

Package ‘NBAMSeq’

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Type Package

Title Negative Binomial Additive Model for RNA-Seq Data

Version 1.23.0

Description High-throughput sequencing experiments followed by differential expression analysis is a widely used approach to detect genomic biomarkers. A fundamental step in differential expression analysis is to model the association between gene counts and covariates of interest. NBAMSeq a flexible statistical model based on the generalized additive model and allows for information sharing across genes in variance estimation.

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URL <https://github.com/reese3928/NBAMSeq>

BugReports <https://github.com/reese3928/NBAMSeq/issues>

Encoding UTF-8

Imports DESeq2, mgcv(>= 1.8-24), BiocParallel, genefilter, methods, stats,

Depends R (>= 3.6), SummarizedExperiment, S4Vectors

Suggests knitr, rmarkdown, testthat, ggplot2

RoxygenNote 6.1.0

VignetteBuilder knitr

biocViews RNASeq, DifferentialExpression, GeneExpression, Sequencing, Coverage

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| | |
|-------------|---------------------------------------|
| makeExample | <i>Make an example NBAMSeqDataSet</i> |
|-------------|---------------------------------------|

Description

This function makes an example NBAMSeqDataSet

Usage

```
makeExample(n = 200, m = 30)
```

Arguments

| | |
|---|-------------------|
| n | number of genes |
| m | number of samples |

Value

a NBAMSeqDataSet object

References

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

Examples

```
gsd = makeExample()
```

`makeplot`*Making plots to visualize nonlinear associations*

Description

This function makes plots to visualize nonlinear associations.

Usage

```
makeplot(object, phenoname, genename, ...)
```

Arguments

| | |
|------------------------|---|
| <code>object</code> | a NBAMSeqDataSet object |
| <code>phenoname</code> | the name of nonlinear variable to be visualized |
| <code>genename</code> | the name of gene to be visualized |
| <code>...</code> | additional arguments provided to plot.gam |

Value

the plot made by `plot.gam()` function

Examples

```
gsd = makeExample(n = 3, m = 10)
gsd = NBAMSeq(gsd)
makeplot(gsd, "pheno", "gene3", main = "gene10")
```

`NBAMSeq`*Differential expression analysis based on negative binomial additive model*

Description

This function performs differential expression analysis based on negative binomial additive model.

Usage

```
NBAMSeq(object, gamma = 2.5, parallel = FALSE, fitlin = FALSE,
        BPPARAM = bpparam(), ...)
```

Arguments

| | |
|----------|--|
| object | a NBAMSeqDataSet object |
| gamma | a number greater or equal to 1. Increase gamma to create smoother models. Default gamma is 2.5. See gam for details. |
| parallel | either TRUE or FALSE indicating whether parallel should be used. Default is FALSE |
| fitlin | either TRUE or FALSE indicating whether linear model should be fitted. Default is FALSE |
| BPPARAM | an argument provided to bplapply . See register for details. |
| ... | additional arguments provided to gam |

Value

a NBAMSeqDataSet object

References

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

Examples

```
gsd = makeExample(n = 3, m = 10)
gsd = NBAMSeq(gsd)
```

 NBAMSeq-methods

Accessor functions and replace methods for NBAMSeqDataSet object

Description

Accessor functions and replace methods for NBAMSeqDataSet object

For `getDesign()`: accessor to the design formula

For `getsf()`: accessor to the size factors

Replace methods for NBAMSeqDataSet object

For `setsf()`: replace size factors

Usage

```
getDesign(theObject)
```

```
## S4 method for signature 'NBAMSeqDataSet'
getDesign(theObject)
```

```
getsf(theObject)
```

```
## S4 method for signature 'NBAMSeqDataSet'  
getsf(theObject)  
  
setsf(theObject) <- value  
  
## S4 replacement method for signature 'NBAMSeqDataSet,numeric'  
setsf(theObject) <- value
```

Arguments

theObject a NBAMSeqDataSet object
value the values to be included in the object

Value

For getDesign(): design formula
For getsf(): size factor
For setsf(): NBAMSeq object

References

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

Examples

```
## For getDesign() ##  
gsd = makeExample()  
design_gsd = getDesign(gsd)  
## For getsf() ##  
gsd = makeExample()  
sf = getsf(gsd)  
## For setsf() ##  
n = 100  
m = 50  
gsd = makeExample(n = n, m = m)  
sf = sample(1:5, m, replace = TRUE)  
setsf(gsd) = sf
```

NBAMSeqDataSet

NBAMSeqDataSet constructor

Description

NBAMSeqDataSet constructor

Usage

```
NBAMSeqDataSet(countData, colData, design, ...)
```

Arguments

```
countData      a matrix or data frame contains gene count
colData        a DataFrame or data.frame
design          a mgcv type design. e.g. ~ s(pheno) or ~ s(pheno) + var1 + var2
...           optional arguments passed to SummarizedExperiment
```

Value

a NBAMSeqDataSet object

Examples

```
n = 100 ## n stands for number of genes
m = 20  ## m stands for sample size
countData = matrix(rnbinom(n*m, mu=100, size=1/3), ncol = m)
mode(countData) = "integer"
colnames(countData) = paste0("sample", 1:m)
rownames(countData) = paste0("gene", 1:n)
pheno = runif(m, 20, 80)
colData = data.frame(pheno = pheno)
rownames(colData) = paste0("sample", 1:m)
gsd = NBAMSeqDataSet(countData = countData,
colData = colData, design = ~s(pheno))
```

NBAMSeqDataSet-class *NBAMSeqDataSet class*

Description

NBAMSeqDataSet is a class inherited from [SummarizedExperiment](#). It is used to store the count matrix, colData, and design formula in differential expression analysis.

Slots

design a mgcv-type design formula

References

Martin Morgan, Valerie Obenchain, Jim Hester and Hervé Pagès (2018). SummarizedExperiment: SummarizedExperiment container. R package version 1.12.0.

| | |
|---------|---------------------------|
| results | <i>Pulling out result</i> |
|---------|---------------------------|

Description

This function pulls out result from NBAMSeqDataSet object returned by [NBAMSeq](#)

Usage

```
results(object, name, contrast, indepfilter = TRUE, alpha = 0.1,
        pAdjustMethod = "BH", parallel = FALSE, BPPARAM = bpparam(), ...)
```

Arguments

| | |
|---------------|---|
| object | a NBAMSeqDataSet object returned by NBAMSeq |
| name | the name of nonlinear variable or continuous linear variable |
| contrast | a character of length 3. 1st element: name of factor variable; 2nd element: name of numerator level; 3rd element: name of denominator level. contrast = c("group", "treatment", "control") means comparing treatment vs control for group variable. |
| indepfilter | either TRUE or FALSE indicating whether independent filtering should be performed. Default is TRUE. |
| alpha | significant threshold for declaring genes as differentially expressed. Default is 0.1. |
| pAdjustMethod | pvalue adjustment method. Default is "BH". See p.adjust for details. |
| parallel | either TRUE or FALSE indicating whether parallel should be used. Default is FALSE. |
| BPPARAM | an argument provided to bplapply . See register for details. |
| ... | additional arguments provided to pvalueAdjustment function in DESeq2. See results for details. |

Value

a DataFrame which contains the result

References

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

Examples

```
gsd = makeExample(n = 3, m = 10)
gsd = NBAMSeq(gsd)
res = results(gsd, name = "pheno")
```

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