

Package ‘edgeR’

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Title Empirical analysis of digital gene expression data in R

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biocViews

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Description Differential expression analysis of RNA-seq and digital gene expression profiles with biological replication. Uses empirical Bayes estimation and exact tests based on the negative binomial distribution. Also useful for differential signal analysis with other types of genome-scale count data.

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Description

edgeR is a package for the analysis of digital gene expression data arising from RNA sequencing technologies such as SAGE, CAGE, Tag-seq or RNA-seq, with emphasis on testing for differential expression.

Particular strengths of the package include the ability to estimate biological variation between replicate libraries, and to conduct exact tests of significance which are suitable for small counts. The package is able to make use of even minimal numbers of replicates.

An extensive User's Guide is available, and can be opened by typing `edgeRUsersGuide()` at the R prompt. Detailed help pages are also provided for each individual function.

The *edgeR* package implements original statistical methodology described in the publications below.

Author(s)

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References

Robinson MD and Smyth GK (2007). Moderated statistical tests for assessing differences in tag abundance. *Bioinformatics* 23, 2881-2887

Robinson MD and Smyth GK (2008). Small-sample estimation of negative binomial dispersion, with applications to SAGE data. *Biostatistics*, 9, 321-332

Robinson MD, McCarthy DJ and Smyth GK (2010). *edgeR*: a Bioconductor package for differential expression analysis of digital gene expression data. *Bioinformatics* 26, 139-140

McCarthy, DJ, Chen, Y, Smyth, GK (2012). Differential expression analysis of multifactor RNA-Seq experiments with respect to biological variation. *Nucleic Acids Research* 40, 4288-4297.

Lund, SP, Nettleton, D, McCarthy, DJ, Smyth, GK (2012). Detecting differential expression in RNA-sequence data using quasi-likelihood with shrunken dispersion estimates. *Statistical Applications in Genetics and Molecular Biology*. (Accepted 31 July 2012)

adjustedProfileLik	<i>Adjusted Profile Likelihood for the Negative Binomial Dispersion Parameter</i>
--------------------	---

Description

Compute adjusted profile-likelihoods for estimating the dispersion parameters of genewise negative binomial glms.

Usage

```
adjustedProfileLik(dispersion, y, design, offset, adjust=TRUE)
```

Arguments

dispersion	numeric scalar or vector of dispersions.
y	numeric matrix of counts.
design	numeric matrix giving the design matrix.
offset	numeric matrix of same size as y giving offsets for the log-linear models. Can be a scalar or a vector of length $\text{ncol}(y)$, in which case it is expanded out to a matrix.
adjust	logical, if TRUE then Cox-Reid adjustment is made to the log-likelihood, if FALSE then the log-likelihood is returned without adjustment.

Details

For each row of data, compute the adjusted profile-likelihood for estimating the dispersion parameter of the negative binomial glm. The adjusted profile likelihood is described by McCarthy et al (2012), and is based on the method of Cox and Reid (1987).

The adjusted profile likelihood is an approximate log-likelihood for the dispersion parameter, conditional on the estimated values of the coefficients in the NB log-linear models. The conditional likelihood approach is a technique for adjusting the likelihood function to allow for the fact that nuisance parameters have to be estimated in order to evaluate the likelihood. When estimating the dispersion, the nuisance parameters are the coefficients in the linear model.

This implementation calls the LAPACK library to perform the Cholesky decomposition during adjustment estimation.

Value

vector of adjusted profile log-likelihood values, one for each row of y.

Author(s)

Yunshun Chen, Gordon Smyth, Aaron Lun

References

Cox, DR, and Reid, N (1987). Parameter orthogonality and approximate conditional inference. *Journal of the Royal Statistical Society Series B* 49, 1-39.

McCarthy, DJ, Chen, Y, Smyth, GK (2012). Differential expression analysis of multifactor RNA-Seq experiments with respect to biological variation. *Nucleic Acids Research* 40, 4288-4297. <http://nar.oxfordjournals.org/content/40/10/4288>

See Also

[glmFit](#)

Examples

```
y <- matrix(rnbinom(1000, mu=10, size=2), ncol=4)
design <- matrix(1, 4, 1)
dispersion <- 0.5
apl <- adjustedProfileLik(dispersion, y, design, offset=0)
apl
```

as.data.frame

Turn a TopTags Object into a Dataframe

Description

Turn a TopTags object into a data.frame.

Usage

```
## S3 method for class 'TopTags'
as.data.frame(x, row.names = NULL, optional = FALSE, ...)
```

Arguments

x	an object of class TopTags
row.names	NULL or a character vector giving the row names for the data frame. Missing values are not allowed.
optional	logical. If TRUE, setting row names and converting column names (to syntactic names) is optional.
...	additional arguments to be passed to or from methods.

Details

This method combines all the components of x which have a row for each tag (transcript) into a data.frame.

Value

A data.frame.

Author(s)

Gordon Smyth

See Also

[as.data.frame](#) in the base package.

as.matrix

Turn a DGEList Object into a Matrix

Description

Turn a digital gene expression object into a numeric matrix by extracting the count values.

Usage

```
## S3 method for class 'DGEList'  
as.matrix(x,...)
```

Arguments

x an object of class DGEList.
... additional arguments, not used for these methods.

Details

This method extracts the matrix of counts.

This involves loss of information, so the original data object is not recoverable.

Value

A numeric matrix.

Author(s)

Gordon Smyth

See Also

[as.matrix](#) in the base package or [as.matrix.RGList](#) in the limma package.

bin.dispersion	<i>Estimate Common Dispersion for Negative Binomial GLMs in Bins of Genes Sorted by Overall Abundance</i>
----------------	---

Description

Estimates the common dispersion parameter for each of a number of bins of data for a DGE dataset. Genes are sorted into bins based on overall expression level. For multiple-group (one-way layout) experimental designs, conditional maximum likelihood (CML) methods can be used. For general experimental designs the binned common dispersions we can use Cox-Reid approximate conditional inference, Pearson or deviance estimators for a negative binomial generalized linear model.

Usage

```
binCMLDispersion(y, nbins=50)
binGLMDispersion(y, design, offset=NULL, min.n=100, method="CoxReid", abundance=NULL, ...)
```

Arguments

y	an object that contains the raw counts for each library (the measure of expression level); it can either be a matrix of counts, or a DGEList object with (at least) elements counts (table of unadjusted counts) and samples (data frame containing information about experimental group, library size and normalization factor for the library size)
nbins	scalar, the number of bins for which to compute common dispersions. Default is 50 bins.
design	numeric matrix giving the design matrix for the GLM that is to be fit.
min.n	scalar, the minimum number of genes to be included in each bin.
offset	(optional) numeric scalar, vector or matrix giving the offset (in addition to the log of the effective library size) that is to be included in the NB GLM for the transcripts. If a scalar, then this value will be used as an offset for all transcripts and libraries. If a vector, it should be have length equal to the number of libraries, and the same vector of offsets will be used for each transcript. If a matrix, then each library for each transcript can have a unique offset, if desired. Default is NULL. If NULL, then offset is $\log(\text{lib.size})$ if y is a matrix or $\log(y\$samples\$lib.size * y\$samples\$norm.factors)$ if y is a DGEList object.
method	method used to estimated the dispersion. Argument passed to estimateGLMCommonDisp , which calls the functions to do the computations. Possible values are "CoxReid", "Pearson" or "deviance".
abundance	numeric vector giving abundance of each gene
...	other arguments are passed to lower-level functions.

Details

To obtain estimates of the common dispersion parameters conditional maximum likelihood ([estimateCommonDisp](#)) is used for binCMLDispersion and one of Cox-Reid approximate conditional inference ([dispCoxReid](#)), the deviance ([dispDeviance](#)) or Pearson ([dispPearson](#)) estimates are used for binGLMDispersion. Genes are assigned to bins using the [cutWithMinN](#) function to obtain bins spread over the abundance range of the genes while ensuring that each bin has a minimum number of genes, thus permitting reliable estimation of the common dispersion for each bin.

If there are fewer than `min.n` rows of `y`, then one bin is used. The number of bins is limited to 1000.

Value

Returns a list with two components:

dispersion	numeric vector providing the common dispersion for each bin
abundance	numeric vector providing the average abundance (expression level) of genes in each bin

Author(s)

Gordon Smyth, Davis McCarthy

References

Cox, DR, and Reid, N (1987). Parameter orthogonality and approximate conditional inference. *Journal of the Royal Statistical Society Series B* 49, 1-39.

See Also

[estimateGLMCommonDisp](#), [dispCoxReid](#), [dispPearson](#), [dispDeviance](#)

Examples

```
y <- matrix(rnbinom(1000,mu=10,size=10),ncol=4)
d <- DGEList(counts=y,group=c(1,1,2,2),lib.size=c(1000:1003))
design <- model.matrix(~group, data=d$samples) # Define the design matrix for the full model
bindisp.CML <- binCMLDispersion(d, nbins=50)
bindisp.GLM <- binGLMDispersion(d, design, min.n=10)
```

binomTest

Exact Binomial Tests for Comparing Two Digital Libraries

Description

Computes p-values for differential abundance for each tag between two digital libraries, conditioning on the total count for each tag. The counts in each group as a proportion of the whole are assumed to follow a binomial distribution.

Usage

```
binomTest(y1, y2, n1=sum(y1), n2=sum(y2), p=n1/(n1+n2))
```

Arguments

<code>y1</code>	integer vector giving counts in first library. Non-integer values are rounded to the nearest integer.
<code>y2</code>	integer vector giving counts in second library. Of same length as <code>x</code> . Non-integer values are rounded to the nearest integer.
<code>n1</code>	total number of tags in first library. Non-integer values are rounded to the nearest integer. Not required if <code>p</code> is supplied.

n2	total number of tags in second library. Non-integer values are rounded to the nearest integer. Not required if p is supplied.
p	expected proportion of y1 to the total under the null hypothesis.

Details

This function can be used to compare two libraries from SAGE, RNA-Seq, ChIP-Seq or other sequencing technologies with respect to technical variation.

An exact two-sided binomial test is computed for each tag. This test is closely related to Fisher's exact test for 2x2 contingency tables but, unlike Fisher's test, it conditions on the total number of counts for each tag. The null hypothesis is that the expected counts are in the same proportions as the library sizes, i.e., that the binomial probability for the first library is $n1/(n1+n2)$.

The two-sided rejection region is chosen analogously to Fisher's test. Specifically, the rejection region consists of those values with smallest probabilities under the null hypothesis.

When the counts are reasonably large, the binomial test, Fisher's test and Pearson's chisquare all give the same results. When the counts are smaller, the binomial test is usually to be preferred in this context.

This function replaces the earlier `sage.test` functions in the `statmod` and `sagenhaft` packages. It produces the same results as `binom.test` in the `stats` package, but is much faster.

Value

Numeric vector of p-values.

Author(s)

Gordon Smyth

References

http://en.wikipedia.org/wiki/Binomial_test
http://en.wikipedia.org/wiki/Fisher's_exact_test
http://en.wikipedia.org/wiki/Serial_analysis_of_gene_expression
<http://en.wikipedia.org/wiki/RNA-Seq>

See Also

[sage.test](#) (statmod package), [binom.test](#) (stats package)

Examples

```
binomTest(c(0,5,10),c(0,30,50),n1=10000,n2=15000)
# Univariate equivalents:
binom.test(5,5+30,p=10000/(10000+15000))$p.value
binom.test(10,10+50,p=10000/(10000+15000))$p.value
```

 calcNormFactors

Calculate Normalization Factors to Align Columns of a Count Matrix

Description

Calculate normalization factors to scale the raw library sizes.

Usage

```
calcNormFactors(object, method=c("TMM","RLE","upperquartile"), refColumn = NULL,
  logratioTrim = .3, sumTrim = 0.05, doWeighting=TRUE, Acutoff=-1e10, p=0.75)
```

Arguments

object	either a matrix of raw (read) counts or a DGEList object
method	method to use to calculate the scale factors
refColumn	column to use as reference for method="TMM"
logratioTrim	amount of trim to use on log-ratios ("M" values) for method="TMM"
sumTrim	amount of trim to use on the combined absolute levels ("A" values) for method="TMM"
doWeighting	logical, whether to compute (asymptotic binomial precision) weights for method="TMM"
Acutoff	cutoff on "A" values to use before trimming for method="TMM"
p	percentile (between 0 and 1) of the counts that is aligned when method="upperquartile"

Details

method="TMM" is the weighted trimmed mean of M-values (to the reference) proposed by Robinson and Oshlack (2010), where the weights are from the delta method on Binomial data. If refColumn is unspecified, the library whose upper quartile is closest to the mean upper quartile is used.

method="RLE" is the scaling factor method proposed by Anders and Huber (2010). We call it "relative log expression", as median library is calculated from the geometric mean of all columns and the median ratio of each sample to the median library is taken as the scale factor.

method="upperquartile" is the upper-quartile normalization method of Bullard et al (2010), in which the scale factors are calculated from the 75% quantile of the counts for each library, after removing transcripts which are zero in all libraries. This idea is generalized here to allow scaling by any quantile of the distributions.

For symmetry, normalization factors are adjusted to multiply to 1. The effective library size is then the original library size multiplied by the scaling factor.

Value

If a matrix is given for object, the output is a vector with length ncol(object) giving the relative normalization factors. If a DGEList object is given for object, the output is a DGEList object containing the normalization factors in the samples\$norm.factors element.

Author(s)

Mark Robinson, Gordon Smyth

References

Anders, S, Huber, W (2010). Differential expression analysis for sequence count data *Genome Biology* 11, R106.

Bullard JH, Purdom E, Hansen KD, Dudoit S. (2010) Evaluation of statistical methods for normalization and differential expression in mRNA-Seq experiments. *BMC Bioinformatics* 11, 94. A scaling normalization method for differential expression analysis of RNA-seq data.

Robinson MD, Oshlack A (2010). *Genome Biology* 11, R25.

Examples

```
y <- matrix( rpois(1000, lambda=5), nrow=200 )
calcNormFactors(y)
```

```
commonCondLogLikDerDelta
```

Conditional Log-Likelihoods in Terms of Delta

Description

Common conditional log-likelihood parameterized in terms of delta ($\phi / (\phi + 1)$)

Usage

```
commonCondLogLikDerDelta(y, delta, der = 0)
```

Arguments

y	list with elements comprising the matrices of count data (or pseudocounts) for the different groups
delta	delta ($\phi / (\phi + 1)$) parameter of negative binomial
der	derivative, either 0 (the function), 1 (first derivative) or 2 (second derivative)

Details

The common conditional log-likelihood is constructed by summing over all of the individual tag conditional log-likelihoods. The common conditional log-likelihood is taken as a function of the dispersion parameter (ϕ), and here parameterized in terms of delta ($\phi / (\phi + 1)$). The value of delta that maximizes the common conditional log-likelihood is converted back to the ϕ scale, and this value is the estimate of the common dispersion parameter used by all tags.

Value

numeric scalar of function/derivative evaluated at given delta

Author(s)

Davis McCarthy

See Also

[estimateCommonDisp](#) is the user-level function for estimating the common dispersion parameter.

Examples

```
counts<-matrix(rnbinom(20,size=1,mu=10),nrow=5)
d<-DGEList(counts=counts,group=rep(1:2,each=2),lib.size=rep(c(1000:1001),2))
y<-splitIntoGroups(d)
ll1<-commonCondLogLikDerDelta(y,delta=0.5,der=0)
ll2<-commonCondLogLikDerDelta(y,delta=0.5,der=1)
```

condLogLikDerSize	<i>Conditional Log-Likelihood of the Dispersion for a Single Group of Replicate Libraries</i>
-------------------	---

Description

Derivatives of the negative-binomial log-likelihood with respect to the dispersion parameter for each tag/transcript, conditional on the mean count, for a single group of replicate libraries of the same size.

Usage

```
condLogLikDerSize(y, r, der=1L)
condLogLikDerDelta(y, delta, der=1L)
```

Arguments

y	matrix of counts, all counts in each row having the same population mean
r	numeric vector or scalar, size parameter of negative binomial distribution, equal to 1/dispersion
delta	numeric vector or scalar, delta parameter of negative binomial, equal to dispersion/(1+dispersion)
der	integer specifying derivative required, either 0 (the function), 1 (first derivative) or 2 (second derivative)

Details

The library sizes must be equalized before running this function. This function carries out the actual mathematical computations for the conditional log-likelihood and its derivatives, calculating the conditional log-likelihood for each tag/transcript. Derivatives are with respect to either the size or the delta parametrization of the dispersion.

Value

vector of function/derivative evaluations, one for each transcript, with respect to

Author(s)

Mark Robinson, Davis McCarthy, Gordon Smyth

Examples

```
y <- matrix(rnbinom(10,size=1,mu=10),nrow=5)
condLogLikDerSize(y,r=1,der=1)
condLogLikDerDelta(y,delta=0.5,der=1)
```

cpm

*Calculate Counts per Million from DGEList or Matrix Object***Description**

Returns counts per million from a DGEList or matrix object by dividing raw counts by library size (which can be normalized) and multiplying by one million.

Usage

```
cpm(x, normalized.lib.sizes=FALSE)
```

Arguments

x either a matrix of counts or a DGEList object with (at least) elements counts (table of unadjusted counts) and samples (data frame containing information about experimental group, library size and normalization factor for the library size)

normalized.lib.sizes logical, should the library sizes (total sum of counts for each library) be normalized using the norm.factors component of the DGEList object? Ignored (with a warning) if x is a count matrix.

Details

A convenience function to compute the counts per million for plotting and comparing libraries on a convenient scale. Essentially just does the calculation $1e06 * t(x) / \text{lib.size}$ to produce counts per million, where x is a matrix of counts and the lib.size can be the total sum of counts in each library or a normalized version of this using TMM normalization or equivalent method.

Value

getPriorN returns a numeric scalar

Author(s)

Davis McCarthy, Gordon Smyth

See Also

[DGEList](#) for more information about the DGEList class.

Examples

```
# generate raw counts from NB, create list object
y<-matrix(rnbinom(20,size=1,mu=10),nrow=5)
cpm(y)
d<-DGEList(counts=y,group=rep(1:2,each=2),lib.size=rep(c(1000:1001),2))
# When applied to a DGEList object, x$samples$lib.size is used
cpm(d)
# As x$samples$lib.size here is very different from colSums(y), cpm(y) and cpm(d) give very different results for the count
```

`cutWithMinN`*Cut numeric vector into non-empty intervals*

Description

Discretizes a numeric vector. Divides the range of `x` into intervals, so that each interval contains a minimum number of values, and codes the values in `x` according to which interval they fall.

Usage

```
cutWithMinN(x, intervals=2, min.n=1)
```

Arguments

<code>x</code>	numeric vector.
<code>intervals</code>	number of intervals (greater than or equal to 2).
<code>min.n</code>	minimum number of values in any interval.

Details

This function strikes a compromise between the base functions `cut`, which by default cuts a vector into equal length intervals, and `quantile`, which is suited to finding equally populated intervals.

Value

A list with components:

<code>group</code>	integer vector of same length as <code>x</code> indicating which interval each value belongs to.
<code>breaks</code>	numeric vector of length <code>intervals+1</code> giving the left and right limits of each interval.

Author(s)

Gordon Smyth

See Also

[cut](#), [quantile](#).

Examples

```
x <- c(1,2,3,4,5,6,7,100)
cutWithMinN(x,3,min.n=1)
```

decideTestsDGE *Multiple Testing Across Genes and Contrasts*

Description

Classify a series of related differential expression statistics as up, down or not significant. A number of different multiple testing schemes are offered which adjust for multiple testing down the genes as well as across contrasts for each gene.

Usage

```
decideTestsDGE(object, adjust.method="BH", p.value=0.05)
```

Arguments

object	deDGElist object, output from <code>exactTest</code> , or DGELRT object, output from DGELRT, from which p-values for differential expression and log-fold change values may be extracted.
adjust.method	character string specifying p-value adjustment method. Possible values are "none", "BH", "fdr" (equivalent to "BH"), "BY" and "holm". See p.adjust for details.
p.value	numeric value between 0 and 1 giving the desired size of the test

Details

These functions implement multiple testing procedures for determining whether each log-fold change in a matrix of log-fold changes should be considered significantly different from zero.

Value

An object of class `TestResults` (see [TestResults](#)). This is essentially a numeric matrix with elements -1, 0 or 1 depending on whether each DE p-value is classified as significant with negative log-fold change, not significant or significant with positive log-fold change, respectively.

Author(s)

Davis McCarthy, Gordon Smyth

See Also

Adapted from [decideTests](#) in the `limma` package.

DGEExact-class	<i>differential expression of Digital Gene Expression data - class</i>
----------------	--

Description

A simple list-based class for storing results of differential expression analysis for DGE data

Slots/List Components

Objects of this class contain the following list components:

table: data frame containing the log-concentration (i.e. expression level), the log-fold change in expression between the two groups/conditions and the exact p-value for differential expression, for each tag.

comparison: vector giving the two experimental groups/conditions being compared.

genes: a data frame containing information about each transcript (can be NULL).

Methods

This class inherits directly from class list so any operation appropriate for lists will work on objects of this class. DGEExact objects also have a show method.

Author(s)

Mark Robinson, Davis McCarthy

DGEGLM-class	<i>Digital Gene Expression Generalized Linear Model results - class</i>
--------------	---

Description

A simple list-based class for storing results of a GLM fit to each tag/gene in a DGE dataset.

Slots/List Components

Objects of this class contain the following list components:

coefficients: matrix containing the coefficients computed from fitting the model defined by the design matrix to each gene/tag in the dataset.

df.residual: vector containing the residual degrees of freedom for the model fit to each tag/gene in the dataset.

deviance: vector giving the deviance from the model fit to each tag/gene.

design: design matrix for the full model from the likelihood ratio test.

offset: scalar, vector or matrix of offset values to be included in the GLMs for each tag/gene.

samples: data frame containing information about the samples comprising the dataset.

genes: data frame containing information about the genes or tags for which we have DGE data (can be NULL if there is no information available).

dispersion: scalar or vector providing the value of the dispersion parameter used in the negative binomial GLM for each tag/gene.

lib.size: vector providing the effective library size for each sample in the dataset.

weights: matrix of weights used in the GLM fitting for each tag/gene.

fitted.values: the fitted (expected) values—here they are counts—from the GLM for each tag/gene.

abundance: vector of gene/tag abundances (expression level), on the log2 scale, computed from the mean count for each gene/tag after scaling count by normalized library size.

Methods

This class inherits directly from class list so any operation appropriate for lists will work on objects of this class. DGEGLM objects also have a show method.

Author(s)

Davis McCarthy

DGEList

DGEList Constructor

Description

Creates a DGEList object from a table of counts (rows=features, columns=samples), group indicator for each column, library size (optional) and a table of feature annotation (optional).

Usage

```
DGEList(counts = matrix(0, 0, 0), lib.size = NULL, norm.factors = NULL,
        group = rep.int(1,ncol(counts)), genes = NULL, remove.zeros = FALSE)
```

Arguments

counts	numeric matrix containing the read counts.
lib.size	numeric vector containing the total to normalize against for each sample (optional)
norm.factors	numeric vector containing normalization factors (optional, defaults to all 1)
group	vector giving the experimental group/condition for each sample/library
genes	data frame containing annotation information for the tags/transcripts/genes for which we have count data (optional).
remove.zeros	logical, whether to remove rows that have 0 total count; default is FALSE so as to retain all information in the dataset

Details

If no lib.size argument is passed to the constructor, the column totals are used.

The optional genes argument supplies a data.frame of annotation for each row or feature.

Value

a DGEList object

Author(s)

Mark Robinson, Davis McCarthy, Gordon Smyth

See Also

[DGEList-class](#)

Examples

```
y <- matrix(rnbinom(10000,mu=5,size=2),ncol=4)
d <- DGEList(counts=y, group=rep(1:2,each=2))
d$samples
```

DGEList-class

Digital Gene Expression data - class

Description

A simple list-based class for storing read counts from digital gene expression technologies and other important information for the analysis of DGE data.

Slots/List Components

Objects of this class contain (at least) the following list components:

counts: numeric matrix containing the read counts.

samples: data.frame containing the library size and group labels.

Methods

This class inherits directly from class list so any operation appropriate for lists will work on objects of this class. DGEList objects also have a show method.

Author(s)

Mark Robinson

See Also

[DGEList](#)

DGELRT-class	<i>Digital Gene Expression Likelihood Ratio Test data and results - class</i>
--------------	---

Description

A simple list-based class for storing results of a GLM-based differential expression analysis for DGE data, with evidence for differential expression assessed using a likelihood ratio test.

Slots/List Components

Objects of this class contain the following list components:

table: data frame containing the log-concentration (i.e. expression level), the log-fold change in expression between the two groups/conditions and the exact p-value for differential expression, for each tag.

coefficients.full: matrix containing the coefficients computed from fitting the full model (fit using glmFit and a given design matrix) to each gene/tag in the dataset.

coefficients.null: matrix containing the coefficients computed from fitting the null model to each gene/tag in the dataset. The null model is the model to which the full model is compared, and is fit using glmFit and dropping selected column(s) (i.e. coefficient(s)) from the design matrix for the full model.

design: design matrix for the full model from the likelihood ratio test.

...: if the argument y to glmLRT (which produces the DGELRT object) was itself a DGEList object, then the DGELRT will contain all of the elements of y, except for the table of counts and the table of pseudocounts.

Methods

This class inherits directly from class list so any operation appropriate for lists will work on objects of this class. DGELRT objects also have a show method.

Author(s)

Davis McCarthy

dglmStdResid	<i>Visualize the mean-variance relationship in DGE data using standardized residuals</i>
--------------	--

Description

Appropriate modelling of the mean-variance relationship in DGE data is important for making inferences about differential expression. However, the standard approach to visualizing the mean-variance relationship is not appropriate for general, complicated experimental designs that require generalized linear models (GLMs) for analysis. Here are functions to compute standardized residuals from a Poisson GLM and plot them for bins based on overall expression level of tags as a way to visualize the mean-variance relationship. A rough estimate of the dispersion parameter can also be obtained from the standardized residuals.

Usage

```
dglmStdResid(y, design, dispersion=0, offset=0, nbins=100, make.plot=TRUE,
             xlab="Mean", ylab="Ave. binned standardized residual", ...)
getDispersions(binned.object)
```

Arguments

<code>y</code>	numeric matrix of counts, each row represents one tag, each column represents one DGE library.
<code>design</code>	numeric matrix giving the design matrix of the GLM. Assumed to be full column rank.
<code>dispersion</code>	numeric scalar or vector giving the dispersion parameter for each GLM. Can be a scalar giving one value for all tags, or a vector of length equal to the number of tags giving tag-wise dispersions.
<code>offset</code>	numeric vector or matrix giving the offset that is to be included in the log-linear model predictor. Can be a vector of length equal to the number of libraries, or a matrix of the same size as <code>y</code> .
<code>nbins</code>	scalar giving the number of bins (formed by using the quantiles of the genewise mean expression levels) for which to compute average means and variances for exploring the mean-variance relationship. Default is 100 bins
<code>make.plot</code>	logical, whether or not to plot the mean standardized residual for binned data (binned on expression level). Provides a visualization of the mean-variance relationship. Default is TRUE.
<code>xlab</code>	character string giving the label for the x-axis. Standard graphical parameter. If left as the default, then the x-axis label will be set to "Mean".
<code>ylab</code>	character string giving the label for the y-axis. Standard graphical parameter. If left as the default, then the y-axis label will be set to "Ave. binned standardized residual".
<code>...</code>	further arguments passed on to plot
<code>binned.object</code>	list object, which is the output of <code>dglmStdResid</code> .

Details

This function is useful for exploring the mean-variance relationship in the data. Raw or pooled variances cannot be used for complex experimental designs, so instead we can fit a Poisson model using the appropriate design matrix to each tag and use the standardized residuals in place of the pooled variance (as in `plotMeanVar`) to visualize the mean-variance relationship in the data. The function will plot the average standardized residual for observations split into `nbins` bins by overall expression level. This provides a useful summary of how the variance of the counts change with respect to average expression level (abundance). A line showing the Poisson mean-variance relationship (mean equals variance) is always shown to illustrate how the genewise variances may differ from a Poisson mean-variance relationship. A log-log scale is used for the plot.

The function `mgglmLS` is used to fit the Poisson models to the data. This code is fast for fitting models, but does not compute the value for the leverage, technically required to compute the standardized residuals. Here, we approximate the standardized residuals by replacing the usual denominator of $(1 - \text{leverage})$ by $(1 - p/n)$, where n is the number of observations per tag (i.e. number of libraries) and p is the number of parameters in the model (i.e. number of columns in the full-rank design matrix).

Value

dglmStdResid produces a mean-variance plot based on standardized residuals from a Poisson model fit for each tag for the DGE data. dglmStdResid returns a list with the following elements:

ave.means	vector of the average expression level within each bin of observations
ave.std.resid	vector of the average standardized Poisson residual within each bin of tags
bin.means	list containing the average (mean) expression level (given by the fitted value from the given Poisson model) for observations divided into bins based on amount of expression
bin.std.resid	list containing the standardized residual from the given Poisson model for observations divided into bins based on amount of expression
means	vector giving the fitted value for each observed count
standardized.residuals	vector giving approximate standardized residual for each observed count
bins	list containing the indices for the observations, assigning them to bins
nbins	scalar giving the number of bins used to split up the observed counts
ngenes	scalar giving the number of genes/tags in the dataset
nlibs	scalar giving the number of libraries in the dataset

getDispersions computes the dispersion from the standardized residuals and returns a list with the following components:

bin.dispersion	vector giving the estimated dispersion value for each bin of observed counts, computed using the average standardized residual for the bin
bin.dispersion.used	vector giving the actual estimated dispersion value to be used. Some computed dispersions using the method in this function can be negative, which is not allowed. We use the dispersion value from the nearest bin of higher expression level with positive dispersion value in place of any negative dispersions.
dispersion	vector giving the estimated dispersion for each observation, using the binned dispersion estimates from above, so that all of the observations in a given bin get the same dispersion value.

Author(s)

Davis McCarthy

See Also

[plotMeanVar](#), [plotMDS.DGEList](#), [plotSmear](#) and [maPlot](#) provide more ways of visualizing DGE data.

Examples

```
y <- matrix(rnbinom(1000,mu=10,size=2),ncol=4)
design <- model.matrix(~c(0,0,1,1)+c(0,1,0,1))
binned <- dglmStdResid(y, design, dispersion=0.5)
```

```
getDispersions(binned)$bin.dispersion.used # Look at the estimated dispersions for the bins
```

dim	<i>Retrieve the Dimensions of a DGEList, DGEEexact, DGEGLM, DGELRT or TopTags Object</i>
-----	--

Description

Retrieve the number of rows (transcripts) and columns (libraries) for an DGEList, DGEEexact or TopTags Object.

Usage

```
## S3 method for class 'DGEList'  
dim(x)  
## S3 method for class 'DGEList'  
length(x)
```

Arguments

x an object of class DGEList, DGEEexact, TopTags, DGEGLM or DGELRT

Details

Digital gene expression data objects share many analogies with ordinary matrices in which the rows correspond to transcripts or genes and the columns to arrays. These methods allow one to extract the size of microarray data objects in the same way that one would do for ordinary matrices.

A consequence is that row and column commands `nrow(x)`, `ncol(x)` and so on also work.

Value

Numeric vector of length 2. The first element is the number of rows (genes) and the second is the number of columns (arrays).

Author(s)

Gordon Smyth, Davis McCarthy

See Also

[dim](#) in the base package.

[02.Classes](#) gives an overview of data classes used in LIMMA.

Examples

```
M <- A <- matrix(11:14,4,2)  
rownames(M) <- rownames(A) <- c("a","b","c","d")  
colnames(M) <- colnames(A) <- c("A1","A2")  
MA <- new("MAList",list(M=M,A=A))  
dim(M)  
ncol(M)  
nrow(M)  
length(M)
```

dimnames	<i>Retrieve the Dimension Names of a DGEList Object</i>
----------	---

Description

Retrieve the dimension names of a digital gene expression data object.

Usage

```
## S3 method for class 'DGEList'  
dimnames(x)  
## S3 replacement method for class 'DGEList'  
dimnames(x) <- value
```

Arguments

x	an object of class DGEList, DGEEexact, DGEGLM or TopTags
value	a possible value for dimnames(x): see dimnames

Details

The dimension names of a microarray object are the same as those of the most important matrix component of that object.

A consequence is that rownames and colnames will work as expected.

Value

Either NULL or a list of length 2. If a list, its components are either NULL or a character vector the length of the appropriate dimension of x.

Author(s)

Gordon Smyth

See Also

[dimnames](#) in the base package.

[02.Classes](#) gives an overview of data classes used in LIMMA.

dispBinTrend	<i>Estimate Dispersions with an Abundance-Dependent Trend for Negative Binomial GLMs</i>
--------------	--

Description

Estimate a dispersion parameter for each of many negative binomial generalized linear models by computing the common dispersion for genes sorted into bins based on overall abundance and then using splines or a loess fit to interpolate a dispersion value for each gene, dependent on overall abundance of the gene.

Usage

```
dispBinTrend(y, design=NULL, offset=NULL, df = 5, span=2/3, min.n=400, method.bin="CoxReid",
             method.trend="spline", trace=0, abundance=NULL, ...)
```

Arguments

y	numeric matrix of counts
design	numeric matrix giving the design matrix for the GLM that is to be fit.
offset	numeric scalar, vector or matrix giving the offset (in addition to the log of the effective library size) that is to be included in the NB GLM for the transcripts. If a scalar, then this value will be used as an offset for all transcripts and libraries. If a vector, it should be have length equal to the number of libraries, and the same vector of offsets will be used for each transcript. If a matrix, then each library for each transcript can have a unique offset, if desired. In <code>adjustedProfileLik</code> the offset must be a matrix with the same dimension as the table of counts.
df	scalar integer, the degrees of freedom for the natural cubic splines fit, used to determine the placement of the knots (number of knots is $df - 1$).
span	scalar between 0 and 1, passed to <code>loess</code> to determine the amount of smoothing for the loess fit.
min.n	scalar integer, minimum number of genes in each of the bins.
method.bin	character, passed to <code>binGLMDispersion</code> , to specify the method used to compute the common dispersion within each bin of genes. Default is "CoxReid", other options are "Pearson" and "deviance".
method.trend	character, specifies method to produce a smooth fit through the binned common dispersions in order to interpolate the trended dispersions. Default is "spline" to use natural cubic splines, other option is "loess" to use a loess fit.
trace	logical, should iteration information be output?
abundance	numeric vector giving abundance of each gene
...	option arguments to be passed to lower-level function <code>binGLMDispersion</code> .

Details

This function takes the binned common dispersion and abundance from `binGLMDispersion` and fits a smooth curve through these binned values using either natural cubic splines or loess. From this smooth curve it predicts the dispersion value for each gene based on the gene's overall abundance. This results in estimates for the NB dispersion parameter which have a dependence on the overall expression level of the gene, and thus have an abundance-dependent trend. This function is called by `estimateGLMTrendedDisp`.

Value

list with the following components:

abundance	numeric vector containing the overall abundance for each gene
dispersion	numeric vector giving the trended dispersion estimate for each gene
bin.abundance	numeric vector of length equal to nbins giving the average (mean) abundance for each bin
bin.dispersion	numeric vector of length equal to nbins giving the estimated common dispersion for each bin

Author(s)

Davis McCarthy and Gordon Smyth

References

Cox, DR, and Reid, N (1987). Parameter orthogonality and approximate conditional inference. *Journal of the Royal Statistical Society Series B* 49, 1-39.

See Also

[binGLMDispersion](#), [estimateGLMTrendedDisp](#)

Examples

```

ntags <- 1000
nlibs <- 4
means <- seq(5,10000,length.out=ntags)
y <- matrix(rnbinom(ntags*nlibs,mu=rep(means,nlibs),size=0.1*means),nrow=ntags,ncol=nlibs)
keep <- rowSums(y) > 0
y <- y[keep,]
group <- factor(c(1,1,2,2))
lib.size <- colSums(y)
design <- model.matrix(~group) # Define the design matrix for the full model
disp <- dispBinTrend(y, design, offset=log(lib.size), min.n=100, span=0.3)
plot(disp$abundance, disp$dispersion)

```

dispCoxReid

Estimate Common Dispersion for Negative Binomial GLMs

Description

Estimate a common dispersion parameter across multiple negative binomial generalized linear models.

Usage

```

dispCoxReid(y, design, offset=NULL, interval=c(0,4), tol=1e-5, min.row.sum=5, subset=10000)
dispDeviance(y, design, offset=NULL, interval=c(0,4), tol=1e-5, min.row.sum=5, subset=10000,
             robust=FALSE, trace=FALSE)
dispPearson(y, design, offset=NULL, interval=c(0,4), tol=1e-5, min.row.sum=5, subset=10000,
            robust=FALSE, trace=FALSE)

```

Arguments

y	numeric matrix of counts. A glm is fitted to each row.
design	numeric design matrix, as for glmFit .
offset	numeric vector or matrix of offsets for the log-linear models, as for glmFit .
interval	numeric vector of length 2 giving allowable values for the dispersion, passed to optimize.
tol	the desired accuracy, see optimize or uniroot.
min.row.sum	integer. Only rows with at least this number of counts are used.
subset	integer, number of rows to use in the calculation. Rows used are chosen evenly spaced by abundance.
trace	logical, should iteration information be output?
robust	logical, should a robust estimator be used?

Details

These are low-level (non-object-orientated) functions called by estimateGLMCommonDisp.

dispCoxReid maximizes the Cox-Reid adjusted profile likelihood (Cox and Reid, 1987). dispPearson sets the average Pearson goodness of fit statistics to its (asymptotic) expected value. This is also known as the *pseudo-likelihood* estimator. dispDeviance sets the average residual deviance statistic to its (asymptotic) expected values. This is also known as the *quasi-likelihood* estimator.

Robinson and Smyth (2008) and McCarthy et al (2011) showed that the Pearson (pseudo-likelihood) estimator typically under-estimates the true dispersion. It can be seriously biased when the number of libraries (ncol(y) is small. On the other hand, the deviance (quasi-likelihood) estimator typically over-estimates the true dispersion when the number of libraries is small. Robinson and Smyth (2008) and McCarthy et al (2011) showed the Cox-Reid estimator to be the least biased of the three options.

dispCoxReid uses optimize to maximize the adjusted profile likelihood, while dispDeviance and dispPearson use uniroot to solve the estimating equation. The robust options use an order statistic instead the mean statistic, and have the effect that a minority of tags with very large (outlier) dispersions should have limited influence on the estimated value.

Value

Numeric vector of length one giving the estimated common dispersion.

Author(s)

Gordon Smyth

References

Cox, DR, and Reid, N (1987). Parameter orthogonality and approximate conditional inference. *Journal of the Royal Statistical Society Series B* 49, 1-39.

Robinson MD and Smyth GK (2008). Small-sample estimation of negative binomial dispersion, with applications to SAGE data. *Biostatistics*, 9, 321-332

McCarthy, DJ, Chen, Y, Smyth, GK (2012). Differential expression analysis of multifactor RNA-Seq experiments with respect to biological variation. *Nucleic Acids Research*. <http://nar.oxfordjournals.org/content/early/2012/02/06/nar.gks042> (Published online 28 January 2012)

See Also

[estimateGLMCommonDisp](#), [optimize](#), [uniroot](#)

Examples

```
ntags <- 100
nlibs <- 4
y <- matrix(rnbinom(ntags*nlibs,mu=10,size=10),nrow=ntags,ncol=nlibs)
group <- factor(c(1,1,2,2))
lib.size <- rowSums(y)
design <- model.matrix(~group) # Define the design matrix for the full model
disp <- dispCoxReid(y, design, offset=log(lib.size), subset=100)
```

dispCoxReidInterpolateTagwise

Estimate Tagwise Dispersion for Negative Binomial GLMs by Cox-Reid Adjusted Profile Likelihood

Description

Estimate tagwise dispersion parameters across multiple negative binomial generalized linear models using weighted Cox-Reid Adjusted Profile-likelihood and cubic spline interpolation over a tagwise grid.

Usage

```
dispCoxReidInterpolateTagwise(y, design, offset=NULL, dispersion, trend=TRUE, abundance=NULL,
  min.row.sum=5, prior.df=20, span=0.3, grid.npts=11, grid.range=c(-6,6))
```

Arguments

y	numeric matrix of counts
design	numeric matrix giving the design matrix for the GLM that is to be fit.
offset	numeric scalar, vector or matrix giving the offset (in addition to the log of the effective library size) that is to be included in the NB GLM for the transcripts. If a scalar, then this value will be used as an offset for all transcripts and libraries. If a vector, it should be have length equal to the number of libraries, and the same vector of offsets will be used for each transcript. If a matrix, then each library for each transcript can have a unique offset, if desired. In <code>adjustedProfileLik</code> the offset must be a matrix with the same dimension as the table of counts.
dispersion	numeric scalar or vector giving the dispersion(s) towards which the tagwise dispersion parameters are shrunk.
trend	logical, whether abundance-dispersion trend is used for smoothing.
abundance	numeric scalar or vector giving the tagwise log-abundance measure for each tag. If null, the abundance is then evaluated by <code>mglmOneGroup</code>
min.row.sum	numeric scalar giving a value for the filtering out of low abundance tags. Only tags with total sum of counts above this value are used. Low abundance tags can adversely affect the estimation of the common dispersion, so this argument allows the user to select an appropriate filter threshold for the tag abundance.

prior.df	numeric scalar, prior degsmoothing parameter that indicates the weight to give to the common likelihood compared to the individual tag's likelihood; default <code>getPriorN(object)</code> gives a value for <code>prior.n</code> that is equivalent to giving the common likelihood 20 prior degrees of freedom in the estimation of the tag/genewise dispersion.
span	numeric parameter between 0 and 1 specifying proportion of data to be used in the local regression moving window. Larger numbers give smoother fits.
grid.npts	numeric scalar, the number of points at which to place knots for the spline-based estimation of the tagwise dispersion estimates.
grid.range	numeric vector of length 2, giving relative range, in terms of $\log_2(\text{dispersion})$, on either side of trendline for each tag for spline grid points.

Details

In the edgeR context, `dispCoxReidInterpolateTagwise` is a low-level function called by `estimateGLMTagwiseDisp`. `dispCoxReidInterpolateTagwise` calls the function `maximizeInterpolant` to fit cubic spline interpolation over a tagwise grid.

Value

`dispCoxReidInterpolateTagwise` produces a vector of tagwise dispersions having the same length as the number of genes in the count data.

Author(s)

Yunshun Chen, Gordon Smyth

References

Cox, DR, and Reid, N (1987). Parameter orthogonality and approximate conditional inference. *Journal of the Royal Statistical Society Series B* 49, 1-39.

McCarthy, DJ, Chen, Y, Smyth, GK (2012). Differential expression analysis of multifactor RNA-Seq experiments with respect to biological variation. *Nucleic Acids Research* 40, 4288-4297. <http://nar.oxfordjournals.org/content/40/10/4288>

See Also

[estimateGLMTagwiseDisp](#), [maximizeInterpolant](#)

Examples

```
y <- matrix(rnbinom(1000, mu=10, size=2), ncol=4)
design <- matrix(1, 4, 1)
dispersion <- 0.5
d <- dispCoxReidInterpolateTagwise(y, design, dispersion=dispersion)
d
```

 dispCoxReidSplineTrend *Estimate Dispersion Trend for Negative Binomial GLMs*

Description

Estimate trended dispersion parameters across multiple negative binomial generalized linear models using Cox-Reid adjusted profile likelihood.

Usage

```
dispCoxReidSplineTrend(y, design, offset=NULL, df = 5, subset=10000, method.optim="Nelder-Mead", trace=
dispCoxReidPowerTrend(y, design, offset=NULL, subset=10000, method.optim="Nelder-Mead", trace=0)
```

Arguments

y	numeric matrix of counts
design	numeric matrix giving the design matrix for the GLM that is to be fit.
offset	numeric scalar, vector or matrix giving the offset (in addition to the log of the effective library size) that is to be included in the NB GLM for the transcripts. If a scalar, then this value will be used as an offset for all transcripts and libraries. If a vector, it should be have length equal to the number of libraries, and the same vector of offsets will be used for each transcript. If a matrix, then each library for each transcript can have a unique offset, if desired. In <code>adjustedProfileLik</code> the offset must be a matrix with the same dimension as the table of counts.
df	integer giving the degrees of freedom of the spline function, see <code>ns</code> in the <code>splines</code> package.
subset	integer, number of rows to use in the calculation. Rows used are chosen evenly spaced by abundance using <code>cutWithMinN</code> .
method.optim	the method to be used in <code>optim</code> . See <code>optim</code> for more detail.
trace	logical, should iteration information be output?

Details

In the `edgeR` context, these are low-level functions called by `estimateGLMTrendedDisp`.

`dispCoxReidSplineTrend` and `dispCoxReidPowerTrend` fit abundance trends to the tagwise dispersions. `dispCoxReidSplineTrend` fits a regression spline whereas `dispCoxReidPowerTrend` fits a log-linear trend of the form $a \cdot \exp(\text{abundance})^b + c$. In either case, `optim` is used to maximize the adjusted profile likelihood (Cox and Reid, 1987).

Value

List containing numeric vectors dispersion and abundance containing the estimated dispersion and abundance for each transcript. The vectors are of the same length as `nrow(y)`.

Author(s)

Yunshun Chen, Davis McCarthy, Gordon Smyth

References

Cox, DR, and Reid, N (1987). Parameter orthogonality and approximate conditional inference. *Journal of the Royal Statistical Society Series B* 49, 1-39.

See Also

[estimateGLMTrendedDisp](#)

Examples

```
design <- matrix(1,4,1)
y <- matrix((rbinom(400,mu=100,size=5)),100,4)
d1 <- dispCoxReidSplineTrend(y, design, df=3)
d2 <- dispCoxReidPowerTrend(y, design)
with(d2,plot(abundance,sqrt(dispersion)))
```

edgeRUsersGuide

View edgeR User's Guide

Description

Finds the location of the edgeR User's Guide and optionally opens it.

Usage

```
edgeRUsersGuide(view=TRUE)
```

Arguments

view logical, should the document be opened using the default PDF document reader?

Details

The function `vignette("edgeR")` will find the short edgeR Vignette which describes how to obtain the Limma User's Guide. The User's Guide is not itself a true vignette because it is not automatically generated using [Sweave](#) during the package build process. This means that it cannot be found using `vignette`, hence the need for this special function.

If the operating system is other than Windows, then the PDF viewer used is that given by `Sys.getenv("R_PDFVIEWER")`. The PDF viewer can be changed using `Sys.putenv(R_PDFVIEWER=)`.

Value

Character string giving the file location. If `view=TRUE`, the PDF document reader is started and the User's Guide is opened, as a side effect.

Author(s)

Gordon Smyth

See Also

[system](#)

Examples

```
# To get the location:
edgeRUsersGuide(view=FALSE)
# To open in pdf viewer:
## Not run: edgeRUsersGuide()
```

equalizeLibSizes

Equalize Library Sizes by Quantile-to-Quantile Normalization

Description

Adjusts counts so that the effective library sizes are equal, preserving fold-changes between groups and preserving biological variability within each group.

Usage

```
equalizeLibSizes(object, dispersion=0, common.lib.size)
```

Arguments

object	DGEList object
dispersion	numeric scalar or vector of dispersion parameters; if a scalar, then a common dispersion parameter is used for all tags
common.lib.size	numeric scalar, the library size to normalize to; default is the geometric mean of the original effective library sizes

Details

Thus function implements the quantile-quantile normalization method of Robinson and Smyth (2008). It computes normalized counts, or pseudo-counts, used by `exactTest` and `estimateCommonDisp`.

Note that the output common library size is a theoretical quantity. The column sums of the normalized counts, while to be exactly equal, nor are they intended to be. However, the expected counts for each tag are equal under the null hypothesis of no differential expression.

Value

A list with components

pseudo.counts	numeric matrix of normalized pseudo-counts
common.lib.size	normalized library size

Author(s)

Mark Robinson, Davis McCarthy, Gordon Smyth

References

Robinson MD and Smyth GK (2008). Small-sample estimation of negative binomial dispersion, with applications to SAGE data. *Biostatistics*, 9, 321-332.

See Also[q2qbinom](#)**Examples**

```

ngenes <- 1000
nlibs <- 2
counts <- matrix(0,ngenes,nlibs)
colnames(counts) <- c("Sample1", "Sample2")
counts[,1] <- rpois(ngenes,lambda=10)
counts[,2] <- rpois(ngenes,lambda=20)
summary(counts)
y <- DGEList(counts=counts)
out <- equalizeLibSizes(y)
summary(out$pseudo.counts)

```

estimateCommonDisp	<i>Estimate Common Negative Binomial Dispersion by Conditional Maximum Likelihood</i>
--------------------	---

Description

Maximizes the negative binomial conditional common likelihood to give the estimate of the common dispersion across all tags.

Usage

```
estimateCommonDisp(object, tol=1e-06, rowsum.filter=5, verbose=FALSE)
```

Arguments

object	DGEList object
tol	the desired accuracy, passed to optimize
rowsum.filter	numeric scalar giving a value for the filtering out of low abundance tags in the estimation of the common dispersion. Only tags with total sum of counts above this value are used in the estimation of the common dispersion.
verbose	logical, if TRUE estimated dispersion and BCV will be printed to standard output.

Details

Implements the method of Robinson and Smyth (2008) for estimating a common dispersion parameter by conditional maximum likelihood. The method of conditional maximum likelihood assumes that library sizes are equal, which is not true in general, so pseudocounts (counts adjusted so that the library sizes are equal) need to be calculated. The function `equalizeLibSizes` is called to adjust the counts using a quantile-to-quantile method, but this requires a fixed value for the common dispersion parameter. To obtain a good estimate for the common dispersion, pseudocounts are calculated under the Poisson model (dispersion is zero) and these pseudocounts are used to give an estimate of the common dispersion. This estimate of the common dispersion is then used to recalculate the pseudocounts, which are used to provide a final estimate of the common dispersion.

Value

Returns object with the following added components:

common.dispersion	estimate of the common dispersion.
pseudo.counts	numeric matrix of quantile-quantile normalized counts. These are counts adjusted so that the library sizes are equal, while preserving differences between groups and variability within each group.
pseudo.lib.size	the common library size to which the counts have been adjusted

Author(s)

Mark Robinson, Davis McCarthy, Gordon Smyth

References

Robinson MD and Smyth GK (2008). Small-sample estimation of negative binomial dispersion, with applications to SAGE data. *Biostatistics*, 9, 321-332

See Also

[equalizeLibSizes](#)

Examples

```
# True dispersion is 1/5=0.2
y <- matrix(rnbinom(1000,mu=10,size=5),ncol=4)
d <- DGEList(counts=y,group=c(1,1,2,2),lib.size=c(1000:1003))
d <- estimateCommonDisp(d, verbose=TRUE)
```

estimateExonGenewiseDisp

Estimate Genewise Dispersions from Exon-Level Count Data

Description

Estimate a dispersion value for each gene from exon-level count data by collapsing exons into the genes to which they belong.

Usage

```
estimateExonGenewiseDisp(y, geneID, group=NULL)
```

Arguments

y	either a matrix of exon-level counts or a DGEList object with (at least) elements counts (table of counts summarized at the exon level) and samples (data frame containing information about experimental group, library size and normalization factor for the library size). Each row of y should represent one exon.
geneID	vector of length equal to the number of rows of y, which provides the gene identifier for each exon in y. These identifiers are used to group the relevant exons into genes for the gene-level analysis of splice variation.

group factor supplying the experimental group/condition to which each sample (column of y) belongs. If NULL (default) the function will try to extract if from y, which only works if y is a DGEList object.

Details

This function can be used to compute genewise dispersion estimates (for an experiment with a one-way, or multiple group, layout) from exon-level count data. `estimateCommonDisp` and `estimateTagwiseDisp` are used to do the computation and estimation, and the default arguments for those functions are used.

Value

`estimateExonGenewiseDisp` returns a vector of genewise dispersion estimates, one for each unique geneID.

Author(s)

Davis McCarthy, Gordon Smyth

See Also

[estimateCommonDisp](#) and related functions for estimating the dispersion parameter for the negative binomial model.

Examples

```
# generate exon counts from NB, create list object
y<-matrix(rnbinom(40,size=1,mu=10),nrow=10)
d<-DGEList(counts=y,group=rep(1:2,each=2))
genes <- rep(c("gene.1","gene.2"), each=5)
estimateExonGenewiseDisp(d, genes)
```

estimateGLMCommonDisp

Estimate Common Dispersion for Negative Binomial GLMs

Description

Estimates a common negative binomial dispersion parameter for a DGE dataset with a general experimental design.

Usage

```
## S3 method for class 'DGEList'
estimateGLMCommonDisp(y, design=NULL, offset=NULL, method="CoxReid", verbose=FALSE, ...)
## Default S3 method:
estimateGLMCommonDisp(y, design=NULL, offset=NULL, method="CoxReid", verbose=FALSE, ...)
```

Arguments

y	object containing read counts, as for glmFit .
design	numeric design matrix, as for glmFit .
offset	numeric vector or matrix of offsets for the log-linear models, as for glmFit .
method	method for estimating the dispersion. Possible values are "CoxReid", "Pearson" or "deviance".
verbose	logical, if TRUE estimated dispersion and BCV will be printed to standard output.
...	other arguments are passed to lower-level functions. See dispCoxReid , dispPearson and dispDeviance for details.

Details

This function calls [dispCoxReid](#), [dispPearson](#) or [dispDeviance](#) depending on the method specified. See [dispCoxReid](#) for details of the three methods and a discussion of their relative performance.

Value

The default method returns a numeric vector of length 1 containing the estimated dispersion.

The DGEList method returns the same DGEList y as input but with common.dispersion as an added component.

Author(s)

Gordon Smyth, Davis McCarthy, Yunshun Chen

References

McCarthy, DJ, Chen, Y, Smyth, GK (2012). Differential expression analysis of multifactor RNA-Seq experiments with respect to biological variation. *Nucleic Acids Research* 40, 4288-4297. <http://nar.oxfordjournals.org/content/40/10/4288>

See Also

[dispCoxReid](#), [dispPearson](#), [dispDeviance](#)

[estimateGLMTrendedDisp](#) for trended dispersion and [estimateGLMTagwiseDisp](#) for tagwise dispersions in the context of a generalized linear model.

[estimateCommonDisp](#) for common dispersion or [estimateTagwiseDisp](#) for tagwise dispersion in the context of a multiple group experiment (one-way layout).

Examples

```
# True dispersion is 1/size=0.1
y <- matrix(rnbinom(1000,mu=10,size=10),ncol=4)
d <- DGEList(counts=y,group=c(1,1,2,2))
design <- model.matrix(~group, data=d$samples)
d1 <- estimateGLMCommonDisp(d, design, verbose=TRUE)

# Compare with classic CML estimator:
d2 <- estimateCommonDisp(d, verbose=TRUE)
```

```
# See example(glmFit) for a different example
```

```
estimateGLMTagwiseDisp
```

Empirical Bayes Tagwise Dispersions for Negative Binomial GLMs

Description

Compute an empirical Bayes estimate of the negative binomial dispersion parameter for each tag or transcript, with expression levels specified by a log-linear model.

Usage

```
## S3 method for class 'DGEList'
estimateGLMTagwiseDisp(y, design=NULL, offset=NULL, dispersion=NULL,
  trend=!is.null(y$trended.dispersion), prior.df=20, span=NULL, ...)
## Default S3 method:
estimateGLMTagwiseDisp(y, design=NULL, offset=NULL, dispersion, trend=TRUE,
  prior.df=20, span=NULL, ...)
```

Arguments

y	matrix of counts or a DGEList object.
design	numeric design matrix, as for glmFit .
trend	logical. Should the prior be the trended dispersion (TRUE) or the common dispersion (FALSE)?
offset	offset matrix for the log-linear model, as for glmFit . Defaults to the log-effective library sizes.
dispersion	common or trended dispersion estimates, used as an initial estimate for the tag-wise estimates. By default uses values stored in the DGEList object.
prior.df	prior degrees of freedom.
span	width of the smoothing window, in terms of proportion of the data set. Default value decreases with the number of tags.
...	other arguments are passed to dispCoxReidInterpolateTagwise .

Details

This function implements the empirical Bayes strategy proposed by McCarthy et al (2012) for estimating the tagwise negative binomial dispersions. The experimental conditions are specified by design matrix allowing for multiple explanatory factors. The empirical Bayes posterior is implemented as a conditional likelihood with tag-specific weights, and the conditional likelihood is computed using Cox-Reid approximate conditional likelihood (Cox and Reid, 1987).

The prior degrees of freedom determines the weight given to the global dispersion trend. The larger the prior degrees of freedom, the more the tagwise dispersions are squeezed towards the global trend.

This function calls the lower-level function [dispCoxReidInterpolateTagwise](#).

Value

estimateGLMTagwiseDisp.DGEList produces a DGEList object, which contains the tagwise dispersion parameter estimate for each tag for the negative binomial model that maximizes the Cox-Reid adjusted profile likelihood. The tagwise dispersions are simply added to the DGEList object provided as the argument to the function.

estimateGLMTagwiseDisp.default returns a vector of the tagwise dispersion estimates.

Author(s)

Gordon Smyth, Davis McCarthy

References

Cox, DR, and Reid, N (1987). Parameter orthogonality and approximate conditional inference. *Journal of the Royal Statistical Society Series B* 49, 1-39.

McCarthy, DJ, Chen, Y, Smyth, GK (2012). Differential expression analysis of multifactor RNA-Seq experiments with respect to biological variation. *Nucleic Acids Research* 40, 4288-4297. <http://nar.oxfordjournals.org/content/40/10/4288>

See Also

[estimateGLMCommonDisp](#) for common dispersion and [estimateGLMTrendedDisp](#) for trended dispersion in the context of a generalized linear model.

[estimateCommonDisp](#) for common dispersion or [estimateTagwiseDisp](#) for tagwise dispersion in the context of a multiple group experiment (one-way layout).

Examples

```
y <- matrix(rnbinom(1000,mu=10,size=10),ncol=4)
d <- DGEList(counts=y,group=c(1,1,2,2),lib.size=c(1000:1003))
design <- model.matrix(~group, data=d$samples) # Define the design matrix for the full model
d <- estimateGLMTrendedDisp(d, design, min.n=10)
d <- estimateGLMTagwiseDisp(d, design)
summary(d$tagwise.dispersion)
```

estimateGLMTrendedDisp

Estimate Trended Dispersion for Negative Binomial GLMs

Description

Estimates the dispersion parameter for each transcript (tag) with a trend that depends on the overall level of expression for the transcript for a DGE dataset for general experimental designs by using Cox-Reid approximate conditional inference for a negative binomial generalized linear model for each transcript (tag) with the unadjusted counts and design matrix provided.

Usage

```
## S3 method for class 'DGEList'
estimateGLMTrendedDisp(y, design=NULL, offset=NULL, method="auto", ...)
## Default S3 method:
estimateGLMTrendedDisp(y, design=NULL, offset=NULL, method="auto", ...)
```

Arguments

y	an object that contains the raw counts for each library (the measure of expression level); it can either be a matrix of counts, or a DGEList object with (at least) elements counts (table of unadjusted counts) and samples (data frame containing information about experimental group, library size and normalization factor for the library size)
design	numeric design matrix, as for glmFit .
method	method (low-level function) used to estimated the trended dispersions. Possible values are "auto" (default, switch to "bin.spline" method if the number of tags is great than 200 and "power" method otherwise), "bin.spline", "bin.loess" (which both result in a call to dispBinTrend), "power" (call to dispCoxReidPowerTrend), or "spline" (call to dispCoxReidSplineTrend).
offset	numeric scalar, vector or matrix giving the offset (in addition to the log of the effective library size) that is to be included in the NB GLM for the transcripts. If a scalar, then this value will be used as an offset for all transcripts and libraries. If a vector, it should be have length equal to the number of libraries, and the same vector of offsets will be used for each transcript. If a matrix, then each library for each transcript can have a unique offset, if desired. In adjustedProfileLik the offset must be a matrix with the same dimension as the table of counts. Default is NULL; if object is a DGEList and offset is NULL then offset will be calculated automatically from codey\$samples.
...	other arguments are passed to lower-level functions. See dispBinTrend , dispCoxReidPowerTrend and dispCoxReidSplineTrend for details.

Details

This is a wrapper function for the lower-level functions that actually carry out the dispersion estimation calculations. Provide a convenient, object-oriented interface for users.

Value

When the input object is a DGEList, estimateGLMTrendedDisp produces a DGEList object, which contains the estimates of the trended dispersion parameter for the negative binomial model according to the method applied.

When the input object is a numeric matrix, the output of one of the lower-level functions dispBinTrend, dispCoxReidPowerTrend or dispCoxReidSplineTrend is returned.

Author(s)

Gordon Smyth, Davis McCarthy, Yunshun Chen

References

Cox, DR, and Reid, N (1987). Parameter orthogonality and approximate conditional inference. *Journal of the Royal Statistical Society Series B* 49, 1-39.

McCarthy, DJ, Chen, Y, Smyth, GK (2012). Differential expression analysis of multifactor RNA-Seq experiments with respect to biological variation. *Nucleic Acids Research* 40, 4288-4297. <http://nar.oxfordjournals.org/content/40/10/4288>

See Also

[dispBinTrend](#), [dispCoxReidPowerTrend](#) and [dispCoxReidSplineTrend](#) for details on how the calculations are done.

[estimateGLMCommonDisp](#) for common dispersion and [estimateGLMTagwiseDisp](#) for (trended) tagwise dispersion in the context of generalized linear models.

[estimateCommonDisp](#) for common dispersion or [estimateTagwiseDisp](#) for tagwise dispersion in the context of a multiple group experiment (one-way layout).

Examples

```
y <- matrix(rnbinom(1000,mu=10,size=10),ncol=4)
d <- DGEList(counts=y,group=c(1,1,2,2),lib.size=c(1000:1003))
design <- model.matrix(~group, data=d$samples) # Define the design matrix for the full model
disp <- estimateGLMTrendedDisp(d, design, min.n=10)
```

estimateTagwiseDisp	<i>Estimate Empirical Bayes Tagwise Dispersion Values</i>
---------------------	---

Description

Estimates tagwise dispersion values by an empirical Bayes method based on weighted conditional maximum likelihood.

Usage

```
estimateTagwiseDisp(object, prior.df=20, trend="movingave", span=NULL, method="grid",
  grid.length=11, grid.range=c(-6,6), tol=1e-06, verbose=FALSE)
```

Arguments

object	object of class DGEList containing (at least) the elements counts (table of raw counts), group (factor indicating group), lib.size (numeric vector of library sizes) and pseudo.alt (numeric matrix of quantile-adjusted pseudocounts calculated under the alternative hypothesis of a true difference between groups; recommended to use the DGEList object provided as the output of estimateCommonDisp)
prior.df	numeric scalar, parameter that indicates the prior degrees of freedom. It is used in calculating prior.n. If NULL, prior.n is then calculated using prior degrees of freedom of 20.
trend	method for estimating dispersion trend. Possible values are "none", "movingave" and "loess".
span	width of the smoothing window, as a proportion of the data set.
method	method for maximizing the posterior likelihood. Possible values are "grid" for interpolation on grid points or "optimize" to call the function of the same name.
grid.length	for method="grid", the number of points on which the interpolation is applied for each tag.
grid.range	for method="grid", the range of the grid points around the trend on a log2 scale.
tol	for method="optimize", the tolerance for Newton-Rhapon iterations.
verbose	logical, if TRUE then diagnostic output is produced during the estimation process.

Details

This function implements the empirical Bayes strategy proposed by Robinson and Smyth (2007) for estimating the tagwise negative binomial dispersions. The experimental design is assumed to be a oneway layout with one or more experimental groups. The empirical Bayes posterior is implemented as a conditional likelihood with tag-specific weights.

The prior values for the dispersions are determined by a global trend. The individual tagwise dispersions are then squeezed towards this trend. The prior degrees of freedom determines the weight given to the prior. The larger the prior degrees of freedom, the more the tagwise dispersions are squeezed towards the global trend. If the number of libraries is large, the prior becomes less important and the tagwise dispersion are determined more by the individual tagwise data.

If `trend="none"`, then the prior dispersion is just a constant, the common dispersion. Otherwise, the trend is determined by a moving average (`trend="movingave"`) or loess smoother applied to the tagwise conditional log-likelihood. `method="loess"` applies a loess curve of degree 0 as implemented in [loessByCol](#).

`method="optimize"` is not recommended for routine use as it is very slow. It is included for testing purposes.

Value

An object of class `DGEList` with the same components as for [estimateCommonDisp](#) plus the following:

<code>prior.n</code>	estimate of the prior weight, i.e. the smoothing parameter that indicates the weight to put on the common likelihood compared to the individual tag's likelihood; <code>prior.n</code> of 10 means that the common likelihood is given 10 times the weight of the individual tag/gene's likelihood in the estimation of the tag/genewise dispersion
<code>tagwise.dispersion</code>	tag- or gene-wise estimates of the dispersion parameter

Author(s)

Mark Robinson, Davis McCarthy, Yunshun Chen and Gordon Smyth

References

Robinson, MD, and Smyth, GK (2007). Moderated statistical tests for assessing differences in tag abundance. *Bioinformatics* 23, 2881-2887. <http://bioinformatics.oxfordjournals.org/content/23/21/2881>

See Also

[estimateCommonDisp](#) is usually run before `estimateTagwiseDisp`.

[movingAverageByCol](#) and [loessByCol](#) implement the moving average or loess smoothers.

Examples

```
# See exactTest
```

estimateTrendedDisp *Estimate Empirical Bayes Trended Dispersion Values*

Description

Estimates trended dispersion values by an empirical Bayes method.

Usage

```
estimateTrendedDisp(object, method="bin.spline", df=5, span=2/3)
```

Arguments

object	object of class DGEList containing (at least) the elements counts (table of raw counts), group (factor indicating group), lib.size (numeric vector of library sizes) and pseudo.alt (numeric matrix of quantile-adjusted pseudocounts calculated under the alternative hypothesis of a true difference between groups; recommended to use the DGEList object provided as the output of estimateCommonDisp)
method	method used to estimated the trended dispersions. Possible values are "spline", and "loess".
df	integer giving the degrees of freedom of the spline function if "spline" method is used, see ns in the splines package. Default is 5.
span	scalar, passed to loess to determine the amount of smoothing for the loess fit when "loess" method is used. Default is 2/3.

Details

This function takes the binned common dispersion and abundance, and fits a smooth curve through these binned values using either natural cubic splines or loess. From this smooth curve it predicts the dispersion value for each gene based on the gene's overall abundance. This results in estimates for the NB dispersion parameter which have a dependence on the overall expression level of the gene, and thus have an abundance-dependent trend.

Value

An object of class DGEList with the same components as for [estimateCommonDisp](#) plus the trended dispersion estimates for each gene or tag.

Author(s)

Yunshun Chen and Gordon Smyth

See Also

[estimateCommonDisp](#) estimates a common value for the dispersion parameter for all tags/genes - should generally be run before estimateTrendedDisp.

Examples

```

y <- matrix(rnbinom(6000, mu=100, size=10), 1000, 6)
group <- c(0,0,0,1,1,1)
d <- DGEList(y, group=group)
d <- estimateCommonDisp(d)
d <- estimateTrendedDisp(d)

```

exactTest

Exact Tests for Differences between Two Groups of Negative-Binomial Counts

Description

Compute genewise exact tests for differences in the means between two groups of negative-binomially distributed counts.

Usage

```

exactTest(object, pair=1:2, dispersion="auto", rejection.region="doubletail",
          big.count=900, prior.count.total=0.5)
exactTestDoubleTail(y1, y2, dispersion=0, big.count=900)
exactTestBySmallP(y1, y2, dispersion=0, big.count=900)
exactTestByDeviance(y1, y2, dispersion=0, big.count=900)
exactTestBetaApprox(y1, y2, dispersion=0)

```

Arguments

object	an object of class DGEList .
pair	vector of length two, either numeric or character, providing the pair of groups to be compared; if a character vector, then should be the names of two groups (e.g. two levels of <code>object\$samples\$group</code>); if numeric, then groups to be compared are chosen by finding the levels of <code>object\$samples\$group</code> corresponding to those numeric values and using those levels as the groups to be compared; if NULL, then first two levels of <code>object\$samples\$group</code> (a factor) are used. Note that the first group listed in the pair is the baseline for the comparison—so if the pair is <code>c("A", "B")</code> then the comparison is B - A, so genes with positive log-fold change are up-regulated in group B compared with group A (and vice versa for genes with negative log-fold change).
dispersion	either a numeric vector of dispersions or a character string indicating that dispersions should be taken from the data object. If a numeric vector, then can be either of length one or of length equal to the number of tags. Allowable character values are "common", "trended", "tagwise" or "auto". Default behavior ("auto" is to use most complex dispersions found in data object.
rejection.region	type of rejection region for two-sided exact test. Possible values are "doubletail", "smallp" or "deviance".
big.count	count size above which asymptotic beta approximation will be used.
prior.count.total	prior count used to shrink log-fold-changes. Larger values produce more shrinkage.

y1	numeric matrix of counts for the first the two experimental groups to be tested for differences. Rows correspond to genes or transcripts and columns to libraries. Libraries are assumed to be equal in size - e.g. adjusted pseudocounts from the output of equalizeLibSizes .
y2	numeric matrix of counts for the second of the two experimental groups to be tested for differences. Rows correspond to genes or transcripts and columns to libraries. Libraries are assumed to be equal in size - e.g. adjusted pseudocounts from the output of equalizeLibSizes . Must have the same number of rows as y1.

Details

The functions test for differential expression between two groups of count libraries. They implement the exact test proposed by Robinson and Smyth (2008) for a difference in mean between two groups of negative binomial random variables. The functions accept two groups of count libraries, and a test is performed for each row of data. For each row, the test is conditional on the sum of counts for that row. The test can be viewed as a generalization of the well-known exact binomial test, implemented in the function `binom.test` in the `stats` package, but generalized to overdispersed counts.

The low level functions `exactTestDoubleTail`, `exactTestBetaApprox`, `exactTestBySmallP` and `exactTestByDeviance` all assume that the libraries have been normalized to have the same size (expected column sum under the null hypothesis). The higher level function `exactTest` is intended to be called by users. This has a more object-orientated flavor and produces an object containing all the necessary components for downstream analysis. `exactTest` equalizes the library sizes using [equalizeLibSizes](#) before calling one of the low level functions.

The functions `exactTestDoubleTail`, `exactTestBySmallP` and `exactTestByDeviance` correspond to different ways to define the two-sided rejection region when the two groups have different numbers of samples. `exactTestBySmallP` implements the method of small probabilities as proposed by Robinson and Smyth (2008). This method corresponds to `binom.test` when the dispersion is near zero, but gives poor results when the dispersion is very large. `exactTestDoubleTail` computes two-sided p-values by doubling the smaller tail probability. `exactTestByDeviance` uses the deviance goodness of fit statistics to define the rejection region, and is therefore equivalent to a conditional likelihood ratio test. This has good statistical properties but is relatively slow to compute. For general remarks on different types of rejection regions for exact tests see Gibbons and Pratt (1975).

`exactTestBetaApprox` implements an asymptotic beta distribution approximation to the conditional count distribution.

Value

`exactTestDoubleTail` and friends produce a numeric vector of genewise p-values, one for each row of y1 and y2.

`exactTest` produces an object of class `DGEEexact` containing the following components:

table	data frame containing columns for the log ₂ -fold-change, logFC, the average log ₂ -counts-per-million, logCPM, and the two-sided p-value PValue
comparison	character vector giving the names of the two groups being compared
genes	optional data frame containing annotation for transcript; taken from object

Author(s)

Mark Robinson, Davis McCarthy, Gordon Smyth

References

Robinson MD and Smyth GK (2008). Small-sample estimation of negative binomial dispersion, with applications to SAGE data. *Biostatistics*, 9, 321-332.

Gibbons, JD and Pratt, JW (1975). P-values: interpretation and methodology. *The American Statistician* 29, 20-25.

See Also

[equalizeLibSizes](#), [binomTest](#)

Examples

```
# generate raw counts from NB, create list object
y <- matrix(rnbinom(80,size=1/0.2,mu=10),nrow=20,ncol=4)
rownames(y) <- paste("Gene",1:nrow(y),sep=".")
group <- factor(c(1,1,2,2))
d <- DGEList(counts=y,group=group,lib.size=rep(1000,4))

# estimate dispersions and find differences in expression
d <- estimateCommonDisp(d, verbose=TRUE)
d <- estimateTagwiseDisp(d)
de <- exactTest(d)
topTags(de)

# same example using low level exactTest function directly
p.value <- exactTestDoubleTail(y[,1:2],y[,3:4],dispersion=0.2)
```

expandAsMatrix

expandAsMatrix

Description

Expand scalar or vector to a matrix.

Usage

```
expandAsMatrix(x, dim)
```

Arguments

x	scalar, vector or matrix. If a vector, length must match one of the output dimensions.
dim	required dimension for the output matrix.

Details

This function expands a row or column vector to be a matrix. It is used internally in edgeR to convert offsets to a matrix.

Value

Numeric matrix of dimension dim.

Author(s)

Gordon Smyth

See Also[mglmLS](#).**Examples**

```
expandAsMatrix(1:3,c(4,3))
expandAsMatrix(1:4,c(4,3))
```

`getCounts`*Extract Specified Component of a DGEList Object*

Description

`getCounts(y)` returns the matrix of read counts `y$counts`.

`getOffset(y)` returns offsets for the log-linear predictor account for sequencing depth and possibly other normalization factors. Specifically it returns the matrix `y$offset` if it is non-null, otherwise it returns the log product of `lib.size` and `norm.factors` from `y$samples`.

`getDispersion(y)` returns the most complex dispersion estimates (common, trended or tagwise) found in `y`.

Usage

```
getCounts(y)
getOffset(y)
getDispersion(y)
```

Arguments

`y` DGEList object containing (at least) the elements `counts` (table of raw counts), `group` (factor indicating group) and `lib.size` (numeric vector of library sizes)

Value

`getCounts` returns the matrix of counts. `getOffset` returns a numeric matrix or vector. `getDispersion` returns vector of dispersion values.

Author(s)

Mark Robinson, Davis McCarthy, Gordon Smyth

See Also[DGEList-class](#)

Examples

```
# generate raw counts from NB, create list object
y <- matrix(rnbinom(20,size=5,mu=10),5,4)
d <- DGEList(counts=y, group=c(1,1,2,2), lib.size=1001:1004)
getCounts(d)
getOffset(d)
d <- estimateCommonDisp(d)
getDispersion(d)
```

getPriorN

Get a Recommended Value for Prior N from DGEList Object

Description

Returns the lib.size component of the samples component of DGEList object multiplied by the norm.factors component

Usage

```
getPriorN(y, design=NULL, prior.df=20)
```

Arguments

y	a DGEList object with (at least) elements counts (table of unadjusted counts) and samples (data frame containing information about experimental group, library size and normalization factor for the library size)
design	numeric matrix (optional argument) giving the design matrix for the GLM that is to be fit. Must be of full column rank. If provided design is used to determine the number of parameters to be fit in the statistical model and therefore the residual degrees of freedom. If left as the default (NULL) then the y\$samples\$group element of the DGEList object is used to determine the residual degrees of freedom.
prior.df	numeric scalar giving the weight, in terms of prior degrees of freedom, to be given to the common parameter likelihood when estimating tagwise dispersion estimates.

Details

When estimating tagwise dispersion values using [estimateTagwiseDisp](#) or [estimateGLMTagwiseDisp](#) we need to decide how much weight to give to the common parameter likelihood in order to smooth (or stabilize) the dispersion estimates. The best choice of value for the prior.n parameter varies between datasets depending on the number of samples in the dataset and the complexity of the model to be fit. The value of prior.n should be inversely proportional to the residual degrees of freedom. We have found that choosing a value for prior.n that is equivalent to giving the common parameter likelihood 20 degrees of freedom generally gives a good amount of smoothing for the tagwise dispersion estimates. This function simply recommends an appropriate value for prior.n—to be used as an argument for [estimateTagwiseDisp](#) or [estimateGLMTagwiseDisp](#)—given the experimental design at hand and the chosen prior degrees of freedom.

Value

getPriorN returns a numeric scalar

Author(s)

Davis McCarthy, Gordon Smyth

See Also

[DGEList](#) for more information about the DGEList class. [as.matrix.DGEList](#).

Examples

```
# generate raw counts from NB, create list object
y<-matrix(rnbinom(20,size=1,mu=10),nrow=5)
d<-DGEList(counts=y,group=rep(1:2,each=2),lib.size=rep(c(1000:1001),2))
getPriorN(d)
```

 glmFit

Genewise Negative Binomial Generalized Linear Models

Description

Fit a negative binomial generalized log-linear model to the read counts for each gene or transcript. Conduct genewise statistical tests for a given coefficient or coefficient contrast.

Usage

```
## S3 method for class 'DGEList'
glmFit(y, design=NULL, dispersion=NULL, offset=NULL, weights=NULL, lib.size=NULL,
       prior.count.total=0.5, start=NULL, method="auto", ...)
glmLRT(glmfit, coef=ncol(glmfit$design), contrast=NULL)
glmQLFTest(glmfit, coef=ncol(glmfit$design), contrast=NULL, abundance.trend=TRUE)
```

Arguments

- | | |
|------------|--|
| y | an object that contains the raw counts for each library (the measure of expression level); alternatively, a matrix of counts, or a DGEList object with (at least) elements counts (table of unadjusted counts) and samples (data frame containing information about experimental group, library size and normalization factor for the library size) |
| design | numeric matrix giving the design matrix for the tagwise linear models. Must be of full column rank. Defaults to a single column of ones, equivalent to treating the columns as replicate libraries. |
| dispersion | numeric scalar or vector providing the value for the dispersion parameter that is used in fitting the GLM for each transcript. Can be a common value for all tags, or a vector of values can provide a unique dispersion value for each tag. If NULL will be extracted from y, with order of precedence: tagwise dispersion, trended dispersions, common dispersion. |
| offset | numeric matrix of same size as y giving offsets for the log-linear models. Can be a scalar or a vector of length $\text{ncol}\{y\}$, in which case it is expanded out to a matrix. |
| weights | optional numeric matrix giving prior weights for the observations (for each library and transcript) to be used in the GLM calculations. Not supported by methods "linesearch" or "levenberg". |

lib.size	numeric vector of length <code>ncol(y)</code> giving library sizes. Only used if <code>offset=NULL</code> , in which case <code>offset</code> is set to <code>log(lib.size)</code> . Defaults to <code>colSums(y)</code> .
prior.count.total	the total number of prior counts to be added to each row of data to shrink the estimated log-fold-changes towards zero.
start	optional numeric matrix of initial estimates for the linear model coefficients.
method	which fitting algorithm to use. Possible values are "auto", "linesearch", "levenberg" or "simple".
...	other arguments are passed to lower-level functions, for example to <code>mglmLS</code> .
glmfit	a DGEGLM object, usually output from <code>glmFit</code> .
coef	integer or character vector indicating which coefficients of the linear model are to be tested equal to zero. Values must be columns or column names of design. Defaults to the last coefficient. Ignored if <code>contrast</code> is specified.
contrast	numeric vector or matrix specifying one or more contrasts of the linear model coefficients to be tested equal to zero. Number of rows must equal to the number of columns of design. If specified, then takes precedence over <code>coef</code> .
abundance.trend	logical, whether to allow an abundance-dependent trend when estimating the prior values for the quasi-likelihood multiplicative dispersion parameter.

Details

`glmFit` and `glmLRT` implement generalized linear model (glm) methods developed by McCarthy et al (2012).

`glmFit` fits genewise negative binomial glms, all with the same design matrix but possibly different dispersions, offsets and weights. When the design matrix defines a one-way layout, or can be re-parametrized to a one-way layout, the glms are fitting very quickly using `mglmOneGroup`. Otherwise the default fitting method, implemented in `mglmLS`, is a parallelized line search algorithm described by McCarthy et al (2012). Other possible fitting methods are `mglmLevenberg` and `mglmSimple`.

Positive `prior.count.total` cause the returned coefficients to be shrunk in such a way that fold-changes between the treatment conditions are decreased. In particular, infinite fold-changes are avoided. Larger values cause more shrinkage. The returned coefficients are affected but not the likelihood ratio tests or p-values.

`glmLRT` conducts likelihood ratio tests for one or more coefficients in the linear model. If `coef` is used, the null hypothesis is that all the coefficients indicated by `coef` are equal to zero. If `contrast` is non-null, then the null hypothesis is that the specified contrast of the coefficients is equal to zero. For example, a contrast of `c(0,1,-1)`, assuming there are three coefficients, would test the hypothesis that the second and third coefficients are equal.

`glmQLFTest` implements the quasi-likelihood method of Lund et al (2012). It behaves the same as `glmLRT` except that it replaces likelihood ratio tests with quasi-likelihood F-tests for coefficients in the linear model. This function calls the `limma` function `squeezeVar` to conduct empirical Bayes smoothing of the genewise multiplicative dispersions. Note that the `QuasiSeq` package provides an alternative implementation of Lund et al (2012), with slightly different glm, trend and FDR methods.

Value

`glmFit` produces an object of class `DGEGLM` containing components `counts`, `samples`, `genes` and `abundance` from `y` plus the following new components:

design	design matrix as input.
weights	matrix of weights as input.
df.residual	numeric vector of residual degrees of freedom, one for each tag.
offset	numeric matrix of linear model offsets.
dispersion	vector of dispersions used for the fit.
coefficients	numeric matrix of estimated coefficients from the glm fits, on the natural log scale, of size nrow(y) by ncol(design).
fitted.values	matrix of fitted values from glm fits, same number of rows and columns as y.
deviance	numeric vector of deviances, one for each tag.

glmLRT and glmQFTest produce objects of class DGELRT with the same components as for glmfit plus the following:

table	data frame with the same rows as y containing the log2-fold changes, test statistics and p-values, ready to be displayed by topTags..
comparison	character string describing the coefficient or the contrast being tested.

The data frame table contains the following columns:

logFC	log2-fold change of expression between conditions being tested.
logCPM	average log2-counts per million, the average taken over all libraries in y.
LR	likelihood ratio statistics (only for glmLRT).
F	F-statistics (only for glmQFTest).
PValue	p-values.

Author(s)

Davis McCarthy and Gordon Smyth

References

- McCarthy, DJ, Chen, Y, Smyth, GK (2012). Differential expression analysis of multifactor RNA-Seq experiments with respect to biological variation. *Nucleic Acids Research* 40, 4288-4297. <http://nar.oxfordjournals.org/content/40/10/4288>
- Lund, SP, Nettleton, D, McCarthy, DJ, Smyth, GK (2012). Detecting differential expression in RNA-sequence data using quasi-likelihood with shrunken dispersion estimates. *Statistical Applications in Genetics and Molecular Biology*. (Accepted 31 July 2012)

See Also

Low-level computations are done by [mgmOneGroup](#), [mgmLS](#), [mgmLevenberg](#) or [mgmSimple](#). [topTags](#) displays results from glmLRT or glmQLFTest.

The QuasiSeq package gives an alternative implementation of glmQLFTest based on the same statistical ideas.

Examples

```

nlibs <- 3
ntags <- 100
dispersion.true <- 0.1

# Make first transcript respond to covariate x
x <- 0:2
design <- model.matrix(~x)
beta.true <- cbind(Beta1=2,Beta2=c(2,rep(0,ntags-1)))
mu.true <- 2^(beta.true %*% t(design))

# Generate count data
y <- rbinom(ntags*nlibs,mu=mu.true,size=1/dispersion.true)
y <- matrix(y,ntags,nlibs)
colnames(y) <- c("x0","x1","x2")
rownames(y) <- paste("Gene",1:ntags,sep="")
d <- DGEList(y)

# Normalize
d <- calcNormFactors(d)

# Fit the NB GLMs
fit <- glmFit(d, design, dispersion=dispersion.true)

# Likelihood ratio tests for trend
results <- glmLRT(fit, coef=2)
topTags(results)

# Estimate the dispersion (may be unreliable with so few tags)
d <- estimateGLMCommonDisp(d, design, verbose=TRUE)

```

gof

*Goodness of Fit Tests for Multiple GLM Fits***Description**

Conducts deviance goodness of fit tests for each fit in a DGEGLM object

Usage

```
gof(glmfit, pcutoff=0.1, adjust="holm", plot=FALSE, main="qq-plot of genewise goodness of fit", ...)
```

Arguments

glmfit	DGEGLM object containing results from fitting NB GLMs to genes in a DGE dataset. Output from glmFit.
pcutoff	scalar giving the cut-off value for the Holm-adjusted p-value. Genes with Holm-adjusted p-values lower than this cutoff value are flagged as ‘dispersion outlier’ genes.
adjust	method used to adjust goodness of fit p-values for multiple testing.
plot	logical, if TRUE a qq-plot is produced.
main	character, title for the plot.
...	other arguments are passed to qqnorm.

Details

If `plot=TRUE`, produces a plot similar to Figure 2 of McCarthy et al (2012).

Value

This function returns a list with the following components:

<code>gof.statistics</code>	numeric vector of deviance statistics, which are the statistics used for the goodness of fit test
<code>gof.pvalues</code>	numeric vector of p-values providing evidence of poor fit; computed from the chi-square distribution on the residual degrees of freedom from the GLM fits.
<code>outlier</code>	logical vector indicating whether or not each gene is a ‘dispersion outlier’ (i.e., the model fit is poor for that gene indicating that the dispersion estimate is not good for that gene).
<code>df</code>	scalar, the residual degrees of freedom from the GLM fit for which the goodness of fit statistics have been computed. Also the degrees of freedom for the goodness of fit statistics for the LR (chi-square) test for significance.

Author(s)

Davis McCarthy and Gordon Smyth

References

McCarthy, DJ, Chen, Y, Smyth, GK (2012). Differential expression analysis of multifactor RNA-Seq experiments with respect to biological variation. *Nucleic Acids Research* 40, 4288-4297 <http://nar.oxfordjournals.org/content/40/10/4288>

See Also

[qqnorm](#).

[glmFit](#) for more information on fitting NB GLMs to DGE data.

Examples

```
nlibs <- 3
ntags <- 100
dispersion.true <- 0.1

# Make first transcript respond to covariate x
x <- 0:2
design <- model.matrix(~x)
beta.true <- cbind(Beta1=2,Beta2=c(2,rep(0,ntags-1)))
mu.true <- 2^(beta.true %*% t(design))

# Generate count data
y <- rbinom(ntags*nlibs,mu=mu.true,size=1/dispersion.true)
y <- matrix(y,ntags,nlibs)
colnames(y) <- c("x0","x1","x2")
rownames(y) <- paste("Gene",1:ntags,sep="")
d <- DGEList(y)

# Normalize
d <- calcNormFactors(d)
```

```
# Fit the NB GLMs
fit <- glmFit(d, design, dispersion=dispersion.true)
# Check how good the fit is for each gene
gof(fit)
```

goodTuring

Good-Turing Frequency Estimation

Description

Non-parametric empirical Bayes estimates of the frequencies of observed (and unobserved) species.

Usage

```
goodTuring(x, conf=1.96)
goodTuringPlot(x)
goodTuringProportions(counts)
```

Arguments

x	numeric vector of non-negative integers, representing the observed frequency of each species.
conf	confidence factor, as a quantile of the standard normal distribution, used to decide for what values the log-linear relationship between frequencies and frequencies of frequencies is acceptable.
counts	matrix of counts

Details

Observed counts are assumed to be Poisson distributed. Using a non-parametric empirical Bayes strategy, the algorithm evaluates the posterior expectation of each species mean given its observed count. The posterior means are then converted to proportions. In the empirical Bayes step, the counts are smoothed by assuming a log-linear relationship between frequencies and frequencies of frequencies. The fundamentals of the algorithm are from Good (1953). Gale and Sampson (1995) proposed a simplified algorithm with a rule for switching between the observed and smoothed frequencies, and it is Gale and Sampson's simplified algorithm that is implemented here. The number of zero values in x are not used in the algorithm, but is returned by this function.

Sampson gives a C code version on his webpage at <http://www.grsampson.net/RGoodTur.html> which gives identical results to this function.

goodTuringPlot plots log-probability (i.e., log frequencies of frequencies) versus log-frequency.

goodTuringProportions runs goodTuring on each column of data, then uses the results to predict the proportion of each tag in each library.

Value

goodTuring returns a list with components

count	observed frequencies, i.e., the unique positive values of x
n	frequencies of frequencies

n0 frequency of zero, i.e., number of zeros found in x
 proportion estimated proportion of each species given its count
 P0 estimated combined proportion of all undetected species

goodTuringProportions returns a matrix of proportions of the same size as counts.

Author(s)

Aaron Lun and Gordon Smyth, adapted from Sampson's C code from <http://www.grsampson.net/RGoodTur.html>

References

Gale, WA, and Sampson, G (1995). Good-Turing frequency estimation without tears. *Journal of Quantitative Linguistics* 2, 217-237.

Examples

```
# True means of observed species
lambda <- rnbinom(10000,mu=2,size=1/10)
lambda <- lambda[lambda>1]

# Observed frequencies
Ntrue <- length(lambda)
x <- rpois(Ntrue, lambda=lambda)
freq <- goodTuring(x)
goodTuringPlot(x)
```

 loessByCol

Locally Weighted Mean By Column

Description

Smooth columns of matrix by non-robust loess curves of degree 0.

Usage

```
loessByCol(y, x=NULL, span=0.5)
locfitByCol(y, x=NULL, weights=1, span=0.5, degree=0)
```

Arguments

y numeric matrix of response variables.
 x numeric covariate vector of length nrow(y), defaults to equally spaced.
 span width of the smoothing window, in terms of proportion of the data set. Larger values produce smoother curves.
 weights relative weights of each observation, one for each covariate value.
 degree degree of local polynomial fit

Details

Fits a loess curve with degree 0 to each column of the response matrix, using the same covariate vector for each column. The smoothed column values are tricube-weighted means of the original values.

locfitByCol uses the locfit.raw function of the locfit package.

Value

A list containing a numeric matrix with smoothed columns and a vector of leverages for each covariate value.

locfitByCol returns a numeric matrix.

Author(s)

Aaron Lun for loessByCol, replacing earlier R code by Davis McCarthy. Gordon Smyth for locfitByCol.

See Also

[loess](#)

Examples

```
y <- matrix(rnorm(100*3), nrow=100, ncol=3)
head(y)
out <- loessByCol(y)
head(out$fitted.values)
```

maPlot

Plots Log-Fold Change versus Log-Concentration (or, M versus A) for Count Data

Description

To represent counts that were low (e.g. zero in 1 library and non-zero in the other) in one of the two conditions, a 'smear' of points at low A value is presented.

Usage

```
maPlot(x, y, logAbundance=NULL, logFC=NULL, normalize=FALSE, plot.it=TRUE,
smearWidth=1, col=NULL, allCol="black", lowCol="orange", deCol="red",
de.tags=NULL, smooth.scatter=FALSE, lowess=FALSE, ...)
```

Arguments

x vector of counts or concentrations (group 1)

y vector of counts or concentrations (group 2)

logAbundance vector providing the abundance of each tag on the log₂ scale. Purely optional (default is NULL), but in combination with logFC provides a more direct way to create an MA-plot if the log-abundance and log-fold change are available.

logFC	vector providing the log-fold change for each tag for a given experimental contrast. Default is NULL, only to be used together with logAbundance as both need to be non-null for their values to be used.
normalize	logical, whether to divide x and y vectors by their sum
plot.it	logical, whether to produce a plot
smearWidth	scalar, width of the smear
col	vector of colours for the points (if NULL, uses allCol and lowCol)
allCol	colour of the non-smearred points
lowCol	colour of the smearred points
deCol	colour of the DE (differentially expressed) points
de.tags	indices for tags identified as being differentially expressed; use exactTest to identify DE genes
smooth.scatter	logical, whether to produce a 'smooth scatter' plot using the KernSmooth::smoothScatter function or just a regular scatter plot; default is FALSE, i.e. produce a regular scatter plot
lowess	logical, indicating whether or not to add a lowess curve to the MA-plot to give an indication of any trend in the log-fold change with log-concentration
...	further arguments passed on to plot

Details

The points to be smearred are identified as being equal to the minimum in one of the two groups. The smear is created by using random uniform numbers of width smearWidth to the left of the minimum A value.

Value

a plot to the current device (if plot.it=TRUE), and invisibly returns the M (logFC) and A (log-Conc) values used for the plot, plus identifiers w and v of genes for which M and A values, or just M values, respectively, were adjusted to make a nicer looking plot.

Author(s)

Mark Robinson, Davis McCarthy

See Also

[plotSmear](#)

Examples

```
y <- matrix(rnbinom(10000,mu=5,size=2),ncol=4)
maPlot(y[,1], y[,2])
```

maximizeInterpolant *Maximize a function given a table of values by spline interpolation.*

Description

Maximize a function given a table of values by spline interpolation.

Usage

```
maximizeInterpolant(x, y)
```

Arguments

x numeric vector of the inputs of the function.
y numeric matrix of function values at the values of x. Columns correspond to x values and each row corresponds to a different function to be maximized.

Details

Calculates the cubic spline interpolant for each row the method of Forsythe et al (1977) using the function `fmm_spline` from `splines.c` in the `stats` package). Then calculates the derivatives of the spline segments adjacent to the input with the maximum function value. This allows identification of the maximum of the interpolating spline.

Value

numeric vector of input values at which the function maximums occur.

Author(s)

Aaron Lun, improving on earlier code by Gordon Smyth

References

Forsythe, G. E., Malcolm, M. A. and Moler, C. B. (1977). *Computer Methods for Mathematical Computations*, Prentice-Hall.

Examples

```
x <- seq(0,1,length=10)
y <- rnorm(10,1,1)
maximizeInterpolant(x,y)
```

maximizeQuadratic	<i>Maximize a function given a table of values by quadratic interpolation.</i>
-------------------	--

Description

Maximize a function given a table of values by quadratic interpolation.

Usage

```
maximizeQuadratic(y, x=1:ncol(y))
```

Arguments

y	numeric matrix of response values.
x	numeric matrix of inputs of the function of same dimension as y. If a vector, must be a row vector of length equal to ncol(y).

Details

For each row of y, finds the three x values bracketing the maximum of y, interpolates a quadratic polynomial through these y for these three values and solves for the location of the maximum of the polynomial.

Value

numeric vector of length equal to nrow(y) giving the x-value at which y is maximized.

Author(s)

Yunshun Chen and Gordon Smyth

See Also

[maximizeInterpolant](#)

Examples

```
y <- matrix(rnorm(5*9),5,9)
maximizeQuadratic(y)
```

 meanvar

Explore the mean-variance relationship for DGE data

Description

Appropriate modelling of the mean-variance relationship in DGE data is important for making inferences about differential expression. Here are functions to compute tag/gene means and variances, as well as looking at these quantities when data is binned based on overall expression level.

Usage

```
plotMeanVar(object, meanvar=NULL, show.raw.vars=FALSE, show.tagwise.vars=FALSE,
  show.binned.common.disp.vars=FALSE, show.ave.raw.vars=TRUE, scalar=NULL,
  NBline=FALSE, nbins=100, log.axes="xy", xlab=NULL, ylab=NULL, ...)
binMeanVar(x, group, nbins=100, common.dispersion=FALSE, object=NULL)
```

Arguments

- | | |
|------------------------------|--|
| object | DGEList object containing the raw data and dispersion value. According to the method desired for computing the dispersion, either <code>estimateCommonDisp</code> and (possibly) <code>estimateTagwiseDisp</code> should be run on the DGEList object before using <code>plotMeanVar</code> . The argument <code>object</code> must be supplied in the function <code>binMeanVar</code> if common dispersion values are to be computed for each bin. |
| meanvar | list (optional) containing the output from <code>binMeanVar</code> or the returned value of <code>plotMeanVar</code> . Providing this object as an argument will save time in computing the tag/gene means and variances when producing a mean-variance plot. |
| show.raw.vars | logical, whether or not to display the raw (pooled) gene/tag variances on the mean-variance plot. Default is FALSE. |
| show.tagwise.vars | logical, whether or not to display the estimated genewise/tagwise variances on the mean-variance plot. Default is FALSE. |
| show.binned.common.disp.vars | logical, whether or not to compute the common dispersion for each bin of tags and show the variances computed from those binned common dispersions and the mean expression level of the respective bin of tags. Default is FALSE. |
| show.ave.raw.vars | logical, whether or not to show the average of the raw variances for each bin of tags plotted against the average expression level of the tags in the bin. Averages are taken on the square root scale as regular arithmetic means are likely to be upwardly biased for count data, whereas averaging on the square scale gives a better summary of the mean-variance relationship in the data. The default is TRUE. |
| scalar | vector (optional) of scaling values to divide counts by. Would expect to have this the same length as the number of columns in the count matrix (i.e. the number of libraries). |
| NBline | logical, whether or not to add a line on the graph showing the mean-variance relationship for a NB model with common dispersion. |

nbins	scalar giving the number of bins (formed by using the quantiles of the genewise mean expression levels) for which to compute average means and variances for exploring the mean-variance relationship. Default is 100 bins
log.axes	character vector indicating if any of the axes should use a log scale. Default is "xy", which makes both y and x axes on the log scale. Other valid options are "x" (log scale on x-axis only), "y" (log scale on y-axis only) and "" (linear scale on x- and y-axis).
xlab	character string giving the label for the x-axis. Standard graphical parameter. If left as the default NULL, then the x-axis label will be set to "logConc".
ylab	character string giving the label for the y-axis. Standard graphical parameter. If left as the default NULL, then the x-axis label will be set to "logConc".
...	further arguments passed on to plot
x	matrix of count data, with rows representing tags/genes and columns representing samples
group	factor giving the experimental group or condition to which each sample (i.e. column of x or element of y) belongs
common.dispersion	logical, whether or not to compute the common dispersion for each bin of tags.

Details

This function is useful for exploring the mean-variance relationship in the data. Raw variances are, for each gene, the pooled variance of the counts from each sample, divided by a scaling factor (by default the effective library size). The function will plot the average raw variance for tags split into nbins bins by overall expression level. The averages are taken on the square-root scale as for count data the arithmetic mean is upwardly biased. Taking averages on the square-root scale provides a useful summary of how the variance of the gene counts change with respect to expression level (abundance). A line showing the Poisson mean-variance relationship (mean equals variance) is always shown to illustrate how the genewise variances may differ from a Poisson mean-variance relationship. Optionally, the raw variances and estimated tagwise variances can also be plotted. Estimated tagwise variances can be calculated using either qCML estimates of the tagwise dispersions (estimateTagwiseDisp) or Cox-Reid conditional inference estimates (CRDisp). A log-log scale is used for the plot.

Value

plotMeanVar produces a mean-variance plot for the DGE data using the options described above. plotMeanVar and binMeanVar both return a list with the following components:

avemeans	vector of the average expression level within each bin of genes, with the average taken on the square-root scale
avevars	vector of the average raw pooled gene-wise variance within each bin of genes, with the average taken on the square-root scale
bin.means	list containing the average (mean) expression level for genes divided into bins based on amount of expression
bin.vars	list containing the pooled variance for genes divided into bins based on amount of expression
means	vector giving the mean expression level for each gene
vars	vector giving the pooled variance for each gene
bins	list giving the indices of the tags in each bin, ordered from lowest expression bin to highest

Author(s)

Davis McCarthy

See Also[plotMDS.DGEList](#), [plotSmear](#) and [maPlot](#) provide more ways of visualizing DGE data.**Examples**

```

y <- matrix(rnbinom(1000,mu=10,size=2),ncol=4)
d <- DGEList(counts=y,group=c(1,1,2,2),lib.size=c(1000:1003))
plotMeanVar(d) # Produce a straight-forward mean-variance plot
meanvar <- plotMeanVar(d, show.raw.vars=TRUE) # Produce a mean-variance plot with the raw variances shown and sa

## If we want to show estimated tagwise variances on the plot, we must first estimate them!
d <- estimateCommonDisp(d) # Obtain an estimate of the dispersion parameter
d <- estimateTagwiseDisp(d) # Obtain tagwise dispersion estimates
plotMeanVar(d, meanvar=meanvar, show.tagwise.vars=TRUE, NBlind=TRUE) # Use previously saved object to speed up
## We could also estimate common/tagwise dispersions using the Cox-Reid methods with an appropriate design matrix

```

mglm

Fit Negative Binomial Generalized Linear Model to Multiple Response Vectors

Description

Fit the same log-link negative binomial or Poisson generalized linear model (GLM) to each row of a matrix of counts.

Usage

```

mglmLS(y, design, dispersion=0, offset=0, coef.start=NULL, tol=1e-5, maxit=50, trace=FALSE)
mglmOneGroup(y, dispersion=0, offset=0, maxit=50, trace=FALSE, tol=1e-6)
mglmOneWay(y, design=NULL, dispersion=0, offset=0, maxit=50, trace=FALSE)
mglmSimple(y, design, dispersion=0, offset=0, weights=NULL)
mglmLevenberg(y, design, dispersion=0, offset=0, coef.start=NULL, start.method="null",
              tol=1e-06, maxit=200)
deviances.function(dispersion)
designAsFactor(design)

```

Arguments

y	numeric matrix containing the negative binomial counts. Rows for tags and columns for libraries.
design	numeric matrix giving the design matrix of the GLM. Assumed to be full column rank.
dispersion	numeric scalar or vector giving the dispersion parameter for each GLM. Can be a scalar giving one value for all tags, or a vector of length equal to the number of tags giving tag-wise dispersions.
offset	numeric vector or matrix giving the offset that is to be included in the log-linear model predictor. Can be a scalar, a vector of length equal to the number of libraries, or a matrix of the same size as y.

weights	numeric vector or matrix of non-negative quantitative weights. Can be a vector of length equal to the number of libraries, or a matrix of the same size as <i>y</i> .
coef.start	numeric matrix of starting values for the linear model coefficients. Number of rows should agree with <i>y</i> and number of columns should agree with design.
start.method	method used to generate starting values when <code>coef.stat=NULL</code> . Possible values are "null" to start from the null model of equal expression levels or "y" to use the data as starting value for the mean.
tol	numeric scalar giving the convergence tolerance. For <code>mglmOneGroup</code> , convergence is judged successful when the step size falls below <code>tol</code> in absolute size.
maxit	scalar giving the maximum number of iterations for the Fisher scoring algorithm.
trace	logical, whether or not to information should be output at each iteration.

Details

The functions `mglmLS`, `mglmOneGroup` and `mglmSimple` all fit negative binomial generalized linear models, with the same design matrix but possibly different dispersions, offsets and weights, to a series of response vectors. `mglmLS` and `mglmOneGroup` are vectorized in R for fast execution, while `mglmSimple` simply makes tagwise calls to `glm.fit` in the stats package. The functions are all low-level functions in that they operate on atomic objects such as matrices. They are used as work-horses by higher-level functions in the edgeR package, especially by `glmFit`.

`mglmOneGroup` fit the null model, with intercept term only, to each response vector. In other words, it treats the libraries as belonging to one group. It implements Fisher scoring with a score-statistic stopping criterion for each tag. Excellent starting values are available for the null model, so this function seldom has any problems with convergence. It is used by other edgeR functions to compute the overall abundance for each tag.

`mglmLS` fits an arbitrary log-linear model to each response vector. It implements a vectorized approximate scoring algorithm with a likelihood derivative stopping criterion for each tag. A simple line search strategy is used to ensure that the residual deviance is reduced at each iteration. This function is the work-horse of other edgeR functions such as `glmFit` and `glmLRT`.

`mglmSimple` is not vectorized, and simply makes tag-wise calls to `glm.fit`. This has the advantage that it accesses all the usual information generated by `glm.fit`. Unfortunately, `glm.fit` does not always converge, and the tag-wise fitting is relatively slow.

`mglmLevenberg` implements a Levenberg-Marquardt modification of the `glm` scoring algorithm to prevent divergence, and is implemented in C++.

All these functions treat the dispersion parameter of the negative binomial distribution as a known input.

`deviances.function` simply chooses the appropriate deviance function to use given a scalar or vector of dispersion parameters. If the dispersion values are zero, then the Poisson deviance function is returned; if the dispersion values are positive, then the negative binomial deviance function is returned.

Value

`mglmOneGroup` produces a vector of length equal to the number of tags/genes (number of rows of *y*) providing the single coefficient from the GLM fit for each tag/gene. This can be interpreted as a measure of the 'average expression' level of the tag/gene.

`mglmLS` produces a list with the following components:

coefficients	matrix of estimated coefficients for the linear models
--------------	--

fitted.values	matrix of fitted values
fail	vector of indices of tags that fail the line search, in that the maximum number of step-halvings in exceeded
not.converged	vector of indices of tags that exceed the iteration limit before satisfying the convergence criterion

mglmSimple produces a list with the following components:

coefficients	matrix of estimated coefficients for the linear models
df.residual	vector of residual degrees of freedom for the linear models
deviance	vector of deviances for the linear models
design	matrix giving the experimental design that was used for each of the linear models
offset	scalar, vector or matrix of offset values used for the linear models
dispersion	scalar or vector of the dispersion values used for the linear model fits
weights	matrix of final weights for the observations from the linear model fits
fitted.values	matrix of fitted values
error	logical vector, did the fit fail?
converged	logical vector, did the fit converge?

deviances.function returns a function to calculate the deviance as appropriate for the given values of the dispersion.

designAsFactor returns a factor of length equal to nrow(design).

Author(s)

Davis McCarthy, Yunshun Chen, Gordon Smyth, Aaron Lun

References

McCarthy, DJ, Chen, Y, Smyth, GK (2012). Differential expression analysis of multifactor RNA-Seq experiments with respect to biological variation. *Nucleic Acids Research* 40, 4288-4297. <http://nar.oxfordjournals.org/content/40/10/4288>

See Also

[glmFit](#), for more object-orientated GLM modelling for DGE data.

Examples

```

y <- matrix(rnbinom(1000,mu=10,size=2),ncol=4)
lib.size <- colSums(y)
dispersion <- 0.1

logAveCPM <- mglmOneGroup(y, dispersion=dispersion, offset=log(lib.size))
summary(logAveCPM)

## Fit the NB GLM to the counts with a given design matrix
f1 <- factor(c(1,1,2,2))
f2 <- factor(c(1,2,1,2))
x <- model.matrix(~f1+f2)
fit <- mglmLS(y, x, dispersion=dispersion, offset=log(lib.size))
head(fit$coefficients)

```

movingAverageByCol *Moving Average Smoother of Matrix Columns*

Description

Apply a moving average smoother to the columns of a matrix.

Usage

```
movingAverageByCol(x, width=5, full.length=TRUE)
```

Arguments

x	numeric matrix
width	integer, width of window of rows to be averaged
full.length	logical value, should output have same number of rows as input?

Details

If full.length=TRUE, narrower windows are used at the start and end of each column to make a column of the same length as input. If FALSE, all values are averager of width input values, so the number of rows is less than input.

Value

Numeric matrix containing smoothed values. If full.length=TRUE, of same dimension as x. If full.length=FALSE, has width-1 fewer rows than x.

Author(s)

Gordon Smyth

Examples

```
x <- matrix(rpois(20,lambda=5),10,2)
movingAverageByCol(x,3)
```

normalizeChIPtoInput *Normalize ChIP-Seq Read Counts to Input and Test for Enrichment*

Description

Normalize ChIP-Seq read counts to input control values, then test for significant enrichment relative to the control.

Usage

```
normalizeChIPtoInput(input, response, dispersion=0.01, niter=6, loss="p", plot=FALSE, verbose=FALSE, ...)
calcNormOffsetsforChIP(input, response, dispersion=0.01, niter=6, loss="p", plot=FALSE, verbose=FALSE, .
```

Arguments

input	numeric vector of non-negative input values, not necessarily integer.
response	vector of non-negative integer counts of some ChIP-Seq mark for each gene or other genomic feature.
dispersion	negative binomial dispersion, must be positive.
niter	number of iterations.
loss	loss function to be used when fitting the response counts to the input: "p" for cumulative probabilities or "z" for z-value.
plot	if TRUE, a plot of the fit is produced.
verbose	if TRUE, working estimates from each iteration are output.
...	other arguments are passed to the plot function.

Details

normalizeChIPtoInput identifies significant enrichment for a ChIP-Seq mark relative to input values. The ChIP-Seq mark might be for example transcriptional factor binding or an epigenetic mark. The function works on the data from one sample. Replicate libraries are not explicitly accounted for, and would normally be pooled before using this function.

ChIP-Seq counts are assumed to be summarized by gene or similar genomic feature of interest.

This function makes the assumption that a non-negligible proportion of the genes, say 25% or more, are not truly marked by the ChIP-Seq feature of interest. Unmarked genes are further assumed to have counts at a background level proportional to the input. The function aligns the counts to the input so that the counts for the unmarked genes behave like a random sample. The function estimates the proportion of marked genes, and removes marked genes from the fitting process. For this purpose, marked genes are those with a Holm-adjusted mid-p-value less than 0.5.

The read counts are treated as negative binomial. The dispersion parameter is not estimated from the data; instead a reasonable value is assumed to be given.

calcNormOffsetsforChIP returns a numeric matrix of offsets, ready for linear modelling.

Value

normalizeChIPtoInput returns a list with components

p.value	numeric vector of p-values for enrichment.
scaling.factor	factor by which input is scaled to align with response counts for unmarked genes.
prop.enriched	proportion of marked genes, as internally estimated

calcNormOffsetsforChIP returns a numeric matrix of offsets.

Author(s)

Gordon Smyth

`plotBCV`*Plot Biological Coefficient of Variation*

Description

Plot genewise biological coefficient of variation (BCV) against gene abundance (in log2 counts per million).

Usage

```
plotBCV(object, xlab="logCPM", ylab="Biological coefficient of variation", pch=16, cex=0.2, ...)
```

Arguments

<code>object</code>	a DGEList object.
<code>xlab</code>	label for the x-axis.
<code>ylab</code>	label for the y-axis.
<code>pch</code>	the plotting symbol. See points for more details.
<code>cex</code>	plot symbol expansion factor. See points for more details.
<code>...</code>	any other arguments are passed to plot.

Details

The BCV is the square root of the negative binomial dispersion. This function displays the common, trended and tagwise BCV estimates.

Value

A plot is created on the current graphics device.

Author(s)

Davis McCarthy, Yunshun Chen, Gordon Smyth

Examples

```
BCV.true <- 0.1
y <- DGEList(matrix(rnbinom(6000, size = 1/BCV.true^2, mu = 10),1000,6))
y <- estimateCommonDisp(y)
y <- estimateTrendedDisp(y)
y <- estimateTagwiseDisp(y)
plotBCV(y)
```

plotExonUsage *Create a Plot of Exon Usage from Exon-Level Count Data*

Description

Create a plot of exon usage for a given gene by plotting the (un)transformed counts for each exon, coloured by experimental group.

Usage

```
plotExonUsage(y, geneID, group=NULL, transform="none", counts.per.million=TRUE, legend.coords=NULL)
```

Arguments

y	either a matrix of exon-level counts, a list containing a matrix of counts for each exon or a DGEList object with (at least) elements counts (table of counts summarized at the exon level) and samples (data frame containing information about experimental group, library size and normalization factor for the library size). Each row of y should represent one exon.
geneID	character string giving the name of the gene for which exon usage is to be plotted.
group	factor supplying the experimental group/condition to which each sample (column of y) belongs. If NULL (default) the function will try to extract if from y, which only works if y is a DGEList object.
transform	character, supplying the method of transformation to be applied to the exon counts, if any. Options are "none" (original counts are preserved), "sqrt" (square-root transformation) and "log2" (log2 transformation). Default is "none".
counts.per.million	logical, if TRUE then counts per million (as determined from total library sizes) will be plotted for each exon, if FALSE the raw read counts will be plotted. Using counts per million effectively normalizes for different read depth among the different samples, which can make the exon usage plots easier to interpret.
legend.coords	optional vector of length 2 giving the x- and y-coordinates of the legend on the plot. If NULL (default), the legend will be automatically placed near the top right corner of the plot.
...	optional further arguments to be passed on to plot.

Details

This function produces a simple plot for comparing exon usage between different experimental conditions for a given gene.

Value

plotExonUsage (invisibly) returns the transformed matrix of counts for the gene being plotted and produces a plot to the current device.

Author(s)

Davis McCarthy, Gordon Smyth

See Also

[spliceVariants](#) for methods to detect genes with evidence for alternative exon usage.

Examples

```
# generate exon counts from NB, create list object
y<-matrix(rnbinom(40,size=1,mu=10),nrow=10)
rownames(y) <- rep(c("gene.1", "gene.2"), each=5)
d<-DGEList(counts=y,group=rep(1:2,each=2))
plotExonUsage(d, "gene.1")
```

plotMDS.DGEList *Multidimensional scaling plot of digital gene expression profiles*

Description

Calculate distances between RNA-seq or DGE libraries, then produce a multidimensional scaling plot. Distances on the plot represent coefficient of variation of expression between samples for the top genes that best distinguish the samples.

Usage

```
## S3 method for class 'DGEList'
plotMDS(x, top=500, labels=colnames(x), col=NULL, cex=1, dim.plot=c(1,2),
        ndim=max(dim.plot), xlab=paste("Dimension",dim.plot[1]), ylab=paste("Dimension",dim.plot[2]), ...)
```

Arguments

x	numeric matrix or DGEList object.
top	number of top genes used to calculate pairwise distances.
labels	character vector of sample names or labels. If x has no column names, then defaults the index of the samples.
col	numeric or character vector of colors for the plotting characters. See text for possible values.
cex	numeric vector of plot symbol expansions. See text for possible values.
dim.plot	which two dimensions should be plotted, numeric vector of length two.
ndim	number of dimensions in which data is to be represented
xlab	title for the x-axis
ylab	title for the y-axis
...	any other arguments are passed to plot.

Details

This function is a variation on the usual multidimensional scaling (or principle coordinate) plot, in that a distance measure particularly appropriate for the digital gene expression (DGE) context is used. A set of top genes are chosen that have largest biological variation between the libraries (those with largest tagwise dispersion treating all libraries as one group). Then the distance between each pair of libraries (columns) is the biological coefficient of variation (square root of the common dispersion) between those two libraries alone, using the top genes.

If `x` is a `DGEList`, then the library sizes and normalization factors found in the object are used. If `x` is a matrix, then library sizes are computed from the column sums, but no other normalization is done.

The number top of top genes chosen for this exercise should roughly correspond to the number of differentially expressed genes with materially large fold-changes. The default setting of 500 genes is widely effective and suitable for routine use, but a smaller value might be chosen for when the samples are distinguished by a specific focused molecular pathway. Very large values (greater than 1000) are not usually so effective.

This function can be slow when there are many libraries.

Value

A plot is created on the current graphics device.

An object of class "MDS" is invisibly returned. This is a list containing the following components:

<code>distance.matrix</code>	numeric matrix of pairwise distances between columns of <code>x</code>
<code>cmdscale.out</code>	output from the function <code>cmdscale</code> given the distance matrix
<code>dim.plot</code>	dimensions plotted
<code>x</code>	x-coordinates of plotted points
<code>y</code>	y-coordinates of plotted points

Author(s)

Yunshun Chen, Mark Robinson and Gordon Smyth

See Also

[plotMDS](#), [cmdscale](#), [as.dist](#)

Examples

```
# Simulate DGE data for 1000 genes(tags) and 6 samples.
# Samples are in two groups
# First 200 genes are differentially expressed in second group

ngenes <- 1000
nlib <- 6
counts <- matrix(rnbinom(ngenes*nlib, size=1/10, mu=20),ngenes,nlib)
rownames(counts) <- paste("Gene",1:ngenes)
group <- gl(2,3,labels=c("Grp1","Grp2"))
counts[1:200,group=="Grp2"] <- counts[1:200,group=="Grp2"] + 10
y <- DGEList(counts,group=group)
y <- calcNormFactors(y)

# without labels, indexes of samples are plotted.
```

```
col <- as.numeric(group)
mds <- plotMDS(y, top=200, col=col)

# or labels can be provided, here group indicators:
plotMDS(mds, col=col, labels=group)
```

plotSmear	<i>Plots log-Fold Change versus log-Concentration (or, M versus A) for Count Data</i>
-----------	---

Description

Both of these functions plot the log-fold change (i.e. the log of the ratio of expression levels for each tag between two experimental groups) against the log-concentration (i.e. the overall average expression level for each tag across the two groups). To represent counts that were low (e.g. zero in 1 library and non-zero in the other) in one of the two conditions, a 'smear' of points at low A value is presented in plotSmear.

Usage

```
plotSmear(object, pair=NULL, de.tags=NULL, xlab="logCPM", ylab="logFC", pch=19,
          cex=0.2, smearWidth=0.5, panel.first=grid(), smooth.scatter=FALSE, lowess=FALSE, ...)
```

Arguments

object	DGEList, DGEEexact or DGELRT object containing data to produce an MA-plot.
pair	pair of experimental conditions to plot (if NULL, the first two conditions are used)
de.tags	rownames for tags identified as being differentially expressed; use exactTest to identify DE genes
xlab	x-label of plot
ylab	y-label of plot
pch	scalar or vector giving the character(s) to be used in the plot; default value of 19 gives a round point.
cex	character expansion factor, numerical value giving the amount by which plotting text and symbols should be magnified relative to the default; default cex=0.2 to make the plotted points smaller
smearWidth	width of the smear
panel.first	an expression to be evaluated after the plot axes are set up but before any plotting takes place; the default grid() draws a background grid to aid interpretation of the plot
smooth.scatter	logical, whether to produce a 'smooth scatter' plot using the KernSmooth::smoothScatter function or just a regular scatter plot; default is FALSE, i.e. produce a regular scatter plot
lowess	logical, indicating whether or not to add a lowess curve to the MA-plot to give an indication of any trend in the log-fold change with log-concentration
...	further arguments passed on to plot

Details

plotSmear is a more sophisticated and superior way to produce an 'MA plot'. plotSmear resolves the problem of plotting tags that have a total count of zero for one of the groups by adding the 'smear' of points at low A value. The points to be smeared are identified as being equal to the minimum estimated concentration in one of the two groups. The smear is created by using random uniform numbers of width smearWidth to the left of the minimum A. plotSmear also allows easy highlighting of differentially expressed (DE) tags.

Value

A plot to the current device

Author(s)

Mark Robinson, Davis McCarthy

See Also

[maPlot](#)

Examples

```
y <- matrix(rnbinom(10000,mu=5,size=2),ncol=4)
d <- DGEList(counts=y, group=rep(1:2,each=2), lib.size=colSums(y))
rownames(d$counts) <- paste("tag",1:nrow(d$counts),sep=".")
d <- estimateCommonDisp(d)
plotSmear(d)

# find differential expression
de <- exactTest(d)

# highlighting the top 500 most DE tags
de.tags <- rownames(topTags(de, n=500)$table)
plotSmear(d, de.tags=de.tags)
```

predFC

Predictive log fold changes

Description

Computes estimated coefficients for a generalised linear model with log-fold-changes shrunk towards zero.

Usage

```
## S3 method for class 'DGEList'
predFC(y, design=NULL, prior.count.total=0.5, offset=NULL, dispersion=NULL)
## Default S3 method:
predFC(y, design=NULL, prior.count.total=0.5, offset=log(colSums(y)), dispersion=0)
```

Arguments

<code>y</code>	a matrix of counts or a <code>DGEList</code> object
<code>design</code>	the design matrix for the experiment
<code>prior.count.total</code>	the total prior number of counts to be added to the data. Larger values produce more shrinkage.
<code>offset</code>	numeric vector or matrix giving the offset in the log-linear model predictor, as for <code>glmFit</code> . Usually equal to log library sizes.
<code>dispersion</code>	the negative binomial dispersion

Details

This function adds `prior.count.total` counts to each row of `y`. The counts are added in such a way that any log-fold-change that was zero prior to augmentation remains zero and non-zero log-fold-changes are shrunk towards zero.

The prior counts can be viewed as equivalent to a prior belief that the log-fold changes are small, and the output can be viewed as posterior log-fold-changes from this Bayesian viewpoint. The output coefficients are called *predictive* log fold-changes because, depending on the prior, they may be a better prediction of the true log fold-changes than the raw estimates.

This is done small count is added to each library in proportion to the library sizes. A larger amount is added to counts from larger libraries, so that any log-fold-change that was zero prior to augmentation remains zero. The specific estimates the predictive or posterior log-fold-changes for count data. If there are 2 groups in the experiment, $n=2$ for each group, the total prior count is 1, and the library sizes are equal, then in effect 0.5 of a count is added to each group, or 0.25 to each library. This prior count is the same for all genes or tags in the data, with the result that genes with low counts will be dampened more severely and genes with a large number of counts in each library will hardly be affected by the addition of a small count to each group.

In order to get the predictive log-fold-changes, a generalised linear model is fitted to the augmented data, and the coefficients outputted in the form of a matrix.

If `offset=NULL`, the offset will be taken from the `DGEList` object or computed from the column sums.

If `dispersion=NULL`, the dispersion used for the glm will be dependent on what is in the `DGEList` object; it is prioritised in the following manner: tagwise, trended, common and finally if no dispersion estimate is found it will set the dispersion to 0.

If `design=NULL`, then the function returns a matrix of the same size as `y` containing log₂ counts-per-million, with zero values for the counts avoided. This equivalent to choosing `design` to be the identity matrix with the same number of columns as `y`.

Value

Numeric matrix of linear model coefficients (if `design` is given) or logCPM (if `design=NULL`) on the log₂ scale.

Author(s)

Belinda Phipson, Gordon Smyth

See Also

[glmFit](#), [exactTest](#)

Examples

```
# generate counts for a two group experiment with n=2 in each group and 100 genes
dispersion <- 0.1
y <- matrix(rnbinom(400,size=1/dispersion,mu=4),nrow=100)
y <- DGEList(y,group=c(1,1,2,2))
design <- model.matrix(~group, data=y$samples)

#estimate the predictive log fold changes
predlfc<-predFC(y,design,dispersion=dispersion,prior.count=4)
logfc <- predFC(y,design,dispersion=dispersion,prior.count=0)
logfc.truncated <- pmax(pmin(logfc,100),-100)

#plot predFC's vs logFC's
plot(predlfc[,2],logfc.truncated[,2],xlab="Predictive log fold changes",ylab="Raw log fold changes")
abline(a=0,b=1)
```

q2qnbinom

*Quantile to Quantile Mapping between Negative-Binomial Distributions***Description**

Interpolated quantile to quantile mapping between negative-binomial distributions with the same dispersion but different means. The Poisson distribution is a special case.

Usage

```
q2qpois(x, input.mean, output.mean)
q2qnbinom(x, input.mean, output.mean, dispersion=0)
```

Arguments

x	numeric matrix of counts.
input.mean	numeric matrix of population means for x. If a vector, then of the same length as nrow(x).
output.mean	numeric matrix of population means for the output values. If a vector, then of the same length as nrow(x).
dispersion	numeric scalar, vector or matrix giving negative binomial dispersion values.

Details

This function finds the quantile with the same left and right tail probabilities relative to the output mean as x has relative to the input mean. q2qpois is equivalent to q2qnbinom with dispersion=0.

In principle, q2qnbinom gives similar results to calling pnbinom followed by qnbinom as in the example below. However this function avoids infinite values arising from rounding errors and does appropriate interpolation to return continuous values.

q2qnbinom is called by [equalizeLibSizes](#) to perform quantile-to-quantile normalization.

Value

numeric matrix of same dimensions as x, with output.mean as the new nominal population mean.

Author(s)

Gordon Smyth

See Also[equalizeLibSizes](#)**Examples**

```
x <- 15
input.mean <- 10
output.mean <- 20
dispersion <- 0.1
q2qnbinom(x,input.mean,output.mean,dispersion)

# Similar in principle:
qnbinom(pnbinom(x,mu=input.mean,size=1/dispersion),mu=output.mean,size=1/dispersion)
```

readDGE

*Read and Merge a Set of Files Containing DGE Data***Description**

Reads and merges a set of text files containing digital gene expression data.

Usage

```
readDGE(files, path=NULL, columns=c(1,2), group=NULL, labels=NULL, ...)
```

Arguments

files	character vector of filenames, or alternatively a data.frame with a column containing the file names of the files containing the libraries of counts and, optionally, columns containing the group to which each library belongs, descriptions of the other samples and other information.
path	character string giving the directory containing the files. The default is the current working directory.
columns	numeric vector stating which two columns contain the tag names and counts, respectively
group	vector, or preferably a factor, indicating the experimental group to which each library belongs. If group is not NULL, then this argument overrides any group information included in the files argument.
labels	character vector giving short names to associate with the libraries. Defaults to the file names.
...	other are passed to read.delim

Details

Each file is assumed to contain digital gene expression data for one sample (or library), with transcript identifiers in the first column and counts in the second column. Transcript identifiers are assumed to be unique and not repeated in any one file. By default, the files are assumed to be tab-delimited and to contain column headings. The function forms the union of all transcripts and creates one big table with zeros where necessary.

Value

DGEList object

Author(s)

Mark Robinson and Gordon Smyth

See Also

[DGEList](#) provides more information about the DGEList class and the function DGEList, which can also be used to construct a DGEList object, if readDGE is not required to read in and construct a table of counts from separate files.

Examples

```
# Read all .txt files from current working directory

## Not run: files <- dir(pattern="*\\.txt$")
RG <- readDGE(files)
## End(Not run)
```

spliceVariants

Identify Genes with Splice Variants

Description

Identify genes exhibiting evidence for splice variants (alternative exon usage/transcript isoforms) from exon-level count data using negative binomial generalized linear models.

Usage

```
spliceVariants(y, geneID, dispersion=NULL, group=NULL, estimate.genewise.disp=TRUE, trace=FALSE)
```

Arguments

y	either a matrix of exon-level counts or a DGEList object with (at least) elements counts (table of counts summarized at the exon level) and samples (data frame containing information about experimental group, library size and normalization factor for the library size). Each row of y should represent one exon.
geneID	vector of length equal to the number of rows of y, which provides the gene identifier for each exon in y. These identifiers are used to group the relevant exons into genes for the gene-level analysis of splice variation.

dispersion	scalar (in future a vector will also be allowed) supplying the negative binomial dispersion parameter to be used in the negative binomial generalized linear model.
group	factor supplying the experimental group/condition to which each sample (column of y) belongs. If NULL (default) the function will try to extract if from y, which only works if y is a DGEList object.
estimate.genewise.disp	logical, should genewise dispersions (as opposed to a common dispersion value) be computed if the dispersion argument is NULL?
trace	logical, whether or not verbose comments should be printed as function is run. Default is FALSE.

Details

This function can be used to identify genes showing evidence of splice variation (i.e. alternative splicing, alternative exon usage, transcript isoforms). A negative binomial generalized linear model is used to assess evidence, for each gene, given the counts for the exons for each gene, by fitting a model with an interaction between exon and experimental group and comparing this model (using a likelihood ratio test) to a null model which does not contain the interaction. Genes that show significant evidence for an interaction between exon and experimental group by definition show evidence for splice variation, as this indicates that the observed differences between the exon counts between the different experimental groups cannot be explained by consistent differential expression of the gene across all exons. The function `topTags` can be used to display the results of `spliceVariants` with genes ranked by evidence for splice variation.

Value

`spliceVariants` returns a `DGEEexact` object, which contains a table of results for the test of differential splicing between experimental groups (alternative exon usage), a data frame containing the gene identifiers for which results were obtained and the dispersion estimate(s) used in the statistical models and testing.

Author(s)

Davis McCarthy, Gordon Smyth

See Also

[estimateExonGenewiseDisp](#) for more information about estimating genewise dispersion values from exon-level counts. [DGEList](#) for more information about the `DGEList` class. [topTags](#) for more information on displaying ranked results from `spliceVariants`. [estimateCommonDisp](#) and related functions for estimating the dispersion parameter for the negative binomial model.

Examples

```
# generate exon counts from NB, create list object
y<-matrix(rnbinom(40,size=1,mu=10),nrow=10)
d<-DGEList(counts=y,group=rep(1:2,each=2))
genes <- rep(c("gene.1","gene.2"), each=5)
disp <- 0.2
spliceVariants(d, genes, disp)
```

splitIntoGroups	<i>Split the Counts or Pseudocounts from a DGEList Object According To Group</i>
-----------------	--

Description

Split the counts from a DGEList object according to group, creating a list where each element consists of a numeric matrix of counts for a particular experimental group. Given a pair of groups, split pseudocounts for these groups, creating a list where each element is a matrix of pseudocounts for a particular group.

Usage

```
splitIntoGroups(object)
splitIntoGroupsPseudo(pseudo, group, pair)
```

Arguments

object	DGEList, object containing (at least) the elements counts (table of raw counts), group (factor indicating group) and lib.size (numeric vector of library sizes)
pseudo	numeric matrix of quantile-adjusted pseudocounts to be split
group	factor indicating group to which libraries/samples (i.e. columns of pseudo belong; must be same length as ncol(pseudo))
pair	vector of length two stating pair of groups to be split for the pseudocounts

Value

splitIntoGroups outputs a list in which each element is a matrix of count counts for an individual group. splitIntoGroupsPseudo outputs a list with two elements, in which each element is a numeric matrix of (pseudo-)count data for one of the groups specified.

Author(s)

Davis McCarthy

Examples

```
# generate raw counts from NB, create list object
y<-matrix(rnbinom(80,size=1,mu=10),nrow=20)
d<-DGEList(counts=y,group=rep(1:2,each=2),lib.size=rep(c(1000:1001),2))
rownames(d$counts)<-paste("tagno",1:nrow(d$counts),sep=".")
z1<-splitIntoGroups(d)

z2<-splitIntoGroupsPseudo(d$counts,d$group,pair=c(1,2))
```

subsetting

*Subset DGEList, DGEGLM, DGEEExact and DGELRT Objects***Description**

Extract a subset of a DGEList, DGEGLM, DGEEExact or DGELRT object.

Usage

```
## S3 method for class 'DGEList'
object[i, j, ...]
## S3 method for class 'DGEGLM'
object[i, j, ...]
## S3 method for class 'DGEEExact'
object[i, j, ...]
## S3 method for class 'DGELRT'
object[i, j, ...]
```

Arguments

object	object of class DGEList, DGEGLM, DGEEExact or DGELRT, respectively
i,j	elements to extract. i subsets the tags or genes while j subsets the libraries. Note, columns of DGEGLM, DGEEExact and DGELRT objects cannot be subsetted.
...	not used

Details

i,j may take any values acceptable for the matrix components of object of class DGEList. See the [Extract](#) help entry for more details on subsetting matrices. For DGEGLM, DGEEExact and DGELRT objects, only rows (i.e. i) may be subsetted.

Value

An object of class DGEList, DGEGLM, DGEEExact or DGELRT as appropriate, holding data from the specified subset of tags/genes and libraries.

Author(s)

Davis McCarthy, Gordon Smyth

See Also

[Extract](#) in the base package.

Examples

```
d <- matrix(rnbinom(16,size=1,mu=10),4,4)
rownames(d) <- c("a","b","c","d")
colnames(d) <- c("A1","A2","B1","B2")
d <- DGEList(counts=d,group=factor(c("A","A","B","B")))
d[1:2,]
d[1:2,2]
```

```
d[,2]
d <- estimateCommonDisp(d)
results <- exactTest(d)
results[1:2,]
# NB: cannot subset columns for DGEEExact objects
```

systematicSubset	<i>Take a systematic subset of indices.</i>
------------------	---

Description

Take a systematic subset of indices stratified by a ranking variable.

Usage

```
systematicSubset(n, order.by)
```

Arguments

n	integer giving the size of the subset.
order.by	numeric vector of the values by which the indices are ordered.

Value

systematicSubset returns a vector of size n.

Author(s)

Gordon Smyth

See Also

[order](#)

Examples

```
y <- rnorm(100, 1, 1)
systematicSubset(20, y)
```

`thinCounts`*Binomial or Multinomial Thinning of Counts*

Description

Reduce the size of Poisson-like counts by binomial thinning.

Usage

```
thinCounts(x, prob=NULL, target.size=min(colSums(x)))
```

Arguments

<code>x</code>	numeric vector or array of non-negative integers.
<code>prob</code>	numeric scalar or vector of same length as <code>x</code> , the expected proportion of the events to keep.
<code>target.size</code>	integer scale or vector of the same length as <code>NCOL{x}</code> , the desired total column counts. Must be not greater than column sum of <code>x</code> . Ignored if <code>prob</code> is not <code>NULL</code> .

Details

If `prob` is not `NULL`, then this function calls `rbinom` with `size=x` and `prob=prob` to generate the new counts. This is classic binomial thinning. The new column sums are random, with expected values determined by `prob`.

If `prob` is `NULL`, then this function does multinomial thinning of the counts to achieve specified column totals. The default behavior is to thin the columns to have the same column sum, equal to the smallest column sum of `x`.

If the elements of `x` are Poisson, then binomial thinning produces new Poisson random variables with expected values reduced by factor `prob`. If the elements of each column of `x` are multinomial, then multinomial thinning produces a new multinomial observation with a reduced sum.

Value

A vector or array of the same dimensions as `x`, with thinned counts.

Author(s)

Gordon Smyth

Examples

```
x <- rpois(10,lambda=10)
thinCounts(x,prob=0.5)
```

topTags

*Table of the Top Differentially Expressed Tags***Description**

Extracts the top DE tags in a data frame for a given pair of groups, ranked by p-value or absolute log-fold change.

Usage

```
topTags(object, n=10, adjust.method="BH", sort.by="p.value")
```

Arguments

object	a DGEEexact object (output from exactTest) or a DGELRT object (output from glmLRT), containing the (at least) the elements table: a data frame containing the log-concentration (i.e. expression level), the log-fold change in expression between the two groups/conditions and the p-value for differential expression, for each tag. If it is a DGEEexact object, then topTags will also use the comparison element, which is a vector giving the two experimental groups/conditions being compared. The object may contain other elements that are not used by topTags.
n	scalar, number of tags to display/return
adjust.method	character string stating the method used to adjust p-values for multiple testing, passed on to p.adjust
sort.by	character string, indicating whether tags should be sorted by p-value ("p.value") or absolute log-fold change ("logFC"); default is to sort by p-value.

Value

an object of class TopTags containing the following elements for the top n most differentially expressed tags as determined by sort.by.

table	a data frame containing the elements logConc, the log-average concentration/abundance for each tag in the two groups being compared, logFC, the log-abundance ratio, i.e. fold change, for each tag in the two groups being compared, p.value, exact p-value for differential expression using the NB model, adj.p.val, the p-value adjusted for multiple testing as found using p.adjust using the method specified
comparison	a vector giving the names of the two groups being compared

There is a show method for this class.

Author(s)

Mark Robinson, Davis McCarthy, Gordon Smyth

References

Robinson MD, Smyth GK (2008). Small-sample estimation of negative binomial dispersion, with applications to SAGE data. *Biostatistics* 9, 321-332.

Robinson MD, Smyth GK (2007). Moderated statistical tests for assessing differences in tag abundance. *Bioinformatics* 23, 2881-2887.

See Also

[exactTest](#), [glmLRT](#), [p.adjust](#).

Analogous to [topTable](#) in the limma package.

Examples

```
# generate raw counts from NB, create list object
y <- matrix(rnbinom(80,size=1,mu=10),nrow=20)
d <- DGEList(counts=y,group=rep(1:2,each=2),lib.size=rep(c(1000:1001),2))
rownames(d$counts) <- paste("tag",1:nrow(d$counts),sep=".")

# estimate common dispersion and find differences in expression
# here we demonstrate the 'exact' methods, but the use of topTags is
# the same for a GLM analysis
d <- estimateCommonDisp(d)
de <- exactTest(d)

# look at top 10
topTags(de)
# Can specify how many tags to view
tp <- topTags(de, n=15)
# Here we view top 15
tp
# Or order by fold change instead
topTags(de,sort.by="logFC")
```

Tu102

Raw Data for Several SAGE Libraries from the Zhang 1997 Science Paper.

Description

SAGE dataset for 2 tumour samples, 2 normal samples.

Usage

```
data(Tu102)
```

Format

Data frames with 22713, 18794, 16270 and 17703 observations (for Tu102, Tu98, NC2, NC1, respectively) on the following 2 variables.

Tag_Sequence a character vector

Count a numeric vector

Source

Zhang et al. (1997) Gene Expression Profiles in Normal and Cancer Cells. *Science*, 276, 1268-72.

 weightedCondLogLikDerDelta

Weighted Conditional Log-Likelihood in Terms of Delta

Description

Weighted conditional log-likelihood parameterized in terms of delta ($\phi / (\phi+1)$) for a given tag/gene - maximized to find the smoothed (moderated) estimate of the dispersion parameter

Usage

```
weightedCondLogLikDerDelta(y, delta, tag, prior.n=10, ntags=nrow(y[[1]]), der=0)
```

Arguments

y	list with elements comprising the matrices of count data (or pseudocounts) for the different groups
delta	delta ($\phi / (\phi+1)$) parameter of negative binomial
tag	tag/gene at which the weighted conditional log-likelihood is evaluated
prior.n	smoothing parameter that indicates the weight to put on the common likelihood compared to the individual tag's likelihood; default 10 means that the common likelihood is given 10 times the weight of the individual tag/gene's likelihood in the estimation of the tag/genewise dispersion
ntags	numeric scalar number of tags/genes in the dataset to be analysed
der	derivative, either 0 (the function), 1 (first derivative) or 2 (second derivative)

Details

This function computes the weighted conditional log-likelihood for a given tag, parameterized in terms of delta. The value of delta that maximizes the weighted conditional log-likelihood is converted back to the phi scale, and this value is the estimate of the smoothed (moderated) dispersion parameter for that particular tag. The delta scale for convenience (delta is bounded between 0 and 1).

Value

numeric scalar of function/derivative evaluated for the given tag/gene and delta

Author(s)

Mark Robinson, Davis McCarthy

Examples

```
counts<-matrix(rnbinom(20,size=1,mu=10),nrow=5)
d<-DGEList(counts=counts,group=rep(1:2,each=2),lib.size=rep(c(1000:1001),2))
y<-splitIntoGroups(d)
ll1<-weightedCondLogLikDerDelta(y,delta=0.5,tag=1,prior.n=10,der=0)
ll2<-weightedCondLogLikDerDelta(y,delta=0.5,tag=1,prior.n=10,der=1)
```

zscoreNBinom	<i>Z-score Equivalents of Negative Binomial Deviate</i>
--------------	---

Description

Compute z-score equivalents of negative binomial random deviates.

Usage

```
zscoreNBinom(q, size, mu)
```

Arguments

q	numeric vector or matrix giving negative binomial random values.
size	negative binomial size parameter (>0).
mu	mean of negative binomial distribution (>0).

Details

This function computes the mid-p value of q, then converts to the standard normal deviate with the same cumulative probability distribution value.

Care is taken to do the computations accurately in both tails of the distributions.

Value

Numeric vector or matrix giving equivalent deviates from a standard normal distribution.

Author(s)

Gordon Smyth

See Also

[pnbinom](#), [qnorm](#) in the stats package.

Examples

```
zscoreNBinom(c(0,10,100), mu=10, size=1/10)
```

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