## COBB 2010 - Foundations of Computational Biology

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## Homework 6 - Due Thursday, October 1, 2020 at 1:45PM EDT

Arabidopsis is a commonly used model organism. There is a preliminary result of the sequencing analysis of Arabidopsis thaliana, and please conduct further analysis associating the questions.

- 1. Please read the following data into a data frame, and get read of NA values. **Data source**: https://raw.githubusercontent.com/jmzeng1314/my-R/master/DEG\_scripts/tair/DESeq2\_DEG.Day1-Day0.txt
- 2. By applying org. At. tair. db package. Based on the manual, transfer the original gene\_id into gene symbol.
  - one may notice some original gene ids in the file have no corresponding gene symbols. This is because these genes have little biological meaning or they have not been annotated yet. Try to count the number of these ids.
  - Meanwhile, some original ids have more than one gene symbol due to they were named by the different research centers. Complete the gene id convert with an output like below and export the data frame into a .csv file:

```
gene_id
            baseMean log2FoldChange ...
                                                   symbol
AT2G33830 1938.15972
                          -2.560609 ...
                                              AtDRM2/DRM2
AT2G33750
                         -19.989301 ...
                                              ATPUP2/PUP2
             9.78974
AT3G54500 2238.31465
                           2.720430 ...
                                                     LNK2
AT2G46830 169.50508
                           3.527082 ...
                                              AtCCA1/CCA1
AT1G28330 1636.02224
                           -1.493341 ... AtDRM1/DRM1/DYL1
AT1G48598
            39.84136
                          -9.779030 ...
                                                 CPuORF31
```

- 3. Find out the top 10 significantly expressed genes by setting padj value less than 0.05.
  - Generate a barplot of baseMean of the top 10 most significantly expressed genes that have symbol names. To make it visually compared easily, one may want to use logarithmic value, and show gene symbols instead of gene ids.
  - Generate a dot plot of the log2FoldChange data in increasing order, and show the error bar as log folder change standard error (lfcSE).
- 4. Isolate all of the deferentially expressed genes (padj < 0.05) into a new data frame, and count the number of significantly up/down-regulated genes.
- 5. Plot the relationship between log2FoldChange and -log(padj), with highlighting the significantly up/down-regulated genes in the figure (Volcano plot).

Please complete the above tasks and generate a report with R markdown, and show all the code written in chunks (it is unnecessary to show outcomes of each chunk, one can decide to show which part). One should submit a *.rmd* file.