

# Package: pctax (via r-universe)

October 23, 2024

**Type** Package

**Title** Professional Comprehensive Omics Data Analysis

**Version** 0.1.3

**Description** Provides a comprehensive suite of tools for analyzing omics data. It includes functionalities for alpha diversity analysis, beta diversity analysis, differential abundance analysis, community assembly analysis, visualization of phylogenetic tree, and functional enrichment analysis. With a progressive approach, the package offers a range of analysis methods to explore and understand the complex communities. It is designed to support researchers and practitioners in conducting in-depth and professional omics data analysis.

**License** GPL-3

**Encoding** UTF-8

**Roxygen** list(markdown = TRUE)

**RoxygenNote** 7.2.3

**Depends** R (>= 4.2.0)

**LazyData** true

**Imports** pcutils (>= 0.2.5), dplyr, ggplot2 (>= 3.2.0), vegan, magrittr, grDevices, RColorBrewer, ggrepel, reshape2, stats, tibble, utils, ggpubr, ggnewscale, ade4, scales

**Suggests** picante, NST, permute, aplot, pheatmap, MASS, Rtsne, mixOmics, geosphere, phyloseq, phyloseqGraphTest, plotly, umap, Hmisc, minpack.lm, bbmle, snow, foreach, doSNOW, patchwork, tidytree, ggtree, ggtreeExtra, vctrs, zoo, ape, DESeq2, limma, ALDEx2, Mfuzz, edgeR, methods, randomForest, knitr, rmarkdown, MetaNet, showtext, jsonlite, prettydoc, readxl, clipr, zetadiv, ggforce, AnnotationDbi, org.Hs.eg.db, linkET, pairwiseAdonis, sankeyD3, Biobase, GEOquery

**VignetteBuilder** knitr

**BugReports** <https://github.com/Asa12138/pctax/issues>

**URL** <https://github.com/Asa12138/pctax>

**ByteCompile** true

**biocViews** Microbiome, Software, Visualization

**Repository** <https://asa12138.r-universe.dev>

**RemoteUrl** <https://github.com/asa12138/pctax>

**RemoteRef** HEAD

**RemoteSha** 11e21a768d7067657927ea7fb6c46869518e820f

## Contents

add_strip . . . . .	4
add_tax . . . . .	4
ALDEX . . . . .	5
all_ec_info . . . . .	6
ann_tree . . . . .	6
aor . . . . .	7
as.b_dist . . . . .	9
as.dist.b_dist . . . . .	10
a_diversity . . . . .	10
bbtt . . . . .	11
before_tree . . . . .	12
b_analyse . . . . .	12
b_NTII . . . . .	14
b_res_3d . . . . .	14
check_taxonkit . . . . .	15
cor_net . . . . .	16
df2tree . . . . .	16
df2tree1 . . . . .	17
diff_da . . . . .	17
download_taxonkit_dataset . . . . .	18
envfitt . . . . .	19
gene2id . . . . .	19
geo_sim . . . . .	20
get_diff_type . . . . .	21
gp_dis_density . . . . .	21
grap_p_test . . . . .	22
install_taxonkit . . . . .	23
kwtest . . . . .	23
load_N_data . . . . .	24
mat_dist . . . . .	24
micro_sbatch . . . . .	25
multi_bar . . . . .	26
myRDA . . . . .	27
m_group_env . . . . .	28
name_or_id2df . . . . .	29
ncm . . . . .	30

nst . . . . .	31
nti_rc . . . . .	32
pc_otu . . . . .	33
pc_tax1 . . . . .	34
pc_valid . . . . .	34
permanova . . . . .	35
plot.a_res . . . . .	36
plot.b_res . . . . .	37
plot.g_test . . . . .	38
plot.pro_res . . . . .	39
plot.time_cm . . . . .	39
plot_element_cycle . . . . .	40
plot_N_cycle . . . . .	41
plot_two_tree . . . . .	42
pre_fastp . . . . .	44
pre_GEO . . . . .	44
pre_tax_table . . . . .	45
print.pc_otu . . . . .	46
procrustes_analyse . . . . .	46
rarefaction . . . . .	47
rare_curve_sample . . . . .	47
rare_curve_species . . . . .	48
RCbray1 . . . . .	49
RDA_plot . . . . .	50
sangji_plot . . . . .	52
suijisenlin . . . . .	53
summary.pc_otu . . . . .	53
sunburst . . . . .	54
taxonkit_filter . . . . .	55
taxonkit_lca . . . . .	56
taxonkit_lineage . . . . .	57
taxonkit_list . . . . .	58
taxonkit_name2taxid . . . . .	59
taxonkit_reformat . . . . .	60
tax_lca . . . . .	62
time_by_cm . . . . .	63
volcano_p . . . . .	64
z_diversity . . . . .	65
z_diversity_decay . . . . .	66

---

add_strip	<i>add strips for a tree plot</i>
-----------	-----------------------------------

---

**Description**

add strips for a tree plot

**Usage**

```
add_strip(trp, some_tax, flat_n = 5, strip_params = NULL)
```

**Arguments**

trp	tree plot from ggtree
some_tax	some tax you want to add strip
flat_n	flat the text when taxa number more than flat_n.
strip_params	parameters parse to <a href="#">geom_strip</a>

**Value**

tree plot

**Examples**

```
data(otutab, package = "pcutils")
# run yourself
if (interactive()) {
  ann_tree(taxonomy, otutab) -> tree
  easy_tree(tree) -> p
  some_tax <- table(taxonomy$Phylum) %>%
    sort(decreasing = TRUE) %>%
    head(5) %>%
    names()
  add_strip(p, some_tax)
}
```

---

add_tax	<i>Add taxonomy for a pc_otu object</i>
---------	---

---

**Description**

Add taxonomy for a pc\_otu object

**Usage**

```
add_tax(pc, taxonomy)
```

**Arguments**

pc a pc\_otu object  
 taxonomy a taxomomy data.frame, look out the rownames of taxonomy and otutab should matched!

**Value**

pc\_otu

**Examples**

```
data(otutab, package = "pcutils")
pc_tax1 <- pc_otu(otutab, metadata)
pc_tax1 <- add_tax(pc_tax1, taxonomy)
```

---

ALDEX

*ALDEX*

---

**Description**

ALDEX

**Usage**

```
ALDEX(otutab, group_df)
```

**Arguments**

otutab otutab  
 group\_df a dataframe with rowname same to dist and one group column

**Value**

diff

**References**

<https://cloud.tencent.com/developer/article/1621879>

**Examples**

```
if (requireNamespace("ALDEX2")) {
  data(otutab, package = "pcutils")
  ALDEX(otutab, metadata["Group"]) -> res
  res %>%
    dplyr::top_n(9, -glm.eBH) %>%
    .[, "tax"] -> sig
  data.frame(t(otutab[sig, ])) %>% pcutils::group_box(., "Group", metadata)
}
```

---

all_ec_info	<i>all element cycle information.</i>
-------------	---------------------------------------

---

**Description**

all element cycle information.

**Format**

a list contains four tables.

**ec\_node** chemicals

**ec\_link** reactions

**ec\_gene** genes

**ec\_path** reactions labels

---

ann_tree	<i>Annotate a tree</i>
----------	------------------------

---

**Description**

Annotate a tree

Easy way to plot a phylogenetic tree

**Usage**

```
ann_tree(f_tax, otutab = NULL, level = ncol(f_tax))
```

```
easy_tree(  
  tree,  
  highlight = "Phylum",  
  colorfill = "color",  
  topN = NULL,  
  pal = NULL,  
  name_prefix = FALSE,  
  basic_params = NULL,  
  add_abundance = TRUE,  
  color_name = "abundance",  
  add_tiplab = TRUE,  
  fontsize = NULL  
)
```

**Arguments**

f_tax	taxonomy dataframe
otutab	otutab, rowname==rowname(taxonomy)
level	1~7
tree	result from ann_tree
highlight	highlight which level, one of tree\$level
colorfill	"color" or "fill"
topN	topN to show
pal	color pal
name_prefix	keep the prefix like "k_" or "p_" in the label? Default: FALSE
basic_params	parameters parse to <a href="#">ggtree</a>
add_abundance	logical
color_name	color name
add_tiplab	logical
fontsize	tip label fontsize

**Value**

a treedata  
a ggplot

**Examples**

```
if (interactive()) {
  data(otutab, package = "pcutils")
  ann_tree(taxonomy, otutab) -> tree
  # run yourself
  easy_tree(tree, add_abundance = FALSE) -> p
  p
}
```

---

aor

*Calculate Abundance-occupancy\_relationship*


---

**Description**

Calculate Abundance-occupancy\_relationship  
Plot a AOR

**Usage**

```
aor(otutab, ...)  
  
## S3 method for class 'data.frame'  
aor(  
  otutab,  
  top_r = 0.7,  
  ocup_n = ceiling(0.8 * ncol(otutab)),  
  special_n = ceiling(0.1 * ncol(otutab)),  
  ...  
)  
  
## S3 method for class 'AOR'  
plot(x, ...)
```

**Arguments**

otutab	otutab
...	add
top_r	percentage of top relative abundance
ocup_n	percentage of top occupied
special_n	how many occupancy define as specialists
x	AOR object

**Value**

AOR  
ggplot

**References**

Barberán, A., Bates, S. T., Casamayor, E. & Fierer, N. (2012) Using network analysis to explore co-occurrence patterns in soil microbial communities.

**Examples**

```
data(otutab, package = "pcutils")  
aor(otutab) -> AOR  
plot(AOR)
```



---

as.b_dist	<i>Transfer dist to b_dist</i>
-----------	--------------------------------

---

### Description

Transfer dist to b\_dist

Plot dist

Plot b\_dist

### Usage

```
as.b_dist(dist, group_df = NULL)

## S3 method for class 'dist'
plot(x, group_df = NULL, ...)

## S3 method for class 'b_dist'
plot(x, mode = 1, c_group = "inter", ...)
```

### Arguments

dist	a dist object
group_df	a dataframe with rowname same to dist and one group column
x	a b_dist
...	additional
mode	1~3
c_group	"inter" or "intra" or both to plot

### Value

a b\_dist with annotation by group

a pheatmap

a ggplot or pheatmap

### Examples

```
data(otutab, package = "pcutils")
mat_dist(otutab) %>% as.b_dist(., group_df = metadata["Group"]) -> aa
plot(aa)
plot(aa, mode = 2)
```

---

as.dist.b_dist	<i>Transfer b_dist to dist</i>
----------------	--------------------------------

---

**Description**

Transfer b\_dist to dist

**Usage**

```
## S3 method for class 'b_dist'
as.dist(m, diag = FALSE, upper = FALSE)
```

**Arguments**

m	a b_dist object
diag	logical value indicating whether the diagonal of the distance matrix should be printed by print.dist.
upper	logical value indicating whether the upper triangle of the distance matrix should be printed by print.dist.

**Value**

dist

---

a_diversity	<i>Calculate a_diversity of otutab</i>
-------------	--

---

**Description**

Calculate a\_diversity of otutab

**Usage**

```
a_diversity(otutab, ...)

## S3 method for class 'data.frame'
a_diversity(
  otutab,
  method = c("richness", "shannon"),
  tree = NULL,
  digits = 4,
  ...
)

## S3 method for class 'pc_otu'
```

```
a_diversity(otutab, method = "all", tbl = "otutab", ...)

## S3 method for class 'numeric'
a_diversity(otutab, ...)
```

### Arguments

otutab	numeric
...	pass to <code>a_diversity.data.frame</code>
method	one of "all", "richness", "chao1", "ace", "gc", "shannon", "simpson", "pd", "pielou"
tree	a iphylo object match the rownames of otutab
digits	maintance how many digits
tbl	which table

### Value

a `a_res` object

### Examples

```
data(otutab, package = "pcutils")
a_diversity(otutab) -> a_res
plot(a_res, "Group", metadata)
```

---

bbtt	<i>ggdotchart for diff analysis</i>
------	-------------------------------------

---

### Description

ggdotchart for diff analysis

### Usage

```
bbtt(res, pvalue = "glm.eBH", topN = 20)
```

### Arguments

res	result of ALDEX or kwtest
pvalue	the name of pvaule
topN	topN

### Value

ggplot

---

before_tree	<i>Before df2tree check</i>
-------------	-----------------------------

---

**Description**

Before df2tree check

**Usage**

```
before_tree(f_tax)
```

**Arguments**

f_tax	table
-------	-------

**Value**

table

**Examples**

```
wrong_taxdf <- data.frame(
  kingdom = c(rep(c("A", "B"), each = 4), "C", NA),
  "phylum" = c("A", "a", "b", "c", "c", "c", "d", NA, NA, "e")
)
before_tree(wrong_taxdf)
```

---

b_analyse	<i>Beta_diversity Ordination: dimensionality reduction</i>
-----------	--

---

**Description**

Species abundance data can be preprocessed with Hellinger transformation or chord transformation data before PCA analysis. Because the Hellinger distance or chord distance with-without data is equal to  $\sqrt{2}\sqrt{1 - \text{Ochiai similarity}}$ , therefore, the sorting diagram (type 1 scale) of PCA analysis after Hellinger transformation or chord transformation with-without data is internal sample. The distance between the squares is the Ochiai distance.  $\sqrt{2}\sqrt{1 - \text{Ochiai similarity}}$  is a distance measure, which is also suitable for the analysis of species data. The processed data is then used for pca without norm.

**Usage**

```

b_analyse(otutab, ...)

## S3 method for class 'data.frame'
b_analyse(
  otutab,
  norm = TRUE,
  method = c("pca"),
  group = NULL,
  dist = "bray",
  ndim = 2,
  scale = FALSE,
  ...
)

```

**Arguments**

otutab	an otutab data.frame, samples are columns, taxa are rows.
...	add
norm	should normalized or not? (hellinger)
method	one of "pca", "pcoa", "ca", "dca", "nmds", "plsda", "tsne", "umap", "lda", "all"
group	if needed, give a group vector
dist	if use pcoa or nmds, your can choose a dist method (default: bray) or input a distance matrix.
ndim	how many dimension be kept? (default:2). 3 for b_res_3d()
scale	scale, default: FALSE

**Value**

b\_res object

**References**

<https://www.jianshu.com/p/9694c0b6302d> <https://zhuanlan.zhihu.com/p/25501130>

**Examples**

```

data(otutab, package = "pcutils")
b_analyse(otutab, method = "pca") -> b_res
plot(b_res, "Group", metadata)

```

---

b\_NTI1

*Calculate beta\_NTI*


---

**Description**

Calculate beta\_NTI

**Usage**

```
b_NTI1(  
  phylo,  
  otutab,  
  beta.reps = 9,  
  weighted = TRUE,  
  threads = 1,  
  verbose = TRUE  
)
```

**Arguments**

phylo	a phylo object
otutab	otutab
beta.reps	how many simulation performed?
weighted	logical
threads	use how many threads to calculate (default:4)
verbose	verbose

**Value**

a dist: b\_NTI

---

b\_res\_3d

*3D plot for b\_res*


---

**Description**

3D plot for b\_res

**Usage**

```
b_res_3d(b_res, Group, metadata = NULL, ...)
```

**Arguments**

b_res	a b_res object
Group	group vector for color
metadata	metadata contain Group
...	add

**Value**

plotly list

**Examples**

```
if (requireNamespace("plotly")) {  
  data(otutab, package = "pcutils")  
  b_analyse(otutab, method = "pca", ndim = 3) -> b_res  
  b_res_3d(b_res, "Group", metadata)  
}
```

---

check\_taxonkit

*Check taxonkit*

---

**Description**

Check taxonkit

**Usage**

```
check_taxonkit(print = TRUE)
```

**Arguments**

print	print
-------	-------

**Value**

taxonkit path

**See Also**

Other Rtaxonkit: [download\\_taxonkit\\_dataset\(\)](#), [install\\_taxonkit\(\)](#), [name\\_or\\_id2df\(\)](#), [taxonkit\\_filter\(\)](#), [taxonkit\\_lca\(\)](#), [taxonkit\\_lineage\(\)](#), [taxonkit\\_list\(\)](#), [taxonkit\\_name2taxid\(\)](#), [taxonkit\\_reformat\(\)](#)

---

cor_net	<i>Correlation network, species-interaction network for omics</i>
---------	---

---

**Description**

Correlation network, species-interaction network for omics

**Usage**

```
cor_net()
```

**Value**

No value

---

df2tree	<i>From a dataframe to construct a phylo</i>
---------	--

---

**Description**

NOTE: this function will do before\_tree first.

**Usage**

```
df2tree(data, edge_df = FALSE)
```

**Arguments**

data	dataframe
edge_df	if the data is edge_df ?

**Value**

phylo object

**Examples**

```
data(otutab, package = "pcutils")
df2tree(taxonomy) -> tax_tree
print(tax_tree)
# check all nodes matched!
if (requireNamespace("picante")) {
  picante::match.phylo.comm(tax_tree, t(otutab)) -> nn
  nrow(nn$comm) == nrow(t(otutab))
}
```



---

df2tree1	<i>From a dataframe to construct a phylo (save nwk)</i>
----------	---

---

**Description**

NOTE: this function will transfer all space to \_

**Usage**

```
df2tree1(taxa)
```

**Arguments**

taxa	dataframe
------	-----------

**Value**

phylo object

**Examples**

```
data(otutab, package = "pcutils")
df2tree(taxonomy) -> tax_tree
print(tax_tree)
```

---

diff_da	<i>Difference analysis</i>
---------	----------------------------

---

**Description**

Difference analysis

**Usage**

```
diff_da(  
  otutab,  
  group_df,  
  ctrl = NULL,  
  method = "deseq2",  
  log = TRUE,  
  add_mini = NULL,  
  ...  
)
```

**Arguments**

otutab	otutab
group_df	a dataframe with rowname same to dist and one group column
ctrl	the control group, one level of groups
method	one of "deseq2", "edger", "limma", "t.test", "wilcox.test"
log	do log transfer for limma?
add_mini	add_mini when calculate the logFC. e.g (10+0.1)/(0+0.1), default 0.5*min(abundance)
...	other parameters

**Value**

a dataframe

**Examples**

```
if (requireNamespace("limma")) {
  data(otutab, package = "pcutils")
  diff_da(otutab, metadata["Group"], method = "limma") -> res
  volcano_p(res)
  volcano_p(res, mode = 2)
}
```

---

download\_taxonkit\_dataset

*Download taxonkit dataset*

---

**Description**

Download taxonkit dataset

**Usage**

```
download_taxonkit_dataset(make_sure = FALSE, taxdump_tar_gz = NULL)
```

**Arguments**

make_sure	make sure to do this
taxdump_tar_gz	your download taxdump_tar_gz file from <a href="https://ftp.ncbi.nih.gov/pub/taxonomy/taxdump.tar.gz">https://ftp.ncbi.nih.gov/pub/taxonomy/taxdump.tar.gz</a>

**Value**

No value

**See Also**

Other Rtaxonkit: [check\\_taxonkit\(\)](#), [install\\_taxonkit\(\)](#), [name\\_or\\_id2df\(\)](#), [taxonkit\\_filter\(\)](#), [taxonkit\\_lca\(\)](#), [taxonkit\\_lineage\(\)](#), [taxonkit\\_list\(\)](#), [taxonkit\\_name2taxid\(\)](#), [taxonkit\\_reformat\(\)](#)

---

envfitt	<i>Envfit test for RDA result</i>
---------	-----------------------------------

---

**Description**

Envfit test for RDA result

**Usage**

```
envfitt(phy.rda, env, ...)
```

**Arguments**

phy.rda	a rda result
env	environmental factors
...	add

**Value**

g\_test object

**See Also**

[envfit](#)

**Examples**

```
data(otutab, package = "pcutils")
env <- metadata[, 6:10]
# RDA
myRDA(otutab, env) -> phy.rda
envfitt(phy.rda, env) -> envfit_res
plot(envfit_res)
```

---

gene2id	<i>Gene symbolid transfer to entrezIDs (human gene)</i>
---------	---

---

**Description**

Gene symbolid transfer to entrezIDs (human gene)

**Usage**

```
gene2id(genes)
```

**Arguments**

genes                    gene symbols e.g:ASGR2

**Value**

gene entrezIDs dataframe

**Examples**

```
if (requireNamespace("AnnotationDbi") && requireNamespace("org.Hs.eg.db")) {
  genes <- c(
    "ASGR2", "BEST1", "SIGLEC16", "ECRP", "C1QC", "TCN2", "RNASE2",
    "DYSF", "C1QB", "FAM20A", "FCGR1A", "CR1", "HP", "VSIG4", "EGR1"
  )
  gene2id(genes) -> geneid
}
```

---

geo\_sim

*Lm for sample similarity and geographical distance*

---

**Description**

Lm for sample similarity and geographical distance

**Usage**

```
geo_sim(otutab, geo, method = "bray", spe_nwk = NULL, ...)
```

**Arguments**

otutab                    an otutab data.frame, samples are columns, taxa are rows.  
 geo                        a two-columns dataframe, first is latitude, second is longitude  
 method                    Dissimilarity index, partial match to "bray", "euclidean"...see [vegdist](#); [unifrac](#)  
 spe\_nwk                    a phylo tree if use unifrac...  
 ...                        additional

**Value**

a ggplot

**References**

Graco-Roza, C. et al. (2022) Distance decay 2.0 - A global synthesis of taxonomic and functional turnover in ecological communities. *Glob Ecol Biogeogr* 31, 1399–1421.

**Examples**

```

if (requireNamespace("NST") && requireNamespace("geosphere")) {
  library(ggplot2)
  data(otutab, package = "pcutils")
  metadata[, c("lat", "long")] -> geo
  geo_sim(otutab, geo) -> geo_res
  pcutils::my_lm(geo_res[4], "dis.geo", geo_res)
}

```

---

get_diff_type	<i>Get mean and type</i>
---------------	--------------------------

---

**Description**

Get mean and type

**Usage**

```
get_diff_type(otutab, group_df)
```

**Arguments**

otutab	otutab
group_df	a dataframe with rowname same to dist and one group column

**Value**

No value

---

gp_dis_density	<i>Group inter-intra density</i>
----------------	----------------------------------

---

**Description**

Group inter-intra density

**Usage**

```
gp_dis_density(otutab, group)
```

**Arguments**

otutab	an otutab data.frame, samples are columns, taxa are rows.
group	group vector

**Value**

ggplot

**Examples**

```
data(otutab, package = "pcutils")
gp_dis_density(otutab, metadata["Group"])
```

---

grap_p_test	<i>Performs graph-based permutation tests</i>
-------------	---

---

**Description**

Performs graph-based permutation tests

**Usage**

```
grap_p_test(otutab, metadata, group = "Group", nperm = 999, ...)
```

**Arguments**

otutab	an otutab data.frame, samples are columns, taxa are rows.
metadata	metadata
group	one group name in columns of metadata
nperm	numbers of permutations to perform
...	additional

**Value**

ggplot

**Examples**

```
if (requireNamespace("phyloseqGraphTest") && requireNamespace("phyloseq")) {
  data(otutab, package = "pcutils")
  grap_p_test(otutab, metadata, "Group")
}
```

---

install_taxonkit	<i>Install taxonkit</i>
------------------	-------------------------

---

**Description**

Install taxonkit

**Usage**

```
install_taxonkit(make_sure = FALSE, taxonkit_tar_gz = NULL)
```

**Arguments**

make_sure	make sure to do this
taxonkit_tar_gz	your download taxonkit_tar_gz file from <a href="https://github.com/shenwei356/taxonkit/releases/">https://github.com/shenwei356/taxonkit/releases/</a>

**Value**

No value

**See Also**

Other Rtaxonkit: [check\\_taxonkit\(\)](#), [download\\_taxonkit\\_dataset\(\)](#), [name\\_or\\_id2df\(\)](#), [taxonkit\\_filter\(\)](#), [taxonkit\\_lca\(\)](#), [taxonkit\\_lineage\(\)](#), [taxonkit\\_list\(\)](#), [taxonkit\\_name2taxid\(\)](#), [taxonkit\\_reformat\(\)](#)

---

kwtest	<i>KW test</i>
--------	----------------

---

**Description**

KW test

**Usage**

```
kwtest(otutab, group_df, method = "kruskal.test")
```

**Arguments**

otutab	otutab
group_df	a dataframe with rowname same to dist and one group column
method	"kruskal.test", see <a href="#">compare_means</a>

**Value**

res

**Examples**

```
data(otutab, package = "pcutils")
kwtest(otutab, metadata["Group"]) -> res
bbtt(res, pvalue = "p.format")
```

---

load_N_data	<i>Load N-cycle data</i>
-------------	--------------------------

---

**Description**

Load N-cycle data

**Usage**

```
load_N_data()
```

**Value**

list

**References**

Tu, Q., Lin, L., Cheng, L., Deng, Y. & He, Z. (2019) NCycDB: a curated integrative database for fast and accurate metagenomic profiling of nitrogen cycling genes. *Bioinformatics* 35, 1040–1048.  
 Kuypers, M. M. M., Marchant, H. K. & Kartal, B. (2018) The microbial nitrogen-cycling network. *Nat Rev Microbiol* 16, 263–276.

---

mat_dist	<i>Calculate distance for otutab</i>
----------	--------------------------------------

---

**Description**

Calculate distance for otutab

**Usage**

```
mat_dist(otutab, method = "bray", spe_nwk = NULL)
```

**Arguments**

otutab	an otutab data.frame, samples are columns, taxa are rows.
method	Dissimilarity index, partial match to "bray", "euclidean"...see <a href="#">vegdist</a> ; <a href="#">unifrac</a>
spe_nwk	a phylo tree if use unifrac...



**Value**

dist

**Examples**

```
data(otutab, package = "pcutils")
mat_dist(otutab)
```

---

micro_sbatch	<i>Microbiome sbatch</i>
--------------	--------------------------

---

**Description**

Microbiome sbatch

**Usage**

```
micro_sbatch(
  work_dir = "/share/home/jianglab/pengchen/work/asthma/",
  step = "fastp",
  all_sample_num = 40,
  array = 1,
  partition = "cpu",
  cpus_per_task = 1,
  mem_per_cpu = "2G"
)
```

**Arguments**

work_dir	work_dir
step	"fastp", "rm_human", "megahit", "prodigal", "salmon-quant", ...
all_sample_num	all sample number
array	array number
partition	partition
cpus_per_task	cpus_per_task
mem_per_cpu	mem_per_cpu, "2G"

**Value**

No value

---

`multi_bar`*Difference analysis*

---

**Description**

Difference analysis

**Usage**

```
multi_bar(  
  otutab,  
  group_df,  
  mode = 1,  
  text_df = NULL,  
  text_x = NULL,  
  text_angle = -90,  
  errorbar = "bottom"  
)
```

**Arguments**

<code>otutab</code>	<code>otutab</code>
<code>group_df</code>	a dataframe with rowname same to dist and one group column
<code>mode</code>	1~2
<code>text_df</code>	<code>text_df</code>
<code>text_x</code>	<code>text_x</code>
<code>text_angle</code>	<code>text_angle</code>
<code>errorbar</code>	top, bottom, none

**Value**

a data.frame

**Examples**

```
data(otutab, package = "pcutils")  
multi_bar(otutab[1:10, ], metadata["Group"])
```

---

myRDA

*RDA*

---

### Description

RDA

### Usage

```
myRDA(  
  otutab,  
  env,  
  norm = TRUE,  
  scale = FALSE,  
  choose_var = FALSE,  
  direction = "forward",  
  nperm = 499,  
  verbose = TRUE,  
  method = "rda",  
  dist = "bray"  
)
```

```
myCCA(  
  otutab,  
  env,  
  norm = TRUE,  
  scale = FALSE,  
  choose_var = FALSE,  
  nperm = 499,  
  verbose = TRUE  
)
```

```
myCAP(  
  otutab,  
  env,  
  norm = TRUE,  
  scale = FALSE,  
  choose_var = FALSE,  
  nperm = 499,  
  verbose = TRUE,  
  dist = "bray"  
)
```

### Arguments

otutab	an otutab data.frame, samples are columns, taxa are rows.
env	environmental factors

norm	should normalize? (default:TRUE)
scale	should scale species? (default:FALSE)
choose_var	should choose variables? use forward step
direction	The direction of the stepwise selection, "both", "forward" or "backward", default is "forward"
nperm	number of permutation
verbose	verbose
method	"rda", "cca", "cap", "dbrda"
dist	The name of the dissimilarity (or distance) index for "cap" or "dbrda", for <a href="#">vegdist</a>

**Value**

rda/cca

**See Also**

[vegdist](#); [unifrac](#)

**Examples**

```
data(otutab, package = "pcutils")
env <- metadata[, 6:10]
# RDA
myRDA(otutab, env) -> phy.rda
RDA_plot(phy.rda, "Group", metadata)
```

---

m\_group\_env

*Multi-table test with env*

---

**Description**

Multi-table test with env

Plot mant\_g object

**Usage**

```
m_group_env(g_otutab, env)
```

```
## S3 method for class 'mant_g'
```

```
plot(x, ...)
```

**Arguments**

g\_otutab multi-otutabs with first column is group

env environmental factors

x mant\_g object

... add

**Value**

a mant\_g object  
a ggplot

**Examples**

```
if (requireNamespace("linkET")) {
  data(otutab, package = "pcutils")
  cbind(group = rep(c("a", "b", "c"), c(200, 100, 185)), otutab) -> g_otutab
  metadata[, 3:8, drop = FALSE] -> env
  m_group_env(g_otutab, env) -> mant_g
  plot(mant_g)
}
```

---

name\_or\_id2df

*Transfer taxon name or taxid to the lineage dataframe*


---

**Description**

Transfer taxon name or taxid to the lineage dataframe

**Usage**

```
name_or_id2df(
  name_or_id,
  mode = "name",
  add_prefix = TRUE,
  fill_miss_rank = TRUE,
  data_dir = NULL
)
```

**Arguments**

name_or_id	name or taxid
mode	"id" or "name"
add_prefix	add_prefix
fill_miss_rank	fill_miss_rank
data_dir	directory containing nodes.dmp and names.dmp (default "/Users/asa/.taxonkit")

**Value**

dataframe

**See Also**

Other Rtaxonkit: [check\\_taxonkit\(\)](#), [download\\_taxonkit\\_dataset\(\)](#), [install\\_taxonkit\(\)](#), [taxonkit\\_filter\(\)](#), [taxonkit\\_lca\(\)](#), [taxonkit\\_lineage\(\)](#), [taxonkit\\_list\(\)](#), [taxonkit\\_name2taxid\(\)](#), [taxonkit\\_reformat\(\)](#)

**Examples**

```
## Not run:
name_or_id2df(c("Homo sapiens", "Akkermansia muciniphila ATCC BAA-835"))

## End(Not run)
```

ncm

*Sloan Neutral Model***Description**

Sloan Neutral Model  
 Plot ncm\_res

**Usage**

```
ncm(otutab, model = "nls")

## S3 method for class 'ncm_res'
plot(
  x,
  mycols = c(Above = "#069870", Below = "#e29e02", In = "#1e353a"),
  text_position = NULL,
  ...
)
```

**Arguments**

otutab	an otutab data.frame, samples are columns, taxa are rows.
model	fit method, one of "nls", "mle"
x	a ncm_res object
mycols	mycols
text_position	text_position
...	add

**Value**

ncm\_res  
 ggplot

**References**

Sloan, W. TRUE. et al. (2006) Quantifying the roles of immigration and chance in shaping prokaryote community structure. *Environmental Microbiology* 8, 732–740.

**Examples**

```
if (requireNamespace("Hmisc") && requireNamespace("minpack.lm")) {
  data(otutab, package = "pcutils")
  ncm(otutab) -> ncm_res
  plot(ncm_res)
}
```

---

nst	<i>Calculate NST for each group</i>
-----	-------------------------------------

---

**Description**

Calculate NST for each group

**Usage**

```
nst(otutab, group_df, threads = 1, file = NULL, rep = 20, save = FALSE)
```

**Arguments**

otutab	an otutab data.frame, samples are columns, taxa are rows.
group_df	a dataframe with rowname and one group column
threads	default:4
file	filename to save
rep	repeat numbers: suggest 999
save	save the file

**Value**

a b\_dist object, dis is MSTij

**References**

Ning, D., Deng, Y., Tiedje, J. M. & Zhou, J. (2019) A general framework for quantitatively assessing ecological stochasticity. *Proceedings of the National Academy of Sciences* 116, 16892–16898.

**Examples**

```
if (requireNamespace("NST")) {
  library(ggplot2)
  data(otutab, package = "pcutils")
  nst(otutab, metadata["Group"]) -> nst_res
  plot(nst_res, c_group = "intra") + geom_hline(yintercept = 0.5, lty = 2) + ylab("NST")
}
```

---

nti\_rc *Calculate b\_NTI and RC\_bray for each group*

---

### Description

Calculate b\_NTI and RC\_bray for each group  
 Plot NTI\_RC object

### Usage

```
nti_rc(
  otutab,
  phylo,
  group_df,
  threads = 1,
  file = NULL,
  rep = 20,
  save = FALSE
)

## S3 method for class 'NTI_RC'
plot(x, ...)
```

### Arguments

otutab	an otutab data.frame, samples are columns, taxa are rows.
phylo	a phylo object
group_df	a dataframe with rowname and one group column
threads	default:4
file	filename to save
rep	repeat numbers: suggest 999
save	save the file
x	NTI_RC object
...	pass to <a href="#">stackplot</a>

### Value

a b\_dist object, dis is MSTij  
 ggplot

### References

Ning, D., Deng, Y., Tiedje, J. M. & Zhou, J. (2019) A general framework for quantitatively assessing ecological stochasticity. *Proceedings of the National Academy of Sciences* 116, 16892–16898.



**Examples**

```
if (requireNamespace("NST") && requireNamespace("pctax")) {  
  data(otutab, package = "pcutils")  
  pctax::df2tree(taxonomy) -> phylo  
  nti_rc(otutab, phylo, metadata["Group"]) -> nti_res  
  plot(nti_res)  
}
```

---

pc\_otu

*Create a pc\_otu class object*

---

**Description**

Create a pc\_otu class object

**Usage**

```
pc_otu(otutab = data.frame(), metadata = data.frame(), taxonomy = NULL, ...)
```

**Arguments**

otutab	an otutab data.frame, samples are columns, taxa are rows.
metadata	a metadata data.frame, samples are rows
taxonomy	a taxonomy data.frame, look out the rowname of taxonomy and otutab should matched!
...	add

**Value**

pc\_otu

**Examples**

```
data(otutab, package = "pcutils")  
pc_tax1 <- pc_otu(otutab, metadata)  
print(pc_tax1)
```

---

pc_tax1	<i>test data (pc_otu class) for pc_tax package.</i>
---------	---

---

**Description**

an otutab, metadata and a taxonomy table.

**Format**

a pc\_otu contains an otutab, metadata and a taxonomy table.

**tbls** contains otutable rawdata

**metas** contains metadata

**otus** contains taxonomy table

---

pc_valid	<i>Judge pc_otu is valid or not</i>
----------	-------------------------------------

---

**Description**

Judge pc\_otu is valid or not

**Usage**

```
pc_valid(pc)
```

**Arguments**

pc            a pc\_otu object

**Value**

logical

---

permanova                      *Permanova between a otutab and a variable*

---

### Description

Permanova between a otutab and a variable

Permanova between a otutab and a variable (added two)

### Usage

```
permanova(
  otutab,
  envs,
  norm = TRUE,
  each = TRUE,
  method = "adonis",
  dist = "bray",
  two = FALSE,
  nperm = 999,
  ...
)
```

```
permanova(
  otutab,
  envs,
  norm = TRUE,
  each = TRUE,
  method = "adonis",
  dist = "bray",
  two = FALSE,
  nperm = 999,
  ...
)
```

### Arguments

otutab	an otutab data.frame, samples are columns, taxa are rows.
envs	factors need to test
norm	should normalize?(default:TRUE)
each	test factor one by one, rather than whole
method	adonis/mrpp/anosim/mantel
dist	if use pcoa or nmDS, you can choose a dist method (default: bray)
two	two by two adonis test
nperm	numbers of permutations to perform
...	additional

**Value**

a `g_test` object with these columns

group	the test group or factor
r	relationship
r2	model R-square
p_value	model test p_value
sig	whether significant

**References**

[https://blog.csdn.net/qq\\_42458954/article/details/110390488](https://blog.csdn.net/qq_42458954/article/details/110390488)

**Examples**

```
data(otutab, package = "pcutils")
permanova(otutab, metadata[, c(2:10)]) -> adonis_res
print(adonis_res)
plot(adonis_res)
```

---

plot.a_res	<i>Plot a_res object</i>
------------	--------------------------

---

**Description**

Plot a\_res object

**Usage**

```
## S3 method for class 'a_res'
plot(x, group, metadata, ...)
```

**Arguments**

x	a a_res object
group	one of colname of metadata
metadata	metadata
...	additional parameters for <a href="#">group_box</a> or <a href="#">my_lm</a>

**Value**

patchwork object, you can change theme with &

**See Also**

[a\\_diversity](#)

---

plot.b\_res

*Plot a b\_res*


---

### Description

Plot a b\_res

### Usage

```
## S3 method for class 'b_res'
plot(
  x,
  Group,
  metadata = NULL,
  Group2 = NULL,
  mode = 1,
  bi = FALSE,
  Topn = 10,
  rate = 1,
  margin = FALSE,
  margin_label = TRUE,
  permanova_res = NULL,
  text_param = list(),
  box_margin = TRUE,
  box_param = list(),
  pal = NULL,
  sample_label = TRUE,
  stat_ellipse = TRUE,
  coord_fix = FALSE,
  bi_text_size = 3,
  ...
)
```

### Arguments

x	a b_res object
Group	group vector for color
metadata	metadata contain Group
Group2	mapping point shape
mode	plot mode: 1~3
bi	plot variables segments?
Topn	how many variables to show?
rate	segments length rate
margin	plot the margin boxplot?

margin_label	margin_label, TRUE
permanova_res	permanova result
text_param	text_param for <a href="#">annotate</a>
box_margin	margin plot box or density?
box_param	box_param for <a href="#">group_box</a>
pal	colors for group
sample_label	plot the labels of samples?
stat_ellipse	plot the stat_ellipse?
coord_fix	fix the coordinates y/x ratio
bi_text_size	biplot text size
...	add

**Value**

a ggplot

**See Also**

[b\\_analyse](#)

---

plot.g\_test

*Plot g\_test*

---

**Description**

Plot g\_test

**Usage**

```
## S3 method for class 'g_test'
plot(x, ...)
```

**Arguments**

x	a g_test object
...	add

**Value**

ggplot

**See Also**

[permanova](#)

---

plot.pro_res	<i>Plot pro_res</i>
--------------	---------------------

---

**Description**

Plot pro\_res

**Usage**

```
## S3 method for class 'pro_res'
plot(x, group, metadata = NULL, pal = NULL, ...)
```

**Arguments**

x	pro_res
group	group
metadata	metadata
pal	pal
...	add

**Value**

a ggplot

---

plot.time_cm	<i>Plot time_cm</i>
--------------	---------------------

---

**Description**

Plot time\_cm

**Usage**

```
## S3 method for class 'time_cm'
plot(x, mem_thr = 0.6, ...)
```

**Arguments**

x	time_cm
mem_thr	membership threshold
...	add

**Value**

ggplot

---

plot\_element\_cycle      *Plot element cycle*

---

### Description

Plot element cycle

### Usage

```
plot_element_cycle(
  cycle = "Nitrogen cycle",
  anno_df = NULL,
  only_anno = FALSE,
  cell_fill = NA,
  cell_color = "orange",
  use_chinese = FALSE,
  chemical_size = 7,
  chemical_bold = TRUE,
  chemical_color = "black",
  chemical_label = TRUE,
  reaction_width = 1,
  reaction_arrow_size = 4,
  reaction_arrow_closed = TRUE,
  gene_or_ko = "gene",
  gene_size = 3,
  gene_x_offset = 0.3,
  gene_y_offset = 0.15,
  gene_label = TRUE,
  gene_color = NULL,
  gene_bold = TRUE,
  gene_italic = TRUE,
  gene_label_fill = "white"
)
```

### Arguments

cycle	one of c("Carbon cycle", "Nitrogen cycle", "Phosphorus cycle", "Sulfur cycle", "Iron cycle")
anno_df	anno_df, columns should contains Gene or KO and Group
only_anno	only show genes in anno_df?
cell_fill	cell fill color
cell_color	cell border color
use_chinese	use chinese label?
chemical_size	chemical text size
chemical_bold	chemical text bold



chemical\_color chemical text color  
 chemical\_label chemical text in geom\_label or geom\_text?  
 reaction\_width reaction line width  
 reaction\_arrow\_size  
                   reaction arrow size  
 reaction\_arrow\_closed  
                   reaction arrow closed?  
 gene\_or\_ko "gene" or "ko"  
 gene\_size gene text size  
 gene\_x\_offset gene\_x\_offset  
 gene\_y\_offset gene\_y\_offset  
 gene\_label gene text in geom\_label or geom\_text?  
 gene\_color gene text color  
 gene\_bold gene text bold?  
 gene\_italic gene text italic?  
 gene\_label\_fill  
                   gene label fill color

**Value**

ggplot

**Examples**

```
if (requireNamespace("ggforce")) plot_element_cycle()
```

---

plot_N_cycle	<i>Plot the N-cycling pathway and genes</i>
--------------	---

---

**Description**

Plot the N-cycling pathway and genes

**Usage**

```
plot_N_cycle(
  my_N_genes = NULL,
  just_diff = FALSE,
  path_col = NULL,
  type_col = c(up = "red", down = "blue", none = NA),
  fill_alpha = 0.5,
  arrow_size = 0.1,
  line_width = 1,
  title = "Nitrogen cycling",
  legend.position = c(0.85, 0.15)
)
```

**Arguments**

my\_N\_genes        dataframe, "Gene\_families", "type" should in colnames of my\_N\_genes  
 just\_diff        logical, just plot the different genes?  
 path\_col         colors of pathways  
 type\_col         colors of types  
 fill\_alpha       alpha, default 0.5  
 arrow\_size      arrow\_size, default 0.1  
 line\_width      line\_width, default 1  
 title            title, default "Nitrogen cycling"  
 legend.position   default c(0.85,0.15)

**Value**

ggplot

**Examples**

```

N_data <- load_N_data()
my_N_genes <- data.frame(
  `Gene_families` = sample(N_data$N_genes$Gene_families, 10, replace = FALSE),
  change = rnorm(10), check.names = FALSE
)
my_N_genes <- dplyr::mutate(my_N_genes,
  type = ifelse(change > 0, "up", ifelse(change < 0, "down", "none"))
)
plot_N_cycle(my_N_genes, just_diff = FALSE, fill_alpha = 0.2)
# ggsave(filename = "test.pdf", width = 14, height = 10)

```

---

plot\_two\_tree

*Plot two trees in one plot*

---

**Description**

Plot two trees in one plot

**Usage**

```

plot_two_tree(
  tree1,
  tree2,
  edge_df = NULL,
  tree2_x = 10,
  filter_link = FALSE,
  tree1_param = list(),

```

```

tree2_param = list(),
line_param = list(),
tree1_tip = FALSE,
tip1_param = list(),
tree2_tip = FALSE,
tip2_param = list(),
tree1_highlight = NULL,
highlight1_param = list(),
highlight1_scale = NULL,
tree2_highlight = NULL,
highlight2_param = list(),
highlight2_scale = ggplot2::scale_fill_hue(na.value = NA)
)

```

### Arguments

tree1	phylo object
tree2	phylo object
edge_df	dataframe with edge information, containing "from" and "to" columns
tree2_x	x position of tree2
filter_link	filter the link between tree1 and tree2
tree1_param	parameters for <a href="#">geom_tree</a>
tree2_param	parameters for <a href="#">geom_tree</a>
line_param	parameters for <a href="#">geom_line</a>
tree1_tip	tree tip label
tip1_param	parameters for <a href="#">geom_tiplab</a>
tree2_tip	tree tip label
tip2_param	parameters for <a href="#">geom_tiplab</a>
tree1_highlight	tree1 highlight data.frame
highlight1_param	parameters for <a href="#">geom_highlight</a>
highlight1_scale	scale_fill_ for highlight1
tree2_highlight	tree2 highlight data.frame
highlight2_param	parameters for <a href="#">geom_highlight</a>
highlight2_scale	scale_fill_ for highlight2

### Value

ggplot object

**Examples**

```

if (requireNamespace("ggtree")) {
  data(otutab, package = "pcutils")
  df2tree(taxonomy[1:50, ]) -> tax_tree
  df2tree(taxonomy[51:100, ]) -> tax_tree2
  link <- data.frame(from = sample(tax_tree$tip.label, 20), to = sample(tax_tree2$tip.label, 20))
  plot_two_tree(tax_tree, tax_tree2, link)
}

```

---

pre_fastp	<i>Prepare the result from fastp (.json file)</i>
-----------	---

---

**Description**

Prepare the result from fastp (.json file)

**Usage**

```
pre_fastp(jsonfiles, prefix = c("Raw", "Clean"))
```

**Arguments**

jsonfiles	the directory contains .json file
prefix	default c("Raw","Clean"), for the before filtering and after filtering.

**Value**

data.frame

---

pre_GEO	<i>prepare the GEO data</i>
---------	-----------------------------

---

**Description**

prepare the GEO data

**Usage**

```
pre_GEO(my_id, GEO_dir = "GEO_data", file = NULL)
```

**Arguments**

my_id	GEOid
GEO_dir	GEO download dir
file	the downloaded file

**Value**

```
list(meta = meta, GSE_expr = GSE_expr)
```

---

```
pre_tax_table      Complete a taxonomy table
```

---

**Description**

Complete a taxonomy table

**Usage**

```
pre_tax_table(
  tax_table,
  tax_levels = c("k", "p", "c", "o", "f", "g", "s", "st"),
  na_tax = "Unclassified|uncultured|Ambiguous|Unknown|unknown|metagenome|Unassig",
  ignore.case = TRUE,
  na_repalce = "Unknown"
)
```

**Arguments**

tax_table	taxonomy table
tax_levels	a vector whose length longer than ncol(taxdf), use to be prefix. Default: c("k", "p", "c", "o", "f", "g", "s", "st")
na_tax	grepl some words and turn to na_repalce, default: "Unclassified uncultured Ambiguous Unknown unkno
ignore.case	ignore.case for na_tax
na_repalce	defalut: Unknown

**Value**

a good taxonomy table

**References**

MicrobiotaProcess

**Examples**

```
taxmat <- matrix(sample("onelevel", 7 * 2, replace = TRUE), nrow = 2, ncol = 7) %>% as.data.frame()
colnames(taxmat) <- c("Kingdom", "Phylum", "Class", "Order", "Family", "Genus", "Species")
pre_tax_table(taxmat)
```

---

print.pc_otu	<i>Print</i>
--------------	--------------

---

**Description**

Print

**Usage**

```
## S3 method for class 'pc_otu'
print(x, ...)
```

**Arguments**

x	pc_otu
...	add

**Value**

No value

---

procrustes_analyse	<i>Procrustes Rotation of Two Configurations and PROTEST</i>
--------------------	--

---

**Description**

Procrustes Rotation of Two Configurations and PROTEST

**Usage**

```
procrustes_analyse(b_res1, b_res2, nperm = 999, ...)
```

**Arguments**

b_res1	Target matrix
b_res2	Matrix to be rotated
nperm	numbers of permutations to perform
...	additional

**Value**

pro\_res

**Examples**

```
data(otutab, package = "pcutils")
b_analyse(otutab, method = "pca") -> b_res1
b_analyse(otutab * abs(rnorm(10)), method = "pca") -> b_res2
pro_res <- procrustes_analyse(b_res1, b_res2)
plot(pro_res, "Group", metadata)
```

---

rarefaction	<i>Rarefy a otutab</i>
-------------	------------------------

---

**Description**

Rarefy a otutab

**Usage**

```
rarefaction(otutab, sample = NULL)
```

**Arguments**

otutab	otutab
sample	number

**Value**

a rarefied otutab

**Examples**

```
data(otutab, package = "pcutils")
rarefaction(otutab)
```

---

rare_curve_sample	<i>Rare the sample</i>
-------------------	------------------------

---

**Description**

Rare the sample  
Plot a rare curve

**Usage**

```
rare_curve_sample(otutab, rep = 30, count_cutoff = 1)

## S3 method for class 'rare_res'
plot(x, ...)
```

**Arguments**

otutab	otutab
rep	repeats number
count_cutoff	cutoff to be 0
x	AOR object
...	add

**Value**

ggplot  
ggplot

**Examples**

```
data(otutab, package = "pcutils")
a <- rare_curve_sample(otutab)
plot(a)
```

---

rare\_curve\_species     *Rare the species*

---

**Description**

Rare the species

**Usage**

```
rare_curve_species(  
  otutab,  
  step = 2000,  
  method = "richness",  
  mode = 2,  
  reps = 3,  
  threads = 1,  
  verbose = TRUE  
)
```

**Arguments**

otutab	otutab
step	default 2000
method	one of "richness", "chao1", "ace", "gc", "shannon", "simpson", "pd", "pielou"
mode	1 for little table, 2 for big
reps	reps
threads	use how many threads to calculate (default:1)
verbose	verbose



**Value**

ggplot

**Examples**

```
data(otutab, package = "pcutils")
a <- rare_curve_species(otutab, mode = 1)
plot(a)
```

---

RCbray1

*Calculate RCbray-curtis*

---

**Description**

Calculate RCbray-curtis

**Usage**

```
RCbray1(  
  otutab,  
  reps = 9,  
  threads = 1,  
  classic_metric = TRUE,  
  split_ties = TRUE  
)
```

**Arguments**

otutab	otutab
reps	how many simulation performed?
threads	use how many threads to calculate (default:4)
classic_metric	standardizes the metric to range from -1 to 1
split_ties	adds half of the number of null observations that are equal to the observed number of shared species to the calculation- this is highly recommended

**Details**

Parallelized version of the Raup-Crick algorithm for "abundance" data (Stegen et al. 2013).

**Value**

a dist

**Examples**

```

if (requireNamespace("picante")) {
  data(otutab, package = "pcutils")
  df2tree(taxonomy) -> phylo
  b_NTI1(phylo, otutab) -> bnti_res
  RCbray1(otutab, reps = 9) -> rc_res

  data.frame(
    type = factor(c("Homo_S", "Heter_S", "Homo_D", "D_limit", "Undominated"),
      levels = c("Homo_S", "Heter_S", "Homo_D", "D_limit", "Undominated")
    ),
    number = c(
      sum(bnti_res < (-2)), sum(bnti_res > 2),
      sum((abs(bnti_res) < 2) & (abs(rc_res) < 0.95)),
      sum((abs(bnti_res) < 2) & (rc_res < (-0.95))),
      sum((abs(bnti_res) < 2) & (rc_res > 0.95))
    )
  ) -> com_pro
  pcutils::gghuan(com_pro, reorder = FALSE)
}

```

---

RDA\_plot

*Plot RDA res*


---

**Description**

Plot RDA res

**Usage**

```

RDA_plot(
  phy.rda,
  Group,
  metadata = NULL,
  Group2 = NULL,
  env_rate = 1,
  mode = 1,
  tri = FALSE,
  Topn = 10,
  rate = 1,
  margin = FALSE,
  box = TRUE,
  pal = NULL,
  sample_label = TRUE,
  stat_ellipse = TRUE,
  coord_fix = FALSE,
  bi_text_size = 3,

```

```
    env_text_param = NULL,  
    ...  
  )
```

### Arguments

phy.rda	rda/cca object
Group	group vector for color
metadata	metadata contain Group
Group2	mapping point shape
env_rate	default 1
mode	plot mode: 1~3
tri	plot variables segments?
Topn	how many variables to show?
rate	segments length rate
margin	plot the margin boxplot?
box	margin plot box or density?
pal	colors for group
sample_label	plot the labels of samples?
stat_ellipse	plot the stat_ellipse?
coord_fix	fix the coordinates y/x ratio
bi_text_size	biplot text size
env_text_param	parameters pass to <a href="#">geom_text</a>
...	add

### Value

ggplot

### See Also

[myRDA](#)

---

`sangji_plot`*Plot a sankey*

---

### Description

Plot a sankey

### Usage

```
sangji_plot(  
  tree,  
  top_N = 5,  
  notshow = c(),  
  intermediate = FALSE,  
  width = 3000,  
  height = 500,  
  ...  
)
```

### Arguments

<code>tree</code>	result from <a href="#">ann_tree</a>
<code>top_N</code>	each level has <code>top_N</code>
<code>notshow</code>	some words you don't want to show
<code>intermediate</code>	logical, show the intermediate rank
<code>width</code>	width
<code>height</code>	height
<code>...</code>	look for parameters in <a href="#">sankeyNetwork</a>

### Value

html widget

### Examples

```
if (requireNamespace("sankeyD3") && requireNamespace("tidytree")) {  
  data(otutab, package = "pcutils")  
  ann_tree(taxonomy[, c(1, 5, 6, 7)], otutab) -> tree  
  sangji_plot(tree)  
}
```

---

suijisenlin	<i>RandomForest</i>
-------------	---------------------

---

**Description**

RandomForest

**Usage**

```
suijisenlin(otutab, group_df, topN = 10)
```

**Arguments**

otutab	otutab
group_df	a dataframe with rowname same to dist and one group column
topN	default: 10

**Value**

diff

**Examples**

```
if (requireNamespace("randomForest")) {
  data(otutab, package = "pcutils")
  suijisenlin(otutab, metadata["Group"]) -> rf_res
}
```

---

summary.pc_otu	<i>Summary pc_otu</i>
----------------	-----------------------

---

**Description**

Summary pc\_otu

**Usage**

```
## S3 method for class 'pc_otu'
summary(object, ...)
```

**Arguments**

object	pc_otu
...	add

**Value**

No value

**Examples**

```
data("pc_tax1")
summary(pc_tax1)
```

---

sunburst

*Plot a sunburst*

---

**Description**

Plot a sunburst

**Usage**

```
sunburst(tree)
```

**Arguments**

tree            result from [ann\\_tree](#)

**Value**

sunburst

**See Also**

[sangji\\_plot\(\)](#)

**Examples**

```
if (requireNamespace("plotly")) {
  data(otutab, package = "pcutils")
  ann_tree(taxonomy[, c(1, 5, 6, 7)], otutab) -> tree
  sunburst(tree)
}
```

---

taxonkit_filter	<i>Filter TaxIDs based on Taxonomic Ranks</i>
-----------------	---

---

### Description

This function uses the "taxonkit filter" command to filter TaxIDs based on taxonomic ranks.

### Usage

```
taxonkit_filter(
  file_path,
  black_list = NULL,
  discard_noranks = FALSE,
  discard_root = FALSE,
  equal_to = NULL,
  higher_than = NULL,
  lower_than = NULL,
  rank_file = NULL,
  root_taxid = NULL,
  save_predictable_norank = FALSE,
  taxid_field = NULL,
  text = FALSE,
  data_dir = NULL
)
```

### Arguments

file_path	The path to the input file containing TaxIDs. Or file text (text=TRUE)
black_list	A character vector specifying the ranks to discard.
discard_noranks	Logical value indicating whether to discard all ranks without order (default is FALSE).
discard_root	Logical value indicating whether to discard the root taxid (default is FALSE).
equal_to	A character vector specifying the ranks for which TaxIDs should be equal to.
higher_than	The rank above which the TaxIDs should be (exclusive).
lower_than	The rank below which the TaxIDs should be (exclusive).
rank_file	The path to a user-defined ordered taxonomic ranks file.
root_taxid	The root taxid (default is 1).
save_predictable_norank	Logical value indicating whether to save some special ranks without order when using lower_than (default is FALSE).
taxid_field	The field index of the taxid in the input file (default is 1).
text	logical
data_dir	directory containing nodes.dmp and names.dmp (default "/Users/asa/.taxonkit")

**Value**

A character vector containing the output of the "taxonkit filter" command.

**See Also**

Other Rtaxonkit: [check\\_taxonkit\(\)](#), [download\\_taxonkit\\_dataset\(\)](#), [install\\_taxonkit\(\)](#), [name\\_or\\_id2df\(\)](#), [taxonkit\\_lca\(\)](#), [taxonkit\\_lineage\(\)](#), [taxonkit\\_list\(\)](#), [taxonkit\\_name2taxid\(\)](#), [taxonkit\\_reformat\(\)](#)

**Examples**

```
## Not run:
taxids2 <- system.file("extdata/taxids2.txt", package = "pctax")
taxonkit_filter(taxids2, lower_than = "genus")

## End(Not run)
```

---

taxonkit\_lca

*Compute Lowest Common Ancestor (LCA) of TaxIDs*


---

**Description**

This function uses the "taxonkit lca" command to compute the Lowest Common Ancestor (LCA) of TaxIDs.

**Usage**

```
taxonkit_lca(
  file_path,
  buffer_size = "1M",
  separator = " ",
  skip_deleted = FALSE,
  skip_unfound = FALSE,
  taxids_field = NULL,
  text = FALSE,
  data_dir = NULL
)
```

**Arguments**

file_path	The path to the input file containing TaxIDs. Or file text (text=TRUE)
buffer_size	The size of the line buffer (supported units: K, M, G).
separator	The separator for TaxIDs.
skip_deleted	Whether to skip deleted TaxIDs and compute with the remaining ones.
skip_unfound	Whether to skip unfound TaxIDs and compute with the remaining ones.
taxids_field	The field index of TaxIDs. Input data should be tab-separated (default 1).
text	logical
data_dir	directory containing nodes.dmp and names.dmp (default "/Users/asa/.taxonkit")



**Value**

A character vector containing the computed LCAs.

**See Also**

Other Rtaxonkit: [check\\_taxonkit\(\)](#), [download\\_taxonkit\\_dataset\(\)](#), [install\\_taxonkit\(\)](#), [name\\_or\\_id2df\(\)](#), [taxonkit\\_filter\(\)](#), [taxonkit\\_lineage\(\)](#), [taxonkit\\_list\(\)](#), [taxonkit\\_name2taxid\(\)](#), [taxonkit\\_reformat\(\)](#)

**Examples**

```
## Not run:
taxonkit_lca("239934, 239935, 349741", text = TRUE, separator = ", ")

## End(Not run)
```

---

taxonkit_lineage	<i>Retrieve Taxonomic Lineage using taxonkit</i>
------------------	--

---

**Description**

Retrieve Taxonomic Lineage using taxonkit

**Usage**

```
taxonkit_lineage(
  file_path,
  delimiter = ";",
  no_lineage = FALSE,
  show_lineage_ranks = FALSE,
  show_lineage_taxids = FALSE,
  show_name = FALSE,
  show_rank = FALSE,
  show_status_code = FALSE,
  taxid_field = 1,
  text = FALSE,
  data_dir = NULL
)
```

**Arguments**

file_path	The path to the input file with taxonomic IDs. Or file text (text=TRUE)
delimiter	The field delimiter in the lineage (default ";").
no_lineage	Logical, indicating whether to exclude lineage information (default: FALSE).
show_lineage_ranks	Logical, indicating whether to append ranks of all levels in the lineage (default: FALSE).

show_lineage_taxids	Logical, indicating whether to append lineage consisting of taxids (default: FALSE).
show_name	Logical, indicating whether to append scientific name (default: FALSE).
show_rank	Logical, indicating whether to append rank of taxids (default: FALSE).
show_status_code	Logical, indicating whether to show status code before lineage (default: FALSE).
taxid_field	The field index of taxid. Input data should be tab-separated (default: 1).
text	logical,
data_dir	directory containing nodes.dmp and names.dmp (default "/Users/asa/.taxonkit")

**Value**

A character vector containing the taxonomic lineage information.

**See Also**

Other Rtaxonkit: [check\\_taxonkit\(\)](#), [download\\_taxonkit\\_dataset\(\)](#), [install\\_taxonkit\(\)](#), [name\\_or\\_id2df\(\)](#), [taxonkit\\_filter\(\)](#), [taxonkit\\_lca\(\)](#), [taxonkit\\_list\(\)](#), [taxonkit\\_name2taxid\(\)](#), [taxonkit\\_reformat\(\)](#)

**Examples**

```
## Not run:
taxonkit_lineage("9606\n63221", show_name = TRUE, show_rank = TRUE, text = TRUE)

## End(Not run)
```

---

taxonkit_list	<i>Taxonkit list</i>
---------------	----------------------

---

**Description**

This function uses Taxonkit to perform the "list" operation, which retrieves information about taxa based on their TaxIDs.

**Usage**

```
taxonkit_list(
  ids,
  indent = " ",
  json = FALSE,
  show_name = FALSE,
  show_rank = FALSE,
  data_dir = NULL
)
```

### Arguments

ids	A character vector of TaxIDs to retrieve information for.
indent	The indentation string to use for pretty-printing the output. Default is " ".
json	Logical value indicating whether to output the result in JSON format. Default is FALSE.
show_name	Logical value indicating whether to show the scientific names of taxa. Default is FALSE.
show_rank	Logical value indicating whether to show the ranks of taxa. Default is FALSE.
data_dir	directory containing nodes.dmp and names.dmp (default "/Users/asa/.taxonkit")

### Value

The output of the Taxonkit list operation.

### See Also

Other Rtaxonkit: [check\\_taxonkit\(\)](#), [download\\_taxonkit\\_dataset\(\)](#), [install\\_taxonkit\(\)](#), [name\\_or\\_id2df\(\)](#), [taxonkit\\_filter\(\)](#), [taxonkit\\_lca\(\)](#), [taxonkit\\_lineage\(\)](#), [taxonkit\\_name2taxid\(\)](#), [taxonkit\\_reformat\(\)](#)

### Examples

```
## Not run:
taxonkit_list(ids = c(9605), indent = "-", show_name = TRUE, show_rank = TRUE)

## End(Not run)
```

---

taxonkit\_name2taxid     *Convert Taxonomic Names to TaxIDs*

---

### Description

This function uses the "taxonkit taxonkit\_name2taxid" command to convert taxonomic names to corresponding taxonomic IDs (TaxIDs).

### Usage

```
taxonkit_name2taxid(
  file_path,
  name_field = NULL,
  sci_name = FALSE,
  show_rank = FALSE,
  text = FALSE,
  data_dir = NULL
)
```

**Arguments**

file_path	The path to the input file containing taxonomic names. Or file text (text=TRUE)
name_field	The field index of the taxonomic name in the input file (default is 1).
sci_name	Logical value indicating whether to search only for scientific names (default is FALSE).
show_rank	Logical value indicating whether to show the taxonomic rank in the output (default is FALSE).
text	Logical
data_dir	directory containing nodes.dmp and names.dmp (default "/Users/asa/.taxonkit")

**Value**

A character vector containing the output of the "taxonkit\_name2taxid" command.

**See Also**

Other Rtaxonkit: [check\\_taxonkit\(\)](#), [download\\_taxonkit\\_dataset\(\)](#), [install\\_taxonkit\(\)](#), [name\\_or\\_id2df\(\)](#), [taxonkit\\_filter\(\)](#), [taxonkit\\_lca\(\)](#), [taxonkit\\_lineage\(\)](#), [taxonkit\\_list\(\)](#), [taxonkit\\_reformat\(\)](#)

**Examples**

```
## Not run:
names <- system.file("extdata/name.txt", package = "pctax")
taxonkit_name2taxid(names, name_field = 1, sci_name = FALSE, show_rank = FALSE)
"Homo sapiens" %>% taxonkit_name2taxid(text = TRUE)

## End(Not run)
```

---

taxonkit_reformat	<i>Reformat Taxonomic Lineage using taxonkit</i>
-------------------	--

---

**Description**

Reformat Taxonomic Lineage using taxonkit

**Usage**

```
taxonkit_reformat(
  file_path,
  delimiter = NULL,
  add_prefix = FALSE,
  prefix_kingdom = "K__",
  prefix_phylum = "p__",
  prefix_class = "c__",
  prefix_order = "o__",
```

```

    prefix_family = "f__",
    prefix_genus = "g__",
    prefix_species = "s__",
    prefix_subspecies = "t__",
    prefix_strain = "T__",
    fill_miss_rank = FALSE,
    format_string = "",
    miss_rank_repl_prefix = "unclassified ",
    miss_rank_repl = "",
    miss_taxid_repl = "",
    output_ambiguous_result = FALSE,
    lineage_field = 2,
    taxid_field = NULL,
    pseudo_strain = FALSE,
    trim = FALSE,
    text = FALSE,
    data_dir = NULL
)

```

### Arguments

<code>file_path</code>	The path to the input file with taxonomic lineages. Or file text ( <code>text=TRUE</code> )
<code>delimiter</code>	The field delimiter in the input lineage (default ";").
<code>add_prefix</code>	Logical, indicating whether to add prefixes for all ranks (default: <code>FALSE</code> ).
<code>prefix_kingdom</code>	The prefix for kingdom, used along with <code>-add-prefix</code> (default: "K__").
<code>prefix_phylum</code>	The prefix for phylum, used along with <code>-add-prefix</code> (default: "p__").
<code>prefix_class</code>	The prefix for class, used along with <code>-add-prefix</code> (default: "c__").
<code>prefix_order</code>	The prefix for order, used along with <code>-add-prefix</code> (default: "o__").
<code>prefix_family</code>	The prefix for family, used along with <code>-add-prefix</code> (default: "f__").
<code>prefix_genus</code>	The prefix for genus, used along with <code>-add-prefix</code> (default: "g__").
<code>prefix_species</code>	The prefix for species, used along with <code>-add-prefix</code> (default: "s__").
<code>prefix_subspecies</code>	The prefix for subspecies, used along with <code>-add-prefix</code> (default: "t__").
<code>prefix_strain</code>	The prefix for strain, used along with <code>-add-prefix</code> (default: "T__").
<code>fill_miss_rank</code>	Logical, indicating whether to fill missing rank with lineage information of the next higher rank (default: <code>FALSE</code> ).
<code>format_string</code>	The output format string with placeholders for each rank.
<code>miss_rank_repl_prefix</code>	The prefix for estimated taxon level for missing rank (default: "unclassified ").
<code>miss_rank_repl</code>	The replacement string for missing rank.
<code>miss_taxid_repl</code>	The replacement string for missing taxid.
<code>output_ambiguous_result</code>	Logical, indicating whether to output one of the ambiguous result (default: <code>FALSE</code> ).

lineage_field	The field index of lineage. Input data should be tab-separated (default: 2).
taxid_field	The field index of taxid. Input data should be tab-separated. It overrides -i/-lineage-field.
pseudo_strain	Logical, indicating whether to use the node with lowest rank as strain name (default: FALSE).
trim	Logical, indicating whether to not fill missing rank lower than current rank (default: FALSE).
text	logical
data_dir	directory containing nodes.dmp and names.dmp (default "/Users/asa/.taxonkit")

**Value**

A character vector containing the reformatted taxonomic lineages.

**See Also**

Other Rtaxonkit: [check\\_taxonkit\(\)](#), [download\\_taxonkit\\_dataset\(\)](#), [install\\_taxonkit\(\)](#), [name\\_or\\_id2df\(\)](#), [taxonkit\\_filter\(\)](#), [taxonkit\\_lca\(\)](#), [taxonkit\\_lineage\(\)](#), [taxonkit\\_list\(\)](#), [taxonkit\\_name2taxid\(\)](#)

**Examples**

```
## Not run:
# Use taxid
taxids2 <- system.file("extdata/taxids2.txt", package = "pctax")
reformatted_lineages <- taxonkit_reformat(taxids2,
  add_prefix = TRUE, taxid_field = 1, fill_miss_rank = TRUE
)
reformatted_lineages
taxonomy <- strsplit2(reformatted_lineages, "\t")
taxonomy <- strsplit2(taxonomy$V2, ";")

# Use lineage result
taxonkit_lineage("9606\n63221", show_name = TRUE, show_rank = TRUE, text = TRUE) %>%
  taxonkit_reformat(text = TRUE)

## End(Not run)
```

---

tax\_lca

---

*Calculate the lowest common ancestor (LCA) of a set of taxa*


---

**Description**

Calculate the lowest common ancestor (LCA) of a set of taxa

**Usage**

```
tax_lca(df)
```

**Arguments**

df                    a data frame with taxonomic information, with columns representing taxonomic levels

**Value**

character

**Examples**

```
df <- data.frame(
  A = c("a", "a", "a", "a"),
  B = c("x", "x", "y", "y"),
  C = c("1", "1", "2", "3"),
  stringsAsFactors = FALSE
)
tax_lca(df)
```

---

time\_by\_cm

*Time series analysis*

---

**Description**

Time series analysis

**Usage**

```
time_by_cm(otu_time, n_cluster = 6, min.std = 0)
```

**Arguments**

otu\_time            otutab hebing by a time variable  
n\_cluster           number of clusters  
min.std             min.std

**Value**

time\_cm

**Examples**

```
if (interactive()) {
  data(otutab, package = "pcutils")
  otu_time <- pcutils::hebing(otutab, metadata$Group)
  time_by_cm(otu_time, n_cluster = 4) -> time_cm_res
  plot(time_cm_res)
}
```

---

`volcano_p`*Volcano plot for difference analysis*

---

**Description**

Volcano plot for difference analysis

**Usage**

```
volcano_p(  
  res,  
  logfc = 1,  
  adjp = 0.05,  
  text = TRUE,  
  repel = TRUE,  
  mode = 1,  
  number = FALSE  
)
```

**Arguments**

<code>res</code>	result of <code>diff_da</code> which have colnames: <code>tax</code> , <code>log2FoldChange</code> , <code>padj</code> , <code>compare</code> , <code>sig</code>
<code>logfc</code>	<code>log_fold_change</code> threshold
<code>adjp</code>	<code>adjust_p_value</code> threshold
<code>text</code>	<code>text</code> , TRUE
<code>repel</code>	<code>repel</code> , TRUE
<code>mode</code>	1:normal; 2:multi_contrast
<code>number</code>	show the tax number

**Value**

ggplot

**See Also**

[diff\\_da](#)



---

z_diversity	<i>Calculate Zeta Diversity</i>
-------------	---------------------------------

---

## Description

This function calculates Zeta diversity for each group in the provided otutab.

This function plots the Zeta diversity results obtained from the `z_diversity` function.

## Usage

```
z_diversity(otutab, group_df = NULL, zetadiv_params = list())

## S3 method for class 'zeta_res'
plot(x, lm_model = c("exp", "pl")[1], ribbon = FALSE, text = TRUE, ...)
```

## Arguments

<code>otutab</code>	A matrix or data frame containing OTU (Operational Taxonomic Unit) counts.
<code>group_df</code>	A data frame containing group information.
<code>zetadiv_params</code>	Additional parameters to be passed to the <code>Zeta.decline.mc</code> function from the <code>zetadiv</code> package.
<code>x</code>	Zeta diversity results obtained from <code>z_diversity</code> function.
<code>lm_model</code>	The linear model to be used for fitting ('exp' or 'pl').
<code>ribbon</code>	Logical, whether to add a ribbon to the plot for standard deviation.
<code>text</code>	Logical, whether to add R-squared and p-value text annotations.
<code>...</code>	Additional arguments to be passed to <code>ggplot2</code> functions.

## Value

`zeta_res`  
A `ggplot` object.

## Examples

```
if (requireNamespace("zetadiv")) {
  data(otutab, package = "pcutils")
  zeta_result <- z_diversity(otutab, metadata["Group"], zetadiv_params = list(sam = 10))
  plot(zeta_result, lm_model = "exp", text = TRUE)
}
```

---

`z_diversity_decay`      *Calculate Zeta Diversity with Distance*

---

### Description

This function calculates Zeta diversity for each group in the provided otutab.

### Usage

```
z_diversity_decay(otutab, xy_df, group_df = NULL, zetadiv_params = list())
```

```
## S3 method for class 'zeta_decay'
plot(x, ribbon = TRUE, ...)
```

### Arguments

<code>otutab</code>	A matrix or data frame containing OTU (Operational Taxonomic Unit) counts.
<code>xy_df</code>	Site coordinates.
<code>group_df</code>	A data frame containing group information.
<code>zetadiv_params</code>	Additional parameters to be passed to the <code>Zeta.ddecay</code> function from the <code>zetadiv</code> package.
<code>x</code>	Zeta diversity results obtained from <code>z_diversity_decay</code> function.
<code>ribbon</code>	Logical, whether to add a ribbon to the plot for standard deviation.
<code>...</code>	Additional arguments to be passed to <code>ggplot2</code> functions.

### Value

`zeta_decay`

A `ggplot` object.

### Examples

```
if (requireNamespace("zetadiv")) {
  data(otutab, package = "pcutils")
  zeta_decay_result <- z_diversity_decay(otutab, metadata[, c("lat", "long")],
    metadata["Group"],
    zetadiv_params = list(sam = 10)
  )
  plot(zeta_decay_result)
}
```

# Index

## \* Rtaxonkit

- check\_taxonkit, 15
  - download\_taxonkit\_dataset, 18
  - install\_taxonkit, 23
  - name\_or\_id2df, 29
  - taxonkit\_filter, 55
  - taxonkit\_lca, 56
  - taxonkit\_lineage, 57
  - taxonkit\_list, 58
  - taxonkit\_name2taxid, 59
  - taxonkit\_reformat, 60
- a\_diversity, 10, 36
- add\_strip, 4
- add\_tax, 4
- ALDEX, 5
- all\_ec\_info, 6
- ann\_tree, 6, 52, 54
- annotate, 38
- aor, 7
- as.b\_dist, 9
- as.dist.b\_dist, 10
- b\_analyse, 12, 38
- b\_NTII1, 14
- b\_res\_3d, 14
- bbtt, 11
- before\_tree, 12
- check\_taxonkit, 15, 18, 23, 29, 56–60, 62
- compare\_means, 23
- cor\_net, 16
- df2tree, 16
- df2tree1, 17
- diff\_da, 17, 64
- download\_taxonkit\_dataset, 15, 18, 23, 29, 56–60, 62
- easy\_tree (ann\_tree), 6
- envfit, 19
- envfitt, 19
- gene2id, 19
- geo\_sim, 20
- geom\_highlight, 43
- geom\_line, 43
- geom\_strip, 4
- geom\_text, 51
- geom\_tiplab, 43
- geom\_tree, 43
- get\_diff\_type, 21
- ggtree, 7
- gp\_dis\_density, 21
- grap\_p\_test, 22
- group\_box, 36, 38
- install\_taxonkit, 15, 18, 23, 29, 56–60, 62
- kwtest, 23
- load\_N\_data, 24
- m\_group\_env, 28
- mat\_dist, 24
- micro\_sbatch, 25
- multi\_bar, 26
- my\_lm, 36
- myCAP (myRDA), 27
- myCCA (myRDA), 27
- myRDA, 27, 51
- name\_or\_id2df, 15, 18, 23, 29, 56–60, 62
- ncm, 30
- nst, 31
- nti\_rc, 32
- pc\_otu, 33
- pc\_tax1, 34
- pc\_valid, 34
- permanova, 35, 38
- plot.a\_res, 36

plot.AOR (aor), 7  
plot.b\_dist (as.b\_dist), 9  
plot.b\_res, 37  
plot.dist (as.b\_dist), 9  
plot.g\_test, 38  
plot.mant\_g (m\_group\_env), 28  
plot.ncm\_res (ncm), 30  
plot.NTI\_RC (nti\_rc), 32  
plot.pro\_res, 39  
plot.rare\_res (rare\_curve\_sample), 47  
plot.time\_cm, 39  
plot.zeta\_decay (z\_diversity\_decay), 66  
plot.zeta\_res (z\_diversity), 65  
plot\_element\_cycle, 40  
plot\_N\_cycle, 41  
plot\_two\_tree, 42  
pre\_fastp, 44  
pre\_GEO, 44  
pre\_tax\_table, 45  
print.pc\_otu, 46  
procrustes\_analyse, 46  
  
rare\_curve\_sample, 47  
rare\_curve\_species, 48  
rarefaction, 47  
RCbray1, 49  
RDA\_plot, 50  
  
sangji\_plot, 52  
sangji\_plot(), 54  
sankeyNetwork, 52  
stackplot, 32  
suijisenlin, 53  
summary.pc\_otu, 53  
sunburst, 54  
  
tax\_lca, 62  
taxonkit\_filter, 15, 18, 23, 29, 55, 57–60, 62  
taxonkit\_lca, 15, 18, 23, 29, 56, 56, 58–60, 62  
taxonkit\_lineage, 15, 18, 23, 29, 56, 57, 57, 59, 60, 62  
taxonkit\_list, 15, 18, 23, 29, 56–58, 58, 60, 62  
taxonkit\_name2taxid, 15, 18, 23, 29, 56–59, 59, 62  
taxonkit\_reformat, 15, 18, 23, 29, 56–60, 60  
time\_by\_cm, 63  
  
unifrac, 20, 24, 28  
vegdist, 20, 24, 28  
volcano\_p, 64  
  
z\_diversity, 65  
z\_diversity\_decay, 66