Package ‘EDASeq’

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Title Exploratory Data Analysis and Normalization for RNA-Seq

Description Numerical and graphical summaries of RNA-Seq read data.
   Within-lane normalization procedures to adjust for GC-content effect (or other gene-level effects) on read counts: loess robust local regression, global-scaling, and full-quantile normalization (Risso et al., 2011). Between-lane normalization procedures to adjust for distributional differences between lanes (e.g., sequencing depth): global-scaling and full-quantile normalization (Bullard et al., 2010).

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EDASeq-package

Exploratory Data Analysis and Normalization for RNA-Seq data

Description

Numerical summaries and graphical representations of some key features of the data along with implementations of both within-lane normalization methods for GC content bias and between-lane normalization methods to adjust for sequencing depth and possibly other differences in distribution.

Details

The SeqExpressionSet class is used to store gene-level counts along with sample information. It extends the virtual class eSet. See the help page of the class for details.

"Read-level" information is managed via the FastqFileList and BamFileList classes of Rsamtools.

Most used graphic tools for the FastqFileList and BamFileList objects are: 'barplot', 'plotQuality', 'plotNtFrequency'. For SeqExpressionSet objects are: 'biasPlot', 'meanVarPlot', 'MDPlot'.

To perform gene-level normalization use the functions 'withinLaneNormalization' and 'betweenLaneNormalization'.

See the package vignette for a typical Exploratory Data Analysis example.
Author(s)

Davide Risso and Sandrine Dudoit. Maintainer: Davide Risso <risso.davide@gmail.com>

References


barplot-methods  Methods for Function barplot in Package EDASeq

Description

High-level functions to produce barplots of some complex objects.

Methods

signature(height = "BamFile") Usage: barplot(height,strata=c("rname","strand")) It produces a barplot of the total number of reads in each chromosome (if "rname") or strand.

signature(height = "BamFileList") It produces a barplot of the total number of reads in each object in height. If unique=TRUE is specified, it stratified the total by uniquely/non-uniquely mapped reads.

signature(height = "FastqFileList") It produces a barplot of the total number of reads in each object in height.

betweenLaneNormalization-methods  Methods for Function betweenLaneNormalization in Package EDASeq

Description

Between-lane normalization for sequencing depth and possibly other distributional differences between lanes.

Usage

betweenLaneNormalization(x, which=c("median","upper","full"), offset=FALSE, round=TRUE)
Arguments

- **x**: A numeric matrix representing the counts or a SeqExpressionSet object.
- **which**: Method used to normalized. See the details section and the reference below for details.
- **offset**: Should the normalized value be returned as an offset leaving the original counts unchanged?
- **round**: If TRUE the normalization returns rounded values (pseudo-counts). Ignored if offset=TRUE.

Details

This method implements three normalizations described in Bullard et al. (2010). The methods are:

- **median**: a scaling normalization that forces the median of each lane to be the same.
- **upper**: the same but with the upper quartile.
- **full**: a non linear full quantile normalization, in the spirit of the one used in microarrays.

Methods

- `signature(x = "matrix")`: It returns a matrix with the normalized counts if offset=FALSE or with the offset if offset=TRUE.
- `signature(x = "SeqExpressionSet")`: It returns a linkS4class{SeqExpressionSet} with the normalized counts in the normalizedCounts slot and with the offset in the offset slot (if offset=TRUE).

Author(s)

Davide Risso.

References


Examples

```r
library(yeastRNASeq)
data(geneLevelData)
data(yeastGC)
sub <- intersect(rownames(geneLevelData), names(yeastGC))
mat <- as.matrix(geneLevelData[sub, ])
data <- newSeqExpressionSet(mat,
```
biasBoxplot-methods

Methods for Function biasBoxplot in Package EDASeq

Description

biasBoxplot produces a boxplot representing the distribution of a quantity of interest (e.g. gene counts, log-fold-changes, ...) stratified by a covariate (e.g. gene length, GC-content, ...).

Usage

biasBoxplot(x, y, num.bins, ...)

Arguments

x A numeric vector with the quantity of interest (e.g. gene counts, log-fold-changes, ...)
y A numeric vector with the covariate of interest (e.g. gene length, GC-content, ...)
num.bins A numeric value specifying the number of bins in which to stratify y. Default to 10.
... See par

Methods

signature(x = "numeric", y = "numeric", num.bins = "numeric") It plots a line representing the regression of every column of the matrix x on the numeric covariate y. One can pass the usual graphical parameters as additional arguments (see par).

Examples

library(yeastRNASeq)
data(geneLevelData)
data(yeastGC)

sub <- intersect(rownames(geneLevelData), names(yeastGC))

mat <- as.matrix(geneLevelData[sub,])
data <- newSeqExpressionSet(mat,
    phenoData=AnnotatedDataFrame(
        data.frame(conditions=factor(c("mut", "mut", "wt", "wt")),
                    row.names=colnames(geneLevelData)),
        featureData=AnnotatedDataFrame(data.frame(gc=yeastGC[sub])))

norm <- betweenLaneNormalization(data, which="full", offset=FALSE)
biasPlot-methods

Methods for Function biasPlot in Package EDASeq

Description

biasPlot produces a plot of the lowess regression of the counts on a covariate of interest, typically the GC-content or the length of the genes.

Methods

signature(x = "matrix", y = "numeric") It plots a line representing the regression of every column of the matrix x on the numeric covariate y. One can pass the usual graphical parameters as additional arguments (see par).

signature(x = "SeqExpressionSet", y = "character") It plots a line representing the regression of every lane in x on the covariate specified by y. y must be one of the column of the featureData slot of the x object. One can pass the usual graphical parameters as additional arguments (see par). The parameter color_code (optional) must be a number specifying the column of phenoData to be used for color-coding. By default it is color-coded according to the first column of phenoData. If legend=TRUE and col is not specified a legend with the information stored in phenoData is added.

Examples

library(yeastRNASeq)
data(geneLevelData)
data(yeastGC)

sub <- intersect(rownames(geneLevelData), names(yeastGC))

mat <- as.matrix(geneLevelData[sub,])
data <- newSeqExpressionSet(mat,
    phenoData=AnnotatedDataFrame(data.frame(conditions=factor(c("mut", "mut", "wt", "wt")), row.names=colnames(geneLevelData))),
    featureData=AnnotatedDataFrame(data.frame(gc=yeastGC[sub])))

biasPlot(data,"gc",ylim=c(0,5),log=TRUE)
Methods

signature(x = "FastqQuality")  It plots the distribution of the quality per read position.
signature(x = "SeqExpressionSet")  It plots the distribution of the log counts in each lane of x.

getGeneLengthAndGCContent

Description
Automatically retrieves gene length and GC-content information from Biomart or org.db packages.

Usage
getGeneLengthAndGCContent(id, org, mode=c("biomart", "org.db"))

Arguments

id  Character vector of one or more ENSEMBL or ENTREZ gene IDs.
org  Organism three letter code, e.g. 'hsa' for 'Homo sapiens'. See also: http://www.genome.jp/kegg/catalog/org_list.html; In org.db mode, this can be also a specific genome assembly, e.g. 'hg38' or 'sac-Cer3'.
mode  Mode to retrieve the information. Defaults to 'biomart'. See Details.

Details
The 'biomart' mode is based on functionality from the biomaRt package and retrieves the required information from the BioMart database. This is available for all ENSEMBL organisms and is typically most current, but can be time-consuming when querying several thousand genes at a time.

The 'org.db' mode uses organism-based annotation packages from Bioconductor. This is much faster than the 'biomart' mode, but is only available for selected model organism currently supported by BioC annotation functionality.

Results for the same gene ID(s) can differ between both modes as they are based on different sources for the underlying genome assembly. While the 'biomart' mode uses the latest ENSEMBL version, the 'org.db' mode uses BioC annotation packages typically built from UCSC.
Value

A numeric matrix with two columns: gene length and GC-content.

Author(s)

Ludwig Geistlinger <Ludwig.Geistlinger@bio.ifi.lmu.de>

See Also

getSequence to retrieve a genomic sequence from BioMart, genes to extract genomic coordinates from a TxB object, getSeq to extract genomic sequences from a BSgenome object, alphabetFrequency to calculate nucleotide frequencies.

Examples

geneLengthAndGCContent("ENSG0000012048", "hsa")
Examples

```r
library(yeastRNASeq)
data(geneLevelData)
data(yeastGC)

sub <- intersect(rownames(geneLevelData), names(yeastGC))

mat <- as.matrix(geneLevelData[sub,])
data <- newSeqExpressionSet(mat,
    phenoData=AnnotatedDataFrame(
        data.frame(conditions=factor(c("mut", "mut", "wt", "wt")),
            row.names=colnames(geneLevelData)),
        featureData=AnnotatedDataFrame(data.frame(gc=yeastGC[sub]))))

MDPlot(data,c(1,3))
```

Description

meanVarPlot produces a smoothScatter plot of the mean variance relation.

Methods

signature(x = "SeqExpressionSet") It takes as additional argument log, which if true consider
the logarithm of the counts before computing mean and variance. To avoid missing values, we
consider the maximum between 0 and the log of the counts. Along with the scatter plot the
function plots a line representing the lowess fit.

newSeqExpressionSet Function to create a new SeqExpressionSet object.

Description

User-level function to create new objects of the class SeqExpressionSet.

Usage

```r
newSeqExpressionSet(counts,
    normalizedCounts = matrix(data=NA, nrow=nrow(counts), ncol=ncol(counts), dimnames=dimnames(counts)),
    offset = matrix(data=0, nrow=nrow(counts), ncol=ncol(counts), dimnames=dimnames(counts)),
    phenoData = annotatedDataFrameFrom(counts, FALSE),
    featureData = annotatedDataFrameFrom(counts, TRUE),
    ...)
```
Arguments

- `counts`: A matrix containing the counts for an RNA-Seq experiment. One column for each lane and one row for each gene.
- `normalizedCounts`: A matrix with the same dimensions of `counts` with the normalized counts.
- `offset`: A matrix with the same dimensions of `counts` defining the offset (usually useful for normalization purposes). See the package vignette for a discussion on the offset.
- `phenoData`: A data.frame or `AnnotatedDataFrame` with sample information, such as biological condition, library preparation protocol, flow-cell,...
- `featureData`: A data.frame or `AnnotatedDataFrame` with feature information, such as gene length, GC-content, ...
- `...`: Other arguments will be passed to the constructor inherited from `eSet`.

Value

An object of class `SeqExpressionSet`.

Author(s)

Davide Risso

See Also

`SeqExpressionSet`

Examples

```r
counts <- matrix(data=0, nrow=100, ncol=4)
for(i in 1:4) {
  counts[, i] <- rpois(100, lambda=50)
}
cond <- c(rep("A", 2), rep("B", 2))

counts <- newSeqExpressionSet(counts, phenoData=data.frame(conditions=cond))
```

Description

High-level function to produce plots given one `BamFileList` object and one `FastqFileList` object.
Methods

signature(x = "BamFileList", y = "FastqFileList") It produce a barplot of the percentage of mapped reads. If strata=TRUE it stratifies the bars according to the unique/non-unique mapped reads. To be meaningful, x should be a set of aligned reads and y a set of raw reads on the same samples.

Methods for Function plotNtFrequency in Package EDASeq

Description

Plots the nucleotide frequencies per position.

Methods

signature(x = "ShortRead")
signature(x = "BamFile")
It plots the nucleotide frequencies per position, averaging all the reads in x.

Methods for Function plotPCA in Package EDASeq

Description

plotPCA produces a Principal Component Analysis (PCA) plot of the counts in object

Usage

## S4 method for signature 'matrix'
plotPCA(object, k=2, labels=TRUE, isLog=FALSE, ...)
## S4 method for signature 'SeqExpressionSet'
plotPCA(object, k=2, labels=TRUE, ...)

Arguments

object Either a numeric matrix or a SeqExpressionSet object containing the gene expression.
k The number of principal components to be plotted.
labels Logical. If TRUE, and k=2, it plots the colnames of object as point labels.
isLog Logical. Set to TRUE if the data are already on the log scale.
... See par
Details

The Principal Component Analysis (PCA) plot is a useful diagnostic plot to highlight differences in the distribution of replicate samples, by projecting the samples into a lower dimensional space.

If there is strong differential expression between two classes, one expects the samples to cluster by class in the first few Principal Components (PCs) (usually 2 or 3 components are enough). This plot also highlights possible batch effects and/or outlying samples.

Methods

signature(x = "matrix")
signature(x = "SeqExpressionSet")

Examples

library(yeastRNASeq)
data(geneLevelData)

mat <- as.matrix(geneLevelData)
data <- newSeqExpressionSet(mat, phenoData=AnnotatedDataFrame(data.frame(conditions=factor(c("mut", "mut", "wt", "wt")), row.names=colnames(geneLevelData))))

plotPCA(data, col=rep(1:2, each=2))

plotQuality-methods

Methods for Function plotQuality in Package EDASeq

Description

plotQuality produces a plot of the quality of the reads.

Methods

signature(x = "BamFileList") It produces a plot that summarizes the per-base mean quality of the reads of each BAM file in x.
signature(x = "BamFile") It produces a boxplot of the per-base distribution of the quality scores of the reads in x.
signature(x = "FastqFileList") It produces a plot that summarizes the per-base mean quality of the reads of each FASTQ file in x.
Details

Since FASTQ files can be very long, it can be very expensive to process a whole file. One way to avoid this, is to consider a subset of the file and then plot the quality of the subset. As long as one assumes that the subset is random, this is a good approximation. The function `FastqSampler` of ShortRead can be used for this. See its help page for an example.

Description

`plotRLE` produces a Relative Log Expression (RLE) plot of the counts in `x`.

Usage

`plotRLE(x, ...)`

Arguments

- `x` Either a numeric matrix or a `SeqExpressionSet` object containing the gene expression.
- `...` See `par`

Details

The Relative Log Expression (RLE) plot is a useful diagnostic plot to visualize the differences between the distributions of read counts across samples. It shows the boxplots of the log-ratios of the gene-level read counts of each sample to those of a reference sample (defined as the median across the samples). Ideally, the distributions should be centered around the zero line and as tight as possible. Clear deviations indicate the need for normalization and/or the presence of outlying samples.

Methods

- `signature(x = "matrix")`
- `signature(x = "SeqExpressionSet")`

Examples

```r
library(yeastRNASeq)
data(geneLevelData)
mat <- as.matrix(geneLevelData)
data <- newSeqExpressionSet(mat, phenoData=AnnotatedDataFrame(data.frame(conditions=factor(c("mut", "mut", "wt", "wt")),
```
SeqExpressionSet-class

"SeqExpressionSet" class for collections of short reads

Description

This class represents a collection of digital expression data (usually counts from RNA-Seq technology) along with sample information.

Objects from the Class

Objects of this class can be created from a call to the `newSeqExpressionSet` constructor.

Extends

Class `eSet`, directly. Class `VersionedBiobase`, by class `eSet`, distance 2. Class `Versioned`, by class `eSet`, distance 3.

Slots

Inherited from `eSet`:

- `assayData` Contains matrices with equal dimensions, and with column number equal to `nrow(phenoData)`. `assayData` must contain a matrix `counts` with rows representing features (e.g., genes) and columns representing samples. The optional matrices `normalizedCounts` and `offset` can be added to represent a normalization in terms of pseudo-counts or offset, respectively, to be used for subsequent analyses. See the vignette for details. Class: `AssayData-class`.

- `phenoData` Sample information. For compatibility with DESeq, there should be at least the column `conditions`. See `eSet` for details.

- `featureData` Feature information. It is recommended to include at least length and GC-content information. This slot is used for `withinLaneNormalization`. See `eSet` for details.

- `experimentData` See `eSet`

- `annotation` See `eSet`

- `protocolData` See `link{eSet}`
Methods

See eSet for inherited methods. Additional methods:

counts signature(object = "SeqExpressionSet"): returns the counts matrix.
counts<- signature(object = "SeqExpressionSet"): method to replace the counts matrix.
normCounts signature(object = "SeqExpressionSet"): returns the normalizedCounts matrix.
normCounts<- signature(object = "SeqExpressionSet"): method to replace the normalizedCounts matrix.
offst signature(object = "SeqExpressionSet"): returns the offset matrix.
offst<- signature(object = "SeqExpressionSet"): method to replace the offset slot.
boxplot signature(x = "SeqExpressionSet"): produces a boxplot of the log counts.
meanVarPlot signature(x = "SeqExpressionSet"): produces a smoothScatter plot of the mean variance relation. See meanVarPlot for details.
biasPlot signature(x = "SeqExpressionSet", y = "character"): produces a plot of the lowess regression of the counts on some covariate of interest (usually GC-content or length). See biasPlot for details.
withinLaneNormalization signature(x = "SeqExpressionSet", y = "missing"): within lane normalization for GC-content (or other lane specific) bias. See withinLaneNormalization for details.
betweenLaneNormalization signature(x = "SeqExpressionSet"): between lane normalization for sequencing depth and possibly other distributional differences between lanes. See betweenLaneNormalization for details.

Author(s)

Davide Risso <risso.davide@gmail.com>

See Also
eSet, newSeqExpressionSet, biasPlot, withinLaneNormalization, betweenLaneNormalization

Examples

takeMethods(class = "SeqExpressionSet", where = getNamespace("EDASeq"))
counts <- matrix(data = 0, nrow = 100, ncol = 4)
for(i in 1:4) {
counts[,i] <- rpois(100, lambda = 50)
}
cond <- c(rep("A", 2), rep("B", 2))
data <- newSeqExpressionSet(counts, phenoData = AnnotatedDataFrame(data.frame(conditions = cond)))
head(counts(data))
boxplot(data, col = as.numeric(pData(data)[,1]) + 1)
**withinLaneNormalization**

Methods for Function withinLaneNormalization in Package EDASeq

### Description

Within-lane normalization for GC-content (or other lane-specific) bias.

### Usage

```r
withinLaneNormalization(x, y, which=c("loess","median","upper","full"), offset=FALSE, num.bins=10, round=TRUE)
```

### Arguments

- **x**: A numeric matrix representing the counts or a `SeqExpressionSet` object.
- **y**: A numeric vector representing the covariate to normalize for (if `x` is a matrix) or a character vector with the name of the covariate (if `x` is a `SeqExpressionSet` object). Usually it is the GC-content.
- **which**: Method used to normalized. See the details section and the reference below for details.
- **offset**: Should the normalized value be returned as an offset leaving the original counts unchanged?
- **num.bins**: The number of bins used to stratify the covariate for median, upper and full methods. Ignored if loess. See the reference for a discussion on the number of bins.
- **round**: If TRUE the normalization returns rounded values (pseudo-counts). Ignored if offset=TRUE.

### Details

This method implements four normalizations described in Risso et al. (2011).

The **loess** normalization transforms the data by regressing the counts on `y` and subtracting the loess fit from the counts to remove the dependence.

The **median**, **upper** and **full** normalizations are based on the stratification of the genes based on `y`. Once the genes are stratified in `num.bins` strata, the methods work as follows.

- **median**: scales the data to have the same median in each bin.
- **upper**: the same but with the upper quartile.
- **full**: forces the distribution of each stratum to be the same using a non linear full quantile normalization, in the spirit of the one used in microarrays.
Methods

signature(x = "matrix", y = "numeric") It returns a matrix with the normalized counts if offset=FALSE or with the offset if offset=TRUE.

signature(x = "SeqExpressionSet", y = "character") It returns a SeqExpressionSet with the normalized counts in the normalizedCounts slot and with the offset in the offset slot (if offset=TRUE).

Author(s)

Davide Risso.

References


Examples

```r
library(yeastRNASeq)
data(geneLevelData)
data(yeastGC)

sub <- intersect(rownames(geneLevelData), names(yeastGC))

mat <- as.matrix(geneLevelData[sub, ])
data <- newSeqExpressionSet(mat,
                          phenoData=AnnotatedDataFrame(
                            data.frame(conditions=factor(c("mut", "mut", "wt", "wt")),
                                       row.names=colnames(geneLevelData))),
                          featureData=AnnotatedDataFrame(data.frame(gc=yeastGC[sub])))

norm <- withinLaneNormalization(data, "gc", which="full", offset=FALSE)
```

<table>
<thead>
<tr>
<th>yeastGC</th>
<th>GC-content of S. Cerevisiae genes</th>
</tr>
</thead>
</table>

Description

This data set gives the GC-content (proportion of G and C) of the genes of *S. Cerevisiae*, from SGD release 64 annotation.

Format

A vector containing 6717 observations.
Source

SGD release 64: http://www.yeastgenome.org

<table>
<thead>
<tr>
<th>yeastLength</th>
<th>Length of <em>S. Cerevisiae</em> genes</th>
</tr>
</thead>
</table>

Description

This data set gives the length (in base pairs) of the genes of *S. Cerevisiae*, from SGD release 64 annotation.

Format

A vector containing 6717 observations.

Source

SGD release 64: http://www.yeastgenome.org
Index

* classes
  SeqExpressionSet-class, 14

* datasets
  yeastGC, 17
  yeastLength, 18

* methods
  barplot-methods, 3
  betweenLaneNormalization-methods, 3
  biasBoxplot-methods, 5
  biasPlot-methods, 6
  boxplot-methods, 7
  MDPlot-methods, 8
  meanVarPlot-methods, 9
  plot-methods, 10
  plotNtFrequency-methods, 11
  plotPCA-methods, 11
  plotQuality-methods, 12
  plotRLE-methods, 13
  withinLaneNormalization-methods, 16

alphabetFrequency, 8
AnnotatedDataFrame, 10

BamFileList, 2
barplot,BamFile-method (barplot-methods), 3
barplot,BamFileList-method (barplot-methods), 3
barplot,FastqFileList-method (barplot-methods), 3
barplot-methods, 3
betweenLaneNormalization, 15
betweenLaneNormalization (betweenLaneNormalization-methods), 3
betweenLaneNormalization,SeqExpressionSet-method (betweenLaneNormalization-methods), 3
betweenLaneNormalization-methods, 3
biasBoxplot (biasBoxplot-methods), 5
biasBoxplot,numeric,numeric-method (biasBoxplot-methods), 5
biasBoxplot,numeric-method (biasBoxplot-methods), 5
biasPlot-methods, 5
biasPlot, 15
biasPlot (biasPlot-methods), 6
biasPlot,numeric-method (biasPlot-methods), 6
biasPlot,SeqExpressionSet,character-method (biasPlot-methods), 6
biasPlot-methods, 6
boxplot,FastqQuality-method (boxplot-methods), 7
boxplot,SeqExpressionSet-method (boxplot-methods), 7
boxplot-methods, 7
counts,SeqExpressionSet-method (SeqExpressionSet-class), 14
counts<-,SeqExpressionSet-method (SeqExpressionSet-class), 14

EDASeq (EDASeq-package), 2
EDASeq-package, 2
eSet, 2, 10, 14, 15
exprs,SeqExpressionSet-method (SeqExpressionSet-class), 14
exprs<-,SeqExpressionSet,ANY-method (SeqExpressionSet-class), 14

FastqFileList, 2
FastqSampler, 13
genesis, 8
getGeneLengthAndGCContent, 7
getSeq, 8
getSequence, 8
initialize, SeqExpressionSet-method (SeqExpressionSet-class), 14
lowess, 6, 9, 15
MDPlot (MDPlot-methods), 8
MDPlot, matrix, numeric-method (MDPlot-methods), 8
MDPlot, SeqExpressionSet, numeric-method (MDPlot-methods), 8
MDPlot-methods, 8
meanVarPlot, 15
meanVarPlot (meanVarPlot-methods), 9
meanVarPlot, SeqExpressionSet-method (meanVarPlot-methods), 9
meanVarPlot-methods, 9
newSeqExpressionSet, 9, 14, 15
normCounts (SeqExpressionSet-class), 14
normCounts, SeqExpressionSet-method (SeqExpressionSet-class), 14
normCounts<- (SeqExpressionSet-class), 14
normCounts<-, SeqExpressionSet-method (SeqExpressionSet-class), 14
offst (SeqExpressionSet-class), 14
offst, SeqExpressionSet-method (SeqExpressionSet-class), 14
offst<- (SeqExpressionSet-class), 14
offst<-, SeqExpressionSet-method (SeqExpressionSet-class), 14
par, 5, 6, 8, 11, 13
plot, BamFileList, FastqFileList-method (plot-methods), 10
plot-methods, 10
plotNtFrequency (plotNtFrequency-methods), 11
plotNtFrequency-methods, 11
plotPCA (plotPCA-methods), 11
plotPCA, matrix-method (plotPCA-methods), 11
plotPCA, SeqExpressionSet-method (plotPCA-methods), 11
plotPCA-methods, 11
plotQuality (plotQuality-methods), 12
plotQuality, BamFile-method (plotQuality-methods), 12
plotQuality, BamFileList-method (plotQuality-methods), 12
plotQuality, FastqFileList-method (plotQuality-methods), 12
plotQuality-methods, 12
plotRLE (plotRLE-methods), 13
plotRLE, matrix-method (plotRLE-methods), 13
plotRLE, SeqExpressionSet-method (plotRLE-methods), 13
plotRLE-methods, 13
Rsamtools, 2
SeqExpressionSet, 2, 4, 8–11, 13, 16, 17
SeqExpressionSet-class, 14
smoothScatter, 15
withinLaneNormalization, 14, 15
withinLaneNormalization (withinLaneNormalization-methods), 16
withinLaneNormalization, matrix, numeric-method (withinLaneNormalization-methods), 16
withinLaneNormalization, SeqExpressionSet, character-method (withinLaneNormalization-methods), 16
withinLaneNormalization-methods, 16
yeastGC, 17
yeastLength, 18