

# Quality assessment with arrayQualityMetrics

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## Introduction

The function `arrayQualityMetrics` can be used on *AffyBatch* for Affymetrix data sets, *ExpressionSet* in the case of non Affymetrix one colour experiments, *NChannelSet* for dual colour experiments and *BeadLevelList* for Illumina bead arrays. `arrayQualityMetrics` produces a *HTML* report as an output.

## 1 Data preparation

### 1.1 Data import

`arrayQualityMetrics` can be used on unnormalized or normalized data sets. In this example, we will produce a report on a normalized *NChannelSet*. We first need to load the data.

```
> library("Biobase")
> library("CC14")
> data("CC14")
```

### 1.2 Normalization

We can normalize the data using the variance stabilization method available in the package *vsn*.

```
> library("vsn")
> nCC14 = justvsn(CC14, subsample=2000)
```

### 1.3 Data specificities

Some of the quality metrics provided by the package are performed using specific information about the features of the arrays. For an optimal use of the package, the data can be prepared accordingly to the following conventions.

**X and Y coordinates of the spots** To plot the images of the arrays in the case of *ExpressionSet* and *NChannelSet*, `arrayQualityMetrics` needs the coordinates of the spots on the chip. Two columns corresponding to the row and column numbers of the features are thus required in the *featureData*. These columns should be named "X" for rows and "Y" for columns. If the arrays are splitted into blocks, then the function `addXYfromGAL` should be executed prior to `arrayQualityMetrics` to convert the rows and columns of the blocks in absolute "X" and "Y" on the array.

```
> featureData(nCC14)$X = featureData(nCC14)$Row
> featureData(nCC14)$Y = featureData(nCC14)$Column
```

**GC content of the reporters** If the GC content of the reporters is known, then it is possible to include it as percentages in the *featureData* of the *NChannelSet* under the column name "GC". Then a study of the GC content effect on intensities of the arrays can be performed.

**Mapping of the reporters** The report can also include a study of the effect of the target mapping of the reporters. Thus a *featureData* column named "HasTarget" should include logical "TRUE" if the reporter matches for a coding mRNA and "FALSE" if not.

```
> featureData(nCC14)$hasTarget = (regexpr("^NM", featureData(nCC14)$Name) > 0)
```

**Covariate** The *phenoData* can also have a column containing a group name (up to 12 groups) for each sample accordingly to a factor of interest. This *phenoData* will be used to provide side bars on the heatmap. In the case of the *CCl4* dataset, the RNA hybridized to the arrays can be of good, medium or poor quality accordingly to its RIN number (see *CCl4* vignette).

```
> datapath = system.file("extdata", package="CCl4")
> p = read.AnnotatedDataFrame("samplesInfo.txt", path=datapath)
> cond = paste(p$RIN.Cy3,p$RIN.Cy5,sep="/")
> poor = grep(cond,pattern="2.5")
> medium = grep(cond,pattern="^5/1/5")
> good = grep(cond,pattern="9.7")
> cov = rep(0, length = nrow(p))
> cov[good] = "Good"
> cov[medium] = "Medium"
> cov[poor] = "Poor"
> phenoData(nCC14)$RNAintegrity = cov
```

## 2 Report production

To produce a report, the function `arrayQualityMetrics` is called with the following arguments:

- *expressionset*: is an object of class *ExpressionSet*, *AffyBatch*, *NChannelSet* or *BeadLevelList*.
- *outdir*: is the directory in which the result files are created.
- *force*: if TRUE, if *outdir* already exists, it will be overwritten.
- *do.logtransform*: if TRUE, the data are log transformed before the analysis.
- *split.plots*: if the number of studied arrays is more than 50 it is advised to define a number of experiments to represent on the density plots.
- *intgroup*: is the name of the column in the *phenoData* that contains the information about a covariate of interest to be shown as a side bar on the heatmap. The default name to this column is 'Covariate'.

```
> library("arrayQualityMetrics")
> arrayQualityMetrics(expressionset = nCC14,
+                      outdir = "CCl4",
+                      force = TRUE,
+                      do.logtransform = FALSE,
+                      split.plots = FALSE,
+                      intgroup = "RNAintegrity")
```

A report named `QMreport.html` is produced in the subdirectory *CCl4*. It contains text illustrated by `.png` files. Each `.png` is linked to corresponding `.pdf` files in order to provide high quality images.

## Session Info

```
> toLatex(sessionInfo())
```

- R version 2.7.1 (2008-06-23), x86\_64-unknown-linux-gnu
- Locale: LC\_CTYPE=en\_US;LC\_NUMERIC=C;LC\_TIME=en\_US;LC\_COLLATE=en\_US;LC\_MONETARY=C;LC\_MESSAGES=
- Base packages: base, datasets, graphics, grDevices, grid, methods, splines, stats, tools, utils
- Other packages: affy 1.18.2, affyio 1.8.1, affyPLM 1.16.0, annotate 1.18.0, AnnotationDbi 1.2.2, arrayQualityMetrics 1.6.1, beadarray 1.8.0, Biobase 2.0.1, CCI4 1.0.7, DBI 0.2-4, gcrma 2.12.1, genefilter 1.20.0, geneplotter 1.18.0, lattice 0.17-12, latticeExtra 0.5-1, limma 2.14.5, match-probes 1.12.0, preprocessCore 1.2.1, RColorBrewer 1.0-2, RSQLite 0.6-9, simpleaffy 2.16.0, survival 2.34-1, vsn 3.6.0, xtable 1.5-2
- Loaded via a namespace (and not attached): KernSmooth 2.22-22