

# Package ‘CoDaSeq’

August 16, 2016

**Type** Package

**Title** Compositional Data Analysis of High Throughput Sequencing

**Version** 0.99.1

**Date** 2016-05-21

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**Description** A set of common functions for the analysis of high throughput sequencing count data

**License** file LICENSE

**NeedsCompilation** no

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codaSeq.clr	<i>Center Log-Ratio Function.</i>
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## Description

Returns a matrix of center log-ratio transformed data with samples by row. Equivalent to  $\log(x/gx)$  for every value where  $gx$  is the geometric mean of the vector  $X$ .

## Usage

```
codaSeq.clr <- function(x, samples.by.row=TRUE)
```

## Arguments

**x** A matrix or dataframe with samples by rows or columns.  
**samples.by.row** TRUE if samples are by row, FALSE if samples are by column.

**Details**

Natural log is used for biplots and other exploratory analyses.

**Value**

returns a matrix of clr tranformed values with samples in the rows and variables in columns

**Author(s)**

Greg Gloor, Jean Macklaim, Wallace Chan

**References**

Please use the citation given by `citation(package="CoDaSeq")`

**See Also**

[codaSeq.filter](#), [codaSeq.outlier](#), [codaSeq.rarefy](#), [codaSeq.propr.phismy](#), [codaSeq.propr.aldex.phi](#)

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codaSeq.filter	<i>Filter compositional dataset for 0 values and abundance.</i>
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**Description**

Returns a reduced able of counts where the samples must contain a minimum number of reads, and OTUs must be found with a minimum abundance in all remaining samples.

**Usage**

```
codaSeq.filter <- function(x, y=tax.vector, min.reads=5000, min.prop=0.001, max.prop=0.025,
  min.occurrence=0, samples.by.row=TRUE)
```

**Arguments**

x	A matrix or dataframe containing a count table.
min.reads	The minimum reads per sample. Default=5000.
min.prop	The minimum proportional abundance of a read in any sample. Default=0.001.
max.prop	The maximum proportional abundance of a read in any sample. Default=0.025.
min.fraction	The minimum sample proportion of non-0 reads for each variable.
sample.by.row	True if rows contain samples, false if rows contain variables.

**Details**

Filters min/max.prop first, min.fraction second, min/max.prop third. Requires numeric data only.

**Value**

Returns a dataframe with the following information:

data.0	
data.1	
data.2	Returns a reduced vector with filtered samples by rows.

**Author(s)**

Greg Gloor, Jean Macklai, Wallace Chan

**References**

Please use the citation given by `citation(package="CoDaSeq")`

**See Also**

[codaSeq.clr](#), [codaSeq.outlier](#), [codaSeq.rarefy](#), [codaSeq.propr.phismy](#), [codaSeq.propr.aldex.phi](#)

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codaSeq.outlier	<i>Identifying sample Outliers.</i>
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**Description**

Returns a list of proportional contribution to group variance, sample names that are outliers, and sample names that are not outliers.

**Usage**

```
codaSeq.outlier <- function(x, plot.me=TRUE, col=rgb(1,0,0,0.3))
```

**Arguments**

x	A matrix or data frame with clr transformed values, with sample by row.
plot.me	A logical value determining if a histogram should be plotted of the variance contribution per sample.
col	RGB values for your selection of colour.

**Details**

Samples must be grouped. This approach makes no sense across groups. If you do not know if you have natural groups, ignore this step and exam your data by PCA. Outliers are defined as those contributing greater than the median plus twice the interquartile range of the sample variance to the total.

**Value**

Returns list

sample.var	Proportional variance contributions for each sample.
bad	Rownames of outlier samples.
good	Rownames of non-outlier samples.

**Author(s)**

Greg Gloor, Jean Macklai, Wallace Chan

**References**

Please use the citation given by `citation(package="CoDaSeq")`

**See Also**

[codaSeq.clr](#), [codaSeq.filter](#), [codaSeq.rarefy](#), [codaSeq.propr.phismy](#), [codaSeq.propr.aldex.phi](#)

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propr.aldex.phi

*Expected Value of Phi From Dirichlet Log-Ratio Distribution*

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**Description**

Returns data frame of the lower-triangle of symmetrical phi metric, where the value of phi is the expected value of a number of Dirichlet Monte-Carlo replicates of the data. This reduces the problem of 0-count and low-count features being highly variable because their values range wildly and so the expected value is always large.

**Usage**

```
propr.aldex.phi <- function(aldex.clr)
```

**Details**

Requires `aldex.clr` function from ALDEx2 Package. Param `aldex.clr` is an S3 object from the `aldex.clr` function. We ignore all the other measures that are used for trouble-shooting phi. The `sma.df` function in particular is very time and memory intensive

**Value**

<code>sym.phi</code>	Calculated sum of phi values through all DIR MR instances.
<code>lt</code>	Indice of correct size.
<code>lt.int</code>	Indice of correct size.
<code>sma.df</code>	Dataframe to hold info.
<code>sma.df\$phi</code>	Dataframe to hold the lower triangle because the matrix is symmetrical

**Author(s)**

Greg Gloor, Jean Macklaim, Wallace Chan

**References**

Please use the citation given by `citation(package="CoDaSeq")`

**See Also**

[codaSeq.clr](#), [codaSeq.filter](#), [codaSeq.rarefy](#), [codaSeq.outlier](#), [codaSeq.propr.phismy](#)

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propr.phisym	<i>Symmetric Phi Statistics</i>
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## Description

Returns a matrix where element (i,j) is the symmetric phi statistic between columns i and j of X.

## Usage

```
propr.phisym <- function (X)
```

## Arguments

X                      A matrix or data frame of centered log ratio transformation.

## Details

X should be the result of a centered log-ratio transformation.

## Author(s)

Greg Gloor, Jean Macklai, Wallace Chan

## References

Please use the citation given by `citation(package="CoDaSeq")`

## See Also

[codaSeq.clr](#), [codaSeq.filter](#), [codaSeq.outlier](#), [codaSeq.rarefy](#), [codaSeq.propr.aldex.phi](#)

## Examples

```
N <- 10
# Number of observations
# Make a data frame with columns a and b roughly proportional
# and columns c and d roughly proportional
X <- data.frame(a=(1:N), b=(1:N) * rnorm(N, 10, 0.1),
                c=(N:1), d=(N:1) * rnorm(N, 10, 1.0))
round(propr.phisym(clr(X)),2)
```

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