

WT

Marginal Blastozone A (distal) vs Marginal Blastzone in B (leaflet, mid) region

```
library(edgeR)
```

Read in Data

Read in raw count data per gene.

```
counts <- read.delim("../sam2countsResults.tsv",row.names=1)

#check the file
head(counts)
summary(counts)
colnames(counts)
#need to convert NA to 0 counts
counts[is.na(counts)] <- 0
```

Subset per DE experiment

I am going to start by subsetting the particular treatments I am looking at.

```
colnames(counts)
```

```
## [1] "tf2ambr1"      "tf2ambr3"      "tf2ambr4"      "tf2ambr6"
## [5] "tf2aother1"    "tf2aother2"    "tf2aother4"    "tf2aother7"
## [9] "tf2bmbr2"      "tf2bmbr5"      "tf2bmbr6"      "tf2bother1"
## [13] "tf2bother3"    "tf2bother4"    "tf2bother6"    "tf2cmbr1.4"
## [17] "tf2cmbr3"      "tf2cmbr6"      "tf2cmbr7"      "tf2cother2"
## [21] "tf2cother5"    "tf2cother6"    "tf2cother7"    "wtambr2"
## [25] "wtambr4"       "wtambr5"       "wtaother1"     "wtaother5"
## [29] "wtaother6"     "wtaother7"     "wtaother8"     "wtbmbr2"
## [33] "wtbmbr3"       "wtbmbr6"       "wtbmbr8"       "wtbother1.4"
## [37] "wtbother3"     "wtbother5"     "wtbother8"     "wtcmbr10"
## [41] "wtcmbr1.4.6"   "wtcmbr2"       "wtcmbr3"       "wtcmbr7"
## [45] "wtcmbr9"       "wtcother1.3.4" "wtcother2"     "wtcother6"
```

```
WTambrVSbmbr <- counts[,c(24:26,32:34)]
head(WTambrVSbmbr)
```

```
##           wtambr2 wtambr4 wtambr5 wtbmbr2 wtbmbr3 wtbmbr6
## Solyc00g005040.2.1      0      2      8      2      4      3
## Solyc00g005050.2.1      0      6      6     20      5     18
## Solyc00g005060.1.1      0      0      1      1      2      1
```

```
## Solyc00g005070.1.1      24      3      9      14      6      12
## Solyc00g005080.1.1      9      15     19     25     15     27
## Solyc00g005150.1.1      0      1      2      0      0      3
```

```
colnames(WTambrVSbmbr)
```

```
## [1] "wtambr2" "wtambr4" "wtambr5" "wtbmbr2" "wtbmbr3" "wtbmbr6"
```

```
group <- c(rep("wtambr", 3), rep("wtbmbr", 3))
d <- DGEList(counts=WTambrVSbmbr,group=group)
```

```
d$samples
```

```
##           group lib.size norm.factors
## wtambr2 wtambr   395165           1
## wtambr4 wtambr   792542           1
## wtambr5 wtambr   632686           1
## wtbmbr2 wtbmbr  1355352           1
## wtbmbr3 wtbmbr  1213142           1
## wtbmbr6 wtbmbr  1598917           1
```

```
cpm.d <- cpm(d)
d <- d[rowSums(cpm.d>5)>=3,]
d <- estimateCommonDisp(d,verbose=T)
```

```
## Disp = 0.3611 , BCV = 0.6009
```

```
d <- calcNormFactors(d)
d <- estimateCommonDisp(d)
DEtest <- exactTest(d,pair=c("wtambr","wtbmbr"))
head(DEtest$table)
```

```
##           logFC logCPM    PValue
## Solyc00g005050.2.1  0.3859  3.275 4.967e-01
## Solyc00g005070.1.1 -2.6718  3.844 4.260e-04
## Solyc00g005080.1.1 -1.1181  4.285 1.122e-01
## Solyc00g005440.1.1  0.4769  4.563 4.648e-01
## Solyc00g005840.2.1 -0.7871  4.646 2.347e-01
## Solyc00g006470.1.1 -5.5457 11.831 3.440e-13
```

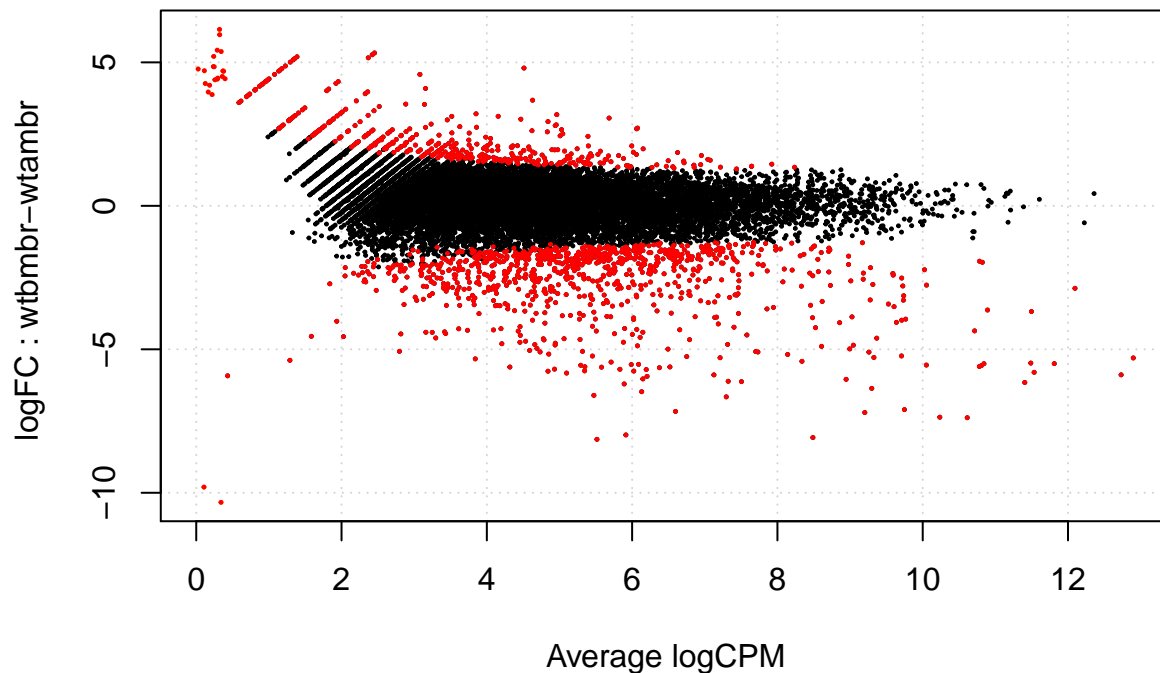
```
sum(DEtest$table$PValue<.05)
```

```
## [1] 1345
```

```
summary(decideTestsDGE(DEtest,p.value=.05))
```

```
##      [,1]
## -1    381
##  0   13403
##  1      69
```

```
sig.genes <- rownames(DEtest$table[DEtest$table$PValue<0.05,])
plotSmea(d,de.tags=sig.genes)
```



Subset by all the ones with a significant score

```
results.sig <- subset(DEtest$table, DEtest$table$PValue < 0.05)
```

What are the genes that are misexpressed? For this we need to add some annotation

```
annotation1<- read.delim("../ITAG2.3_all_Arabidopsis_ITAG_annotations.tsv", header=FALSE) #Changed to
colnames(annotation1)<- c("ITAG", "SGN_annotation")
annotation2<- read.delim("../ITAG2.3_all_Arabidopsis_annotated.tsv")
annotation <- merge (annotation1,annotation2, by =1,1, all.x=TRUE)
head(annotation)
```

```
##          ITAG
## 1 Solyc00g005000.2.1
## 2 Solyc00g005020.1.1
## 3 Solyc00g005040.2.1
## 4 Solyc00g005050.2.1
## 5 Solyc00g005060.1.1
## 6 Solyc00g005070.1.1
##
## 1 Aspartic proteinase nepenthesin I (AHRD V1 ***- A9ZMF9_NEPAL); contains Interpro domain(s) IPR001
## 2                                     Unknown
## 3 Potassium channel (AHRD V1 ***- DOEM91_9ROSI); contains Interpro domain(s) IPR000595 Cyclic n
## 4                                     Arabinogalactan protein (AHRD V1 :
## 5                                     Unknown
## 6                                     Unknown
##          AGI symbol
## 1 AT3G20015 <NA>
```

```
## 2      <NA>    <NA>
## 3 AT5G46240    KAT1
## 4 AT5G11680    <NA>
## 5      <NA>    <NA>
## 6      <NA>    <NA>
##
## 1 pepsin A; similar to aspartyl protease family protein [Arabidopsis thaliana] (TAIR:AT3G18490.1); s
## 2
## 3
## 4
## 5
## 6
##      X..identity alignment.length e.value bit.score percent.query.align
## 1      63.76              447 7e-148      520              89.94
## 2      NA                NA      NA      NA              NA
## 3      66.02              103 2e-37      150              85.71
## 4      76.96              204 1e-88      322              98.98
## 5      NA                NA      NA      NA              NA
## 6      NA                NA      NA      NA              NA
```

```
results.annotated <- merge(results.sig,annotation,by.x="row.names",by.y="ITAG",all.x=T,sort=F)
```

Write table with results

```
write.table(results.annotated,"wtambr_wtbmbr_DE1.txt",sep="\t",row.names=F)
```