

Package ‘pcaExplorer’

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Type Package

Title Interactive Visualization of RNA-seq Data Using a Principal Components Approach

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Description This package provides functionality for interactive visualization of RNA-seq datasets based on Principal Components Analysis. The methods provided allow for quick information extraction and effective data exploration. A Shiny application encapsulates the whole analysis.

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Imports DESeq2, SummarizedExperiment, mosdef (>= 1.1.0), GenomicRanges, IRanges, S4Vectors, genefilter, ggplot2 (>= 2.0.0), heatmaply, plotly, scales, NMF, plyr, topGO, limma, GOstats, GO.db, AnnotationDbi, shiny (>= 0.12.0), shinydashboard, shinyBS, ggrepel, DT, shinyAce, threejs, biomaRt, pheatmap, knitr, rmarkdown, base64enc, tidyr, grDevices, methods

Suggests testthat, BiocStyle, markdown, airway, org.Hs.eg.db, htmltools

URL <https://github.com/federicomarini/pcaExplorer>,
<https://federicomarini.github.io/pcaExplorer/>

BugReports <https://github.com/federicomarini/pcaExplorer/issues>

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correlatePCs	<i>Principal components (cor)relation with experimental covariates</i>
--------------	--

Description

Computes the significance of (cor)relations between PCA scores and the sample experimental covariates, using Kruskal-Wallis test for categorial variables and the `cor.test` based on Spearman's correlation for continuous variables

Usage

```
correlatePCs(pcaobj, coldata, pcs = 1:4)
```

Arguments

<code>pcaobj</code>	A <code>prcomp</code> object
<code>coldata</code>	A <code>data.frame</code> object containing the experimental covariates
<code>pcs</code>	A numeric vector, containing the corresponding PC number

Value

A `data.frame` object with computed p values for each covariate and for each principal component

Examples

```
library(DESeq2)
dds <- makeExampleDESeqDataSet_multifac(betaSD_condition = 3, betaSD_tissue = 1)
r1t <- DESeq2::rlogTransformation(dds)
pcaobj <- prcomp(t(assay(r1t)))
correlatePCs(pcaobj, colData(dds))
```

deprecated

Deprecated functions in `pcaExplorer`

Description

Functions that are on their way to the function afterlife. Their successors are also listed.

Arguments

... Ignored arguments.

Details

The successors of these functions are likely coming after the rework that led to the creation of the `mosdef` package. See more into its documentation for more details.

Value

All functions throw a warning, with a deprecation message pointing towards its descendent (if available).

Transitioning to the `mosdef` framework

- `topG0table()` is now being replaced by the more flexible `mosdef::run_topG0()` function

Author(s)

Federico Marini

Examples

```
# try(topG0table())
```

distro_expr	<i>Plot distribution of expression values</i>
-------------	---

Description

Plot distribution of expression values

Usage

```
distro_expr(rld, plot_type = "density")
```

Arguments

rld	A <code>DESeq2::DESeqTransform()</code> object.
plot_type	Character, choose one of boxplot, violin or density. Defaults to density

Value

A plot with the distribution of the expression values

Examples

```
dds <- makeExampleDESeqDataSet_multifac(betaSD_condition = 3, betaSD_tissue = 1)
rld <- DESeq2::rlogTransformation(dds)
distro_expr(rld)
```

geneprofiler	<i>Extract and plot the expression profile of genes</i>
--------------	---

Description

Extract and plot the expression profile of genes

Usage

```
geneprofiler(se, genelist = NULL, intgroup = "condition", plotZ = FALSE)
```

Arguments

se	A <code>DESeq2::DESeqDataSet()</code> object, or a <code>DESeq2::DESeqTransform()</code> object.
genelist	An array of characters, including the names of the genes of interest of which the profile is to be plotted
intgroup	A factor, needs to be in the colnames of <code>colData(se)</code>
plotZ	Logical, whether to plot the scaled expression values. Defaults to FALSE

Value

A plot of the expression profile for the genes

Examples

```
dds <- makeExampleDESeqDataSet_multifac(betaSD_condition = 3, betaSD_tissue = 1)
r1t <- DESeq2::rlogTransformation(dds)
geneprofiler(r1t, paste0("gene", sample(1:1000, 20)))
geneprofiler(r1t, paste0("gene", sample(1:1000, 20)), plotZ = TRUE)
```

genespca

Principal components analysis on the genes

Description

Computes and plots the principal components of the genes, eventually displaying the samples as in a typical biplot visualization.

Usage

```
genespca(  
  x,  
  ntop,  
  choices = c(1, 2),  
  arrowColors = "steelblue",  
  groupNames = "group",  
  biplot = TRUE,  
  scale = 1,  
  pc.biplot = TRUE,  
  obs.scale = 1 - scale,  
  var.scale = scale,  
  groups = NULL,  
  ellipse = FALSE,  
  ellipse.prob = 0.68,  
  labels = NULL,  
  labels.size = 3,  
  alpha = 1,  
  var.axes = TRUE,  
  circle = FALSE,  
  circle.prob = 0.69,  
  varname.size = 4,  
  varname.adjust = 1.5,  
  varname.abbrev = FALSE,  
  returnData = FALSE,  
  coordEqual = FALSE,  
  scaleArrow = 1,  
  useRownamesAsLabels = TRUE,  
  point_size = 2,  
  annotation = NULL  
)
```

Arguments

x A `DESeq2::DESeqTransform()` object, with data in `assay(x)`, produced for example by either `DESeq2::rlog()` or `DESeq2::varianceStabilizingTransformation()`

<code>ntop</code>	Number of top genes to use for principal components, selected by highest row variance
<code>choices</code>	Vector of two numeric values, to select on which principal components to plot
<code>arrowColors</code>	Vector of character, either as long as the number of the samples, or one single value
<code>groupNames</code>	Factor containing the groupings for the input data. Is efficiently chosen as the (interaction of more) factors in the <code>colData</code> for the object provided
<code>biplot</code>	Logical, whether to additionally draw the samples labels as in a biplot representation
<code>scale</code>	Covariance biplot (<code>scale = 1</code>), form biplot (<code>scale = 0</code>). When <code>scale = 1</code> , the inner product between the variables approximates the covariance and the distance between the points approximates the Mahalanobis distance.
<code>pc.biplot</code>	Logical, for compatibility with <code>biplot.princomp()</code>
<code>obs.scale</code>	Scale factor to apply to observations
<code>var.scale</code>	Scale factor to apply to variables
<code>groups</code>	Optional factor variable indicating the groups that the observations belong to. If provided the points will be colored according to groups
<code>ellipse</code>	Logical, draw a normal data ellipse for each group
<code>ellipse.prob</code>	Size of the ellipse in Normal probability
<code>labels</code>	optional Vector of labels for the observations
<code>labels.size</code>	Size of the text used for the labels
<code>alpha</code>	Alpha transparency value for the points (0 = transparent, 1 = opaque)
<code>var.axes</code>	Logical, draw arrows for the variables?
<code>circle</code>	Logical, draw a correlation circle? (only applies when <code>prcomp</code> was called with <code>scale = TRUE</code> and when <code>var.scale = 1</code>)
<code>circle.prob</code>	Size of the correlation circle in Normal probability
<code>varname.size</code>	Size of the text for variable names
<code>varname.adjust</code>	Adjustment factor the placement of the variable names, ' <code>>= 1</code> ' means farther from the arrow
<code>varname.abbrev</code>	Logical, whether or not to abbreviate the variable names
<code>returnData</code>	Logical, if <code>TRUE</code> returns a <code>data.frame</code> for further use, containing the selected principal components for custom plotting
<code>coordEqual</code>	Logical, default <code>FALSE</code> , for allowing brushing. If <code>TRUE</code> , plot using equal scale cartesian coordinates
<code>scaleArrow</code>	Multiplicative factor, usually <code>>=1</code> , only for visualization purposes, to allow for distinguishing where the variables are plotted
<code>useRownamesAsLabels</code>	Logical, if <code>TRUE</code> uses the row names as labels for plotting
<code>point_size</code>	Size of the points to be plotted for the observations (genes)
<code>annotation</code>	A <code>data.frame</code> object, with <code>row.names</code> as gene identifiers (e.g. ENSEMBL ids) and a column, <code>gene_name</code> , containing e.g. HGNC-based gene symbols

Details

The implementation of this function is based on the beautiful ggbiplot package developed by Vince Vu, available at <https://github.com/vqv/ggbiplot>. The adaptation and additional parameters are tailored to display typical genomics data such as the transformed counts of RNA-seq experiments

Value

An object created by ggplot, which can be assigned and further customized.

Examples

```
library(DESeq2)
dds <- makeExampleDESeqDataSet_multifac(betaSD_condition = 3, betaSD_tissue = 1)
rlt <- rlogTransformation(dds)
groups <- colData(dds)$condition
groups <- factor(groups, levels = unique(groups))
cols <- scales::hue_pal()(2)[groups]
genespca(rlt, ntop=100, arrowColors = cols, groupNames = groups)

groups_multi <- interaction(as.data.frame(colData(rlt)[, c("condition", "tissue")]))
groups_multi <- factor(groups_multi, levels = unique(groups_multi))
cols_multi <- scales::hue_pal()(length(levels(groups_multi)))[factor(groups_multi)]
genespca(rlt, ntop = 100, arrowColors = cols_multi, groupNames = groups_multi)
```

get_annotation

Get an annotation data frame from biomaRt

Description

Get an annotation data frame from biomaRt

Usage

```
get_annotation(dds, biomaRt_dataset, idtype)
```

Arguments

dds	A <code>DESeq2::DESeqDataSet()</code> object
biomaRt_dataset	A biomaRt dataset to use. To see the list, type <code>mart = useMart('ensembl')</code> , followed by <code>listDatasets(mart)</code> .
idtype	Character, the ID type of the genes as in the row names of dds, to be used for the call to <code>biomaRt::getBM()</code>

Value

A data frame for ready use in `pcaExplorer`, retrieved from biomaRt.

Examples

```
library("airway")
data("airway", package = "airway")
airway
dds_airway <- DESeq2::DESeqDataSetFromMatrix(assay(airway),
                                           colData = colData(airway),
                                           design = ~dex+cell)

## Not run:
get_annotation(dds_airway, "hsapiens_gene_ensembl", "ensembl_gene_id")

## End(Not run)
```

get_annotation_orgdb *Get an annotation data frame from org db packages*

Description

Get an annotation data frame from org db packages

Usage

```
get_annotation_orgdb(dds, orgdb_species, idtype, key_for_genenames = "SYMBOL")
```

Arguments

dds	A DESeq2::DESeqDataSet() object
orgdb_species	Character string, named as the org.XX.eg.db package which should be available in Bioconductor
idtype	Character, the ID type of the genes as in the row names of dds, to be used for the call to AnnotationDbi::mapIds()
key_for_genenames	Character, corresponding to the column name for the key in the orgDb package containing the official gene name (often called gene symbol). This parameter defaults to "SYMBOL", but can be adjusted in case the key is not found in the annotation package (e.g. for org.Sc.sgd.db).

Value

A data frame for ready use in `pcaExplorer`, retrieved from the org db packages

Examples

```
library("airway")
data("airway", package = "airway")
airway
dds_airway <- DESeq2::DESeqDataSetFromMatrix(assay(airway),
                                           colData = colData(airway),
                                           design = ~dex+cell)

anno_df <- get_annotation_orgdb(dds_airway, "org.Hs.eg.db", "ENSEMBL")
head(anno_df)
```

hi_loadings	<i>Extract genes with highest loadings</i>
-------------	--

Description

Extract genes with highest loadings

Usage

```
hi_loadings(
  pcaobj,
  whichpc = 1,
  topN = 10,
  exprTable = NULL,
  annotation = NULL,
  title = "Top/bottom loadings"
)
```

Arguments

pcaobj	A prcomp object
whichpc	An integer number, corresponding to the principal component of interest
topN	Integer, number of genes with top and bottom loadings
exprTable	A matrix object, e.g. the counts of a <code>DESeq2::DESeqDataSet()</code> . If not NULL, returns the counts matrix for the selected genes
annotation	A data.frame object, with row.names as gene identifiers (e.g. ENSEMBL ids) and a column, gene_name, containing e.g. HGNC-based gene symbols
title	The title of the plot

Value

A ggplot2 object, or a matrix, if exprTable is not null

Examples

```
dds <- makeExampleDESeqDataSet_multifac(betaSD = 3, betaSD_tissue = 1)
r1t <- DESeq2::rlogTransformation(dds)
pcaobj <- prcomp(t(SummarizedExperiment::assay(r1t)))
hi_loadings(pcaobj, topN = 20)
hi_loadings(pcaobj, topN = 10, exprTable = dds)
hi_loadings(pcaobj, topN = 10, exprTable = counts(dds))
```

limmaquickpca2go	<i>Functional interpretation of the principal components, based on simple overrepresentation analysis</i>
------------------	---

Description

Extracts the genes with the highest loadings for each principal component, and performs functional enrichment analysis on them using the simple and quick routine provided by the `limma` package

Usage

```
limmaquickpca2go(
  se,
  pca_ngenes = 10000,
  inputType = "ENSEMBL",
  organism = "Mm",
  loadings_ngenes = 500,
  background_genes = NULL,
  scale = FALSE,
  ...
)
```

Arguments

<code>se</code>	A <code>DESeq2::DESeqTransform()</code> object, with data in <code>assay(se)</code> , produced for example by either <code>DESeq2::rlog()</code> or <code>DESeq2::varianceStabilizingTransformation()</code>
<code>pca_ngenes</code>	Number of genes to use for the PCA
<code>inputType</code>	Input format type of the gene identifiers. Defaults to <code>ENSEMBL</code> , that then will be converted to <code>ENTREZ</code> ids. Can assume values such as <code>ENTREZID</code> , <code>GENENAME</code> or <code>SYMBOL</code> , like it is normally used with the <code>select</code> function of <code>AnnotationDbi</code>
<code>organism</code>	Character abbreviation for the species, using <code>org.XX.eg.db</code> for annotation
<code>loadings_ngenes</code>	Number of genes to extract the loadings (in each direction)
<code>background_genes</code>	Which genes to consider as background.
<code>scale</code>	Logical, defaults to <code>FALSE</code> , scale values for the PCA
<code>...</code>	Further parameters to be passed to the <code>goana</code> routine

Value

A nested list object containing for each principal component the terms enriched in each direction. This object is to be thought in combination with the displaying feature of the main `pcaExplorer()` function

Examples

```
library("airway")
library("DESeq2")
library("limma")
data("airway", package = "airway")
```

```

airway
dds_airway <- DESeqDataSet(airway, design = ~ cell + dex)
## Not run:
rld_airway <- rlogTransformation(dds_airway)
goquick_airway <- limmaquickpca2go(rld_airway,
                                   pca_ngenes = 10000,
                                   inputType = "ENSEMBL",
                                   organism = "Hs")

## End(Not run)

```

```
makeExampleDESeqDataSet_multifac
```

Make a simulated DESeqDataSet for two or more experimental factors

Description

Constructs a simulated dataset of Negative Binomial data from different conditions. The fold changes between the conditions can be adjusted with the `betaSD_condition` and the `betaSD_tissue` arguments.

Usage

```

makeExampleDESeqDataSet_multifac(
  n = 1000,
  m = 12,
  betaSD_condition = 1,
  betaSD_tissue = 3,
  interceptMean = 4,
  interceptSD = 2,
  dispMeanRel = function(x) 4/x + 0.1,
  sizeFactors = rep(1, m)
)

```

Arguments

<code>n</code>	number of rows (genes)
<code>m</code>	number of columns (samples)
<code>betaSD_condition</code>	the standard deviation for condition betas, i.e. $\beta \sim N(0, \text{betaSD})$
<code>betaSD_tissue</code>	the standard deviation for tissue betas, i.e. $\beta \sim N(0, \text{betaSD})$
<code>interceptMean</code>	the mean of the intercept betas (log2 scale)
<code>interceptSD</code>	the standard deviation of the intercept betas (log2 scale)
<code>dispMeanRel</code>	a function specifying the relationship of the dispersions on $2^{\text{trueIntercept}}$
<code>sizeFactors</code>	multiplicative factors for each sample

Details

This function is designed and inspired following the proposal of `DESeq2::makeExampleDESeqDataSet()` from the DESeq2 package. Credits are given to Mike Love for the nice initial implementation

Value

a `DESeq2::DESeqDataSet()` with true dispersion, intercept for two factors (condition and tissue) and beta values in the metadata columns. Note that the true betas are provided on the log2 scale.

Examples

```
dds <- makeExampleDESeqDataSet_multifac(betaSD_condition = 3, betaSD_tissue = 1)
dds
dds2 <- makeExampleDESeqDataSet_multifac(betaSD_condition = 1, betaSD_tissue = 4)
dds2
```

pair_corr

Pairwise scatter and correlation plot of counts

Description

Pairwise scatter and correlation plot of counts

Usage

```
pair_corr(df, log = FALSE, method = "pearson", use_subset = TRUE)
```

Arguments

df	A data frame, containing the (raw/normalized/transformed) counts
log	Logical, whether to convert the input values to log2 (with addition of a pseudo-count). Defaults to FALSE.
method	Character string, one of pearson (default), kendall, or spearman as in cor
use_subset	Logical value. If TRUE, only 1000 values per sample will be used to speed up the plotting operations.

Value

A plot with pairwise scatter plots and correlation coefficients

Examples

```
library("airway")
data("airway", package = "airway")
airway
dds_airway <- DESeq2::DESeqDataSetFromMatrix(assay(airway),
                                             colData = colData(airway),
                                             design = ~dex+cell)
pair_corr(counts(dds_airway)[1:100, ]) # use just a subset for the example
```

pca2go

*Functional interpretation of the principal components***Description**

Extracts the genes with the highest loadings for each principal component, and performs functional enrichment analysis on them using routines and algorithms from the topGO package

Usage

```
pca2go(
  se,
  pca_ngenes = 10000,
  annotation = NULL,
  inputType = "geneSymbol",
  organism = "Mm",
  ensToGeneSymbol = FALSE,
  loadings_ngenes = 500,
  background_genes = NULL,
  scale = FALSE,
  return_ranked_gene_loadings = FALSE,
  annopkg = NULL,
  ...
)
```

Arguments

se	A <code>DESeq2::DESeqTransform()</code> object, with data in <code>assay(se)</code> , produced for example by either <code>DESeq2::rlog()</code> or <code>DESeq2::varianceStabilizingTransformation()</code>
pca_ngenes	Number of genes to use for the PCA
annotation	A <code>data.frame</code> object, with <code>row.names</code> as gene identifiers (e.g. ENSEMBL ids) and a column, <code>gene_name</code> , containing e.g. HGNC-based gene symbols
inputType	Input format type of the gene identifiers. Will be used by the routines of topGO
organism	Character abbreviation for the species, using <code>org.XX.eg.db</code> for annotation
ensToGeneSymbol	Logical, whether to expect ENSEMBL gene identifiers, to convert to gene symbols with the annotation provided
loadings_ngenes	Number of genes to extract the loadings (in each direction)
background_genes	Which genes to consider as background.
scale	Logical, defaults to FALSE, scale values for the PCA
return_ranked_gene_loadings	Logical, defaults to FALSE. If TRUE, simply returns a list containing the top ranked genes with hi loadings in each PC and in each direction
annopkg	String containing the name of the organism annotation package. Can be used to override the <code>organism</code> parameter, e.g. in case of alternative identifiers used in the annotation package (Arabidopsis with TAIR)
...	Further parameters to be passed to the topGO routine

Value

A nested list object containing for each principal component the terms enriched in each direction. This object is to be thought in combination with the displaying feature of the main `pcaExplorer()` function

Examples

```
library("airway")
library("DESeq2")
data("airway", package = "airway")
airway
dds_airway <- DESeqDataSet(airway, design= ~ cell + dex)
## Not run:
rld_airway <- rlogTransformation(dds_airway)
# constructing the annotation object
anno_df <- data.frame(gene_id = rownames(dds_airway),
                      stringsAsFactors = FALSE)
library("AnnotationDbi")
library("org.Hs.eg.db")
anno_df$gene_name <- mapIds(org.Hs.eg.db,
                           keys = anno_df$gene_id,
                           column = "SYMBOL",
                           keytype = "ENSEMBL",
                           multiVals = "first")
rownames(anno_df) <- anno_df$gene_id
bg_ids <- rownames(dds_airway)[rowSums(counts(dds_airway)) > 0]
library(topGO)
pca2go_airway <- pca2go(rld_airway,
                       annotation = anno_df,
                       organism = "Hs",
                       ensToGeneSymbol = TRUE,
                       background_genes = bg_ids)

## End(Not run)
```

pcaExplorer

Explore a dataset from a PCA perspective

Description

Launch a Shiny App for interactive exploration of a dataset from the perspective of Principal Components Analysis

Usage

```
pcaExplorer(
  dds = NULL,
  dst = NULL,
  countmatrix = NULL,
  coldata = NULL,
  pca2go = NULL,
  annotation = NULL,
```

```

    runLocal = TRUE
  )

```

Arguments

dds	A <code>DESeq2::DESeqDataSet()</code> object. If not provided, then a <code>countmatrix</code> and a <code>coldata</code> need to be provided. If none of the above is provided, it is possible to upload the data during the execution of the Shiny App
dst	A <code>DESeq2::DESeqTransform()</code> object. Can be computed from the <code>dds</code> object if left <code>NULL</code> . If none is provided, then a <code>countmatrix</code> and a <code>coldata</code> need to be provided. If none of the above is provided, it is possible to upload the data during the execution of the Shiny App
countmatrix	A count matrix, with genes as rows and samples as columns. If not provided, it is possible to upload the data during the execution of the Shiny App
coldata	A <code>data.frame</code> containing the info on the covariates of each sample. If not provided, it is possible to upload the data during the execution of the Shiny App
pca2go	An object generated by the <code>pca2go()</code> function, which contains the information on enriched functional categories in the genes that show the top or bottom loadings in each principal component of interest. If not provided, it is possible to compute live during the execution of the Shiny App
annotation	A <code>data.frame</code> object, with <code>row.names</code> as gene identifiers (e.g. ENSEMBL ids) and a column, <code>gene_name</code> , containing e.g. HGNC-based gene symbols
runLocal	A logical indicating whether the app is to be run locally or remotely on a server, which determines how documentation will be accessed.

Value

A Shiny App is launched for interactive data exploration

Examples

```

library("airway")
data("airway", package = "airway")
airway
dds_airway <- DESeq2::DESeqDataSetFromMatrix(assay(airway),
                                           colData = colData(airway),
                                           design = ~dex+cell)

## Not run:
rld_airway <- DESeq2::rlogTransformation(dds_airway)

pcaExplorer(dds_airway, rld_airway)

pcaExplorer(countmatrix = counts(dds_airway), coldata = colData(dds_airway))

pcaExplorer() # and then upload count matrix, covariate matrix (and eventual annotation)

## End(Not run)

```

pcaExplorer-pkg	<i>pcaExplorer: analyzing time-lapse microscopy imaging, from detection to tracking</i>
-----------------	---

Description

pcaExplorer provides functionality for interactive visualization of RNA-seq datasets based on Principal Components Analysis. The methods provided allow for quick information extraction and effective data exploration. A Shiny application encapsulates the whole analysis.

Details

pcaExplorer provides functionality for interactive visualization of RNA-seq datasets based on Principal Components Analysis. The methods provided allow for quick information extraction and effective data exploration. A Shiny application encapsulates the whole analysis.

Author(s)

Federico Marini <marinif@uni-mainz.de>, 2016
Maintainer: Federico Marini <marinif@uni-mainz.de>

See Also

Useful links:

- <https://github.com/federicomarini/pcaExplorer>
- <https://federicomarini.github.io/pcaExplorer/>
- Report bugs at <https://github.com/federicomarini/pcaExplorer/issues>

pcaPlot	<i>Sample PCA plot for transformed data</i>
---------	---

Description

Plots the results of PCA on a 2-dimensional space

Usage

```
pcaPlot(  
  x,  
  intgroup = NULL,  
  ntop = 500,  
  returnData = FALSE,  
  title = NULL,  
  pcX = 1,  
  pcY = 2,  
  text_labels = TRUE,  
  point_size = 3,  
  ellipse = TRUE,  
  ellipse.prob = 0.95  
)
```

Arguments

x	A <code>DESeq2::DESeqTransform()</code> object, with data in <code>assay(x)</code> , produced for example by either <code>DESeq2::rlog()</code> or <code>DESeq2::varianceStabilizingTransformation()/DESeq2::</code>
intgroup	Interesting groups: a character vector of names in <code>colData(x)</code> to use for grouping. Defaults to <code>NULL</code> , which would then select the first column of the <code>colData</code> slot
ntop	Number of top genes to use for principal components, selected by highest row variance
returnData	logical, if <code>TRUE</code> returns a data.frame for further use, containing the selected principal components and <code>intgroup</code> covariates for custom plotting
title	The plot title
pcX	The principal component to display on the x axis
pcY	The principal component to display on the y axis
text_labels	Logical, whether to display the labels with the sample identifiers
point_size	Integer, the size of the points for the samples
ellipse	Logical, whether to display the confidence ellipse for the selected groups
ellipse.prob	Numeric, a value in the interval <code>[0;1)</code>

Value

An object created by `ggplot`, which can be assigned and further customized.

Examples

```
dds <- makeExampleDESeqDataSet_multifac(betaSD_condition = 3, betaSD_tissue = 1)
rlt <- DESeq2::rlogTransformation(dds)
pcaplot(rlt, ntop = 200)
```

pcaplot3d

Sample PCA plot for transformed data

Description

Plots the results of PCA on a 3-dimensional space, interactively

Usage

```
pcaplot3d(
  x,
  intgroup = "condition",
  ntop = 500,
  returnData = FALSE,
  title = NULL,
  pcX = 1,
  pcY = 2,
  pcZ = 3,
  text_labels = TRUE,
  point_size = 3
)
```

Arguments

<code>x</code>	A <code>DESeq2::DESeqTransform()</code> object, with data in <code>assay(x)</code> , produced for example by either <code>DESeq2::rlog()</code> or <code>DESeq2::varianceStabilizingTransformation()</code>
<code>intgroup</code>	Interesting groups: a character vector of names in <code>colData(x)</code> to use for grouping
<code>ntop</code>	Number of top genes to use for principal components, selected by highest row variance
<code>returnData</code>	logical, if TRUE returns a data.frame for further use, containing the selected principal components and <code>intgroup</code> covariates for custom plotting
<code>title</code>	The plot title
<code>pcX</code>	The principal component to display on the x axis
<code>pcY</code>	The principal component to display on the y axis
<code>pcZ</code>	The principal component to display on the z axis
<code>text_labels</code>	Logical, whether to display the labels with the sample identifiers
<code>point_size</code>	Integer, the size of the points for the samples

Value

A html-based visualization of the 3d PCA plot

Examples

```
dds <- makeExampleDESeqDataSet_multifac(betaSD_condition = 3, betaSD_tissue = 1)
r1t <- DESeq2::rlogTransformation(dds)
pcaplot3d(r1t, ntop = 200)
```

pcascree

Scree plot of the PCA on the samples

Description

Produces a scree plot for investigating the proportion of explained variance, or alternatively the cumulative value

Usage

```
pcascree(obj, type = c("pev", "cev"), pc_nr = NULL, title = NULL)
```

Arguments

<code>obj</code>	A <code>prcomp</code> object
<code>type</code>	Display absolute proportions or cumulative proportion. Possible values: "pev" or "cev"
<code>pc_nr</code>	How many principal components to display max
<code>title</code>	Title of the plot

Value

An object created by `ggplot`, which can be assigned and further customized.

Examples

```

dds <- makeExampleDESeqDataSet_multifac(betaSD_condition = 3, betaSD_tissue = 1)
r1t <- DESeq2::rlogTransformation(dds)
pcaobj <- prcomp(t(SummarizedExperiment::assay(r1t)))
pcascree(pcaobj, type = "pev")
pcascree(pcaobj, type = "cev", title = "Cumulative explained proportion of variance - Test dataset")

```

plotPCcorrs	<i>Plot significance of (cor)relations of covariates VS principal components</i>
-------------	--

Description

Plots the significance of the (cor)relation of each covariate vs a principal component

Usage

```
plotPCcorrs(pccorrs, pc = 1, logp = TRUE)
```

Arguments

pccorrs	A data.frame object generated by correlatePCs
pc	An integer number, corresponding to the principal component of interest
logp	Logical, defaults to TRUE, displays the $-\log_{10}$ of the pvalue instead of the p value itself

Value

A base plot object

Examples

```

library(DESeq2)
dds <- makeExampleDESeqDataSet_multifac(betaSD_condition = 3, betaSD_tissue = 1)
r1t <- rlogTransformation(dds)
pcaobj <- prcomp(t(assay(r1t)))
res <- correlatePCs(pcaobj, colData(dds))
plotPCcorrs(res)

```

topGOTable

Extract functional terms enriched in the DE genes, based on topGO

Description

A wrapper for extracting functional GO terms enriched in the DE genes, based on the algorithm and the implementation in the topGO package

Usage

```
topGOTable(
  DEgenes,
  BGgenes,
  ontology = "BP",
  annot = annFUN.org,
  mapping = "org.Mm.eg.db",
  geneID = "symbol",
  topTablerows = 200,
  fullNamesInRows = TRUE,
  addGeneToTerms = TRUE,
  plotGraph = FALSE,
  plotNodes = 10,
  writeOutput = FALSE,
  outputFile = "",
  topGO_method2 = "elim",
  do_padj = FALSE
)
```

Arguments

DEgenes	A vector of (differentially expressed) genes
BGgenes	A vector of background genes, e.g. all (expressed) genes in the assays
ontology	Which Gene Ontology domain to analyze: BP (Biological Process), MF (Molecular Function), or CC (Cellular Component)
annot	Which function to use for annotating genes to GO terms. Defaults to annFUN.org
mapping	Which org.XX.eg.db to use for annotation - select according to the species
geneID	Which format the genes are provided. Defaults to symbol, could also be entrez or ENSEMBL
topTablerows	How many rows to report before any filtering
fullNamesInRows	Logical, whether to display or not the full names for the GO terms
addGeneToTerms	Logical, whether to add a column with all genes annotated to each GO term
plotGraph	Logical, if TRUE additionally plots a graph on the identified GO terms
plotNodes	Number of nodes to plot
writeOutput	Logical, if TRUE additionally writes out the result to a file
outputFile	Name of the file the result should be written into

topGO_method2	Character, specifying which of the methods implemented by topGO should be used, in addition to the classic algorithm. Defaults to elim
do_padj	Logical, whether to perform the adjustment on the p-values from the specific topGO method, based on the FDR correction. Defaults to FALSE, since the assumption of independent hypotheses is somewhat violated by the intrinsic DAG-structure of the Gene Ontology Terms

Details

Allowed values assumed by the topGO_method2 parameter are one of the following: elim, weight, weight01, lea, parentchild. For more details on this, please refer to the original documentation of the topGO package itself

Value

A table containing the computed GO Terms and related enrichment scores

Examples

```
library("airway")
library("DESeq2")
data("airway", package = "airway")
airway
dds_airway <- DESeqDataSet(airway, design= ~ cell + dex)
# Example, performing extraction of enriched functional categories in
# detected significantly expressed genes
## Not run:
dds_airway <- DESeq(dds_airway)
res_airway <- results(dds_airway)
library("AnnotationDbi")
library("org.Hs.eg.db")
res_airway$symbol <- mapIds(org.Hs.eg.db,
                           keys = row.names(res_airway),
                           column = "SYMBOL",
                           keytype = "ENSEMBL",
                           multiVals = "first")
res_airway$entrez <- mapIds(org.Hs.eg.db,
                           keys = row.names(res_airway),
                           column = "ENTREZID",
                           keytype = "ENSEMBL",
                           multiVals = "first")
resOrdered <- as.data.frame(res_airway[order(res_airway$padj),])
de_df <- resOrdered[resOrdered$padj < .05 & !is.na(resOrdered$padj),]
de_symbols <- de_df$symbol
bg_ids <- rownames(dds_airway)[rowSums(counts(dds_airway)) > 0]
bg_symbols <- mapIds(org.Hs.eg.db,
                    keys = bg_ids,
                    column = "SYMBOL",
                    keytype = "ENSEMBL",
                    multiVals = "first")

library(topGO)
topgoDE_airway <- topGOtable(de_symbols, bg_symbols,
                             ontology = "BP",
                             mapping = "org.Hs.eg.db",
                             geneID = "symbol")
```

```
## End(Not run)
```

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