Package 'DiffExp'

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Title Runs different methods of differential expression analysis			
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Description Runs different methods of differential expression analysis			
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R topics documented:			
get_contrast_condition_B_minus_A_factors_interaction get_contrast_vector opts_diffexp opts_prepro opts_prepro preprocess_object run_deseq2 run_edgeR run_limma tcga_brca			
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```
\label{lem:contrast_condition_B_minus_A_factors_interaction} Build\ a\ list\ of\ character\ contrasts\ of\ B-A\ for\ an\ interacting\ variable.
```

Description

Build a list of character contrasts of B-A for an interacting variable.

Usage

```
get_contrast_condition_B_minus_A_factors_interaction(
  condition,
  interaction,
  cond_name = "condition",
  inte_name = "interaction",
  level_B_cond = NULL,
  level_A_cond = NULL,
  cond_main_effect = T,
  inte_main_effect = F
)
```

Arguments

```
condition
                  the factor vector of condition values
interaction
                  the factor vector of interaction values
                  the name of the condition variable
cond_name
                  the name of the interaction variable
inte_name
                  the B level to be contrasted with A
level_B_cond
level_A_cond
                  the A level to be contrasted with B
cond_main_effect
                  is the condition a main effect
inte_main_effect
                  is the interaction a main effect
```

Author(s)

Yoann Pradat

Description

Produce a contrast vector

Usage

```
get_contrast_vector(contrast, design, data)
```

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Arguments

contrast a character vector specifying the contrast beween refined beta names (see refine_beta_names design a formula specifying the design a data.frame used for building the model matrix

Value

a character vector of refined beta names

Author(s)

Yoann Pradat

References

internal

opts_diffexp

Define the parameters specific to each differential analysis method.

Description

Define the parameters specific to each differential analysis method.

Usage

```
opts_diffexp(
  alpha = 0.1,
  ncores = 6,
  save_table = T,
  only_significant = T,
  folder_results = "./results",
  use_deseq2 = T,
  use_edgeR = T,
  use_limma = F,
   ...
)
```

Arguments

fdr level when adjusting for multiple testing. alpha number of cores available for doing parallel computations. Used in DESeq. ncores save_table boolean to decide whether to save tables in txt files or not only_significant boolean to decide whether only significant (FDR) variables are kept in the results tables or not use_deseq2 boolean to choose to run DESeq2. use_edgeR boolean to choose to run edgeR. use_limma boolean to choose to run limma. extra parameters added to the configuration list . . .

opts_prepro

Value

a list

Author(s)

Yoann Pradat

opts_prepro

Define the options for preprocess_object

Description

See edgeR::filterByExp for more details about the filtering options and edgeR::calcNormFactors for mode details about the normalization options.

Usage

```
opts_prepro(
  design = NULL,
  min_count = 0,
  min_total_count = 15,
  large_n = 10,
  min_prop = 0.7,
  norm_factors_method = c("TMM", "TMMwsp", "RLE", "upperquartile", "none"),
  ...
)
```

Arguments

Value

a list

Author(s)

Yoann Pradat

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Description

Perform preprocessing steps.

Usage

```
preprocess_object(object, opts)
```

Arguments

object a SummarizedExperiment object

opts a named list of options. See opts_prepro

Value

a SummarizedExperiment object

Author(s)

Yoann Pradat

run_deseq2 Run DESEQ2 algorithm.

Description

Run DESEQ2 algorithm.

Usage

```
run_deseq2(object, design = NULL, contrasts, opts_algo, opts_comm)
```

Arguments

object a SummarizedExperiment object

design a formula specifying the design for the model matrix of DESeq2. Any variable appearing should be present in the colData of object

contrasts a character vector specifying the contrasts (one or multiple beta coefficient) to be used for making tests and building results table.

opts_algo a named list of options specific to DESeq2

opts_comm a summarizedExperiment object

a formula specifying the design for the model matrix of DESeq2. Any variable appearing should be present in the colData of object

a character vector specifying the contrasts (one or multiple beta coefficient) to be used for making tests and building results table.

Value

a dataframe of results

run_edgeR

Author(s)

Yoann Pradat

References

Love, M.I., Huber, W., Anders, S. (2014) Moderated estimation of fold change and dispersion for RNA-seq data with DESeq2. Genome Biology, 15:550. https://doi.org/10.1186/s13059-014-0550-8

run_edgeR	Run edgeR algorithm.
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Description

Run edgeR algorithm.

Usage

```
run_edgeR(object, design = NULL, contrasts, opts_algo, opts_comm)
```

Arguments

object	a SummarizedExperiment object
design	a formula specifying the design for the model matrix of DESeq2. Any variable appearing should be present in the colData of object
contrasts	a character vector specifying the contrasts (one or multiple beta coefficient) to be used for making tests and building results table.
opts_algo	a named list of options specific to edgeR
opts_comm	a named list of options common to all methods

Value

a dataframe of results

Author(s)

Yoann Pradat

References

Robinson MD, McCarthy DJ, Smyth GK (2010). "edgeR: a Bioconductor package for differential expression analysis of digital gene expression data." Bioinformatics, 26(1), 139-140. https://doi.org/10.1093/bioinformatics/btp616

McCarthy DJ, Chen Y, Smyth GK (2012). "Differential expression analysis of multifactor RNA-Seq experiments with respect to biological variation." Nucleic Acids Research, 40(10), 4288-4297. https://doi.org/10.1093/nar/gks042

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run_limma

Run limma algorithm.

Description

Run limma algorithm.

Usage

run_limma(object)

Author(s)

Yoann Pradat

References

Ritchie ME, Phipson B, Wu D, Hu Y, Law CW, Shi W, Smyth GK (2015). "limma powers differential expression analyses for RNA-sequencing and microarray studies." Nucleic Acids Research, 43(7), e47. https://doi.org/10.1093/nar/gkv007

Law, C.W., Chen, Y., Shi, W. et al. voom: precision weights unlock linear model analysis tools for RNA-seq read counts. Genome Biol 15, R29 (2014). https://doi.org/10.1186/gb-2014-15-2-r29

tcga_brca

RNA-seq and clinical data for TCGA BRCA Stage I patients.

Description

Data from the reanalysis of RNA-seq files by Zheng et al. 2019. The clinical data were taken from the GDC data portal and the cBio data portal. Both the RNA-seq data and the clinical data are public access. The samples were chosen so that

- 1. all samples are Stage I
- 2. all samples are Female
- 3. all samples have complete clinical data

Usage

```
data(tcga_brca)
```

Format

A SummarizedExperiment object with 58,288 genes and 98 samples.

assays A SimpleList with just one "counts" matrix colData A DFrame with 98 rows and 8 columns rowData A DFrame with 58,288 rows and 2 columns metadata An empty list elementMetadata Alias for rowData

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Details

This selection was performed only to keep this example data set small and may not be suitable for meaningful analyses.

Source

RNA-seq source: https://stanfordmedicine.app.box.com/s/lu703xuaulfz02vgd2lunxnvt4mfvo3q.cBioPortal BRCA Pan-cancer Atlas https://www.cbioportal.org/study/summary?id=brca_tcga_pan_can_atlas_2018. GDC legacy archive clinical BRCA https://portal.gdc.cancer.gov/legacy-archive/files/735bc5ff-86d1-421a-8693-6e6f92055563

References

Hong Zheng, Kevin Brennan, Mikel Hernaez, Olivier Gevaert, Benchmark of long non-coding RNA quantification for RNA sequencing of cancer samples, GigaScience, Volume 8, Issue 12, December 2019, giz145, https://doi.org/10.1093/gigascience/giz145

Examples

data(tcga_brca)

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