

# Package ‘tinyarray’

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**Type** Package

**Title** Expression Data Analysis and Visualization

**Version** 2.2.7

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**Description** Gene Expression Omnibus(GEO) and The Cancer Genome Atlas(TCGA) are common bioinformatics public databases. We integrate the regular analysis and charts for expression data, to analyze and display the data concisely and intuitively.

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**Encoding** UTF-8

**LazyData** true

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**URL** <https://github.com/xjsun1221/tinyarray>

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<i>box_surv</i>	<i>box_surv</i>
-----------------	-----------------

---

### Description

draw box plot for a hub gene expression matrix

### Usage

```
box_surv(exp_hub, exprSet_hub, meta)
```

### Arguments

<i>exp_hub</i>	an expression matrix for hubgenes
<i>exprSet_hub</i>	a tumor expression set for hubgenes
<i>meta</i>	meta data corresponds to expression set

### Value

patchwork result for hub genes boxplot and survival plot

### Author(s)

Xiaojie Sun

### See Also

[exp\\_boxplot](#); [exp\\_surv](#)

### Examples

```
k = box_surv(log2(exp_hub1+1), exprSet_hub1, meta1); k[[1]]
```

cod                      *cod*

---

**Description**

An expression matrix form TCGA

**Usage**

cod

**Format**

An object of class `matrix` (inherits from `array`) with 100 rows and 512 columns.

**Examples**

```
cod
```

---

cor.full                      *cor.test for all variables*

---

**Description**

cor.test for all variables(each two columns)

**Usage**

```
cor.full(x)
```

**Arguments**

x                      A numeric matrix or data.frame

**Value**

a data.frame with cor.test p.value and estimate

**Author(s)**

Xiaojie Sun

**See Also**

[cor.one](#)

**Examples**

```
x = iris[,-5]
cor.full(x)
```

---

`cor.one`*cor.test for one variable with all variables*

---

**Description**

cor.test for all variables(each two columns)

**Usage**

```
cor.one(x, var)
```

**Arguments**

x	A numeric matrix or data.frame
var	your chosen variable,only one.

**Value**

A data.frame with cor.test p.value and estimate

**Author(s)**

Xiaojie Sun

**See Also**

[cor.full](#)

**Examples**

```
x = iris[,-5]
cor.one(x,"Sepal.Width")
```

---

deg	<i>deg</i>
-----	------------

---

**Description**

limma differential analysis result for GSE42872

**Usage**

deg

**Format**

An object of class `data.frame` with 18591 rows and 10 columns.

**Examples**

```
head(deg)
```

---

deseq_data	<i>deseq_data</i>
------------	-------------------

---

**Description**

DEseq2 differential analysis result

**Usage**

deseq\_data

**Format**

An object of class `data.frame` with 552 rows and 6 columns.

**Examples**

```
head(deseq_data)
```

---

double_enrich	<i>draw enrichment bar plots for both up and down genes</i>
---------------	---

---

**Description**

draw enrichment bar plots for both up and down genes,for human only.

**Usage**

```
double_enrich(deg, n = 10, color = c("#2874C5", "#f87669"))
```

**Arguments**

deg	a data.frame contains at least two columns:"ENTREZID" and "change"
n	how many terms will you perform for up and down genes respectively
color	color for bar plot

**Value**

a list with kegg and go bar plot according to up and down genes enrichment result.

**Author(s)**

Xiaojie Sun

**See Also**

[quick\\_enrich](#)

**Examples**

```
double_enrich(deg)
```

---

draw_boxplot	<i>draw boxplot for expression</i>
--------------	------------------------------------

---

**Description**

draw boxplot for expression

**Usage**

```
draw_boxplot(
  exp,
  group_list,
  method = "kruskal.test",
  sort = TRUE,
  drop = FALSE,
  width = 0.5,
  pvalue_cutoff = 0.05,
  xlab = "Gene",
  ylab = "Expression",
  grouplab = "Group",
  p.label = FALSE,
  add_error_bar = FALSE,
  color = c("#2874C5", "#f87669", "#e6b707", "#868686", "#66C2A5", "#FC8D62",
            "#8DA0CB", "#E78AC3", "#A6D854", "#FFD92F", "#E5C494", "#B3B3B3")
)
```

**Arguments**

exp	A numeric matrix
group_list	A factor with duplicated character or factor
method	one of kruskal.test,aov,t.test and wilcox.test
sort	whether the boxplot will be sorted
drop	whether to discard insignificant values
width	width of boxplot and error bar
pvalue_cutoff	if drop = TRUE,genes with p-values below the threshold will be drawn
xlab	title of the x axis
ylab	title of the y axis
grouplab	title of group legend
p.label	whether to show p value in the plot
add_error_bar	whether to add error bar
color	color vector

**Value**

a boxplot according to exp and grouped by group.

**Author(s)**

Xiaojie Sun

**See Also**

[draw\\_heatmap](#);[draw\\_volcano](#);[draw\\_venn](#)



**Examples**

```

draw_boxplot(t(iris[,1:4]),iris$Species)
exp <- matrix(rnorm(60),nrow = 10)
colnames(exp) <- paste0("sample",1:6)
rownames(exp) <- paste0("gene",1:10)
exp[,4:6] = exp[,4:6] +10
exp[1:4,1:4]
group_list <- factor(rep(c("A","B"),each = 3))
draw_boxplot(exp,group_list)
draw_boxplot(exp,group_list,color = c("grey","red"))

```

---

draw_heatmap	<i>draw a heatmap plot</i>
--------------	----------------------------

---

**Description**

print a heatmap plot for expression matrix and group by group\_list paramter, exp will be scaled.

**Usage**

```

draw_heatmap(
  n,
  group_list,
  scale_before = FALSE,
  n_cutoff = 3,
  cluster_cols = TRUE,
  legend = FALSE,
  show_rownames = FALSE,
  annotation_legend = FALSE,
  split_column = FALSE,
  show_column_title = FALSE,
  color = (grDevices::colorRampPalette(c("#2fa1dd", "white", "#f87669")))(100),
  color_an = c("#2fa1dd", "#f87669", "#e6b707", "#868686", "#92C5DE", "#F4A582",
    "#66C2A5", "#FC8D62", "#8DA0CB", "#E78AC3", "#A6D854", "#FFD92F", "#E5C494",
    "#B3B3B3"),
  scale = TRUE,
  main = NA
)

```

**Arguments**

n	A numeric matrix
group_list	A factor with duplicated character or factor
scale_before	deprecated parameter
n_cutoff	3 by defalut , scale before plot and set a cutoff,usually 2 or 1.6
cluster_cols	if F,heatmap will nor cluster in column

legend	logical,show legend or not
show_rownames	logical,show rownames or not
annotation_legend	logical,show annotation legend or not
split_column	split column by group_list
show_column_title	show column title or not
color	color for heatmap
color_an	color for column annotation
scale	logical,scale the matrix or not
main	the title of the plot

**Value**

a heatmap plot according to exp and grouped by group.

**Author(s)**

Xiaojie Sun

**See Also**

[draw\\_pca](#);[draw\\_volcano](#);[draw\\_venn](#)

**Examples**

```
#example data
exp = matrix(abs(rnorm(60,sd = 16)),nrow = 10)
exp[,4:6] <- exp[,4:6]+20
colnames(exp) <- paste0("sample",1:6)
rownames(exp) <- paste0("gene",1:10)
exp[1:4,1:4]
group_list = factor(rep(c("A","B"),each = 3))
draw_heatmap(exp,group_list)
#use iris
n = t(iris[,1:4]);colnames(n) = 1:150
group_list = iris$Species
draw_heatmap(n,group_list)
draw_heatmap(n,group_list,color = colorRampPalette(c("green","black","red"))(100),
             color_an = c("red","blue","pink") )
```

---

draw_heatmap2	<i>draw heatmap plots</i>
---------------	---------------------------

---

### Description

print heatmap plots for expression matrix and group by group\_list paramter

### Usage

```
draw_heatmap2(
  exp,
  group_list,
  deg,
  heat_union = TRUE,
  heat_id = 1,
  gene_number = 200,
  show_rownames = FALSE,
  scale_before = FALSE,
  n_cutoff = 3,
  cluster_cols = TRUE,
  annotation_legend = FALSE,
  legend = FALSE,
  color = (grDevices::colorRampPalette(c("#2fa1dd", "white", "#f87669")))(100),
  color_an = c("#2fa1dd", "#f87669", "#e6b707", "#868686", "#92C5DE", "#F4A582",
    "#66C2A5", "#FC8D62", "#8DA0CB", "#E78AC3", "#A6D854", "#FFD92F", "#E5C494",
    "#B3B3B3")
)
```

### Arguments

exp	A numeric matrix
group_list	A factor with duplicated character or factor
deg	a data.frame created by Differential analysis
heat_union	logical ,use union or intersect DEGs for heatmap
heat_id	id of heatmap,1 for all DEGs,2 for head and tail,3 for top n DEGs
gene_number	how many DEGs will heatmap show .
show_rownames	logical,show rownames or not
scale_before	deprecated parameter
n_cutoff	3 by defalut , scale before plot and set a cutoff,usually 2 or 1.6
cluster_cols	if F,heatmap will nor cluster in column
annotation_legend	
	logical,show annotation legend or not
legend	logical,show legend or not
color	color vector
color_an	color for column annotation

**Value**

a heatmap plot according to exp and grouped by group.

**Author(s)**

Xiaojie Sun

**See Also**

[draw\\_pca](#); [draw\\_volcano](#); [draw\\_venn](#)

**Examples**

```
gse = "GSE474"
geo = geo_download(gse, destdir=tempdir(), by_annoprobe = FALSE)
geo$exp[1:4, 1:4]
geo$exp=log2(geo$exp+1)
group_list=ifelse(stringr::str_detect(geo$pd$title, "MObese"), "MObese",
  ifelse(stringr::str_detect(geo$pd$title, "NonObese"), "NonObese", "Obese"))
group_list=factor(group_list, levels = c("NonObese", "Obese", "MObese"))
find_anno(geo$gpl)
ids <- AnnoProbe::idmap(geo$gpl, destdir = tempdir())
deg = multi_deg(geo$exp, group_list, ids, adjust = FALSE)
draw_heatmap2(geo$exp, group_list, deg)
```

---

draw\_KM

*draw\_KM*

---

**Description**

draw KM-plot with two or more group

**Usage**

```
draw_KM(
  meta,
  group_list,
  time_col = "time",
  event_col = "event",
  legend.title = "Group",
  legend.labs = levels(group_list),
  color = c("#2874C5", "#f87669", "#e6b707", "#868686", "#92C5DE", "#F4A582",
    "#66C2A5", "#FC8D62", "#8DA0CB", "#E78AC3", "#A6D854", "#FFD92F", "#E5C494",
    "#B3B3B3")
)
```

**Arguments**

meta	survival data with time and event column
group_list	A factor with duplicated character or factor
time_col	colname of time
event_col	colname of event
legend.title	legend title
legend.labs	character vector specifying legend labels
color	color vector

**Value**

a KM-plot

**Author(s)**

Xiaojie Sun

**Examples**

```
require("survival")
x = survival::lung
draw_KM(meta = x,
        group_list = x$sex, event_col = "status")
```

---

draw\_pca

*draw PCA plots*

---

**Description**

do PCA analysis and print a PCA plot

**Usage**

```
draw_pca(
  exp,
  group_list,
  color = c("#2874C5", "#f87669", "#e6b707", "#868686", "#92C5DE", "#F4A582",
            "#66C2A5", "#FC8D62", "#8DA0CB", "#E78AC3", "#A6D854", "#FFD92F", "#E5C494",
            "#B3B3B3"),
  addEllipses = TRUE,
  style = "default",
  color.label = "Group"
)
```

**Arguments**

exp	A numeric matrix
group_list	A factor with duplicated character or factor
color	color vector
addEllipses	logical,add ellipses or not
style	plot style,"default","ggplot2"and "3D"
color.label	color legend label

**Value**

a pca plot according to exp and grouped by group.

**Author(s)**

Xiaojie Sun

**See Also**

[draw\\_heatmap](#);[draw\\_volcano](#);[draw\\_venn](#)

**Examples**

```
draw_pca(t(iris[,1:4]),iris$Species)
exp <- matrix(rnorm(60),nrow = 10)
colnames(exp) <- paste0("sample",1:6)
rownames(exp) <- paste0("gene",1:10)
exp[1:4,1:4]
group_list <- factor(rep(c("A","B"),each = 3))
draw_pca(exp,group_list)
draw_pca(exp,group_list,color = c("blue","red"))
```

---

draw\_tsne

*draw\_tsne*

---

**Description**

draw tsne plot with annotation by ggplot2

**Usage**

```
draw_tsne(
  exp,
  group_list,
  perplexity = 30,
  color = c("#2874C5", "#f87669", "#e6b707", "#868686", "#92C5DE", "#F4A582",
    "#66C2A5", "#FC8D62", "#8DA0CB", "#E78AC3", "#A6D854", "#FFD92F", "#E5C494",
```

```

    "#B3B3B3"),
    color.label = "group",
    addEllipses = TRUE
  )

```

### Arguments

exp	A numeric matrix
group_list	A factor with duplicated character or factor
perplexity	numeric; perplexity parameter for Rtsne
color	color vector
color.label	color legend label
addEllipses	logical, add ellipses or not

### Value

a ggplot object

### Author(s)

Xiaojie Sun

### Examples

```

exp <- matrix(rnorm(10000), nrow = 50)
colnames(exp) <- paste0("sample", 1:200)
rownames(exp) <- paste0("gene", 1:50)
exp[1:4, 1:4]
exp[, 1:100] = exp[, 1:100] + 10
group_list <- factor(rep(c("A", "B"), each = 100))
draw_tsne(exp, group_list)

```

---

draw\_venn

*draw a venn plot*

---

### Description

print a venn plot for deg result created by three packages

### Usage

```

draw_venn(
  x,
  name,
  color = c("#2874C5", "#f87669", "#e6b707", "#868686", "#66C2A5", "#FC8D62",
    "#8DA0CB", "#E78AC3", "#A6D854", "#FFD92F", "#E5C494", "#B3B3B3")
)

```

**Arguments**

x                    a list for plot  
name                main of the plot  
color                color vector

**Value**

a venn plot according to x, y and.z named "name" paramter

**Author(s)**

Xiaojie Sun

**See Also**

[draw\\_pca](#);[draw\\_volcano](#);[draw\\_heatmap](#)

**Examples**

```
x = list(Deseq2=sample(1:100,30),edgeR = sample(1:100,30),limma = sample(1:100,30))
draw_venn(x,"test")
draw_venn(x,"test",color = c("darkgreen", "darkblue", "#B2182B"))
```

---

draw\_volcano

*draw a volcano plot*

---

**Description**

print a volcano plot for Differential analysis result in data.frame format.

**Usage**

```
draw_volcano(  
  deg,  
  lab = NA,  
  pvalue_cutoff = 0.05,  
  logFC_cutoff = 1,  
  pkg = 1,  
  adjust = FALSE,  
  symmetry = FALSE,  
  color = c("#2874C5", "grey", "#f87669")  
)
```



**Arguments**

deg	a data.frame created by Differential analysis
lab	label for x axis in volcano plot
pvalue_cutoff	Cutoff value of pvalue,0.05 by default.
logFC_cutoff	Cutoff value of logFC,1 by default.
pkg	a integer ,means which Differential analysis packages you used,we support three packages by now, 1,2,3,4 respectively means "DESeq2","edgeR","limma(voom)","limma"
adjust	a logical value, would you like to use adjusted pvalue to draw this plot,FAISE by default.
symmetry	a logical value ,would you like to get your plot symmetrical
color	color vector

**Value**

a volcano plot according to logFC and P.value(or adjust P.value)

**Author(s)**

Xiaojie Sun

**See Also**

[draw\\_heatmap](#);[draw\\_pca](#);[draw\\_venn](#)

**Examples**

```
head(deseq_data)
draw_volcano(deseq_data)
draw_volcano(deseq_data,pvalue_cutoff = 0.01,logFC_cutoff = 2)
draw_volcano(deseq_data,color = c("darkgreen", "darkgrey", "#B2182B"))
```

---

draw\_volcano2

*draw\_volcano2*

---

**Description**

print one or more volcano plot for Differential analysis result in data.frame format.

**Usage**

```
draw_volcano2(
  deg,
  pkg = 4,
  lab,
  pvalue_cutoff = 0.05,
  logFC_cutoff = 1,
  adjust = FALSE,
  symmetry = TRUE,
  color = c("#2874C5", "grey", "#f87669")
)
```

**Arguments**

deg	a data.frame created by Differential analysis
pkg	a integer ,means which Differential analysis packages you used,we support three packages by now, 1,2,3,4 respectively means "DESeq2","edgeR","limma(voom)","limma"
lab	label for x axis in volcano plot
pvalue_cutoff	Cutoff value of pvalue,0.05 by default.
logFC_cutoff	Cutoff value of logFC,1 by default.
adjust	a logical value, would you like to use adjusted pvalue to draw this plot,FAISE by default.
symmetry	a logical value ,would you like to get your plot symmetrical
color	color vector

**Value**

one or more volcano plot

**Author(s)**

Xiaojie Sun

**See Also**

[geo\\_download](#);[draw\\_volcano](#);[draw\\_venn](#)

**Examples**

```
#two group
gse = "GSE42872"
geo = geo_download(gse,destdir=tempdir(),by_annoprobe = FALSE)
group_list = rep(c("A","B"),each = 3)
ids = AnnoProbe::idmap('GPL6244',destdir = tempdir())
deg = get_deg(geo$exp,group_list,ids)
draw_volcano2(deg)
#multigroup
```

```
gse = "GSE474"
geo = geo_download(gse, destdir=tempdir(), by_anno = FALSE)
geo$exp[1:4, 1:4]
geo$exp=log2(geo$exp+1)
group_list=ifelse(stringr::str_detect(geo$pd$title, "MObese"), "MObese",
  ifelse(stringr::str_detect(geo$pd$title, "NonObese"), "NonObese", "Obese"))
group_list=factor(group_list, levels = c("NonObese", "Obese", "MObese"))
find_anno(geo$gpl)
ids <- AnnoProbe::idmap(geo$gpl, destdir = tempdir())
deg = multi_deg(geo$exp, group_list, ids, adjust = FALSE)
draw_volcano2(deg)
draw_volcano2(deg, color = c("darkgreen", "grey", "darkred"))
```

---

dumd

*count unique values in every columns for data.frame*

---

## Description

in geo analysis, this function can help you simplify pdata, delete columns with unique values, which can't be used as group vector

## Usage

```
dumd(x)
```

## Arguments

x                    A data.frame.

## Value

The simple data.frame of columns unique values count in x

## Examples

```
dumd(iris)
data(ToothGrowth)
x = ToothGrowth
dumd(ToothGrowth)
```

---

edges_to_nodes	<i>edges_to_nodes</i>
----------------	-----------------------

---

**Description**

get nodes from edges

**Usage**

```
edges_to_nodes(edges)
```

**Arguments**

edges            data.frame

**Value**

nodes data.frame

**Author(s)**

Xiaojie Sun

**See Also**

[interaction\\_to\\_edges](#)

**Examples**

```
df = data.frame(a = c("gene1", "gene2", "gene3"),
  b = c("d, f, a, b",
  "c, e, g",
  "a, b, d"))
edges = interaction_to_edges(df)
nodes = edges_to_nodes(edges)
```

---

exists_anno_list	<i>exists_anno_list</i>
------------------	-------------------------

---

**Description**

AnnoProbe supported GPLs

**Usage**

```
exists_anno_list
```

**Format**

An object of class character of length 175.

**Examples**

```
exists_anno_list
```

---

<code>exprSet_hub1</code>	<i>exprSet_hub1</i>
---------------------------	---------------------

---

**Description**

An expression matrix from TCGA,tumor samples only

**Usage**

```
exprSet_hub1
```

**Format**

An object of class data.frame with 8 rows and 177 columns.

**Examples**

```
exprSet_hub1[1:4,1:4]
```

---

<code>exp_boxplot</code>	<i>exp_boxplot</i>
--------------------------	--------------------

---

**Description**

draw box plot for a hub gene expression matrix

**Usage**

```
exp_boxplot(exp_hub, color = c("grey", "red"))
```

**Arguments**

<code>exp_hub</code>	an expression matrix for hubgenes
<code>color</code>	color for boxplot

**Value**

box plots list for all genes in the matrix

**Author(s)**

Xiaojie Sun

**See Also**[exp\\_surv](#); [box\\_surv](#)**Examples**

```
k = exp_boxplot(log2(exp_hub1+1));k[[1]]
```

---

exp_hub1	<i>exp_hub1</i>
----------	-----------------

---

**Description**

An expression matrix from TCGA

**Usage**

exp\_hub1

**Format**An object of class `matrix` (inherits from `array`) with 8 rows and 350 columns.**Examples**

```
exp_hub1[1:4,1:4]
```

---

exp_surv	<i>exp_surv</i>
----------	-----------------

---

**Description**

draw surv plot for a hub gene expression matrix for tumor samples

**Usage**

```
exp_surv(exprSet_hub, meta, color = c("grey", "red"))
```

**Arguments**

exprSet_hub	a tumor expression set for hubgenes
meta	meta data corresponds to expression set
color	color for boxplot

**Value**

survival plots list for all genes

**Author(s)**

Xiaojie Sun

**See Also**

[exp\\_boxplot](#); [box\\_surv](#); [draw\\_venn](#)

**Examples**

```
tmp = exp_surv(exprSet_hub1, meta1)
patchwork::wrap_plots(tmp)
```

---

find_anno	<i>find annotation package or files</i>
-----------	---

---

**Description**

find gpl annotation package or files

**Usage**

```
find_anno(gpl, install = FALSE, update = FALSE)
```

**Arguments**

gpl	a gpl accession
install	whether to install and library the package
update	whether to update the package

**Value**

a list with deg data.frame, volcano plot and a list with DEGs.

**Author(s)**

Xiaojie Sun

**See Also**

[geo\\_download](#)

**Examples**

```
find_anno("GPL570")
```

---

genes	<i>genes</i>
-------	--------------

---

**Description**

some gene entriz ids

**Usage**

genes

**Format**

An object of class character of length 511.

**Examples**

```
genes
```

---

geo_download	<i>geo_download</i>
--------------	---------------------

---

**Description**

download gse data and get informations

**Usage**

```
geo_download(
  gse,
  by_annoprobe = TRUE,
  simpd = TRUE,
  colon_remove = FALSE,
  destdir = getwd()
)
```

**Arguments**

gse	gse assession number
by_annoprobe	getGEO or geoChina
simpd	get simplified pdata,drop out columns with all same values
colon_remove	whether to remove duplicated columns with colons
destdir	The destination directory for data downloads.



**Value**

a list with exp,pd and gpl

**Author(s)**

Xiaojie Sun

**See Also**

[find\\_anno](#)

**Examples**

```
gse = "GSE42872"  
a = geo_download(gse,by_annoprobe = FALSE,destdir=tempdir())
```

---

get\_cgs

*get\_cgs*

---

**Description**

extract DEGs from deg data.frame

**Usage**

```
get_cgs(deg)
```

**Arguments**

deg                    a data.frame created by Differential analysis

**Value**

a list with upgenes,downgenes,diffgenes.

**Author(s)**

Xiaojie Sun

**See Also**

[geo\\_download](#);[draw\\_volcano](#);[draw\\_venn](#)

**Examples**

```

#two group
gse = "GSE42872"
geo = geo_download(gse,destdir=tempdir(),by_annoprobe = FALSE)
group_list = rep(c("A","B"),each = 3)
ids = AnnoProbe::idmap('GPL6244',destdir=tempdir())
deg = get_deg(geo$exp,group_list,ids)
cgs = get_cgs(deg)
#mutigroup
gse = "GSE474"
geo = geo_download(gse,destdir=tempdir(),by_annoprobe = FALSE)
geo$exp[1:4,1:4]
geo$exp=log2(geo$exp+1)
group_list=ifelse(stringr::str_detect(geo$pd$title,"MObese"),"MObese",
ifelse(stringr::str_detect(geo$pd$title,"NonObese"),"NonObese","Obese"))
group_list=factor(group_list,levels = c("NonObese","Obese","MObese"))
find_anno(geo$gpl)
ids = AnnoProbe::idmap(geo$gpl,destdir = tempdir())
deg = multi_deg(geo$exp,group_list,ids,adjust = FALSE)
cgs = get_cgs(deg)

```

---

get\_deg

*get\_deg*


---

**Description**

do differential analysis according to expression set and group information

**Usage**

```

get_deg(
  exp,
  group_list,
  ids,
  logFC_cutoff = 1,
  pvalue_cutoff = 0.05,
  adjust = FALSE,
  entriz = TRUE
)

```

**Arguments**

exp	A numeric matrix
group_list	A factor with duplicated character or factor
ids	a data.frame with 2 columns,including probe_id and symbol
logFC_cutoff	Cutoff value of logFC,1 by default.

pvalue\_cutoff Cutoff value of pvalue,0.05 by default.  
adjust a logical value, would you like to use adjusted pvalue to draw this plot,FAISE by default.  
entriz whether convert symbols to entriz ids

**Value**

a deg data.frame with 10 columns

**Author(s)**

Xiaojie Sun

**See Also**

[multi\\_deg](#); [get\\_deg\\_all](#)

**Examples**

```
gse = "GSE42872"  
geo = geo_download(gse,destdir=tempdir(),by_annoprobe = FALSE)  
Group = rep(c("control","treat"),each = 3)  
Group = factor(Group)  
find_anno(geo$gpl)  
ids <- AnnoProbe::idmap(geo$gpl,destdir = tempdir())  
deg = get_deg(geo$exp,Group,ids)  
head(deg)
```

---

get\_deg\_all

*get\_deg\_all*

---

**Description**

do diffiencial analysis according to exprission set and group information

**Usage**

```
get_deg_all(  
  exp,  
  group_list,  
  ids,  
  logFC_cutoff = 1,  
  pvalue_cutoff = 0.05,  
  adjust = FALSE,  
  entriz = TRUE,  
  scale_before = FALSE,  
  n_cutoff = 3,
```

```

cluster_cols = TRUE,
annotation_legend = FALSE,
show_rownames = FALSE,
legend = FALSE,
lab = NA,
pkg = 4,
symmetry = FALSE,
heat_union = TRUE,
heat_id = 1,
gene_number = 200,
color_volcano = c("#2874C5", "grey", "#f87669")
)

```

### Arguments

exp	A numeric matrix
group_list	A factor with duplicated character or factor
ids	a data.frame with 2 columns,including probe_id and symbol
logFC_cutoff	Cutoff value of logFC,1 by default.
pvalue_cutoff	Cutoff value of pvalue,0.05 by default.
adjust	a logical value, would you like to use adjusted pvalue to draw this plot,FAISE by default.
entriz	logical , if TRUE ,convert symbol to entriz id.
scale_before	deprecated parameter
n_cutoff	3 by defalut , scale before plot and set a cutoff,usually 2 or 1.6
cluster_cols	if F,heatmap will nor cluster in column
annotation_legend	logical,show annotation legend or not
show_rownames	logical,show rownames or not
legend	logical,show legend or not
lab	label for x axis in volcano plot
pkg	a integer ,means which Differential analysis packages you used,we support three packages by now, 1,2,3,4 respectively means "DESeq2","edgeR","limma(voom)","limma"
symmetry	a logical value ,would you like to get your plot symmetrical
heat_union	logical ,use union or intersect DEGs for heatmap
heat_id	id of heatmap,1 for all DEGs,2 for head and tail,3 for top n DEGs
gene_number	how many DEGs will heatmap show .
color_volcano	color for volcano plot

### Value

a list with deg data.frame, volcano plot ,pca plot ,heatmap and a list with DEGs.

**Author(s)**

Xiaojie Sun

**See Also**[get\\_deg;multi\\_deg\\_all](#)**Examples**

```
gse = "GSE42872"
geo = geo_download(gse,destdir=tempdir(),by_annoprobe = FALSE)
group_list = rep(c("A","B"),each = 3)
group_list = factor(group_list)
find_anno(geo$gpl)
ids <- AnnoProbe::idmap(geo$gpl,destdir = tempdir())
dcp = get_deg_all(geo$exp,group_list,ids)
head(dcp$deg)
dcp$plots
```

---

*ggheat**ggheat*

---

**Description**

draw heatmap plot with annotation by ggplot2

**Usage**

```
ggheat(
  dat,
  group,
  cluster = FALSE,
  color = c("#2874C5", "white", "#f87669"),
  legend_color = c("#2874C5", "#f87669", "#e6b707", "#868686", "#66C2A5", "#FC8D62",
    "#8DA0CB", "#E78AC3", "#A6D854", "#FFD92F", "#E5C494", "#B3B3B3"),
  show_rownames = TRUE,
  show_colnames = TRUE,
  groupname = "group",
  expname = "exp",
  fill_mid = TRUE
)
```

**Arguments**

<code>dat</code>	expression matrix for plot
<code>group</code>	group for expression colnames
<code>cluster</code>	logical, cluster or not, default F
<code>color</code>	color for heatmap
<code>legend_color</code>	color for legend
<code>show_rownames</code>	logical, show rownames in plot or not, default T
<code>show_colnames</code>	logical, show colnames in plot or not, default T
<code>groupname</code>	name of group legend
<code>expname</code>	name of exp legend
<code>fill_mid</code>	use median value as geom_tile fill midpoint

**Value**

a ggplot object

**Author(s)**

Xiaojie Sun

**Examples**

```
exp_dat = matrix(sample(100:1000,40),ncol = 4)
exp_dat[1:(nrow(exp_dat)/2),] = exp_dat[1:(nrow(exp_dat)/2),]-1000
rownames(exp_dat) = paste0("sample",1:nrow(exp_dat))
colnames(exp_dat) = paste0("gene",1:ncol(exp_dat))
group = rep(c("A","B"),each = nrow(exp_dat)/2)
group = factor(group,levels = c("A","B"))
ggheat(exp_dat,group)
ggheat(exp_dat,group,cluster = TRUE)
ggheat(exp_dat,group,cluster = TRUE,show_rownames = FALSE,
       show_colnames = FALSE,groupname = "risk",expname = "expression")
```

---

hypertest

*hypertest*

---

**Description**

make hypertest for given lncRNA and mRNA common miRNAs

**Usage**

```
hypertest(lnc, pc, deMIR = NULL, lnctarget, pctarget)
```

**Arguments**

lnc	lncRNA names
pc	mRNA names
deMIR	miRNA names , default NULL
lnctarget	a data.frame with two column,lncRNA in the first column ,miRNA in the second column
pctarget	a data.frame with two column,mRNA in the first column ,miRNA in the second column

**Value**

a data.frame with hypertest result

**Author(s)**

Xiaojie Sun

**See Also**

[plcortest](#)

**Examples**

```
# to update
```

---

```
interaction_to_edges  interaction_to_edges
```

---

**Description**

split interactions by sep paramter,return edges data.frame

**Usage**

```
interaction_to_edges(df, a = 1, b = 2, sep = ",")
```

**Arguments**

df	interactions data.frame
a	column to replicate
b	column to split
sep	a character string to separate b column

**Value**

a new data.frame with two column ,one interaction by one rows

**Author(s)**

Xiaojie Sun

**See Also**[edges\\_to\\_nodes](#)**Examples**

```
df = data.frame(a = c("gene1", "gene2", "gene3"),
               b = c("d, f, a, b",
                   "c, e, g",
                   "a, b, d"))
interaction_to_edges(df)
```

---

`intersect_all`*intersect\_all*

---

**Description**

calculate intersect set for two or more elements

**Usage**`intersect_all(...)`**Arguments**`...` some vectors or a list with some vectors**Value**

vector

**Author(s)**

Xiaojie Sun

**See Also**[union\\_all](#)**Examples**

```
x1 = letters[1:4]
x2 = letters[3:6]
x3 = letters[3:4]
re = intersect_all(x1, x2, x3)
re2 = intersect_all(list(x1, x2, x3))
re3 = union_all(x1, x2, x3)
```



---

lnc_anno	<i>lnc_anno</i>
----------	-----------------

---

**Description**

annotation for TCGA expression matrix(lncRNA),form genecode v22 gtf file.

**Usage**

```
lnc_anno
```

**Format**

An object of class data.frame with 14826 rows and 3 columns.

**Examples**

```
head(lnc_anno)
```

---

lnc_annotv23	<i>lnc_annotv23</i>
--------------	---------------------

---

**Description**

annotation for TCGA and gtex expression matrix(lncRNA),form genecode v23 gtf file.

**Usage**

```
lnc_annotv23
```

**Format**

An object of class data.frame with 14852 rows and 3 columns.

**Examples**

```
head(lnc_annotv23)
```

---

make_tcga_group	<i>make_tcga_group</i>
-----------------	------------------------

---

**Description**

make tcga group for given tcga expression matrix

**Usage**

```
make_tcga_group(exp)
```

**Arguments**

exp                    TCGA or TCGA\_Gtex expression set from gdc or xena

**Value**

a group factor with normal and tumor ,correspond to colnames for expression matrix

**Author(s)**

Xiaojie Sun

**See Also**

[sam\\_filter](#); [match\\_exp\\_cl](#)

**Examples**

```
k = make_tcga_group(exp_hub1); table(k)
```

---

match_exp_cl	<i>match_exp_cl</i>
--------------	---------------------

---

**Description**

match exp and clinical data from TCGA

**Usage**

```
match_exp_cl(exp, cl, id_column = "id", sample_centric = TRUE)
```

**Arguments**

exp            TCGA expression set  
 cl             TCGA clinical data.frame  
 id\_column     which column contains patient ids, column number or column name.  
 sample\_centric logical,default T,keep all samples from the same patients.if FALSE,keep only one tumor sample for one patient.

**Value**

a transformed clinical data.frame with sample ids.

**Author(s)**

Xiaojie Sun

**See Also**

[make\\_tcga\\_group;sam\\_filter](#)

**Examples**

```

a = match_exp_cl(exp_hub1,meta1[,2:4],"X_PATIENT")
exp_matched = a[[1]]
cl_matched = a[[2]]
b = match_exp_cl(exp_hub1,meta1[,2:4],"X_PATIENT",sample_centric = FALSE)
exp_matched = b[[1]]
cl_matched = b[[2]]

```

---

meta1

*meta1*

---

**Description**

clinical messages for some TCGA patients,correspond to exprSet\_hub1

**Usage**

```
meta1
```

**Format**

An object of class data.frame with 177 rows and 4 columns.

**Examples**

```
head(meta1)
```

---

`mRNA_anno``mRNA_anno`

---

**Description**

annotation for TCGA and gtex expression matrix(mRNA),form genecode v22 gtf file.

**Usage**`mRNA_anno`**Format**

An object of class data.frame with 19814 rows and 3 columns.

**Examples**`head(mRNA_anno)`

---

`mRNA_annot23``mRNA_annot23`

---

**Description**

annotation for TCGA and gtex expression matrix(mRNA),form genecode v23 gtf file.

**Usage**`mRNA_annot23`**Format**

An object of class data.frame with 19797 rows and 3 columns.

**Examples**`head(mRNA_annot23)`

---

`multi_deg`*multi\_deg*

---

**Description**

do differential analysis according to expression set and group information

**Usage**

```
multi_deg(  
  exp,  
  group_list,  
  ids,  
  logFC_cutoff = 1,  
  pvalue_cutoff = 0.05,  
  adjust = FALSE,  
  entriz = TRUE  
)
```

**Arguments**

<code>exp</code>	A numeric matrix
<code>group_list</code>	A factor with duplicated character or factor
<code>ids</code>	a data.frame with 2 columns,including probe_id and symbol
<code>logFC_cutoff</code>	Cutoff value of logFC,1 by default.
<code>pvalue_cutoff</code>	Cutoff value of pvalue,0.05 by default.
<code>adjust</code>	a logical value, would you like to use adjusted pvalue to draw this plot,FAISE by default.
<code>entriz</code>	whether convert symbols to entriz ids

**Value**

a deg data.frame with 10 columns

**Author(s)**

Xiaojie Sun

**See Also**

[get\\_deg](#); [multi\\_deg\\_all](#)

**Examples**

```

gse = "GSE474"
geo = geo_download(gse,destdir=tempdir(),by_annoProbe = FALSE)
geo$exp[1:4,1:4]
geo$exp=log2(geo$exp+1)
group_list=ifelse(stringr::str_detect(geo$pd$title,"MObese"),
"MObese",ifelse(stringr::str_detect(geo$pd$title,"NonObese"),
"NonObese","Obese"))
group_list=factor(group_list,levels = c("NonObese","Obese","MObese"))
find_anno(geo$gpl)
ids <- AnnoProbe::idmap(geo$gpl,destdir = tempdir())
deg = multi_deg(geo$exp,group_list,ids,adjust = FALSE)
names(deg)
head(deg[[1]])
head(deg[[2]])
head(deg[[3]])

```

---

multi\_deg\_all

*multi\_deg\_all*


---

**Description**

do diffiencial analysis according to exprission set and group information

**Usage**

```

multi_deg_all(
  exp,
  group_list,
  ids,
  logFC_cutoff = 1,
  pvalue_cutoff = 0.05,
  adjust = FALSE,
  entriz = TRUE,
  scale_before = FALSE,
  n_cutoff = 3,
  cluster_cols = TRUE,
  annotation_legend = FALSE,
  show_rownames = FALSE,
  legend = FALSE,
  lab = NA,
  pkg = 4,
  symmetry = FALSE,
  heat_union = TRUE,
  heat_id = 1,
  gene_number = 200,

```

```

    color_volcano = c("#2874C5", "grey", "#f87669")
  )

```

### Arguments

exp	A numeric matrix
group_list	A factor with duplicated character or factor
ids	a data.frame with 2 columns,including probe_id and symbol
logFC_cutoff	Cutoff value of logFC,1 by default.
pvalue_cutoff	Cutoff value of pvalue,0.05 by default.
adjust	a logical value, would you like to use adjusted pvalue to draw this plot,FAISE by default.
entriz	whether convert symbols to entriz ids
scale_before	deprecated parameter
n_cutoff	3 by defalut , scale before plot and set a cutoff,usually 2 or 1.6
cluster_cols	if F,heatmap will nor cluster in column
annotation_legend	logical,show annotation legend or not
show_rownames	logical,show rownames or not
legend	logical,show legend or not
lab	label for x axis in volcano plot
pkg	a integer ,means which Differential analysis packages you used,we support three packages by now, 1,2,3,4 respectively means "DESeq2","edgeR","limma(voom)","limma"
symmetry	a logical value ,would you like to get your plot symmetrical
heat_union	logical ,use union or intersect DEGs for heatmap
heat_id	id of heatmap,1 for all DEGs,2 for head and tail,3 for top n DEGs
gene_number	how many DEGs will heatmap show .
color_volcano	color for volcano

### Value

a list with deg data.frame, volcano plot and a list with DEGs.

### Author(s)

Xiaojie Sun

### See Also

[geo\\_download](#);[draw\\_volcano](#);[draw\\_venn](#)

**Examples**

```

gse = "GSE474"
geo = geo_download(gse, destdir=tempdir(), by_annoProbe = FALSE)
geo$exp[1:4, 1:4]
geo$exp=log2(geo$exp+1)
group_list=ifelse(stringr::str_detect(geo$pd$title, "MObese"), "MObese",
ifelse(stringr::str_detect(geo$pd$title, "NonObese"), "NonObese", "Obese"))
group_list=factor(group_list, levels = c("NonObese", "Obese", "MObese"))
find_anno(geo$gpl)
ids = AnnoProbe::idmap(geo$gpl, destdir = tempdir())
dcp = multi_deg_all(geo$exp,
group_list, ids, adjust = FALSE)
dcp[[3]]

```

---

pkg\_all

*pkg\_all*

---

**Description**

bioconductor annotation packages for GPLs

**Usage**

pkg\_all

**Format**

An object of class data.frame with 85 rows and 3 columns.

**Examples**

```
head(pkg_all)
```

---

plcortest

*plcortest*

---

**Description**

make cor.test for given lncRNA and mRNA

**Usage**

```
plcortest(lnc_exp, mRNA_exp, cor_cutoff = 0)
```



**Arguments**

lnc\_exp            lncRNA expression set  
mRNA\_exp        mRNA expression set which nrow equal to lncRNA\_exp  
cor\_cutoff       cor estimate cut\_off, default 0

**Value**

a list with cor.test result, names are lncRNAs, element are mRNAs

**Author(s)**

Xiaojie Sun

**See Also**

[hypertest](#)

**Examples**

```
# to update
```

---

point\_cut            *point\_cut*

---

**Description**

calculate cut point for multiple genes

**Usage**

```
point_cut(exprSet_hub, meta)
```

**Arguments**

exprSet\_hub       a tumor expression set for hubgenes  
meta               meta data corresponds to expression set

**Value**

a vector with cutpoint for genes

**Author(s)**

Xiaojie Sun

**See Also**

[surv\\_KM](#); [surv\\_cox](#)

**Examples**

```
point_cut(exprSet_hub1,meta1)
```

---

quick\_enrich

*quick\_enrich*

---

**Description**

do diffiencial analysis according to exprission set and group information,for human only

**Usage**

```
quick_enrich(genes, kkgofile = "kkgofile.Rdata", destdir = getwd())
```

**Arguments**

genes	a gene symbol or entrizid vector
kkgofile	Rdata filename for kegg and go result
destdir	destdir to save kkgofile

**Value**

enrichment results and dotplots

**Author(s)**

Xiaojie Sun

**See Also**

[double\\_enrich](#)

**Examples**

```
head(genes)
g = quick_enrich(genes,destdir = tempdir())
names(g)
g[[1]][1:4,1:4]
g[[3]]
g[[4]]
```

---

sam_filter	<i>sam_filter</i>
------------	-------------------

---

**Description**

drop duplicated samples from the same patients

**Usage**

```
sam_filter(exp)
```

**Arguments**

exp                    TCGA or TCGA\_Gtex expression set from gdc or xena

**Value**

a transformed expression set without duplicated samples

**Author(s)**

Xiaojie Sun

**See Also**

[make\\_tcga\\_group;match\\_exp\\_cl](#)

**Examples**

```
cod[1:4,1:4]
dim(cod)
cod2 = sam_filter(cod)
dim(cod2)
g = make_tcga_group(cod);table(g)
library(stringr)
table(!duplicated(str_sub(colnames(cod[,g=="tumor"]),1,12)))
```

---

surv_cox	<i>surv_cox</i>
----------	-----------------

---

### Description

calculate cox p values and HR for genes

### Usage

```
surv_cox(
  exprSet_hub,
  meta,
  cut.point = FALSE,
  pvalue_cutoff = 0.05,
  HRkeep = "all",
  continuous = FALSE,
  min_gn = 0.1
)
```

### Arguments

exprSet_hub	a tumor expression set for hubgenes
meta	meta data corresponds to expression set
cut.point	logical , use cut_point or not, if FALSE,use median by default
pvalue_cutoff	p value cut off ,0.05 by default
HRkeep	one of "all","protect"or"risk"
continuous	logical, gene expression or gene expression group
min_gn	Depending on the expression of a gene, there may be a large difference in the number of samples between the two groups, and if a smaller group of samples is less than 10 percent (default) of all, the gene will be discarded

### Value

a matrix with gene names ,cox p value and HR

### Author(s)

Xiaojie Sun

### See Also

[point\\_cut](#);[surv\\_KM](#)

### Examples

```
surv_cox(exprSet_hub1,meta1,cut.point = TRUE,HRkeep = "all")
```

---

surv_KM	<i>surv_KM</i>
---------	----------------

---

**Description**

calculate log\_rank test p values for genes

**Usage**

```
surv_KM(  
  exprSet_hub,  
  meta,  
  cut.point = FALSE,  
  pvalue_cutoff = 0.05,  
  min_gn = 0.1  
)
```

**Arguments**

exprSet_hub	a tumor expression set for hubgenes
meta	meta data corresponds to expression set
cut.point	logical , use cut_point or not, if FALSE,use median by default
pvalue_cutoff	p value cut off ,0.05 by default
min_gn	Depending on the expression of a gene, there may be a large difference in the number of samples between the two groups, and if a smaller group of samples is less than 10 percent (default) of all, the gene will be discarded

**Value**

a vector with gene names and log\_rank p value

**Author(s)**

Xiaojie Sun

**See Also**

[point\\_cut](#); [surv\\_cox](#)

**Examples**

```
surv_KM(exprSet_hub1,meta1)  
surv_KM(exprSet_hub1,meta1,cut.point = TRUE)
```

---

trans_array	<i>trans_array</i>
-------------	--------------------

---

**Description**

transform rownames for microarray expression matrix

**Usage**

```
trans_array(exp, ids, from = "probe_id", to = "symbol")
```

**Arguments**

exp	TCGA or TCGA_Gtex expression set from gdc or xena
ids	data.frame with original rownames and new rownames
from	colname for original rownames
to	colname for new rownames

**Value**

a transformed expression set with new rownames

**Author(s)**

Xiaojie Sun

**See Also**

[trans\\_exp](#)

**Examples**

```
exp = matrix(1:50, nrow = 10)
rownames(exp) = paste0("g", 1:10)
ids = data.frame(probe_id = paste0("g", 1:10),
                 symbol = paste0("G", c(1:9, 9)))
trans_array(exp, ids)
```

---

trans_exp	<i>trans_exp</i>
-----------	------------------

---

### Description

transform rownames of TCGA or TCGA\_Gtex expression set from gdc or xena, from ensembl id to gene symbol

### Usage

```
trans_exp(exp, mrna_only = FALSE, lncrna_only = FALSE, gtex = FALSE)
```

### Arguments

exp	TCGA or TCGA_Gtex expression set from gdc or xena
mrna_only	only keep mrna rows in result
lncrna_only	only keep lncrna rows in result
gtex	logical, whether including Gtex data

### Value

a transformed expression set with symbol

### Author(s)

Xiaojie Sun

### See Also

[trans\\_array](#)

### Examples

```
exp = matrix(rnorm(1000), ncol = 10)
rownames(exp) = sample(mRNA_annot23$gene_id, 100)
colnames(exp) = c(paste0("TCGA", 1:5), paste0("GTEx", 1:5))
k = trans_exp(exp)
```

---

t_choose	<i>t_choose</i>
----------	-----------------

---

### Description

choose differential expressed genes by simple t.test

### Usage

```
t_choose(  
  genes,  
  exp,  
  group_list,  
  up_only = FALSE,  
  down_only = FALSE,  
  pvalue_cutoff = 0.05  
)
```

### Arguments

genes	a vector with some genes
exp	A numeric matrix
group_list	A factor with duplicated character or factor
up_only	keep up genes in the result only
down_only	keep down genes in the result only
pvalue_cutoff	p value cut off ,0.05 by default

### Value

a vector with differential expressed genes

### Author(s)

Xiaojie Sun

### Examples

```
exp = matrix(rnorm(1000),ncol = 10)  
rownames(exp) = sample(mRNA_annov23$gene_id,100)  
colnames(exp) = c(paste0("TCGA",1:5),paste0("GTEx",1:5))  
exp2 = trans_exp(exp)  
exp2[,1:5] = exp2[,1:5]+10  
group_list = rep(c("A","B"),each = 5)  
genes = sample(rownames(exp2),3)  
t_choose(genes,exp2,group_list)
```



---

union_all	<i>union_all</i>
-----------	------------------

---

**Description**

calculate union set for two or more elements

**Usage**

```
union_all(...)
```

**Arguments**

... some vectors or a list with some vectors

**Value**

vector

**Author(s)**

Xiaojie Sun

**See Also**

[intersect\\_all](#)

**Examples**

```
x1 = letters[1:4]
x2 = letters[3:6]
x3 = letters[3:4]
re =intersect_all(x1,x2,x3)
re2 = intersect_all(list(x1,x2,x3))
re3 = union_all(x1,x2,x3)
```

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