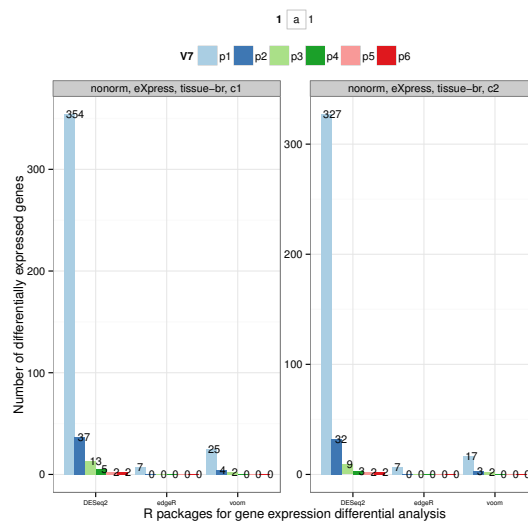
The R package *eXpress* is solely used to quantify the abundance of differentially expressed genes from **NOT normalized** reads and aligned with **Bowtie 1**.

```
read.table("./data/summary.br.nonnorm.44062.txt") %>%
```

```

ggplot(aes(
  x = V1,
  y = V10,
  fill = V7)) +
  geom_bar(stat = "identity",
    position = "dodge") +
  geom_text(aes(x = V1,
    y = V10,
    ymax = V10,
    label = V10,
    size = 1,
    hjust = 0),
    position = position_dodge(width = 1)) +
  facet_wrap(~ V4 + V5 + V6 + V8,
    ncol = 2,
    scales = "free") +
  scale_fill_brewer(type = "qual", palette = "Paired") +
  theme_bw() +
  theme(legend.position = "top",
    axis.text.x = element_text(vjust = .5,
      size = 6)) +
  labs(x = "R packages for gene expression differential analysis",
    y = "Number of differentially expressed genes")

```



29 The R package *eXpress* is solely used to quantify the abundance of differentially expressed genes from  
 30 **normalized** reads and aligned with **Bowtie 1**.  
 31

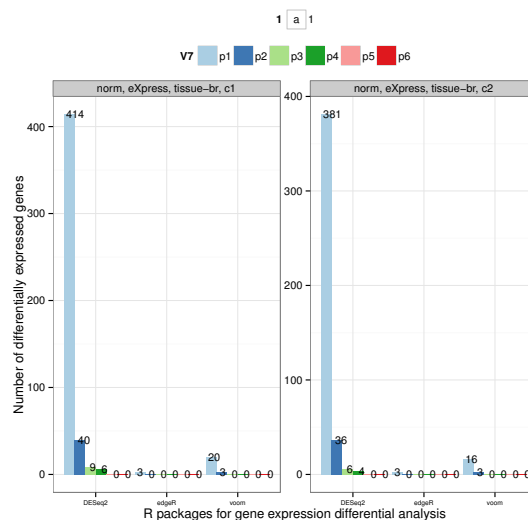
```
read.table("./data/summary.br.norm.44060.txt") %>%
```



```

ggplot(aes(
  x = V1,
  y = V10,
  fill = V7)) +
  geom_bar(stat = "identity",
    position = "dodge") +
  geom_text(aes(x = V1,
    y = V10,
    ymax = V10,
    label = V10,
    size = 1,
    hjust = 0),
    position = position_dodge(width = 1)) +
  facet_wrap(~ V4 + V5 + V6 + V8,
    ncol = 2,
    scales = "free") +
  scale_fill_brewer(type = "qual", palette = "Paired") +
  theme_bw() +
  theme(legend.position = "top",
    axis.text.x = element_text(vjust = .5,
      size = 6)) +
  labs(x = "R packages for gene expression differential analysis",
    y = "Number of differentially expressed genes")

```



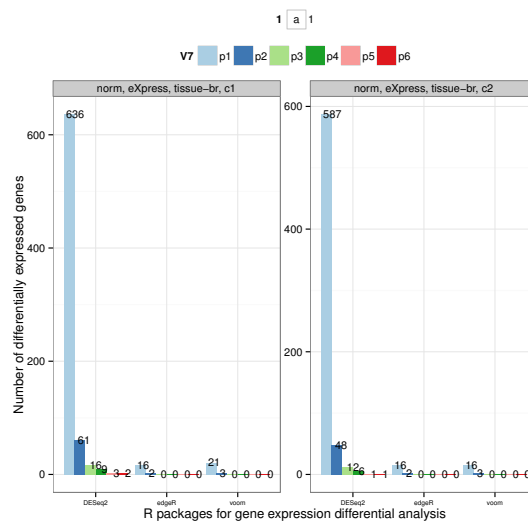
The R package *eXpress* is solely used to quantify the abundance of differentially expressed genes from **normalized reads** and aligned with **Bowtie 2**.

```
read.table("./data/summary.br.norm.44061.txt") %>%
```

```

ggplot(aes(
  x = V1,
  y = V10,
  fill = V7)) +
  geom_bar(stat = "identity",
    position = "dodge") +
  geom_text(aes(x = V1,
    y = V10,
    ymax = V10,
    label = V10,
    size = 1,
    hjust = 0),
    position = position_dodge(width = 1)) +
  facet_wrap(~ V4 + V5 + V6 + V8,
    ncol = 2,
    scales = "free") +
  scale_fill_brewer(type = "qual", palette = "Paired") +
  theme_bw() +
  theme(legend.position = "top",
    axis.text.x = element_text(vjust = .5,
      size = 6)) +
  labs(x = "R packages for gene expression differential analysis",
    y = "Number of differentially expressed genes")

```



## 2 A linear representation of gene expression

Only selected genes can be represented as follow.

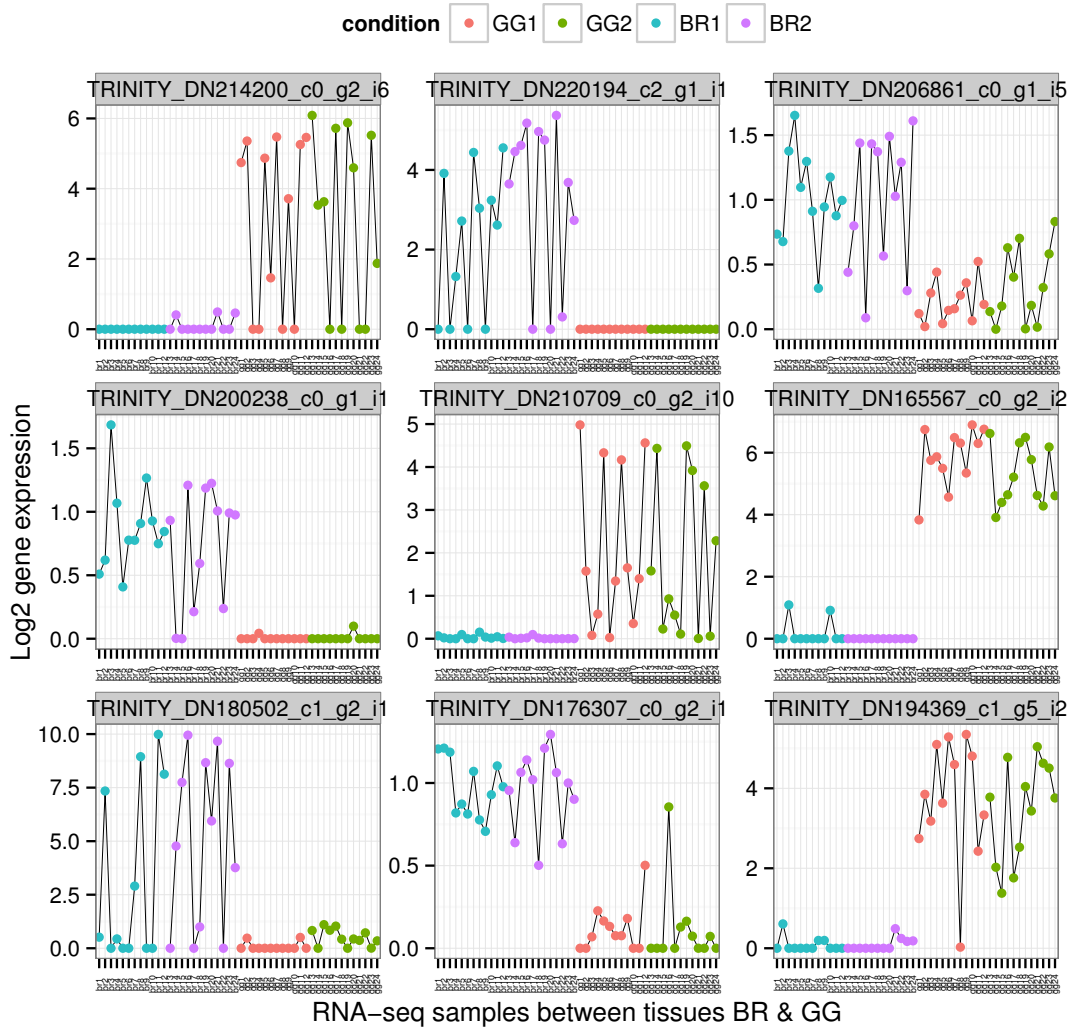
† Not more than 10 genes

```
dat <- t(read.table("./data/test.txt"))
```

```

dat <- data.frame(dat,
                  sample = rownames(dat),
                  condition = gl(4, 12, 48,
                                labels=c("GG1", "GG2", "BR1", "BR2")))
dat %>%
  gather("genes", "expression", 1:(dim(dat)[2]-2)) %>%
  ggplot(aes(x = factor(sample,
                        c(paste("br", seq(1, 24), sep=""),
                          paste("gg", seq(1, 24), sep=""))),
            y = expression,
            group = condition)) +
  theme_bw() +
  geom_line(size = .2) +
  geom_point(aes(x = factor(sample),
                       y = expression,
                       colour = condition)) +
  facet_wrap(~ genes,
             ncol = 3,
             scales = "free") +
  labs(x = "RNA-seq samples between tissues BR & GG",
       y = "Log2 gene expression") +
  theme(legend.position = "top",
        axis.text.x = element_text(angle = 90,
                                     vjust = .5,
                                     size = 4)) +
  scale_fill_brewer(type = "qual", palette = "Paired",
                    name = "Oyster tissues and Diet conditions")

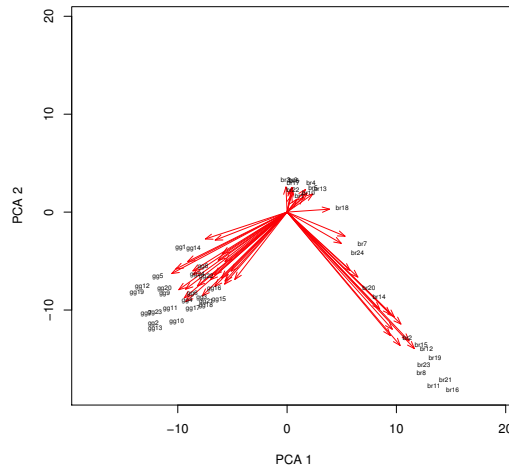
```



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39 Principal component analysis on testing data.

```
dat <- read.table("./data/test.txt")
p = prcomp(dat, retx=T)
scores = p$x
loadings <- p$rotation
sd <- p$sdev
plot(scores[,1], scores[,2],
      xlab="PCA 1", ylab="PCA 2",
      type="n", xlim=c(min(scores[,1:2]),
                        max(scores[,1:2])),
          ylim=c(min(scores[,1:2]),
                  max(scores[,1:2]))),
      arrows(0,0,loadings[,1]*50,loadings[,2]*50,
             length=0.1,angle=20, col="red")
text(loadings[,1]*50*1.3,loadings[,2]*50*1.3,
      rownames(loadings), col="black", cex=0.5)
```



### 3 Identifying foreign transcripts in oyster cells

Microscopy protocols identified the presence of a form of parasitic worm in healthy oyster tissues. As a result we might have a parasitic contamination of our RNA-seq reads. Fortunately, this can give a chance to extract the foreign RNA reads, which are already sequenced, assemble them so we identify later on the parasitic species.

We gathered 5 genomes of parasitic worms [first link](#), [second link](#). We used 2 separate strategies to assemble contigs specific for each genome, i- using genome-guided assembly by mapping reads to genome or ii- mapping contigs directly to genome.

#### 3.1 Genome-guided assembly

We used bwa to map reads to each genome separately, then use only the proper mapped mates to assemble a special transcriptome using trinity. Those selected reads were then aligned to each of the five new transcriptomes separately. Abundance of reads and differentially expressed transcripts were finally identified.

1. Clonorchis sinensis
2. Opisthorchis viverrini
3. Schistosoma haematobium
4. Schistosoma japonicum
5. Schistosoma mansoni

All previous genomes, fasta files, and some annotations can be found at [NCBI](#).

```
p1 <- read.table("./data/summary.eXpress.134221.txt")
```

```

p3 <- read.table("./data/summary.eXpress.134239.txt")
p4 <- read.table("./data/summary.eXpress.134248.txt")
head(p1)

      V1      V2      V3 V4 V5      V6 V7 V8
1 DESeq2 eXpress tissue p1 c1 134221 198 198
2 DESeq2 eXpress tissue p1 c2 134221 134 134
3 DESeq2 eXpress tissue p2 c1 134221 99 99
4 DESeq2 eXpress tissue p2 c2 134221 80 80
5 DESeq2 eXpress tissue p3 c1 134221 72 72
6 DESeq2 eXpress tissue p3 c2 134221 59 59

p1$V9 <- c("P1")
p3$V9 <- c("P3")
p4$V9 <- c("P4")
rbind(p1,p3,p4) %>%
  ggplot(aes(x = V9,
             y = V7,
             fill = V4)) +
  theme_bw() +
  geom_bar(stat = "identity",
           position = "dodge") +
  facet_wrap(~ V1 + V3 + V5,
             ncol = 6,
             scales = "free") +
  labs(x = "Worm parasites assembled transcriptomes",
       y = "Number of differentially expressed contigs") +
  theme(legend.position = "top",
        axis.text.x = element_text(vjust = .5, size = 6),
        axis.text.y = element_text(vjust = .5, size = 6),
        strip.text = element_text(size = 4),
        axis.ticks = element_line(size = .2),
        axis.ticks.length = unit(.05, "cm")) +
  scale_fill_brewer(type = "qual", palette = "Paired",
                   name = "P-value")

```



60

## 61 4 System Information

62 The version number of R and packages loaded for generating the vignette were:

```
###save(list=ls(pattern=".*|. *" ), file="PD.Rdata")
```

# **sessionInfo()**

R version 3.2.1 (2015-06-18)

Platform: x86\_64-unknown-linux-gnu (64-bit)

Running under: elementary OS Luna

## locale:

[1] LC_CTYPE=en_US.UTF-8	LC_NUMERIC=C
[3] LC_TIME=en_US.UTF-8	LC_COLLATE=en_US.UTF-8
[5] LC_MONETARY=en_US.UTF-8	LC_MESSAGES=en_US.UTF-8
[7] LC_PAPER=en_US.UTF-8	LC_NAME=en_US.UTF-8
[9] LC_ADDRESS=en_US.UTF-8	LC_TELEPHONE=en_US.UTF-8
[11] LC_MEASUREMENT=en_US.UTF-8	LC_IDENTIFICATION=en_US.UTF-8

## attached base packages:

[1] grid	stats	graphics	grDevices	utils	datasets
[7] methods	base				

## other attached packages:

[1] scales_0.2.5	tidyr_0.2.0	dplyr_0.4.2
[4] latticeExtra_0.6-26	RColorBrewer_1.1-2	glmnet_2.0-2
[7] foreach_1.4.2	Matrix_1.2-1	leaps_2.9
[10] caret_6.0-47	ggplot2_1.0.1	lattice_0.20-31
[13] xlsx_0.5.7	xlsxjars_0.6.1	rJava_0.9-6
[16] knitr_1.10.5	RevoUtilsMath_3.2.1	

## loaded via a namespace (and not attached):

[1] Rcpp_0.11.6	compiler_3.2.1	formatR_1.2
[4] nloptr_1.0.4	plyr_1.8.3	highr_0.5
[7] iterators_1.0.7	tools_3.2.1	digest_0.6.8
[10] lme4_1.1-8	evaluate_0.7	nlme_3.1-121
[13] gtable_0.1.2	mgcv_1.8-6	DBI_0.3.1
[16] parallel_3.2.1	brglm_0.5-9	SparseM_1.6
[19] proto_0.3-10	BradleyTerry2_1.0-6	stringr_1.0.0
[22] gtools_3.5.0	nnet_7.3-10	R6_2.0.1
[25] minqa_1.2.4	reshape2_1.4.1	car_2.0-25
[28] magrittr_1.5	codetools_0.2-11	MASS_7.3-41
[31] splines_3.2.1	assertthat_0.1	pbkrtest_0.4-2
[34] colorspace_1.2-6	labeling_0.3	quantreg_5.11
[37] stringi_0.5-5	lazyeval_0.1.10	munsell_0.4.2