

Primer: Preparing NChannelSet objects with differential expression scores

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This document exemplifies the processing of differential expression data using small, simulated datasets shipped with the gCMAP package. To see real-life examples with data from available from public databases, please refer to the documentation of the gCMAPWeb companion package.

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. The `bigmemory` and `bigmemoryExtras` packages can be used to create `BigMatrix` objects, allowing methods to subset large datasets without having to load them fully into memory first.

Note: at the time of writing, the `bigmemory` package was only available for Unix and Mac OS X operating systems but not for Windows. Windows users can take advantage of `gCMAP`'s functionality but datasets must be fully loaded into memory first.

If the `bigmemory` and `bigmemoryExtras` packages are available and 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. Note: this section will only create the expected `big.matrix` files on disk if the `bigmemory` and `bigmemoryExtras` packages can be loaded. Otherwise, a standard Rdata object is created and a warning is issued.

```
> ## 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/RtmpuomIg8/file37ed2cbd20dc"

> 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] "file37ed2cbd20dc.rdata"

```

If the `bigmemoryExtras` package is available, it generated a `BigMatrix` objects containing pointers to three files in the temporary directory, one for each channel (identified by their suffices). If the package is unavailable, a standard `eSet` is saved to disk, which will be read fully into memory upon reload.

To demonstrate the use of disk-based `NChannelSet` objects, we will first delete the object from the current R workspace and reload it from disk.

Accessing the complete matrix in the `assayData` slots, e.g. for the "z" channel, returns another `BigMatrix` object with `assayData` slot pointing 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") )

[1] "matrix"

> 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. In addition, 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 3.2.0 RC (2015-04-08 r68161)
Platform: x86_64-apple-darwin10.8.0 (64-bit)
Running under: OS X 10.6.8 (Snow Leopard)
```

```
locale:
```

```
[1] en_US.UTF-8/en_US.UTF-8/en_US.UTF-8/C/en_US.UTF-8/en_US.UTF-8
```

```
attached base packages:
```

```
[1] stats4    parallel  stats      graphics  grDevices  utils      datasets
[8] methods   base
```

```
other attached packages:
```

```
[1] DESeq_1.20.0      lattice_0.20-31    locfit_1.5-9.1
[4] gCMAP_1.12.0      limma_3.24.0       GSEABase_1.30.0
[7] graph_1.46.0      annotate_1.46.0     XML_3.98-1.1
[10] AnnotationDbi_1.30.0 GenomeInfoDb_1.4.0 IRanges_2.2.0
[13] S4Vectors_0.6.0   Biobase_2.28.0     BiocGenerics_0.14.0
```

```
loaded via a namespace (and not attached):
```

```
[1] splines_3.2.0      GSEAlm_1.28.0      xtable_1.7-4        Category_2.34.0
[5] tools_3.2.0        grid_3.2.0          DBI_0.3.1            genefilter_1.50.0
[9] survival_2.38-1    RBGL_1.44.0         Matrix_1.2-0         geneplotter_1.46.0
[13] RColorBrewer_1.1-2 RSQLite_1.0.0
```