

# Package ‘clusterProfiler’

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**Type** Package

**Title** A Universal Enrichment Tool for Interpreting Omics Data

**Version** 4.19.7

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**Description** A universal tool for interpreting functional characteristics of omics data.

It supports Over-Representation Analysis (ORA) and Gene Set Enrichment Analysis (GSEA) for both coding and non-coding genomics data of thousands of species.

It provides a unified and tidy interface to access, manipulate, and visualize enrichment results. A key capability is the simultaneous analysis and comparison of datasets from multiple treatments or time points.

Furthermore, it integrates Large Language Model (LLM) capabilities to provide automated and insightful interpretation of enrichment results.

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**VignetteBuilder** quarto

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**License** Artistic-2.0

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**BugReports** <https://github.com/YuLab-SMU/clusterProfiler/issues>

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clusterProfiler-package

*clusterProfiler: A Universal Enrichment Tool for Interpreting Omics Data*

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## Description

A universal tool for interpreting functional characteristics of omics data. It supports Over-Representation Analysis (ORA) and Gene Set Enrichment Analysis (GSEA) for both coding and non-coding genomics data of thousands of species. It provides a unified and tidy interface to access, manipulate, and visualize enrichment results. A key capability is the simultaneous analysis and comparison of datasets from multiple treatments or time points. Furthermore, it integrates Large Language Model (LLM) capabilities to provide automated and insightful interpretation of enrichment results.

## Author(s)

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## See Also

Useful links:

- <https://yulab-smu.top/contribution-knowledge-mining/>
- Report bugs at <https://github.com/YuLab-SMU/clusterProfiler/issues>

---

append\_kegg\_category    *append\_kegg\_category*

---

**Description**

add KEGG pathway category information

**Usage**

```
append_kegg_category(x)
```

**Arguments**

x                    KEGG enrichment result

**Details**

This function appends the KEGG pathway category information to KEGG enrichment result (either output of 'enrichKEGG' or 'gseKEGG')

**Value**

update KEGG enrichment result with category information

**Author(s)**

Guangchuang Yu

---

bitr

*bitr*

---

**Description**

Biological Id TRanslator

**Usage**

```
bitr(geneID, fromType, toType, OrgDb, drop = TRUE)
```

**Arguments**

geneID	input gene id
fromType	input id type
toType	output id type
OrgDb	annotation db
drop	drop NA or not

**Value**

data.frame

**Author(s)**

Guangchuang Yu

---

`bitr_kegg`*bitr\_kegg*

---

**Description**

convert biological ID using KEGG API

**Usage**`bitr_kegg(geneID, fromType, toType, organism, drop = TRUE)`**Arguments**

<code>geneID</code>	input gene id
<code>fromType</code>	input id type
<code>toType</code>	output id type
<code>organism</code>	supported organism, can be search using <code>search_kegg_organism</code> function
<code>drop</code>	drop NA or not

**Value**

data.frame

**Author(s)**

Guangchuang Yu

---

`browseKEGG`*browseKEGG*

---

**Description**

open KEGG pathway with web browser

**Usage**`browseKEGG(x, pathID)`**Arguments**

<code>x</code>	an instance of <code>enrichResult</code> or <code>gseaResult</code>
<code>pathID</code>	pathway ID

**Value**

url

**Author(s)**

Guangchuang Yu

---

`compareCluster`*Compare gene clusters functional profile*

---

**Description**

Given a list of gene set, this function will compute profiles of each gene cluster.

**Usage**

```
compareCluster(  
  geneClusters,  
  fun = "enrichGO",  
  data = "",  
  source_from = NULL,  
  ...  
)
```

**Arguments**

<code>geneClusters</code>	a list of entrez gene id. Alternatively, a formula of type <code>Entrez~group</code> or a formula of type <code>Entrez   logFC ~ group</code> for "gseGO", "gseKEGG" and "GSEA".
<code>fun</code>	One of "groupGO", "enrichGO", "enrichKEGG", "enrichDO" or "enrichPathway". Users can also supply their own function.
<code>data</code>	if <code>geneClusters</code> is a formula, the data from which the clusters must be extracted.
<code>source_from</code>	If using a custom function in "fun", provide the source package as a string here. Otherwise, the function will be obtained from the global environment.
<code>...</code>	Other arguments.

**Value**

A `clusterProfResult` instance.

**Author(s)**

Guangchuang Yu <https://yulab-smu.top>

**See Also**

[`compareClusterResult-class`], [`groupGO`], [`enrichGO`], [`enrichKEGG`], [`enrichDO`][`DOSE::enrichDO`], [`enrichPathway`][`ReactomePA::enrichPathway`]

**Examples**

```
## Not run:
data(gcSample)
xx <- compareCluster(gcSample, fun="enrichKEGG",
                     organism="hsa", pvalueCutoff=0.05)
as.data.frame(xx)
# plot(xx, type="dot", caption="KEGG Enrichment Comparison")
dotplot(xx)

## formula interface
mydf <- data.frame(Entrez=c('1', '100', '1000', '100101467',
                           '100127206', '100128071'),
                  logFC = c(1.1, -0.5, 5, 2.5, -3, 3),
                  group = c('A', 'A', 'A', 'B', 'B', 'B'),
                  othergroup = c('good', 'good', 'bad', 'bad', 'good', 'bad'))
xx.formula <- compareCluster(Entrez~group, data=mydf,
                             fun='groupGO', OrgDb='org.Hs.eg.db')
as.data.frame(xx.formula)

## formula interface with more than one grouping variable
xx.formula.twogroups <- compareCluster(Entrez~group+othergroup, data=mydf,
                                       fun='groupGO', OrgDb='org.Hs.eg.db')
as.data.frame(xx.formula.twogroups)

## End(Not run)
```

---

 DataSet

*Datasets gcSample contains a sample of gene clusters.*


---

**Description**

Datasets gcSample contains a sample of gene clusters.

Datasets kegg\_species contains kegg species information

Datasets kegg\_category contains kegg pathway category information

Datasets DE\_GSE8057 contains differential expressed genes obtained from GSE8057 dataset

---

 download\_KEGG

*download\_KEGG*


---

**Description**

download the latest version of KEGG pathway/module

**Usage**

```
download_KEGG(species, keggType = "KEGG", keyType = "kegg")
```

**Arguments**

species	species
keggType	one of 'KEGG' or 'MKEGG'
keyType	supported keyType, see <code>bitr_kegg</code>

**Value**

list

**Author(s)**

Guangchuang Yu

---

dropGO

*dropGO*

---

**Description**

drop GO term of specific level or specific terms (mostly too general).

**Usage**

```
dropGO(x, level = NULL, term = NULL)
```

**Arguments**

x	an instance of 'enrichResult' or 'compareClusterResult'
level	GO level
term	GO term

**Value**

modified version of x

**Author(s)**

Guangchuang Yu

enrichDAVID

*enrichDAVID***Description**

enrichment analysis by DAVID

**Usage**

```
enrichDAVID(
  gene,
  idType = "ENTREZ_GENE_ID",
  universe,
  minGSSize = 10,
  maxGSSize = 500,
  annotation = "GOTERM_BP_FAT",
  pvalueCutoff = 0.05,
  pAdjustMethod = "BH",
  qvalueCutoff = 0.2,
  species = NA,
  david.user
)
```

**Arguments**

gene	input gene
idType	id type
universe	background genes. If missing, the all genes listed in the database (eg TERM2GENE table) will be used as background.
minGSSize	minimal size of genes annotated for testing
maxGSSize	maximal size of genes annotated for testing
annotation	david annotation
pvalueCutoff	adjusted pvalue cutoff on enrichment tests to report
pAdjustMethod	one of "holm", "hochberg", "hommel", "bonferroni", "BH", "BY", "fdr", "none"
qvalueCutoff	qvalue cutoff on enrichment tests to report as significant. Tests must pass i) pvalueCutoff on unadjusted pvalues, ii) pvalueCutoff on adjusted pvalues and iii) qvalueCutoff on qvalues to be reported.
species	species
david.user	david user

**Value**

A enrichResult instance

**Author(s)**

Guangchuang Yu

enricher

*enricher***Description**

A universal enrichment analyzer

**Usage**

```
enricher(
  gene,
  pvalueCutoff = 0.05,
  pAdjustMethod = "BH",
  universe = NULL,
  minGSSize = 10,
  maxGSSize = 500,
  qvalueCutoff = 0.2,
  gson = NULL,
  TERM2GENE,
  TERM2NAME = NA
)
```

**Arguments**

gene	a vector of gene id
pvalueCutoff	adjusted pvalue cutoff on enrichment tests to report
pAdjustMethod	one of "holm", "hochberg", "hommel", "bonferroni", "BH", "BY", "fdr", "none"
universe	background genes. If missing, the all genes listed in the database (eg TERM2GENE table) will be used as background.
minGSSize	minimal size of genes annotated for testing
maxGSSize	maximal size of genes annotated for testing
qvalueCutoff	qvalue cutoff on enrichment tests to report as significant. Tests must pass i) pvalueCutoff on unadjusted pvalues, ii) pvalueCutoff on adjusted pvalues and iii) qvalueCutoff on qvalues to be reported.
gson	a GSON object, if not NULL, use it as annotation data.
TERM2GENE	user input annotation of TERM TO GENE mapping, a data.frame of 2 column with term and gene. Only used when gson is NULL.
TERM2NAME	user input of TERM TO NAME mapping, a data.frame of 2 column with term and name. Only used when gson is NULL.

**Value**

A enrichResult instance

**Author(s)**

Guangchuang Yu <https://yulab-smu.top>

---

enrichGO	<i>GO Enrichment Analysis of a gene set. Given a vector of genes, this function will return the enrichment GO categories after FDR control.</i>
----------	---

---

### Description

GO Enrichment Analysis of a gene set. Given a vector of genes, this function will return the enrichment GO categories after FDR control.

### Usage

```
enrichGO(
  gene,
  OrgDb,
  keyType = "ENTREZID",
  ont = "MF",
  pvalueCutoff = 0.05,
  pAdjustMethod = "BH",
  universe,
  qvalueCutoff = 0.2,
  minGSSize = 10,
  maxGSSize = 500,
  readable = FALSE,
  pool = FALSE
)
```

### Arguments

gene	a vector of entrez gene id.
OrgDb	OrgDb
keyType	keytype of input gene
ont	One of "BP", "MF", and "CC" subontologies, or "ALL" for all three.
pvalueCutoff	adjusted pvalue cutoff on enrichment tests to report
pAdjustMethod	one of "holm", "hochberg", "hommel", "bonferroni", "BH", "BY", "fdr", "none"
universe	background genes. If missing, the all genes listed in the database (eg TERM2GENE table) will be used as background.
qvalueCutoff	qvalue cutoff on enrichment tests to report as significant. Tests must pass i) pvalueCutoff on unadjusted pvalues, ii) pvalueCutoff on adjusted pvalues and iii) qvalueCutoff on qvalues to be reported.
minGSSize	minimal size of genes annotated by Ontology term for testing.
maxGSSize	maximal size of genes annotated for testing
readable	whether mapping gene ID to gene Name
pool	If ont='ALL', whether pool 3 GO sub-ontologies

### Value

An enrichResult instance.

**Author(s)**

Guangchuang Yu <https://yulab-smu.top>

**See Also**

[enrichResult-class], [compareCluster]

**Examples**

```
## Not run:
  data(geneList, package = "DOSE")
  de <- names(geneList)[1:100]
  yy <- enrichG0(de, 'org.Hs.eg.db', ont="BP", pvalueCutoff=0.01)
  head(yy)

## End(Not run)
```

---

enrichKEGG

*KEGG Enrichment Analysis of a gene set. Given a vector of genes, this function will return the enrichment KEGG categories with FDR control.*

---

**Description**

KEGG Enrichment Analysis of a gene set. Given a vector of genes, this function will return the enrichment KEGG categories with FDR control.

**Usage**

```
enrichKEGG(
  gene,
  organism = "hsa",
  keyType = "kegg",
  pvalueCutoff = 0.05,
  pAdjustMethod = "BH",
  universe,
  minGSSize = 10,
  maxGSSize = 500,
  qvalueCutoff = 0.2,
  use_internal_data = FALSE
)
```

**Arguments**

gene	a vector of entrez gene id.
organism	supported organism listed in ' <a href="https://www.genome.jp/kegg/catalog/org_list.html">https://www.genome.jp/kegg/catalog/org_list.html</a> '
keyType	one of "kegg", 'ncbi-geneid', 'ncbi-proteinid' and 'uniprot'
pvalueCutoff	adjusted pvalue cutoff on enrichment tests to report
pAdjustMethod	one of "holm", "hochberg", "hommel", "bonferroni", "BH", "BY", "fdr", "none"
universe	background genes. If missing, the all genes listed in the database (eg TERM2GENE table) will be used as background.

minGSSize      minimal size of genes annotated by Ontology term for testing.  
 maxGSSize      maximal size of genes annotated for testing  
 qvalueCutoff    qvalue cutoff on enrichment tests to report as significant. Tests must pass i)  
                   pvalueCutoff on unadjusted pvalues, ii) pvalueCutoff on adjusted pvalues  
                   and iii) qvalueCutoff on qvalues to be reported.  
 use\_internal\_data  
                   logical, use KEGG.db or latest online KEGG data

**Value**

A enrichResult instance.

**Author(s)**

Guangchuang Yu <https://yulab-smu.top>

**See Also**

[enrichResult-class], [compareCluster]

**Examples**

```
## Not run:
data(geneList, package='DOSE')
de <- names(geneList)[1:100]
yy <- enrichKEGG(de, pvalueCutoff=0.01)
head(yy)

## End(Not run)
```

---

enrichMKEGG	<i>KEGG Module Enrichment Analysis of a gene set. Given a vector of genes, this function will return the enrichment KEGG Module categories with FDR control.</i>
-------------	--

---

**Description**

KEGG Module Enrichment Analysis of a gene set. Given a vector of genes, this function will return the enrichment KEGG Module categories with FDR control.

**Usage**

```
enrichMKEGG(
  gene,
  organism = "hsa",
  keyType = "kegg",
  pvalueCutoff = 0.05,
  pAdjustMethod = "BH",
  universe,
  minGSSize = 10,
  maxGSSize = 500,
  qvalueCutoff = 0.2
)
```

**Arguments**

gene	a vector of entrez gene id.
organism	supported organism listed in 'https://www.genome.jp/kegg/catalog/org_list.html'
keyType	one of "kegg", 'ncbi-geneid', 'ncbi-proteinid' and 'uniprot'
pvalueCutoff	adjusted pvalue cutoff on enrichment tests to report
pAdjustMethod	one of "holm", "hochberg", "hommel", "bonferroni", "BH", "BY", "fdr", "none"
universe	background genes. If missing, the all genes listed in the database (eg TERM2GENE table) will be used as background.
minGSSize	minimal size of genes annotated by Ontology term for testing.
maxGSSize	maximal size of genes annotated for testing
qvalueCutoff	qvalue cutoff on enrichment tests to report as significant. Tests must pass i) pvalueCutoff on unadjusted pvalues, ii) pvalueCutoff on adjusted pvalues and iii) qvalueCutoff on qvalues to be reported.

**Value**

A enrichResult instance.

---

enrichPC

*enrichPC*


---

**Description**

ORA analysis for Pathway Commons

**Usage**

```
enrichPC(gene, ...)
```

**Arguments**

gene	a vector of genes (either hgnc symbols or uniprot IDs)
...	additional parameters, see also the parameters supported by the enricher() function

**Details**

This function performs over-representation analysis using Pathway Commons

**Value**

A enrichResult instance

---

`enrichWP`*enrichWP*

---

**Description**

ORA analysis for WikiPathways

**Usage**

```
enrichWP(gene, organism, ...)
```

**Arguments**

<code>gene</code>	a vector of entrez gene id
<code>organism</code>	supported organisms, which can be accessed via the <code>get_wp_organisms()</code> function
<code>...</code>	additional parameters, see also the parameters supported by the <code>enricher()</code> function

**Details**

This function performs over-representation analysis using WikiPathways

**Value**

A `enrichResult` instance

**Author(s)**

Guangchuang Yu

---

`getPPI`*getPPI*

---

**Description**

`getPPI`

**Usage**

```
getPPI(  
  x,  
  ID = 1,  
  taxID = "auto",  
  required_score = NULL,  
  network_type = "functional",  
  add_nodes = 0,  
  show_query_node_labels = 0,  
  output = "igraph"  
)
```

**Arguments**

x	an ‘enrichResult‘ object or a vector of proteins, e.g. ‘c("PTCH1", "TP53", "BRCA1", "BRCA2")‘
ID	ID or index to extract genes in the enriched term(s) if ‘x‘ is an ‘enrichResult‘ object
taxID	NCBI taxon identifiers (e.g. Human is 9606, see: [STRING organisms](https://string-db.org/cgi/input.pl?input_page_active_form=organisms)).
required_score	threshold of significance to include a interaction, a number between 0 and 1000 (default depends on the network)
network_type	network type: functional (default), physical
add_nodes	adds a number of proteins with to the network based on their confidence score (default:1)
show_query_node_labels	when available use submitted names in the preferredName column when (0 or 1) (default:0)
output	one of ‘data.frame‘ or ‘igraph‘

**Details**

[Getting the STRING network interactions](https://string-db.org/cgi/help.pl?sessionId=btsvnCeNrBk7).

**Value**

a ‘data.frame‘ or an ‘igraph‘ object

**Author(s)**

Yonghe Xia and modified by Guangchuang Yu

---

getTaxID

*getTaxID*

---

**Description**

Convert species scientific name to taxonomic ID

**Usage**

```
getTaxID(species)
```

**Arguments**

species	scientific name of a species
---------	------------------------------

**Value**

taxonomic ID

**Author(s)**

Guangchuang Yu

---

`getTaxInfo`*getTaxInfo*

---

**Description**

Query taxonomy information from 'stringdb' or 'ensembl' web services

**Usage**

```
getTaxInfo(species, source = "stringdb")
```

**Arguments**

species	scientific name of a species
source	one of 'stringdb' or 'ensembl'

**Value**

a 'data.frame' of query information

**Author(s)**

Guangchuang Yu

---

`get_wp_organisms`*get\_wp\_organism*

---

**Description**

list supported organism of WikiPathways

**Usage**

```
get_wp_organisms()
```

**Details**

This function extracts information from 'https://data.wikipathways.org/current/gmt/' and lists all supported organisms

**Value**

supported organism list

**Author(s)**

Guangchuang Yu

---

Gff2GeneTable

*Gff2GeneTable*

---

**Description**

read GFF file and build gene information table

**Usage**

```
Gff2GeneTable(gffFile, compress = TRUE)
```

**Arguments**

<code>gffFile</code>	GFF file
<code>compress</code>	compress file or not

**Details**

given a GFF file, this function extracts information from it and save it in working directory

**Value**

file save.

**Author(s)**

Yu Guangchuang

---

go2ont

*go2ont*

---

**Description**

convert goid to ontology (BP, CC, MF)

**Usage**

```
go2ont(goid)
```

**Arguments**

<code>goid</code>	a vector of GO IDs
-------------------	--------------------

**Value**

data.frame

**Author(s)**

Guangchuang Yu

---

`go2term`*go2term*

---

**Description**

convert goid to descriptive term

**Usage**

```
go2term(goid)
```

**Arguments**

`goid` a vector of GO IDs

**Value**

data.frame

**Author(s)**

Guangchuang Yu

---

`gofilter`*gofilter*

---

**Description**

filter GO enriched result at specific level

**Usage**

```
gofilter(x, level = 4)
```

**Arguments**

`x` output from `enrichGO` or `compareCluster`  
`level` GO level

**Value**

updated object

**Author(s)**

Guangchuang Yu

---

groupGO	<i>Functional Profile of a gene set at specific GO level. Given a vector of genes, this function will return the GO profile at a specific level.</i>
---------	--

---

### Description

Functional Profile of a gene set at specific GO level. Given a vector of genes, this function will return the GO profile at a specific level.

### Usage

```
groupGO(  
  gene,  
  OrgDb,  
  keyType = "ENTREZID",  
  ont = "CC",  
  level = 2,  
  readable = FALSE  
)
```

### Arguments

gene	a vector of entrez gene id.
OrgDb	OrgDb
keyType	key type of input gene
ont	One of "MF", "BP", and "CC" subontologies.
level	Specific GO Level.
readable	if readable is TRUE, the gene IDs will mapping to gene symbols.

### Value

A groupGOResult instance.

### Author(s)

Guangchuang Yu <https://yulab-smu.top>

### See Also

[groupGOResult-class], [compareCluster]

### Examples

```
data(gcSample)  
yy <- groupGO(gcSample[[1]], 'org.Hs.eg.db', ont="BP", level=2)  
head(summary(yy))  
#plot(yy)
```

---

groupGOResult-class    *Class "groupGOResult" This class represents the result of functional Profiles of a set of gene at specific GO level.*

---

### Description

Class "groupGOResult" This class represents the result of functional Profiles of a set of gene at specific GO level.

### Slots

result GO classification result  
 ontology Ontology  
 level GO level  
 organism one of "human", "mouse" and "yeast"  
 gene Gene IDs  
 readable logical flag of gene ID in symbol or not.

### Author(s)

Guangchuang Yu <https://yulab-smu.top>

### See Also

[compareClusterResult], [compareCluster], [groupGO]

---

GSEA

*GSEA*

---

### Description

a universal gene set enrichment analysis tools

### Usage

```
GSEA(  

  geneList,  

  exponent = 1,  

  minGSSize = 10,  

  maxGSSize = 500,  

  pvalueCutoff = 0.05,  

  pAdjustMethod = "BH",  

  gson = NULL,  

  TERM2GENE,  

  TERM2NAME = NA,  

  verbose = TRUE,  

  nPerm = 1000,  

  method = "multilevel",
```

```

    adaptive = FALSE,
    minPerm = 101,
    maxPerm = 1e+05,
    pvalThreshold = 0.1,
    ...
)

```

### Arguments

geneList	order ranked geneList
exponent	weight of each step
minGSSize	minimal size of each geneSet for analyzing
maxGSSize	maximal size of genes annotated for testing
pvalueCutoff	adjusted pvalue cutoff
pAdjustMethod	one of "holm", "hochberg", "hommel", "bonferroni", "BH", "BY", "fdr", "none"
gson	a GSON object, if not NULL, use it as annotation data.
TERM2GENE	user input annotation of TERM TO GENE mapping, a data.frame of 2 column with term and gene. Only used when gson is NULL.
TERM2NAME	user input of TERM TO NAME mapping, a data.frame of 2 column with term and name. Only used when gson is NULL.
verbose	logical
nPerm	The number of permutations.
method	method of calculating the pvalue, one of "multilevel", "monte carlo" and "fgsea"
adaptive	logical, whether to use adaptive method for calculating pvalue
minPerm	minimal number of permutations for adaptive method
maxPerm	maximal number of permutations for adaptive method
pvalThreshold	pvalue threshold for adaptive method
...	other parameter

### Value

gseaResult object

### Author(s)

Guangchuang Yu <https://yulab-smu.top>

---

gseGO

*gseGO*

---

### Description

Gene Set Enrichment Analysis of Gene Ontology

**Usage**

```

gseGO(
  geneList,
  ont = "BP",
  OrgDb,
  keyType = "ENTREZID",
  exponent = 1,
  minGSSize = 10,
  maxGSSize = 500,
  pvalueCutoff = 0.05,
  pAdjustMethod = "BH",
  verbose = TRUE,
  nPerm = 1000,
  method = "multilevel",
  adaptive = FALSE,
  minPerm = 101,
  maxPerm = 1e+05,
  pvalThreshold = 0.1,
  ...
)

```

**Arguments**

geneList	order ranked geneList
ont	one of "BP", "MF", and "CC" subontologies, or "ALL" for all three.
OrgDb	OrgDb
keyType	keytype of gene
exponent	weight of each step
minGSSize	minimal size of each geneSet for analyzing
maxGSSize	maximal size of genes annotated for testing
pvalueCutoff	pvalue Cutoff
pAdjustMethod	pvalue adjustment method
verbose	print message or not
nPerm	The number of permutations.
method	method of calculating the pvalue, one of "multilevel", "monte carlo" and "fgsea"
adaptive	logical, whether to use adaptive method for calculating pvalue
minPerm	minimal number of permutations for adaptive method
maxPerm	maximal number of permutations for adaptive method
pvalThreshold	pvalue threshold for adaptive method
...	other parameter

**Value**

gseaResult object

**Author(s)**

Yu Guangchuang

gseKEGG

*gseKEGG***Description**

Gene Set Enrichment Analysis of KEGG

**Usage**

```

gseKEGG(
  geneList,
  organism = "hsa",
  keyType = "kegg",
  exponent = 1,
  minGSSize = 10,
  maxGSSize = 500,
  pvalueCutoff = 0.05,
  pAdjustMethod = "BH",
  verbose = TRUE,
  use_internal_data = FALSE,
  nPerm = 1000,
  method = "multilevel",
  adaptive = FALSE,
  minPerm = 101,
  maxPerm = 1e+05,
  pvalThreshold = 0.1,
  ...
)

```

**Arguments**

geneList	order ranked geneList
organism	supported organism listed in ' <a href="https://www.genome.jp/kegg/catalog/org_list.html">https://www.genome.jp/kegg/catalog/org_list.html</a> '
keyType	one of "kegg", 'ncbi-geneid', 'ncib-proteinid' and 'uniprot'
exponent	weight of each step
minGSSize	minimal size of each geneSet for analyzing
maxGSSize	maximal size of genes annotated for testing
pvalueCutoff	pvalue Cutoff
pAdjustMethod	pvalue adjustment method
verbose	print message or not
use_internal_data	logical, use KEGG.db or latest online KEGG data
nPerm	The number of permutations.
method	method of calculating the pvalue, one of "multilevel", "monte carlo" and "fgsea"
adaptive	logical, whether to use adaptive method for calculating pvalue
minPerm	minimal number of permutations for adaptive method
maxPerm	maximal number of permutations for adaptive method
pvalThreshold	pvalue threshold for adaptive method
...	other parameter

**Value**

gseaResult object

**Author(s)**

Yu Guangchuang

gseMKEGG

*gseMKEGG***Description**

Gene Set Enrichment Analysis of KEGG Module

**Usage**

```

gseMKEGG(
  geneList,
  organism = "hsa",
  keyType = "kegg",
  exponent = 1,
  minGSSize = 10,
  maxGSSize = 500,
  pvalueCutoff = 0.05,
  pAdjustMethod = "BH",
  verbose = TRUE,
  nPerm = 1000,
  method = "multilevel",
  adaptive = FALSE,
  minPerm = 101,
  maxPerm = 1e+05,
  pvalThreshold = 0.1,
  ...
)

```

**Arguments**

geneList	order ranked geneList
organism	supported organism listed in ' <a href="https://www.genome.jp/kegg/catalog/org_list.html">https://www.genome.jp/kegg/catalog/org_list.html</a> '
keyType	one of "kegg", 'ncbi-geneid', 'ncib-proteinid' and 'uniprot'
exponent	weight of each step
minGSSize	minimal size of each geneSet for analyzing
maxGSSize	maximal size of genes annotated for testing
pvalueCutoff	pvalue Cutoff
pAdjustMethod	pvalue adjustment method
verbose	print message or not
nPerm	The number of permutations.
method	method of calculating the pvalue, one of "multilevel", "monte carlo" and "fgsea"

adaptive	logical, whether to use adaptive method for calculating pvalue
minPerm	minimal number of permutations for adaptive method
maxPerm	maximal number of permutations for adaptive method
pvalThreshold	pvalue threshold for adaptive method
...	other parameter

**Value**

gseaResult object

**Author(s)**

Yu Guangchuang

---

gsePC

*gsePC*

---

**Description**

GSEA analysis for Pathway Commons

**Usage**

```
gsePC(geneList, ...)
```

**Arguments**

geneList	a ranked gene list
...	additional parameters, see also the parameters supported by the GSEA() function

**Details**

This function performs GSEA using Pathway Commons

**Value**

A gseaResult instance

---

gseWP

*gseWP*


---

**Description**

GSEA analysis for WikiPathways

**Usage**

```
gseWP(geneList, organism, ...)
```

**Arguments**

geneList	ranked gene list
organism	supported organisms, which can be accessed via the <code>get_wp_organisms()</code> function
...	additional parameters, see also the parameters supported by the <code>GSEA()</code> function

**Details**

This function performs GSEA using WikiPathways

**Value**

A `gseaResult` instance

**Author(s)**

Guangchuang Yu

---

gson\_GO

*gson\_KEGG*


---

**Description**

download the latest version of KEGG pathway and stored in a 'GSON' object

**Usage**

```
gson_GO(OrgDb, keytype = "ENTREZID", ont = "BP")
```

**Arguments**

OrgDb	OrgDb
keytype	keytype of genes.
ont	one of "BP", "MF", "CC", and "ALL"

**Value**

a 'GSON' object

---

`gson_GO_local`                      *Build a gson object that annotate Gene Ontology*

---

**Description**

Build a gson object that annotate Gene Ontology

**Usage**

```
gson_GO_local(data, ont = c("ALL", "BP", "CC", "MF"), species = NULL, ...)
```

**Arguments**

<code>data</code>	a two-column data.frame of original GO annotation. The columns are "gene_id" and "go_id".
<code>ont</code>	type of GO annotation, which is "ALL", "BP", "MF", or "CC". default: "ALL".
<code>species</code>	name of species. Default: NULL.
<code>...</code>	pass to 'gson::gson()' constructor.

**Value**

a 'gson' instance

**Examples**

```
data = data.frame(gene_id = "gene1",
                  go_id = c("GO:0035492", "GO:0009764", "GO:0031040", "GO:0033714", "GO:0036349"))
gson_GO_local(data, species = "E. coli")
```

---

`gson_KEGG`                              *gson\_KEGG*

---

**Description**

download the latest version of KEGG pathway and stored in a 'GSON' object

**Usage**

```
gson_KEGG(species, KEGG_Type = "KEGG", keyType = "kegg")
```

**Arguments**

<code>species</code>	species
<code>KEGG_Type</code>	one of "KEGG" and "MKEGG"
<code>keyType</code>	one of "kegg", 'ncbi-geneid', 'ncib-proteinid' and 'uniprot'.

**Value**

a 'GSON' object

**Author(s)**

Guangchuang Yu

gson\_KEGG\_mapper

*Build KEGG annotation for novel species using KEGG Mapper***Description**

KEGG Mapper service can annotate protein sequences for novel species with KO database, and KO annotation need to be converted into Pathway or Module annotation, which can then be used in ‘clusterProfiler‘

**Usage**

```
gson_KEGG_mapper(
  file,
  format = c("BLAST", "Ghost", "Kofam"),
  type = c("pathway", "module"),
  species = NULL,
  ...
)
```

**Arguments**

file	the name of the file which comes from the KEGG Mapper service, see Details for file format
format	string indicate format of KEGG Mapper result
type	string indicate annotation database
species	your species, NULL if ignored
...	pass to gson::gson()

**Details**

File is a two-column dataset with K numbers in the second column, optionally preceded by the user’s identifiers in the first column. This is consistent with the output files of automatic annotation servers, BlastKOALA, GhostKOALA, and KofamKOALA. KOALA (KEGG Orthology And Links Annotation) is KEGG’s internal annotation tool for K number assignment of KEGG GENES using SSEARCH computation. BlastKOALA and GhostKOALA assign K numbers to the user’s sequence data by BLAST and GHOSTX searches, respectively, against a nonredundant set of KEGG GENES. KofamKOALA is a new member of the KOALA family available at GenomeNet using the HMM profile search, rather than the sequence similarity search, for K number assignment. see <https://www.kegg.jp/blastkoala/>, <https://www.kegg.jp/ghostkoala/> and <https://www.genome.jp/tools/kofamkoala/> for more information.

**Value**

a gson instance

**Examples**

```
## Not run:
file = system.file('extdata', "kegg_mapper_blast.txt", package='clusterProfiler')
gson_KEGG_mapper(file, format = "BLAST", type = "pathway")

## End(Not run)
```

---

gson\_WP

*gson\_WP*


---

**Description**

Download the latest version of WikiPathways data and stored in a 'GSON' object

**Usage**

```
gson_WP(organism)
```

**Arguments**

organism supported organism, which can be accessed via the `get_wp_organisms()` function.

---

idType

*idType*


---

**Description**

list ID types supported by annoDb

**Usage**

```
idType(OrgDb = "org.Hs.eg.db")
```

**Arguments**

OrgDb annotation db

**Value**

character vector

**Author(s)**

Guangchuang Yu

---

infer_model_id	<i>Infer Model ID</i>
----------------	-----------------------

---

**Description**

Maps bare model names to the aisdk 'provider:model' format for backward compatibility. Emits a warning when guessing and suggests the explicit form. If the model already contains a colon, it is returned as-is.

**Usage**

```
infer_model_id(model)
```

**Arguments**

model	A model string, either bare (e.g., "deepseek-chat") or fully qualified (e.g., "deepseek:deepseek-chat").
-------	--

**Value**

A string in 'provider:model' format.

---

interpret	<i>Interpret Enrichment Results Using LLMs</i>
-----------	--

---

**Description**

Functions for interpreting functional enrichment analysis results using Large Language Models. Supports single-call interpretation, multi-agent deep analysis, and hierarchical cluster strategies.

Built on top of aisdk's 'generate\_object()' for reliable structured output, and the Agent/Session system for multi-agent pipelines.

Sends enrichment results along with optional experimental context to an LLM to generate a structured biological interpretation, hypothesis, and narrative suitable for a publication.

**Usage**

```
interpret(
  x,
  context = NULL,
  n_pathways = 20,
  model = NULL,
  task = "interpretation",
  prior = NULL,
  add_ppi = FALSE,
  gene_fold_change = NULL,
  max_tokens = 8192,
  temperature = 0.3,
  verbose = FALSE
)
```

**Arguments**

x	An enrichment result object ('enrichResult', 'gseaResult', 'compareClusterResult', or a 'data.frame' with pathway columns).
context	A string describing the experimental background (e.g., "scRNA-seq of mouse myocardial infarction at day 3").
n_pathways	Number of top significant pathways to include. Default 20.
model	Optional LLM model. When 'NULL' (default), uses the aisdk package-wide default model configured via 'aisdk::set_model()'. You can also supply a model ID in 'provider:model' format (e.g., "deepseek:deepseek-chat", "gemini:gemini-2.5-flash") or a 'LanguageModelV1' object. Bare model names are supported with a warning (e.g., "deepseek-chat").
task	Task type: "interpretation" (default), "cell_type"/"annotation", or "phenotype"/"phenotyping".
prior	Optional prior knowledge or preliminary annotation to guide the task.
add_ppi	Logical, whether to query STRING PPI network data. Default FALSE.
gene_fold_change	Named numeric vector of log fold changes for expression context.
max_tokens	Maximum tokens for the LLM response. Default 8192. Some models (especially reasoning models) may need much higher values (e.g., 16384 or more) to produce complete structured output.
temperature	Sampling temperature. Default 0.3.
verbose	Logical, whether to print debug messages showing raw API responses, token usage, and JSON parsing details. Default FALSE. Equivalent to setting 'options(aisdk.debug = TRUE)' for the call.

**Details**

Uses 'generate\_object()' internally for reliable structured output with automatic JSON repair, eliminating manual parsing failures.

**Value**

An 'interpretation' object (list) with task-specific fields. For "interpretation": overview, key\_mechanisms, hypothesis, narrative, etc. For "annotation": cell\_type, confidence, reasoning, markers, etc. For "phenotype": phenotype, confidence, reasoning, key\_processes, etc.

**Examples**

```
## Not run:
# Basic usage with a data frame
df <- data.frame(
  ID = c("GO:0006915", "GO:0008284"),
  Description = c("apoptotic process", "positive regulation of proliferation"),
  GeneRatio = c("10/100", "20/100"),
  p.adjust = c(0.01, 0.02),
  geneID = c("TP53/BAX", "MYC/CCND1/CDK4")
)
res <- interpret(df,
  model = "deepseek:deepseek-chat",
  context = "Cancer proliferation study"
)
# Reuse aisdk's global default model
```

```
# aisdk::set_model("openai:gpt-4o-mini")
# res <- interpret(df, context = "Cancer proliferation study")
print(res)

## End(Not run)
```

---

interpret\_agent

*Interpret enrichment results using a multi-agent pipeline (Deep Mode)*


---

## Description

Employs three specialized AI agents in sequence for rigorous interpretation:

1. Agent Cleaner: Filters noise and selects relevant pathways.
2. Agent Detective: Identifies key regulators and functional modules.
3. Agent Synthesizer: Produces a coherent biological narrative.

## Usage

```
interpret_agent(
  x,
  context = NULL,
  n_pathways = 50,
  model = NULL,
  add_ppi = FALSE,
  gene_fold_change = NULL,
  max_tokens = 8192,
  temperature = 0.3,
  verbose = FALSE
)
```

## Arguments

x	An enrichment result object.
context	A string describing the experimental background.
n_pathways	Number of top pathways to consider initially. Default 50.
model	Optional LLM model. When 'NULL' (default), uses the aisdk package-wide default model configured via 'aisdk::set_model()'. You can also supply a model ID in 'provider:model' format or a 'LanguageModelV1' object. Bare model names are supported with a warning.
add_ppi	Logical, whether to query PPI data. Default FALSE.
gene_fold_change	Named numeric vector of log fold changes.
max_tokens	Maximum tokens per agent call. Default 8192.
temperature	Sampling temperature. Default 0.3.
verbose	Logical, whether to print debug messages. Default FALSE.

## Details

Uses aisdk's Agent and Session system for shared context across agents.

**Value**

An ‘interpretation’ object with deep analysis fields plus regulatory\_drivers, refined\_network, and network\_evidence from the detective agent.

**Examples**

```
## Not run:
res <- interpret_agent(df,
  model = "openai:gpt-4o",
  context = "scRNA-seq of mouse MI day 3"
)
print(res)

## End(Not run)
```

---

```
interpret_hierarchical
```

*Interpret enrichment results using a hierarchical strategy*

---

**Description**

First interprets major clusters to establish lineage context, then interprets sub-clusters with hierarchical constraints from the major cluster annotations.

**Usage**

```
interpret_hierarchical(
  x_minor,
  x_major,
  mapping,
  model = NULL,
  task = "cell_type",
  max_tokens = 8192,
  temperature = 0.3
)
```

**Arguments**

x_minor	Enrichment result for sub-clusters.
x_major	Enrichment result for major clusters.
mapping	A named vector mapping sub-cluster IDs to major cluster IDs.
model	Optional LLM model. When ‘NULL’ (default), uses the aisdk package-wide default model configured via ‘aisdk::set_model()’. You can also supply a model ID in ‘provider:model’ format or a ‘LanguageModelV1’ object. Bare model names are supported with a warning.
task	Task type, default "cell_type".
max_tokens	Maximum tokens. Default 8192.
temperature	Sampling temperature. Default 0.3.

**Value**

An ‘interpretation\_list’ object.

---

ko2name	<i>ko2name</i>
---------	----------------

---

**Description**

convert ko ID to descriptive name

**Usage**

```
ko2name(ko)
```

**Arguments**

ko                    ko ID

**Value**

data.frame

**Author(s)**

guangchuang yu

---

merge_result	<i>merge_result</i>
--------------	---------------------

---

**Description**

merge a list of enrichResult objects to compareClusterResult

**Usage**

```
merge_result(enrichResultList)
```

**Arguments**

enrichResultList  
                  a list of enrichResult objects

**Value**

a compareClusterResult instance

**Author(s)**

Guangchuang Yu

---

plot.interpretation    *plot*

---

### Description

plot

### Usage

```
## S3 method for class 'interpretation'
plot(x, layout = "nicely", ...)
```

### Arguments

x	An 'interpretation' object.
layout	Graph layout, default is "nicely".
...	Additional arguments passed to 'ggplot2::ggplot'.

---

plotGOgraph            *plotGOgraph*

---

### Description

plot GO graph

### Usage

```
plotGOgraph(
  x,
  firstSigNodes = 10,
  useInfo = "all",
  sigForAll = TRUE,
  useFullNames = TRUE,
  ...
)
```

### Arguments

x	output of enrichGO or gseGO
firstSigNodes	number of significant nodes (rectangle nodes in the graph)
useInfo	additional info
sigForAll	if TRUE the score/p-value of all nodes in the DAG is shown, otherwise only score will be shown
useFullNames	logical
...	additional parameter of showSigOfNodes, please refer to topGO

**Value**

GO DAG graph

**Author(s)**

Guangchuang Yu

---

read.gmt.pc	<i>read.gmt.pc</i>
-------------	--------------------

---

**Description**

Parse gmt file from Pathway Common

**Usage**

```
read.gmt.pc(gmtfile, output = "data.frame")
```

**Arguments**

gmtfile	A gmt file
output	one of 'data.frame' or 'GSON'

**Details**

This function parse gmt file downloaded from Pathway common

**Value**

A data.frame or A GSON object depends on the value of 'output'

---

reexports	<i>Objects exported from other packages</i>
-----------	---

---

**Description**

These objects are imported from other packages. Follow the links below to see their documentation.

**dplyr** [arrange](#), [filter](#), [group\\_by](#), [mutate](#), [n](#), [rename](#), [select](#), [slice](#), [summarise](#)

**enrichit** [geneID](#), [geneInCategory](#), [gsfilter](#), [setReadable](#)

**enrichplot** [cnetplot](#), [dotplot](#), [emapplot](#), [goplot](#), [gseaplot](#), [heatplot](#), [ridgeplot](#)

**GOSemSim** [buildGOMap](#), [get\\_organism](#), [read.blast2go](#), [read.gaf](#)

**gson** [read.gmt](#), [read.gmt.wp](#)

**magrittr** [%<>%](#), [%>%](#)

search\_kegg\_organism    *search\_kegg\_organism*

---

**Description**

search kegg organism, listed in [https://www.genome.jp/kegg/catalog/org\\_list.html](https://www.genome.jp/kegg/catalog/org_list.html)

**Usage**

```
search_kegg_organism(  
  str,  
  by = "scientific_name",  
  ignore.case = FALSE,  
  use_internal_data = TRUE  
)
```

**Arguments**

str                    string  
by                    one of 'kegg.code', 'scientific\_name' and 'common\_name'  
ignore.case        TRUE or FALSE  
use\_internal\_data    logical, use kegg\_species.rda or latest online KEGG data

**Value**

data.frame

**Author(s)**

Guangchuang Yu

---

simplify                    *simplify method*

---

**Description**

simplify output from enrichGO and gseGO by removing redundancy of enriched GO terms

simplify output from compareCluster by removing redundancy of enriched GO terms

**Usage**

```
## S4 method for signature 'enrichResult'
simplify(
  x,
  cutoff = 0.7,
  by = "p.adjust",
  select_fun = min,
  measure = "Wang",
  semData = NULL
)

## S4 method for signature 'gseaResult'
simplify(
  x,
  cutoff = 0.7,
  by = "p.adjust",
  select_fun = min,
  measure = "Wang",
  semData = NULL
)

## S4 method for signature 'compareClusterResult'
simplify(
  x,
  cutoff = 0.7,
  by = "p.adjust",
  select_fun = min,
  measure = "Wang",
  semData = NULL
)
```

**Arguments**

x	output of enrichGO
cutoff	similarity cutoff
by	feature to select representative term, selected by 'select_fun' function
select_fun	function to select feature passed by 'by' parameter
measure	method to measure similarity
semData	GOSemSimDATA object

**Value**

updated enrichResult object  
updated compareClusterResult object

**Author(s)**

Guangchuang Yu  
Gwang-Jin Kim and Guangchuang Yu

**References**

issue #28 <https://github.com/GuangchuangYu/clusterProfiler/issues/28>

issue #162 <https://github.com/GuangchuangYu/clusterProfiler/issues/162>

---

uniprot_get	<i>uniprot_get</i>
-------------	--------------------

---

**Description**

retrieves annotation data from uniprot

**Usage**

```
uniprot_get(taxID)
```

**Arguments**

taxID            taxonomy ID

**Value**

gene table data frame

**Author(s)**

guangchuang yu

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