Package ‘scater’
March 29, 2017

Type Package
Maintainer Davis McCarthy <davis@ebi.ac.uk>
Author Davis McCarthy
Version 1.2.0
Date 2016-10-14
License GPL (>= 2)
Title Single-cell analysis toolkit for gene expression data in R
Description A collection of tools for doing various analyses of single-cell RNA-seq gene expression data, with a focus on quality control.
Depends R (>= 3.3), Biobase, ggplot2, methods
Imports biomaRt, BiocGenerics, data.table, dplyr, edgeR, ggbeeswarm, grid, limma, matrixStats, parallel, plyr, reshape2, rhdf5, rjson, shiny, shinydashboard, stats, tximport, viridis
Suggests BiocStyle, cowplot, cluster, destiny, knitr, monocle, mvoutlier, rmarkdown, Rtsne, testthat, magrittr
VignetteBuilder knitr
LazyData true
biocViews SingleCell, RNASEq, QualityControl, Preprocessing, Normalization, Visualization, DimensionReduction, Transcriptomics, GeneExpression, Sequencing, Software, DataImport, DataRepresentation, Infrastructure
RoxygenNote 5.0.1
NeedsCompilation yes
URL https://github.com/davismcc/scater
BugReports https://github.com/davismcc/scater/issues

R topics documented:

  scater-package ........................................... 3
  arrange ................................................. 3
  bootstraps ............................................... 4
  calcIsExprs ........................................... 5
R topics documented:
calculateFPKM ......................................................... 6
calculateQCMetrics .................................................. 6
calculateTPM .......................................................... 9
cellNames< .............................................................. 10
cellPairwiseDistances ................................................. 11
counts ................................................................. 12
cpm ................................................................. 13
data<.SCESet,AnnotatedDataFrame-method ......................... 14
featurePairwiseDistances ............................................. 15
filter ................................................................. 16
findImportantPCs ...................................................... 17
fkkm ................................................................. 18
fromCellDataSet ...................................................... 19
getBMFeatureAnnos ................................................... 20
getExprs .............................................................. 21
get_exprs ............................................................. 22
isOutlier .............................................................. 23
is_exprs .............................................................. 24
mergeSCESet .......................................................... 25
multiplot ............................................................... 26
mutate ................................................................. 27
newSCESet ............................................................. 27
nexprs ................................................................. 29
normaliseExprs ......................................................... 30
normalize ............................................................. 32
norm_counts .......................................................... 33
norm_cpm ............................................................. 34
norm_exprs ............................................................ 35
norm_fkkm ............................................................. 36
norm_tpm .............................................................. 37
pData<.SCESet,AnnotatedDataFrame-method ......................... 38
plot ................................................................. 39
plotDiffusionMap ...................................................... 40
plotExplanatoryVariables ............................................ 43
plotExpression ........................................................ 44
plotExprsFreqVsMean ................................................. 46
plotExprsVsTxLength .................................................. 48
plotFeatureData ....................................................... 49
plotHighestExprs ...................................................... 50
plotMDS ............................................................... 51
plotMetadata .......................................................... 53
plotPCA ............................................................... 54
plotPhenoData ........................................................ 56
plotPlatePosition ..................................................... 57
plotQC ................................................................. 59
plotReducedDim ....................................................... 60
plotTSNE ............................................................. 61
readKallistoResults .................................................. 63
readKallistoResultsOneSample ...................................... 64
readSalmonResults .................................................... 65
readSalmonResultsOneSample ....................................... 66
readTxResults ........................................................ 67
scater-package

Description

scater provides a class and numerous functions for the quality control, normalisation and visualisation of single-cell RNA-seq expression data.

Details

In particular, scater provides easy generation of quality control metrics and simple functions to visualise quality control metrics and their relationships.

arrange

Arrange rows of pData(object) by variables.

Description

The SCESet returned will have cells ordered by the corresponding variable in pData(object).

Usage

arrange(object, ...)

## S4 method for signature 'SCESet'
arrange(object, ...)

arrange.SCESet(object, ...)
Arguments

object  A SCESet object.

...  Additional arguments to be passed to dplyr::arrange to act on pData(object).

Examples

data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
example_sceset <- arrange(example_sceset, Cell_Cycle)

bootstraps  

Accessor and replacement for bootstrap results in an SCESet object

Description

SCESet objects can contain an of bootstrap expression values (for example, as generated by the kallisto software for quantifying feature abundance). These functions conveniently access and replace the ‘bootstrap’ slot with the value supplied, which must be an matrix of the correct size, namely the same number of rows and columns as the SCESet object as a whole.

Usage

bootstraps(object)

bootstraps(object) <- value

bootstraps.SCESet(object)

### S4 method for signature 'SCESet'
bootstraps(object)

### S4 replacement method for signature 'SCESet,array'
bootstraps(object) <- value

Arguments

object  a SCESet object.

value  an array of class "numeric" containing bootstrap expression values

Value

If accessing bootstraps slot of an SCESet, then an array with the bootstrap values, otherwise an SCESet object containing new bootstrap values.

Author(s)

Davis McCarthy
calcIsExprs

Examples

data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
bootstraps(example_sceset)

Description

Calculate which features are expressed in which cells using a threshold on observed counts, transcripts-per-million, counts-per-million, FPKM, or defined expression levels.

Usage

calcIsExprs(object, lowerDetectionLimit = NULL, exprs_data = "counts")

Arguments

object

an SCESet object with expression and/or count data.

lowerDetectionLimit

numeric scalar giving the minimum expression level for an expression observation in a cell for it to qualify as expressed.

exprs_data

character scalar indicating whether the count data ("counts"), the transformed expression data ("exprs"), transcript-per-million ("tpm"), counts-per-million ("cpm") or FPKM ("fpkm") should be used to define if an observation is expressed or not.

Value

a logical matrix indicating whether or not a feature in a particular cell is expressed.

Examples

data("sc_example_counts")
data("sc_example_cell_info")
example_sceset <- newSCESet(countData = sc_example_counts)
is_exprs(example_sceset) <- calcIsExprs(example_sceset, lowerDetectionLimit = 1, exprs_data = "exprs")
calculateFPKM  

Calculate fragments per kilobase of exon per million reads mapped (FPKM)

Description

Calculate fragments per kilobase of exon per million reads mapped (FPKM) values for expression from counts for a set of features.

Usage

calculateFPKM(object, effective_length)

Arguments

object  
an SCESet object

effective_length  
vector of class "numeric" providing the effective length for each feature in the SCESet object

Value

Matrix of FPKM values.

Examples

data("sc_example_counts")
data("sc_example_cell_info")
example_sceset <- newSCESet(countData = sc_example_counts)
effective_length <- rep(1000, 2000)
fpkm(example_sceset) <- calculateFPKM(example_sceset, effective_length)


calculateQCMetrics  

Calculate QC metrics

Description

Calculate QC metrics

Usage

calculateQCMetrics(object, feature_controls = NULL, cell_controls = NULL, nmads = 5, pct_feature_controls_threshold = 80)
**calculateQCMetrics**

**Arguments**

- **object**
  - An SCESet object containing expression values and experimental information. Must have been appropriately prepared.

- **feature_controls**
  - A named list containing one or more vectors (character vector of feature names, logical vector, or a numeric vector of indices are all acceptable) used to identify feature controls (for example, ERCC spike-in genes, mitochondrial genes, etc).

- **cell_controls**
  - A character vector of cell (sample) names, or a logical vector, or a numeric vector of indices used to identify cell controls (for example, blank wells or bulk controls).

- **nmads**
  - A numeric scalar giving the number of median absolute deviations to be used to flag potentially problematic cells based on total_counts (total number of counts for the cell, or library size) and total_features (number of features with non-zero expression). For total_features, cells are flagged for filtering only if total_features is nmads below the median. Default value is 5.

- **pct_feature_controls_threshold**
  - A numeric scalar giving a threshold for percentage of expression values accounted for by feature controls. Used as to flag cells that may be filtered based on high percentage of expression from feature controls.

**Details**

Calculate useful quality control metrics to help with pre-processing of data and identification of potentially problematic features and cells.

The following QC metrics are computed:

- **total_counts**: Total number of counts for the cell (aka “library size”)
- **log10_total_counts**: Total counts on the log10-scale
- **total_features**: The number of endogenous features (i.e. not control features) for the cell that have expression above the detection limit (default detection limit is zero)
- **filter_on_depth**: Would this cell be filtered out based on its log10-depth being (by default) more than 5 median absolute deviations from the median log10-depth for the dataset?
- **filter_on_coverage**: Would this cell be filtered out based on its coverage being (by default) more than 5 median absolute deviations from the median coverage for the dataset?
- **filter_on_pct_counts_feature_controls**: Should the cell be filtered out on the basis of having a high percentage of counts assigned to control features? Default threshold is 80 percent (i.e. cells with more than 80 percent of counts assigned to feature controls are flagged).
- **counts_feature_controls**: Total number of counts for the cell that come from (one or more sets of user-defined) control features. Defaults to zero if no control features are indicated. If more than one set of feature controls are defined (for example, ERCC and MT genes are defined as controls), then this metric is produced for all sets, plus the union of all sets (so here, we get columns counts_feature_controls_ERCC, counts_feature_controls_MT and counts_feature_controls).
- **log10_counts_feature_controls**: Just as above, the total number of counts from feature controls, but on the log10-scale. Defaults to zero (i.e. \(\log_{10}(0 + 1)\), offset to avoid negative infinite values) if no feature control are indicated.
- **pct_counts_feature_controls**: Just as for the counts described above, but expressed as a percentage of the total counts. Defined for all control sets and their union, just like the raw counts. Defaults to zero if no feature controls are defined.
**filter_on_pct_counts_feature_controls**: Would this cell be filtered out on the basis that the percentage of counts from feature controls is higher than a defined threshold (default is 80%)? Just as with `counts_feature_controls`, this is defined for all control sets and their union.

**pct_counts_top_50_features**: What percentage of the total counts is accounted for by the 50 highest-count features? Also computed for the top 100 and top 200 features, with the obvious changes to the column names. Note that the top “X” percentage will not be computed if the total number of genes is less than “X”.

**pct_dropout**: Percentage of features that are not “detectably expressed”, i.e. have expression below the `lowerDetectionLimit` threshold.

**counts_endogenous_features**: Total number of counts for the cell that come from endogenous features (i.e. not control features). Defaults to ‘depth’ if no control features are indicated.

**log10_counts_endogenous_features**: Total number of counts from endogenous features on the log10-scale. Defaults to all counts if no control features are indicated.

**n_detected_feature_controls**: Number of defined feature controls that have expression greater than the threshold defined in the object (that is, they are “detectably expressed”; see `object@lowerDetectionLimit` to check the threshold). As with other metrics for feature controls, defined for all sets of feature controls (set names appended as above) and their union. So we might commonly get columns `n_detected_feature_controls_ERCC`, `n_detected_feature_controls_MT` and `n_detected_feature_controls` (ERCC and MT genes detected).

**is_cell_control**: Has the cell been defined as a cell control? If more than one set of cell controls are defined (for example, blanks and bulk libraries are defined as cell controls), then this metric is produced for all sets, plus the union of all sets (so we could typically get columns `is_cell_control_Blank`, `is_cell_control_Bulk`, and `is_cell_control`, the latter including both blanks and bulks as cell controls).

These cell-level QC metrics are added as columns to the “phenotypeData” slot of the `SCESet` object so that they can be inspected and are readily available for other functions to use. Furthermore, wherever “counts” appear in the above metrics, the same metrics will also be computed for “exprs”, “tpm” and “fpkm” values (if TPM and FPKM values are present in the `SCESet` object), with the appropriate term replacing “counts” in the name. The following feature-level QC metrics are also computed:

**mean_exprs**: The mean expression level of the gene/feature.

**exprs_rank**: The rank of the feature’s mean expression level in the cell.

**n_cells_exprs**: The number of cells for which the expression level of the feature is above the detection limit (default detection limit is zero).

**total_feature_counts**: The total number of counts assigned to that feature across all cells.

**log10_total_feature_counts**: Total feature counts on the log10-scale.

**pct_total_counts**: The percentage of all counts that are accounted for by the counts assigned to the feature.

**pct_dropout**: The percentage of all cells that have no detectable expression (i.e. `is_exprs(object)` is `FALSE`) for the feature.

**is_feature_control**: Is the feature a control feature? Default is ‘FALSE’ unless control features are defined by the user. If more than one feature control set is defined (as above), then a column of this type is produced for each control set (e.g. here, `is_feature_control_ERCC` and `is_feature_control_MT`) as well as the column named `is_feature_control`, which indicates if the feature belongs to any of the control sets.
These feature-level QC metrics are added as columns to the “featureData” slot of the SCESet object so that they can be inspected and are readily available for other functions to use. As with the cell-level metrics, wherever “counts” appear in the above, the same metrics will also be computed for “exprs”, “tpm” and “fpkm” values (if TPM and FPKM values are present in the SCESet object), with the appropriate term replacing “counts” in the name.

Value
an SCESet object

Examples

data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data=sc_example_cell_info)
rownames(pd) <- pd$Cell
example_sceset <- newSCESet(countData=sc_example_counts, phenoData=pd)
example_sceset <- calculateQCMetrics(example_sceset)

## with a set of feature controls defined
example_sceset <- calculateQCMetrics(example_sceset, feature_controls = 1:40)

## with a named set of feature controls defined
example_sceset <- calculateQCMetrics(example_sceset,
    feature_controls = list(ERCC = 1:40))

--
calculateTPM  Calculate transcripts-per-million (TPM)

Description
Calculate transcripts-per-million (TPM) values for expression from counts for a set of features.

Usage
calculateTPM(object, effective_length = NULL, calc_from = "counts")

Arguments

object an SCESet object

effective_length vector of class "numeric" providing the effective length for each feature in the SCESet object

calc_from character string indicating whether to compute TPM from "counts", "norm_counts", "fpkm" or "norm_fpkm". Default is to use "counts", in which case the effective_length argument must be supplied.

Value
Matrix of TPM values.
Examples

data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
exmaple_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
effective_length <- rep(1000, 2000)
tpm(example_sceset) <- calculateTPM(example_sceset, effective_length,
calc_from = "counts")

## calculate from FPKM
fpkm(example_sceset) <- calculateFPKM(example_sceset, effective_length)
tpm(example_sceset) <- calculateTPM(example_sceset, effective_length,
calc_from = "fpkm")

cellNames<- Get or set cell names from an SCESet object

Description

Get or set cell names from an SCESet object

Usage

cellNames(object) <- value
cellNames(object)

## S4 replacement method for signature 'SCESet,vector'
cellNames(object)<-value

Arguments

object An SCESet object.
value a vector of cell names to apply to the SCESet object.

Details

Simply a wrapper to sampleNames.

Value

A vector of cell names.

Author(s)

Davis McCarthy
Examples

data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
cellNames(example_sceset)

data("sc_example_counts")
data("sc_example_cell_info")
example_sceset <- newSCESet(countData = sc_example_counts)
cellNames(example_sceset) <- 1:ncol(example_sceset)

cellPairwiseDistances

**cellPairwiseDistances in an SCESet object**

Description

SCESet objects can contain a matrix of pairwise distances between cells. These functions conveniently access and replace the cell pairwise distances with the value supplied, which must be a matrix of the correct size. The function cellDist is simply shorthand for cellPairwiseDistances.

Usage

cellPairwiseDistances(object)
cellPairwiseDistances(object) <- value
cellDist(object)
cellDist(object) <- value
cellPairwiseDistances.SCESet(object)
## S4 method for signature 'SCESet'
cellPairwiseDistances(object)
cellDistSCESet(object)
## S4 method for signature 'SCESet'
cellDist(object)
## S4 method for signature 'SCESet'
cellDist(object)
## S4 replacement method for signature 'SCESet,matrix'
cellPairwiseDistances(object) <- value
## S4 replacement method for signature 'SCESet,dist'
cellPairwiseDistances(object) <- value
## S4 replacement method for signature 'SCESet,matrix'
cellDist(object) <- value
## S4 replacement method for signature 'SCESet,dist'
cellDist(object) <- value
counts

Arguments

object a SCESet object.
value a matrix of class "numeric" containing cell pairwise distances

Value

An SCESet object containing new cell pairwise distances matrix.

Author(s)

Davis McCarthy

Examples

data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
cellPairwiseDistances(example_sceset)

counts Accessors for the 'counts' element of an SCESet object.

Description

The counts element holds the count data as a matrix of non-negative integer count values, one row for each feature (gene, exon, region, etc), and one column for each cell. It is an element of the assayData slot of the SCESet object.

Usage

## S4 method for signature 'SCESet'
counts(object)

## S4 replacement method for signature 'SCESet,matrix'
counts(object) <- value

## S4 method for signature 'SCESet'
counts(object)

## S4 replacement method for signature 'SCESet,matrix'
counts(object) <- value

Arguments

object a SCESet object.
value an integer matrix

Author(s)

Davis McCarthy
**Examples**

```r
data("sc_example_counts")
data("sc_example_cell_info")
example_sceset <- newSCESet(countData = sc_example_counts)
counts(example_sceset)
```

<table>
<thead>
<tr>
<th>cpm</th>
<th>Accessors for the 'cpm' (counts per million) element of an SCESet object.</th>
</tr>
</thead>
</table>

**Description**

The `cpm` element of the arrayData slot in an SCESet object holds a matrix containing counts-per-million values. It has the same dimensions as the `exprs` and `counts` elements, which hold the transformed expression data and count data, respectively.

**Usage**

```r
cpm(object)
```

```r
cpm(object) <- value
```

```r
## S4 method for signature 'SCESet'
cpm(object)
```

```r
## S4 replacement method for signature 'SCESet, matrix'
cpm(object) <- value
```

```r
## S4 method for signature 'SCESet'
cpm(object)
```

```r
## S4 replacement method for signature 'SCESet, matrix'
cpm(object) <- value
```

**Arguments**

- `object` a SCESet object.
- `value` a matrix of class "numeric"

**Value**

a matrix of counts-per-million values

**Author(s)**

Davis McCarthy
Examples

data("sc_example_counts")
data("sc_example_cell_info")
example_sceset <- newSCESet(countData = sc_example_counts)
cpm(example_sceset)[1:10, 1:6]

Description

SCESet objects contain feature information (inherited from the ExpressionSet class). This function conveniently replaces the feature data with the value supplied, which must be an AnnotatedDataFrame.

Usage

## S4 replacement method for signature 'SCESet,AnnotatedDataFrame'
fData(object) <- value

## S4 replacement method for signature 'SCESet,data.frame'
fData(object) <- value

Arguments

object An SCESet object.
value an AnnotatedDataFrame with updated featureData to replace existing

Value

A matrix of expression count data, where rows correspond to features (e.g. genes) and columns correspond to cells.

Examples

## Not run:
data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
fData(example_sceset)

## End(Not run)
**featurePairwiseDistances**

*featurePairwiseDistances in an SCESet object*

**Description**

SCESet objects can contain a matrix of pairwise distances between features (e.g. genes, transcripts). These functions conveniently access and replace the gene pairwise distances with the value supplied, which must be a matrix of the correct size. The function `featDist` is simply shorthand for `featurePairwiseDistances`.

**Usage**

```r
featurePairwiseDistances(object)

featurePairwiseDistances(object) <- value

featDist(object)

featDist(object) <- value

featurePairwiseDistancesSCESet(object)

## S4 method for signature 'SCESet'
featurePairwiseDistances(object)

featDistSCESet(object)

## S4 method for signature 'SCESet'
featDist(object)

## S4 replacement method for signature 'SCESet,matrix'
featurePairwiseDistances(object) <- value

## S4 replacement method for signature 'SCESet,dist'
featurePairwiseDistances(object) <- value

## S4 replacement method for signature 'SCESet,matrix'
featDist(object) <- value

## S4 replacement method for signature 'SCESet,dist'
featDist(object) <- value
```

**Arguments**

- `object` a SCESet object.
- `value` a matrix of class "numeric" containing feature pairwise distances

**Value**

An SCESet object containing new feature pairwise distances matrix.
**filter**

Return SCESet with cells matching conditions.

---

**Description**

Subsets the columns (cells) of a SCESet based on matching conditions in the rows of pData(object).

**Usage**

```r
filter(object, ...)
```

## S4 method for signature 'SCESet'

```r
filter(object, ...)
```

```r
filter.SCESet(object, ...)
```

**Arguments**

- `object` A SCESet object.
- `...` Additional arguments to be passed to dplyr::filter to act on pData(object).

**Examples**

```r
data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
exmaple_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
featurePairwiseDistances(example_sceset)

example_sceset_treat1 <- filter(example_sceset, Treatment == "treat1")
```
findImportantPCs

Find most important principal components for a given variable

Description

Find most important principal components for a given variable

Usage

findImportantPCs(object, variable = "total_features",
plot_type = "pcs-vs-vars", exprs_values = "exprs", ntop = 500,
feature_set = NULL, scale_features = TRUE, theme_size = 10)

Arguments

object an SCESet object containing expression values and experimental information. Must have been appropriately prepared.
variable character scalar providing a variable name (column from pData(object)) for which to determine the most important PCs.
plot_type character string, indicating which type of plot to produce. Default, "pairs-pcs" produces a pairs plot for the top 5 PCs based on their R-squared with the variable of interest. A value of "pcs-vs-vars" produces plots of the top PCs against the variable of interest.
exprs_values which slot of the assayData in the object should be used to define expression? Valid options are "counts" (default), "tpm", "fpkm" and "exprs", or anything else in the object added manually by the user.
ntop numeric scalar indicating the number of most variable features to use for the PCA. Default is 500, but any ntop argument is overridden if the feature_set argument is non-NULL.
feature_set character, numeric or logical vector indicating a set of features to use for the PCA. If character, entries must all be in featureNames(object). If numeric, values are taken to be indices for features. If logical, vector is used to index features and should have length equal to nrow(object).
scale_features logical, should the expression values be standardised so that each feature has unit variance? Default is TRUE.
theme_size numeric scalar providing base font size for ggplot theme.

Details

Plot the top 5 or 6 most important PCs (depending on the plot_type argument for a given variable. Importance here is defined as the R-squared value from a linear model regressing each PC onto the variable of interest.

Value

a ggplot plot object
Examples

data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
rownames(pd) <- pd$Cell
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
drop_genes <- apply(exprs(example_sceset), 1, function(x) {var(x) == 0})
example_sceset <- example_sceset[!drop_genes, ]
example_sceset <- calculateQCMetrics(example_sceset)
findImportantPCs(example_sceset, variable="total_features")

fpkm

Accessors for the 'fpkm' (fragments per kilobase of exon per million reads mapped) element of an SCESet object.

Description

The fpkm element of the arrayData slot in an SCESet object holds a matrix containing fragments per kilobase of exon per million reads mapped (FPKM) values. It has the same dimensions as the 'exprs' and 'counts' elements, which hold the transformed expression data and count data, respectively.

Usage

fpkm(object)

fpkm(object) <- value

## S4 method for signature 'SCESet'
fpkm(object)

## S4 replacement method for signature 'SCESet,matrix'
fpkm(object) <- value

## S4 method for signature 'SCESet'
fpkm(object)

## S4 replacement method for signature 'SCESet,matrix'
fpkm(object) <- value

Arguments

object a SCESet object.
value a matrix of class "numeric"

Value

a matrix of FPKM values

Author(s)

Davis McCarthy
Examples

```r
data("sc_example_counts")
data("sc_example_cell_info")
example_sceset <- newSCESet(countData = sc_example_counts)
fpkm(example_sceset)
```

Description

Convert a `CellDataSet` to an `SCESet`

Usage

```r
fromCellDataSet(cds, exprs_values = "tpm", logged = FALSE)
```

Arguments

- **cds**
  - A `CellDataSet` from the `monocle` package
- **exprs_values**
  - What should `exprs(cds)` be mapped to in the `SCESet`? Should be one of "exprs", "tpm", "fpkm", "counts"
- **logged**
  - logical, if a value is supplied for the exprsData argument, are the expression values already on the log2 scale, or not?

Value

An object of class `SCESet`

Examples

```r
data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
if (requireNamespace("monocle")) {
  # cds <- toCellDataset(example_sceset) # not run
  # sceset <- fromCellDataSet(cds) # not run
}
```
getBMFeatureAnnos

Get feature annotation information from Biomart

Description

Use the bioMaRt package to add feature annotation information to an SCESet.

Usage

getBMFeatureAnnos(object, filters = "ensembl_transcript_id",
attributes = c("ensembl_transcript_id", "ensembl_gene_id", "mgi_symbol",
"chromosome_name", "transcript_biotype", "transcript_start", "transcript_end",
"transcript_count"), feature_symbol = "mgi_symbol",
feature_id = "ensembl_gene_id", biomart = "ENSEMBL_MART_ENSEMBL",
dataset = "mmusculus_gene_ensembl", host = "www.ensembl.org")

Arguments

object an SCESet object
filters character vector defining the "filters" terms to pass to the bioMaRt::getBM function.
attributes character vector defining the bioMaRt attributes to pass to the attributes argument of getBM.
feature_symbol character string defining the bioMaRt attribute to be used to define the symbol to be used for each feature (which appears as the feature_symbol in fData(object), subsequently). Default is "mgi_symbol", gene symbols for mouse. This should be changed if the organism is not Mus musculus!
feature_id character string defining the bioMaRt attribute to be used to define the ID to be used for each feature (which appears as the feature_id in fData(object), subsequently). Default is "ensembl_gene_id", Ensembl gene IDs for mouse. This should be changed if the organism is not Mus musculus!
biomart character string defining the bioMaRt to be used. Default is "ENSEMBL_MART_ENSEMBL".
dataset character string defining the bioMaRt dataset to use. Default is "mmusculus_gene_ensembl", which should be changed if the organism is not the mouse!
host optional character string argument which can be used to select a particular "host" from bioMaRt to use. Useful for accessing archived versions of bioMaRt data. Default is "www.ensembl.org", in which case the current version of the bioMaRt (now hosted by Ensembl) is used.

Details

See the documentation for the bioMaRt package, specifically for the functions useMart and getBM, for information on what are permitted values for the filters, attributes, biomart, dataset and host arguments.

Value

an SCESet object
getExprs

Examples

## Not run:
object <- getBMFeatureAnnos(object)

## End(Not run)

text

getExprs

Retrieve a representation of gene expression

Description

Gene expression can be summarised in a variety of ways, e.g. as TPM, FPKM or as raw counts. Many internal methods and external packages rely on accessing a generic representation of expression without worrying about the particulars. Scater allows the user to set `object@useForExprs` to the preferred type (either "exprs", "TPM", "fpkm" or "counts") and that particular representation will be returned by calls to `getExprs`. Note if such representation is not defined, this method returns `NULL`.

Usage

getExprs(object)

Arguments

object  
An object of type SCESet

Value

A matrix representation of expression corresponding to `object@useForExprs`.

Examples

data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd, useForExprs = "exprs")
all(exprs(example_sceset) == getExprs(example_sceset)) # TRUE
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd, useForExprs = "counts")
all(exprs(example_sceset) == getExprs(example_sceset)) # FALSE
all(counts(example_sceset) == getExprs(example_sceset)) # TRUE
get_exprs

**Generic accessor for expression data from an SCESet object.**

**Description**

Access by name a matrix of expression values, one row for each feature (gene, exon, region, etc), and one column for each cell stored an element of the assayData slot of the SCESet object.

**Usage**

```r
get_exprs(object, exprs_values)
```

## S4 method for signature 'SCESet'

```r
get_exprs(object, exprs_values)
```

## S4 method for signature 'SCESet'

```r
get_exprs(object, exprs_values = "exprs")
```

**Arguments**

- `object` a SCESet object.
- `exprs_values` character string indicating which values should be used as the expression values for this plot. Valid arguments are "tpm" (default; transcripts per million), "norm_tpm" (normalised TPM values), "fpkm" (FPKM values), "norm_fpkm" (normalised FPKM values), "counts" (counts for each feature), "norm_counts", "cpm" (counts-per-million), "norm_cpm" (normalised counts-per-million), "exprs" (whatever is in the ‘exprs’ slot of the SCESet object; default), "norm_exprs" (normalised expression values) or "stand_exprs" (standardised expression values) or any other slots that have been added to the "assayData" slot by the user.

**Value**

a matrix of expression values

**Author(s)**

Davis McCarthy

**Examples**

```r
data("sc_example_counts")
data("sc_example_cell_info")
example_sceset <- newSCESet(countData = sc_example_counts)
get_exprs(example_sceset, "counts")

## new slots can be defined and accessed
set_exprs(example_sceset, "scaled_counts") <- t(t(counts(example_sceset)) / colSums(counts(example_sceset)))
get_exprs(example_sceset, "scaled_counts"[1:6, 1:6])
```
isOutlier

Identify if a cell is an outlier based on a metric

Description

Convenience function to determine which values for a metric are outliers based on median-absolute-deviation (MAD).

Usage

isOutlier(metric, nmads = 5, type = c("both", "lower", "higher"),
    log = FALSE, na.rm = FALSE)

Arguments

metric numeric or integer vector of values for a metric
nmads scalar, number of median-absolute-deviations away from median required for a
    value to be called an outlier
type character scalar, choice indicate whether outliers should be looked for at both
tails (default: "both") or only at the lower end ("lower") or the higher end ("higher")
log logical, should the values of the metric be transformed to the log10 scale before
    computing median-absolute-deviation for outlier detection?
na.rm logical, should NA (missing) values be removed before computing median and
    median-absolute-deviation values? If FALSE then return values for median and
    median-absolute-deviation will be NA if any value is NA.

Value

a logical vector of the same length as the metric argument

Examples

data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data=sc_example_cell_info)
rownames(pd) <- pd$Cell
example_sceset <- newSCESet(countData=sc_example_counts, phenoData=pd)
example_sceset <- calculateQCMetrics(example_sceset)
## with a set of feature controls defined
example_sceset <- calculateQCMetrics(example_sceset, feature_controls = 1:40)
isOutlier(example_sceset$total_counts, nmads = 3)
Description

The `is_exprs` element holds a logical matrix indicating whether or not each observation is above the defined lowerDetectionLimit in the SCESet object. It has the same dimensions as the `exprs` and `counts` elements, which hold the transformed expression data and count data, respectively.

Usage

```r
is_exprs(object)

is_exprs(object) <- value

## S4 method for signature 'SCESet'
is_exprs(object)

## S4 replacement method for signature 'SCESet,matrix'
is_exprs(object) <- value

## S4 method for signature 'SCESet'
is_exprs(object)

## S4 replacement method for signature 'SCESet,matrix'
is_exprs(object) <- value
```

Arguments

- `object`: a SCESet object.
- `value`: an integer matrix

Value

A logical matrix indicating if observations are "expressed" or not

Author(s)

Davis McCarthy

Examples

```r
data("sc_example_counts")
data("sc_example_cell_info")
example_sceset <- newSCESet(countData = sc_example_counts)
is_exprs(example_sceset)
```
mergeSCESet  Merge SCESet objects

Description
Merge two SCESet objects that have the same features but contain different cells/samples.

Usage
mergeSCESet(x, y, fdata_cols_x = 1:ncol(fData(x)),
fdata_cols_y = fdata_cols_x, pdata_cols_x = NULL, pdata_cols_y = NULL)

Arguments
x  an SCESet object
y  an SCESet object
fdata_cols_x  a logical or numeric vector indicating which columns of featureData for x are
shared between x and y and should feature in the returned merged SCESet. Default is all columns of fData(x).
fdata_cols_y  a logical or numeric vector indicating which columns of featureData for y are
shared between x and y and should feature in the returned merged SCESet. Default is fdata_cols_x.
pdata_cols_x  a logical or numeric vector indicating which columns of phenoData of x should
be retained.
pdata_cols_y  a logical or numeric vector indicating which columns of phenoData of y should
be retained.

Details
Existing cell-cell pairwise distances and feature-feature pairwise distances will not be valid for a
merged SCESet so these are set to NULL in the returned object. Similarly experimentData will
need to be added anew to the merged SCESet returned.

Value
a merged SCESet object combining data and metadata from x and y

Examples
data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
mergeSCESet(example_sceset[, 1:20], example_sceset[, 21:40])

## with specification of columns of fData
example_sceset <- calculateQCMetrics(example_sceset)
mergeSCESet(example_sceset[, 1:20], example_sceset[, 21:40], fdata_cols_x = c(1, 7))

## with specification of columns of pData
mergeSCESet(example_sceset[, 1:20], example_sceset[, 21:40], pdata_cols_x = 1:6)
multiplot

Multiple plot function for ggplot2 plots

Description

Place multiple `ggplot` plots on one page.

Usage

```r
multiplot(..., plotlist = NULL, cols = 1, layout = NULL)
```

Arguments

- `...`: `ggplot` objects can be passed in ... or to `plotlist` (as a list of `ggplot` objects)
- `plotlist`: a list of `ggplot` objects
- `cols`: numeric scalar giving the number of columns in the layout
- `layout`: a matrix specifying the layout. If present, `cols` is ignored.

Details

If the layout is something like `matrix(c(1,2,3,3), nrow=2, byrow=TRUE)`, then plot 1 will go in the upper left, 2 will go in the upper right, and 3 will go all the way across the bottom. There is no way to tweak the relative heights or widths of the plots with this simple function. It was adapted from [http://www.cookbook-r.com/Graphs/Multiple_graphs_on_one_page_(ggplot2)/](http://www.cookbook-r.com/Graphs/Multiple_graphs_on_one_page_(ggplot2)/)

Value

a `ggplot` plot object

Examples

```r
library(ggplot2)
## This example uses the ChickWeight dataset, which comes with ggplot2
## First plot
p1 <- ggplot(ChickWeight, aes(x = Time, y = weight, colour = Diet, group = Chick)) +
  geom_line() +
  ggtitle("Growth curve for individual chicks")
## Second plot
p2 <- ggplot(ChickWeight, aes(x = Time, y = weight, colour = Diet)) +
  geom_point(alpha = .3) +
  geom_smooth(alpha = .2, size = 1) +
  ggtitle("Fitted growth curve per diet")
## Third plot
p3 <- ggplot(subset(ChickWeight, Time == 21), aes(x = weight, colour = Diet)) +
  geom_density() +
  ggtitle("Final weight, by diet")
## Fourth plot
p4 <- ggplot(subset(ChickWeight, Time == 21), aes(x = weight, fill = Diet)) +
  geom_histogram(colour = "black", binwidth = 50) +
  facet_grid(Diet ~ .) +
  ggtitle("Final weight, by diet") +
```
mutate

Add new variables to pData(object).

Description

Adds new columns to pData(object) preserving existing variables.

Usage

mutate(object, ...)

## S4 method for signature 'SCESet'
mutate(object, ...)

mutate.SCESet(object, ...)

Arguments

object

A SCESet object.

...  

Additional arguments to be passed to dplyr::mutate to act on pData(object).

Examples

data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
example_sceset <- mutate(example_sceset, is_quiescent = Cell_Cycle == "G0")

newSCESet

Create a new SCESet object.

Description

Scater requires that all data be housed in SCESet objects. SCESet extends Bioconductor’s ExpressionSet class, and the same basic interface is supported. newSCESet() expects a matrix of expression values as its first argument, with rows as features (usually genes) and columns as cells. Per-feature and per-cell metadata can be supplied with the featureData and phenoData arguments, respectively. Use of these optional arguments is strongly encouraged. The SCESet also includes a slot ‘counts’ to store an object containing raw count data.
newSCESet

Usage

newSCESet(exprsData = NULL, countData = NULL, tpmData = NULL,
          fpkmData = NULL, cpmData = NULL, phenoData = NULL, featureData = NULL,
          experimentData = NULL, is_exprsData = NULL,
          cellPairwiseDistances = dist(vector()),
          featurePairwiseDistances = dist(vector()), lowerDetectionLimit = 0,
          logExprsOffset = 1, logged = FALSE, useForExprs = "exprs")

Arguments

eqrsData expression data matrix for an experiment
countData data matrix containing raw count expression values
tpmData matrix of class "numeric" containing transcripts-per-million (TPM) expression values
fpkmData matrix of class "numeric" containing fragments per kilobase of exon per million reads mapped (FPKM) expression values
cpmData matrix of class "numeric" containing counts per million (CPM) expression values (optional)
phenoData data frame containing attributes of individual cells
featureData data frame containing attributes of features (e.g. genes)
experimentData MIAME class object containing metadata data and details about the experiment and dataset.
is_exprsData matrix of class "logical", indicating whether or not each observation is above the lowerDetectionLimit.
cellPairwiseDistances object of class "dist" (or a class that extends "dist") containing cell-cell distance or dissimilarity values.
featurePairwiseDistances object of class "dist" (or a class that extends "dist") containing feature-feature distance or dissimilarity values.
lowerDetectionLimit the minimum expression level that constitutes true expression (defaults to zero and uses count data to determine if an observation is expressed or not).
logExprsOffset numeric scalar, providing the offset used when doing log2-transformations of expression data to avoid trying to take logs of zero. Default offset value is 1.
logged logical, if a value is supplied for the exprsData argument, are the expression values already on the log2 scale, or not?
useForExprs character string, either 'exprs' (default), 'tpm', 'counts' or 'fpkm' indicating which expression representation both internal methods and external packages should use when performing analyses.

Details

SCESet objects store a matrix of expression values. These values are typically transcripts-per-million (tpm), counts-per-million (cpm), fragments per kilobase per million mapped (FPKM) or some other output from a program that calculates expression values from RNA-Seq reads. We recommend that expression values on the log2 scale are used for the 'exprs' slot in the SCESet. For example, you may wish to store raw tpm values in the 'tpm' slot and log2(tpm + 1) values
in the 'exprs' slot. However, expression values could also be values from a single cell qPCR run or some other type of assay. The newSCESet function can also accept raw count values. In this case see calculateTPM and calculateFPKM for computing TPM and FPKM expression values, respectively, from counts. The function cpm from the package edgeR to can be used to compute log2(counts-per-million), if desired.

An SCESet object has to have the 'exprs' slot defined, so if the exprsData argument is NULL, then this function will define 'exprs' with the following order of precedence: log2(TPM + logExprsOffset), if tpmData is defined; log2(FPKM + logExprsOffset) if fpkmData is defined; otherwise log2(counts-per-million + logExprsOffset) are used. The cpm function from the edgeR package is used to compute cpm. Note that for many analyses counts-per-million are not recommended, and if possible transcripts-per-million should be used.

In many downstream functions you will likely find it most convenient if the 'exprs' values are on the log2-scale, so this is recommended.

Value

a new SCESet object

Examples

data("sc_example_counts")
data("sc_example_cell_info")

pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
example_sceset

df <- as.data.frame(example_sceset)

nexprs

Count the number of expressed genes per cell

Description

An efficient internal function that avoids the need to construct 'is_exprs_mat' by counting the number of expressed genes per cell on the fly.

Usage

nexprs(object, threshold = NULL, subset.row = NULL, byrow = FALSE)

Arguments

object an SCESet object
threshold numeric scalar providing the value above which observations are deemed to be expressed. Defaults to object@lowerDetectionLimit.
subset.row logical or character vector indicating which rows (i.e. features/genes) to subset and calculate 'is_exprs_mat' for.
byrow logical scalar indicating if TRUE to count expressing cells per feature (i.e. gene) and if FALSE to count expressing features (i.e. genes) per cell.

Value

a numeric vector of the same length as the number of features if byrow argument is TRUE and the same length as the number of cells if byrow is FALSE
normaliseExprs

Normalise expression expression levels for an SCESet object

Description

Compute normalised expression values from an SCESet object and return the object with the normalised expression values added.

Usage

normaliseExprs(object, method = "none", design = NULL, feature_set = NULL, exprs_values = "counts", return_norm_as_exprs = TRUE, ...)

normalizeExprs(...)

Arguments

object an SCESet object.
method character string giving method to be used to calculate normalisation factors. Passed to calcNormFactors.
design design matrix defining the linear model to be fitted to the normalised expression values. If not NULL, then the residuals of this linear model fit are used as the normalised expression values.
feature_set character, numeric or logical vector indicating a set of features to use for the PCA. If character, entries must all be in featureNames(object). If numeric, values are taken to be indices for features. If logical, vector is used to index features and should have length equal to nrow(object).
exprs_values character string indicating which slot of the assayData from the SCESet object should be used as expression values. Valid options are 'counts', the count values, 'exprs' the expression slot, 'tpm' the transcripts-per-million slot or 'fpkm' the FPKM slot.
return_norm_as_exprs logical, should the normalised expression values be returned to the exprs slot of the object? Default is TRUE. If FALSE, values in the exprs slot will be left untouched. Regardless, normalised expression values will be returned to the norm_exprs slot of the object.
...

Examples

data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data=sc_example_cell_info)
rownames(pd) <- pd$Cell
example_sceset <- newSCESet(countData=sc_example_counts, phenoData=pd)
nexprs(example_sceset)[1:10]
nexprs(example_sceset, byrow = TRUE)[1:10]
normaliseExprs

Details

This function allows the user to compute normalised expression values from an SCESet object. The 'raw' values used can be the values in the 'counts' (default), 'exprs', 'tpm' or 'fpkm' slot of the SCESet. Normalised expression values are added to the 'norm_exprs' slot of the object. Normalised expression values are on the log2-scale, with an offset defined by the logExprsOffset slot of the SCESet object. If the 'exprs_values' argument is one of 'counts', 'tpm' or 'fpkm', then a corresponding slot with normalised values is added: 'norm_counts', 'norm_tpm' or 'norm_fpkm', as appropriate. If 'exprs_values' argument is 'counts' a 'norm_cpm' slot is also added, containing normalised counts-per-million values.

Normalisation is done relative to a defined feature set, if desired, which defines the 'library size' by which expression values are divided. If no feature set is defined, then all features are used. A normalisation size factor can be computed (optionally), which internally uses calcNormFactors. Thus, any of the methods available for calcNormFactors can be used: "TMM", "RLE", "upperquartile" or "none". See that function for further details. Library sizes are multiplied by size factors to obtain a 'normalised library size' before normalisation.

If the user wishes to remove the effects of certain explanatory variables, then the 'design' argument can be defined. The design argument must be a valid design matrix, for example as produced by model.matrix, with the relevant variables. A linear model is then fitted using lmFit on expression values after any size-factor and library size normalisation as described above. The returned normalised expression values are then the residuals from the linear model fit.

After normalisation, normalised expression values can be accessed with the norm_exprs function (with corresponding accessor functions for counts, tpm, fpkm, cpm). These functions can also be used to assign normalised expression values produced with external tools to an SCESet object.

normalizeExprs is exactly the same as normaliseExprs, provided for those who prefer North American spelling.

Value

an SCESet object

Author(s)

Davis McCarthy

Examples

data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
keep_gene <- rowSums(counts(example_sceset)) > 0
example_sceset <- example_sceset[keep_gene,]

## Apply TMM normalisation taking into account all genes
example_sceset <- normaliseExprs(example_sceset, method = "TMM")

## Scale counts relative to a set of control features (here the first 100 features)
example_sceset <- normaliseExprs(example_sceset, method = "none",
feature_set = 1:100)
normalize

Normalize an SCESet object using pre-computed size factors

Description

Compute normalised expression values from an SCESet object using the size factors stored in the object. Return the object with the normalised expression values added.

Usage

normalize.SCESet(object, exprs_values = "counts", logExprsOffset = NULL, recompute_cpm = TRUE, return_norm_as_exprs = TRUE)

## S4 method for signature 'SCESet'
normalize(object, exprs_values = "counts",
logExprsOffset = NULL, recompute_cpm = TRUE, return_norm_as_exprs = TRUE)

normalise(...)

Arguments

object an SCESet object.

exprs_values character string indicating which slot of the assayData from the SCESet object should be used as expression values. Valid options are 'counts', the count values, 'exprs' the expression slot, 'tpm' the transcripts-per-million slot or 'fpkm' the FPKM slot.

logExprsOffset scalar numeric value giving the offset to add when taking log2 of normalised values to return as expression values. If NULL (default), then the value from object@logExprsOffset is used.

recompute_cpm logical, should the counts-per-million values be recomputed after normalising with the stored size factors in the object and stored in cpm(object) in the returned object?

return_norm_as_exprs logical, should the normalised expression values be returned to the exprs slot of the object? Default is TRUE. If FALSE, values in the exprs slot will be left untouched. Regardless, normalised expression values will be returned in the norm_exprs(object) slot.

... arguments passed to normalize when calling normalise.

Details

normalize is exactly the same as normalise, the option provided for those who have a preference for North American or British/Australian spelling.

Value

an SCESet object
norm_counts

Author(s)

Davis McCarthy and Aaron Lun

Examples

data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
keep_gene <- rowSums(counts(example_sceset)) > 0
example_sceset <- example_sceset[keep_gene,]

## Apply TMM normalisation taking into account all genes
example_sceset <- normaliseExprs(example_sceset, method = "TMM")
## Scale counts relative to a set of control features (here the first 100 features)
example_sceset <- normaliseExprs(example_sceset, method = "none",
feature_set = 1:100)

## normalize the object using the saved size factors
example_sceset <- normalize(example_sceset)

norm_counts Accessors for the 'norm_counts' element of an SCESet object.

Description

The norm_counts element holds normalised count data as a matrix of non-negative values, one row for each feature (gene, exon, region, etc), and one column for each cell. It is an element of the assayData slot of the SCESet object.

Usage

norm_counts(object)
norm_counts(object) <- value

## S4 method for signature 'SCESet'
norm_counts(object)

## S4 replacement method for signature 'SCESet,matrix'
norm_counts(object) <- value

## S4 method for signature 'SCESet'
norm_counts(object)

## S4 replacement method for signature 'SCESet,matrix'
norm_counts(object) <- value

Arguments

object a SCESet object.
value an integer matrix
norm_cpm

Value
a matrix of normalised count data

Author(s)
Davis McCarthy

Examples
data("sc_example_counts")
data("sc_example_cell_info")
example_sceset <- newSCESet(countData = sc_example_counts)
norm_counts(example_sceset)

norm_cpm

Accessors for the 'norm_cpm' (normalised counts per million) element of an SCESet object.

Description
The norm_cpm element of the arrayData slot in an SCESet object holds a matrix containing normalised counts-per-million values. It has the same dimensions as the 'exprs' and 'counts' elements, which hold the transformed expression data and count data, respectively.

Usage
norm_cpm(object)
norm_cpm(object) <- value

## S4 method for signature 'SCESet'
norm_cpm(object)

## S4 replacement method for signature 'SCESet,matrix'
norm_cpm(object) <- value

## S4 method for signature 'SCESet'
norm_cpm(object)

## S4 replacement method for signature 'SCESet,matrix'
norm_cpm(object) <- value

Arguments

object a SCESet object.
value a matrix of class "numeric"

Value
a matrix of normalised counts-per-million data
Examples

data("sc_example_counts")
data("sc_example_cell_info")
example_sceset <- newSCESet(countData=sc_example_counts)
norm_cpm(example_sceset)


norm_exprs  Accessors for the 'norm_exprs' (normalised expression) element of an SCESet object.

Description

The `norm_exprs` element of the `arrayData` slot in an SCESet object holds a matrix containing normalised expression values. It has the same dimensions as the 'exprs' and 'counts' elements, which hold the transformed expression data and count data, respectively.

Usage

norm_exprs(object)
norm_exprs(object) <- value

### S4 method for signature 'SCESet'
norm_exprs(object)

### S4 replacement method for signature 'SCESet, matrix'
norm_exprs(object) <- value

### S4 method for signature 'SCESet'
norm_exprs(object)

### S4 replacement method for signature 'SCESet, matrix'
norm_exprs(object) <- value

Arguments

object  a SCESet object.
value   an integer matrix

Details

The default for normalised expression values is mean-centred and variance-standardised expression data from the `exprs` slot of the SCESet object. The function `normaliseExprs` (or `normalizeExprs`) provides more options and functionality for normalising expression data.
Value

a matrix of normalised expression data

Author(s)

Davis McCarthy

Examples

data("sc_example_counts")
data("sc_example_cell_info")
example_sceset <- newSCESet(countData = sc_example_counts)
norm_exprs(example_sceset)

Description

The `norm_fpkm` element of the arrayData slot in an SCESet object holds a matrix containing normalised fragments per kilobase of exon per million reads mapped (FPKM) values. It has the same dimensions as the 'exprs' and 'counts' elements, which hold the transformed expression data and count data, respectively.

Usage

```r
norm_fpkm(object)
norm_fpkm(object) <- value

## S4 method for signature 'SCESet'
norm_fpkm(object)

## S4 replacement method for signature 'SCESet,matrix'
norm_fpkm(object) <- value

## S4 method for signature 'SCESet'

## S4 replacement method for signature 'SCESet,matrix'
norm_fpkm(object) <- value
```

Arguments

- **object**: a SCESet object.
- **value**: a matrix of class "numeric"

Value

a matrix of normalised FPKM data
norm_tpm

Author(s)
Davis McCarthy

Examples
data("sc_example_counts")
data("sc_example_cell_info")
example_sceset <- newSCESet(countData = sc_example_counts)
norm_fpkm(example_sceset)

norm_tpm

Accessors for the 'norm_tpm' (transcripts per million) element of an SCESet object.

Description
The norm_tpm element of the arrayData slot in an SCESet object holds a matrix containing normalised transcripts-per-million values. It has the same dimensions as the 'exprs' and 'counts' elements, which hold the transformed expression data and count data, respectively.

Usage
norm_tpm(object)
norm_tpm(object) <- value

## S4 method for signature 'SCESet'
norm_tpm(object)

## S4 replacement method for signature 'SCESet,matrix'
norm_tpm(object) <- value

Arguments
object a SCESet object.
value a matrix of class "numeric"

Value
a matrix of normalised transcripts-per-million data

Author(s)
Davis McCarthy
Examples

data("sc_example_counts")
data("sc_example_cell_info")
example_sceset <- newSCESet(countData = sc_example_counts)
norm_tpm(example_sceset)

Description

SCESet objects contain phenotype information (inherited from the ExpressionSet class). This function conveniently replaces the phenotype data with the value supplied, which must be an AnnotatedDataFrame.

Usage

## S4 replacement method for signature 'SCESet,AnnotatedDataFrame'
pData(object) <- value

## S4 replacement method for signature 'SCESet,data.frame'
pData(object) <- value

Arguments

object An SCESet object.
value an AnnotatedDataFrame with updated phenoData to replace existing

Value

A matrix of expression count data, where rows correspond to features (e.g. genes) and columns correspond to cells.

Examples

## Not run:
data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
pData(example_sceset)

## End(Not run)
**Description**

Plot the relative proportion of the library accounted for by the most highly expressed features for each cell for an SCESet dataset.

**Usage**

```r
## S4 method for signature 'SCESet,ANY'
plot(x, y, ...)

plotSCESet(x, block1 = NULL, block2 = NULL, colour_by = NULL,
          nfeatures = 500, exprs_values = "tpm", ncol = 3, linewidth = 1.5,
          theme_size = 10)
```

**Arguments**

- `x`: an SCESet object
- `y`: optional argument for generic `plot` functions, not used for plotting an SCESet object
- `...`: arguments passed to `plotSCESet`
- `block1`: character string defining the column of `pData(object)` to be used as a factor by which to separate the cells into blocks (separate panels) in the plot. Default is `NULL`, in which case there is no blocking.
- `block2`: character string defining the column of `pData(object)` to be used as a factor by which to separate the cells into blocks (separate panels) in the plot. Default is `NULL`, in which case there is no blocking.
- `colour_by`: character string defining the column of `pData(object)` to be used as a factor by which to colour the points in the plot.
- `nfeatures`: numeric scalar indicating the number of features to include in the plot.
- `exprs_values`: character string indicating which values should be used as the expression values for this plot. Valid arguments are "tpm" (default; transcripts per million), "cpm" (counts per million), "fpkm" (FPKM values), "counts" (counts for each feature) or "exprs" (whatever is in the 'exprs' slot of the SCESet object; if already on the log2 scale, as indicated by the `logged` slot of the object, then `exprs` values are set to the power of 2 (so they are back on the raw scale they were on) before making the plot).
- `ncol`: number of columns to use for `facet_wrap` if only one block is defined.
- `linewidth`: numeric scalar giving the "size" parameter (in `ggplot2` parlance) for the lines plotted. Default is 1.5.
- `theme_size`: numeric scalar giving font size to use for the plotting theme
Details

Plots produced by this function are intended to provide an overview of large-scale differences between cells. For each cell, the features are ordered from most-expressed to least-expressed and the cumulative proportion of the total expression for the cell is computed across the top n features. These plots can flag cells with a very high proportion of the library coming from a small number of features; such cells are likely to be problematic for analyses. Using the colour and blocking arguments can flag overall differences in cells under different experimental conditions or affected by different batch and other variables.

Value

a ggplot plot object

Examples

```r
## Set up an example SCESet
data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)

plot(example_sceset, exprs_values = "exprs")
plot(example_sceset, exprs_values = "exprs", colour_by = "Cell_Cycle")
plot(example_sceset, exprs_values = "exprs", block1 = "Treatment", colour_by = "Cell_Cycle")
plot(example_sceset, exprs_values = "exprs", block1 = "Treatment", block2 = "Mutation_Status", colour_by = "Cell_Cycle")
# What happens if chosen expression values are not available?
plot(example_sceset, block1 = "Treatment", colour_by = "Cell_Cycle")
```

plotDiffusionMap

Plot a diffusion map for an SCESet object

Description

Produce a diffusion map plot of two components for an SCESet dataset.

Usage

```r
plotDiffusionMap(object, 
plotDiffusionMapSCESet(object, ntop = 500, ncomponents = 2,
```
Arguments

**object**
an SCESet object

... further arguments passed to `DiffusionMap`

**ntop**
numeric scalar indicating the number of most variable features to use for the diffusion map. Default is 500, but any ntop argument is overrided if the feature_set argument is non-NULL.

**ncomponents**
numeric scalar indicating the number of principal components to plot, starting from the first diffusion map component. Default is 2. If ncomponents is 2, then a scatterplot of component 1 vs component 2 is produced. If ncomponents is greater than 2, a pairs plots for the top components is produced. NB: computing many components for the diffusion map can become time consuming.

**exprs_values**
character string indicating which values should be used as the expression values for this plot. Valid arguments are “tpm” (default; transcripts per million), “norm_tpm” (normalised TPM values), “fpkm” (FPKM values), “norm_fpkm” (normalised FPKM values), “counts” (counts for each feature), “norm_counts”, “cpm” (counts-per-million), “norm_cpm” (normalised counts-per-million), “exprs” (whatever is in the ‘exprs’ slot of the SCESet object; default), “norm_exprs” (normalised expression values) or “stand_exprs” (standardised expression values) or any other named element of the assayData slot of the SCESet object that can be accessed with the get_exprs function.

**colour_by**
character string defining the column of pData(object) to be used as a factor by which to colour the points in the plot.

**shape_by**
character string defining the column of pData(object) to be used as a factor by which to define the shape of the points in the plot.

**size_by**
character string defining the column of pData(object) to be used as a factor by which to define the size of points in the plot.

**feature_set**
character, numeric or logical vector indicating a set of features to use for the diffusion map. If character, entries must all be in featureNames(object). If numeric, values are taken to be indices for features. If logical, vector is used to index features and should have length equal to nrow(object).

**return_SCESet**
logical, should the function return an SCESet object with principal component values for cells in the reducedDimension slot. Default is FALSE, in which case a ggplot object is returned.

**scale_features**
logical, should the expression values be standardised so that each feature has unit variance? Default is TRUE.

**draw_plot**
logical, should the plot be drawn on the current graphics device? Only used if return_SCESet is TRUE, otherwise the plot is always produced.

**theme_size**
numeric scalar giving default font size for plotting theme (default is 10).

**rand_seed**
(optional) numeric scalar that can be passed to set.seed to make plots reproducible.

**sigma**
argument passed to `DiffusionMap`

**distance**
argument passed to `DiffusionMap`
legend character, specifying how the legend(s) be shown? Default is "auto", which hides legends that have only one level and shows others. Alternatives are "all" (show all legends) or "none" (hide all legends).

Details

The function DiffusionMap is used internally to compute the diffusion map.

Value

If return_SCESet is TRUE, then the function returns an SCESet object, otherwise it returns a ggplot object.

References


See Also

destiny

Examples

## Set up an example SCESet
data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
drop_genes <- apply(exprs(example_sceset), 1, function(x) {var(x) == 0})
example_sceset <- example_sceset[!drop_genes, ]

## Examples plotting diffusion maps
plotDiffusionMap(example_sceset)
plotDiffusionMap(example_sceset, colour_by = "Cell_Cycle")
plotDiffusionMap(example_sceset, colour_by = "Cell_Cycle", shape_by = "Treatment")
plotDiffusionMap(example_sceset, colour_by = "Cell_Cycle", shape_by = "Treatment", size_by = "Mutation_Status")
plotDiffusionMap(example_sceset, shape_by = "Treatment", size_by = "Mutation_Status")
plotDiffusionMap(example_sceset, feature_set = 1:100, colour_by = "Treatment", shape_by = "Mutation_Status")

plotDiffusionMap(example_sceset, shape_by = "Treatment", return_SCESet = TRUE)
plotExplanatoryVariables

Plot explanatory variables ordered by percentage of phenotypic variance explained

Description
Plot explanatory variables ordered by percentage of phenotypic variance explained

Usage
plotExplanatoryVariables(object, method = "density", exprs_values = "exprs",
nvars_to_plot = 10, min_marginal_r2 = 0, variables = NULL,
return_object = FALSE, theme_size = 10, ...)

Arguments
object an SCESet object containing expression values and experimental information. Must have been appropriately prepared.
method character scalar indicating the type of plot to produce. If "density", the function produces a density plot of R-squared values for each variable when fitted as the only explanatory variable in a linear model. If "pairs", then the function produces a pairs plot of the explanatory variables ordered by the percentage of feature expression variance (as measured by R-squared in a marginal linear model) explained.
exprs_values which slot of the assayData in the object should be used to define expression? Valid options are "exprs" (default), "tpm", "fpkm", "cpm", and "counts".
nvars_to_plot integer, the number of variables to plot in the pairs plot. Default value is 10.
min_marginal_r2 numeric scalar giving the minimal value required for median marginal R-squared for a variable to be plotted. Only variables with a median marginal R-squared strictly larger than this value will be plotted.
variables optional character vector giving the variables to be plotted. Default is NULL, in which case all variables in pData(object) are considered and the nvars_to_plot variables with the highest median marginal R-squared are plotted.
return_object logical, should an SCESet object with median marginal R-squared values added to varMetadata(object) be returned?
theme_size numeric scalar giving font size to use for the plotting theme
... parameters to be passed to pairs.

Details
If the method argument is "pairs", then the function produces a pairs plot of the explanatory variables ordered by the percentage of feature expression variance (as measured by R-squared in a marginal linear model) explained by variable. Median percentage R-squared is reported on the plot for each variable. Discrete variables are coerced to a factor and plotted as integers with jittering. Variables with only one unique value are quietly ignored.
plotExpression

Value

A ggplot object

Examples

```r
data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
rownames(pd) <- pd$Cell
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
drop_genes <- apply(exprs(example_sceset), 1, function(x) {var(x) == 0})
example_sceset <- example_sceset[!drop_genes, ]
example_sceset <- calculateQCMetrics(example_sceset)
vars <- names(pData(example_sceset))[-c(2:3, 5:14)]
plotExplanatoryVariables(example_sceset, variables = vars)
```

Description

Plot expression values for a set of features (e.g. genes or transcripts)

Usage

```r
plotExpression(object, ...)

plotExpressionSCESet(object, features, x = NULL, exprs_values = "exprs",
colour_by = NULL, shape_by = NULL, size_by = NULL, ncol = 2,
xlab = NULL, show_median = FALSE, show_violin = TRUE,
show_smooth = FALSE, alpha = 0.6, theme_size = 10,
log2_values = FALSE, size = NULL, scales = "fixed", se = TRUE,
jitter = "swarm")

plotExpressionDefault(object, aesth, ncol = 2, xlab = NULL, ylab = NULL,
show_median = FALSE, show_violin = TRUE, show_smooth = FALSE,
alpha = 0.6, size = NULL, scales = "fixed", one_facet = FALSE,
se = TRUE, jitter = "swarm")
```

Arguments

- `object`: an SCESet object containing expression values and experimental information. Must have been appropriately prepared. For the `plotExpressionDefault` method, the object argument is a data.frame in 'long' format providing expression values for a set of features to plot, plus metadata used in the `aesth` argument, but this is not meant to be a user-level operation.
plotExpression

... optional arguments (from those listed above) passed to `plotExpressionSCESet` or `plotExpressionDefault`

features a character vector of feature names or Boolean vector or numeric vector of indices indicating which features should have their expression values plotted

x character string providing a column name of `pData(object)` or a feature name (i.e. gene or transcript) to plot on the x-axis in the expression plot(s). If a feature name, then expression values for the feature will be plotted on the x-axis for each subplot.

exprs_values character string indicating which values should be used as the expression values for this plot. Valid arguments are "tpm" (default: transcripts per million), "norm_tpm" (normalised TPM values), "fpkm" (FPKM values), "norm_fpkm" (normalised FPKM values), "counts" (counts for each feature), "norm_counts", "cpm" (counts-per-million), "norm_cpm" (normalised counts-per-million), "exprs" (whatever is in the ‘exprs’ slot of the SCESet object; default), "norm_exprs" (normalised expression values) or "stand_exprs" (standardised expression values) or any other slots that have been added to the "assayData" slot by the user.

colour_by optional character string supplying name of a column of `pData(object)` which will be used as a variable by which to colour expression values on the plot.

shape_by optional character string supplying name of a column of `pData(object)` which will be used as a variable to define the shape of points for expression values on the plot.

size_by optional character string supplying name of a column of `pData(object)` which will be used as a variable to define the size of points for expression values on the plot.

ncol number of columns to be used for the panels of the plot

xlab label for x-axis; if NULL (default), then x will be used as the x-axis label

show_median logical, show the median for each group on the plot

show_violin logical, show a violin plot for the distribution for each group on the plot

show_smooth logical, show a smoothed fit through the expression values on the plot

alpha numeric value between 0 (completely transparent) and 1 (completely solid) defining how transparent plotted points (cells) should be. Points are jittered horizontally if the x-axis value is categorical rather than numeric to avoid overplotting.

theme_size numeric scalar giving default font size for plotting theme (default is 10)

log2_values should the expression values be transformed to the log2-scale for plotting (with an offset of 1 to avoid logging zeroes)?

size numeric scalar optionally providing size for points if size_by argument is not given. Default is NULL, in which case `ggplot2` default is used.

scales character scalar, should scales be fixed ("fixed"), free ("free"), or free in one dimension ("free_x"; "free_y", the default). Passed to the scales argument in the facet_wrap function from the ggplot2 package.

se logical, should standard errors be shown (default `TRUE`) for the smoothed fit through the cells. (Ignored if show_smooth is `FALSE`).

jitter character scalar to define whether points are to be jittered ("jitter") or presented in a "beeswarm" style (if "swarm"; default). "Beeswarm" style usually looks more attractive, but for datasets with a large number of cells, or for dense plots, the jitter option may work better.

aesth an aes object to use in the call to `ggplot`. 
ylab character string defining a label for the y-axis (y-axes) of the plot.

one_facet logical, should expression values for features be plotted in one facet instead of multiple facets, one per feature? Default if \( x = \text{NULL} \).

### Examples

```r
## prepare data
data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
example_sceset <- calculateQCMetrics(example_sceset)

## default plot
plotExpression(example_sceset, 1:15)
plotExpression(example_sceset, 1:15, jitter = "jitter")

## plot expression against an x-axis value
plotExpression(example_sceset, 1:6, "Mutation_Status")

## explore options
plotExpression(example_sceset, 1:6, x="Mutation_Status", exprs_values="exprs",
colour_by="Cell_Cycle", show_violin=TRUE, show_median=TRUE)
plotExpression(example_sceset, 1:6, x="Mutation_Status", exprs_values="counts",
colour_by="Cell_Cycle", show_violin=TRUE, show_median=TRUE)

## plot expression against expression values for Gene_0004
plotExpression(example_sceset, 1:4, "Gene_0004")
plotExpression(example_sceset, 1:4, "Gene_0004", show_smooth = TRUE)
plotExpression(example_sceset, 1:4, "Gene_0004", show_smooth = TRUE, se = FALSE)
```

### Description

Plot frequency of expression against mean expression level

### Usage

```r
plotExprsFreqVsMean(object, feature_set = NULL, feature_controls = NULL,
shape = 1, alpha = 0.7, show_smooth = TRUE, se = TRUE, ...)
```
plotExprsFreqVsMean

Arguments

object
an SCESet object.

feature_set
character, numeric or logical vector indicating a set of features to plot. If character, entries must all be in featureNames(object). If numeric, values are taken to be indices for features. If logical, vector is used to index features and should have length equal to nrow(object). If NULL, then the function checks if feature controls are defined. If so, then only feature controls are plotted, if not, then all features are plotted.

feature_controls
character, numeric or logical vector indicating a set of features to be used as feature controls for computing technical dropout effects. If character, entries must all be in featureNames(object). If numeric, values are taken to be indices for features. If logical, vector is used to index features and should have length equal to nrow(object). If NULL, then the function checks if feature controls are defined. If so, then these feature controls are used.

shape
(optional) numeric scalar to define the plotting shape.

alpha
(optional) numeric scalar (in the interval 0 to 1) to define the alpha level (transparency) of plotted points.

show_smooth
logical, should a smoothed fit through feature controls (if available; all features if not) be shown on the plot? Lowess used if a small number of feature controls. For details see geom_smooth.

se
logical, should standard error (confidence interval) be shown for smoothed fit?

... further arguments passed to plotMetadata (should only be size, if anythin).

Details

This function plots gene expression frequency versus mean expression level, which can be useful to assess the effects of technical dropout in the dataset. We fit a non-linear least squares curve for the relationship between expression frequency and mean expression and use this to define the number of genes above high technical dropout and the numbers of genes that are expressed in at least 50 of genes to be treated as feature controls can be specified, otherwise any feature controls previously defined are used.

Value

a ggplot plot object

Examples

data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data=sc_example_cell_info)
rownames(pd) <- pd$Cell
ex_sceset <- newSCESet(countData=sc_example_counts, phenoData=pd)
ex_sceset <- calculateQCMetrics(ex_sceset)
plotExprsFreqVsMean(ex_sceset)

ex_sceset <- calculateQCMetrics(
  ex_sceset, feature_controls = list(controls1 = 1:20,
                                    controls2 = 500:1000),
  cell_controls = list(set_1 = 1:5,
                       set_2 = 31:40))
Description

Plot expression values from an SCESet object against transcript length values defined in the SCESet object or supplied as an argument.

Usage

```r
plotExprsVsTxLength(object, tx_length = "median_feat_eff_len", exprs_values = "exprs", colour_by = NULL, shape_by = NULL, size_by = NULL, xlab = NULL, show_exprs_sd = FALSE, show_smooth = FALSE, alpha = 0.6, theme_size = 10, log2_values = FALSE, size = NULL, se = TRUE)
```

Arguments

- `object`: an SCESet object
- `tx_length`: transcript lengths to plot on the x-axis. Can be one of: (1) the name of a column of fData(object) containing the transcript length values, or (2) the name of an element of assayData(object) containing a matrix of transcript length values, or (3) a numeric vector of length equal to the number of rows of object (number of features).
- `exprs_values`: character string indicating which values should be used as the expression values for this plot. Valid arguments are "tpm" (default; transcripts per million), "norm_tpm" (normalised TPM values), "fpkm" (FPKM values), "norm_fpkm" (normalised FPKM values), "counts" (counts for each feature), "norm_counts", "cpm" (counts-per-million), "norm_cpm" (normalised counts-per-million), "exprs" (whatever is in the 'exprs' slot of the SCESet object; default), "norm_exprs" (normalised expression values) or "stand_exprs" (standardised expression values) or any other slots that have been added to the "assayData" slot by the user.
- `colour_by`: optional character string supplying name of a column of fData(object) which will be used as a variable by which to colour expression values on the plot.
- `shape_by`: optional character string supplying name of a column of fData(object) which will be used as a variable to define the shape of points for expression values on the plot.
- `size_by`: optional character string supplying name of a column of fData(object) which will be used as a variable to define the size of points for expression values on the plot.
- `xlab`: label for x-axis; if NULL (default), then x will be used as the x-axis label
- `show_exprs_sd`: logical, show the standard deviation of expression values for each feature on the plot
- `show_smooth`: logical, show a smoothed fit through the expression values on the plot
plotFeatureData

Plot feature (gene) data from an SCESet object

Description

Plot feature (gene) data from an SCESet object

alpha  numeric value between 0 (completely transparent) and 1 (completely solid) defining how transparent plotted points (cells) should be. Points are jittered horizontally if the x-axis value is categorical rather than numeric to avoid overplotting.

theme_size  numeric scalar giving default font size for plotting theme (default is 10)

log2_values  should the expression values be transformed to the log2-scale for plotting (with an offset of 1 to avoid logging zeroes)?

size  numeric scalar optionally providing size for points if size_by argument is not given. Default is NULL, in which case ggplot2 default is used.

se  logical, should standard errors be shown (default TRUE) for the smoothed fit through the cells. (Ignored if show_smooth is FALSE).

Value

a ggplot object

Examples

data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
fd <- new("AnnotatedDataFrame", data =
data.frame(gene_id = rownames(sc_example_counts),
  feature_id = paste("feature", rep(1:500, each = 4), sep = ","),
  median_tx_length = rnorm(2000, mean = 5000, sd = 500))
rownames(fd) <- rownames(sc_example_counts)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd,
  featureData = fd)

plotExprsVsTxLength(example_sceset, "median_tx_length")
plotExprsVsTxLength(example_sceset, "median_tx_length", show_smooth = TRUE)
plotExprsVsTxLength(example_sceset, "median_tx_length", show_smooth = TRUE,
  show_exprs_sd = TRUE)

## using matrix of tx length values in assayData(object)
set_exprs(example_sceset, "tx_len") <-
matrix(rnorm(ncol(example_sceset) * nrow(example_sceset), mean = 5000, sd = 500),
nrow = nrow(example_sceset))
plotExprsVsTxLength(example_sceset, "tx_len", show_smooth = TRUE,
  show_exprs_sd = TRUE)

## using a vector of tx length values
plotExprsVsTxLength(example_sceset, rnorm(2000, mean = 5000, sd = 500))
plotHighestExprs

Usage

plotFeatureData(object, aesth = aes_string(x = "n_cells_exprs", y = "prop_total_counts"), theme_size = 10, ...)

Arguments

object an SCESet object containing expression values and experimental information. Must have been appropriately prepared.

aesth aesthetics function call to pass to ggplot. This function expects at least x and y variables to be supplied. The default is to produce a density plot of number of cells expressing the feature (requires calculateQC_metrics to have been run on the SCESet object prior).

theme_size numeric scalar giving default font size for plotting theme (default is 10).

Value a ggplot plot object

Examples
data("sc_example_counts")
data("sc_example_cell_info")

pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)

eample_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)

eample_sceset <- calculateQCMetrics(example_sceset)

plotFeatureData(example_sceset, aesth=aes(x=n_cells_exprs, y=pct_total_counts))

plotHighestExprs

Plot the features with the highest expression values

Description

Plot the features with the highest expression values

Usage

plotHighestExprs(object, col_by_variable = "total_features", n = 50, drop_features = NULL, exprs_values = "counts", feature_names_to_plot = NULL)
**plotMDS**

**Arguments**

- **object**: an SCESet object containing expression values and experimental information. Must have been appropriately prepared.
- **col_by_variable**: variable name (must be a column name of pData(object)) to be used to assign colours to cell-level values.
- **n**: numeric scalar giving the number of the most expressed features to show. Default value is 50.
- **drop_features**: a character, logical or numeric vector indicating which features (e.g. genes, transcripts) to drop when producing the plot. For example, control genes might be dropped to focus attention on contribution from endogenous rather than synthetic genes.
- **exprs_values**: which slot of the assayData in the object should be used to define expression? Valid options are "counts" (default), "tpm", "fpkm" and "exprs".
- **feature_names_to_plot**: character scalar indicating which column of the featureData slot in the object is to be used for the feature names displayed on the plot. Default is NULL, in which case featureNames(object) is used.

**Details**

Plot the percentage of counts accounted for by the top n most highly expressed features across the dataset.

**Value**

a ggplot plot object

**Examples**

```r
data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
rownames(pd) <- pdCell
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
rownames(example_sceset) <- calculateQMetrics(example_sceset, feature_controls = 1:500)
plotHighestExprs(example_sceset, col_by_variable="total_features")
plotHighestExprs(example_sceset, col_by_variable="Mutation_Status")
```

**Description**

`# Produce an MDS plot from the cell pairwise distance data in an SCESet dataset.`
plotMDS

Usage

plotMDS(object, ...)

plotMDSSCESet(object, ncomponents = 2, colour_by = NULL, shape_by = NULL, size_by = NULL, return_SCESet = FALSE, draw_plot = TRUE, theme_size = 10, legend = "auto")

## S4 method for signature 'SCESet'
plotMDS(object, ncomponents = 2, colour_by = NULL, shape_by = NULL, size_by = NULL, return_SCESet = FALSE, draw_plot = TRUE, theme_size = 10, legend = "auto")

Arguments

object an SCESet object

... arguments passed to S4 plotMDS method

ncomponents numeric scalar indicating the number of principal components to plot, starting from the first principal component. Default is 2. If ncomponents is 2, then a scatterplot of PC2 vs PC1 is produced. If ncomponents is greater than 2, a pairs plots for the top components is produced. NB: computing more than two components for t-SNE can become very time consuming.

colour_by character string defining the column of pData(object) to be used as a factor by which to colour the points in the plot.

shape_by character string defining the column of pData(object) to be used as a factor by which to define the shape of the points in the plot.

size_by character string defining the column of pData(object) to be used as a factor by which to define the size of points in the plot.

return_SCESet logical, should the function return an SCESet object with principal component values for cells in the reducedDimension slot. Default is FALSE, in which case a ggplot object is returned.

draw_plot logical, should the plot be drawn on the current graphics device? Only used if return_SCESet is TRUE, otherwise the plot is always produced.

theme_size numeric scalar giving default font size for plotting theme (default is 10).

legend character, specifying how the legend(s) be shown? Default is "auto", which hides legends that have only one level and shows others. Alternatives are "all" (show all legends) or "none" (hide all legends).

Details

The function cmdscale is used internally to compute the multidimensional scaling components to plot.

Value

If return_SCESet is TRUE, then the function returns an SCESet object, otherwise it returns a ggplot object.
Examples

```r
## Set up an example SCESet
data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
exs <- newSCESet(countData = sc_example_counts, phenoData = pd)
drop_genes <- apply(exprs(exs), 1, function(x) {var(x) == 0})
exs <- exs[!drop_genes, ]
exs <- calculateQCMetrics(exs)

## define cell-cell distances
cellDist(exs) <- as.matrix(dist(t(exprs(exs))))

## Examples plotting
plotMDS(exs)
plotMDS(exs, colour_by = "Cell_Cycle")
plotMDS(exs, colour_by = "Cell_Cycle", shape_by = "Treatment")

## define cell-cell distances differently
cellDist(exs) <- as.matrix(dist(t(counts(exs)), method = "canberra"))
plotMDS(exs, colour_by = "Cell_Cycle", shape_by = "Treatment", size_by = "Mutation_Status")
```

plotMetadata

Plot metadata for cells or features

Description

Plot metadata for cells or features

Usage

```r
plotMetadata(object, aesth = aes_string(x = "log10(total_counts)", y = "total_features"), shape = NULL, alpha = NULL, size = NULL, theme_size = 10)
```

Arguments

- **object**: a data.frame (or object that can be coerced to such) object containing metadata in columns to plot.
- **aesth**: aesthetics function call to pass to ggplot. This function expects at least x and y variables to be supplied. The default is to plot total_features against log10(total_counts).
- **shape**: numeric scalar to define the plotting shape. Ignored if shape is included in the aesth argument.
- **alpha**: numeric scalar (in the interval 0 to 1) to define the alpha level (transparency) of plotted points. Ignored if alpha is included in the aesth argument.
- **size**: numeric scalar to define the plotting size. Ignored if size is included in the aesth argument.
- **theme_size**: numeric scalar giving default font size for plotting theme (default is 10)
plotPCA

Plot PCA for an SCESet object

Description

Produce a principal components analysis (PCA) plot of two or more principal components for an SCESet dataset.

Usage

plotPCASCESet(object, ntop = 500, ncomponents = 2, exprs_values = "exprs", colour_by = NULL, shape_by = NULL, size_by = NULL, feature_set = NULL, return_SCESet = FALSE, scale_features = TRUE, draw_plot = TRUE, pca_data_input = "exprs", selected_variables = NULL, detect_outliers = FALSE, theme_size = 10, legend = "auto")

# S4 method for signature 'SCESet'
plotPCA(object, ntop = 500, ncomponents = 2, exprs_values = "exprs", colour_by = NULL, shape_by = NULL, size_by = NULL, feature_set = NULL, return_SCESet = FALSE, scale_features = TRUE, draw_plot = TRUE, pca_data_input = "exprs", selected_variables = NULL, detect_outliers = FALSE, theme_size = 10, legend = "auto")

Arguments

object an SCESet object
ntop numeric scalar indicating the number of most variable features to use for the PCA. Default is 500, but any ntop argument is overrided if the feature_set argument is non-NULL.
ncomponents numeric scalar indicating the number of principal components to plot, starting from the first principal component. Default is 2. If ncomponents is 2, then a scatterplot of PC2 vs PC1 is produced. If ncomponents is greater than 2, a pairs plots for the top components is produced.

exprs_values character string indicating which values should be used as the expression values for this plot. Valid arguments are “tpm” (default; transcripts per million), “norm_tpm” (normalised TPM values), “fpkm” (FPKM values), “norm_fpkm” (normalised FPKM values), “counts” (counts for each feature), “norm_counts”, “cpm” (counts-per-million), “norm_cpm” (normalised counts-per-million), “exprs” (whatever is in the ‘exprs’ slot of the SCESet object; default), “norm_exprs” (normalised expression values) or “stand_exprs” (standardised expression values) or any other named element of the assayData slot of the SCESet object that can be accessed with the get_exprs function.

colour_by character string defining the column of pData(object) to be used as a factor by which to colour the points in the plot.

shape_by character string defining the column of pData(object) to be used as a factor by which to define the shape of the points in the plot.

size_by character string defining the column of pData(object) to be used as a factor by which to define the size of points in the plot.

feature_set character, numeric or logical vector indicating a set of features to use for the PCA. If character, entries must all be in featureNames(object). If numeric, values are taken to be indices for features. If logical, vector is used to index features and should have length equal to nrow(object).

return_SCESet logical, should the function return an SCESet object with principal component values for cells in the reducedDimension slot. Default is FALSE, in which case a ggplot object is returned.

scale_features logical, should the expression values be standardised so that each feature has unit variance? Default is TRUE.

draw_plot logical, should the plot be drawn on the current graphics device? Only used if return_SCESet is TRUE, otherwise the plot is always produced.

cpyca_data_input character argument defining which data should be used as input for the PCA. Possible options are “exprs” (default), which uses expression data to produce a PCA at the cell level; “pdata” which uses numeric variables from pData(object) to do PCA at the cell level; and “fdata” which uses numeric variables from fData(object) to do PCA at the feature level.

selected_variables character vector indicating which variables in pData(object) to use for the phenotype-data based PCA. Ignored if the argument pca_data_input is anything other than “pdata”.

detect_outliers logical, should outliers be detected in the PC plot? Only an option when pca_data_input argument is “pdata”. Default is FALSE.

theme_size numeric scalar giving default font size for plotting theme (default is 10).

legend character, specifying how the legend(s) be shown? Default is “auto”, which hides legends that have only one level and shows others. Alternatives are “all” (show all legends) or “none” (hide all legends).

... further arguments passed to plotPCASCESet
Details

The function `prcomp` is used internally to do the PCA. The function checks whether the object has standardised expression values (by looking at `stand_exprs(object)`). If yes, the existing standardised expression values are used for the PCA. If not, then standardised expression values are computed using `scale` (with feature-wise unit variances or not according to the `scale_features` argument), added to the object and PCA is done using these new standardised expression values.

If the arguments `detect_outliers` and `return_SCESet` are both TRUE, then the element `$outlier` is added to the pData (phenotype data) slot of the SCESet object. This element contains indicator values about whether or not each cell has been designated as an outlier based on the PCA. These values can be accessed for filtering low quality cells with, for example, `example_sceset$outlier`.

Value

either a ggplot plot object or an SCESet object

Examples

```r
## Set up an example SCESet
data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
drop_genes <- apply(exprs(example_sceset), 1, function(x) {var(x) == 0})
example_sceset <- example_sceset[!drop_genes, ]

## Examples plotting PC1 and PC2
plotPCA(example_sceset)
plotPCA(example_sceset, colour_by = "Cell_Cycle")
plotPCA(example_sceset, colour_by = "Cell_Cycle", shape_by = "Treatment")
plotPCA(example_sceset, colour_by = "Cell_Cycle", shape_by = "Treatment", size_by = "Mutation_Status")
plotPCA(example_sceset, shape_by = "Treatment", size_by = "Mutation_Status")
plotPCA(example_sceset, feature_set = 1:100, colour_by = "Treatment", shape_by = "Mutation_Status")

## experiment with legend
example_subset <- example_sceset[, example_sceset$Treatment == "treat1"]
plotPCA(example_subset, colour_by = "Cell_Cycle", shape_by = "Treatment", legend = "all")

plotPCA(example_sceset, shape_by = "Treatment", return_SCESet = TRUE)

## Examples plotting more than 2 PCs
plotPCA(example_sceset, ncomponents = 8)
plotPCA(example_sceset, ncomponents = 4, colour_by = "Treatment", shape_by = "Mutation_Status")
```

plotPhenoData

Plot phenotype data from an SCESet object

Description

Plot phenotype data from an SCESet object
Usage

plotPhenoData(object, aesth = aes_string(x = "log10(total_counts)", y = "total_features"), theme_size = 10, ...)

Arguments

object an SCESet object containing expression values and experimental information. Must have been appropriately prepared.

aesth aesthetics function call to pass to ggplot. This function expects at least x and y variables to be supplied. The default is to plot total_features against log10(total_counts).

theme_size numeric scalar giving default font size for plotting theme (default is 10).

... arguments passed to plotMetadata.

Details

Plot phenotype data from an SCESet object. If one variable is supplied then a density plot will be returned. If both variables are continuous (numeric) then a scatter plot will be returned. If one variable is discrete and one continuous then a violin plot with jittered points overlaid will be returned. If both variables are discrete then a jitter plot will be produced. The object returned is a ggplot object, so further layers and plotting options (titles, facets, themes etc) can be added.

Value

a ggplot plot object

Examples

data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
example_sceset <- calculateQCMetrics(example_sceset)
plotPhenoData(example_sceset, aesth = aes_string(x = "log10(total_counts)", y = "total_features", colour = "Mutation_Status"))

plotPlatePosition

Plot cells in plate positions

Description

Plots cells in their position on a plate, coloured by phenotype data or feature expression.

Usage

plotPlatePosition(object, plate_position = NULL, colour_by = NULL, x_position = NULL, y_position = NULL, theme_size = 24, legend = "auto")
**Arguments**

*object* an SCESet object. If `object$plate_position` is not NULL, then this will be used to define each cell’s position on the plate, unless the `plate_position` argument is specified.

*plate_position* optional character vector providing a position on the plate for each cell (e.g. A01, B12, etc, where letter indicates row and number indicates column). Specifying this argument overrides any plate position information extracted from the SCESet object.

*colour_by* character string defining the column of `pData(object)` to be used as a factor by which to colour the points in the plot.

*x_position* numeric vector providing x-axis positions for the cells (ignored if `plate_position` is not NULL)

*y_position* numeric vector providing y-axis positions for the cells (ignored if `plate_position` is not NULL)

*theme_size* numeric scalar giving default font size for plotting theme (default is 10).

*legend* character, specifying how the legend(s) be shown? Default is "auto", which hides legends that have only one level and shows others. Alternatives are "all" (show all legends) or "none" (hide all legends).

**Details**

This function expects plate positions to be given in a character format where a letter indicates the row on the plate and a numeric value indicates the column. So each cell has a plate position such as "A01", "B12", "K24" and so on. From these plate positions, the row is extracted as the letter, and the column as the numeric part. If `object$plate_position` or the `plate_position` argument are used to define plate positions, then positions should be provided in this format. Alternatively, numeric values to be used as x- and y-coordinates by supplying both the `x_position` and `y_position` arguments to the function.

**Examples**

```r
## prepare data
data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
example_sceset <- calculateQCMetrics(example_sceset)

## define plate positions
example_sceset$plate_position <- paste0(rep(LETTERS[1:5], each = 8), rep(formatC(1:8, width = 2, flag = "0"), 5))

## plot plate positions
plotPlatePosition(example_sceset, colour_by = "Mutation_Status")
plotPlatePosition(example_sceset, colour_by = "Gene_0004")
```
**plotQC**

 Produce QC diagnostic plots

**Description**

Produce QC diagnostic plots

**Usage**

```r
plotQC(object, type = "highest-expression", ...)```

**Arguments**

- `object` an SCESet object containing expression values and experimental information. Must have been appropriately prepared.
- `type` character scalar providing type of QC plot to compute: "highest-expression" (showing features with highest expression), "find-pcs" (showing the most important principal components for a given variable), "explanatory-variables" (showing a set of explanatory variables plotted against each other, ordered by marginal variance explained), or "exprs-mean-vs-freq" (plotting the mean expression levels against the frequency of expression for a set of features).
- `...` arguments passed to `plotHighestExprs`, `plotImportantPCs`, `plotExplanatoryVariables` and `plotExprsMeanVsFreq` as appropriate.

**Details**

Display useful quality control plots to help with pre-processing of data and identification of potentially problematic features and cells.

**Value**

a ggplot plot object

**Examples**

```r
data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data=sc_example_cell_info)ownames(pd) <- pd$Cell
example_sceset <- newSCESet(countData=sc_example_counts, phenoData=pd)
drop_genes <- apply(exprs(example_sceset), 1, function(x) {var(x) == 0})
example_sceset <- example_sceset[!drop_genes, ]
example_sceset <- calculateQCMetrics(example_sceset)
plotQC(example_sceset, type="high", col_by_variable="Mutation_Status")
plotQC(example_sceset, type="find", variable="total_features")
vars <- names(pData(example_sceset))[c(2:3, 5:14)]
plotQC(example_sceset, type="expl", variables=vars)
```
plotReducedDim  Plot reduced dimension representation of cells

Description
Plot reduced dimension representation of cells

Usage
plotReducedDim(object, ...)

plotReducedDim.default(df_to_plot, ncomponents = 2, colour_by = NULL,
shape_by = NULL, size_by = NULL, percentVar = NULL, theme_size = 10,
legend = "auto")

plotReducedDim.SCESet(object, ncomponents = 2, colour_by = NULL,
shape_by = NULL, size_by = NULL, theme_size = 10, legend = "auto")

## S4 method for signature 'SCESet'
plotReducedDim(object, ncomponents = 2, colour_by = NULL,
shape_by = NULL, size_by = NULL, theme_size = 10, legend = "auto")

## S4 method for signature 'data.frame'
plotReducedDim(object, ncomponents = 2,
colour_by = NULL, shape_by = NULL, size_by = NULL, percentVar = NULL,
legend = "auto")

Arguments

object  an SCESet object with a non-NULL reducedDimension slot.
...
optional arguments (from those listed above) passed to plotReducedDim.SCESet
or plotReducedDim.default

df_to_plot  data.frame containing a reduced dimension representation of cells and optional
metadata for the plot.
ncomponents  numeric scalar indicating the number of principal components to plot, starting
from the first principal component. Default is 2. If ncomponents is 2, then
a scatterplot of Dimension 2 vs Dimension 1 is produced. If ncomponents is
greater than 2, a pairs plots for the top dimensions is produced.
colour_by  character string defining the column of pData(object) to be used as a factor
by which to colour the points in the plot.
shape_by  character string defining the column of pData(object) to be used as a factor
by which to define the shape of the points in the plot.
size_by  character string defining the column of pData(object) to be used as a factor
by which to define the size of points in the plot.
percentVar  numeric vector giving the proportion of variance in expression explained by
each reduced dimension. Only expected to be used internally in the plotPCA
function.
theme_size  numeric scalar giving default font size for plotting theme (default is 10).
legend character, specifying how the legend(s) be shown? Default is "auto", which hides legends that have only one level and shows others. Alternatives are "all" (show all legends) or "none" (hide all legends).

Details
The function plotReducedDim.default assumes that the first ncomponents columns of df_to_plot contain the reduced dimension components to plot, and that any subsequent columns define factors for colour_by, shape_by and size_by in the plot.

Value
a ggplot plot object

Examples
data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
drop_genes <- apply(exprs(example_sceset), 1, function(x) {var(x) == 0})
example_sceset <- example_sceset[!drop_genes, ]
reducedDimension(example_sceset) <- prcomp(t(exprs(example_sceset)), scale. = TRUE)$x
plotReducedDim(example_sceset)
plotReducedDim(example_sceset, colour_by="Cell_Cycle")
plotReducedDim(example_sceset, colour_by="Cell_Cycle", shape_by="Treatment")
plotReducedDim(example_sceset, colour_by="Cell_Cycle", size_by="Treatment")
plotReducedDim(example_sceset, ncomponents=5)
plotReducedDim(example_sceset, ncomponents=5, colour_by="Cell_Cycle", shape_by="Treatment")

plotTSNE

Plot t-SNE for an SCESet object

Description
Produce a t-distributed stochastic neighbour embedding (t-SNE) plot of two components for an SCESet dataset.

Usage
plotTSNE(object, ...)

## S4 method for signature 'SCESet'
plotTSNE(object, ntop = 500, ncomponents = 2,
 exprs_values = "exprs", colour_by = NULL, shape_by = NULL,
 size_by = NULL, feature_set = NULL, return_SCESet = FALSE,
 scale_features = TRUE, draw_plot = TRUE, theme_size = 10,
 rand_seed = NULL, perplexity = floor(ncol(object)/5), legend = "auto",
 ...)

plotTSNE
### Arguments

- **object**: an SCESet object
- **...**: further arguments passed to `Rtsne`
- **ntop**: numeric scalar indicating the number of most variable features to use for the t-SNE. Default is 500, but any ntop argument is overridden if the feature_set argument is non-NULL.
- **ncomponents**: numeric scalar indicating the number of t-SNE components to plot, starting from the first t-SNE component. Default is 2. If ncomponents is 2, then a scatterplot of component 1 vs component 2 is produced. If ncomponents is greater than 2, a pairs plots for the top components is produced. NB: computing more than two components for t-SNE can become very time consuming.
- **exprs_values**: character string indicating which values should be used as the expression values for this plot. Valid arguments are "tpm" (default; transcripts per million), "norm_tpm" (normalised TPM values), "fpkm" (FPKM values), "norm_fpkm" (normalised FPKM values), "counts" (counts for each feature), "norm_counts", "cpm" (counts-per-million), "norm_cpm" (normalised counts-per-million), "exprs" (whatever is in the 'exprs' slot of the SCESet object; default), "norm_exprs" (normalised expression values) or "stand_exprs" (standardised expression values), or any other named element of the assayData slot of the SCESet object that can be accessed with the `get_exprs` function.
- **colour_by**: character string defining the column of `pData(object)` to be used as a factor by which to colour the points in the plot.
- **shape_by**: character string defining the column of `pData(object)` to be used as a factor by which to define the shape of the points in the plot.
- **size_by**: character string defining the column of `pData(object)` to be used as a factor by which to define the size of points in the plot.
- **feature_set**: character, numeric or logical vector indicating a set of features to use for the t-SNE calculation. If character, entries must all be in `featureNames(object)`. If numeric, values are taken to be indices for features. If logical, vector is used to index features and should have length equal to `nrow(object)`.
- **return_SCESet**: logical, should the function return an SCESet object with principal component values for cells in the reducedDimension slot. Default is `FALSE`, in which case a `ggplot` object is returned.
- **scale_features**: logical, should the expression values be standardised so that each feature has unit variance? Default is `TRUE`.
- **draw_plot**: logical, should the plot be drawn on the current graphics device? Only used if `return_SCESet` is `TRUE`, otherwise the plot is always produced.
- **theme_size**: numeric scalar giving default font size for plotting theme (default is 10).
- **rand_seed**: (optional) numeric scalar that can be passed to `set.seed` to make plots reproducible.
- **perplexity**: numeric scalar value defining the "perplexity parameter" for the t-SNE plot. Passed to `Rtsne` - see documentation for that package for more details.
- **legend**: character, specifying how the legend(s) be shown? Default is "auto", which hides legends that have only one level and shows others. Alternatives are "all" (show all legends) or "none" (hide all legends).

### Details

The function `Rtsne` is used internally to compute the t-SNE.
Value

If `return_SCESet` is TRUE, then the function returns an SCESet object, otherwise it returns a ggplot object.

References


See Also

Rtsne

Examples

```r
## Set up an example SCESet
data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
drop_genes <- apply(exprs(example_sceset), 1, function(x) {var(x) == 0})
example_sceset <- example_sceset[!drop_genes, ]

## Examples plotting PC1 and PC2
plotTSNE(example_sceset, perplexity = 10)
plotTSNE(example_sceset, colour_by = "Cell_Cycle", perplexity = 10)
plotTSNE(example_sceset, colour_by = "Cell_Cycle", shape_by = "Treatment", perplexity = 10)
plotTSNE(example_sceset, colour_by = "Cell_Cycle", shape_by = "Treatment", size_by = "Mutation_Status", perplexity = 10)
plotTSNE(example_sceset, shape_by = "Treatment", size_by = "Mutation_Status", perplexity = 5)
plotTSNE(example_sceset, feature_set = 1:100, colour_by = "Treatment", shape_by = "Mutation_Status", perplexity = 5)
plotTSNE(example_sceset, shape_by = "Treatment", return_SCESet = TRUE, perplexity = 10)
```

---

**readKallistoResults**  
*Read kallisto results from a batch of jobs*

**Description**  
After generating transcript/feature abundance results using kallisto for a batch of samples, read these abundance values into an SCESet object.

**Usage**  
```r
readKallistoResults(kallisto_log = NULL, samples = NULL, directories = NULL, read_h5 = FALSE, kallisto_version = "current", logExprsOffset = 1, verbose = TRUE)
```
**Arguments**

- `kallisto_log` list, generated by `runKallisto`. If provided, then `samples` and `directories` arguments are ignored.
- `samples` character vector providing a set of sample names to use for the abundance results.
- `directories` character vector providing a set of directories containing kallisto abundance results to be read in.
- `read_h5` logical, should the bootstrap results be read in from the HDF5 objects produced by kallisto?
- `kallisto_version` character string indicating whether or not the version of kallisto to be used is "pre-0.42.2" or "current". This is required because the kallisto developers changed the output file extensions and added features in version 0.42.2.
- `logExprsOffset` numeric scalar, providing the offset used when doing log2-transformations of expression data to avoid trying to take logs of zero. Default offset value is 1.
- `verbose` logical, should function provide output about progress?

**Details**

This function expects to find only one set of kallisto abundance results per directory; multiple abundance results in a given directory will be problematic.

**Value**

an SCESet object

**Examples**

```r
## Not run:
kallisto_log <- runKallisto("targets.txt", "transcripts.idx", single_end=FALSE, 
output_prefix="output", verbose=TRUE, n_bootstrap_samples=10)
sceset <- readKallistoResults(kallisto_log)
## End(Not run)
```

---

**readKallistoResultsOneSample**

*Read kallisto results for a single sample into a list*

**Description**

Read kallisto results for a single sample into a list

**Usage**

```r
readKallistoResultsOneSample(directory, read_h5 = FALSE, 
kallisto_version = "current")
```
**Arguments**

directory character string giving the path to the directory containing the kallisto results for the sample.

read_h5 logical, if TRUE then read in bootstrap results from the HDF5 object produced by kallisto.

kallisto_version character string indicating whether or not the version of kallisto to be used is "pre-0.42.2" or "current". This is required because the kallisto developers changed the output file extensions and added features in version 0.42.2.

**Details**

The directory is expected to contain results for just a single sample. Putting more than one sample’s results in the directory will result in unpredictable behaviour with this function. The function looks for the files (with the default names given by kallisto) 'abundance.txt', 'run_info.json' and (if read_h5=TRUE) 'abundance/h5'. If these files are missing, or if results files have different names, then this function will not find them.

**Value**

A list with two elements: (1) a data.frame abundance with columns for 'target_id' (feature, transcript, gene etc), 'length' (feature length), 'eff_length' (effective feature length), 'est_counts' (estimated feature counts), 'tpm' (transcripts per million) and possibly many columns containing bootstrap estimated counts; and (2) a list run_info with details about the kallisto run that generated the results.

**Examples**

```r
# If kallisto results are in the directory "output", then call:
# readKallistoResultsOneSample("output")
```

---

**Description**

After generating transcript/feature abundance results using Salmon for a batch of samples, read these abundance values into an SCESet object.

**Usage**

```r
readSalmonResults(Salmon_log = NULL, samples = NULL, directories = NULL,
                  logExprsOffset = 1, verbose = TRUE)
```

**Arguments**

Salmon_log list, generated by runSalmon. If provided, then samples and directories arguments are ignored.

samples character vector providing a set of sample names to use for the abundance results.
directories character vector providing a set of directories containing Salmon abundance results to be read in.

logExprsOffset numeric scalar, providing the offset used when doing log2-transformations of expression data to avoid trying to take logs of zero. Default offset value is 1.

verbose logical, should function provide output about progress?

Details

This function expects to find only one set of Salmon abundance results per directory; multiple abundance results in a given directory will be problematic.

Value

an SCESet object

Examples

```r
## Not run:
## Define output directories in a vector called here "Salmon_dirs"
## and sample names as "Salmon_samples"
sceset <- readSalmonResults(samples = Salmon_samples,
directories = Salmon_dirs)
## End(Not run)
```

readSalmonResultsOneSample

Read Salmon results for a single sample into a list

Description

Read Salmon results for a single sample into a list

Usage

`readSalmonResultsOneSample(directory)`

Arguments

directory character string giving the path to the directory containing the Salmon results for the sample.

Details

The directory is expected to contain results for just a single sample. Putting more than one sample’s results in the directory will result in unpredictable behaviour with this function. The function looks for the files (with the default names given by Salmon) ‘quant.sf’, ‘stats.tsv’, ‘libFormatCounts.txt’ and the sub-directories ‘logs’ (which contains a log file) and ‘libParams’ (which contains a file detailing the fragment length distribution). If these files are missing, or if results files have different names, then this function will not find them.

This function will work for Salmon v0.7.x and greater, as the name of one of the default output directories was changed from "aux" to "aux_info" in Salmon v0.7.
Value
A list with two elements: (1) a data.frame abundance with columns for 'target_id' (feature, transcript, gene etc), 'length' (feature length), 'est_counts' (estimated feature counts), 'tpm' (transcripts per million); (2) a list, run_info, with metadata about the Salmon run that generated the results, including number of reads processed, mapping percentage, the library type used for the RNA-sequencing, including details about number of reads that did not match the given or inferred library type, details about the Salmon command used to generate the results, and so on.

Examples
```r
## Not run:
# If Salmon results are in the directory "output", then call:
readSalmonResultsOneSample("output")
## End(Not run)
```

`readTxResults`  
**Read transcript quantification data with tximport package**

Description
After generating transcript/feature abundance results using kallisto, Salmon, Sailfish or RSEM for a batch of samples, read these abundance values into an SCESet object.

Usage
`readTxResults(samples = NULL, files = NULL, log = NULL, type = "kallisto", txOut = TRUE, logExprsOffset = 1, verbose = TRUE, ...)`

Arguments

- `samples` character vector providing a set of sample names to use for the abundance results.
- `files` character vector providing a set of filenames containing kallisto abundance results to be read in.
- `log` list (optional), generated by `runKallisto`. If provided, then `samples` and `files` arguments are ignored.
- `type` character, the type of software used to generate the abundances. Options are "kallisto", "salmon", "sailfish", "rsem". This argument is passed to `tximport`.
- `txOut` logical, whether the function should just output transcript-level (default TRUE)
- `logExprsOffset` numeric scalar, providing the offset used when doing log2-transformations of expression data to avoid trying to take logs of zero. Default offset value is 1.
- `verbose` logical, should function provide output about progress?
- `...` optional parameters passed to `tximport`. See documentation for `tximport` for options and details.
Details

Note: tximport does not import bootstrap estimates from kallisto, Salmon, or Sailfish. If you want bootstrap estimates use the readKallistoResults or readSalmonResults functions.

Value

an SCESet object containing the abundance, count and feature length data from the supplied samples.

References


Examples

```r
## this example requires installation of the tximportData package from
## Bioconductor
library(tximportData)

dir <- system.file("extdata", package = "tximportData")
list.files(dir)
samples <- read.table(file.path(dir, "samples.txt"), header = TRUE)
samples
directories <- file.path(dir, "kallisto", samples$run)
names(directories) <- paste0("sample", 1:6)
files <- file.path(directories, "abundance.tsv")
sce_example <- readTxResults(samples = names(directories),
files = files, type = "kallisto")

## for faster reading of results use the read_tsv function from the readr pkg
library(readr)
sce_example <- readTxResults(samples = names(directories),
files = files, type = "kallisto", reader = read_tsv)

## End(Not run)
```

reducedDimension

Reduced dimension representation for cells in an SCESet object

Description

SCESet objects can contain a matrix of reduced dimension coordinates for cells. These functions conveniently access and replace the reduced dimension coordinates with the value supplied, which must be a matrix of the correct size. The function redDim is simply shorthand for reducedDimension.

Usage

```r
reducedDimension(object)

reducedDimension(object) <- value
```
rename

```r
redDim(object)
redDim(object) <- value
reducedDimension.SCESet(object)

## S4 method for signature 'SCESet'
reducedDimension(object)
redDim.SCESet(object)

## S4 method for signature 'SCESet'
redDim(object)

## S4 replacement method for signature 'SCESet,matrix'
reducedDimension(object) <- value

## S4 replacement method for signature 'SCESet,matrix'
redDim(object) <- value
```

**Arguments**

- `object`: a SCESet object.
- `value`: a matrix of class "numeric" containing reduced dimension coordinates for cells.

**Value**

If accessing the `reducedDimension` slot, then the matrix of reduced dimension coordinates. If replacing the `reducedDimension` slot then the new matrix is added to the SCESet object.

**Author(s)**

Davis McCarthy

**Examples**

```r
data("sc_example_counts")
data("sc_example_cell_info")

pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)

example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)

reducedDimension(example_sceset)
```

---

**rename**

*Rename variables of pData(object).*

**Description**

Rename variables of pData(object).
Run kallisto on FASTQ files to quantify feature abundance

### Description

Run the abundance quantification tool kallisto on a set of FASTQ files. Requires kallisto ([http://pachterlab.github.io/kallisto/](http://pachterlab.github.io/kallisto/)) to be installed and a kallisto feature index must have been generated prior to using this function. See the kallisto website for installation and basic usage instructions.

### Usage

```r
runKallisto(targets_file, transcript_index, single_end = TRUE, 
            output_prefix = "output", fragment_length = NULL, 
            fragment_standard_deviation = NULL, n_cores = 2, 
            n_bootstrap_samples = 0, bootstrap_seed = NULL, correct_bias = TRUE, 
            plaintext = FALSE, kallisto_version = "current", verbose = TRUE, 
            dry_run = FALSE, kallisto_cmd = "kallisto")
```

### Arguments

- **targets_file** character string giving the path to a tab-delimited text file with either 2 columns (single-end reads) or 3 columns (paired-end reads) that gives the sample names (first column) and FastQ file names (column 2 and if applicable 3). The file is assumed to have column headers, although these are not used.
- **transcript_index** character string giving the path to the kallisto index to be used for the feature abundance quantification.
- **single_end** logical, are single-end reads used, or paired-end reads?
output_prefix character string giving the prefix for the output folder that will contain the kallisto results. The default is "output" and the sample name (column 1 of targets_file) is appended (preceded by an underscore).

fragment_length scalar integer or numeric giving the estimated average fragment length. Required argument if single_end is TRUE, optional if FALSE (kallisto default for paired-end data is that the value is estimated from the input data).

fragment_standard_deviation scalar numeric giving the estimated standard deviation of read fragment length. Required argument if single_end is TRUE, optional if FALSE (kallisto default for paired-end data is that the value is estimated from the input data).

n_cores integer giving the number of cores (nodes/threads) to use for the kallisto jobs. The package parallel is used. Default is 2 cores.

n_bootstrap_samples integer giving the number of bootstrap samples that kallisto should use (default is 0). With bootstrap samples, uncertainty in abundance can be quantified.

bootstrap_seed scalar integer or numeric giving the seed to use for the bootstrap sampling (default used by kallisto is 42). Optional argument.

correct_bias logical, should kallisto’s option to model and correct abundances for sequence specific bias? Requires kallisto version 0.42.2 or higher.

plaintext logical, if TRUE then bootstrapping results are returned in a plain text file rather than an HDF5 file. https://www.hdfgroup.org/HDF5/

callisto_version character string indicating whether or not the version of kallisto to be used is "pre-0.42.2" or "current". This is required because the kallisto developers changed the output file extensions and added features in version 0.42.2.

verbose logical, should timings for the run be printed?

dry_run logical, if TRUE then a list containing the kallisto commands that would be run and the output directories is returned. Can be used to read in results if kallisto is run outside an R session or to produce a script to run outside of an R session.

callisto_cmd (optional) string giving full command to use to call kallisto, if simply typing "kallisto" at the command line does not give the required version of kallisto or does not work. Default is simply "kallisto". If used, this argument should give the full path to the desired kallisto binary.

Details

A kallisto transcript index can be built from a FASTA file: kallisto index [arguments] FASTA-file. See the kallisto documentation for further details.

Value

A list containing three elements for each sample for which feature abundance has been quantified: (1) kallisto_call, the call used for kallisto, (2) kallisto_log the log generated by kallisto, and (3) output_dir the directory in which the kallisto results can be found.

Examples

## Not run:
## If in kallisto's 'test' directory, then try these calls:
runSalmon

## Generate 'targets.txt' file:
write.table(data.frame(Sample="sample1", File1="reads_1.fastq.gz", File2="reads_1.fastq.gz"),
file="targets.txt", quote=FALSE, row.names=FALSE, sep="\t")
kallisto_log <- runKallisto("targets.txt", "transcripts.idx", single_end=FALSE,
output_prefix="output", verbose=TRUE, n_bootstrap_samples=10,
dry_run = FALSE)

## End(Not run)

runSalmon

Run Salmon on FASTQ files to quantify feature abundance

Description

Run the abundance quantification tool Salmon on a set of FASTQ files. Requires Salmon (https://combine-lab.github.io/salmon/) to be installed and a Salmon transcript index must have been generated prior to using this function. See the Salmon website for installation and basic usage instructions.

Usage

runSalmon(targets_file, transcript_index, single_end = FALSE,
output_prefix = "output", lib_type = "A", n_processes = 2,
n_thread_per_process = 4, n_bootstrap_samples = 0, seqBias = TRUE,
GCbias = TRUE, posBias = FALSE, allowOrphans = FALSE,
advanced_opts = NULL, verbose = TRUE, dry_run = FALSE,
salmon_cmd = "salmon")

Arguments

targets_file character string giving the path to a tab-delimited text file with either 2 columns (single-end reads) or 3 columns (paired-end reads) that gives the sample names (first column) and FastQ file names (column 2 and if applicable 3). The file is assumed to have column headers, although these are not used.

transcript_index character string giving the path to the Salmon index to be used for the feature abundance quantification.

single_end logical, are single-end reads used, or paired-end reads?

output_prefix character string giving the prefix for the output folder that will contain the Salmon results. The default is "output" and the sample name (column 1 of targets_file) is appended (preceded by an underscore).

lib_type scalar, indicating RNA-seq library type. See Salmon documentation for details. Default is "A", for automatic detection.

n_processes integer giving the number of processes to use for parallel Salmon jobs across samples. The package parallel is used. Default is 2 concurrent processes.

n_thread_per_process integer giving the number of threads for Salmon to use per process (to parallelize Salmon for a given sample). Default is 4.

n_bootstrap_samples integer giving the number of bootstrap samples that Salmon should use (default is 0). With bootstrap samples, uncertainty in abundance can be quantified.
runSalmon

seqBias logical, should Salmon's option be used to model and correct abundances for sequence specific bias? Default is TRUE.

gcBias logical, should Salmon's option be used to model and correct abundances for GC content bias? Requires Salmon version 0.7.2 or higher. Default is TRUE.

posBias logical, should Salmon's option be used to model and correct abundances for positional biases? Requires Salmon version 0.7.3 or higher. Default is FALSE.

allowOrphans logical, Consider orphaned reads as valid hits when performing lightweight-alignment. This option will increase sensitivity (allow more reads to map and more transcripts to be detected), but may decrease specificity as orphaned alignments are more likely to be spurious. For more details see Salmon documentation.

advanced_opts character scalar supplying list of advanced option arguments to apply to each Salmon call. For details see Salmon documentation or type salmon quant --help-reads at the command line.

verbose logical, should timings for the run be printed?

dry_run logical, if TRUE then a list containing the Salmon commands that would be run and the output directories is returned. Can be used to read in results if Salmon is run outside an R session or to produce a script to run outside of an R session.

salmon_cmd (optional) string giving full command to use to call Salmon, if simply typing "salmon" at the command line does not give the required version of Salmon or does not work. Default is simply "salmon". If used, this argument should give the full path to the desired Salmon binary.

Details

A Salmon transcript index can be built from a FASTA file: salmon index [arguments] FASTA-file. See the Salmon documentation for further details. This simple wrapper does not give access to all nuances of Salmon usage. For finer-grained usage of Salmon please run it at the command line - results can still be read into R with readSalmonResults.

Value

A list containing three elements for each sample for which feature abundance has been quantified: (1) salmon_call, the call used for Salmon, (2) salmon_log the log generated by Salmon, and (3) output_dir the directory in which the Salmon results can be found.

Examples

```r
## Not run:
## If in Salmon's 'test' directory, then try these calls:
## Generate 'targets.txt' file:
write.table(data.frame(Sample="sample1", File1="reads_1.fastq.gz", File2="reads_1.fastq.gz"),
file="targets.txt", quote=FALSE, row.names=FALSE, sep="\t")
Salmon_log <- runSalmon("targets.txt", "transcripts.idx", single_end=FALSE,
    output_prefix="output", verbose=TRUE, n_bootstrap_samples=10,
    dry_run = FALSE)

## End(Not run)
```
scater_gui  

**Description**

scater shiny app GUI for workflow for less programmatically inclined users or those who would like a quick and easy way to view multiple plots.

**Usage**

```r
cscater_gui(sce_set)
```

**Arguments**

- `sce_set`  
  SCESet object after running `calculateQCMetrics` on it

**Value**

Opens a browser window with an interactive shiny app and visualize all possible plots included in the scater

**Author(s)**

Davis McCarthy and Vladimir Kiselev

**Examples**

```r
data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data=sc_example_cell_info)
rownames(pd) <- pd$Cell
example_sceset <- newSCESet(countData=sc_example_counts, phenoData=pd)
drop_genes <- apply(exprs(example_sceset), 1, function(x) {var(x) == 0})
example_sceset <- example_sceset[!drop_genes, ]
example_sceset <- calculateQCMetrics(example_sceset, feature_controls = 1:40)
## Not run:
scater_gui(example_sceset)
## End(Not run)
```

---

**SCESet**

*The "Single Cell Expression Set" (SCESet) class*

**Description**

S4 class and the main class used by scater to hold single cell expression data. SCESet extends the basic Bioconductor ExpressionSet class.

**Details**

This class is initialized from a matrix of expression values.

Methods that operate on SCESet objects constitute the basic scater workflow.
Slots

- `logged`: Scalar of class "logical", indicating whether or not the expression data in the 'exprs' slot have been log2-transformed or not.
- `logExprsOffset`: Scalar of class "numeric", providing an offset applied to expression data in the 'exprs' slot when undergoing log2-transformation to avoid trying to take logs of zero.
- `lowerDetectionLimit`: Scalar of class "numeric", giving the lower limit for an expression value to be classified as "expressed".
- `cellPairwiseDistances`: Matrix of class "numeric", containing pairwise distances between cells.
- `featurePairwiseDistances`: Matrix of class "numeric", containing pairwise distances between features.
- `reducedDimension`: Matrix of class "numeric", containing reduced-dimension coordinates for cells (generated, for example, by PCA).
- `bootstraps`: Array of class "numeric" that can contain bootstrap estimates of the expression or count values.
- `sc3`: List containing results from consensus clustering from the SC3 package.
- `featureControlInfo`: Data frame of class "AnnotatedDataFrame" that can contain information/metadata about sets of control features defined for the SCESet object. Bootstrap estimates of the expression or count values.
- `useForExprs`: Character string (one of 'exprs','tpm','counts' or 'fpkm') indicating which expression representation both internal methods and external packages should use. Defaults to 'exprs'.

References

Thanks to the Monocle package (github.com/cole-trapnell-lab/monocle-release/) for their CellDataSet class, which provided the inspiration and template for SCESet.

SCESet-subset  Subsetting SCESet Objects

Description

Subset method for SCESet objects, which subsets both the expression data, phenotype data, feature data and other slots in the object.

Usage

```r
## S4 method for signature 'SCESet'
x[1, j, ..., drop = FALSE]
```

Arguments

- `x`: object from which to extract element(s) or in which to replace element(s).
- `i, j, ...`: indices specifying elements to extract or replace. Indices are numeric or character vectors or empty (missing) or NULL. Numeric values are coerced to integer as by `as.integer` (and hence truncated towards zero). Character vectors will be matched to the names of the object (or for matrices/arrays, the dimnames): see `Extract` for further details.
For `[i-indexing only: i, j, ... can be logical vectors, indicating elements/slices to select. Such vectors are recycled if necessary to match the corresponding extent. i, j, ... can also be negative integers, indicating elements/slices to leave out of the selection. When indexing arrays by a single argument i can be a matrix with as many columns as there are dimensions of x; the result is then a vector with elements corresponding to the sets of indices in each row of i. An index value of NULL is treated as if it were integer(0).

drop For matrices and arrays. If TRUE the result is coerced to the lowest possible dimension (see the examples). This only works for extracting elements, not for the replacement. See drop for further details.

Value

an SCESet object

See Also

Extract

sc_example_cell_info  Cell information for the small example single-cell counts dataset to demonstrate capabilities of scater

Description

This data.frame contains cell metadata information for the 40 cells included in the example counts dataset included in the package.

Usage

sc_example_cell_info

Format

a data.frame instance, 1 row per cell.

Value

NULL, but makes available a data frame with cell metadata

Author(s)

Davis McCarthy, 2015-03-05

Source

Wellcome Trust Centre for Human Genetics, Oxford
sc_example_counts

A small example of single-cell counts dataset to demonstrate capabilities of scater

Description
This data set contains counts for 2000 genes for 40 cells. They are from a real experiment, but details have been anonymised.

Usage
sc_example_counts

Format
a matrix instance, 1 row per gene.

Value
NULL, but makes available a matrix of count data

Author(s)
Davis McCarthy, 2015-03-05

Source
Wellcome Trust Centre for Human Genetics, Oxford

set_exprs<- Assignment method for the new elements of an SCESet object.

Description
The assayData slot of an SCESet object holds the expression data matrices. This function makes it convenient to add new transformations of the expression data to the assayData slot.

Usage
set_exprs(object, name) <- value

## S4 replacement method for signature 'SCESet,ANY,matrix'
set_exprs(object, name)<=value

## S4 replacement method for signature 'SCESet,ANY,\'NULL\''
set_exprs(object, name) <- value
sizeFactors

Arguments

object  a SCESet object.
name    character string giving the name of the slot to which the data matrix is to be 
        assigned (can already exist or not).
value   a numeric or integer matrix matching the dimensions of the other elements of 
        the assayData slot of the SCESet object.

Value

NULL, but adds expression data to the SCESet object

Author(s)

Davis McCarthy

Examples

data("sc_example_counts")
data("sc_example_cell_info")
example_sceset <- newSCESet(countData = sc_example_counts)
set_exprs(example_sceset, "scaled_counts") <- t(t(counts(example_sceset)) / 
colSums(counts(example_sceset)))
get_exprs(example_sceset, "scaled_counts")[1:6, 1:6]

## get rid of scaled counts
set_exprs(example_sceset, "scaled_counts") <- NULL

sizeFactors

Accessors size factors of an SCESet object.

Description

For normalisation, library-specific size factors can be defined. Raw values can be divided by the 
appropriate size factors to obtain normalised counts, TPM, etc.

Usage

## S4 method for signature 'SCESet'
sizeFactors(object, type)

## S4 replacement method for signature 'SCESet,numeric'
sizeFactors(object, type) <- value

## S4 replacement method for signature 'SCESet,NULL'
sizeFactors(object, type) <- value

## S4 method for signature 'SCESet'
sizeFactors(object, type = NULL)

## S4 replacement method for signature 'SCESet,numeric'
sizeFactors(object, type = NULL, ...) <- value
## S4 replacement method for signature 'SCESet,`NULL`'

```
sizeFactors(object, type = NULL, ...) <- value
```

### Arguments

- **object**: a SCESet object.
- **type**: optional character argument providing the type or name of the size factors to be accessed or assigned.
- **...**: further arguments passed to the function
- **value**: a vector of class "numeric" or NULL

### Details

The size factors can alternatively be directly accessed from the SCESet object with `object$size_factor_type` (where "type" in the preceding is replaced by the actual type name).

### Author(s)

Davis McCarthy and Aaron Lun

### Examples

```r
data("sc_example_counts")
data("sc_example_cell_info")
example_sceset <- newSCESet(countData = sc_example_counts)
sizeFactors(example_sceset)
sizeFactors(example_sceset, NULL) <- 2 ^ rnorm(ncol(example_sceset))

example_sceset <- calculateQCMetrics(example_sceset,
  feature_controls = list(set1 = 1:40))
sizeFactors(example_sceset, "set1") <- 2 ^ rnorm(ncol(example_sceset))
sizeFactors(example_sceset)
```

---

### stand_exprs

Accessors for the 'stand_exprs' (standardised expression) element of an SCESet object.

### Description

The stand_exprs element of the arrayData slot in an SCESet object holds a matrix containing standardised (mean-centred, variance standardised, by feature) expression values. It has the same dimensions as the 'exprs' and 'counts' elements, which hold the transformed expression data and count data, respectively.
Usage

```r
stand_exprs(object)
stand_exprs(object) <- value
```

```r
## S4 method for signature 'SCESet'
stand_exprs(object)
## S4 replacement method for signature 'SCESet, matrix'
stand_exprs(object) <- value
## S4 method for signature 'SCESet'
stand_exprs(object)
## S4 replacement method for signature 'SCESet, matrix'
stand_exprs(object) <- value
```

Arguments

- `object`: a SCESet object.
- `value`: an integer matrix

Details

The default for normalised expression values is mean-centred and variance-standardised expression data from the `exprs` slot of the SCESet object. The function `normaliseExprs` (or `normalizeExprs`) provides more options and functionality for normalising expression data.

Value

- a matrix of standardised expression data

Author(s)

Davis McCarthy

Examples

```r
data("sc_example_counts")
data("sc_example_cell_info")
example_sceset <- newSCESet(countData = sc_example_counts)
stand_exprs(example_sceset)
```

summariseExprsAcrossFeatures

*Summarise expression values across feature*

Description

Create a new SCESet with counts summarised at a different feature level. A typical use would be to summarise transcript-level counts at gene level.
Usage

```r
summariseExprsAcrossFeatures(object, exprs_values = "tpm",
summarise_by = "feature_id", scaled_tpm_counts = TRUE, lib_size = NULL)
```

Arguments

- `object`: an SCESet object.
- `exprs_values`: character string indicating which slot of the assayData from the SCESet object should be used as expression values. Valid options are 'exprs' the expression slot, 'tpm' the transcripts-per-million slot or 'fpkm' the FPKM slot.
- `summarise_by`: character string giving the column of `fData(object)` that will be used as the features for which summarised expression levels are to be produced. Default is 'feature_id'.
- `scaled_tpm_counts`: logical, should feature-summarised counts be computed from summed TPM values scaled by total library size? This approach is recommended (see [https://f1000research.com/articles/4-1521/v2](https://f1000research.com/articles/4-1521/v2)), so the default is `TRUE` and it is applied if TPM values are available in the object.
- `lib_size`: optional vector of numeric values of same length as the number of columns in the SCESet object providing the total library size (e.g. "count of mapped reads") for each cell/sample.

Details

Only transcripts-per-million (TPM) and fragments per kilobase of exon per million reads mapped (FPKM) expression values should be aggregated across features. Since counts are not scaled by the length of the feature, expression in counts units are not comparable within a sample without adjusting for feature length. Thus, we cannot sum counts over a set of features to get the expression of that set (for example, we cannot sum counts over transcripts to get accurate expression estimates for a gene). See the following link for a discussion of RNA-seq expression units by Harold Pimentel: [https://haroldpimentel.wordpress.com/2014/05/08/what-the-fpkm-a-review-rna-seq-expression-units](https://haroldpimentel.wordpress.com/2014/05/08/what-the-fpkm-a-review-rna-seq-expression-units). For more details about the effects of summarising transcript expression values at the gene level see Sonesen et al, 2016 ([https://f1000research.com/articles/4-1521/v2](https://f1000research.com/articles/4-1521/v2)).

Value

an SCESet object

Examples

```r
data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
fd <- new("AnnotatedDataFrame", data =
data.frame(gene_id = featureNames(example_sceset),
feature_id = paste("feature", rep(1:500, each = 4), sep = ","))
rownames(fd) <- featureNames(example_sceset)
fData(example_sceset) <- fd
effective_length <- rep(c(1000, 2000), times = 1000)
 TPM(example_sceset) <- calculateTPM(example_sceset, effective_length, calc_from = "counts")

example_sceset_summarised <-
```
summariseExprsAcrossFeatures(example_sceset, exprs_values = "tpm")
example_sceset_summarised <-
summariseExprsAcrossFeatures(example_sceset, exprs_values = "counts")
example_sceset_summarised <-
summariseExprsAcrossFeatures(example_sceset, exprs_values = "exprs")

---

toCellDataSet

Convert an SCESet to a CellDataSet

Description

Convert an SCESet to a CellDataSet

Usage

toCellDataSet(sce, exprs_values = "exprs")

Arguments

sce

An SCESet object

exprs_values

What should exprs(cds) be mapped from in the SCESet? Should be one of "exprs", "tpm", "fpkm", "counts"

Value

An object of class SCESet

Examples

data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
if (requireNamespace("monocle") ) {
toCellDataSet(example_sceset)
}

---

tpm

Accessors for the 'tpm' (transcripts per million) element of an SCESet object.

Description

The tpm element of the arrayData slot in an SCESet object holds a matrix containing transcripts-per-million values. It has the same dimensions as the 'exprs' and 'counts' elements, which hold the transformed expression data and count data, respectively.
updateSCESet

Usage

tpm(object)

tpm(object) <- value

## S4 method for signature 'SCESet'

## S4 replacement method for signature 'SCESet,matrix'

## S4 method for signature 'SCESet'

tpm(object)

## S4 replacement method for signature 'SCESet,matrix'

tpm(object) <- value

Arguments

object a SCESet object.
value a matrix of class "numeric"

Value

a matrix of transcripts-per-million data

Author(s)

Davis McCarthy

Examples

data("sc_example_counts")
data("sc_example_cell_info")
example_sceset <- newSCESet(countData = sc_example_counts)
tpm(example_sceset)

updateSCESet

Update an SCESet object to the current version

Description

It can be necessary to update an SCESet produced with an older version of the package to be compatible with the current version of the package.

Usage

updateSCESet(object)
writeSCESet

Arguments

object an SCESet object to be updated

Value

an updated SCESet object

Examples

data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
example_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)
updateSCESet(example_sceset)

writeSCESet

Write an SCESet object to an HDF5 file

Description

Write an SCESet object to an HDF5 file

Usage

writeSCESet(object, file_path, type = "HDF5", overwrite_existing = FALSE)

Arguments

object SCESet object to be writted to file
file_path path to written file containing data from SCESet object
type character string indicating type of output file. Default is "HDF5".
overwrite_existing logical, if a file of the same name already exists should it be overwritten? Default is FALSE.

Details

Currently writing to HDF5 files is supported. The rhdf5 package is used to write data to file and can be used to read data from HDF5 files into R. For further details about the HDF5 data format see https://support.hdfgroup.org/HDF5/.

Value

Return is NULL, having written the SCESet object to file.
Examples

```r
data("sc_example_counts")
data("sc_example_cell_info")
pd <- new("AnnotatedDataFrame", data = sc_example_cell_info)
exemple_sceset <- newSCESet(countData = sc_example_counts, phenoData = pd)

## Not run:
writeSCESet(exemple_sceset, "test.h5")
file.remove("test.h5")

## End(Not run)
```
Index

[,SCESet,ANY,ANY,ANY-method (SCESet-subset), 75
[,SCESet,ANY,ANY-method (SCESet-subset), 75
[,SCESet,ANY-method (SCESet-subset), 75
[,SCESet-method (SCESet-subset), 75

arrange, 3
arrange,SCESet-method (arrange), 3
arrange.SCESet (arrange), 3
as.integer, 75

bootstraps, 4
bootstraps,SCESet-method (bootstraps), 4
bootstraps.SCESet (bootstraps), 4
bootstraps<- (bootstraps), 4
bootstraps<-,SCESet,array-method (bootstraps), 4

calcIsExprs, 5
calcNormFactors, 30, 31
calculateFPKM, 6, 29
calculateQCMetrics, 6, 74
calculateTPM, 9, 29
cellDist (cellPairwiseDistances), 11
cellDist,SCESet-method (cellPairwiseDistances), 11
cellDist<-(cellPairwiseDistances), 11

cellDist,SCESet-method (cellPairwiseDistances), 11
cellDist<-,SCESet,dist-method (cellPairwiseDistances), 11

cellDist<-,SCESet,matrix-method (cellPairwiseDistances), 11
cellDistSCESet (cellPairwiseDistances), 11
cellNames (cellNames<-), 10
cellNames<-, 10
cellNames<-,SCESet,vector-method (cellNames<-), 10

cellPairwiseDistances, 11
cellPairwiseDistances,SCESet-method (cellPairwiseDistances), 11
cellPairwiseDistances.SCESet (cellPairwiseDistances), 11

celPairwiseDistances<-(cellPairwiseDistances), 11
cellPairwiseDistances<-,SCESet,dist-method (cellPairwiseDistances), 11

cellPairwiseDistances<-,SCESet,matrix-method (cellPairwiseDistances), 11

cmdscale, 52
counts, 12
counts,SCESet-method (counts), 12
counts.SCESet (counts), 12
counts<-,SCESet,method (counts), 12

cpm, 13, 29
cpm,SCESet-method (cpm), 13
cpm<-(cpm), 13
cpm<-,SCESet,matrix-method (cpm), 13
cpmSCESet (cpm), 13

destiny, 42
DiffusionMap, 41, 42
drop, 76

Extract, 75, 76

facet_wrap, 45

fData
tData<-,SCESet,AnnotatedDataFrame-method, 14

cellDist<-,SCESet,AnnotatedDataFrame-method, 14

cellDist<-,SCESet,AnnotatedDataFrame-method, 14

cellDist<-,SCESet,AnnotatedDataFrame-method, 14

featDist (featurePairwiseDistances), 15
featDist,SCESet-method (featurePairwiseDistances), 15

featDist<-(featurePairwiseDistances), 15

featDist<-,SCESet,method (featurePairwiseDistances), 15
featDist<-,SCESet,matrix-method
  (featurePairwiseDistances), 15
featDistSCESet
  (featurePairwiseDistances), 15
featurePairwiseDistances, 15
featurePairwiseDistances,SCESet-method
  (featurePairwiseDistances), 15
featurePairwiseDistances<-
  (featurePairwiseDistances), 15
featurePairwiseDistances<-,SCESet,dist-method
  (featurePairwiseDistances), 15
featurePairwiseDistances<-,SCESet,matrix-method
  (featurePairwiseDistances), 15
featurePairwiseDistancesSCESet
  (featurePairwiseDistances), 15
filter, 16
filter,SCESet-method(filter), 16
filter.SCESet
  (filter), 16
findImportantPCs, 17
fpkm, 18
fpkm,SCESet-method
  (fpkm), 18
fpkm.SCESet
  (fpkm), 18
fpkm<-
  (fpkm), 18
fpkm<-,SCESet,matrix-method
  (fpkm), 18
fromCellDataSet, 19
ggplot, 26, 45
ggplot2, 26, 45
geom_smooth, 47
get_exprs, 22
get_exprs,SCESet-method(get_exprs), 22
get_exprs.SCESet
  (get_exprs), 22
getBM, 20
getBMFeatureAnnos, 20
getExprs, 21
is_exprs, 24
is_exprs,SCESet-method(is_exprs), 24
is_exprs<-,SCESet,method
  (is_exprs), 24
isOutlier, 23
lmFit, 31
mergeSCESet, 25
model.matrix, 31
mutate, 27
mutate,SCESet-method(mutate), 27
mutate.SCESet
  (mutate), 27
newSCESet, 27
nexprs, 29
normaliseExprs, 30
normalize, 32
normalize,SCESet-method(normalize), 32
normaliseExprs
  (normaliseExprs), 30
normaliseExprs(normaliseExprs), 30
norm_cpm, 34
norm_cpm,SCESet-method
  (norm_cpm), 34
norm_cpm.SCESet
  (norm_cpm), 34
norm_cpm<-
  (norm_cpm), 34
norm_exprs, 31
norm_exprs,SCESet-method
  (norm_exprs), 35
norm_exprs.SCESet
  (norm_exprs), 35
norm_exprs<-
  (norm_exprs), 35
norm_tpm, 37
norm_tpm,SCESet-method(norm_tpm), 37
norm_tpm.SCESet
  (norm_tpm), 37
norm_tpm<-
  (norm_tpm), 37
normalise(normalize), 32
normalise,SCESet-method(normalize), 32
normaliseExprs, 30
normalise, 32
normalise,SCESet-method(normalize), 32
normalize,SCESet(normalize), 32
normalizeExprs(normaliseExprs), 30
normliseExprs(normaliseExprs), 30
pairs, 43
pData
  (pData<-,SCESet,AnnotatedDataFrame-method), 38
pData,SCESet-method
  (pData<-,SCESet,AnnotatedDataFrame-method), 38
pData<-,SCESet,AnnotatedDataFrame-method
  (pData<-,SCESet,AnnotatedDataFrame-method), 38
pData<-,SCESet,DataFrame-method
  (pData<-,SCESet,AnnotatedDataFrame-method),
stand_exprs<- (stand_exprs), 79
stand_exprs<-, SCESet, matrix-method
(stand_exprs), 79
summariseExprsAcrossFeatures, 80
toCellDataSet, 82
tpm, 82
tpm, SCESet-method (tpm), 82
tpm.SCESet (tpm), 82
tpm<-(tpm), 82
tpm<-, SCESet, matrix-method (tpm), 82
tximport, 67

updateSCESet, 83
writeSCESet, 84