

# Package ‘PathoStat’

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**Type** Package

**Title** PathoStat Statistical Microbiome Analysis Package

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**Description** The purpose of this package is to perform Statistical Microbiome Analysis on metagenomics results from sequencing data samples. In particular, it supports analyses on the PathoScope generated report files. PathoStat provides various functionalities including Relative Abundance charts, Diversity estimates and plots, tests of Differential Abundance, Time Series visualization, and Core OTU analysis.

**URL** <https://github.com/mani2012/PathoStat>

**BugReports** <https://github.com/mani2012/PathoStat/issues>

**License** GPL (>= 2)

**Depends** R (>= 3.4)

**Imports** limma, corpcor, matrixStats, reshape2, scales, ggplot2, rentrez, DT, tidyr, plyr, dplyr, phyloseq, shiny, stats, methods, XML, graphics, utils, BiocStyle, edgeR, DESeq2, ComplexHeatmap, plotly, webshot, vegan, shinyjs, glmnet, gmodels, ROCR, RColorBrewer, knitr, devtools, ape

**Collate** 'pathoStat.R' 'utils.R' 'taxonomy.R' 'biomarker.R'  
'allClasses.R' 'visualization.R' 'differentialAnalysis.R'

**biocViews** Microbiome, Metagenomics, GraphAndNetwork, Microarray, PatternLogic, PrincipalComponent, Sequencing, Software, Visualization, RNASeq

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**VignetteBuilder knitr****git\_url** <https://git.bioconductor.org/packages/PathoStat>**git\_branch** RELEASE\_3\_7**git\_last\_commit** 661f5e8**git\_last\_commit\_date** 2018-06-02**Date/Publication** 2018-10-15**R topics documented:**

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---

Bootstrap\_LOOCV\_LR\_AUC

*Do bootstrap and LOOCV*

---

### Description

Do bootstrap and LOOCV

### Usage

```
Bootstrap_LOOCV_LR_AUC(df, targetVec, nboot = 50)
```

### Arguments

df	Row is sample, column is feature. Required
targetVec	y vector. Required
nboot	number of BOOTSTRAP

### Value

bootstrap loocv result dataframe

### Examples

```
data('iris')
Bootstrap_LOOCV_LR_AUC(iris[,1:4],
c(rep(1,100), rep(0,50)), nboot = 3)
```

Chisq\_Test\_Pam

*Given PAM and disease/control annotation, do Chi-square test for each row of PAM*

---

### Description

Given PAM and disease/control annotation, do Chi-square test for each row of PAM

### Usage

```
Chisq_Test_Pam(pam, label.vec.num, pvalue.cutoff = 0.05)
```

### Arguments

pam	Input data object that contains the data to be tested. Required
label.vec.num	The target binary condition. Required
pvalue.cutoff	choose p-value cut-off

### Value

df.output object

**Examples**

```
tmp <- matrix(rbinom(12,1,0.5), nrow = 3)
rownames(tmp) <- c("a", "b", "c")
Chisq_Test_Pam(tmp, c(1,1,0,0))
```

**findRAfromCount***Return the Relative Abundance (RA) data for the given count OTU table***Description**

Return the Relative Abundance (RA) data for the given count OTU table

**Usage**

```
findRAfromCount(count_otu)
```

**Arguments**

count_otu	Count OTU table
-----------	-----------------

**Value**

ra_otu	Relative Abundance (RA) OTU table
--------	-----------------------------------

**Examples**

```
data_dir <- system.file("data", package = "PathoStat")
infileName <- "pstat_data.rda"
pstat_test <- loadPstat(data_dir, infileName)
ra_otu <- findRAfromCount(phyloseq::otu_table(pstat_test))
```

**findTaxonMat***Find the Taxonomy Information Matrix***Description**

Find the Taxonomy Information Matrix

**Usage**

```
findTaxonMat(names, taxonLevels)
```

**Arguments**

names	Row names of the taxonomy matrix
taxonLevels	Taxon Levels of all tids

**Value**

taxmat	Taxonomy Information Matrix
--------	-----------------------------

**Examples**

```
example_data_dir <- system.file("example/data", package = "PathoStat")
pathoreport_file_suffix <- "-sam-report.tsv"
datlist <- readPathoscopeData(example_data_dir, pathoreport_file_suffix)
dat <- datlist$data
ids <- rownames(dat)
tids <- unlist(lapply(ids, FUN = grepTid))
taxonLevels <- findTaxonomy(tids[1:5])
taxmat <- findTaxonMat(ids[1:5], taxonLevels)
```

findTaxonomy

*Find the taxonomy for each taxon ids***Description**

Find the taxonomy for each taxon ids

**Usage**

```
findTaxonomy(tids)
```

**Arguments**

tids	Given taxonomy ids
------	--------------------

**Value**

taxondata Data with the taxonomy information

**Examples**

```
example_data_dir <- system.file("example/data", package = "PathoStat")
pathoreport_file_suffix <- "-sam-report.tsv"
datlist <- readPathoscopeData(example_data_dir, pathoreport_file_suffix)
dat <- datlist$data
ids <- rownames(dat)
tids <- unlist(lapply(ids, FUN = grepTid))
taxonLevels <- findTaxonomy(tids[1:5])
```

Fisher\_Test\_Pam

*Given PAM and disease/control annotation, do Chi-square test for each row of PAM***Description**

Given PAM and disease/control annotation, do Chi-square test for each row of PAM

**Usage**

```
Fisher_Test_Pam(pam, label.vec.num, pvalue.cutoff = 0.05)
```

**Arguments**

<code>pam</code>	Input data object that contains the data to be tested. Required
<code>label.vec.num</code>	The target binary condition. Required
<code>pvalue.cutoff</code>	choose p-value cut-off

**Value**

`df.output` object

**Examples**

```
tmp <- matrix(rbinom(12,1,0.5), nrow = 3)
rownames(tmp) <- c("a", "b", "c")
Fisher_Test_Pam(tmp, c(1,1,0,0))
```

`formatTaxTable`      *Format taxonomy table for rendering*

**Description**

Format taxonomy table for rendering

**Usage**

```
formatTaxTable(ttable)
```

**Arguments**

<code>ttable</code>	Taxonomy table
---------------------	----------------

**Value**

Formatted table suitable for rendering with. `DT::renderDataTable`

`getShinyInput`      *Getter function to get the shinyInput option*

**Description**

Getter function to get the `shinyInput` option

**Usage**

```
getShinyInput()
```

**Value**

`shinyInput` option

**Examples**

```
getShinyInput()
```

---

`getShinyInputCombat`     *Getter function to get the shinyInputCombat option*

---

### Description

Getter function to get the shinyInputCombat option

### Usage

```
getShinyInputCombat()
```

### Value

shinyInputCombat option

### Examples

```
getShinyInputCombat()
```

---

`getShinyInputOrig`     *Getter function to get the shinyInputOrig option*

---

### Description

Getter function to get the shinyInputOrig option

### Usage

```
getShinyInputOrig()
```

### Value

shinyInputOrig option

### Examples

```
getShinyInputOrig()
```

`getSignatureFromMultipleGlmnet`  
*Use Lasso to do feature selection*

### Description

Use Lasso to do feature selection

### Usage

```
getSignatureFromMultipleGlmnet(df.input, target.vec, nfolds = 10,
                               logisticRegression = TRUE, nRun = 100, alpha = 1)
```

### Arguments

<code>df.input</code>	Row is sample, column is feature. Required
<code>target.vec</code>	y vector. Required
<code>nfolds</code>	glmnet CV nfolds
<code>logisticRegression</code>	doing logistic regression or linear regression.
<code>nRun</code>	number of glmnet runs
<code>alpha</code>	same as in glmnet

### Value

`signature`

### Examples

```
data('iris')
getSignatureFromMultipleGlmnet(iris[,1:4],
                               c(rep(1,100), rep(0,50)), nfolds = 3, nRun = 10)
```

`GET_PAM` *transform cpm counts to presence-absence matrix*

### Description

transform cpm counts to presence-absence matrix

### Usage

```
GET_PAM(df)
```

### Arguments

<code>df</code>	Input data object that contains the data to be tested. Required
-----------------	---

**Value**

df.output object

**Examples**

```
GET_PAM(data.frame(a = c(1,3,0), b = c(0,0.1,2)))
```

---

grepTid

*Greps the tid from the given identifier string*

---

**Description**

Greps the tid from the given identifier string

**Usage**

```
grepTid(id)
```

**Arguments**

id	Given identifier string
----	-------------------------

**Value**

tid string

**Examples**

```
grepTid("ti|700015|org|Coriobacterium_glomerans_PW2")
```

---

loadPathoscopeReports *Loads all data from a set of PathoID reports. For each column in the PathoID report, construct a matrix where the rows are genomes and the columns are samples. Returns a list where each element is named according to the PathoID column. For example, ret[["Final.Best.Hit.Read.Numbers"]]* on the result of this function will get you the final count matrix. Also includes elements "total\_reads" and "total\_genomes" from the first line of the PathoID report.

---

**Description**

Loads all data from a set of PathoID reports. For each column in the PathoID report, construct a matrix where the rows are genomes and the columns are samples. Returns a list where each element is named according to the PathoID column. For example, ret[["Final.Best.Hit.Read.Numbers"]] on the result of this function will get you the final count matrix. Also includes elements "total\_reads" and "total\_genomes" from the first line of the PathoID report.

**Usage**

```
loadPathoscopeReports(reportfiles, nrows = NULL)
```

**Arguments**

reportfiles	Paths to report files
nrows	Option to read first N rows of PathoScope reports

**Value**

Returns a list where each element is named according to the PathoID column. For example, `ret[["Final.Best.Hit.Read.Numbers"]]` on the result of this function will get you the final count matrix. Also includes elements "total\_reads" and "total\_genomes" from the first line of the PathoID report.

**Examples**

```
input_dir <- system.file("example/data", package = "PathoStat")
reportfiles <- list.files(input_dir, pattern = "*-sam-report.tsv",
full.names = TRUE)
```

---

<b>loadPstat</b>	<i>Load the R data(.rda) file with pathostat object</i>
------------------	---

---

**Description**

Load the R data(.rda) file with pathostat object

**Usage**

```
loadPstat(indir = ".", inFile = "pstat_data.rda")
```

**Arguments**

indir	Input Directory of the .rda file
inFile	File name of the .rda file

**Value**

pstat pathostat object (NULL if it does not exist)

**Examples**

```
data_dir <- system.file("data", package = "PathoStat")
inFile <- "pstat_data.rda"
pstat <- loadPstat(data_dir, inFile)
```

<code>log2CPM</code>	<i>Compute log2(counts per mil reads) and library size for each sample</i>
----------------------	--

### Description

Compute log2(counts per mil reads) and library size for each sample

### Usage

```
log2CPM(qcounts, lib.size = NULL)
```

### Arguments

<code>qcounts</code>	quantile normalized counts
<code>lib.size</code>	default is colsums( <code>qcounts</code> )

### Value

list containing log2(quantile counts per mil reads) and library sizes

### Examples

```
log2CPM(matrix(1:12, nrow = 3))
```

<code>LOOAUC_simple_multiple_noplot_one_df</code>	<i>LOOCV</i>
---	--------------

### Description

LOOCV

### Usage

```
LOOAUC_simple_multiple_noplot_one_df(df, targetVec)
```

### Arguments

<code>df</code>	Row is sample, column is feature. Required
<code>targetVec</code>	y vector. Required

### Value

mean auc

### Examples

```
data('iris')
LOOAUC_simple_multiple_noplot_one_df(iris[,1:4],
c(rep(1,100), rep(0,50)))
```

`LOOAUC_simple_multiple_one_df`  
*LOOCV with ROC curve*

### Description

LOOCV with ROC curve

### Usage

```
LOOAUC_simple_multiple_one_df(df, targetVec)
```

### Arguments

<code>df</code>	Row is sample, column is feature. Required
<code>targetVec</code>	y vector. Required

### Value

the ROC

### Examples

```
data('iris')
LOOAUC_simple_multiple_one_df(iris[,1:4],
c(rep(1,100), rep(0,50)))
```

### Description

Contains all currently-supported BatchQC output data classes:

### Details

slots:

- average\_count** a single object of class otu\_tableOrNULL
- besthit\_count** a single object of class otu\_tableOrNULL
- highconf\_count** a single object of class otu\_tableOrNULL
- lowconf\_count** a single object of class otu\_tableOrNULL

---

percent	<i>Compute percentage</i>
---------	---------------------------

---

**Description**

Compute percentage

**Usage**

```
percent(x, digits = 2, format = "f")
```

**Arguments**

x	a number or a vector
digits	how many digit of percentage
format	numeric format, "f" for float

**Value**

the percentage

**Examples**

```
percent.vec <- percent(c(0.9, 0.98))
```

---

phyloseq_to_edgeR	<i>Convert phyloseq OTU count data into DGEList for edgeR package</i>
-------------------	---

---

**Description**

Further details.

**Usage**

```
phyloseq_to_edgeR(physeq, group, method = "RLE", ...)
```

**Arguments**

physeq	(Required).
group	(Required). A character vector or factor giving the experimental group/condition for each sample/library.
method	(Optional).
...	Additional arguments passed on to

**Value**

dispersion

## Examples

```
data_dir_test <- system.file("data", package = "PathoStat")
pstat_test <- loadPstat(indir=data_dir_test,
                        inFileNamE="pstat_data_2_L1.rda")
phyloseq_to_edgeR(pstat_test, group="Sex")
```

**plotHeatmapColor**      *Plot heatmap with color bar*

## Description

Plot heatmap with color bar

## Usage

```
plotHeatmapColor(df.input, condition.vec.1, condition.vec.2, condition.1.name,
                 condition.2.name, do.scale = TRUE, clusterRow = TRUE, clusterCol = TRUE,
                 displayRowLabels = TRUE, displayColumnLabels = TRUE,
                 displayRowDendograms = TRUE, displayColumnDendograms = TRUE,
                 annotationColors = "auto", columnTitle = "Title")
```

## Arguments

<b>df.input</b>	Input data object that contains the data to be plotted. Required
<b>condition.vec.1</b>	color vector. Required
<b>condition.vec.2</b>	color vector 2. Required
<b>condition.1.name</b>	color vector 1 name. Required
<b>condition.2.name</b>	color vector 2 name. Required
<b>do.scale</b>	whether to do row scaling
<b>clusterRow</b>	Cluster the rows. The default is TRUE
<b>clusterCol</b>	Cluster the columns. The default is TRUE
<b>displayRowLabels</b>	Display the row labels on the heatmap. The default is TRUE.
<b>displayColumnLabels</b>	Display the column labels on the heatmap. The default is TRUE
<b>displayRowDendograms</b>	Display the row dendograms on the heatmap. The default is TRUE
<b>displayColumnDendograms</b>	Display the column dendograms on the heatmap. The default is TRUE.
<b>annotationColors</b>	Set of annotation colors for color bar. If null, no color bar is shown. If "auto", then colors will be added automatically. The default is "auto".
<b>columnTitle</b>	Title to be displayed at top of heatmap.

**Value**

ComplexHeatmap object

**Examples**

```
data('iris')
plotHeatmapColor(iris[,1:4],
c(rep(1,100), rep(0,50)),c(rep(0,100), rep(1,50)),
"condition.1.name", "condition.1.name")
```

---

plotPCAPlotly

*Plot PCA*

---

**Description**

Plot PCA

**Usage**

```
plotPCAPlotly(df.input, condition.color.vec,
  condition.color.name = "condition", condition.shape.vec,
  condition.shape.name = "condition", columnTitle = "Title", pc.a = "PC1",
  pc.b = "PC2")
```

**Arguments**

df.input        Input data object that contains the data to be plotted. Required  
condition.color.vec  
                  color vector. Required  
condition.color.name  
                  color variable name. Required  
condition.shape.vec  
                  shape vector. Required  
condition.shape.name  
                  shape variable name. Required  
columnTitle     Title to be displayed at top of heatmap.  
pc.a            pc.1  
pc.b            pc.2

**Value**

the plot

**Examples**

```
data('iris')
plotPCAPlotly(t(iris[,1:4]),
condition.color.vec = c(rep(1,100), rep(0,50)),
condition.shape.vec = c(rep(0,100), rep(1,50)))
```

**plotPCoAPlotly**      *Plot PCoA*

## Description

Plot PCoA

## Usage

```
plotPCoAPlotly(physeq.input, condition.color.vec,
  condition.color.name = "condition", condition.shape.vec,
  condition.shape.name = "condition", method = "bray",
  columnTitle = "Title", pc.a = "Axis.1", pc.b = "Axis.2")
```

## Arguments

physeq.input	Input data object that contains the data to be plotted. Required
condition.color.vec	color vector. Required
condition.color.name	color variable name. Required
condition.shape.vec	shape vector. Required
condition.shape.name	shape variable name. Required
method	which distance metric
columnTitle	Title to be displayed at top of heatmap.
pc.a	pc.1
pc.b	pc.2

## Value

the plot

## Examples

```
data_dir_test <- system.file("data", package = "PathoStat")
pstat_test <- loadPstat(indir=data_dir_test,
  inFileNam="pstat_data_2_L1.rda")
plotPCoAPlotly(pstat_test, condition.color.vec = rbinom(33,1,0.5),
  condition.shape.vec = rbinom(33,1,0.5))
```

---

pstat_data	<i>pathostat object generated from example pathoscope report files</i>
------------	--

---

## Description

This example data consists of 33 samples from a diet study with 11 subjects taking 3 different diets in random order

## Usage

```
pstat
```

## Format

pathostat object extension of phyloseq-class experiment-level object:

**otu\_table** OTU table with 41 taxa and 33 samples

**sample\_data** Sample Data with 33 samples by 18 sample variables

**tax\_table** Taxonomy Table with 41 taxa by 9 taxonomic ranks

**sample\_data** Phylogenetic Tree with 41 tips and 40 internal nodes

## Value

pathostat object

---

readPathoscopeData	<i>Reads the data from PathoScope reports and returns a list of final guess relative abundance and count data</i>
--------------------	---

---

## Description

Reads the data from PathoScope reports and returns a list of final guess relative abundance and count data

## Usage

```
readPathoscopeData(input_dir = ".",
  pathoreport_file_suffix = "-sam-report.tsv", use.input.files = FALSE,
  input.files.path.vec = NULL, input.files.name.vec = NULL)
```

## Arguments

**input\_dir** Directory where the tsv files from PathoScope are located

**pathoreport\_file\_suffix**

PathoScope report files suffix

**use.input.files**

whether input dir to pathoscope files or directly pathoscope files

**input.files.path.vec**

vector of pathoscope file paths

**input.files.name.vec**

vector of pathoscope file names

**Value**

List of final guess relative abundance and count data

**Examples**

```
example_data_dir <- system.file("example/data", package = "PathoStat")
```

---

**runPathoStat**

*Statistical Microbiome Analysis on the pathostat input and generates a html report and produces interactive shiny app plots*

---

**Description**

Statistical Microbiome Analysis on the pathostat input and generates a html report and produces interactive shiny app plots

**Usage**

```
runPathoStat(pstat = NULL, report_dir = ".",
             report_option_binary = "111111111", interactive = TRUE)
```

**Arguments**

pstat	phyloseq extension pathostat object
report_dir	Output report directory path
report_option_binary	9 bits Binary String representing the plots to display and hide in the report
interactive	when TRUE, opens the interactive shinyApp

**Value**

outputfile The output file with all the statistical plots

**Examples**

```
runPathoStat(interactive = FALSE)
```

---

savePstat	<i>Save the pathostat object to R data(.rda) file</i>
-----------	---

---

**Description**

Save the pathostat object to R data(.rda) file

**Usage**

```
savePstat(pstat, outdir = ".", outfileName = "pstat_data.rda")
```

**Arguments**

pstat	pathostat object
outdir	Output Directory of the .rda file
outfileName	File name of the .rda file

**Value**

outfile .rda file

**Examples**

```
data_dir_test <- system.file("data", package = "PathoStat")
pstat_test <- loadPstat(indir=data_dir_test,
                        inFileNam="pstat_data_2_L1.rda")
outfile <- savePstat(pstat_test)
```

---

setShinyInput	<i>Setter function to set the shinyInput option</i>
---------------	---

---

**Description**

Setter function to set the shinyInput option

**Usage**

```
setShinyInput(x)
```

**Arguments**

x	shinyInput option
---	-------------------

**Value**

shinyInput option

**Examples**

```
setShinyInput(NULL)
```

---

`setShinyInputCombat`     *Setter function to set the shinyInputCombat option*

---

## Description

Setter function to set the shinyInputCombat option

## Usage

```
setShinyInputCombat(x)
```

## Arguments

x                shinyInputCombat option

## Value

shinyInputCombat option

## Examples

```
setShinyInputCombat(NULL)
```

---

`setShinyInputOrig`     *Setter function to set the shinyInputOrig option*

---

## Description

Setter function to set the shinyInputOrig option

## Usage

```
setShinyInputOrig(x)
```

## Arguments

x                shinyInputOrig option

## Value

shinyInputOrig option

## Examples

```
setShinyInputOrig(NULL)
```

---

summarizeTable	<i>Summarize sample</i>
----------------	-------------------------

---

## Description

Creates a table of summary metrics

## Usage

```
summarizeTable(pstat)
```

## Arguments

pstat	Input pstat
-------	-------------

## Value

A data.frame object of summary metrics.

## Examples

```
data_dir_test <- system.file("data", package = "PathoStat")
pstat_test <- loadPstat(indir=data_dir_test,
  inFileNam="pstat_data_2_L1.rda")
st.tmp <- summarizeTable(pstat_test)
```

---

---

TranslateIdToTaxLevel	<i>Find the taxonomy for the given taxon id name</i>
-----------------------	--

---

## Description

Find the taxonomy for the given taxon id name

## Usage

```
TranslateIdToTaxLevel(pstat, input.id.vec, tax.level)
```

## Arguments

pstat	pathostat object
input.id.vec	names containing id
tax.level	target taxon level

## Value

target taxon level names

## Examples

```
data_dir_test <- system.file("data", package = "PathoStat")
pstat_test <- loadPstat(indir=data_dir_test,
  infilename="pstat_data_2_L1.rda")
names.new <- TranslateIdToTaxLevel(pstat_test,
  c("ti|862962|org|Bacteroides_fragilis_638R",
  "ti|697329|org|Ruminococcus_albus_7" ),
  "genus")
```

Wilcox_Test_df	<i>Mann-whitney test for a dataframe</i>
----------------	--

## Description

Mann-whitney test for a dataframe

## Usage

```
Wilcox_Test_df(df, label.vec.num, pvalue.cutoff = 0.05)
```

## Arguments

df	Input data object that contains the data to be tested. Required
label.vec.num	The target binary condition. Required
pvalue.cutoff	choose p-value cut-off

## Value

df.output object

## Examples

```
data('iris')
Wilcox_Test_df(t(iris[,1:4]),
  c(rep(1,100), rep(0,50)))
```

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