

# Package ‘lefsr’

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**Type** Package

**Title** R implementation of the LEfSE method for microbiome biomarker discovery

**Description** lefsr is an implementation in R of the popular “LDA Effect Size (LEfSe)” method for microbiome biomarker discovery. It uses the Kruskal-Wallis test, Wilcoxon-Rank Sum test, and Linear Discriminant Analysis to find biomarkers of groups and sub-groups.

**Version** 1.13.4

**License** Artistic-2.0

**Depends** SummarizedExperiment, R (>= 4.0.0)

**Imports** coin, MASS, ggplot2, S4Vectors, stats, methods, utils

**Suggests** knitr, rmarkdown, curatedMetagenomicData, BiocStyle, phyloseq, testthat, pkgdown, covr, withr

**Encoding** UTF-8

**BugReports** <https://github.com/waldronlab/lefsr/issues>

**URL** <https://github.com/waldronlab/lefsr>

**VignetteBuilder** knitr

**biocViews** Software, Sequencing, DifferentialExpression, Microbiome, StatisticalMethod, Classification

**RoxygenNote** 7.3.1

**Roxygen** list(markdown = TRUE)

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lefser	<i>R implementation of the LEfSe method</i>
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## Description

Perform a LEfSe analysis: the function carries out differential analysis between two sample groups for multiple microorganisms and uses linear discriminant analysis to establish their effect sizes. Subclass information for each class can be incorporated into the analysis (see examples). Microorganisms with large differences between two sample groups are identified as biomarkers.

## Usage

```
lefser(
  relab,
  kruskal.threshold = 0.05,
  wilcox.threshold = 0.05,
  lda.threshold = 2,
  groupCol = "GROUP",
  blockCol = NULL,
  assay = 1L,
  trim.names = FALSE,
  checkAbundances = TRUE,
  ...,
  expr
)
```

## Arguments

relab	A <a href="#">SummarizedExperiment</a> with relative abundances in the assay
kruskal.threshold	numeric(1) The p-value for the Kruskal-Wallis Rank Sum Test (default 0.05).

wilcox.threshold	numeric(1) The p-value for the Wilcoxon Rank-Sum Test when 'blockCol' is present (default 0.05).
lda.threshold	numeric(1) The effect size threshold (default 2.0).
groupCol	character(1) Column name in colData(relab) indicating groups, usually a factor with two levels (e.g., c("cases", "controls")); default "GROUP".
blockCol	character(1) Optional column name in colData(relab) indicating the blocks, usually a factor with two levels (e.g., c("adult", "senior")); default NULL.
assay	The i-th assay matrix in the SummarizedExperiment ('relab'; default 1).
trim.names	If TRUE extracts the most specific taxonomic rank of organism.
checkAbundances	logical(1) Whether to check if the assay data in the relab input are relative abundances or counts. If counts are found, a warning will be emitted (default TRUE).
expr	(deprecated) Use relab instead. A <a href="#">SummarizedExperiment</a> with relative abundances in the assay
...	Additional inputs to lower level functions (not used).

### Details

The LefSe method expects relative abundances in the `expr` input. A warning will be emitted if the column sums do not result in 1. Use the `relativeAb` helper function to convert the data in the `SummarizedExperiment` to relative abundances. The `checkAbundances` argument enables checking the data for presence of relative abundances and can be turned off by setting the argument to `FALSE`.

### Value

The function returns a `data.frame` with two columns, which are names of microorganisms and their LDA scores.

### Examples

```
# (1) Using classes only
data(zeller14)
# exclude 'adenoma'
zeller14 <- zeller14[, zeller14$study_condition != "adenoma"]
res_group <- lefser(zeller14, groupCol = "study_condition")
head(res_group)

# (2) Using classes and subclasses
data(zeller14)
# exclude 'adenoma'
zeller14 <- zeller14[, zeller14$study_condition != "adenoma"]
res_block <- lefser(
  zeller14, groupCol = "study_condition", blockCol = "age_category"
)
head(res_block)
```

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lefserPlot	<i>Plots results from lefser function</i>
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### Description

lefserPlot function displays effect sizes for differentially expressed microorganisms and whether they are more abundant in '0' or '1' sample group.

### Usage

```
lefserPlot(df, colors = c("red", "forestgreen"), trim.names = TRUE)
```

### Arguments

df	Data frame produced by lefser.
colors	character(2) The two colors corresponding to class 0 and 1, respectively. Defaults to c("red", "forestgreen").
trim.names	If TRUE extracts the most specific taxonomic rank of organism.

### Value

Function returns plot of effect size scores produced by lefser. Positive scores represent microorganisms with that are more abundant in class '1'. Negative scores represent microorganisms with that are more abundant in class '0'.

### Examples

```
example("lefser")
lefserPlot(res_group)
```

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relativeAb	<i>Utility function to calculate relative abundances</i>
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### Description

The function calculates the column totals and divides each value within the column by the respective column total.

This function calculates the relative abundance of each feature in the SummarizedExperiment object containing count data, expressed as counts per million (CPM)

### Usage

```
relativeAb(se, assay = 1L)
```

**Arguments**

`se` A SummarizedExperiment object with counts  
`assay` The i-th assay matrix in the SummarizedExperiment (`'relab'`; default 1).

**Value**

returns a new SummarizedExperiment object with counts per million calculated and added as a new assay named `rel_abs`.

**Examples**

```
se <- SummarizedExperiment(  
  assays = list(  
    counts = matrix(  
      rep(1, 4), ncol = 1, dimnames = list(LETTERS[1:4], "SAMP")  
    )  
  )  
)  
assay(se)  
assay(relativeAb(se))
```

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zeller14

*Example dataset for lefser*

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**Description**

The ZellerG\_2014 dataset contains microbiome count data for CRC patients and controls. It was for curatedMetagenomicData using the script in the package directory "data-raw".

**Usage**

```
data("zeller14")
```

**Format**

A SummarizedExperiment with 1585 features, 199 samples

**study\_condition** adenoma, control, CRC

**age\_category** adult, senior

**Source**

<https://pubmed.ncbi.nlm.nih.gov/25432777/>

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