## ABSTAT17

IGC, April 10–13, 2017

ABSTAT17 - MT

**EXERCISES:** Multiple Testing

In this exercise we will work with the multtest R package for multiple testing, although there are several packages which have in their procedures multiple testing approaches.

Sandrine Dudoit and colleagues implemented the R multtest package, which is part of the Bioconductor software, to perform multiple testing analyses. See the multtest documentation here.

1. Load the multtest package and explore the mt.teststat function. Load the data golub from the leukemia microarray study of Golub et al. (1999).

```
source("http://bioconductor.org/biocLite.R")
library(Biobase)
biocLite("multtest")
library(multtest)
?mt.teststat

data(golub)
?golub
head(golub)
golub.cl
ncol(golub)
nrow(golub)
```

2. Compute the t-statistic and Wilcoxon statistic and compare their normal QQ plots. Are the Wilcoxon results similar to those for the t-test?

The mt.teststat function calculates a test statistic (for example, the t-statistic or the Wilcoxon) for each row in a data frame.

```
?mt.teststat
welch_t<-mt.teststat(golub,golub.cl,test="t")
head(welch_t)
qqnorm(welch_t)
qqline(welch_t)
wilk<-mt.teststat(golub,golub.cl,test="wilcoxon")
qqnorm(wilk)
qqline(wilk)</pre>
```

3. Calculate the unadjusted p-value corresponding to each t-statistic in the list determined earlier.

```
raw_p_t<-2*(1-pnorm(abs(welch_t)))
hist(raw_p_t)
plot(sort(raw_p_t))
length(raw_p_t)*0.01
length(raw_p_t[raw_p_t<0.01])</pre>
```

4. Calculate the adjusted p-values using the Bonferroni, Holm, and Benjamini-Hochberg methods

The mt.rawp2adjp function computes adjusted p-values for each raw-pvalue.

```
?mt.rawp2adjp
procs = c("Bonferroni", "Holm", "BH")
res = mt.rawp2adjp(raw_p_t, procs)
names(res)
adjp = res$adjp[order(res$index),]
round(adjp,3)
```

5. Look at the number of rejected null hypotheses at successive p-values from alpha = 0.05 to alpha=1, using the Bonferroni, Holm, and Benjamini-Hochberg procedures and compare the results.

The mt.reject function returns the number of rejected hypotheses corresponding to each adjusted p-value for the multiple testing correction procedures you specify.

```
?mt.reject
mt.reject(adjp, c(0.01,0.05))$r
mt.reject(adjp, seq(0,1, 0.05))$r
```

6. Compare de adjusted p-values obtained via packages multtest, qvalue, p.adjust and fdrtool. If possible, print also the FDR.

```
biocLite("fdrtool")
library(fdrtool)
fdr<-fdrtool(raw_p_t,statistic="pvalue")</pre>
names (fdr)
head(fdr$pval)
head(fdr$qval)
head(fdr$lfdr)
# or
biocLite("qvalue")
library(qvalue)
fdr.q <- qvalue(raw_p_t)</pre>
names(fdr.q)
head(fdr.q$pvalues)
head(fdr.q$qvalues)
head(fdr.q$lfdr)
# or
fdr.p <- p.adjust(raw_p_t, "BH")</pre>
head(fdr.p)
```