Package 'EBSeq'

April 5, 2014

Type Package

Title An R package for gene and isoform differential expression analysis of RNA-seq data

Version 1.2.0

Date 2013-6-3

Author Ning Leng

Maintainer Ning Leng <nleng@wisc.edu>

Depends blockmodeling, gplots, R (>= 2.10)

Description Differential Expression analysis at both gene and isoform level using RNA-seq data

License Artistic-2.0

LazyLoad yes

bioViews Statistics, Bioinformatics, DifferentialExpression,MultipleComparisons, RNAseq, HighThroughputSequencing

Collate 'MedianNorm.R' 'GetNg.R' 'beta.mom.R' 'f0.R' 'f1.R'
'Likefun.R' 'LogN.R' 'LogNMulti.R' 'LikefunMulti.R' 'EBTest.R'
'GetPatterns.R' 'EBMultiTest.R' 'GetPP.R' 'PostFC.R'
'GetPPMat.R' 'GetMultiPP.R' 'GetMultiFC.R' 'PlotPostVsRawFC.R'
'crit_fun.R' 'DenNHist.R' 'GetNormalizedMat.R' 'PlotPattern.R'
'PolyFitPlot.R' 'QQP.R' 'QuantileNorm.R' 'RankNorm.R'

BuildVignettes yes

R topics documented:

`EBSeq_NingLeng-package`

EBSeq: RNA-Seq Differential Expression Analysis on both gene and isoform level

---

Description

In 'EBSeq_NingLeng-package,' a Negative Binomial-beta model was built to analyze the RNASeq data. We used the empirical bayes method and EM algrithom.

Details

Package:	EBSeq_NingLeng
Type:	Package
Version:	1.0
Date:	2011-06-13
License:	What license is it under?
LazyLoad:	yes

Author(s)

Ning Leng

Maintainer: Ning Leng <nleng@wisc.edu>

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

See Also

EBTest, EBMultiTest

Examples

```
data(GeneMat)
GeneMat.small = GeneMat[c(1:10,511:550),]
Sizes = MedianNorm(GeneMat.small)
EBOut = EBTest(Data=GeneMat.small,
Conditions=as.factor(rep(c("C1","C2"), each=5)),
sizeFactors=Sizes, maxround=5)
```

---

beta.mom *Fit the beta distribution by method of moments*

Description

'beta.mom' fits the beta distribution by method of moments.

Usage

```
beta.mom(qs.in)
```

Arguments

qs.in	A vector contains the numbers that are assumed to follow a beta distribution.

Value

alpha.hat	Returns the estimation of alpha.
beta.hat	Returns the estimation of beta.

Author(s)

Ning Leng

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

See Also

DenNHist, DenNHistTable

Examples

```
#tmp = rbeta(5, 5, 100)
#param = beta.mom(tmp)
```

`crit_fun` *Calculate the soft threshold for a target FDR*

Description

'crit_fun' calculates the soft threshold for a target FDR.

Usage

```
crit_fun(PPEE, thre)
```

Arguments

`PPEE`	The posterior probabilities of being EE.
`thre`	The target FDR.

Details

Regarding a target FDR alpha, both hard threshold and soft threshold could be used. If the hard threshold is preferred, user could simply take the transcripts with PP(DE) greater than (1-alpha). Using the hard threshold, any DE transcript in the list is with FDR <= alpha.

If the soft threshold is preferred, user could take the transcripts with PP(DE) greater than crit_fun(PPEE, alpha). Using the soft threshold, the list of DE transcripts is with average FDR alpha.

Value

The adjusted FDR threshold of target FDR.

Author(s)

Ning Leng

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

Examples

```
data(GeneMat)
GeneMat.small = GeneMat[c(1:10, 500:600),]
Sizes = MedianNorm(GeneMat.small)
EBOut = EBTest(Data = GeneMat.small,
Conditions = as.factor(rep(c("C1","C2"), each=5)),
sizeFactors = Sizes, maxround = 5)
PP = GetPPMat(EBOut)
DEfound = rownames(PP)[which(PP[,"PPDE"] >= 0.95)]
str(DEfound)

SoftThre = crit_fun(PP[,"PPEE"], 0.05)
DEfound_soft = rownames(PP)[which(PP[,"PPDE"] >= SoftThre)]
```

`DenNHist` *Density plot to compare the empirical q's and the simulated q's from the fitted beta distribution.*

Description

'DenNHist' gives the density plot that compares the empirical q's and the simulated q's from the fitted beta distribution.

Usage

```
DenNHist(EBOut, GeneLevel = F)
```

Arguments

`EBOut`	The output of EBTest or EBMultiTest.
`GeneLevel`	Indicate whether the results are from data at gene level.

Value

For data with n1 conditions and n2 uncertainty groups, n1*n2 plots will be generated. Each plot represents a subset of the data. The empirical estimation of q's will be represented as blue histograms and the density of the fitted beta distribution will be represented as the green line.

Author(s)

Ning Leng

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

See Also

beta.mom, QQP, EBTest, EBMultiTest

Examples

```
data(GeneMat)
GeneMat.small = GeneMat[c(500:1000),]
Sizes = MedianNorm(GeneMat.small)
EBOut = EBTest(Data = GeneMat.small,
Conditions = as.factor(rep(c("C1","C2"), each=5)),
sizeFactors = Sizes, maxround = 5)
par(mfrow = c(2,2))
DenNHist(EBOut)
```

`EBMultiTest` *Using EM algorithm to calculate the posterior probabilities of interested patterns in a multiple condition study*

Description

'EBMultiTest' is built based on the assumption of NB-Beta Empirical Bayes model. It utilizes the EM algorithm to give the posterior probability of the interested patterns.

Usage

```
EBMultiTest(Data, NgVector = NULL, Conditions, AllParti = NULL,
sizeFactors, maxround, Pool = F, NumBin = 1000,
ApproxVal=10^-10, PoolLower=.25, PoolUpper = .75, Print=T)
```

Arguments

`Data`	A data matrix contains expression values for each transcript (gene or isoform level). In which rows should be transcripts and columns should be samples.
`NgVector`	A vector indicates the uncertainty group assignment of each isoform. e.g. if we use number of isoforms in the host gene to define the uncertainty groups, suppose the isoform is in a gene with 2 isoforms, Ng of this isoform should be 2. The length of this vector should be the same as the number of rows in Data. If it's gene level data, Ngvector could be left as NULL.
`Conditions`	A vector indicates the condition in which each sample belongs to.

`AllParti` A matrix indicates the interested patterns. Columns shoule be conditions and rows should be patterns. The matrix could be obtained by the GetPatterns function. If AllParti=NULL, all possible patterns will be used.

`sizeFactors` The normalization factors. It should be a vector with lane specific numbers (the length of the vector should be the same as the number of samples, with the same order as the columns of Data).

`maxround` Number of iterations. The default value is 5. Users should always check the convergency by looking at the Alpha and Beta in output. If the hyper-parameter estimations are not converged in 5 iterations, larger number is suggested.

`Pool` While working without replicates, user could define the Pool = TRUE in the EBTest function to enable pooling.

`NumBin` By defining NumBin = 1000, EBSeq will group the genes with similar means together into 1,000 bins.

`PoolLower, PoolUpper`
With the assumption that only subset of the genes are DE in the data set, we take genes whose FC are in the PoolLower - PoolUpper quantile of the FC's as the candidate genes (default is 25%-75%).

For each bin, the bin-wise variance estimation is defined as the median of the cross condition variance estimations of the candidate genes within that bin.

We use the cross condition variance estimations for the candidate genes and the bin-wise variance estimations of the host bin for the non-candidate genes.

`ApproxVal` The variances of the transcripts with mean < var will be approximated as mean/(1-ApproxVal).

`Print` Whether print the elapsed-time while running the test.

Value

`Alpha` Fitted parameter alpha of the prior beta distribution. Rows are the values for each iteration.

`Beta` Fitted parameter beta of the prior beta distribution. Rows are the values for each iteration.

`P, PFromZ` The bayes estimator of following each pattern of interest. Rows are the values for each iteration.

`Z, PoissonZ` The Posterior Probability of following each pattern of interest for each transcript. (Maybe not in the same order of input).

`RList` The fitted values of r for each transcript.

`MeanList` The mean of each transcript. (across conditions).

`VarList` The variance of each transcript. (across conditions).

`QList` The fitted q values of each transcript within each condition.

`SPMean` The mean of each transcript within each condition (adjusted by the normalization factors).

`SPEstVar` The estimated variance of each transcript within each condition (adjusted by the normalization factors).

PoolVar	The variance of each transcript (The pooled value of within condition EstVar).
DataList	A List of data that grouped with Ng and bias.
PPpattern	The Posterior Probability of following each pattern (columns) for each transcript (rows). Transcripts with expression 0 for all samples are not shown in this matrix.
f	The likelihood of likelihood of prior predictive distribution of being each pattern for each transcript.
AllParti	The matrix describe the patterns.
PPpatternWith0	The Posterior Probability of following each pattern (columns) for each transcript (rows). Transcripts with expression 0 for all samples are shown in this matrix with PP(any_pattrn)=NA.
ConditionOrder	The condition assignment for C1Mean, C2Mean, etc.

Author(s)

Ning Leng

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

See Also

EBTest, GetMultiPP, GetMultiFC

Examples

```
data(MultiGeneMat)
MultiGeneMat.small = MultiGeneMat[201:210,]
Conditions = c("C1","C1","C2","C2","C3","C3")
PosParti = GetPatterns(Conditions)
Parti = PosParti[-3,]
MultiSize = MedianNorm(MultiGeneMat.small)
MultiOut = EBMultiTest(MultiGeneMat.small, NgVector = NULL,
Conditions = Conditions, AllParti = Parti,
sizeFactors = MultiSize, maxround = 5)
MultiPP = GetMultiPP(MultiOut)
```

EBTest *Using EM algorithm to calculate the posterior probabilities of being DE*

Description

Base on the assumption of NB-Beta Empirical Bayes model, the EM algorithm is used to get the posterior probability of being DE.

Usage

```
EBTest(Data, NgVector = NULL, Conditions, sizeFactors, maxround,
Pool = F, NumBin = 1000, ApproxVal = 10^-10, Alpha = NULL,
Beta = NULL, PInput = NULL, RInput = NULL,
PoolLower = .25, PoolUpper = .75, Print = T)
```

Arguments

Data	A data matrix contains expression values for each transcript (gene or isoform level). In which rows should be transcripts and columns should be samples.
NgVector	A vector indicates the uncertainty group assignment of each isoform. e.g. if we use number of isoforms in the host gene to define the uncertainty groups, suppose the isoform is in a gene with 2 isoforms, Ng of this isoform should be 2. The length of this vector should be the same as the number of rows in Data. If it's gene level data, Ngvector could be left as NULL.
Conditions	A factor indicates the condition which each sample belongs to.
sizeFactors	The normalization factors. It should be a vector with lane specific numbers (the length of the vector should be the same as the number of samples, with the same order as the columns of Data).
maxround	Number of iterations. The default value is 5. Users should always check the convergency by looking at the Alpha and Beta in output. If the hyper-parameter estimations are not converged in 5 iterations, larger number is suggested.
Pool	While working without replicates, user could define the Pool = TRUE in the EBTest function to enable pooling.
NumBin	By defining NumBin = 1000, EBSeq will group the genes with similar means together into 1,000 bins.
PoolLower, PoolUpper	With the assumption that only subset of the genes are DE in the data set, we take genes whose FC are in the PoolLower - PoolUpper quantile of the FC's as the candidate genes (default is 25%-75%). For each bin, the bin-wise variance estimation is defined as the median of the cross condition variance estimations of the candidate genes within that bin. We use the cross condition variance estimations for the candidate genes and the bin-wise variance estimations of the host bin for the non-candidate genes.

`ApproxVal` The variances of the transcripts with mean < var will be approximated as mean/(1-ApproxVal).

`Alpha, Beta, PInput, RInput` If the parameters are known and the user doesn't want to estimate them from the data, user could specify them here.

`Print` Whether print the elapsed-time while running the test.

Details

For each transcript gi within condition, the model assumes: X_gis|mu_gi ~ NB (r_gi0 * l_s, q_gi) q_gi|alpha, beta^N_g ~ Beta (alpha, beta^N_g) In which the l_s is the sizeFactors of samples.

The function will test "H0: q_gi^C1 = q_gi^C2" and "H1: q_gi^C1 != q_gi^C2."

Value

`Alpha` Fitted parameter alpha of the prior beta distribution. Rows are the values for each iteration.

`Beta` Fitted parameter beta of the prior beta distribution. Rows are the values for each iteration.

`P, PFromZ` The bayes estimator of being DE. Rows are the values for each iteration.

`Z, PoissonZ` The Posterior Probability of being DE for each transcript(Maybe not in the same order of input).

`RList` The fitted values of r for each transcript.

`MeanList` The mean of each transcript (across conditions).

`VarList` The variance of each transcript (across conditions).

`QListi1` The fitted q values of each transcript within condition 1.

`QListi2` The fitted q values of each transcript within condition 2.

`C1Mean` The mean of each transcript within Condition 1 (adjusted by normalization factors).

`C2Mean` The mean of each transcript within Condition 2 (adjusted by normalization factors).

`C1EstVar` The estimated variance of each transcript within Condition 1 (adjusted by normalization factors).

`C2EstVar` The estimated variance of each transcript within Condition 2 (adjusted by normalization factors).

`PoolVar` The variance of each transcript (The pooled value of within condition EstVar).

`DataList` A List of data that grouped with Ng.

`PPDE` The Posterior Probability of being DE for each transcript (The same order of input).

`f0,f1` The likelihood of the prior predictive distribution of being EE or DE (in log scale).

`AllZeroIndex` The transcript with expression 0 for all samples (which are not tested).

PPMat	A matrix contains posterior probabilities of being EE (the first column) or DE (the second column). Rows are transcripts. Transcripts with expression 0 for all samples are not shown in this matrix.
PPMatWith0	A matrix contains posterior probabilities of being EE (the first column) or DE (the second column). Rows are transcripts. Transcripts with expression 0 for all samples are shown as PP(EE) = PP(DE) = NA in this matrix. The transcript order is exactly the same as the order of the input data.
ConditionOrder	The condition assignment for C1Mean, C2Mean, etc.
Conditions	The input conditions.

Author(s)

Ning Leng

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

See Also

EBMultiTest, PostFC, GetPPMat

Examples

```
data(GeneMat)
str(GeneMat)
GeneMat.small = GeneMat[c(1:10,511:550),]
Sizes = MedianNorm(GeneMat.small)
EBOut = EBTest(Data = GeneMat.small,
Conditions = as.factor(rep(c("C1","C2"), each = 5)),
sizeFactors = Sizes, maxround = 5)
PP = GetPPMat(EBOut)
```

---

f0 *The Prior Predictive Distribution of being EE*

Description

'f0' gives the Prior Predictive Distribution of being EE.

Usage

```
f0(Input, AlphaIn, BetaIn, EmpiricalR, NumOfGroups, log)
```

Arguments

Input	Expression Values.
AlphaIn, BetaIn, EmpiricalR	The parameters estimated from last iteration of EM.
NumOfGroups	How many transcripts within each Ng group.
log	If true, will give the log of the output.

Value

The function will return the prior predictive distribution values of being EE.

Author(s)

Ning Leng

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

See Also

f1

Examples

```
#
#f0(matrix(rnorm(100,100,1),ncol=10), .5, .6,
# matrix(rnorm(100,200,1),ncol=10), 100, TRUE)
```

---

f1 *The Prior Predictive Distribution of being DE*

Description

'f1' gives the Prior Predictive Distribution of DE.

Usage

```
f1(Input1, Input2, AlphaIn, BetaIn, EmpiricalRSP1,
EmpiricalRSP2, NumOfGroup, log)
```

Arguments

Input1	Expressions from Condition1.
Input2	Expressions from Condition2.
AlphaIn, BetaIn, EmpiricalRSP1, EmpiricalRSP2	The parameters estimated from last iteration of EM.
NumOfGroup	How many transcripts within each Ng group.
log	If true, will give the log of the output.

Value

The function will return the prior predictive distribution values of being DE.

Author(s)

Ning Leng

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

See Also

f0

Examples

```
#f1(matrix(rnorm(100,100,1),ncol=10),
# matrix(rnorm(100,100,1),ncol=10), .5, .6,
# matrix(rnorm(100,200,1),ncol=10),
# matrix(rnorm(100,200,1),ncol=10), 100, TRUE)
```

GeneMat *The simulated data for two condition gene DE analysis*

Description

'GeneMat' gives the simulated data for two condition gene DE analysis.

Usage

```
data(GeneMat)
```

Source

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

See Also

IsoList

Examples

```
data(GeneMat)
```

`GetMultiFC` *Calculate the Fold Changes for Multiple Conditions*

Description

'GetMultiFC' calculates the Fold Changes for each pair of conditions in a multiple condition study.

Usage

```
GetMultiFC(EBMultiOut, SmallNum = 0.01)
```

Arguments

`EBMultiOut`	The output of EBMultiTest function.
`SmallNum`	A small number will be added for each transcript in each condition to avoid Inf and NA. Default is 0.01.

Details

Provide the FC (adjusted by the normalization factors) for each pair of comparisons. A small number will be added for each transcript in each condition to avoid Inf and NA. Default is set to be 0.01.

Value

`FCMat`	The FC of each pair of comparison (adjusted by the normalization factors).
`Log2FCMat`	The log 2 FC of each pair of comparison (adjusted by the normalization factors).
`PostFCMat`	The posterior FC of each pair of comparison.
`Log2PostFCMat`	The log 2 posterior FC of each pair of comparison.
`CondMean`	The mean of each transcript within each condition (adjusted by the normalization factors).
`ConditionOrder`	The condition assignment for C1Mean, C2Mean, etc.

Author(s)

Ning Leng

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

See Also

EBMultiTest, PostFC

Examples

```
data(MultiGeneMat)
MultiGeneMat.small = MultiGeneMat[201:210,]

Conditions = c("C1","C1","C2","C2","C3","C3")

PosParti = GetPatterns(Conditions)
Parti = PosParti[-3,]

MultiSize = MedianNorm(MultiGeneMat.small)

MultiOut = EBMultiTest(MultiGeneMat.small,
NgVector=NULL, Conditions=Conditions,
AllParti=Parti, sizeFactors=MultiSize,
maxround=5)

MultiFC = GetMultiFC(MultiOut)
```

GetMultiPP *Posterior Probability of Each Transcript*

Description

'GetMultiPP' generates the Posterior Probability of being each pattern of each transcript based on the EBMultiTest output.

Usage

```
GetMultiPP(EBout)
```

Arguments

EBout	The output of EBMultiTest function.

Value

PP	The poster probabilities of being each pattern.
MAP	Gives the most likely pattern.
Patterns	The Patterns.

Author(s)

Ning Leng

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

See Also

GetPPMat

Examples

```
data(MultiGeneMat)
MultiGeneMat.small = MultiGeneMat[201:210,]

Conditions = c("C1","C1","C2","C2","C3","C3")
PosParti = GetPatterns(Conditions)
Parti = PosParti[-3,]
MultiSize = MedianNorm(MultiGeneMat.small)

MultiOut = EBMultiTest(MultiGeneMat.small,
NgVector=NULL, Conditions=Conditions,
AllParti=Parti, sizeFactors=MultiSize,
maxround=5)
MultiPP = GetMultiPP(MultiOut)
```

GetNg *Ng Vector*

Description

'GetNg' generates the Ng vector for the isoform level data. (While using the number of isoform in the host gene to define the uncertainty groups.)

Usage

```
GetNg(IsoformName, GeneName, TrunThre = 3)
```

Arguments

IsoformName	A vector contains the isoform names.
GeneName	The gene names of the isoforms in IsoformNames (Should be in the same order).
TrunThre	The number of uncertainty groups the user wish to define. The default is 3.

Value

GeneNg	The number of isoforms that are contained in each gene.
GeneNgTrun	The truncated Ng of each gene. (The genes contain more than 3 isoforms are with Ng 3.)
IsoformNg	The Ng of each isoform.
IsoformNgTrun	The truncated Ng of each isoform (could be used to define the uncertainty group assignment).

Author(s)

Ning Leng

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

Examples

```
data(IsoList)

IsoMat = IsoList$IsoMat
IsoNames = IsoList$IsoNames
IsosGeneNames = IsoList$IsosGeneNames
IsoSizes = MedianNorm(IsoMat)
NgList = GetNg(IsoNames, IsosGeneNames)

#IsoNgTrun = NgList$IsoformNgTrun
#IsoEBOut = EBTest(Data = IsoMat, NgVector = IsoNgTrun,
# Conditions = as.factor(rep(c("C1","C2"), each=5)),
# sizeFactors = IsoSizes, maxround = 5)
```

`GetNormalizedMat` *Calculate normalized expression matrix*

Description

'GetNormalizedMat' calculates the normalized expression matrix. (Note: this matrix is only used for visualization etc. EBTes and EBMultiTest request *un-adjusted* expressions and normalization factors.)

Usage

```
GetNormalizedMat(Data, Sizes)
```

Arguments

`Data`	The data matrix with transcripts in rows and lanes in columns.
`Sizes`	A vector contains the normalization factor for each lane.

Value

The function will return a normalized matrix.

Author(s)

Ning Leng

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

Examples

```
data(GeneMat)
str(GeneMat)
Sizes = MedianNorm(GeneMat)
NormData = GetNormalizedMat(GeneMat, Sizes)
```

GetPatterns — *Generate all possible patterns in a multiple condition study*

Description

'GetPatterns' generates all possible patterns in a multiple condition study.

Usage

```
GetPatterns(Conditions)
```

Arguments

Conditions	The names of the Conditions in the study.

Value

A matrix describe all possible patterns.

Author(s)

Ning Leng

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

Examples

```
Conditions = c("C1","C1","C2","C2","C3","C3")
PosParti = GetPatterns(Conditions)
```

GetPP — *Generate the Posterior Probability of each transcript.*

Description

'GetPP' generates the Posterior Probability of being DE of each transcript based on the EBTest output.

Usage

```
GetPP(EBout)
```

Arguments

EBout	The output of EBTest function.

Value

The poster probabilities of being DE.

Author(s)

Ning Leng

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

See Also

GetPPMat

Examples

```
data(GeneMat)
GeneMat.small = GeneMat[c(1:10,500:550),]
Sizes = MedianNorm(GeneMat.small)
EBOut = EBTest(Data = GeneMat.small,
Conditions = as.factor(rep(c("C1","C2"), each=5)),
sizeFactors = Sizes, maxround = 5)
PPDE = GetPP(EBOut)
str(PPDE)
head(PPDE)
```

GetPPMat *Posterior Probability of Transcripts*

Description

'GetPPMat' generates the Posterior Probability of being each pattern of each transcript based on the EBTest output.

Usage

```
GetPPMat(EBout)
```

Arguments

EBout	The output of EBTest function.

Value

The poster probabilities of being EE (first column) and DE (second column).

Author(s)

Ning Leng

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

Examples

```
data(GeneMat)
GeneMat.small = GeneMat[c(500:550),]
Sizes = MedianNorm(GeneMat.small)
EBOut = EBTest(Data = GeneMat.small,
Conditions = as.factor(rep(c("C1","C2"), each=5)),
sizeFactors = Sizes, maxround = 5)
PP = GetPPMat(EBOut)
str(PP)
head(PP)
```

IsoList *The simulated data for two condition isoform DE analysis*

Description

'IsoList' gives the simulated data for two condition isoform DE analysis.

Usage

```
data(IsoList)
```

Source

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

See Also

GeteMat

Examples

```
data(IsoList)
```

IsoMultiList *The simulated data for multiple condition isoform DE analysis*

Description

'IsoMultiList' gives a set of simulated data for multiple condition isoform DE analysis.

Usage

```
data(IsoMultiList)
```

Source

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

See Also

IsoList

Examples

```
data(IsoMultiList)
```

Likefun *Likelihood Function of the NB-Beta Model*

Description

'Likefun' specifies the Likelihood Function of the NB-Beta Model.

Usage

```
Likefun(ParamPool, InputPool)
```

Arguments

ParamPool	The parameters that will be estimated in EM.
InputPool	The control parameters that will not be estimated in EM.

Value

The function will return the log-likelihood.

Author(s)

Ning Leng

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

Examples

```
#x1 = c(.6,.7,.3)
#Input = matrix(rnorm(100,100,1), ncol=10)
#RIn = matrix(rnorm(100,200,1), ncol=10)
#InputPool = list(Input[,1:5], Input[,6:10], Input,
# rep(.1,100), 1, RIn, RIn[,1:5], RIn[,6:10], 100)
#Likefun(x1, InputPool)
```

`LikefunMulti` *Likelihood Function of the NB-Beta Model In Multiple Condition Test*

Description

'LikefunMulti' specifies the Likelihood Function of the NB-Beta Model In Multiple Condition Test.

Usage

```
LikefunMulti(ParamPool, InputPool)
```

Arguments

`ParamPool`	The parameters that will be estimated in EM.
`InputPool`	The control parameters that will not be estimated in EM.

Value

The function will return the log-likelihood.

Author(s)

Ning Leng

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

Examples

```
#x1 = c(.6,.7,.3)
#Input = matrix(rnorm(100,100,1),ncol=10)
#RIn = matrix(rnorm(100,200,1),ncol=10)
#InputPool = list(list(Input[,1:5],Input[,6:10]),
# Input, cbind(rep(.1, 10), rep(.9,10)), 1,
# RIn, list(RIn[,1:5],RIn[,6:10]),
# 10, rbind(c(1,1),c(1,2)))
#LikefunMulti(x1, InputPool)
```

LogN *The function to run EM (one round) algorithm for the NB-beta model.*

Description

'LogN' specifies the function to run (one round of) the EM algorithm for the NB-beta model.

Usage

```
LogN(Input, InputSP, EmpiricalR, EmpiricalRSP, NumOfEachGroup,
AlphaIn, BetaIn, PIn, NoneZeroLength)
```

Arguments

`Input, InputSP`	The expressions among all the samples.
`NumOfEachGroup`	Number of genes in each Ng group.
`AlphaIn, PIn, BetaIn, EmpiricalR, EmpiricalRSP`	The parameters from the last EM step.
`NoneZeroLength`	Number of Ng groups.

Author(s)

Ning Leng

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

Examples

```
#Input = matrix(rnorm(100,100,1), ncol=10)
#rownames(Input) = paste("g",1:10)
#RIn = matrix(rnorm(100,200,1), ncol=10)
#res = LogN(Input, list(Input[,1:5], Input[,6:10]),
# RIn, list(RIn[,1:5], RIn[,6:10]),
# 10, .6, .7, .3, 1)
```

`LogNMulti` *EM algorithm for the NB-beta model in the multiple condition test*

Description

'LogNMulti' specifies the function to run (one round of) the EM algorithm for the NB-beta model in the multiple condition test.

Usage

```
LogNMulti(Input, InputSP, EmpiricalR, EmpiricalRSP,
NumOfEachGroup, AlphaIn, BetaIn, PIn,
NoneZeroLength, AllParti, Conditions)
```

Arguments

`Input, InputSP`	The expressions among all the samples.
`NumOfEachGroup`	Number of genes in each Ng group.
`AlphaIn, PIn, BetaIn, EmpiricalR, EmpiricalRSP`	The parameters from the last EM step.
`NoneZeroLength`	Number of Ng groups.
`AllParti`	The patterns of interests.
`Conditions`	The condition assignment for each sample.

Author(s)

Ning Leng

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

Examples

```
#

#Input = matrix(rnorm(100,100,1),ncol=10)
#rownames(Input) = paste("g",1:10)
#RIn = matrix(rnorm(100,200,1), ncol=10)
#res = LogNMulti(Input, list(Input[,1:5], Input[,6:10]),
# RIn, list(RIn[,1:5], RIn[,6:10]), 10, .6, .7,
# c(.3,.7), 1, rbind(c(1,1), c(1,2)),
# as.factor(rep(c("C1","C2"), each=5)))
```

`MedianNorm` *Median Normalization*

Description

'MedianNorm' specifies the median normalization function from Anders et. al., 2010.

Usage

```
MedianNorm(Data)
```

Arguments

`Data`	The data matrix with transcripts in rows and lanes in columns.

Value

The function will return a vector contains the normalization factor for each lane.

Author(s)

Ning Leng

References

Simon Anders and Wolfgang Huber. Differential expression analysis for sequence count data. Genome Biology (2010) 11:R106 (open access)

See Also

QuantileNorm

Examples

```
data(GeneMat)
Sizes = MedianNorm(GeneMat)
#EBOut = EBTest(Data = GeneMat,
# Conditions = as.factor(rep(c("C1","C2"), each=5)),
# sizeFactors = Sizes, maxround = 5)
```

MultiGeneMat *The simulated data for multiple condition gene DE analysis*

Description

'MultiGeneMat' generates a set of the simulated data for multiple condition gene DE analysis.

Usage

```
data(MultiGeneMat)
```

Source

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

See Also

GeneMat

Examples

```
data(MultiGeneMat)
```

PlotPattern *Visualize the patterns*

Description

'PlotPattern' generates the visualized patterns before the multiple condition test.

Usage

```
PlotPattern(Patterns)
```

Arguments

Patterns	The output of GetPatterns function.

Value

A heatmap to visualize the patterns of interest.

Author(s)

Ning Leng

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

Examples

```
Conditions = c("C1","C1","C2","C2","C3","C3")
Patterns = GetPatterns(Conditions)
PlotPattern(Patterns)
```

PlotPostVsRawFC *Plot Posterior FC vs FC*

Description

'PlotPostVsRawFC' helps the users visualize the posterior FC vs FC in a two condition study.

Usage

```
PlotPostVsRawFC(EBOut, FCOut)
```

Arguments

EBOut	The output of EBMultiTest function.
FCOut	The output of PostFC function.

Value

A figure shows fold change vs posterior fold change.

Author(s)

Ning Leng

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

See Also

PostFC

Examples

```
data(GeneMat)
GeneMat.small = GeneMat[c(500:600),]
Sizes = MedianNorm(GeneMat.small)
EBOut = EBTest(Data = GeneMat.small,
Conditions = as.factor(rep(c("C1","C2"), each=5)),
sizeFactors = Sizes, maxround = 5)
FC = PostFC(EBOut)
PlotPostVsRawFC(EBOut,FC)
```

PolyFitPlot *Fit the mean-var relationship using polynomial regression*

Description

'PolyFitPlot' fits the mean-var relationship using polynomial regression.

Usage

```
PolyFitPlot(X, Y, nterms, xname = "Estimated Mean",
yname = "Estimated Var", pdfname = "",
xlim =  c(-1,5), ylim = c(-1,7), ChangeXY = F,
col = "red")
```

Arguments

X	The first group of values want to be fitted by the polynomial regression (e.g Mean of the data).
Y	The second group of values want to be fitted by the polynomial regression (e.g. variance of the data). The length of Y should be the same as the length of X.
nterms	How many polynomial terms want to be used.
xname	Name of the x axis.
yname	Name of the y axis.
pdfname	Name of the plot.
xlim	The x limits of the plot.
ylim	The y limits of the plot.
ChangeXY	If ChangeXY is setted to be TRUE, X will be treated as the dependent variable and Y will be treated as the independent one. Default is FALSE.
col	Color of the fitted line.

Value

The PolyFitPlot function provides a smooth scatter plot of two variables and their best fitting line of polynomial regression.

Author(s)

Ning Leng

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

Examples

```
data(IsoList)
str(IsoList)
IsoMat = IsoList$IsoMat
IsoNames = IsoList$IsoNames
IsosGeneNames = IsoList$IsosGeneNames
IsoSizes = MedianNorm(IsoMat)
NgList = GetNg(IsoNames, IsosGeneNames)

IsoNgTrun = NgList$IsoformNgTrun
#IsoEBOut = EBTest(Data = IsoMat.small,
# NgVector = IsoNgTrun,
# Conditions = as.factor(rep(c("C1","C2"), each=5)),
# sizeFactors = IsoSizes, maxround = 5)

#par(mfrow=c(2,2))
#PolyFitValue = vector("list",3)

#for(i in 1:3)
# PolyFitValue[[i]] = PolyFitPlot(IsoEBOut$C1Mean[[i]],
# IsoEBOut$C1EstVar[[i]], 5)

#PolyAll = PolyFitPlot(unlist(IsoEBOut$C1Mean),
# unlist(IsoEBOut$C1EstVar), 5)

#lines(log10(IsoEBOut$C1Mean[[1]][PolyFitValue[[1]]$sort]),
# PolyFitValue[[1]]$fit[PolyFitValue[[1]]$sort],
# col="yellow", lwd=2)
#lines(log10(IsoEBOut$C1Mean[[2]][PolyFitValue[[2]]$sort]),
# PolyFitValue[[2]]$fit[PolyFitValue[[2]]$sort],
# col="pink", lwd=2)
#lines(log10(IsoEBOut$C1Mean[[3]][PolyFitValue[[3]]$sort]),
# PolyFitValue[[3]]$fit[PolyFitValue[[3]]$sort],
# col="green", lwd=2)

#legend("topleft",c("All Isoforms","Ng = 1","Ng = 2","Ng = 3"),
# col = c("red","yellow","pink","green"),
```

```
# lty=1, lwd=3, box.lwd=2)
```

PostFC *Calculate the posterior fold change for each transcript across conditions*

Description

'PostFC' calculates the posterior fold change for each transcript across conditions.

Usage

```
PostFC(EBoutput, SmallNum = 0.01)
```

Arguments

EBoutput	The ourput from function EBTest.
SmallNum	A small number will be added for each transcript in each condition to avoid Inf and NA. Default is 0.01.

Value

Provide both FC and posterior FC across two conditions. FC is calculated as (MeanC1+SmallNum)/(MeanC2+SmallNum). And Posterior FC is calculated as:

Post alpha P_a_C1 = alpha + r_C1 * n_C1

Post beta P_b_C1 = beta + Mean_C1 * n_C1

P_q_C1 = P_a_C1 / (P_a_C1 + P_b_C1)

Post FC = ((1-P_q_C1)/P_q_c1) / ((1-P_q_c2)/P_q_c2)

PostFC	The posterior FC across two conditions.
RealFC	The FC across two conditions (adjusted by the normalization factors).
Direction	The diretion of FC calculation.

Author(s)

Ning Leng

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

See Also

EBTest, GetMultiFC

Examples

```
data(GeneMat)
GeneMat.small = GeneMat[c(500:550),]
Sizes = MedianNorm(GeneMat.small)
EBOut = EBTest(Data = GeneMat.small,
Conditions = as.factor(rep(c("C1","C2"), each=5)),
sizeFactors = Sizes, maxround = 5)
FC=PostFC(EBOut)
```

QQP *The Quantile-Quantile Plot to compare the empirical q's and simulated q's from fitted beta distribution*

Description

'QQP' gives the Quantile-Quantile Plot to compare the empirical q's and simulated q's from fitted beta distribution.

Usage

```
QQP(EBOut, GeneLevel = F)
```

Arguments

EBOut	The output of EBTest or EBMultiTest.
GeneLevel	Indicate whether the results are from data at gene level.

Value

For data with n1 conditions and n2 uncertainty groups, n1*n2 plots will be generated. Each plot represents a subset of the data.

Author(s)

Ning Leng

References

Ning Leng, John A. Dawson, James A. Thomson, Victor Ruotti, Anna I. Rissman, Bart M.G. Smits, Jill D. Haag, Michael N. Gould, Ron M. Stewart, and Christina Kendziorski. EBSeq: An empirical Bayes hierarchical model for inference in RNA-seq experiments. Bioinformatics (2013)

See Also

EBTest, EBMultiTest, DenNHist

Examples

```
data(GeneMat)
GeneMat.small = GeneMat[c(500:1000),]
Sizes = MedianNorm(GeneMat.small)
EBOut = EBTest(Data = GeneMat.small,
Conditions = as.factor(rep(c("C1","C2"), each=5)),
sizeFactors = Sizes, maxround = 5)
par(mfrow=c(2,2))
QQP(EBOut)
```

QuantileNorm *Quantile Normalization*

Description

'QuantileNorm' gives the quantile normalization.

Usage

```
QuantileNorm(Data, Quantile)
```

Arguments

Data	The data matrix with transcripts in rows and lanes in columns.
Quantile	The quantile the user wishs to use. Should be a number between 0 and 1.

Details

Use a quantile point to normalize the data.

Value

The function will return a vector contains the normalization factor for each lane.

Author(s)

Ning Leng

References

Bullard, James H., et al. Evaluation of statistical methods for normalization and differential expression in mRNA-Seq experiments. BMC bioinformatics 11.1 (2010): 94.

See Also

MedianNorm

Examples

```
data(GeneMat)
Sizes = QuantileNorm(GeneMat,.75)
#EBOut = EBTest(Data = GeneMat,
# Conditions = as.factor(rep(c("C1","C2"), each=5)),
# sizeFactors = Sizes, maxround = 5)
```

RankNorm *Rank Normalization*

Description

'RankNorm' gives the rank normalization.

Usage

```
RankNorm(Data)
```

Arguments

Data	The data matrix with transcripts in rows and lanes in columns.

Value

The function will return a matrix contains the normalization factor for each lane and each transcript.

Author(s)

Ning Leng

See Also

MedianNorm, QuantileNorm

Examples

```
data(GeneMat)
Sizes = RankNorm(GeneMat)
# Run EBSeq
# EBres = EBTest(Data = GeneData, NgVector = rep(1,10^4),
# Vect5End = rep(1,10^4), Vect3End = rep(1,10^4),
# Conditions = as.factor(rep(c(1,2), each=5)),
# sizeFactors = Sizes, maxround=5)
```

Index