

# Primer: Preparing NChannelSet objects with differential expression scores

July 30, 2013

## 1 Differential expression analysis

The `gCMAP` package offers a the `generate_gCMAP_NChannelSet` function to process multiple instances differential expression experiments with two classes (e.g. cases vs controls). For microarray data, the `limma` package is used to calculate a moderated t-statistic (default). Optionally, a standard t-test can be computed instead. For RNAseq data, the `DESeq` package is used instead.

Data preprocessing differs considerably between different technologies and array platforms and needs to be performed beforehand. Normalized microarray data and accompanying annotation is passed to `generate_gCMAP_NChannelSet` as a list of `ExpressionSet` objects, RNAseq data can be passed as a list of `CountDataSet` objects instead.

To generate a set of 3 example `CountDataSets` , we use the `makeExampleCountDataSet` function from the `DESeq` package.

```
> library(gCMAP)
> library(DESeq)
> set.seed( 123 )
> cds.list <- lapply( 1:3, function(n) {
+   cds <- makeExampleCountDataSet()
+   featureNames(cds) <- paste("gene",1:10000, sep="_")
+   cds
+ })
> names(cds.list) <- paste("Instance", 1:3, sep="")
> sapply(cds.list, dim)
```

	Instance1	Instance2	Instance3
Features	10000	10000	10000
Samples	5	5	5

```
> sapply(cds.list, function(n) pData(n)$condition )
```

	Instance1	Instance2	Instance3
[1,]	"A"	"A"	"A"
[2,]	"A"	"A"	"A"
[3,]	"B"	"B"	"B"
[4,]	"B"	"B"	"B"
[5,]	"B"	"B"	"B"

By default, each `CountDataset` object contains counts for 10000 genes from five samples. Each sample is assigned to one of two conditions, A or B, in the `phenoData` slot of the `CountDataSet` . The `pData`

column containing group membership information ( e.g. "condition" ) is provided as the `control_perturb_col` parameter. The levels associated with control and treatment groups are specified as "control" and "perturb" character strings.

Each of the three `CountDataSet` instances is analyzed individually by `generate_gCMAP_NChannelSet` . To assemble the results into a single `NChannelSet`, the input `ExpressionSet` or `CountDataSet` objects must contain measurements for the same features (e.g. the vectors returned by "featureNames" must be identical across all instances).

To include information about the instances in the `NChannelSet`, a 'sample.annotation' data.frame can be provided, containing exactly one row for each element of the input list of `ExpressionSet` / `CountDataSet` objects.

```
> ## this step takes a little time
> cde <- generate_gCMAP_NChannelSet(cds.list,
+                               uids=1:3,
+                               sample.annotation=NULL,
+                               platform.annotation="Entrez",
+                               control_perturb_col="condition",
+                               control="A",
+                               perturb="B")
> channelNames(cde)

[1] "exprs" "log_fc" "mod_fc" "p"      "z"
```

For array data, a `NChannelSet` with slots "exprs", "z", "p", and "log\_fc" is returned, containing the average intensity across all samples within the instance, z-scores, (raw) p-values and log2 fold changes, respectively. If count data is processed, an additional "mod\_fc" channel is returned, providing the moderated fold change, calculated after performing variance-stabilising transformation across all input instances. (Please consult the `DESeq` vignette for details.)

## 1.1 Storing assayData as BigMatrix objects on disk

When large numbers of instances are processed, the resulting `NChannelSet` objects can require large amounts of memory. If a file name is provided via the "big.matrix" parameter, `generate_gCMAP_NChannelSet` uses the `BigMatrix` package to store data from each channel on disk. In the future, individual channels and / or subsets of the datasets can then be loaded without requiring the full object to be read into memory again.

To highlight this functionality, we derive three (arbitrary) instances from the `sample.ExpressionSet` object available from the `Biobase` package, process them and store the results in a temporary directory.

```
> ## list of ExpressionSets
> data("sample.ExpressionSet") ## from Biobase
> es.list <- list( sample.ExpressionSet[,1:4],
+                 sample.ExpressionSet[,5:8],
+                 sample.ExpressionSet[,9:12])
> ## three instances
> names(es.list) <- paste( "Instance", 1:3, sep=".")
> storage.file <- tempfile()
> storage.file ## filename prefix for BigMatrices

[1] "/tmp/RtmpM1HJUD/file47b71ce46e0d"

> de <- generate_gCMAP_NChannelSet(
+   es.list,
+   1:3,
```

```

+   platform.annotation = annotation(es.list[[1]]),
+   control_perturb_col="type",
+   control="Control",
+   perturb="Case",
+   big.matrix=storage.file)
> channelNames(de)

[1] "exprs" "log_fc" "p"      "z"

> head( assayDataElement(de, "z") )

           1          2          3
AFFX-MurIL2_at -1.36808562  0.04333555 -0.7255849
AFFX-MurIL10_at  1.56254427 -0.69203457  0.1589525
AFFX-MurIL4_at  -0.65915229 -0.85080055  0.1804448
AFFX-MurFAS_at  -0.31745996  0.43936805  0.2813885
AFFX-BioB-5_at  -0.08767134  0.15619365 -0.2836740
AFFX-BioB-M_at  -0.32253278  0.82819990 -0.5521458

> dir(dirname( storage.file ))

[1] "file47b71ce46e0d.rdata"          "file47b71ce46e0d_exprs"
[3] "file47b71ce46e0d_exprs.desc.rds"  "file47b71ce46e0d_log_fc"
[5] "file47b71ce46e0d_log_fc.desc.rds" "file47b71ce46e0d_p"
[7] "file47b71ce46e0d_p.desc.rds"     "file47b71ce46e0d_z"
[9] "file47b71ce46e0d_z.desc.rds"     "libloc_184_942ba54a84f5acd9.rds"
[11] "libloc_190_5c44d2ef563190b3.rds"

```

For each channel, three files were generated in the temporary directory, identified by their suffices. To demonstrate the use of these disk-based `NChannelSet` objects, we will first delete the object from the current R session and reload it from disk.

Accessing the complete matrix in the `assayData` slots, e.g. for the "z" channel, returns a `BigMatrix` object - a pointer to the associated file on disk. Upon subsetting, only the requested part of the dataset is loaded into memory.

```

> ## remove de object from R session and reload
> rm( de )
> de <-get( load( paste( storage.file, "rdata", sep=".") ) )
> class( assayDataElement(de, "z") ) ## Bigmatrix

[1] "BigMatrix"
attr(,"package")
[1] "bigmemoryExtras"

> assayDataElement(de, "z")[1:10,] ## load subset

           1          2          3
AFFX-MurIL2_at -1.36808562  0.04333555 -0.7255849
AFFX-MurIL10_at  1.56254427 -0.69203457  0.1589525
AFFX-MurIL4_at  -0.65915229 -0.85080055  0.1804448
AFFX-MurFAS_at  -0.31745996  0.43936805  0.2813885
AFFX-BioB-5_at  -0.08767134  0.15619365 -0.2836740
AFFX-BioB-M_at  -0.32253278  0.82819990 -0.5521458

```

```

AFFX-BioB-3_at -0.30488232 1.79473755 0.4374636
AFFX-BioC-5_at -0.29368831 0.34488031 0.0982909
AFFX-BioC-3_at 0.05507180 -1.89130218 0.2943413
AFFX-BioDn-5_at 0.78669240 0.74946863 1.0688364

```

The `memorize` function reads the complete `NChannelSet` into memory. Alternatively, one or more selected channels can be specified with the `'name'` parameter.

```

> ## read z-score channel into memory
> dem <- memorize( de, name="z" )
> channelNames(dem)

[1] "z"

> class( assayDataElement(dem, "z") ) ## matrix

[1] "matrix"

> sessionInfo()

```

```

R version 3.0.1 (2013-05-16)
Platform: x86_64-unknown-linux-gnu (64-bit)

```

```

locale:
 [1] LC_CTYPE=en_US.UTF-8      LC_NUMERIC=C
 [3] LC_TIME=en_US.UTF-8      LC_COLLATE=C
 [5] LC_MONETARY=en_US.UTF-8  LC_MESSAGES=en_US.UTF-8
 [7] LC_PAPER=C               LC_NAME=C
 [9] LC_ADDRESS=C            LC_TELEPHONE=C
[11] LC_MEASUREMENT=en_US.UTF-8 LC_IDENTIFICATION=C

```

```

attached base packages:
[1] parallel stats graphics grDevices utils datasets methods
[8] base

```

```

other attached packages:
 [1] DESeq_1.12.0      lattice_0.20-15    locfit_1.5-9.1
 [4] gCMAP_1.4.1      limma_3.16.7      GSEABase_1.22.0
 [7] graph_1.38.3     annotate_1.38.0    AnnotationDbi_1.22.6
[10] Biobase_2.20.1   BiocGenerics_0.6.0

```

```

loaded via a namespace (and not attached):
 [1] DBI_0.2-7         GSEAlm_1.20.0     IRanges_1.18.2
 [4] Matrix_1.0-12    RColorBrewer_1.0-5 RSQlite_0.11.4
 [7] XML_3.98-1.1     biganalytics_1.1.1 bigmemory_4.4.3
[10] bigmemory.sri_0.1.2 bigmemoryExtras_1.2.0 genefilter_1.42.0
[13] geneplotter_1.38.0 grid_3.0.1        splines_3.0.1
[16] stats4_3.0.1     survival_2.37-4   tools_3.0.1
[19] xtable_1.7-1

```