

Package ‘ideal’

October 16, 2018

Type Package

Title Interactive Differential Expression AnaLysis

Version 1.4.0

Date 2018-04-20

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Description This package provides functions for an Interactive Differential Expression AnaLysis of RNA-sequencing datasets, to extract quickly and effectively information downstream the step of differential expression. A Shiny application encapsulates the whole package.

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LazyData TRUE

Depends topGO

Imports DESeq2, SummarizedExperiment, GenomicRanges, IRanges, S4Vectors, ggplot2 (>= 2.0.0), d3heatmap, pheatmap, pcaExplorer, IHW, gplots, UpSetR, goseq, stringr, plyr, dplyr, limma, GOstats, GO.db, AnnotationDbi, shiny (>= 0.12.0), shinydashboard, shinyBS, DT, rrentrez, rintrojs, knitr, rmarkdown, shinyAce, BiocParallel, grDevices, methods

Suggests testthat, BiocStyle, airway, org.Hs.eg.db, TxDb.Hsapiens.UCSC.hg38.knownGene, DEFormats, edgeR

URL <https://github.com/federicomarini/ideal>

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

biocViews GeneExpression, DifferentialExpression, RNASeq, Sequencing, Visualization, QualityControl, GUI, GeneSetEnrichment, ReportWriting

VignetteBuilder knitr

RoxygenNote 6.0.1

git_url <https://git.bioconductor.org/packages/ideal>

git_branch RELEASE_3_7

git_last_commit 3cc6d98

git_last_commit_date 2018-04-30

Date/Publication 2018-10-15

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deseqresult2DEgenes *Generate a tidy table with the DE genes from the results of DESeq*

Description

Generate a tidy table with the DE genes from the results of DESeq

Usage

```
deseqresult2DEgenes(deseqresult, FDR = 0.05)
```

Arguments

deseqresult	A DESeqResults object
FDR	Numeric value, the significance level for thresholding adjusted p-values

Value

A "tidy" data.frame with only genes marked as differentially expressed

Examples

```
# with simulated data...
library(DESeq2)
dds <- DESeq2::makeExampleDESeqDataSet(n=100, m=8, betaSD = 2)
dds <- DESeq(dds)
res <- results(dds)
deseqresult2DEgenes(res)
```

deseqresult2tbl *Generate a tidy table with the results of DESeq*

Description

Generate a tidy table with the results of DESeq

Usage

```
deseqresult2tbl(deseqresult)
```

Arguments

deseqresult A [DESeqResults](#) object

Value

A "tidy" data.frame with all genes

Examples

```
# with simulated data...
library(DESeq2)
dds <- DESeq2::makeExampleDESeqDataSet(n=100, m=8, betaSD = 1)
dds <- DESeq2::DESeq(dds)
res <- DESeq2::results(dds)
deseqresult2tbl(res)
```

ggplotCounts *Plot normalized counts for a gene*

Description

Plot for normalized counts of a single gene, with jittered points superimposed on the boxplot

Usage

```
ggplotCounts(dds, gene, intgroup = "condition", annotation_obj = NULL,
            transform = TRUE)
```

Arguments

dds A [DESeqDataSet](#) object.

gene A character, specifying the name of the gene to plot

intgroup Interesting groups: a character vector of names in colData(dds) to use for grouping

<code>annotation_obj</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. Optional.
<code>transform</code>	Logical value, corresponding whether to have log scale y-axis or not. Defaults to TRUE.

Details

Note: this function relies on the `plotCounts` function of DESeq2, therefore pseudocounts of 0.5 are added to each point

Value

An object created by `ggplot`

Examples

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

ggplotCounts(dds_airway,
             gene = "ENSG00000103196", # CRISPLD2 in the original publication
             intgroup = "dex")
```

Description

A wrapper for extracting functional GO terms enriched in a list of (DE) genes, based on the algorithm and the implementation in the `goseq` package

Usage

```
goseqTable(de.genes, assayed.genes, genome = "hg38", id = "ensGene",
           testCats = c("GO:BP", "GO:MF", "GO:CC"), FDR_GO_cutoff = 1, nTop = 200,
           orgDbPkg = "org.Hs.eg.db", addGeneToTerms = TRUE)
```

Arguments

<code>de.genes</code>	A vector of (differentially expressed) genes
<code>assayed.genes</code>	A vector of background genes, e.g. all (expressed) genes in the assays
<code>genome</code>	A string identifying the genome that genes refer to, as in the <code>goseq</code> function
<code>id</code>	A string identifying the gene identifier used by genes, as in the <code>goseq</code> function

testCats	A vector specifying which categories to test for over representation amongst DE genes - can be any combination of "GO:CC", "GO:BP", "GO:MF" & "KEGG"
FDR_GO_cutoff	Numeric value for subsetting the results
nTop	Number of categories to extract, and optionally process for adding genes to the respective
orgDbPkg	Character string, named as the org.XX.eg.db package which should be available in Bioconductor
addGeneToTerms	Logical, whether to add a column with all genes annotated to each GO term

Details

Note: the feature length retrieval is based on the [goseq](#) function, and requires that the corresponding TxDb packages are installed and available

Value

A table containing the computed GO Terms and related enrichment scores

Examples

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

res_subset <- deseqresult2DEgenes(res_airway)[1:100,]
myde <- res_subset$id
myassayed <- rownames(res_airway)

## Not run:
mygo <- goseqTable(myde,
                     myassayed,
                     testCats = "GO:BP",
                     addGeneToTerms = FALSE)
head(mygo)

## End(Not run)
```

Description

ideal makes differential expression analysis interactive, easy and reproducible. This function launches the main application included in the package.

Usage

```
ideal(dds_obj = NULL, res_obj = NULL, annotation_obj = NULL,
      countmatrix = NULL, expdesign = NULL)
```

Arguments

<code>dds_obj</code>	A <code>DESeqDataSet</code> object. If not provided, then a <code>countmatrix</code> and a <code>expdesign</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
<code>res_obj</code>	A <code>DESeqResults</code> object. If not provided, it can be computed during the execution of the application
<code>annotation_obj</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. If not provided, it can be constructed during the execution via the <code>org.eg.XX.db</code> packages - these need to be installed
<code>countmatrix</code>	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
<code>expdesign</code>	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

Value

A Shiny App is launched for interactive data exploration and differential expression analysis

Examples

```
# with simulated data...
library(DESeq2)
dds <- DESeq2::makeExampleDESeqDataSet(n=100, m=8)
cm <- counts(dds)
cd <- colData(dds)

# with the well known airway package...
library(airway)
data(airway)
airway
dds_airway <- DESeq2::DESeqDataSetFromMatrix(assay(airway),
                                              colData = colData(airway),
                                              design=~cell+dex)

## Not run:

ideal()
ideal(dds)
ideal(dds_airway)

dds_airway <- DESeq2::DESeq(dds_airway)
res_airway <- DESeq2::results(dds_airway)
ideal(dds_airway, res_airway)

## End(Not run)
```

ideal-pkg*ideal: Interactive Differential Expression Analysis*

Description

ideal makes differential expression analysis interactive, easy and reproducible. The analysis of RNA-seq datasets is guided by the Shiny app as main component of the package, which also provides a wide set of functions to efficiently extract information from the existing data. The app can be also deployed on a Shiny server, to allow its usage without any installation on the user's side.

Details

ideal makes differential expression analysis interactive, easy and reproducible. The analysis of RNA-seq datasets is guided by the Shiny app as main component of the package, which also provides a wide set of functions to efficiently extract information from the existing data. The app can be also deployed on a Shiny server, to allow its usage without any installation on the user's side.

Author(s)

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plot_ma*MA-plot from base means and log fold changes*

Description

MA-plot from base means and log fold changes, in the ggplot2 framework, with additional support to annotate genes if provided.

Usage

```
plot_ma(res_obj, FDR = 0.05, point_alpha = 0.2, sig_color = "red",
        annotation_obj = NULL, hlines = NULL, title = NULL,
        xlab = "mean of normalized counts - log10 scale", ylim = NULL,
        add_rug = TRUE, intgenes = NULL, intgenes_color = "steelblue",
        labels_intgenes = TRUE)
```

Arguments

<code>res_obj</code>	A DESeqResults object
<code>FDR</code>	Numeric value, the significance level for thresholding adjusted p-values
<code>point_alpha</code>	Alpha transparency value for the points (0 = transparent, 1 = opaque)
<code>sig_color</code>	Color to use to mark differentially expressed genes. Defaults to red
<code>annotation_obj</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. Optional
<code>hlines</code>	The y coordinate (in absolute value) where to draw horizontal lines, optional

title	A title for the plot, optional
xlab	X axis label, defaults to "mean of normalized counts - log10 scale"
ylim	Vector of two numeric values, Y axis limits to restrict the view
add_rug	Logical, whether to add rug plots in the margins
intgenes	Vector of genes of interest. Gene symbols if a symbol column is provided in res_obj , or else the identifiers specified in the row names
intgenes_color	The color to use to mark the genes on the main plot.
labels_intgenes	Logical, whether to add the gene identifiers/names close to the marked plots

Details

The genes of interest are to be provided as gene symbols if a **symbol** column is provided in **res_obj**, or else by using the identifiers specified in the row names

Value

An object created by ggplot

Examples

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

plot_ma(res_airway, FDR = 0.05, hlines = 1)

plot_ma(res_airway, FDR = 0.1,
        intgenes = c("ENSG00000103196", # CRISPLD2
                    "ENSG00000120129", # DUSP1
                    "ENSG00000163884", # KLF15
                    "ENSG00000179094") # PER1
      )
```

Description

Volcano plot for log fold changes and log p-values in the ggplot2 framework, with additional support to annotate genes if provided.

Usage

```
plot_volcano(res_obj, FDR = 0.05, ylim_up = NULL, vlines = NULL,
             title = NULL, intgenes = NULL, intgenes_color = "steelblue",
             labels_intgenes = TRUE)
```

Arguments

res_obj	A DESeqResults object
FDR	Numeric value, the significance level for thresholding adjusted p-values
ylim_up	Numeric value, Y axis upper limits to restrict the view
vlines	The x coordinate (in absolute value) where to draw vertical lines, optional
title	A title for the plot, optional
intgenes	Vector of genes of interest. Gene symbols if a symbol column is provided in res_obj, or else the identifiers specified in the row names
intgenes_color	The color to use to mark the genes on the main plot.
labels_intgenes	Logical, whether to add the gene identifiers/names close to the marked plots

Details

The genes of interest are to be provided as gene symbols if a symbol column is provided in res_obj, or else b< using the identifiers specified in the row names

Value

An object created by ggplot

Examples

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

plot_volcano(res_airway)
```

Description

This function tries to guess which separator was used in a text delimited file

Usage

```
sepguesser(file, sep_list = c(",","\\t",";"," "))
```

Arguments

- file** The name of the file which the data are to be read from
sep_list A vector containing the candidates for being identified as separators. Defaults to c(", ", "\t", ";", " ")

Value

A character value, corresponding to the guessed separator. One of "," (comma), "\t" (tab), ";" (semicolon), " " (whitespace)

Examples

```
sepguesser(system.file("extdata/design_commas.txt", package = "ideal"))
sepguesser(system.file("extdata/design_semicolons.txt", package = "ideal"))
sepguesser(system.file("extdata/design_spaces.txt", package = "ideal"))
mysep <- sepguesser(system.file("extdata/design_tabs.txt", package = "ideal"))

# to be used for reading in the same file, without having to specify the sep
```

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