

# Package ‘mpress’

November 7, 2018

**Type** Package

**Title** Microbiome Power Estimation with Sampling and Simulation

**Version** 0.9.2

**Description** Given a phyloseq representation of a microbiome complete with metadata, MPrESS returns the estimated power to detect differences between samples with different metadata values either by sampling or by simulation if there are insufficient samples.

**Depends** R(>= 3.4.0), ggplot2, phytools, phyloseq(>= 1.16), vegan, HMP, ape, DESeq2

**License** GPL-2

**LazyData** true

**RoxygenNote** 6.0.1

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mpress-class	<i>An S4 class representing the MPrESS data and output</i>
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## Description

An S4 class representing the MPrESS data and output

## Slots

- `dist.mst` The distance matrix showing the distances between different genomes
- `phy` The phylogenetic tree in newick format
- `rank` The ranks of the respective genomes with lower getting higher priority in being called as a medoid
- `cluster` A vector giving the numeric cluster ID for each genome

h The threshold variable used to make the clusters

medoids A vector giving the medoid for each cluster

gmm A data.frame containing all the gaussian distributions used to find the threshold when available

gmm.orig A data.frame containing all the gaussian distributions prior to cleaning. Used to recalculate the threshold when needed

annotation A data.frame containing the annotations foreach genome with the genomes as the rows and the annotations with the columns. The

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plot.mpress	<i>plot.mpress makes a ggplot with the p-value and power for all the samples investigated an mpress object</i>
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## Description

plot.mpress makes a ggplot with the p-value and power for all the samples investigated an mpress object

## Usage

```
plot.mpress(x)
```

## Examples

```
library(mpress);
#Loading in the microbiome files
data(ChinaData);
data(SpainData);

#Use summary() to examine the data loaded

#Use plot() to see the plot of the power and p-value in the different sample numbers
plot(spain.ibs.power)
```

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power.est	<i>power.calc</i>
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## Description

power.calc() calculates the number of samples to detect a difference between microbiomes with two different metadata values.

## Usage

```
power.est(in.phyloseq, metadata.var, metadata.vals, start.n = 5,
  alpha = 0.05, beta = 0.95, test.type = "permanova",
  dist.metric = "bray", deseq.val = 0, n.rep = 100, burn.in = 10,
  verbose = T, binom = F, switch.val = 0, seed.val = 0)
```

## Arguments

<code>in.phyloseq</code>	Required. The collection of microbiome samples to use. Required to be a class phyloseq data
<code>metadata.var</code>	Required. The column name in the metadata values that is being compared for the power.
<code>metadata.vals</code>	Required. Vector with the values to be compared.
<code>alpha</code>	Alpha value used to test the microbiome differences. Default is 0.05
<code>beta</code>	Beta value used to test proportion of estimated samples that contain significant differences. Default is 0.95
<code>test.type</code>	String giving the statistic to test the between the microbiome samples. Default is "permanova"
<code>dist.metric</code>	String giving the distance metric to use between the microbiome samples. Default "bray" (Bray-Curtis). Available options can be seen with distanceMethodList.
<code>deseq.val</code>	Value to use for the DESeq2 run to trim the taxa. if the value is 0, use all taxa (Default); if the value is between 0 and 1, all taxa with an FDR below the value will be selected; otherwise, if the value is $\geq 1$ , that number of top taxa by FDR will be selected
<code>n.rep</code>	Maximum number of replicates to run before final check if above the beta threshold. Default = 100.
<code>burn.in</code>	Value to burning in for the binomial test. Default is 10
<code>verbose</code>	Flag to prints intermediate values to the screen. Default = F
<code>binom</code>	Flag to use the binomial test to stop low mapping values. Default = F
<code>switch.val</code>	sample number at which to automatically swap between sampling and simulation. Default = 0, or only swap once sample number exceeds the number of samples
<code>seed.val</code>	Value to set seed value for troubleshooting. Default = 0 (not set)
<code>start.</code>	The sample size at which to start the tests. Default is 5.

## Value

Returns a class MPrESS variable with the sample number in addition to runtime information for use in verification

## Examples

```
#The following data is from Chavda et al 2016 which phylotyped Enterobacter genomes

#Additional printing and plotting options are available with plot() and print(). For more
#The following data is from Zhang et al 2015 using Microbiome data from multiple sites in
# Our example uses the data underpinning the tree shown in Figure 2

library(mpress);
#Loading in the Chinese metadata file
data(ChinaData);
data(SpainData);

#Use summary() to examine the data loaded
```

```
summary(china.power)

#Other microbiomes: china.trim from data(ChinaData): unnamed OTUs removed
#                      spain.ibs.trim from

#Use plot() to see the plot of the power and p-value in the different sample numbers
plot(china.power)
```

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print.mpress	<i>print.mpress makes a table with the runtime information for the samples investigated an mpress object</i>
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### Description

print.mpress makes a table with the runtime information for the samples investigated an mpress object

### Usage

```
## S3 method for class 'mpress'
print(x)
```

### Examples

```
library(mpress);
#Loading in the microbiome files
data(ChinaData);
data(SpainData);

#Use summary() to examine the data loaded

#Use print() to see the table of the run info in the different sample numbers
print(spain.ibs.power)
```

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summary.mpress	<i>summary.mpress writes the power value and the estimation type to the screen</i>
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### Description

summary.mpress writes the power value and the estimation type to the screen

### Usage

```
## S3 method for class 'mpress'
summary(x)
```

**Examples**

```
library(mpress);  
#Loading in the microbiome files  
data(ChinaData);  
data(SpainData);  
  
summary(china.trim.power)
```

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