

# Package ‘EBSEA’

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**Type** Package

**Title** Exon Based Strategy for Expression Analysis of genes

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**Description** Calculates differential expression of genes based on exon counts of genes obtained from RNA-seq sequencing data.

**License** GPL-2

**biocViews** Software, DifferentialExpression, GeneExpression, Sequencing

**Imports** edgeR, limma, gtools, graphics, stats

**NeedsCompilation** no

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EBSEA

*Exon Based Startegy for Expression Analysis of genes*

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## Description

EBSEA takes as input unnormalized counts of exons, normalizes them and then performs a two group comparison of the samples to detect differentially expressed between the groups. Both paired or unpaired comparison are supported. It calculates fold changes, p-values and false discovery rate of the genes between the groups.

**Usage**

```
EBSEA(countData, group, paired = FALSE, effects = NULL, plot = FALSE)
```

**Arguments**

<code>countData</code>	A datafram of exon count data
<code>group</code>	A vector indicating the sample groups in the experiment
<code>paired</code>	A logical indicating whether the samples are paired or unpaired. Default: FALSE
<code>effects</code>	A vector indicating the paired samples.
<code>plot</code>	A logical indicating whether a volcano plot is visualized. Default: FALSE

**Value**

EBSEA returns a list of two dataframes. ExonTable is a datafram that contains exon statistics including log fold change, p-values, adjusted p-values, average expression and fold change. GeneTable is a datafram that contains the corresponding fold change, log fold change, p-values and false discovery rate.

**References**

Laiho, A., & Elo, L. L. (2014). A note on an exon-based strategy to identify differentially expressed genes in RNA-seq experiments. PloS One, 9(12), e115964.

**See Also**

[visualizeGenes](#)

**Examples**

```
data(origCounts)
group <- c('Group1', 'Group1', 'Group1', 'Group2', 'Group2', 'Group2', 'Group2')
result <- EBSEA(origCounts, group)
```

`filterCounts`

*Filter Count Data*

**Description**

The exons are filtered based on their expression levels so that each exon has a cpm (count per million) of more than 1 in user defined percent of the samples.

**Usage**

```
filterCounts(x, noOfSamples)
```

**Arguments**

- |             |  |
|-------------|--|
| x           | A numeric dataframe of counts in the sample with gene and exon number as the row names and samples as the column names |
| noOfSamples | Percentage of the number of samples that should have cpm greater than 1.   |

**Value**

A dataframe of filtered counts of exons

**See Also**

[EBSEA](#)

**Examples**

```
data(origCounts)
res <- filterCounts(origCounts, 20)
```

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filterGenes

*Filter Gene List*

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**Description**

The differentially expressed genes are filtered based on the FC and FDR provided by the user. The default thresholds are FC => 1.25 and fdr <= 0.01

**Usage**

```
filterGenes(x, fc = 1.25, fdr = 0.01)
```

**Arguments**

- |     |  |
|-----|--|
| x   | A dataframe containing the gene statistics returned by EBSEA.        |
| fc  | A fold change threshold for the genes to be filtered. Default: 1.25. |
| fdr | A FDR threshold for the genes to be filtered. Default: 0.01.         |

**Value**

A list containing upregulated and downregulated genes.

**See Also**

[EBSEA](#)

## Examples

```
data(origCounts)
group <- c('Group1', 'Group1', 'Group1', 'Group2', 'Group2', 'Group2', 'Group2')
result <- EBSEA(origCounts, group)
filteredGenes <- filterGenes(result$GeneTable)
```

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origCounts

*Subset of Pasilla Dataset*

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## Description

origCounts consists of a subset of the exon counts from the pasilla dataset.

## Usage

```
data("origCounts")
```

## Format

A data frame with 1000 observations on the following 7 variables.

```
treated1fb a numeric vector
treated2fb a numeric vector
treated3fb a numeric vector
untreated1fb a numeric vector
untreated2fb a numeric vector
untreated3fb a numeric vector
untreated4fb a numeric vector
```

## Value

Dataset

## See Also

[EBSEA](#)

## Examples

```
data(origCounts)
```

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visualizeGenes	<i>Visualize Gene</i>
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## Description

Plots for each exon of the gene entered by the user, the mean of the counts and the fold changes.

## Usage

```
visualizeGenes(gene, group, countData, result)
```

## Arguments

gene	Gene Name. The gene name should be the from the genes in count data.
group	A vector indicating the sample group in the experiment.
countData	A data frame of the original exon count data.
result	Results returned by EBSEA.

## Value

A plot of mean counts and fold changes of exons of a gene.

## See Also

[EBSEA](#)

## Examples

```
data(origCounts)
group <- c('Group1', 'Group1', 'Group1', 'Group2', 'Group2', 'Group2', 'Group2')
result <- EBSEA(origCounts, group)
visualizeGenes('FBgn0000017', group, origCounts, result)
```

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