

**Outline: bringing together what we've covered so far with an example**

**read in data**

**see if it makes sense**

**clean up or filter it**

**do some statistical tests**

**explore the results**

**manuscript()**

**Today's example starts with pre-processed data**

**we're using a specialised program to preprocess array data**

**gcos**

**dChip**

**RMAexpress**

**text file, one column per chip**

## read in data

you really want informative row and column names

you don't want extraneous columns

```
kk <- read.delim('data.txt',sep='\t',row.names=1)
```

```
str(kk)
```

```
kk <- as.matrix(kk)
```

## explore the data: what does it look like?

```
image(kk)
```

```
image(is.na(kk))
```

```
plot(density(kk))
```

```
plot(density(log(kk),na.rm=T))
```

```
mean(kk)
```

```
rowmeans<-apply(kk,1,mean,na.rm=T)
```

### explore the data: any odd men out?

```
colmeans<-apply(kk,2,mean,na.rm=T)
```

```
(kkcor<-cor(kk))
```

```
image(kkcor)
```

```
plot.cor(kkcor)
```

```
min(kkcor)
```

### filtering the data

get rid of uninteresting or problem data

reduce number of tests

various ways of generating a logical vector

```
high <- rowmeans > 300
```

```
complete <- apply(kk,1, function(x) !any(is.na(x)) )
```

### making factors for statistical tests

```
fac<-strsplit(colnames(kkm),'_')
```

```
fac<-data.frame(fac)
```

```
fac<-t(fac)
```

```
rownames(fac)<-NULL
```

```
colnames(fac)<-c('chip','date','sampno','geno','treat','tissue')
```

```
fac<-data.frame(fac)
```

### try some tests on one row

```
t.test(kkm[1,]~fac$tissue)
```

```
t.test(kkm[1,]~fac$tissue,var.equal=T) ->tt
```

```
str(tt)
```

```
tt$p.value
```

**make a wrapper function and apply it to all the rows**

```
ttrow <- function (x) {  
  tt<-t.test(x ~ fac$tissue)  
  tt$p.value }  
ttispval<-apply(kk,1,ttrow)
```

**look at the p-values**

```
sort(ttispval)[1:10]  
sum(ttispval < .05/1999)  
library(GeneTS)  
fdr.control(ttispval)
```

## exercise for next time

do the same thing using anova

do some other anova tests

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