GS01 0163
Analysis of Microarray Data

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dChip is Tricky?

Reported answers:

444, 594, 595*, 604, 604, 605, 609, 609, 615, 896*.

Answers we got:

610, using dChip2004
604, using dChip2006
900 or so, using PM-MM.

We didn’t check the intersections. Reporting version as well as filters may be important.
justRMA() is tricky

I grouped all 44 CEL files into a single folder and ran justRMA(). The resulting exprSet had 44 columns, even though 2 of the CEL files were U95Av2. The annotation indicated that the arrays were hgu95a.

It trusts you.
adding phenoData is tricky

After putting the 2 U95Av2 files in a different folder, I ran justRMA() again. I also loaded in the sample info file that we’d used for dChip in order to use it as phenoData here.

> zed <- justRMA();
> pD1 <- read.phenoData("krc-sample-info.xls", sep="\t", header=TRUE);
what do we have?

> pData(zed)[1:4,]

          sample
CL2001011101AA.CEL 1
CL2001011102AA.CEL 2
CL2001011104AA.CEL 3
CL2001011105AA.CEL 4

> pData(pD1)[1:4,]

          Scan.name Sample.Name type  Split
1   CL2001011101AA        ALL01   ALL  Training
2   CL2001011104AA        ALL02   ALL  Training
3   CL2001011105AA        ALL03   ALL  Training
4   CL2001011108AA        ALL04   ALL  Training

Note that the order is not the same...
> phenoData(zed) <- pData1;
> pData(zed)[1:4,]

<table>
<thead>
<tr>
<th>Scan.name</th>
<th>Sample.Name</th>
<th>type</th>
<th>Split</th>
</tr>
</thead>
<tbody>
<tr>
<td>CL20010111101AA</td>
<td>ALL01</td>
<td>ALL</td>
<td>Training</td>
</tr>
<tr>
<td>CL20010111104AA</td>
<td>ALL02</td>
<td>ALL</td>
<td>Training</td>
</tr>
<tr>
<td>CL20010111105AA</td>
<td>ALL03</td>
<td>ALL</td>
<td>Training</td>
</tr>
<tr>
<td>CL20010111108AA</td>
<td>ALL04</td>
<td>ALL</td>
<td>Training</td>
</tr>
</tbody>
</table>

> colnames(zed@exprs)[1:4]

[1] "CL20010111101AA.CEL" "CL20010111102AA.CEL"
[3] "CL20010111104AA.CEL" "CL20010111105AA.CEL"

oops...

The columns weren’t reordered when the phenoData changed.
phenoData assignment is trusting

There is very little validity checking involved. You can assign a phenoData object that has the wrong number of rows, or that has an order different from the ordering of the CEL files in the exprs matrix.

You need to check to make sure that the order matches.

If you supply the filenames option to just.rma() or justRMA(), the files will be loaded in the order specified.
how can we reorder things?

One option:

```r
zed2 <- as.character(pData(pD1)$Scan.name);
sampleOrder <- order(charmatch(zed2, colnames(zed@exprs)));
pD2 <- pD1;
pData(pD2) <- pData(pD2)[sampleOrder,];
phenoData(zed) <- pD2;
```
Other checks

if (length(colnames(zed@exprs)) != dim(pData(zed))[1]) {
    print("Houston, we have a problem");
}

Now, given that we have associated a phenoData object with the exprSet in the right way, we want to invoke contrasts using variables from here.

myTs <- MultiTtest(zed, "type");