Package ‘LMGene’
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\texttt{R} topics documented:

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**Genediff**

**Description**

Computes two sets of p-values per gene or probe via gene-by-gene ANOVA, using both the gene-specific MSE and the posterior MSE for each term in the ANOVA. P-values are not adjusted for multiple testing.

Assumes a fixed effects model and that the correct denominator for all comparisons is the MSE.

**Usage**

```r
genediff(eS, model = NULL, method = c("MLE", "MOM", "MOMlog"),
          verbose = TRUE)
```

**Arguments**

- `eS` An ExpressionSet object. Any transformation and normalization of `exprs(eS)` should be conducted prior to use in `genediff`.
- `model` Model used for comparison; see details and `LMGene`.
- `method` Method by which posterior p-values are calculated. Default "MLE".
- `verbose` If TRUE, the prior degrees of freedom and mean reciprocal precision are printed. See details.

**Details**

The argument `eS` must be an ExpressionSet object from the Biobase package. If you have data in a matrix and information about experimental design factors, then you can use `neweS` to convert the data into an ExpressionSet object. Please see `neweS` for more detail.

The `model` argument is an optional character string, constructed like the right-hand side of a formula for `lm`. It specifies which of the variables in the ExpressionSet will be used in the model and whether interaction terms will be included. If `model=NULL`, it uses all variables from the ExpressionSet without interactions. Be careful of using interaction terms with factors; this often leads to overfitting, which will yield an error.

The method argument specifies how the adjusted MSE and degrees of freedom should be calculated for use in computation of the posterior p-values:

"MLE" Default. Calculate adjusted MSE and degrees of freedom by maximum likelihood estimation, as described in Wright and Simon (2003).

"MOM" Calculate adjusted MSE and degrees of freedom by method of moments, as described in Rocke (2003).

"MOMlog" Calculate adjusted MSE and degrees of freedom by method of moments on log scale, as described in Smyth (2004). Uses functions `fitFdist` and `trigammaInv` from the package `limma`. Note that the method of Smyth (2004) is used here to calculate the posterior MSE, but not to directly calculate the posterior p-values.
All three methods assume that the gene-specific MSE’s follow a gamma distribution with mean tau. (NB: Notation and parameterization vary somewhat between each of the source papers.) The mean of the gamma distribution, tau, is modeled with an inverse gamma prior with hyperparameters alpha and beta. Empirical Bayes methods are used to estimate the prior hyperparameters, either by maximum likelihood, method of moments, or method of moments on the log scale. The "posterior MSE" is the posterior mean of the variances given the observed gene-specific MSE’s.

If verbose = TRUE, the function prints the estimated prior degrees of freedom, which equals twice the prior shape parameter alpha, and the estimated prior mean reciprocal precision, or 1/(alpha*beta).

All p-values are calculated from fixed-effects ANOVA F statistics, using either the gene-specific MSE or the posterior MSE as the denominator.

**Value**

A list with components:

- **Gene.Specific** A matrix of p-values calculated using the gene-specific MSE, with one row for each gene/probe and one column for each factor
- **Posterior** A matrix of p-values calculated using the posterior MSE, with one row for each gene/probe and one column for each factor

**Author(s)**

David Rocke, Geun-Cheol Lee, and Blythe Durbin-Johnson

**References**


**See Also**

- `LMGene`, `rowaov`, `neweS`

**Examples**

```r
library(Biobase)
library(LMGene)
# data
data(sample.mat)
data(vlist)
raw.eS <- neweS(sample.mat, vlist)
# glog transform data
trans.eS <- transeS(raw.eS, lambda = 727, alpha = 56)
```
# calculate p-values
pvlist <- genediff(trans.eS)
pvlist$Posterior[1:5,]

---

**glog**  
*Generalized log transformation function*

**Description**

This function transforms the input values by the generalized log function.

**Usage**

```r
glog(y, lambda)
```

**Arguments**

- `y`: A data matrix
- `lambda`: Transformation parameter

**Details**

The glog transformation of a variable `y` is defined as \( \log(y + \sqrt{y^2 + \lambda}) \). Using \( \lambda = 0 \) corresponds to the log transformation, up to a scale factor of 2. (Other, equivalent expressions exist for the glog transformation. See Durbin et al. (2002) and Huber et al. (2002) for further details.)

The input matrix `y` may be modified prior to transformation by subtracting a constant or vector ("alpha"). The parameters `lambda` and `alpha` may be estimated from `tranest`.

**Value**

`yt` A matrix of glog-transformed values

**Author(s)**

David Rocke and Geun-Cheol Lee

**References**


[http://dmrocke.ucdavis.edu](http://dmrocke.ucdavis.edu)

**See Also**

`tranest`, `transeS`
Examples

```r
#library
library(Biobase)
library(LMGene)

#data
data(sample.mat)
sample.mat[1:5,1:4]

GloggedSmpd<-glog(sample.mat-50,500)
GloggedSmpd[1:5,1:4]
```

LMGene

**LMGene main function**

Description

LMGene calls function `genediff` to calculate the unadjusted gene-specific and posterior p-values of all genes and then calculates the FDR-adjusted p-values of all genes. Significant genes for each factor in `model` (based on either the gene-specific or posterior FDR-adjusted p-values) are output.

Usage

```r
LMGene(eS, model = NULL, level = 0.05, posterior = FALSE,
method = c("MLE", "MOM", "MOMlog"))
```

Arguments

- **eS** An `ExpressionSet` object. Any transformation and normalization of `exprs(eS)` should be conducted prior to use in LMGene.
- **model** Specifies model to be used. Default is to use all variables from eS without interactions. See details.
- **level** Significance level
- **posterior** If TRUE, the posterior FDR-adjusted p-values are used in listing significant genes for each factor. Default is to use gene-specific FDR-adjusted p-values.
- **method** Method by which the posterior p-values are calculated. Default is "MLE".

Details

If you have data in a matrix and information about experimental design factors, then you can use `neweS` to convert the data into an `ExpressionSet` object. Please see `neweS` for more detail.

The `level` argument indicates the False Discovery Rate, e.g. level=0.05 means a 5 percent FDR.

The `model` argument is an optional character string, constructed like the right-hand side of a formula for `lm`. It specifies which of the variables in the `ExpressionSet` will be used in the model and whether interaction terms will be included. If `model=NULL`, it uses all variables from the `ExpressionSet` without interactions. Be careful of using interaction terms with factors; this often leads to overfitting, which will yield an error.

See `genediff` for details of method.
Value

`lmres`  
A list with one component for each factor in `model`. Each component consists of a character vector with one element per significant gene. If no genes are significant for a given factor, the component for that factor is set to "No significant genes".

Author(s)

David Rocke and Geun-Cheol Lee

References


[http://dmrocke.ucdavis.edu](http://dmrocke.ucdavis.edu)

See Also

genediff, neweS

Examples

```r
library(Biobase)
library(LMGene)

# data
data(sample.mat)
data(vlist)

raw.eS <- neweS(sample.mat, vlist)

# glog transform data
transeS <- transeS(raw.eS, lambda = 727, alpha = 56)

# Identify significant genes, using an FDR of 1 percent
LMGene(trans.eS, level = 0.01)
```

Description

Lowess normalization function

Usage

```r
lnorm(mat1, span = 0.1)
```

Arguments

- `mat1`: A data matrix to be normalized
- `span`: Lowess smoother span. Larger values give more smoothness.
lnormeS

Details

mat1 must be a p by n matrix, where p is the number of genes and n is the number of arrays or samples.

Value

matnorm1 Normalized matrix

Author(s)

David Rocke and Geun-Cheol Lee

References

http://dmrocke.ucdavis.edu

See Also

lnormeS, norm

Examples

library(Biobase)
library(LMGene)

# data
data(sample.mat)
data(vlist)

raw.eS <- neweS(sample.mat, vlist)

# glog transform data
trans.eS <- transeS(raw.eS, lambda = 727, alpha = 56)

# normalize
normed.exprs <- lnorm(exprs(trans.eS))

lnormeS

Function to apply lowess normalization to an expression set.

Description

Like lnorm, but applies to and returns an ExpressionSet or AffyBatch object instead of a matrix.

Usage

lnormeS(eS, span=0.1)

Arguments

eS An ExpressionSet or AffyBatch object
span Smoothing parameter for lowess. Larger values correspond to more smoothness.
Value

Returns an ExpressionSet with exprs(eS) normalized by \texttt{lnorm}.

Author(s)

John Tillinghast, Blythe Durbin-Johnson

References

\url{http://dmrocke.ucdavis.edu}

See Also

\texttt{lnorm}, \texttt{norm}

Examples

\begin{verbatim}
library(LMGene)
library(Biobase)

data(sample.eS)

# glog transform expression set
trsample.eS <- transeS(sample.eS, 667, 65)

# normalize expression set
normtrsample.eS <- lnormeS(trsample.eS)
\end{verbatim}

\begin{verbatim}
neweS(mat, vlist, vlabel = as.list(names(vlist)))
\end{verbatim}

Arguments

\begin{itemize}
\item \texttt{mat} A data matrix to be converted.
\item \texttt{vlist} A list, each component of which describes a factor in the experimental design.
\item \texttt{vlabel} A list of labels for each component of \texttt{vlist}.
\end{itemize}

Details

Each element of a component of \texttt{vlist} corresponds to a column of \texttt{mat}. See \texttt{vlist} for an example.

Value

\begin{itemize}
\item \texttt{eset} An ExpressionSet object.
\end{itemize}
norm

Author(s)
David Rocke and Geun-Cheol Lee

References
http://dmrocke.ucdavis.edu

See Also
vlist

Examples
library(Biobase)
library(LMGene)

#data
data(sample.mat)
data(vlist)

Smpdt <- neweS(sample.mat,vlist)
data(sample.es)
identical(exprs(sample.es), exprs(Smpdt))
identical(pData(sample.es), pData(Smpdt))

---

norm | Normalization function

Description
This function normalizes a matrix by subtracting the column (sample) mean from each element and adding the grand mean.

Usage
norm(mat1)

Arguments
mat1 A matrix to be normalized

Value
matnorm Normalized matrix

Author(s)
David Rocke and Geun-Cheol Lee

References
http://dmrocke.ucdavis.edu
plotMeanSD

See Also

lnorm

Examples

library(Biobase)
library(LMGene)

#data
data(sample.mat)
data(vlist)

raw.eS <- neweS(sample.mat, vlist)

# glog transform data
trans.eS <- transeS(raw.eS, lambda = 727, alpha = 56)

# normalize
normed.exprs <- norm(exprs(trans.eS))

plotMeanSD

Plotting function for gene means and standard deviations

Description

Plots the row standard deviation of a matrix of expression data against the row mean, or the rank of the row mean.

Usage

plotMeanSD(indata, by.rank = TRUE, line = FALSE, ymax = NULL)

Arguments

indata An object of class matrix, data.frame, ExpressionSet, or AffyBatch
by.rank If TRUE, the row standard deviations are plotted against the ranks of the row means. Otherwise, the row standard deviations are plotted against the row means themselves.
line If TRUE, a lowess smoother line is drawn on the plot.
ymax The upper limit for the plot y-axis. If missing, axis limits are generated automatically by plot.

Details

Generates a scatter plot of the row standard deviations of a matrix of expression data against the row means or ranks of the row means.

Value

NULL
Author(s)

Rachel Chen and Blythe Durbin-Johnson

Examples

library(LMGene)
library(Biobase)
data(sample.eS)
# transform data
trans.eS <- transeS(sample.eS, lambda = 727, alpha = 56)

# plot SD against rank of mean
plotMeanSD(trans.eS, line = TRUE)
plotMeanSD(sample.eS, line = TRUE, ymax = 1000)

Description

Converts an ExpressionSet or AffyBatch object with one row of expression data per probeset into an ExpressionSet or AffyBatch object with one row per probe.

Usage

psmeans(eS, ind)

Arguments

eS An ExpressionSet or AffyBatch object
ind A vector used to indicate which probes go into which probesets.

Details

Each entry of ind corresponds to one probe and tells the number of the probeset it belongs to. See tranestAffyProbeLevel and sample.ind for examples.

Value

Returns an ExpressionSet or AffyBatch object with the expression matrix rows corresponding to probesets instead of individual probes. Elements of the returned ExpressionSet or AffyBatch object are means over each probeset.

Author(s)

John Tillinghast

See Also

tranestAffyProbeLevel, sample.ind
pvadjust

Examples

library(LMGene)
library(Biobase)
data(sample.eS)
data(sample.ind)

# glog transform data
trs.eS <- transeS(sample.eS, 667, 65)

# lowess normalize
ntrs.eS <- lnormeS(trs.eS)

# take means over probesets
genesample.eS <- psmeans(ntrs.eS, sample.ind)

pvadjust(pvlist)

Description

This function converts the given raw p-values into the FDR adjusted p-values using R package 'multtest'.

Usage

pvadjust(pvlist)

Arguments

pvlist A list containing raw p-values

Details

pvlist is the output from genediff containing p-values from gene-specific MSE's and posterior MSE's.

Value

pvlist2 A list with the raw p-values and the newly computed FDR adjusted p-values

Author(s)

David Rocke and Geun-Cheol Lee

References

David M. Rocke (2004), Design and analysis of experiments with high throughput biological assay data, Seminars in Cell & Developmental Biology, 15, 703-713.

http://www.idav.ucdavis.edu/~dmrocke/
rowaov

**See Also**

genediff

**Examples**

```r
#library
library(Biobase)
library(LMGene)

#data
data(sample.mat)
data(vlist)
LoggedSmpd0<-neweS(lnorm(log(sample.mat)),vlist)

pvlist<-genediff(LoggedSmpd0)
pvlist$Posterior[1:5,]

apvlist<-pvadjust(pvlist)
names(apvlist)
apvlist$Posterior.FDR[1:5,]
```

---

**rowaov**

*Gene by gene ANOVA function*

**Description**

Computes the mean squares and degrees of freedom for gene-by-gene ANOVAs.

**Usage**

```r
rowaov(eS, model=NULL)
```

**Arguments**

- `eS`: An ExpressionSet object. Any transformation and normalization of `eS` should be done prior to use in `rowaov`.
- `model`: Model used for comparison. See details and `LMGene`.

**Details**

If you have data in a matrix and information about experimental design factors, then you can use `neweS` to convert the data into an ExpressionSet object. Please see `neweS` for more detail.

The `model` argument is an optional character string, constructed like the right-hand side of a formula for `lm`. It specifies which of the variables in the ExpressionSet will be used in the model and whether interaction terms will be included. If `model=NULL`, it uses all variables from the ExpressionSet without interactions. Be careful of using interaction terms with factors; this often leads to overfitting, which will yield an error.
Value

resmat  A matrix of MSEs and degrees of freedom for all model factors and all genes. The first rows of resmat contain MSE’s for each effect in model, ending with the residual MSE. The remaining rows contain degrees of freedom for each effect in the model, ending with the residual d.f. Each column corresponds to a gene.

Author(s)

David Rocke and Geun-Cheol Lee

References


See Also

genediff, LMGene

Examples

```r
library(Biobase)
library(LMGene)

#data
data(sample.mat)
data(vlist)

raw.eS <- neweS(sample.mat, vlist)

# glog transform data
trans.eS <- transeS(raw.eS, lambda = 727, alpha = 56)

# Perform gene-by-gene anova
resmat <- rowaov(trans.eS)
resmat[,1:3]
```

---

**sample.eS**  Sample array data for LMGene

Description

Sample ExpressionSet class data.

Usage

data(sample.eS)

Format

Formal class ExpressionSet [package Biobase].
**Details**

Identical with `neweS(sample.mat, vlist)`, up to metadata

**Examples**

```r
library(Biobase)
library(LMGene)

#data
data(sample.mat)
data(vlist)

Smpdt <- neweS(sample.mat, vlist)

data(sample.eS)
identical(exprs(sample.eS), exprs(Smpdt))
identical(pData(sample.eS), pData(Smpdt))
```

---

**sample.ind**  
*Sample probeset index vector*

**Description**

Vector indicating which probeset each probe belongs to

**Usage**

`data(sample.ind)`

**Format**

A vector of integers, e.g., c(1,1,2,2,3,3,4,4,...). Length is equal to the number of probes (rows) in `sample.mat`.

**Examples**

```r
data(sample.eS)
data(sample.ind)
trs.eS <- transeS(sample.eS, 667, 65)
ntrs.eS <- lnormeS(trs.eS)
genesample.eS <- psmeans(ntrs.eS, sample.ind)
```
sample.mat  

Sample array data for LMGene package

Description

A matrix of array data

Usage

data(sample.mat)

Format

A matrix measuring 613 rows (probes) by 32 columns (samples).

Examples

library(Biobase)
library(LMGene)

#data
data(sample.mat)
data(vlist)

Smpdt<-neweS(sample.mat,vlist)

data(sample.eS)
identical(exprs(sample.eS), exprs(Smpdt))
identical(pData(sample.eS), pData(Smpdt))

tranest  

Glog transformation parameter estimation function

Description

Estimates parameters for the glog transformation, by maximum likelihood or by minimizing the stability score.

Usage

tranest(eS, ngenes = -1, starting = FALSE, lambda = 1000, alpha = 0,
        gradtol = 1e-3, lowessnorm = FALSE, method=1, mult=FALSE, model=NULL,
        SD = FALSE, rank = TRUE, model.based = TRUE, rep.arrays = NULL)
Arguments

eS: An ExpressionSet object
ngen: Number of genes to be used in parameter estimation. Default is to use all genes unless there are more than 100,000, in which case a subset of 50,000 genes is selected at random.
start: If TRUE, user-specified starting values for lambda and alpha are input to the optimization routine
lambda: Starting value for parameter lambda. Ignored unless start = TRUE
alpha: Starting value for parameter alpha. Ignored unless start = TRUE
gradtol: A positive scalar giving the tolerance at which the scaled gradient is considered close enough to zero to terminate the algorithm
lowessnorm: If TRUE, lowess normalization (using lnorm) is used in calculating the likelihood.
method: Determines optimization method. Default is 1, which corresponds to a Newton-type method (see nlm and details.)
mult: If TRUE, tranest will use a vector alpha with one (possibly different) entry per sample. Default is to use same alpha for every sample. SD and mult may not both be TRUE.
model: Specifies model to be used. Default is to use all variables from eS without interactions. See details.
SD: If TRUE, transformation parameters are estimated by minimizing the stability score rather than by maximum likelihood. See details.
ranks: If TRUE, the stability score is calculated by regressing the replicate standard deviations on the ranks of the gene/row means (rather than on the means themselves). Ignored unless SD = TRUE
model.based: If TRUE, the stability score is calculated using the standard deviations of residuals from the linear model in model. Ignored unless SD = TRUE
rep.arrays: List of sets of replicate arrays. Each element of rep.arrays should be a vector with entries corresponding to arrays (columns) in exprs(eS) conducted under the same experimental conditions, i.e., with identical rows in pData(eS). Ignored unless SD = TRUE and model.based = FALSE

Details

If you have data in a matrix and information about experimental design factors, then you can use neweS to convert the data into an ExpressionSet object. Please see neweS for more detail.

The model argument is an optional character string, constructed like the right-hand side of a formula for lm. It specifies which of the variables in the ExpressionSet will be used in the model and whether interaction terms will be included. If model=NULL, it uses all variables from the ExpressionSet without interactions. Be careful of using interaction terms with factors; this often leads to overfitting, which will yield an error.

The default estimation method is maximum likelihood. The likelihood is derived by assuming that there exist values for lambda and alpha such that the residuals from the linear model in model, fit to glog-transformed data using those values for lambda and alpha, follow a normal distribution. See Durbin and Rocke (2003) for details.

If SD = TRUE, lambda and alpha are estimated by minimizing the stability score rather than by maximum likelihood. The stability score is defined as the absolute value of the slope coefficient from the regression of the replicate/residual standard deviation on the gene/row means, or on the
rank of the gene/row means. If `model.based = TRUE`, the stability score is calculated using the standard deviation of residuals from the linear model in `model`. Otherwise, the stability score is calculated using the pooled standard deviation over sets of replicates in `rep.arrays`. See Wu and Rocke (2009) for details.

Optimization methods in `method` are as follows:

1 = Newton-type method, using `nlm`
2 = Nelder-Mead, using `optim`
3 = BFGS, using `optim`
4 = Conjugate gradients, using `optim`
5 = Simulated annealing, using `optim` (may only be used when `mult = TRUE`)

Value

A list with components:

- `lambda`: Estimate of transformation parameter lambda
- `alpha`: Estimate of transformation parameter alpha

Author(s)

David Rocke, Geun-Cheol Lee, John Tillinghast, Blythe Durbin-Johnson, and Shiquan Wu

References


[http://dmrocke.ucdavis.edu](http://dmrocke.ucdavis.edu)

See Also

- `tranestAffyProbeLevel`, `lnorm`, `glog`

Examples

```r
library(Biobase)
library(LMGene)

data(sample.eS)
tranpar <- tranest(sample.eS, 100)
tranpar
tranpar <- tranest(sample.eS, mult=TRUE)
tranpar
```
tranestAffyProbeLevel  Glog transformation parameter estimation function for probe-level Affymetrix expression data

Description

Estimates parameters for the glog transformation on probe-level Affymetrix expression data, by maximum likelihood or by minimizing the stability score.

Usage

tranestAffyProbeLevel(eS, ngenes = 5000, starting = FALSE, lambda = 1000, alpha = 0, gradtol = 0.001, lowessnorm = FALSE, method = 1, mult = FALSE, model = NULL, SD = FALSE, rank = TRUE, model.based = TRUE, rep.arrays = NULL)

Arguments

eS  An AffyBatch object
ngenes  Number of randomly sampled probesets to be used in estimating the transformation parameter
starting  If TRUE, user-specified starting values for lambda and alpha are input to the optimization routine
lambda  Starting value for parameter lambda. Ignored unless starting = TRUE
alpha  Starting value for parameter alpha. Ignored unless starting = TRUE
gradtol  A positive scalar giving the tolerance at which the scaled gradient is considered close enough to zero to terminate the algorithm
lowessnorm  If TRUE, lowess normalization (using lnorm) is used in calculating the likelihood.
method  Determines optimization method. Default is 1, which corresponds to a Newton-type method (see nlm and details.)
mult  If TRUE, tranest will use a vector alpha with one (possibly different) entry per sample. Default is to use same alpha for every sample. SD and mult may not both be TRUE.
model  Specifies model to be used. Default is to use all variables from eS without interactions. See details.
SD  If TRUE, transformation parameters are estimated by minimizing the stability score. See details.
rank  If TRUE, the stability score is calculated by regressing the replicate standard deviation on the rank of the probe/row means (rather than on the means themselves). Ignored unless SD = TRUE
model.based  If TRUE, the stability score is calculated using the standard deviation of residuals from the linear model in model. Ignored unless SD = TRUE
rep.arrays  List of sets of replicate arrays. Each element of rep.arrays should be a vector with entries corresponding to arrays (columns) in exprs(eS) conducted under the same experimental conditions, i.e., with identical rows in pData(eS). Ignored unless SD = TRUE and model.based = FALSE
Details

The `model` argument is an optional character string, constructed like the right-hand side of a formula for `lm`. It specifies which of the variables in the `ExpressionSet` will be used in the model and whether interaction terms will be included. If `model=NULL`, it uses all variables from the `ExpressionSet` without interactions. Be careful of using interaction terms with factors; this often leads to overfitting, which will yield an error.

The default estimation method is maximum likelihood. The likelihood is derived by assuming that there exist values for `lambda` and `alpha` such that the residuals from the linear model in `model`, fit to glog-transformed data using those values for `lambda` and `alpha`, follow a normal distribution. See Durbin and Rocke (2003) for details.

If `SD = TRUE`, `lambda` and `alpha` are estimated by minimizing the stability score rather than by maximum likelihood. The stability score is defined as the absolute value of the slope coefficient from the regression of the replicate/residual standard deviation on the probe/row means, or on the rank of the probe/row means. If `model.based = TRUE`, the stability score is calculated using the standard deviation of residuals from the linear model in `model`. Otherwise, the stability score is calculated using the pooled standard deviation over sets of replicates in `rep.arrays`. See Wu and Rocke (2009) for details.

A random sample of probsets (of size `ngene`) is sampled from `featureNames(eS)`. Expression data from all probes in the sampled probsets is used in estimating the transformation parameters.

Optimization methods in `method` are as follows:

1 = Newton-type method, using `nlm`
2 = Nelder-Mead, using `optim`
3 = BFGS, using `optim`
4 = Conjugate gradients, using `optim`
5 = Simulated annealing, using `optim` (may only be used when `mult = TRUE`)

Value

A list with components:

- `lambda`: Estimate of transformation parameter `lambda`
- `alpha`: Estimate of transformation parameter `alpha`

Author(s)

Lei Zhou, David Rocke, Geun-Cheol Lee, John Tillinghast, Blythe Durbin-Johnson, and Shiquan Wu

References


http://dmrocke.ucdavis.edu
transeS

Function to apply the glog transform to an expression set.

Description

For each element in the array of expression data, this function applies the glog transform $y \rightarrow \text{glog}(y - \alpha, \lambda)$. If alpha is a vector, it must have one element for each column in exprs(eS).

Usage

transeS(es, lambda, alpha)
Arguments

- **eS**: An ExpressionSet or AffyBatch object
- **lambda**: The parameter lambda to be used in the glog transform.
- **alpha**: The alpha parameter(s) for the glog transform. May be a single number used for all samples, or a vector with one entry per sample.

Details

The glog transformation of a variable $y$ is defined as $\log(y + \sqrt{y^2 + \lambda})$. Using $\lambda = 0$ corresponds to the log transformation, up to a scale factor of 2. (Other, equivalent expressions exist for the glog transformation. See Durbin et al. (2002) and Huber et al. (2002) for further details.)

transeS subtracts a (scalar or vector) parameter $alpha$ prior to application of the glog transformation, resulting in the expression $\log(y - alpha + \sqrt{(y - alpha)^2 + \lambda})$.

The parameters $lambda$ and $alpha$ may be estimated using `tranest`.

Value

Returns an ExpressionSet or AffyBatch object with the expression matrix glog-transformed.

Author(s)

John Tillinghast

References


[http://dmrocke.ucdavis.edu](http://dmrocke.ucdavis.edu)

See Also

- `glog`, `tranest`

Examples

```r
library(LMGene)
library(Biobase)

data(sample.eS)
trsmp.eS <- transeS(sample.eS, 667, 65)
```
### vlist

**Sample experimental/phenotype data for LMGene package**

<table>
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<tr>
<th>Description</th>
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<tbody>
<tr>
<td>List of experimental factors for the sample matrix array data, 'sample.mat'.</td>
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<th>Usage</th>
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<tbody>
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<td>data(vlist)</td>
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<td>library(Biobase)</td>
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<td>library(LMGene)</td>
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```r
#data
data(vlist)

testlist
```
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