

Package ‘dearseq’

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Type Package

Title Differential Expression Analysis for RNA-seq data through a robust variance component test

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Imports CompQuadForm, ggplot2, KernSmooth, matrixStats, methods, parallel, pbapply, stats, statmod

Suggests Biobase, BiocManager, BiocSet, edgeR, DESeq2, GEOquery, GSA, knitr, limma, readxl, rmarkdown, S4Vectors, SummarizedExperiment, testthat, covr

Description Differential Expression Analysis RNA-seq data with variance component score test accounting for data heteroscedasticity through precision weights. Perform both gene-wise and gene set analyses, and can deal with repeated or longitudinal data. Methods are detailed in: Agniel D & Hejblum BP (2017) Variance component score test for time-course gene set analysis of longitudinal RNA-seq data, *Biostatistics*, 18(4):589-604. and Gauthier M, Agniel D, Thiébaud R & Hejblum BP (2019). dearseq: a variance component score test for RNA-Seq differential analysis that effectively controls the false discovery rate, **bioRxiv** 635714.

LazyData true

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biocViews BiomedicalInformatics, CellBiology, DifferentialExpression, DNASEq, GeneExpression, Genetics, GeneSetEnrichment, ImmunoOncology, KEGG, Regression, RNASeq, Sequencing, SystemsBiology, TimeCourse, Transcription, Transcriptomics

BugReports <https://github.com/borishejblum/dearseq/issues>

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dearseq-package	<i>dearseq: Differential Expression Analysis for RNA-seq data through a robust variance component test</i>
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Description

Differential Expression Analysis RNA-seq data with variance component score test accounting for data heteroscedasticity through precision weights. Perform both gene-wise and gene set analyses, and can deal with repeated or longitudinal data. Methods are detailed in: Agniel D & Hejblum BP (2017) Variance component score test for time-course gene set analysis of longitudinal RNA-seq data, *Biostatistics*, 18(4):589-604. and Gauthier M, Agniel D, Thiébaud R & Hejblum BP (2019). *dearseq: a variance component score test for RNA-Seq differential analysis that effectively controls the false discovery rate*, *bioRxiv* 635714.

Details

Analysis of RNA-seq data with variance component score test accounting for data heteroscedasticity through precision weights. Performs gene-wise analysis as well as gene set analysis, including for complex experimental designs such as longitudinal data.

Package: dearseq
Type: Package
Version: 1.0.0
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The two main functions of the dearseq package are `dear_seq` and `dgsa_seq`.

Author(s)

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References

Agniel D & Hejblum BP (2017). Variance component score test for time-course gene set analysis of longitudinal RNA-seq data, *Biostatistics*, 18(4):589-604. DOI: [10.1093/biostatistics/kxx005](https://doi.org/10.1093/biostatistics/kxx005). [arXiv:1605.02351](https://arxiv.org/abs/1605.02351).

Gauthier M, Agniel D, Thiébaud R & Hejblum BP (2019). dearseq: a variance component score test for RNA-Seq differential analysis that effectively controls the false discovery rate, *635714*. DOI: [10.1101/635714v1](https://doi.org/10.1101/635714v1)

See Also

Useful links:

- Report bugs at <https://github.com/borishejblum/dearseq/issues>

baduel_5gs

Small portion of RNA-seq data from plant physiology study.

Description

A subsample of the RNA-seq data from Baduel *et al.* studying Arabidopsis Arenosa physiology.

Usage

```
data(baduel_5gs)
```

Format

3 objects

- design: a design matrix for the 48 measured samples, containing the following variables:
 - SampleName corresponding column names from `expr_norm_corr`
 - Intercept an intercept variable
 - Population a factor identifying the plant population
 - Age_weeks numeric age of the plant at sampling time (in weeks)
 - Replicate a purely technical variable as replicates are not from the same individual over weeks. Should not be used in analysis.
 - Vernalized a logical variable indicating whether the plant had undergone vernalization (exposition to cold and short day photoperiods)
 - Vernalized a binary variable indicating whether the plant belonged to the KA population

- AgeWeeks_Population interaction variable between the AgeWeeks and Population variables
- AgeWeeks_Vernalized interaction variable between the AgeWeeks and Vernalized variables
- Vernalized_Population interaction variable between the Vernalized and Population variables
- AgeWeeks_Vernalized_Population interaction variable between the AgeWeeks, Vernalized and Population variables
- baduel_gmt: a gmt object containing 5 gene sets of interest (see [GSA.read.gmt](#)), which is simply a list with the 3 following components:
 - genesets: a list of n gene identifiers vectors composing each gene set (each gene set is represented as the vector of the gene identifiers composing it)
 - geneset.names: a vector of length n containing the gene set names (i.e. gene sets identifiers)
 - geneset.descriptions: a vector of length n containing gene set descriptions (e.g. textual information on their biological function)
- expr_norm_corr: a numeric matrix containing the normalized batch corrected expression for the 2454 genes included in either of the 5 gene sets of interests

Source

<http://www.ncbi.nlm.nih.gov/bioproject/PRJNA312410>

References

Baduel P, Arnold B, Weisman CM, Hunter B & Bomblies K (2016). Habitat-Associated Life History and Stress-Tolerance Variation in *Arabidopsis Arenosa*. *Plant Physiology*, 171(1):437-51. [10.1104/pp.15.01875](#).

Agniel D & Hejblum BP (2017). Variance component score test for time-course gene set analysis of longitudinal RNA-seq data, *Biostatistics*, 18(4):589-604. [10.1093/biostatistics/kxx005](#). [arXiv:1605.02351](#).

Examples

```
if(interactive()){
data('baduel_5gs')

set.seed(54321)
KAvstBG <- dgsa_seq(exprmat=log2(expr_norm_corr+1),
                   covariates=apply(as.matrix(design[,
c('Intercept', 'Vernalized', 'AgeWeeks', 'Vernalized_Population',
'AgeWeeks_Population')], drop=FALSE)], 2, as.numeric),
                   variables2test =
                     as.matrix(design[, c('PopulationKA'), drop=FALSE]),
                   genesets=baduel_gmt$genesets[c(3,5)],
                   which_test = 'permutation', which_weights = 'loclin',
                   n_perm=1000, preprocessed = TRUE)

set.seed(54321)
Cold <- dgsa_seq(exprmat=log2(expr_norm_corr+1),
                 covariates=apply(as.matrix(design[,
c('Intercept', 'AgeWeeks', 'PopulationKA', 'AgeWeeks_Population')],
drop=FALSE)], 2, as.numeric),
                 variables2test=as.matrix(design[, c('Vernalized',
```

```

    'Vernalized_Population')]],
  genesets=baduel_gmt$genesets[c(3,5)],
  which_test = 'permutation', which_weights = 'loclin',
  n_perm=1000, preprocessed = TRUE)
}

```

dear_seq

Differential expression analysis of RNA-seq data through a variance component test

Description

Wrapper function for gene-by-gene association testing of RNA-seq data

Usage

```

dear_seq(
  exprmat = NULL,
  object = NULL,
  covariates = NULL,
  variables2test,
  sample_group = NULL,
  weights_var2test_condi = TRUE,
  cov_variables2test_eff = NULL,
  which_test = c("permutation", "asymptotic"),
  which_weights = c("loclin", "voom", "none"),
  n_perm = 1000,
  progressbar = TRUE,
  parallel_comp = TRUE,
  nb_cores = parallel::detectCores() - 1,
  preprocessed = FALSE,
  gene_based_weights = FALSE,
  bw = "nrd",
  kernel = c("gaussian", "epanechnikov", "rectangular", "triangular", "biweight",
    "tricube", "cosine", "optcosine"),
  exact = FALSE,
  transform = TRUE,
  padjust_methods = c("BH", "BY", "holm", "hochberg", "hommel", "bonferroni"),
  lowess_span = 0.5,
  R = NULL,
  homogen_traj = FALSE,
  na.rm_dearseq = TRUE
)

```

Arguments

exprmat a numeric matrix of size $G \times n$ containing the raw RNA-seq counts or preprocessed expressions from n samples for G genes. Default is NULL, in which case object must not be NULL.

object	an object that can be either a SummarizedExperiment , an ExpressionSet , a DESeqDataSet , or a DGEList . Default is NULL, in which case exprmat must not be NULL.
covariates	<ul style="list-style-type: none"> • If exprmat is specified as a matrix: then covariates must be a numeric matrix of size $n \times p$ containing the model covariates for n samples (design matrix). Usually, its first column is the intercept (full of 1s). • If object is specified: then covariates must be a character vector of length p containing the colnames of the design matrix given in object. <p>If covariates is NULL (the default), then it is just the intercept.</p>
variables2test	<ul style="list-style-type: none"> • If exprmat is specified as a matrix: a numeric design matrix of size $n \times K$ containing the K variables to be tested. • If object is specified: then variables2test must be a character vector of length K containing the colnames of the design matrix given in object.
sample_group	a vector of length n indicating whether the samples should be grouped (e.g. paired samples or longitudinal data). Coerced to be a factor. Default is NULL in which case no grouping is performed.
weights_var2test_condi	a logical flag indicating whether heteroscedasticity weights computation should be conditional on both the variables to be tested variables2test and on the covariates, or on covariates alone. Default is TRUE in which case conditional means are estimated conditionally on both variables2test and covariates.
cov_variables2test_eff	a matrix of size $K \times K$ containing the covariance matrix of the K random effects. Only used if homogen_traj is FALSE. Default assume diagonal correlation matrix, i.e. independence of random effects.
which_test	a character string indicating which method to use to approximate the variance component score test, either 'permutation' or 'asymptotic'. Default is 'permutation'.
which_weights	a character string indicating which method to use to estimate the mean-variance relationship weights. Possibilities are 'loclin', 'voom' or 'none' (in which case no weighting is performed). Default is 'loclin'. See sp_weights and voom_weights for details.
n_perm	the number of perturbations. Default is 1000
progressbar	logical indicating whether a progressBar should be displayed when computing permutations (only in interactive mode).
parallel_comp	a logical flag indicating whether parallel computation should be enabled. Only Linux and MacOS are supported, this is ignored on Windows. Default is TRUE.
nb_cores	an integer indicating the number of cores to be used when parallel_comp is TRUE. Only Linux and MacOS are supported, this is ignored on Windows. Default is <code>parallel::detectCores() - 1</code> .
preprocessed	a logical flag indicating whether the expression data have already been preprocessed (e.g. log2 transformed). Default is FALSE, in which case y is assumed to contain raw counts and is normalized into $\log(\text{counts})$ per million.
gene_based_weights	a logical flag used for 'loclin' weights, indicating whether to estimate weights at the gene-level, or rather at the observation-level. Default is FALSE, which is what it should be for gene-wise analysis.
bw	a character string indicating the smoothing bandwidth selection method to use. See bandwidth for details. Possible values are 'ucv', 'SJ', 'bcv', 'nrd' or 'nrd0'.

kernel	a character string indicating which kernel should be used. Possibilities are 'gaussian', 'epanechnikov', 'rectangular', 'triangular', 'biweight', 'tricube', 'cosine', 'optcosine'. Default is 'gaussian' (NB: 'tricube' kernel corresponds to the loess method).
exact	a logical flag indicating whether the non-parametric weights accounting for the mean-variance relationship should be computed exactly or extrapolated from the interpolation of local regression of the mean against the variance. Default is FALSE, which uses interpolation (faster computation).
transform	a logical flag used for 'locclin' weights, indicating whether values should be transformed to uniform for the purpose of local linear smoothing. This may be helpful if tail observations are sparse and the specified bandwidth gives suboptimal performance there. Default is TRUE.
padjust_methods	multiple testing correction method used if genesets is a list. Default is 'BH', i.e. Benjamini-Hochberg procedure for controlling the FDR. Other possibilities are: 'holm', 'hochberg', 'hommel', 'bonferroni' or 'BY' (for Benjamini-Yekutieli procedure).
lowess_span	smoother span for the lowess function, between 0 and 1. This gives the proportion of points in the plot which influence the smooth at each value. Larger values give more smoothness. Only used if which_weights is 'vroom'. Default is 0.5.
R	library.size (optional, important to provide if preprocessed = TRUE). Default is NULL
homogen_traj	a logical flag indicating whether trajectories should be considered homogeneous. Default is FALSE in which case trajectories are not only tested for trend, but also for heterogeneity.
na.rm_dearseq	logical: should missing values in y (including NA and NaN) be omitted from the calculations? Default is FALSE.

Value

A list with the following elements:

- `which_test`: a character string carrying forward the value of the 'which_test' argument indicating which test was performed (either 'asymptotic' or 'permutation').
- `preprocessed`: a logical flag carrying forward the value of the 'preprocessed' argument indicating whether the expression data were already preprocessed, or were provided as raw counts and transformed into log-counts per million.
- `n_perm`: an integer carrying forward the value of the 'n_perm' argument indicating the number of perturbations performed (NA if asymptotic test was performed).
- `genesets`: carrying forward the value of the 'genesets' argument defining the gene sets of interest (NULL for gene-wise testing).
- `pval`: computed p-values. A data.frame with one row for each gene set, or for each gene if genesets argument is NULL, and with 2 columns: the first one 'rawPval' contains the raw p-values, the second one contains the FDR adjusted p-values (according to the 'padjust_methods' argument) and is named 'adjPval'.

References

Gauthier M, Agniel D, Thiébaud R & Hejblum BP (2019). dearseq: a variance component score test for RNA-Seq differential analysis that effectively controls the false discovery rate, **bioRxiv** 635714. [DOI: 10.1101/635714v1](https://www.biorxiv.org/content/10.1101/635714v1).

See Also

[sp_weights](#) [vc_test_perm](#) [vc_test_asym](#) [p.adjust](#)

Examples

```
#Monte-Carlo estimation of the proportion of DE genes over `nsims` simulations under the null

#number of runs
nsims <- 2 #100
res <- numeric(nsims)
for(i in 1:nsims){
  n <- 1000 #number of genes
  nr=5 #number of measurements per subject (grouped data)
  ni=50 #number of subjects
  r <- nr*ni #number of measurements
  t <- matrix(rep(1:nr), ni, ncol=1, nrow=r) # the variable to be tested
  sigma <- 0.5
  b0 <- 1

  #under the null:
  b1 <- 0

  #create the matrix of gene expression
  y.tilde <- b0 + b1*t + rnorm(r, sd = sigma)
  y <- t(matrix(rnorm(n*r, sd = sqrt(sigma*abs(y.tilde))), ncol=n, nrow=r) +
        matrix(rep(y.tilde, n), ncol=n, nrow=r))

  #no covariates
  x <- matrix(1, ncol=1, nrow=r)

  #run test
  #asymptotic test with preprocessed grouped data
  res_genes <- dear_seq(exprmat=y, covariates=x, variables2test=t,
                      sample_group=rep(1:ni, each=nr),
                      which_test='asymptotic',
                      which_weights='none', preprocessed=TRUE)

  #proportion of raw p-values>0.05
  mean(res_genes$pvals[, 'rawPval']>0.05)

  #quantiles of raw p-values
  quantile(res_genes$pvals[, 'rawPval'])

  #proportion of raw p-values<0.05 i.e. proportion of DE genes
  res[i] <- mean(res_genes$pvals[, 'rawPval']<0.05)
  message(i)
}

#results
mean(res)

if(interactive()){
  b0 <- 1
  #under the null:
  b1 <- 0
```



```

#create the matrix of gene expression
y.tilde <- b0 + b1*t + rnorm(r, sd = sigma)
y <- t(matrix(rnorm(n*r, sd = sqrt(sigma*abs(y.tilde))), ncol=n, nrow=r) +
        matrix(rep(y.tilde, n), ncol=n, nrow=r))

#run test
#asymptotic test with preprocessed grouped data
res_genes <- dear_seq(exprmat=y, covariates=x, variables2test=t,
                    sample_group=rep(1:ni, each=nr),
                    which_weights='none', preprocessed=TRUE)

#results
summary(res_genes$pvals)
}

```

dgsa_seq

Time-course Gene Set Analysis

Description

Wrapper function for performing gene set analysis of (potentially longitudinal) RNA-seq data

Usage

```

dgsa_seq(
  exprmat = NULL,
  object = NULL,
  covariates = NULL,
  variables2test,
  weights_var2test_condi = TRUE,
  genesets,
  sample_group = NULL,
  cov_variables2test_eff = NULL,
  which_test = c("permutation", "asymptotic"),
  which_weights = c("loclin", "voom", "none"),
  n_perm = 1000,
  progressbar = TRUE,
  parallel_comp = TRUE,
  nb_cores = parallel::detectCores() - 1,
  preprocessed = FALSE,
  gene_based_weights = TRUE,
  bw = "nrd",
  kernel = c("gaussian", "epanechnikov", "rectangular", "triangular", "biweight",
            "tricube", "cosine", "optcosine"),
  exact = FALSE,
  transform = TRUE,
  padjust_methods = c("BH", "BY", "holm", "hochberg", "hommel", "bonferroni"),
  lowess_span = 0.5,
  R = NULL,
  homogen_traj = FALSE,
  na.rm_gsaseq = TRUE,

```

```

    verbose = TRUE
  )

```

Arguments

- exprmat** a numeric matrix of size $G \times n$ containing the raw RNA-seq counts or preprocessed expressions from n samples for G genes. Default is NULL, in which case object must not be NULL.
- object** an object that can be either an [SummarizedExperiment](#), an [ExpressionSet](#), a [DESeqDataSet](#), or a [DGEList](#). Default is NULL, in which case exprmat must not be NULL.
- covariates**
 - If exprmat is specified as a matrix: then covariates must be a numeric matrix of size $n \times p$ containing the model covariates for n samples (design matrix). Usually, its first column is the intercept (full of 1s).
 - If object is specified: then covariates must be a character vector of length p containing the colnames of the design matrix given in object.
 If covariates is NULL (the default), then it is just the intercept.
- variables2test**
 - If exprmat is specified as a matrix: a numeric design matrix of size $n \times K$ containing the K variables to be tested.
 - If object is specified: then variables2test must be a character vector of length K containing the colnames of the design matrix given in object.
- weights_var2test_condi** a logical flag indicating whether heteroscedasticity weights computation should be conditional on both the variable(s) to be tested ϕ_i and on covariate(s) x , or on x alone. Default is TRUE in which case conditional means are estimated conditionally on both x and ϕ_i .
- genesets** Can be either:
 - a vector
 - a list
 - a [BiocSet](#) object
 Can be a vector of index or subscripts that defines which rows of y constitute the investigated gene set (when only 1 gene set is being tested).
 Can also be a list of index (or rownames of y) when several gene sets are tested at once, such as the first element of a [gmt](#) object.
 Finally, can also be a [BiocSet](#) object
 If NULL, then gene-wise p-values are returned.
- sample_group** a vector of length n indicating whether the samples should be grouped (e.g. paired samples or longitudinal data). Coerced to be a factor. Default is NULL in which case no grouping is performed.
- cov_variables2test_eff** a matrix of size $K \times K$ containing the covariance matrix of the K random effects. Only used if `homogen_traj` is FALSE. Default assume diagonal correlation matrix, i.e. independence of random effects.
- which_test** a character string indicating which method to use to approximate the variance component score test, either 'permutation' or 'asymptotic'. Default is 'permutation'.
- which_weights** a character string indicating which method to use to estimate the mean-variance relationship weights. Possibilities are 'loclin', 'voom' or 'none' (in which case no weighting is performed). Default is 'loclin'. See [sp_weights](#) and [voom_weights](#) for details.

n_perm	the number of perturbations. Default is 1000.
progressbar	logical indicating whether a progressBar should be displayed when computing permutations (only in interactive mode).
parallel_comp	a logical flag indicating whether parallel computation should be enabled. Only Linux and MacOS are supported, this is ignored on Windows. Default is TRUE.
nb_cores	an integer indicating the number of cores to be used when parallel_comp is TRUE. Only Linux and MacOS are supported, this is ignored on Windows. Default is <code>parallel::detectCores() - 1</code> .
preprocessed	a logical flag indicating whether the expression data have already been preprocessed (e.g. log2 transformed). Default is FALSE, in which case y is assumed to contain raw counts and is normalized into log(counts) per million.
gene_based_weights	a logical flag used for 'locLin' weights, indicating whether to estimate weights at the gene-level, or rather at the observation-level. Default is TRUE, and weights are then estimated at the gene-level.
bw	a character string indicating the smoothing bandwidth selection method to use. See bandwidth for details. Possible values are 'ucv', 'SJ', 'bcv', 'nrd' or 'nrd0'
kernel	a character string indicating which kernel should be used. Possibilities are 'gaussian', 'epanechnikov', 'rectangular', 'triangular', 'biweight', 'tricube', 'cosine', 'optcosine'. Default is 'gaussian' (NB: 'tricube' kernel corresponds to the loess method).
exact	a logical flag indicating whether the non-parametric weights accounting for the mean-variance relationship should be computed exactly or extrapolated from the interpolation of local regression of the mean against the variance. Default is FALSE, which uses interpolation (faster computation).
transform	a logical flag used for 'locLin' weights, indicating whether values should be transformed to uniform for the purpose of local linear smoothing. This may be helpful if tail observations are sparse and the specified bandwidth gives suboptimal performance there. Default is TRUE.
padjust_methods	multiple testing correction method used if genesets is a list. Default is 'BH', i.e. Benjamini-Hochberg procedure for controlling the FDR. Other possibilities are: 'holm', 'hochberg', 'hommel', 'bonferroni' or 'BY' (for Benjamini-Yekutieli procedure).
lowess_span	smoother span for the lowess function, between 0 and 1. This gives the proportion of points in the plot which influence the smooth at each value. Larger values give more smoothness. Only used if which_weights is 'voom'. Default is 0.5.
R	library size (optional, important to provide if preprocessed = TRUE). Default is NULL
homogen_traj	a logical flag indicating whether trajectories should be considered homogeneous. Default is FALSE in which case trajectories are not only tested for trend, but also for heterogeneity.
na.rm_gsaseq	logical: should missing values in y (including NA and NaN) be omitted from the calculations? Default is TRUE.
verbose	logical: should informative messages be printed during the computation? Default is TRUE.

Value

A list with the following elements:

- `which_test`: a character string carrying forward the value of the `'which_test'` argument indicating which test was performed (either `'asymptotic'` or `'permutation'`).
- `preprocessed`: a logical flag carrying forward the value of the `'preprocessed'` argument indicating whether the expression data were already preprocessed, or were provided as raw counts and transformed into log-counts per million.
- `n_perm`: an integer carrying forward the value of the `'n_perm'` argument indicating the number of perturbations performed (NA if asymptotic test was performed).
- `genesets`: carrying forward the value of the `'genesets'` argument defining the gene sets of interest (NULL for gene-wise testing).
- `pval`: computed p-values. A data frame with one row for each gene set, or for each gene if `genesets` argument is NULL, and with 2 columns: the first one `'rawPval'` contains the raw p-values, the second one contains the FDR adjusted p-values (according to the `'padjust_methods'` argument) and is named `'adjPval'`.

References

- Agniel D & Hejblum BP (2017). Variance component score test for time-course gene set analysis of longitudinal RNA-seq data, *Biostatistics*, 18(4):589-604. [10.1093/biostatistics/kxx005](https://doi.org/10.1093/biostatistics/kxx005). [arXiv:1605.02351](https://arxiv.org/abs/1605.02351).
- Law, C. W., Chen, Y., Shi, W., & Smyth, G. K. (2014). voom: Precision weights unlock linear model analysis tools for RNA-seq read counts. *Genome Biology*, 15(2), R29.

See Also

[sp_weights](#) [vc_test_perm](#) [vc_test_asym](#) [p.adjust](#)

Examples

```
nsims <- 2 #100
res_quant <- list()
for(i in 1:2){
  n <- 2000#0
  nr <- 3
  r <- nr*20 #4*nr#100*nr
  t <- matrix(rep(1:nr), r/nr, ncol=1, nrow=r)
  sigma <- 0.4
  b0 <- 1

  #under the null:
  b1 <- 0

  y.tilde <- b0 + b1*t + rnorm(r, sd = sigma)
  y <- t(matrix(rnorm(n*r, sd = sqrt(sigma*abs(y.tilde))), ncol=n, nrow=r) +
    matrix(rep(y.tilde, n), ncol=n, nrow=r))
  x <- matrix(1, ncol=1, nrow=r)

  #run test
  res <- dgsa_seq(exprmat = y, covariates = x, variables2test = t,
    genesets=lapply(0:9, function(x){x*10+(1:10)}),
    cov_variables2test_eff = matrix(1),
```

```

        sample_group = rep(1:(r/nr), each=nr),
        which_test='asymptotic',
        which_weights='none', preprocessed=TRUE)
res_genes <- dgsa_seq(exprmat = y, covariates = x,
                    variables2test = cbind(t,#, rnorm(r)), #t^2
                    genesets = NULL,
                    cov_variables2test_eff = diag(1),
                    sample_group = rep(1:(r/nr), each=nr),
                    which_test = 'asymptotic',
                    which_weights = 'none', preprocessed = TRUE)
length(res_genes$pvals[, 'rawPval'])
quantile(res_genes$pvals[, 'rawPval'])
res_quant[[i]] <- res_genes$pvals[, 'rawPval']
}

#round(rowMeans(vapply(res_quant, FUN = quantile, FUN.VALUE = rep(1.1, 5)), 3)
#plot(density(unlist(res_quant)))
#mean(unlist(res_quant)<0.05)

if(interactive()){
res_genes <- dgsa_seq(exprmat = y, covariates = x, variables2test = t,
                    genesets = NULL,
                    cov_variables2test_eff = matrix(1),
                    sample_group = rep(1:(r/nr), each=nr),
                    which_test = 'permutation',
                    which_weights = 'none', preprocessed = TRUE,
                    n_perm = 1000, parallel_comp = FALSE)

mean(res_genes$pvals$rawPval < 0.05)
summary(res_genes$pvals$adjPval)
}

```

PBT_gmt

PBT gene sets related to kidney transplant

Description

9 Pathogenesis Based Transcripts (PBT) gene sets specifically related to kidney transplant

Usage

```
data(PBT_gmt)
```

Format

a gmt object containing 9 gene sets specific to kidney transplant (see [GSA.read.gmt](#)), which is simply a list with the 3 following components:

- `genesets`: a list of n gene identifiers vectors composing each gene set (each gene set is represented as the vector of the gene identifiers composing it)
- `geneset.names`: a vector of length n containing the gene set names (i.e. gene sets identifiers)
- `geneset.descriptions`: a vector of length n containing gene set descriptions (e.g. textual information on their biological function)

Source

<http://atagc.med.ualberta.ca/Research/GeneLists>

References

Halloran PF, De Freitas DG, Einecke G, *et al.*, The molecular phenotype of kidney transplants: Personal viewpoint, *Am J Transplant*, 10: 2215-2222, 2010. .

Sellares J, Reeve J, Loupy A, *et al.*, Molecular diagnosis of antibody-mediated rejection in human kidney transplants, *Am J Transplant*, 13:971-983, 2013.

Broin PO, Hayde N, Bao Y, *et al.*, A pathogenesis-based transcript signature in donor-specific antibody-positive kidney transplant patients with normal biopsies, *Genomics Data 2*: 357-60, 2014.

Examples

```
data('PBT_gmt')
PBT_gmt
```

 perm_pe

Exact permutation p-values

Description

Calculates exact p-values for permutation tests when permutations are randomly drawn with replacement. This implementation is based on (slightly adapted) the implementation by Belinda Phipson and Gordon Smyth from the R package *statmod*

Usage

```
perm_pe(nperm_supobs, nperm_eff, total_possible_nperm)
```

Arguments

nperm_supobs number of permutations that yielded test statistics at least as extreme as the observed data. Can be a vector or an array of values.

nperm_eff number of permutations effectively computed.

total_possible_nperm total number of permutations possible.

Value

a vector (or an array, similar to nperm_supobs) of exact p-values

Author(s)

Belinda Phipson and Gordon Smyth (adapted by Boris Hejblum)

References

Phipson B, and Smyth GK (2010). Permutation p-values should never be zero: calculating exact p-values when permutations are randomly drawn. *Statistical Applications in Genetics and Molecular Biology*, Volume 9, Issue 1, Article 39. <http://www.statsci.org/smyth/pubs/PermPValuesPreprint.pdf>

See Also

statmod::permp

Examples

```
perm_pe(10, 100, 1000)
```

plot_weights

Plotting mean-variance fit for precision weights estimation

Description

Display the variability with respect to the level of expression and the associated smoothed estimation of precision weights accounting for heteroscedasticity.

Usage

```
plot_weights(x)
```

Arguments

- x a list (such as outputed by the functions `sp_weights` or `voom_weights`) containing the following components:
- `weights`: a matrix $n \times G$ containing the estimated precision weights
 - `plot_utilities`: a list containing the following elements:
 - `reverse_trans`: a function encoding the reverse function used for smoothing the observations before computing the weights
 - `method`: the weight computation method (either "voom" or "locfin")
 - `smth`: the vector of the smoothed values computed
 - `gene_based`: a logical indicating whether the computed weights are based on average at the gene level or on individual observations
 - `mu`: the transformed observed counts or averages
 - `v`: the observed variability estimates

Value

a `ggplot` object

Author(s)

Boris Hejblum

Examples

```

G <- 10000
n <- 12
p <- 2
y <- sapply(1:n, FUN = function(x){rbinom(n = G, size = 0.07, mu = 200)})
x <- sapply(1:p, FUN = function(x){rnorm(n = n, mean = n, sd = 1)})

if(interactive()){
  w <- sp_weights(y, x, use_phi=FALSE, na.rm = TRUE, gene_based=TRUE)
  plot_weights(w)

  vw <- voom_weights(y, x)
  plot_weights(vw)
}

```

sp_weights

*Non parametric local heteroscedasticity weights***Description**

Computes precision weights that account for heteroscedasticity in RNA-seq count data based on non-parametric local linear regression estimates.

Usage

```

sp_weights(
  y,
  x,
  phi = NULL,
  use_phi = TRUE,
  preprocessed = FALSE,
  gene_based = FALSE,
  bw = c("nrd", "ucv", "SJ", "nrd0", "bcv"),
  kernel = c("gaussian", "epanechnikov", "rectangular", "triangular", "biweight",
    "tricube", "cosine", "optcosine"),
  exact = FALSE,
  transform = TRUE,
  verbose = TRUE,
  na.rm = FALSE
)

```

Arguments

y	a numeric matrix of size G x n containing the raw RNA-seq counts or preprocessed expression from n samples for G genes.
x	a numeric matrix of size n x p containing the model covariate(s) from n samples (design matrix).
phi	a numeric design matrix of size n x K containing the K variable(s) of interest (e.g. bases of time).

use_phi	a logical flag indicating whether conditional means should be conditioned on phi and on covariate(s) x, or on x alone. Default is TRUE in which case conditional means are estimated conditionally on both x and phi.
preprocessed	a logical flag indicating whether the expression data have already been preprocessed (e.g. log2 transformed). Default is FALSE, in which case y is assumed to contain raw counts and is normalized into log(counts) per million.
gene_based	a logical flag indicating whether to estimate weights at the gene-level. Default is FALSE, when weights will be estimated at the observation-level.
bw	a character string indicating the smoothing bandwidth selection method to use. See bandwidth for details. Possible values are 'ucv', 'SJ', 'bcv', 'nrd' or 'nrd0'. Default is 'nrd'.
kernel	a character string indicating which kernel should be used. Possibilities are 'gaussian', 'epanechnikov', 'rectangular', 'triangular', 'biweight', 'tricube', 'cosine', 'optcosine'. Default is 'gaussian' (NB: 'tricube' kernel corresponds to the loess method).
exact	a logical flag indicating whether the non-parametric weights accounting for the mean-variance relationship should be computed exactly or extrapolated from the interpolation of local regression of the mean against the variance. Default is FALSE, which uses interpolation (faster).
transform	a logical flag indicating whether values should be transformed to uniform for the purpose of local linear smoothing. This may be helpful if tail observations are sparse and the specified bandwidth gives suboptimal performance there. Default is TRUE.
verbose	a logical flag indicating whether informative messages are printed during the computation. Default is TRUE.
na.rm	logical: should missing values (including NA and NaN) be omitted from the calculations? Default is FALSE.

Value

a list containing the following components:

- **weights**: a matrix $n \times G$ containing the computed precision weights
- **plot_utilities**: a list containing the following elements:
 - **reverse_trans**: a function encoding the reverse function used for smoothing the observations before computing the weights
 - **method**: the weight computation method ("loclin")
 - **smth**: the vector of the smoothed values computed
 - **gene_based**: a logical indicating whether the computed weights are based on average at the gene level or on individual observations
 - **mu**: the transformed observed counts or averages
 - **v**: the observed variability estimates

Author(s)

Boris Hejblum

See Also

[bandwidth density](#)

Examples

```

set.seed(123)

G <- 10000
n <- 12
p <- 2
y <- sapply(1:n, FUN = function(x){rbinom(n = G, size = 0.07, mu = 200)})

x <- sapply(1:p, FUN = function(x){rnorm(n = n, mean = n, sd = 1)})

w <- sp_weights(y, x, use_phi=FALSE, na.rm = TRUE)

```

vc_test_asym

Computes variance component test statistic for longitudinal

Description

This function computes an approximation of the variance component test based on the asymptotic distribution of a mixture of χ^2 s using Davies method from [davies](#)

Usage

```

vc_test_asym(
  y,
  x,
  indiv = rep(1, nrow(x)),
  phi,
  w,
  Sigma_xi = diag(ncol(phi)),
  genewise_pvals = FALSE,
  homogen_traj = FALSE,
  na.rm = FALSE
)

```

Arguments

y	a numeric matrix of dim g x n containing the raw or normalized RNA-seq counts for g genes from n samples.
x	a numeric design matrix of dim n x p containing the p covariates to be adjusted for
indiv	a vector of length n containing the information for attributing each sample to one of the studied individuals. Coerced to be a factor.
phi	a numeric design matrix of size n x K containing the K longitudinal variables to be tested (typically a vector of time points or functions of time)
w	a vector of length n containing the weights for the n samples, corresponding to the inverse of the diagonal of the estimated covariance matrix of y.
Sigma_xi	a matrix of size K x K containing the covariance matrix of the K random effects corresponding to phi.

genewise_pvals	a logical flag indicating whether gene-wise p-values should be returned. Default is FALSE in which case gene set p-value is computed and returned instead.
homogen_traj	a logical flag indicating whether trajectories should be considered homogeneous. Default is FALSE in which case trajectories are not only tested for trend, but also for heterogeneity.
na.rm	logical: should missing values (including NA and NaN) be omitted from the calculations? Default is FALSE.

Value

A list with the following elements when the set p-value is computed:

- set_score_obs: the approximation of the observed set score
- set_pval: the associated set p-value

or a list with the following elements when gene-wise p-values are computed:

- gene_scores_obs: vector of approximating the observed gene-wise scores
- gene_pvals: vector of associated gene-wise p-values

See Also

[davies](#)

Examples

```
set.seed(123)

##generate some fake data
#####
n <- 100
r <- 12
t <- matrix(rep(1:(r/4)), 4, ncol=1, nrow=r)
sigma <- 0.4
b0 <- 1

#under the null:
b1 <- 0
#under the alternative:
#b1 <- 0.5
y.tilde <- b0 + b1*t + rnorm(r, sd = sigma)
y <- t(matrix(rnorm(n*r, sd = sqrt(sigma*abs(y.tilde))), ncol=n, nrow=r) +
      matrix(rep(y.tilde, n), ncol=n, nrow=r))
x <- matrix(1, ncol=1, nrow=r)

#run test
asymTestRes <- vc_test_asym(y, x, phi=cbind(t, t^2),
                          w=matrix(1, ncol=ncol(y), nrow=nrow(y)),
                          Sigma_xi=diag(2), indiv=1:r, genewise_pvals=TRUE)
plot(density(asymTestRes$gene_pvals))
quantile(asymTestRes$gene_pvals)
```

vc_test_perm	<i>Permutation-based variance component test statistic</i>
--------------	--

Description

This function computes an approximation of the Variance Component test for a mixture of χ^2 s using permutations. This is preferable to the asymptotic approximation for small sample sizes. We rely on exact p-values following Phipson and Smyth, 2010 (see References).

Usage

```
vc_test_perm(
  y,
  x,
  indiv = rep(1, nrow(x)),
  phi,
  w,
  Sigma_xi = diag(ncol(phi)),
  n_perm = 1000,
  progressbar = TRUE,
  parallel_comp = TRUE,
  nb_cores = parallel::detectCores() - 1,
  genewise_pvals = FALSE,
  homogen_traj = FALSE,
  na.rm = FALSE
)
```

Arguments

y	a numeric matrix of dim G x n containing the raw RNA-seq counts for G genes from n samples.
x	a numeric design matrix of dim n x p containing the p covariates to be adjusted for.
indiv	a vector of length n containing the information for attributing each sample to one of the studied individuals. Coerced to be a factor.
phi	a numeric design matrix of size n x K containing the K variables to be tested
w	a vector of length n containing the weights for the n samples.
Sigma_xi	a matrix of size K x K containing the covariance matrix of the K random effects.
n_perm	the number of perturbations. Default is 1000.
progressbar	logical indicating whether a progress bar should be displayed when computing permutations (only in interactive mode).
parallel_comp	a logical flag indicating whether parallel computation should be enabled. Only Linux and MacOS are supported, this is ignored on Windows. Default is TRUE.
nb_cores	an integer indicating the number of cores to be used when parallel_comp is TRUE. Only Linux and MacOS are supported, this is ignored on Windows. Default is parallel::detectCores() - 1.
genewise_pvals	a logical flag indicating whether gene-wise p-values should be returned. Default is FALSE in which case gene-set p-value is computed and returned instead.

- homogen_traj a logical flag indicating whether trajectories should be considered homogeneous. Default is FALSE in which case trajectories are not only tested for trend, but also for heterogeneity.
- na.rm logical: should missing values (including NA and NaN) be omitted from the calculations? Default is FALSE.

Value

A list with the following elements when the set p-value is computed:

- set_score_obs: the approximation of the observed set score
- set_pval: the associated set p-value

or a list with the following elements when gene-wise p-values are computed:

- gene_scores_obs: vector of approximating the observed gene-wise scores
- gene_pvals: vector of associated gene-wise p-values
- ds_fdr: vector of associated gene-wise discrete false discovery rates

References

Phipson B, and Smyth GK (2010). Permutation p-values should never be zero: calculating exact p-values when permutations are randomly drawn. *Statistical Applications in Genetics and Molecular Biology*, Volume 9, Issue 1, Article 39. <http://www.statsci.org/smyth/pubs/PermPValuesPreprint.pdf>

See Also

[davies](#)

Examples

```
set.seed(123)

##generate some fake data
#####
n <- 100
r <- 12
t <- matrix(rep(1:3), 4, ncol=1, nrow=r)
sigma <- 0.4
b0 <- 1

#under the null:
b1 <- 0
#under the alternative:
b1 <- 0.5
y.tilde <- b0 + b1*t + rnorm(r, sd = sigma)
y <- t(matrix(rnorm(n*r, sd = sqrt(sigma*abs(y.tilde))), ncol=n, nrow=r) +
  matrix(rep(y.tilde, n), ncol=n, nrow=r))
x <- matrix(1, ncol=1, nrow=r)

#run test
permTestRes <- vc_test_perm(y, x, phi=t,
  w=matrix(1, ncol=ncol(y), nrow=nrow(y)),
  indiv=rep(1:4, each=3), n_perm=50, #1000,
```

```
permTestRes$set_pval      parallel_comp = FALSE)
```

voom_weights	<i>Precision weights accounting for heteroscedasticity in RNA-seq count data</i>
--------------	--

Description

Implementation of the procedure described in Law *et al.* for estimating precision weights from RNA-seq data.

Usage

```
voom_weights(y, x, preprocessed = FALSE, lowess_span = 0.5, R = NULL)
```

Arguments

y	a matrix of size G x n containing the raw RNA-seq counts or preprocessed expressions from n samples for G genes.
x	a matrix of size n x p containing the model covariates from n samples (design matrix).
preprocessed	a logical flag indicating whether the expression data have already been preprocessed (e.g. log2 transformed). Default is FALSE, in which case y is assumed to contain raw counts and is normalized into log(counts) per million.
lowess_span	smoother span for the lowess function, between 0 and 1. This gives the proportion of points in the plot which influence the smooth at each value. Larger values give more smoothness. Default is 0.5.
R	library.size (optional, important to provide if preprocessed = TRUE). Default is NULL

Value

a vector of length n containing the computed precision weights

Author(s)

Boris Hejblum

References

Law, C. W., Chen, Y., Shi, W., & Smyth, G. K. (2014). voom: Precision weights unlock linear model analysis tools for RNA-seq read counts. *Genome Biology*, 15(2), R29.

See Also

[lowess](#) [approxfun](#) [voom](#)

Examples

```
set.seed(123)

G <- 10000
n <- 12
p <- 2

y <- sapply(1:n, FUN=function(x){rbinom(n=G, size=0.07, mu=200)})
x <- sapply(1:p, FUN=function(x){rnorm(n=n, mean=n, sd=1)})

my_w <- voom_weights(y, x)
plot_weights(my_w)
if (requireNamespace('limma', quietly = TRUE)) {
  w_voom <- limma::voom(counts=y, design=x, plot=TRUE)
  #slightly faster, same results
  all.equal(my_w$weights, w_voom$weights)
}

if(interactive()){
#microbenchmark::microbenchmark(limma::voom(counts=t(y), design=x,
#                                           plot=FALSE), voom_weights(x, y),
#                                 times=30)
}
```

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