

Package ‘MAGeCKFlute’

April 12, 2022

Type Package

Title Integrative Analysis Pipeline for Pooled CRISPR Functional Genetic Screens

Version 1.14.0

Date 2020-10-26

Author Binbin Wang, Wubing Zhang, Feizhen Wu, Wei Li & X. Shirley Liu

Maintainer Wubing Zhang<Watson5bZhang@gmail.com>

Description CRISPR (clustered regularly interspaced short palindrome repeats) coupled with nuclease Cas9 (CRISPR/Cas9) screens represent a promising technology to systematically evaluate gene functions. Data analysis for CRISPR/Cas9 screens is a critical process that includes identifying screen hits and exploring biological functions for these hits in downstream analysis. We have previously developed two algorithms, MAGeCK and MAGeCK-VISPR, to analyze CRISPR/Cas9 screen data in various scenarios. These two algorithms allow users to perform quality control, read count generation and normalization, and calculate beta score to evaluate gene selection performance. In downstream analysis, the biological functional analysis is required for understanding biological functions of these identified genes with different screening purposes. Here, We developed MAGeCKFlute for supporting downstream analysis. MAGeCKFlute provides several strategies to remove potential biases within sgRNA-level read counts and gene-level beta scores. The downstream analysis with the package includes identifying essential, non-essential, and target-associated genes, and performing biological functional category analysis, pathway enrichment analysis and protein complex enrichment analysis of these genes. The package also visualizes genes in multiple ways to benefit users exploring screening data. Collectively, MAGeCKFlute enables accurate identification of essential, non-essential, and targeted genes, as well as their related biological functions. This vignette explains the use of the package and demonstrates typical workflows.

License GPL (>=3)

VignetteBuilder knitr

Depends R (>= 3.5)

Imports Biobase, clusterProfiler (>= 3.16.1), enrichplot, gridExtra, ggplot2, ggrepel, grDevices, grid, reshape2, stats, utils

Suggests biomaRt, BiocStyle, DOSE, dendextend, graphics, knitr, msigdb, pheatmap, png, pathview, scales, sva, testthat,

LazyData TRUE

NeedsCompilation no

biocViews FunctionalGenomics, CRISPR, PooledScreens, QualityControl, Normalization, GeneSetEnrichment, Pathways, Visualization, GeneTarget, KEGG

Encoding UTF-8

RoxygenNote 7.1.1

git_url <https://git.bioconductor.org/packages/MAGeCKFlute>

git_branch RELEASE_3_14

git_last_commit e03a467

git_last_commit_date 2021-10-26

Date/Publication 2022-04-12

R topics documented:

arrangePathview	3
BarView	5
BatchRemove	6
ConsistencyView	7
CutoffCalling	8
DensityDiffView	9
DensityView	10
enrich.GSE	11
enrich.HGT	13
enrich.ORT	14
EnrichAB	16
EnrichAnalyzer	17
EnrichedFilter	18
EnrichedGeneView	19
EnrichedView	20
EnrichSquare	22
FluteMLE	23
FluteRRA	25
getCols	27
getGeneAnn	28
getOrg	29
getOrtAnn	29
gsGetter	30
hclustView	31
HeatmapView	32
IdentBarView	33
IncorporateDepmap	34

MapRatesView	35
MAView	36
noEnrichPlot	38
normalize.loess	38
NormalizeBeta	39
OmitCommonEssential	41
RankView	42
ReadBeta	43
ReadGMT	44
ReadRRA	45
ReadsgRRA	46
ResembleDepmap	46
retrieve_gs	47
ScatterView	48
Selector	50
sgRankView	51
SquareView	52
TransGeneID	54
ViolinView	55
VolcanoView	56

Index	58
--------------	-----------

arrangePathview *Kegg pathway view and arrange grobs on page*

Description

Kegg pathway view and arrange grobs on page.

Usage

```

arrangePathview(
  genelist,
  pathways = c(),
  top = 4,
  ncol = 2,
  title = NULL,
  sub = NULL,
  organism = "hsa",
  output = ".",
  path.archive = ".",
  kegg.native = TRUE,
  verbose = TRUE
)
    
```

Arguments

genelist	a data frame with columns of ENTREZID, Control and Treatment. The columns of Control and Treatment represent gene score in Control and Treatment sample.
pathways	character vector, the KEGG pathway ID(s), usually 5 digit, may also include the 3 letter KEGG species code.
top	integer, specifying how many top enriched pathways to be visualized.
ncol	integer, specifying how many column of figures to be arranged in each page.
title	optional string, or grob.
sub	optional string, or grob.
organism	character, either the kegg code, scientific name or the common name of the target species. This applies to both pathway and gene.data or cpd.data. When KEGG ortholog pathway is considered, species="ko". Default species="hsa", it is equivalent to use either "Homo sapiens" (scientific name) or "human" (common name).
output	Path to save plot to.
path.archive	character, the directory of KEGG pathway data file (.xml) and image file (.png). Users may supply their own data files in the same format and naming convention of KEGG's (species code + pathway id, e.g. hsa04110.xml, hsa04110.png etc) in this directory. Default kegg.dir="." (current working directory).
kegg.native	logical, whether to render pathway graph as native KEGG graph (.png) or using graphviz layout engine (.pdf). Default kegg.native=TRUE.
verbose	Boolean

Value

plot on the current device

Author(s)

Wubing Zhang

Examples

```
file3 = file.path(system.file("extdata", package = "MAGeCKFlute"),
"testdata/mle.gene_summary.txt")
dd = ReadBeta(file3)
colnames(dd)[2:3] = c("Control", "Treatment")
# arrangePathview(dd, c("hsa00534"), title=NULL, sub=NULL, organism="hsa")
```

BarView

Bar plot

Description

Bar plot

Usage

```
BarView(  
  df,  
  x = "x",  
  y = "y",  
  fill = "#FC6665",  
  bar.width = 0.8,  
  position = "dodge",  
  dodge.width = 0.8,  
  main = NA,  
  xlab = NULL,  
  ylab = NA,  
  ...  
)
```

Arguments

df	A data frame.
x	A character, specifying the x-axis.
y	A character, specifying the y-axis.
fill	A character, specifying the fill color.
bar.width	A numeric, specifying the width of bar.
position	"dodge" (default), "stack", "fill".
dodge.width	A numeric, set the width in position_dodge.
main	A character, specifying the figure title.
xlab	A character, specifying the title of x-axis.
ylab	A character, specifying the title of y-axis.
...	Other parameters in geom_bar

Value

An object created by ggplot, which can be assigned and further customized.

Author(s)

Wubing Zhang

Examples

```
mdata = data.frame(group=letters[1:5], count=sample(1:100,5))
BarView(mdata, x = "group", y = "count")
```

BatchRemove

Batch effect removal

Description

Batch effect removal

Usage

```
BatchRemove(
  mat,
  batchMat,
  log2trans = FALSE,
  pca = TRUE,
  positive = FALSE,
  cluster = FALSE,
  outdir = NULL
)
```

Arguments

mat	A data frame, each row is a gene, and each column is a sample.
batchMat	A data frame, the first column should be ‘Samples’(matched colnames of mat) and the second column is ‘Batch’. The remaining columns could be Covariates.
log2trans	Boolean, specifying whether do logarithmic transformation before batch removal.
pca	Boolean, specifying whether return pca plot.
positive	Boolean, specifying whether all values should be positive.
cluster	Boolean, specifying whether perform hierarchical clustering.
outdir	Output directory for hierarchical cluster tree.

Value

A list contains two objects, including data and p.

Author(s)

Wubing Zhang

See Also

[ComBat](#)

Examples

```
edata = matrix(c(rnorm(2000, 5), rnorm(2000, 8)), 1000)
colnames(edata) = paste0("s", 1:4)
batchMat = data.frame(sample = colnames(edata), batch = rep(1:2, each = 2))
edata1 = BatchRemove(edata, batchMat)
print(edata1$p)
```

ConsistencyView

Visualize the estimate cell cycle compared to control.

Description

Estimate cell cycle time in different samples by linear fitting of beta scores.

Usage

```
ConsistencyView(
  beta,
  ctrlname,
  treatname,
  main = NULL,
  filename = NULL,
  width = 5,
  height = 4,
  ...
)
```

Arguments

beta	Data frame, which has columns of ctrlname and other samples.
ctrlname	A character, specifying the names of control samples.
treatname	A character, specifying the name of treatment samples.
main	As in 'plot'.
filename	Figure file name to create on disk. Default filename="NULL", which means no output.
width	As in ggsave.
height	As in ggsave.
...	Other available parameters in ggsave.

Value

An object created by ggplot, which can be assigned and further customized.

Author(s)

Wubing Zhang

Examples

```
file3 = file.path(system.file("extdata", package = "MAGeCKFlute"),
"testdata/mle.gene_summary.txt")
dd = ReadBeta(file3)
ConsistencyView(dd, ctrlname = "dms0", treatname = "plx")
```

CutoffCalling

Quantile of normal distribution.

Description

Compute cutoff from a normal-distributed vector.

Usage

```
CutoffCalling(d, scale = 1)
```

Arguments

d	A numeric vector.
scale	Boolean or numeric, specifying how many standard deviation will be used as cutoff.

Value

A numeric value.

Examples

```
CutoffCalling(rnorm(10000))
```

DensityDiffView	<i>Density plot</i>
-----------------	---------------------

Description

Plot the density of beta score deviations.

Usage

```
DensityDiffView(  
  beta,  
  ctrlname = "Control",  
  treatname = "Treatment",  
  main = NULL,  
  filename = NULL,  
  width = 5,  
  height = 4,  
  ...  
)
```

Arguments

beta	Data frame, including ctrlname and treatname as columns.
ctrlname	A character, specifying the name of control sample.
treatname	A character, specifying the name of treatment sample.
main	As in 'plot'.
filename	Figure file name to create on disk. Default filename="NULL", which means no output.
width	As in ggsave.
height	As in ggsave.
...	Other parameters in ggsave.

Value

An object created by ggplot, which can be assigned and further customized.

Author(s)

Wubing Zhang

Examples

```
file3 = file.path(system.file("extdata", package = "MAGeCKFlute"),
"testdata/mle.gene_summary.txt")
dd = ReadBeta(file3)
# Density plot of beta score deviation between control and treatment
DensityDiffView(dd, ctrlname = "dms0", treatname = "plx")
```

DensityView

Density plot for gene beta scores in Control and Treatment

Description

Plot the density of gene beta scores in two samples.

Usage

```
DensityView(
  beta,
  samples = NULL,
  main = NULL,
  xlab = "Beta Score",
  filename = NULL,
  width = 5,
  height = 4,
  ...
)
```

Arguments

beta	Data frame, including samples as columns.
samples	Character, specifying sample names in beta.
main	As in 'plot'.
xlab	As in 'plot'.
filename	Figure file name to create on disk. Default filename="NULL", which means don't save the figure on disk.
width	As in ggsave.
height	As in ggsave.
...	Other available parameters in ggsave.

Value

An object created by ggplot, which can be assigned and further customized.

Author(s)

Wubing Zhang

See Also[ViolinView](#)**Examples**

```
file3 = file.path(system.file("extdata", package = "MAGeCKFlute"),
"testdata/mle.gene_summary.txt")
dd = ReadBeta(file3)
DensityView(dd, samples=c("dms0", "plx"))
#or
DensityView(dd[, c("dms0", "plx")])
```

enrich.GSE

*Gene set enrichment analysis***Description**

A universal gene set enrichment analysis tools

Usage

```
enrich.GSE(
  geneList,
  keytype = "Symbol",
  type = "GOBP",
  organism = "hsa",
  pvalueCutoff = 0.25,
  limit = c(2, 200),
  gmtpath = NULL,
  by = "fgsea",
  verbose = TRUE,
  ...
)
```

Arguments

geneList	A order ranked numeric vector with geneid as names
keytype	"Entrez", "Ensembl", or "Symbol"
type	Molecular signatures for testing, available datasets include Pathway (KEGG, REACTOME, C2_CP), GO (GOBP, GOCC, GOMF), MSIGDB (C1, C2 (C2_CP (C2_CP_PID, C2_CP_BIOCARTA), C2_CGP), C3 (C3_MIR, C3_TFT), C4, C6, C7, HALLMARK) and Complex (CORUM). Any combination of them are also accessible (e.g. 'GOBP+GOMF+KEGG+REACTOME')

organism	'hsa' or 'mmu'
pvalueCutoff	FDR cutoff
limit	A two-length vector, specifying the minimal and maximal size of gene sets for enrichment analysis
gmtpath	The path to customized gmt file
by	One of 'fgsea' or 'DOSE'
verbose	Boolean
...	Other parameter

Value

An enrichResult instance

Author(s)

Wubing Zhang

See Also

[enrich.HGT](#)

[enrich.ORT](#)

[EnrichAnalyzer](#)

[gseGO](#)

[gseKEGG](#)

[GSEA](#)

[enrichResult-class](#)

Examples

```
data(geneList, package = "DOSE")
## Not run:
  enrichRes = enrich.GSE(geneList, keytype = "entrez")
  head(slot(enrichRes, "result"))

## End(Not run)
```

enrich.HGT

Do enrichment analysis using hypergeometric test

Description

Do enrichment analysis using hypergeometric test

Usage

```
enrich.HGT(
  geneList,
  keytype = "Symbol",
  type = "GOBP",
  organism = "hsa",
  pvalueCutoff = 0.25,
  limit = c(2, 200),
  universe = NULL,
  gmtpath = NULL,
  verbose = TRUE,
  ...
)
```

Arguments

geneList	A numeric vector with gene as names
keytype	"Entrez", "Ensembl", or "Symbol"
type	Molecular signatures for testing, available datasets include Pathway (KEGG, REACTOME, C2_CP), GO (GOBP, GOCC, GOMF), MSIGDB (C1, C2 (C2_CP (C2_CP_PID, C2_CP_BIOCARTA), C2_CGP), C3 (C3_MIR, C3_TFT), C4, C6, C7, HALLMARK) and Complex (CORUM). Any combination of them are also accessible (e.g. 'GOBP+GOMF+KEGG+REACTOME')
organism	'hsa' or 'mmu'
pvalueCutoff	FDR cutoff
limit	A two-length vector, specifying the minimal and maximal size of gene sets for enrichment analysis
universe	A character vector, specifying the background genelist, default is whole genome
gmtpath	The path to customized gmt file
verbose	Boolean
...	Other parameter

Value

An enrichResult instance.

Author(s)

Wubing Zhang

See Also[enrich.GSE](#)[enrich.ORT](#)[EnrichAnalyzer](#)[enrichResult-class](#)**Examples**

```
data(geneList, package = "DOSE")
genes <- geneList[1:300]
enrichRes <- enrich.HGT(genes, type = "KEGG", keytype = "entrez")
head(slot(enrichRes, "result"))
```

`enrich.ORT`*Do enrichment analysis using over-representation test*

Description

Do enrichment analysis using over-representation test

Usage

```
enrich.ORT(
  geneList,
  keytype = "Symbol",
  type = "GOBP",
  organism = "hsa",
  pvalueCutoff = 0.25,
  limit = c(2, 200),
  universe = NULL,
  gmtpath = NULL,
  verbose = TRUE,
  ...
)
```

Arguments

<code>geneList</code>	A numeric vector with gene as names.
<code>keytype</code>	"Entrez" or "Symbol".

type	Molecular signatures for testing, available datasets include Pathway (KEGG, REACTOME, C2_CP), GO (GOBP, GOCC, GOMF), MSIGDB (C1, C2 (C2_CP (C2_CP_PID, C2_CP_BIOCARTA), C2_CGP), C3 (C3_MIR, C3_TFT), C4, C6, C7, HALLMARK) and Complex (CORUM). Any combination of them are also accessible (e.g. 'GOBP+GOMF+KEGG+REACTOME').
organism	'hsa' or 'mmu'.
pvalueCutoff	FDR cutoff.
limit	A two-length vector, specifying the minimal and maximal size of gene sets for enrichment analysis.
universe	A character vector, specifying the background genelist, default is whole genome.
gmtpath	The path to customized gmt file.
verbose	Boolean
...	Other parameter

Value

An enrichedResult instance.

Author(s)

Wubing Zhang

See Also

[enrich.HGT](#)

[enrich.GSE](#)

[EnrichAnalyzer](#)

[enrichGO](#)

[enrichKEGG](#)

[enricher](#)

[enrichResult-class](#)

Examples

```
data(geneList, package = "DOSE")
genes <- geneList[1:100]
enrichedRes <- enrich.ORT(genes, keytype = "entrez")
head(slot(enrichedRes, "result"))
```

Description

Do enrichment analysis for selected genes, in which positive selection and negative selection are termed as GroupA and GroupB

Usage

```
EnrichAB(
  data,
  pvalue = 0.25,
  enrich_method = "ORT",
  organism = "hsa",
  limit = c(1, 120),
  filename = NULL,
  out.dir = ".",
  width = 6.5,
  height = 4,
  verbose = TRUE,
  ...
)
```

Arguments

data	A data frame.
pvalue	Pvalue cutoff.
enrich_method	One of "ORT"(Over-Representing Test) and "HGT"(HyperGemetric test).
organism	"hsa" or "mmu".
limit	A two-length vector (default: c(1, 120)), specifying the min and max size of pathways for enrichent analysis.
filename	Suffix of output file name.
out.dir	Path to save plot to (combined with filename).
width	As in ggsave.
height	As in ggsave.
verbose	Boolean
...	Other available parameters in ggsave.

Value

A list containing enrichment results for each group genes. This list contains eight items, which contain subitems of gridPlot and enrichRes.

Author(s)

Wubing Zhang

EnrichAnalyzer

*Enrichment analysis***Description**

Enrichment analysis

Usage

```

EnrichAnalyzer(
  geneList,
  keytype = "Symbol",
  type = "Pathway+GOBP",
  method = "HGT",
  organism = "hsa",
  pvalueCutoff = 0.25,
  limit = c(2, 200),
  universe = NULL,
  filter = FALSE,
  gmtpath = NULL,
  verbose = TRUE
)

```

Arguments

geneList	A numeric vector with gene as names.
keytype	"Entrez" or "Symbol".
type	Molecular signatures for testing, available datasets include Pathway (KEGG, REACTOME, C2_CP), GO (GOBP, GOCC, GOMF), MSIGDB (C1, C2 (C2_CP (C2_CP_PID, C2_CP_BIOCARTA), C2_CGP), C3 (C3_MIR, C3_TFT), C4, C6, C7, HALLMARK) and Complex (CORUM). Any combination of them are also accessible (e.g. 'GOBP+GOMF+KEGG+REACTOME').
method	One of "ORT"(Over-Representing Test), "GSEA"(Gene Set Enrichment Analysis), and "HGT"(HyperGemetric test).
organism	'hsa' or 'mmu'.
pvalueCutoff	FDR cutoff.
limit	A two-length vector (default: c(2, 200)), specifying the minimal and maximal size of gene sets for enrichment analysis.
universe	A character vector, specifying the background genelist, default is whole genome.
filter	Boolean, specifying whether filter out redundancies from the enrichment results.
gmtpath	The path to customized gmt file.
verbose	Boolean

Value

enrichRes is an enrichResult instance.

Author(s)

Wubing Zhang

See Also

[enrich.GSE](#)
[enrich.ORT](#)
[enrich.HGT](#)
[enrichResult-class](#)

Examples

```
data(geneList, package = "DOSE")
## Not run:
  keggA = EnrichAnalyzer(geneList[1:500], keytype = "entrez")
  head(keggA@result)

## End(Not run)
```

EnrichedFilter

Simplify the enrichment results based on Jaccard index

Description

Simplify the enrichment results based on Jaccard index

Usage

```
EnrichedFilter(enrichment = enrichment, cutoff = 0.8)
```

Arguments

enrichment	A data frame of enrichment result.
cutoff	A numeric, specifying the cutoff of Jaccard index between two pathways.

Value

A data frame.

Author(s)

Yihan Xiao

Examples

```

data(geneList, package = "DOSE")
## Not run:
  enrichRes <- enrich.HGT(geneList, keytype = "entrez")
  EnrichedFilter(enrichRes)

## End(Not run)

```

EnrichedGeneView *Visualize enriched pathways and genes in those pathways*

Description

Visualize enriched pathways and genes in those pathways

Usage

```

EnrichedGeneView(
  enrichment,
  geneList,
  rank_by = "p.adjust",
  top = 5,
  bottom = 0,
  keytype = "Symbol",
  gene_cutoff = c(-log2(1.5), log2(1.5)),
  custom_gene = NULL,
  charLength = 40,
  filename = NULL,
  width = 7,
  height = 5,
  ...
)

```

Arguments

enrichment	A data frame of enrichment result or an <code>enrichResult</code> object.
geneList	A numeric <code>geneList</code> used in enrichment analysis.
rank_by	"p.adjust" or "NES", specifying the indices for ranking pathways.
top	An integer, specifying the number of positively enriched terms to show.
bottom	An integer, specifying the number of negatively enriched terms to show.
keytype	"Entrez" or "Symbol".
gene_cutoff	A two-length numeric vector, specifying cutoff for genes to show.
custom_gene	A character vector (gene names), customizing genes to show.
charLength	Integer, specifying max length of enriched term name to show as coordinate label.

filename	Figure file name to create on disk. Default filename="NULL", which means no output.
width	As in ggsave.
height	As in ggsave.
...	Other available parameters in ggsave.

Value

An object created by ggplot, which can be assigned and further customized.

Author(s)

Wubing Zhang

Examples

```
data(geneList, package = "DOSE")
## Not run:
  enrichRes <- enrich.GSE(geneList, keytype = "Entrez")
  EnrichedGeneView(enrichment=slot(enrichRes, "result"), geneList, keytype = "Entrez")

## End(Not run)
```

EnrichedView

View enriched terms

Description

Grid plot for enriched terms

Usage

```
EnrichedView(
  enrichment,
  rank_by = "pvalue",
  mode = 1,
  subset = NULL,
  top = 0,
  bottom = 0,
  x = "LogFDR",
  charLength = 40,
  filename = NULL,
  width = 7,
  height = 4,
  ...
)
```

Arguments

enrichment	A data frame of enrichment result, with columns of ID, Description, p.adjust and NES.
rank_by	"pvalue" or "NES", specifying the indices for ranking pathways.
mode	1 or 2.
subset	A vector of pathway ids.
top	An integer, specifying the number of upregulated terms to show.
bottom	An integer, specifying the number of downregulated terms to show.
x	Character, "NES", "LogP", or "LogFDR", indicating the variable on the x-axis.
charLength	Integer, specifying max length of enriched term name to show as coordinate lab.
filename	Figure file name to create on disk. Default filename="NULL".
width	As in ggsave.
height	As in ggsave.
...	Other available parameters in ggsave.

Value

An object created by ggplot, which can be assigned and further customized.

Author(s)

Wubing Zhang

See Also

[EnrichedView](#)

Examples

```
data(geneList, package = "DOSE")
## Not run:
  enrichRes = enrich.GSE(geneList, organism="hsa")
  EnrichedView(enrichRes, top = 5, bottom = 5)

## End(Not run)
```

Description

Do enrichment analysis for selected treatment related genes in 9-squares

Usage

```
EnrichSquare(
  beta,
  id = "Gene",
  keytype = "Symbol",
  x = "Control",
  y = "Treatment",
  pvalue = 0.05,
  enrich_method = "ORT",
  organism = "hsa",
  limit = c(1, 120),
  filename = NULL,
  out.dir = ".",
  width = 6.5,
  height = 4,
  verbose = TRUE,
  ...
)
```

Arguments

beta	Data frame, with columns of "Gene", "group", and "Diff".
id	A character, indicating the gene column in the data.
keytype	A character, "Symbol" or "Entrez".
x	A character, indicating the x-axis in the 9-square scatter plot.
y	A character, indicating the y-axis in the 9-square scatter plot.
pvalue	Pvalue cutoff.
enrich_method	One of "ORT"(Over-Representing Test) and "HGT"(HyperGemetric test).
organism	"hsa" or "mmu".
limit	A two-length vector (default: c(1, 120)), specifying the min and max size of pathways for enrichent analysis.
filename	Suffix of output file name. NULL(default) means no output.
out.dir	Path to save plot to (combined with filename).
width	As in ggsave.
height	As in ggsave.
verbose	Boolean.
...	Other available parameters in ggsave.

Value

A list containing enrichment results for each group genes. Each item in the returned list has two sub items:

gridPlot an object created by ggplot, which can be assigned and further customized.
enrichRes a enrichResult instance.

Author(s)

Wubing Zhang

FluteMLE

Downstream analysis based on MAGeCK-MLE result

Description

Integrative analysis pipeline using the gene summary table in MAGeCK MLE results

Usage

```
FluteMLE(  
  gene_summary,  
  treatname,  
  ctrlname = "Depmap",  
  keytype = "Symbol",  
  organism = "hsa",  
  incorporateDepmap = FALSE,  
  cell_lines = NA,  
  lineages = "All",  
  norm_method = "cell_cycle",  
  posControl = NULL,  
  omitEssential = FALSE,  
  top = 10,  
  toplabels = NA,  
  scale_cutoff = 2,  
  limit = c(0, 200),  
  pvalueCutoff = 0.25,  
  enrich_method = "ORT",  
  proj = NA,  
  width = 10,  
  height = 7,  
  outdir = ".",  
  pathview.top = 4,  
  verbose = TRUE  
)
```

Arguments

gene_summary	A data frame or a file path to gene summary file generated by MAGeCK-MLE.
treatname	A character vector, specifying the names of treatment samples.
ctrlname	A character vector, specifying the names of control samples. If there is no controls in your CRISPR screen, you can specify "Depmap" as ctrlname and set 'incorporateDepmap=TRUE'.
keytype	"Entrez" or "Symbol".
organism	"hsa" or "mmu".
incorporateDepmap	Boolean, indicating whether incorporate Depmap data into analysis.
cell_lines	A character vector, specifying the cell lines in Depmap to be considered.
lineages	A character vector, specifying the lineages in Depmap to be considered.
norm_method	One of "none", "cell_cycle" (default) or "loess".
posControl	A character vector, specifying a list of positive control gene symbols.
omitEssential	Boolean, indicating whether omit common essential genes from the downstream analysis.
top	An integer, specifying number of top selected genes to be labeled in rank figure.
toplabels	A character vector, specifying interested genes to be labeled in rank figure.
scale_cutoff	Boolean or numeric, specifying how many standard deviation will be used as cutoff.
limit	A two-length vector, specifying the minimal and maximal size of gene sets for enrichment analysis.
pvalueCutoff	A numeric, specifying pvalue cutoff of enrichment analysis, default 1.
enrich_method	One of "ORT"(Over-Representing Test) and "HGT"(HyperGemetric test).
proj	A character, indicating the prefix of output file name, which can't contain special characters.
width	The width of summary pdf in inches.
height	The height of summary pdf in inches.
outdir	Output directory on disk.
pathview.top	Integer, specifying the number of pathways for pathview visualization.
verbose	Boolean

Details

MAGeCK-MLE can be used to analyze screen data from multi-conditioned experiments. MAGeCK-MLE also normalizes the data across multiple samples, making them comparable to each other. The most important output of MAGeCK MLE is 'gene_summary' file, which includes the beta scores of multiple conditions and the associated statistics. The 'beta score' for each gene describes how the gene is selected: a positive beta score indicates a positive selection, and a negative beta score indicates a negative selection.

The downstream analysis includes identifying essential, non-essential, and target-associated genes, and performing biological functional category analysis and pathway enrichment analysis of these genes. The function also visualizes genes in the context of pathways to benefit users exploring screening data.

Value

All of the pipeline results is output into the `out.dir/MAGeCKFlute_proj`, which includes a pdf file and many folders. The pdf file 'FluteMLE_proj_norm_method.pdf' is the summary of pipeline results. For each section in this pipeline, figures and useful data are outputted to corresponding subfolders.

- QC: Quality control
- Selection: Positive selection and negative selection.
- Enrichment: Enrichment analysis for positive and negative selection genes.
- PathwayView: Pathway view for top enriched pathways.

Author(s)

Wubing Zhang

See Also

[FluteRRA](#)

Examples

```
file3 = file.path(system.file("extdata", package = "MAGeCKFlute"),
"testdata/mle.gene_summary.txt")
## Not run:
# functional analysis for MAGeCK MLE results
FluteMLE(file3, treatname = "plx", ctrlname = "dms0", proj = "PLX")

## End(Not run)
```

FluteRRA

Downstream analysis based on MAGeCK-RRA result

Description

Integrative analysis pipeline using the gene summary table in MAGeCK RRA results

Usage

```
FluteRRA(  
  gene_summary,  
  sgrna_summary = NULL,  
  keytype = "Symbol",  
  organism = "hsa",  
  incorporateDepmap = TRUE,  
  cell_lines = NA,  
  lineages = "All",
```

```

omitEssential = FALSE,
top = 5,
toplabels = NULL,
scale_cutoff = 2,
limit = c(2, 200),
pvalueCutoff = 0.25,
proj = NA,
width = 12,
height = 6,
outdir = ".",
verbose = TRUE
)

```

Arguments

gene_summary	A file path or a data frame of gene summary data.
sgrna_summary	A file path or a data frame of sgRNA summary data.
keytype	"Entrez" or "Symbol".
organism	"hsa" or "mmu".
incorporateDepmap	Boolean, indicating whether incorporate Depmap data into analysis.
cell_lines	A character vector, specifying the cell lines in Depmap to be considered.
lineages	A character vector, specifying the lineages in Depmap to be considered.
omitEssential	Boolean, indicating whether omit common essential genes from the downstream analysis.
top	An integer, specifying number of top selected genes to be labeled in rank figure.
toplabels	A character vector, specifying interested genes to be labeled in rank figure.
scale_cutoff	Boolean or numeric, specifying how many standard deviation will be used as cutoff.
limit	A two-length vector, specifying the minimal and maximal size of gene sets for enrichment analysis.
pvalueCutoff	A numeric, specifying pvalue cutoff of enrichment analysis, default 1.
proj	A character, indicating the prefix of output file name.
width	The width of summary pdf in inches.
height	The height of summary pdf in inches.
outdir	Output directory on disk.
verbose	Boolean

Details

MAGeCK RRA allows for the comparison between two experimental conditions. It can identify genes and sgRNAs are significantly selected between the two conditions. The most important output of MAGeCK RRA is the file 'gene_summary.txt'. MAGeCK RRA will output both the negative score and positive score for each gene. A smaller score indicates higher gene importance. MAGeCK

RRA will also output the statistical value for the scores of each gene. Genes that are significantly positively and negatively selected can be identified based on the p-value or FDR.

The downstream analysis of this function includes identifying positive and negative selection genes, and performing biological functional category analysis and pathway enrichment analysis of these genes.

Value

All of the pipeline results is output into the `out.dir/proj_Results`, which includes a pdf file and a folder named 'RRA'.

Author(s)

Wubing Zhang

See Also

[FluteMLE](#)

Examples

```
file1 = file.path(system.file("extdata", package = "MAGeCKFlute"),
"testdata/rra.gene_summary.txt")
file2 = file.path(system.file("extdata", package = "MAGeCKFlute"),
"testdata/rra.sgrna_summary.txt")
## Not run:
# Run the FluteRRA pipeline
FluteRRA(file1, file2, proj="PLX", organism="hsa", incorporateDepmap = FALSE,
scale_cutoff = 1, outdir = "./")
## End(Not run)
```

getCols

Map values to colors

Description

Map values to colors

Usage

```
getCols(x, palette = 1)
```

Arguments

x	A numeric vector.
palette	diverge, rainbow, sequential

Value

A vector of colors corresponding to input vector.

Author(s)

Wubing Zhang

Examples

```
getCols(1:4)
```

getGeneAnn	<i>Retrieve gene annotations from the NCBI, HNSC, and Uniprot databases.</i>
------------	--

Description

Retrieve gene annotations from the NCBI, HNSC, and Uniprot databases.

Usage

```
getGeneAnn(org = "hsa", update = FALSE)
```

Arguments

org	Character, hsa (default), bta, cfa, mmu, ptr, rno, ssc are optional.
update	Boolean, indicating whether download current annotation.

Value

A data frame.

Author(s)

Wubing Zhang

Examples

```
## Not run:  
ann = getGeneAnn("hsa")  
head(ann)  
  
## End(Not run)
```

getOrg *Get the kegg code of specific mammalia organism.*

Description

Get the kegg code of specific mammalia organism.

Usage

```
getOrg(organism)
```

Arguments

organism	Character, KEGG species code, or the common species name. For all potential values check: <code>data(bods); bods</code> . Default <code>org="hsa"</code> , and can also be "human" (case insensitive).
----------	--

Value

A list containing three elements:

org	species
pkgannotation	package name

Author(s)

Wubing Zhang

Examples

```
ann = getOrg("human")
print(ann$pkg)
```

getOrtAnn *Retrieve reference orthologs annotation.*

Description

Retrieve reference orthologs annotation.

Usage

```
getOrtAnn(fromOrg = "mmu", toOrg = "hsa", update = FALSE)
```

Arguments

fromOrg	Character, hsa (default), bta, cfa, mmu, ptr, rno, ssc are optional.
toOrg	Character, hsa (default), bta, cfa, mmu, ptr, rno, ssc are optional.
update	Boolean, indicating whether download recent annotation from NCBI.

Value

A data frame.

Author(s)

Wubing Zhang

Examples

```
## Not run:
ann = getOrtAnn("mmu", "hsa")
head(ann)

## End(Not run)
```

gsGetter

Extract pathway annotation from GMT file.

Description

Extract pathway annotation from GMT file.

Usage

```
gsGetter(
  gmtpath = NULL,
  type = "All",
  limit = c(0, Inf),
  organism = "hsa",
  update = FALSE
)
```

Arguments

gmtpath	The path to customized gmt file.
type	Molecular signatures for testing, available datasets include Pathway (KEGG, REACTOME, C2_CP:PID, C2_CP:BIOCARTA), GO (GOBP, GOCC, GOMF), MSIGDB (C1, C2 (C2_CP (C2_CP:PID, C2_CP:BIOCARTA), C2_CGP), C3 (C3_MIR, C3_TFT), C4 (C4_CGN, C4_CM), C5 (C5_BP, C5_CC, C5_MF), C6, C7, H) and Complex (CORUM). Any combination of them are also accessible (e.g. 'GOBP+GOMF+KEGG+REACTOME').

limit	A two-length vector, specifying the minimal and maximal size of gene sets to load.
organism	'hsa' or 'mmu'.
update	Boolean, indicating whether update the gene sets from source database.

Value

A three-column data frame.

Author(s)

Wubing Zhang

Examples

```
gene2path = gsGetter(type = "REACTOME+KEGG")
head(gene2path)
```

hclustView

Cluster and view cluster tree

Description

Cluster and view cluster tree

Usage

```
hclustView(
  d,
  method = "average",
  label_cols = NULL,
  bar_cols = NULL,
  main = NA,
  xlab = NA,
  horiz = TRUE,
  ...
)
```

Arguments

d	A dissimilarity structure as produced by dist.
method	The agglomeration method to be used. This should be (an unambiguous abbreviation of) one of "ward.D", "ward.D2", "single", "complete", "average" (= UPGMA), "mcquitty" (= WPGMA), "median" (= WPGMC) or "centroid" (= UPGMC).
label_cols	A vector to be used as label's colors for the dendrogram.

<code>bar_cols</code>	Either a vector or a matrix, which will be plotted as a colored bar.
<code>main</code>	As in 'plot'.
<code>xlab</code>	As in 'plot'.
<code>horiz</code>	Logical indicating if the dendrogram should be drawn horizontally or not.
<code>...</code>	Arguments to be passed to methods, such as graphical parameters (see <code>par</code>).

Value

Plot figure on open device.

Author(s)

Wubing Zhang

Examples

```
label_cols = rownames(USArrests)
hclustView(dist(USArrests), label_cols=label_cols, bar_cols=label_cols)
```

HeatmapView

Draw heatmap

Description

Draw heatmap

Usage

```
HeatmapView(
  mat,
  limit = c(-2, 2),
  na_col = "gray70",
  colPal = rev(colorRampPalette(c("#c12603", "white", "#0073B6"), space = "Lab")(199)),
  filename = NA,
  width = NA,
  height = NA,
  ...
)
```

Arguments

<code>mat</code>	Matrix like object, each row is gene and each column is sample.
<code>limit</code>	Max value in heatmap
<code>na_col</code>	Color for missing values
<code>colPal</code>	<code>colorRampPalette</code> .

filename	File path where to save the picture.
width	Manual option for determining the output file width in inches.
height	Manual option for determining the output file height in inches.
...	Other parameters in pheatmap.

Value

Invisibly a pheatmap object that is a list with components.

Author(s)

Wubing Zhang

Examples

```
file3 = file.path(system.file("extdata", package = "MAGeCKFlute"),
"testdata/mle.gene_summary.txt")
dd = ReadBeta(file3)
gg = cor(dd[,2:ncol(dd)])
HeatmapView(gg, display_numbers = TRUE)
```

IdentBarView

Identical bar plot

Description

Identical bar plot

Usage

```
IdentBarView(
  gg,
  x = "x",
  y = "y",
  fill = c("#CF3C2B", "#394E80"),
  main = NULL,
  xlab = NULL,
  ylab = NULL,
  filename = NULL,
  width = 5,
  height = 4,
  ...
)
```

Arguments

<code>gg</code>	A data frame.
<code>x</code>	A character, indicating column (in <code>countSummary</code>) of x-axis.
<code>y</code>	A character, indicating column (in <code>countSummary</code>) of y-axis.
<code>fill</code>	A character, indicating fill color of all bars.
<code>main</code>	A character, specifying the figure title.
<code>xlab</code>	A character, specifying the title of x-axis.
<code>ylab</code> ,	A character, specifying the title of y-axis.
<code>filename</code>	Figure file name to create on disk. Default filename="NULL", which means don't save the figure on disk.
<code>width</code>	As in <code>ggsave</code> .
<code>height</code>	As in <code>ggsave</code> .
<code>...</code>	Other available parameters in <code>ggsave</code> .

Value

An object created by `ggplot`, which can be assigned and further customized.

Author(s)

Wubing Zhang

Examples

```
file4 = file.path(system.file("extdata", package = "MAGeCKFlute"),
"testdata/countsummary.txt")
countsummary = read.delim(file4, check.names = FALSE)
IdentBarView(countsummary, x="Label", y="Reads")
```

IncorporateDepmap

Incorporate Depmap screen into analysis

Description

Incorporate Depmap screen into analysis

Usage

```
IncorporateDepmap(
  dd,
  symbol = "id",
  cell_lines = NA,
  lineages = "All",
  na.rm = FALSE
)
```

Arguments

<code>dd</code>	A data frame.
<code>symbol</code>	A character, specifying the column name of gene symbols in the data frame.
<code>cell_lines</code>	A character vector, specifying the cell lines in Depmap to be considered.
<code>lineages</code>	A character vector, specifying the lineages in Depmap to be considered.
<code>na.rm</code>	Boolean, indicating whether removing NAs from the results.

Value

A data frame with Depmap column attached.

Author(s)

Wubing Zhang

Examples

```
file1 = file.path(system.file("extdata", package = "MAGeCKFlute"),
"testdata/rra.gene_summary.txt")
gdata = ReadRRA(file1)
head(gdata)
## Not run:
  gdata = IncorporateDepmap(gdata)
  head(gdata)

## End(Not run)
```

MapRatesView

View mapping ratio

Description

View mapping ratio of each sample

Usage

```
MapRatesView(
  countSummary,
  Label = "Label",
  Reads = "Reads",
  Mapped = "Mapped",
  filename = NULL,
  width = 5,
  height = 4,
  ...
)
```

Arguments

countSummary	A data frame, which contains columns of 'Label', 'Reads', and 'Mapped'
Label	A character, indicating column (in countSummary) of sample names.
Reads	A character, indicating column (in countSummary) of total reads.
Mapped	A character, indicating column (in countSummary) of mapped reads.
filename	Figure file name to create on disk. Default filename="NULL", which means don't save the figure on disk.
width	As in ggsave.
height	As in ggsave.
...	Other available parameters in ggsave.

Value

An object created by ggplot, which can be assigned and further customized.

Author(s)

Wubing Zhang

Examples

```
file4 = file.path(system.file("extdata", package = "MAGeCKFlute"),
"testdata/countsummary.txt")
countsummary = read.delim(file4, check.names = FALSE)
MapRatesView(countsummary)
```

MAView

*M*Aplot of gene beta scores

Description

MAplot of gene beta scores in Control vs Treatment

Usage

```
MAView(
  beta,
  ctrlname = "Control",
  treatname = "Treatment",
  main = NULL,
  show.statistics = TRUE,
  add.smooth = TRUE,
  lty = 1,
  smooth.col = "red",
```

```

    plot.method = c("loess", "lm", "glm", "gam"),
    filename = NULL,
    width = 5,
    height = 4,
    ...
)

```

Arguments

beta	Data frame, including ctrlname and treatname as columns.
ctrlname	Character vector, specifying the name of control sample.
treatname	Character vector, specifying the name of treatment sample.
main	As in plot.
show.statistics	Show statistics .
add.smooth	Whether add a smooth line to the plot.
lty	Line type for smooth line.
smooth.col	Color of smooth line.
plot.method	A string specifying the method to fit smooth line, which should be one of "loess" (default), "lm", "glm" and "gam".
filename	Figure file name to create on disk. Default filename="NULL", which means don't save the figure on disk.
width	As in ggsave.
height	As in ggsave.
...	Other available parameters in function 'ggsave'.

Value

An object created by ggplot, which can be assigned and further customized.

Author(s)

Wubing Zhang

Examples

```

file3 = file.path(system.file("extdata", package = "MAGeCKFlute"),
"testdata/mle.gene_summary.txt")
dd = ReadBeta(file3)
MAView(dd, ctrlname = "dms0", treatname = "plx")

```

noEnrichPlot	<i>Blank figure</i>
--------------	---------------------

Description

Blank figure

Usage

```
noEnrichPlot(main = "No enriched terms")
```

Arguments

main	The title of figure.
------	----------------------

Value

An object created by ggplot, which can be assigned and further customized.

Author(s)

Wubing Zhang

normalize.loess	<i>normalize.loess</i>
-----------------	------------------------

Description

Loess normalization method.

Usage

```
normalize.loess(  
  mat,  
  subset = sample(1:(dim(mat)[1]), min(c(5000, nrow(mat)))),  
  epsilon = 10^-2,  
  maxit = 1,  
  log.it = FALSE,  
  verbose = TRUE,  
  span = 2/3,  
  family.loess = "symmetric",  
  ...  
)
```

Arguments

mat	A matrix with columns containing the values of the chips to normalize.
subset	A subset of the data to fit a loess to.
epsilon	A tolerance value (supposed to be a small value - used as a stopping criterion).
maxit	Maximum number of iterations.
log.it	Logical. If TRUE it takes the log2 of mat.
verbose	Logical. If TRUE displays current pair of chip being worked on.
span	Parameter to be passed the function loess
family.loess	Parameter to be passed the function loess . "gaussian" or "symmetric" are acceptable values for this parameter.
...	Any of the options of <code>normalize.loess</code> you would like to modify (described above).

Value

A matrix similar as mat.

Author(s)

Wubing Zhang

See Also

[loess](#)

[NormalizeBeta](#)

Examples

```
file3 = file.path(system.file("extdata", package = "MAGeCKFlute"),
"testdata/mle.gene_summary.txt")
dd = ReadBeta(file3)
beta_loess = normalize.loess(dd[,c("dms0", "plx")])
```

NormalizeBeta

Normalize gene beta scores

Description

Two normalization methods are available. `cell_cycle` method normalizes gene beta scores based on positive control genes in CRISPR screening. `loess` method normalizes gene beta scores using `loess`.

Usage

```
NormalizeBeta(
  beta,
  id = 1,
  method = "cell_cycle",
  posControl = NULL,
  samples = NULL
)
```

Arguments

<code>beta</code>	Data frame.
<code>id</code>	An integer specifying the column of gene.
<code>method</code>	Character, one of 'cell_cycle' (default) and 'loess'. or character string giving the name of the table column containing the gene names.
<code>posControl</code>	A character vector, specifying a list of positive control genes.
<code>samples</code>	Character vector, specifying the sample names in <i>beta</i> columns. If NULL (default), take all <i>beta</i> columns as samples.

Details

In CRISPR screens, cells treated with different conditions (e.g., with or without drug) may have different proliferation rates. So it's necessary to normalize the proliferation rate based on defined positive control genes among samples. After normalization, the beta scores are comparable across samples. loess is another optional normalization method, which is used to normalize array data before.

Value

A data frame with same format as input data *beta*.

Author(s)

Wubing Zhang

Examples

```
file3 = file.path(system.file("extdata", package = "MAGeCKFlute"),
"testdata/mle.gene_summary.txt")
dd = ReadBeta(file3)
#Cell Cycle normalization
dd_essential = NormalizeBeta(dd, samples=c("dms0", "plx"), method="cell_cycle")
head(dd_essential)

#Optional loess normalization (not recommended)
dd_loess = NormalizeBeta(dd, samples=c("dms0", "plx"), method="loess")
head(dd_loess)
```

OmitCommonEssential *Omit common essential genes based on depmap data*

Description

Omit common essential genes based on depmap data

Usage

```
OmitCommonEssential(dd, symbol = "id", lineages = "All", dependency = -0.5)
```

Arguments

dd	A data frame.
symbol	A character, specifying the column name of gene symbols in the data frame.
lineages	A character vector, specifying the lineages used for common essential gene selection.
dependency	A numeric, specifying the threshold for common essential gene selection.

Value

A data frame.

Author(s)

Wubing Zhang

Examples

```
file1 = file.path(system.file("extdata", package = "MAGeCKFlute"),
"testdata/rra.gene_summary.txt")
gdata = ReadRRA(file1)
dim(gdata)
## Not run:
rra.omit = OmitCommonEssential(gdata)
dim(rra.omit)

## End(Not run)
```

RankView

*View the rank of gene points***Description**

Rank all genes according to beta score deviation, and label top and bottom meaningful genes. Some other interested genes can be labeled too.

Usage

```
RankView(
  rankdata,
  genelist = NULL,
  top = 10,
  bottom = 10,
  cutoff = NULL,
  main = NULL,
  filename = NULL,
  width = 5,
  height = 4,
  ...
)
```

Arguments

rankdata	Numeric vector, with gene as names.
genelist	Character vector, specifying genes to be labeled in figure.
top	Integer, specifying number of top genes to be labeled.
bottom	Integer, specifying number of bottom genes to be labeled.
cutoff	Numeric.
main	As in 'plot'.
filename	Figure file name to create on disk. Default filename="NULL", which means no output.
width	As in ggsave.
height	As in ggsave.
...	Other available parameters in function 'ggsave'.

Value

An object created by ggplot, which can be assigned and further customized.

Author(s)

Wubing Zhang

Examples

```
file1 = file.path(system.file("extdata", package = "MAGeCKFlute"),
"testdata/rra.gene_summary.txt")
gdata = ReadRRA(file1)
rankdata = gdata$Score
names(rankdata) = gdata$id
RankView(rankdata)
```

ReadBeta

Read gene beta scores

Description

Read gene beta scores from file or data frame

Usage

```
ReadBeta(gene_summary)
```

Arguments

gene_summary A data frame or a file path to gene summary file generated by MAGeCK-MLE.

Value

A data frame, whose first column is Gene and other columns are comparisons.

Author(s)

Wubing Zhang

Examples

```
file3 = file.path(system.file("extdata", package = "MAGeCKFlute"),
"testdata/mle.gene_summary.txt")
dd = ReadBeta(file3)
head(dd)
```

ReadGMT

ReadGMT

Description

Parse gmt file to a data.frame

write data frame to a gmt file

Usage

```
ReadGMT(gmtpath, limit = c(0, Inf))
```

```
writeGMT(gene2path, gmtfile)
```

Arguments

gmtpath	The path to gmt file.
limit	A integer vector of length two, specifying the limit of geneset size.
gene2path	A data frame. The columns should be Gene, Pathway ID, and Pathway Name.
gmtfile	Path to gmt file.

Value

An data.frame, in which the first column is gene, and the second column is pathway name.

Output gmt file to local folder.

Author(s)

Wubing Zhang

Wubing Zhang

Examples

```
gene2path = gsGetter(type = "Complex")  
# writeGMT(gene2path, "Protein_complex.gmt")
```

ReadRRA	<i>Read gene summary file in MAGeCK-RRA results</i>
---------	---

Description

Read gene summary file in MAGeCK-RRA results

Usage

```
ReadRRA(gene_summary, score = c("lfc", "rra")[1])
```

Arguments

gene_summary	A data frame or a file path to gene summary file generated by MAGeCK-RRA.
score	"lfc" (default) or "rra", specifying the score type.

Details

If the score type is equal to lfc, then LFC will be returned. If the score type is rra, the log10 transformed RRA score will be returned.

Value

A data frame including three columns, including "id", "LFC" and "FDR".

Author(s)

Wubing Zhang

Examples

```
file1 = file.path(system.file("extdata", package = "MAGeCKFlute"),  
"testdata/rra.gene_summary.txt")  
gdata = ReadRRA(file1)  
head(gdata)
```

ReadsgRRA	<i>Read sgRNA summary in MAGeCK-RRA results</i>
-----------	---

Description

Read sgRNA summary in MAGeCK-RRA results

Usage

```
ReadsgRRA(sgRNA_summary)
```

Arguments

sgRNA_summary A file path or a data frame of sgRNA summary data.

Value

A data frame.

Author(s)

Wubing Zhang

Examples

```
file2 = file.path(system.file("extdata", package = "MAGeCKFlute"),
                  "testdata/rra.sgrna_summary.txt")
sgrra = ReadsgRRA(file2)
head(sgrra)
```

ResembleDepmap	<i>Compute the similarity between customized CRISPR screen with Depmap screens</i>
----------------	--

Description

Compute the similarity between customized CRISPR screen with Depmap screens

Usage

```
ResembleDepmap(
  dd,
  symbol = "id",
  score = "Score",
  lineages = "All",
  method = c("pearson", "spearman", "kendall")[1]
)
```

Arguments

dd	A data frame.
symbol	A character, specifying the column name of gene symbols in the data frame.
score	A character, specifying the column name of gene essentiality score in the data frame.
lineages	A character vector, specifying the lineages used for common essential gene selection.
method	A character, indicating which correlation coefficient is to be used for the test. One of "pearson", "kendall", or "spearman".

Value

A data frame with correlation and test p.value.

Author(s)

Wubing Zhang

Examples

```
file1 = file.path(system.file("extdata", package = "MAGeCKFlute"),
"testdata/rra.gene_summary.txt")
gdata = ReadRRA(file1)
## Not run:
rra.omit = OmitCommonEssential(gdata)
depmap_similarity = ResembleDepmap(rra.omit)
head(depmap_similarity)

## End(Not run)
```

retrieve_gs

Update genesets from source database

Description

Update genesets from source database

Usage

```
retrieve_gs(type = c("KEGG", "REACTOME", "CORUM", "GO"), organism = "hsa")
```

Arguments

type	A vector of databases, such as KEGG, REACTOME, CORUM, GO.
organism	'hsa' or 'mmu'.

Value

save data to local library.

Author(s)

Wubing Zhang

ScatterView

Scatter plot

Description

Scatter plot supporting groups.

Usage

```
ScatterView(  
  data,  
  x = "x",  
  y = "y",  
  label = 0,  
  model = c("none", "ninesquare", "volcano", "rank")[1],  
  x_cut = NULL,  
  y_cut = NULL,  
  slope = 1,  
  intercept = NULL,  
  auto_cut = FALSE,  
  auto_cut_x = auto_cut,  
  auto_cut_y = auto_cut,  
  auto_cut_diag = auto_cut,  
  groups = NULL,  
  group_col = NULL,  
  groupnames = NULL,  
  label.top = TRUE,  
  top = 0,  
  toplabels = NULL,  
  display_cut = FALSE,  
  color = NULL,  
  shape = 16,  
  size = 1,  
  alpha = 0.6,  
  main = NULL,  
  xlab = x,  
  ylab = y,  
  legend.position = "none",  
  ...  
)
```


Arguments

<code>data</code>	Data frame.
<code>x</code>	A character, specifying the x-axis.
<code>y</code>	A character, specifying the y-axis.
<code>label</code>	An integer or a character specifying the column used as the label, default value is 0 (row names).
<code>model</code>	One of "none" (default), "ninesquare", "volcano", and "rank".
<code>x_cut</code>	An one or two-length numeric vector, specifying the cutoff used for x-axis.
<code>y_cut</code>	An one or two-length numeric vector, specifying the cutoff used for y-axis.
<code>slope</code>	A numeric value indicating slope of the diagonal cutoff.
<code>intercept</code>	A numeric value indicating intercept of the diagonal cutoff.
<code>auto_cut</code>	Boolean, take 1.5 fold standard deviation as cutoff.
<code>auto_cut_x</code>	Boolean, take 1.5 fold standard deviation as cutoff on x-axis.
<code>auto_cut_y</code>	Boolean, take 1.5 fold standard deviation as cutoff on y-axis.
<code>auto_cut_diag</code>	Boolean, take 1.5 fold standard deviation as cutoff on diagonal.
<code>groups</code>	A character vector specifying groups. Optional groups include "top", "mid", "bottom", "left", "center", "right", "topleft", "topcenter", "topright", "midleft", "midcenter", "midright", "bottomleft", "bottomcenter", "bottomright".
<code>group_col</code>	A vector of colors for specified groups.
<code>groupnames</code>	A vector of group names to show on the legend.
<code>label.top</code>	Boolean, specifying whether label top hits.
<code>top</code>	Integer, specifying the number of top terms in the groups to be labeled.
<code>toplabels</code>	Character vector, specifying terms to be labeled.
<code>display_cut</code>	Boolean, indicating whether display the dashed line of cutoffs.
<code>color</code>	A character, specifying the column name of color in the data frame.
<code>shape</code>	A character, specifying the column name of shape in the data frame.
<code>size</code>	A character, specifying the column name of size in the data frame.
<code>alpha</code>	A numeric, specifying the transparency of the dots.
<code>main</code>	Title of the figure.
<code>xlab</code>	Title of x-axis
<code>ylab</code>	Title of y-axis.
<code>legend.position</code>	Position of legend, "none", "right", "top", "bottom", or a two-length vector indicating the position.
<code>...</code>	Other available parameters in function 'geom_text_repel'.

Value

An object created by `ggplot`, which can be assigned and further customized.

Author(s)

Wubing Zhang

Examples

```
file3 = file.path(system.file("extdata", package = "MAGeCKFlute"),
"testdata/mle.gene_summary.txt")
dd = ReadBeta(file3)
ScatterView(dd, x = "dms0", y = "plx", label = "Gene",
x_cut = 1, y_cut = 1, groups = "topright", top = 5, display_cut = TRUE)
```

Selector	<i>Select signatures from candidate list (according to the consistence in most samples).</i>
----------	--

Description

Select signatures from candidate list (according to the consistence in most samples).

Usage

```
Selector(mat, cutoff = 0, type = "<", select = 0.8)
```

Arguments

mat	Data matrix, each row is candidates (genes), each column is samples.
cutoff	Cutoff to define the signatures.
type	Direction to select signatures.
select	Proportion of samples in which signature is selected.

Value

An list containing two elements, first is selected signature and second is a ggplot object.

Examples

```
mat = matrix(rnorm(1000*30), 1000, 30)
rownames(mat) = paste0("Gene", 1:1000)
colnames(mat) = paste0("Sample", 1:30)
hits = Selector(mat, select = 0.68)
print(hits$p)
```

sgRankView	<i>View sgRNA rank.</i>
------------	-------------------------

Description

View sgRNA rank.

Usage

```
sgRankView(  
  df,  
  gene = NULL,  
  top = 3,  
  bottom = 3,  
  neg_ctrl = NULL,  
  binwidth = 0.3,  
  interval = 0.1,  
  bg.col = "gray90",  
  filename = NULL,  
  width = 5,  
  height = 3.5,  
  ...  
)
```

Arguments

df	A data frame, which contains columns of 'sgrna', 'Gene', and 'LFC'.
gene	Character vector, specifying genes to be plotted.
top	Integer, specifying number of top genes to be plotted.
bottom	Integer, specifying number of bottom genes to be plotted.
neg_ctrl	A vector specifying negative ctrl genes.
binwidth	A numeric value specifying the bar width.
interval	A numeric value specifying the interval length between each bar.
bg.col	A character value specifying the background color.
filename	Figure file name to create on disk. Default filename="NULL", which means no output.
width	As in ggsave.
height	As in ggsave.
...	Other available parameters in function 'ggsave'.

Value

An object created by ggplot.

Author(s)

Yihan Xiao

Examples

```
file2 = file.path(system.file("extdata", package = "MAGeCKFlute"),
                  "testdata/rra.sgrna_summary.txt")
sgrra = ReadsgRRA(file2)
sgRankView(sgrra)
```

 SquareView

Scatter plot of 9-Square

Description

Plot a scatter plot with Control beta score as x-axis and Treatment beta score as y-axis, and colored treatment related genes.

Usage

```
SquareView(
  beta,
  ctrlname = "Control",
  treatname = "Treatment",
  label = 0,
  label.top = TRUE,
  top = 5,
  genelist = c(),
  x_cutoff = NULL,
  y_cutoff = NULL,
  intercept = NULL,
  groups = c("midleft", "topcenter", "midright", "bottomcenter"),
  groupnames = paste0("Group", 1:length(groups)),
  main = NULL,
  filename = NULL,
  width = 6,
  height = 4,
  ...
)
```

Arguments

beta	Data frame, including columns of <i>ctrlname</i> and <i>treatname</i> , with Gene Symbol as rowname.
ctrlname	A character, specifying the names of control samples.
treatname	A character, specifying the name of treatment samples.

label	An integer or a character specifying the column used as the label, default value is 0 (row names).
label.top	Boolean, whether label the top selected genes, default label the top 10 genes in each group.
top	Integer, specifying the number of top selected genes to be labeled. Default is 5.
genelist	Character vector, specifying labeled genes.
x_cutoff	An one or two-length numeric vector, specifying the cutoff used for x-axis.
y_cutoff	An one or two-length numeric vector, specifying the cutoff used for y-axis.
intercept	An one or two-length numeric vector, specifying the intercept of diagonal.
groups	A character vector, specifying which group to be colored. Optional groups include "topleft", "topcenter", "topright", "midleft", "midright", "bottomleft", "bottomcenter", "bottomright".
groupnames	A character vector, specifying group names.
main	As in 'plot'.
filename	Figure file name to create on disk. Default filename="NULL", which means don't save the figure on disk.
width	As in ggsave.
height	As in ggsave.
...	Other available parameters in function 'ggsave'.

Value

An object created by ggplot, which can be assigned and further customized.

Author(s)

Wubing Zhang

See Also

[ScatterView](#)

Examples

```
file3 = file.path(system.file("extdata", package = "MAGeCKFlute"),
"testdata/mle.gene_summary.txt")
dd = ReadBeta(file3)
SquareView(dd, ctrlname = "dms0", treatname = "plx", label = "Gene")
```

TransGeneID

*Gene ID conversion between ENTREZID and SYMBOL***Description**

Gene ID conversion between ENTREZID and SYMBOL

Usage

```

TransGeneID(
  genes,
  fromType = "Symbol",
  toType = "Entrez",
  organism = "hsa",
  fromOrg = organism,
  toOrg = organism,
  ensemblHost = "www.ensembl.org",
  unique = TRUE,
  update = FALSE
)

```

Arguments

genes	A character vector, input genes to be converted.
fromType	The input ID type, one of "entrez", "symbol"(default), "hgnc", "ensembl", "full-name" and "uniprotswissprot"; you can also input other valid attribute names for biomaRt. Look at the code in examples to check valid attributes.
toType	The output ID type, similar to 'fromType'.
organism	"hsa"(default), "mmu", "bta", "cfa", "ptr", "rno", and "ssc" are optional.
fromOrg	"hsa", "mmu", "bta", "cfa", "ptr", "rno", and "ssc" are optional (Only used when transform gene ids between organisms).
toOrg	"hsa"(default), "mmu", "bta", "cfa", "ptr", "rno", and "ssc" are optional (Only used when transform gene ids between organisms).
ensemblHost	String, specifying ensembl host, you can use 'listEnsemblArchives()' to show all available Ensembl archives hosts.
unique	Boolean, specifying whether do one-to-one mapping.
update	Boolean, specifying whether update built-in gene annotation (needs network and takes time).

Value

A character vector, named by unique input gene ids.

Author(s)

Wubing Zhang

Examples

```
TransGeneID("HLA-A", organism="hsa")
TransGeneID("HLA-A", toType = "uniprot", organism="hsa")
TransGeneID("H2-K1", toType="Symbol", fromOrg = "mmu", toOrg = "hsa")
```

ViolinView

Violin plot

Description

Plots the violin of beta scores in Control and Treatment samples.

Usage

```
ViolinView(
  beta,
  samples = NULL,
  main = NULL,
  ylab = "Beta Score",
  filename = NULL,
  width = 5,
  height = 4,
  ...
)
```

Arguments

beta	Data frame, , including samples as columns.
samples	Character, specifying the name of samples to be compared.
main	As in 'plot'.
ylab	As in 'plot'.
filename	Figure file name to create on disk. Default filename="NULL", which means don't save the figure on disk.
width	As in ggsave.
height	As in ggsave.
...	Other available parameters in function 'ggsave'.

Value

An object created by ggplot, which can be assigned and further customized.

Author(s)

Wubing Zhang

See Also[DensityView](#)**Examples**

```
file3 = file.path(system.file("extdata", package = "MAGeCKFlute"),
"testdata/mle.gene_summary.txt")
dd = ReadBeta(file3)
ViolinView(dd, samples=c("dms0", "plx"))
#or
ViolinView(dd[, c("dms0", "plx")])
```

VolcanoView

Volcano View

Description

Volcano plot

Usage

```
VolcanoView(
  df,
  x = "logFC",
  y = "adj.P.Val",
  Label = NA,
  top = 5,
  topnames = NULL,
  x_cutoff = log2(1.5),
  y_cutoff = 0.05,
  mycolour = c("gray80", "#e41a1c", "#377eb8"),
  alpha = 0.6,
  force = 0.1,
  main = NULL,
  xlab = "Log2 Fold Change",
  ylab = "-Log10(Adjust.P)",
  filename = NULL,
  width = 4,
  height = 2.5,
  ...
)
```


Arguments

df	Data frame
x	Colname of df specifying x-axis in Volcano figure, 'logFC' (default).
y	Colname of df specifying y-axis in Volcano figure, 'adj.P.Val' (default), which will be plot after log10 transformation.
Label	Colname of df specifying labeled terms in Volcano figure.
top	Interger, the number of top significant terms to be labeled.
topnames	Character vector, indicating interested terms to be labeled.
x_cutoff	Cutoff of x-axis.
y_cutoff	Cutoff of y-axis.
mycolour	A color vector, specifying colors of non-significant, significant up and down-regulated genes.
alpha	Parameter in ggplot.
force	Parameter for geom_text_repel.
main	Title of volcano figure.
xlab	Label of x-axis in figure.
ylab	Label of y-axis in figure.
filename	Figure file name to create on disk. Default filename="NULL", which means don't save the figure on disk.
width	Width of figure.
height	Height of figure.
...	Other available parameters in ggsave.

Value

An object created by ggplot, which can be assigned and further customized.

Author(s)

Wubing Zhang

Examples

```
file1 = file.path(system.file("extdata", package = "MAGeCKFlute"),
"testdata/rra.gene_summary.txt")
gdata = ReadRRA(file1)
VolcanoView(gdata, x = "Score", y = "FDR", Label = "id")
```

Index

arrangePathview, 3

BarView, 5

BatchRemove, 6

ComBat, 6

ConsistencyView, 7

CutoffCalling, 8

DensityDiffView, 9

DensityView, 10, 56

enrich.GSE, 11, 14, 15, 18

enrich.HGT, 12, 13, 15, 18

enrich.ORT, 12, 14, 14, 18

EnrichAB, 16

EnrichAnalyzer, 12, 14, 15, 17

EnrichedFilter, 18

EnrichedGeneView, 19

EnrichedView, 20, 21

enricher, 15

enrichGO, 15

enrichGSE (enrich.GSE), 11

enrichKEGG, 15

enrichment (EnrichAnalyzer), 17

enrichORT (enrich.ORT), 14

EnrichSquare, 22

FluteMLE, 23, 27

flutemle (FluteMLE), 23

FluteRRA, 25, 25

getCols, 27

getGeneAnn, 28

getOrg, 29

getOrtAnn, 29

GSEA, 12

gseGO, 12

gseKEGG, 12

gsGetter, 30

hclustView, 31

HeatmapView, 32

IdentBarView, 33

IncorporateDepmap, 34

loess, 39

loess.normalize (normalize.loess), 38

MapRatesView, 35

MAView, 36

noEnrichPlot, 38

normalize.loess, 38

NormalizeBeta, 39, 39

normalizebeta (NormalizeBeta), 39

OmitCommonEssential, 41

RankView, 42

rankview (RankView), 42

ReadBeta, 43

readbeta (ReadBeta), 43

ReadGMT, 44

ReadRRA, 45

readrra (ReadRRA), 45

ReadsgRRA, 46

ResembleDepmap, 46

retrieve_gs, 47

RRApipeline (FluteRRA), 25

ScatterView, 48, 53

Selector, 50

sgRankView, 51

SquareView, 52

squareview (SquareView), 52

TransGeneID, 54

transGeneID (TransGeneID), 54

ViolinView, 11, 55

violinview (ViolinView), [55](#)

VolcanoView, [56](#)

writeGMT (ReadGMT), [44](#)