

# Package ‘SGCP’

November 21, 2024

**Type** Package

**Title** SGCP: A semi-supervised pipeline for gene clustering using self-training approach in gene co-expression networks

**Version** 1.6.0

**Description** SGC is a semi-supervised pipeline for gene clustering in gene co-expression networks. SGC consists of multiple novel steps that enable the computation of highly enriched modules in an unsupervised manner. But unlike all existing frameworks, it further incorporates a novel step that leverages Gene Ontology information in a semi-supervised clustering method that further improves the quality of the computed modules.

**License** GPL-3

**Encoding** UTF-8

**Imports** ggplot2, expm, caret, plyr, dplyr, GO.db, annotate, SummarizedExperiment, genefilter, GOstats, RColorBrewer, xtable, Rgraphviz, reshape2, openxlsx, ggridges, DescTools, org.Hs.eg.db, methods, grDevices, stats, RSpectra, graph

**Suggests** knitr, rmarkdown, BiocManager, devtools, BiocStyle

**Depends** R (>= 4.2.0)

**biocViews** GeneExpression, GeneSetEnrichment, NetworkEnrichment, SystemsBiology, Classification, Clustering, DimensionReduction, GraphAndNetwork, NeuralNetwork, Network, mRNAArray, RNASeq, Visualization

**VignetteBuilder** knitr

**NeedsCompilation** no

**URL** <https://github.com/na396/SGCP>

**RoxygenNote** 7.2.1

**LazyData** true

**git\_url** <https://git.bioconductor.org/packages/SGCP>

**git\_branch** RELEASE\_3\_20

**git\_last\_commit** e164401

**git\_last\_commit\_date** 2024-10-29

**Repository** Bioconductor 3.20

**Date/Publication** 2024-11-20

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adjacencyMatrix	<i>Performs network construction step in the SGCP pipeline</i>
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## Description

It creates the adjacency matrix of the gene co-expression network in the SGCP pipeline. Users can specify steps in the following order: calibration, norm, Gaussian kernel, and tom. If calibration is set to TRUE, SGCP performs calibration as the first step (refer to the manuscript for details). If norm is TRUE, each gene is normalized by its L2 norm. The Gaussian kernel metric is then calculated as a mandatory step to determine pairwise gene similarity values. If tom is TRUE, SGCP incorporates second-order node neighborhood information into the network. The pipeline concludes by returning a symmetric adjacency matrix `adja` of size  $m \times n$ , where  $n$  is the number of genes. All values in the adjacency matrix range from 0 to 1, with 1 indicating maximum similarity. The diagonal elements of the matrix are set to zero.

## Usage

```
adjacencyMatrix(expData, calibration = FALSE, norm = TRUE,  
                tom = TRUE, saveAdja = FALSE,  
                adjaNameFile = "adjacency.RData",  
                hm = "adjaHeatMap.png")
```

## Arguments

expData	A dataframe or matrix containing the expression data, where rows correspond to genes and columns to samples.
calibration	Logical, default FALSE. If TRUE, performs calibration step.
norm	Logical, default TRUE. If TRUE, divides each gene (row) by its norm2.
tom	Logical, default TRUE. If TRUE, adds TOM to the network.
saveAdja	Logical, default FALSE. If TRUE, saves the adjacency matrix.
adjaNameFile	String indicating the name of the file for saving the adjacency matrix.
hm	String indicating the name of the file for saving the adjacency matrix heatmap.

## Value

adja	A symmetric matrix of dimension $n * n$ representing the adjacency matrix, where $n$ is the number of genes. Values range in $(0, 1)$ with a zero diagonal.
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## References

[Aghaieabiane, N and Koutis, I \(2022\) SGCP: A semi-supervised pipeline for gene clustering using self-training approach in gene co-expression networks](#)

## See Also

[SGCP Tutorial calibration step information](#)

## Examples

```
## create an adjacency matrix  
GeneExpression <- matrix(runif(1000, 0,1), nrow = 200, ncol = 5)  
diag(GeneExpression) <- 0  
  
## call the function  
adja <- adjacencyMatrix(GeneExpression, hm= NULL)  
head(adja)
```

---

cheng

*Normalized gene expression data from Cheng et al.'s publication on ischemic cardiomyopathy (ICM).*

---

## Description

This dataset contains normalized gene expression data for 1500 genes across 5 samples. It is a subset of a larger dataset related to ischemic cardiomyopathy (ICM), which includes 5000 genes and 57 samples. The normalization was performed using the DESeq method, which utilizes the median ratio of gene counts to achieve normalization.

**Usage**

```
data(cheng)
```

**Format**

An object of class SummarizedExperiment.

**Details**

assays contains the gene expression data and rowData field includes the corresponding gene Entrez IDs. Sample names also are available in colData.

**Source**

<https://www.sciencedirect.com/science/article/pii/S0010482520303061?via%3Dihub>

**Examples**

```
## load cheng dataset
library(SGCP)
library(SummarizedExperiment)

data(cheng)
expData <- assay(cheng)
geneID <- rowData(cheng)
geneID <- geneID$ENTREZID
```

---

clustering

*Perform network clustering step in the SGCP pipeline*

---

**Description**

It performs clustering on the adjacency network of gene co-expression network in SGCP pipeline. Initially, it transforms the  $n \times n$  adjacency matrix into a new dimension  $Y$ . Subsequently, it determines the number of clusters  $k$  using three methods: "relativeGap", "secondOrderGap", and "additiveGap". For each method,  $k$ -means clustering is applied to  $Y$  with the determined  $k$  as input. Conductance indices are computed for the clusters within each method, and the cluster with the smallest conductance index is selected for further analysis. Following this, gene ontology enrichment analysis is performed on the selected clusters to finalize the optimal  $k$ . The pipeline concludes by returning the result of  $k$ -means clustering based on the selected method, along with the transformed matrix  $Y$  and additional information. This step produces the initial clusters.

**Usage**

```
clustering(adjaMat, geneID, annotation_db,
           kopt = NULL, method = NULL,
           func.GO = sum, func.conduct = min,
           maxIter = 1e8, numStart = 1000, eff.egs = TRUE,
           saveOrig = TRUE, n_egvec = 200, sil = FALSE)
```

**Arguments**

<code>adjaMat</code>	A squared symmetric matrix of size $n \times n$ with values in (0, 1) and 0 diagonal. This is the output of the <code>adjacencyMatrix</code> function in SGCP.
<code>geneID</code>	A vector containing gene IDs of size $n$ , where $n$ is the number of genes.
<code>annotation_db</code>	A string indicating the genomic-wide annotation database.
<code>kopt</code>	An integer denoting the optimal number of clusters $k$ chosen by the user (default: NULL).
<code>method</code>	Method for identifying the number of clusters $k$ (default: NULL). Options include "relativeGap", "secondOrderGap", "additiveGap", or NULL.
<code>func.GO</code>	A function for gene ontology validation (default: sum).
<code>func.conduct</code>	A function for conductance validation (default: min).
<code>maxIter</code>	An integer specifying the maximum number of iterations for k-means clustering.
<code>numStart</code>	An integer indicating the number of starts for k-means clustering.
<code>eff.egs</code>	Boolean (default: TRUE). If TRUE, uses <code>eigs_sym</code> to calculate eigenvalues and eigenvectors, which is more efficient than R's default function.
<code>saveOrig</code>	Boolean (default: TRUE). If TRUE, keeps the transformation matrix.
<code>n_egvec</code>	An integer (default: 200) specifying the number of columns of the transformation matrix to retain. Should be less than 200.
<code>sil</code>	Boolean (default: FALSE). If TRUE, calculates silhouette index for each cluster.

**Details**

If `kopt` is not null, SGCP will determine clusters based on the specified `kopt`. Otherwise, if `method` is not NULL, SGCP will select  $k$  using the specified method. If both `geneID` and `annotation_db` are NULL, SGCP will determine the optimal method and its corresponding number of clusters based on conductance validation. It selects a method where the conductance, evaluated by `func.conduct`, is minimized. Alternatively, SGCP defaults to gene ontology validation to find the optimal method and its corresponding clusters. It performs gene ontology enrichment on clusters, selecting the method where the cluster with the minimum conductance index yields the highest `func.GO` over  $\log_{10}$  of the p-values.

**Value**

- `dropped.indices` A vector of dropped gene indices.
- `geneID` A vector of gene IDs.
- `method` Indicates the selected method for number of clusters.
- `k` Selected number of clusters.
- `Y` Transformed matrix with  $2 \times k$  columns.
- `X` Eigenvalues corresponding to  $2 \times k$  columns in `Y`.
- `cluster` An object of class `kmeans`.
- `clusterLabels` A vector containing the cluster label per gene, with a 1-to-1 correspondence to `geneID`.
- `conductance` A list containing mean and median conductance indices for clusters per method. The index in `clusterConductance` field denotes the cluster label and the value shows the conductance index.

- `cvGOdf` A dataframe used for gene ontology validation. For each method, it returns the gene ontology enrichment result on the cluster with the minimum conductance index.
- `cv` A string indicating the validation method for number of clusters:
  - `"cvGO"`: Gene ontology validation used.
  - `"cvConductance"`: Conductance validation used.
  - `"userMethod"`: User-defined method.
  - `"userkopt"`: User-defined `kopt`.
- `clusterNumberPlot` An object of class `ggplot2` for `relativeGap`, `secondOrderGap`, and `additiveGap`.
- `silhouette` A dataframe indicating the silhouette index for genes.
- `original` A list with matrix transformation, corresponding eigenvalues, and `n_egvec`, where the top `n_egvec` columns of the transformation are retained.

## References

[Aghaieabiane, N and Koutis, I \(2022\) SGCP: A semi-supervised pipeline for gene clustering using self-training approach in gene co-expression networks](#)

## See Also

[adjacencyMatrix SGCP Tutorial](#)

## Examples

```
## load cheng dataset
library(SGCP)
library(SummarizedExperiment)

data(cheng)
expData <- assay(cheng)
geneID <- rowData(cheng)
geneID <- geneID$ENTREZID

# to create the adjacency matrix un comment the following
## resAdja <- adjacencyMatrix(expData = expData, hm = NULL)
## resAdja[0:10, 0:5]

# to perform clustering
## library(org.Hs.eg.db)
annotation_db = "org.Hs.eg.db"
## resClus = clustering(adjaMat = resAdja, geneID = geneID,
##                       annotation_db = annotation_db)
```

**Description**

The SGCP pipeline for gene co-expression network construction and analysis integrates multiple steps into a single function. It begins with network construction, where gene expression data and gene IDs are utilized alongside an annotation database to build an adjacency matrix. Next, network clustering identifies initial clusters. Gene ontology enrichment distinguishes genes into remarkable and unremarkable sets, enabling semi-labeling to convert the problem into semi-supervised learning. Remarkable genes serve as the training set for a supervised model, predicting labels for unremarkable genes and producing final modules. Finally, another gene ontology step evaluates module enrichment.

**Usage**

```
ezSGCP(expData, geneID, annotation_db, semilabel = TRUE,
        calib = FALSE, norm = TRUE, tom = TRUE,
        saveAdja = FALSE, adjaNameFile = "adjacency.Rdata",
        hm = "adjaHeatMap.png",
        kopt = NULL, method_k = NULL, f.GO = sum, f.conduct = min,
        maxIteration = 1e8, numberStart = 1000, eff.egs = TRUE,
        saveOrig = TRUE, n_egvec = 100, sil = FALSE,
        dir = c("over", "under"), onto = c("BP", "CC", "MF"),
        hgCut = NULL, condTest = TRUE,
        cutoff = NULL, percent = 0.10, stp = 0.01,
        model = "knn", kn = NULL)
```

**Arguments**

expData	A dataframe or matrix containing the expression data, where rows correspond to genes and columns to samples.
geneID	A vector containing the gene IDs of size n, where n is the number of genes.
annotation_db	A string indicating the genomic-wide annotation database.
semilabel	Logical, default TRUE. If TRUE, performs semilabeling step.
calib	Logical, default FALSE. If TRUE, performs calibration step.
norm	Logical, default TRUE. If TRUE, divides each gene (row) by its norm2.
tom	Logical, default TRUE. If TRUE, adds TOM to the network.
saveAdja	Logical, default FALSE. If TRUE, saves the adjacency matrix.
adjaNameFile	String indicating the name of the file for saving the adjacency matrix.
hm	String indicating the name of the file for saving the adjacency matrix heatmap.
kopt	An integer indicating the optimal number of clusters k chosen by the user, default is NULL.
method_k	Method for identifying the number of clusters k, default NULL. Options are "relativeGap", "secondOrderGap", "additiveGap", or NULL.
f.GO	A function for gene ontology validation, default is sum.
f.conduct	A function for conductance validation, default is min.

<code>maxIteration</code>	An integer indicating the maximum number of iterations for kmeans.
<code>numberStart</code>	An integer indicating the number of starts for kmeans.
<code>eff.egs</code>	Boolean, default TRUE. If TRUE, uses <code>eigs_sym</code> to calculate eigenvalues and eigenvectors, which is more efficient than R's default function.
<code>saveOrig</code>	Boolean, default TRUE. If TRUE, keeps the transformation matrix.
<code>n_egvec</code>	Either "all" or an integer indicating the number of columns of the transformation matrix to keep, default is 100.
<code>sil</code>	Logical, default FALSE. If TRUE, calculates silhouette index for each cluster.
<code>dir</code>	Test direction for GO terms, default <code>c("over", "under")</code> .
<code>onto</code>	GO ontologies to consider, default <code>c("BP", "CC", "MF")</code> .
<code>hgCut</code>	Numeric value in (0,1) as the p-value cutoff for GO terms, default 0.05.
<code>condTest</code>	Logical, default TRUE. If TRUE, performs conditional hypergeometric test.
<code>cutoff</code>	Numeric in (0, 1) default NULL, baseline for GO term significance.
<code>percent</code>	Numeric in (0,1) default 0.1, percentile for finding top GO terms.
<code>stp</code>	Numeric in (0,1) default 0.01, increasing value for percent parameter.
<code>model</code>	Type of classification model, either "knn" (k nearest neighbors) or "lr" (logistic regression).
<code>kn</code>	Integer indicating the number of neighbors in knn, default NULL.

## Details

For clustering step; If `kopt` is not NULL, SGCP finds clusters based on `kopt`. If `method_k` is not NULL, SGCP picks `k` based on the selected method (`"relativeGap"`, `"secondOrderGap"`, `"additiveGap"`). If `geneID` or `annotation_db` is NULL, SGCP determines the optimal method and corresponding number of clusters based on conductance validation. It selects the method where `func.conduct` on its clusters is minimized. Otherwise, SGCP uses gene ontology validation (by default) to find the optimal method and its corresponding number of clusters. It performs gene ontology enrichment on the cluster with the minimum conductance index per method and selects the method that maximizes `func.GO` over  $-\log_{10}$  of p-values.

For semilabeling step; Genes associated with GO terms more significant than `cutoff` value are considered remarkable. If `cutoff` value is NULL, SGCP determines the cutoff based on the significance level of GO terms. SGCP selects the top percent (default 0.1) GO terms from all clusters collectively and considers genes associated with those as remarkable. If all remarkable genes come from a single cluster, SGCP increases the percent by 0.01 until remarkable genes come from at least two clusters.

For semi-supervise step; Remarkable clusters are those that have at least one remarkable gene. SGCP performs semi-supervised classification using the transformed matrix from clustering and gene semilabels from semilabeling function. It uses remarkable genes as the training set to train either a "k nearest neighbor" (knn) or "logistic regression" (lr) model and makes predictions for unremarkable genes to produce the final modules.

## Value

Returns a list with the following fields, depending on the initial call:

- `semilabel` Boolean indicating if semilabeling step was performed.
- `clusterLabels` DataFrame with `geneID` and its corresponding initial and final labels.
- `clustering` List containing clustering information:



- `dropped.indices` Vector of dropped gene indices.
- `geneID` Vector of geneIDs.
- `method` Method selected for number of clusters.
- `k` Selected number of clusters.
- `Y` Transformed matrix with  $2*k$  columns.
- `X` Eigenvalues corresponding to  $2*k$  columns in `Y`.
- `cluster` Object of class `kmeans`.
- `clusterLabels` Vector containing cluster labels for each gene.
- `conductance` List with mean, median, and individual cluster conductance indices. `clusterConductance` field denotes the cluster label and its conductance index.
- `cvGOdf` DataFrame used for gene ontology validation. For each method, shows GO enrichment on the cluster with smallest conductance index.
- `cv` String indicating validation method for number of clusters: "cvGO", "cvConductance", "userMethod", or "userkopt".
- `clusterNumberPlot` Object of class `ggplot2` for visualizing cluster number selection methods.
- `silhouette` DataFrame indicating silhouette indices for genes.
- `original` List with matrix transformation, corresponding eigenvalues, and `n_egvec` top columns of transformation matrix kept.
- `initial.GO` List containing initial gene ontology (GO) information:
  - `GOresults` DataFrame summarizing GO term information. Includes `clusterNum`, `GOtype`, `GOID`, `Pvalue`, `OddsRatio`, `ExpCount`, `Count`, `Size`, and `Term`.
  - `FinalGOTermGenes` List of geneIDs associated with each GO term per cluster.
- `semiLabeling` List containing semilabeling information:
  - `cutoff` Numeric (0,1) indicating selected cutoff for significant GO terms.
  - `geneLabel` DataFrame with `geneID` and its corresponding cluster label if remarkable, otherwise NA.
- `semiSupervised` List containing semi-supervised classification information:
  - `semiSupervised` Object of classification result.
  - `prediction` Vector of predicted labels for unremarkable genes.
  - `FinalLabeling` DataFrame of `geneID` with its corresponding semilabel and final label.
- `final.GO` List containing final gene ontology (GO) information:
  - `GOresults` DataFrame summarizing GO term information. Includes `clusterNum`, `GOtype`, `GOID`, `Pvalue`, `OddsRatio`, `ExpCount`, `Count`, `Size`, and `Term`.
  - `FinalGOTermGenesList` of geneIDs associated with each GO term per cluster.

## References

[Aghaieabiane, N and Koutis, I \(2022\) SGCP: A semi-supervised pipeline for gene clustering using self-training approach in gene co-expression networks](#)

## See Also

[SGCP Tutorial](#)

**Examples**

```

## load cheng dataset
library(SGCP)
library(SummarizedExperiment)
data(cheng)
expData <- assay(cheng)
geneID <- rowData(cheng)
geneID <- geneID$ENTREZID

library(org.Hs.eg.db)

# to call the function uncomment the following
## res <- ezSGCP(expData = expData, geneID = geneID, annotation_db = "org.Hs.eg.db")
## summary(res)
## summary(res$clustering)
## summary(res$initial.GO)
## summary(res$semiLabeling)
## summary(res$semiSupervised)
## summary(res$final.GO)

```

---

geneOntology

*Performs gene ontology enrichment step in the SGCP pipeline.*


---

**Description**

It performs gene ontology enrichment step Gostat package in SGCP pipeline. It takes the entire genes in the input with their labels, along with annotation\_db to perform gene ontology enrichment for each set of genes that have similar label.

**Usage**

```

geneOntology(geneUniv, clusLab, annotation_db,
             direction = c("over", "under"),
             ontology = c("BP", "CC", "MF"), hgCutoff = NULL,
             cond = TRUE)

```

**Arguments**

geneUniv	a vector of all the geneIDs in the expression dataset.
clusLab	a vector of cluster label for each geneID.
annotation_db	a string indicating the genomic wide annotation database.
direction	test direction, default c("over", "under"), for over-represented, or under-represented GO terms.
ontology	GO ontologies, default c("BP", "CC", "MF"), BP: Biological Process, CC: Cellular Component, MF: Molecular Function.

hgCutoff	a numeric value in (0,1) as the p-value cutoff, default 0.05, GO terms smaller than hgCutoff value are kept.
cond	Boolean, default TRUE, if TRUE conditional hypergeometric test is performed.

**Value**

GOresults	a dataframe containing the summary of the information of GOTerms, cluster-Num: indicates the cluster label, GOtype: indicates the test directions plus ontology, GOID: unique GO term id, Pvalue: the p-value of hypergeometric test for the GO term, OddsRatio: the odds ratio of the GO term, ExpCount: expected count value for genes associated the GO term, Count: actual count of the genes associated to the GO term in the cluster, Size: actual size of the genes associated to the GO term in the entire geneIDs, Term: description of the GO term.
FinalGOTermGenes	a list containing the geneIDs of each GOTerms per cluster.

**References**

[Aghaieabiane, N and Koutis, I \(2022\) SGCP: A semi-supervised pipeline for gene clustering using self-training approach in gene co-expression networks](#)

**See Also**

[SGCP Tutorial](#) [Gostat Tutorial](#)

**Examples**

```
library(SGCP)
# load the output of clustering function
data(resClus)

# call the function
library(org.Hs.eg.db)

# to call the geneOntology uncomment the following
## res <- geneOntology(geneUniv = resClus$geneID, clusLab = resClus$clusterLabels,
##                    annotation_db = "org.Hs.eg.db")
## summary(res$GOresults)
## summary(res$FinalGOTermGenes)
```

---

resClus	<i>An example of the output from clustering function in the SGCP pipeline</i>
---------	---

---

**Description**

This is an example of the output from the clustering function, representing the network clustering step in the SGCP pipeline. Initially, the adjacency matrix is generated using the adjacencyMatrix function within the SGCP framework applied to the cheng dataset. This adjacency matrix serves as input to the clustering function, resulting in the clustering outcome stored in resClus

## Usage

```
data(resClus)
```

## Format

An object of class `list` containing the clustering information.

## Details

`resClus` is a list containing the following clustering information:

- `dropped.indices`: A vector of dropped gene indices.
- `geneID`: A vector of gene IDs.
- `method`: Indicates the selected method for determining the number of clusters.
- `k`: The selected number of clusters.
- `Y`: Transformed matrix with  $2*k$  columns.
- `X`: Eigenvalues corresponding to the  $2*k$  columns in `Y`.
- `cluster`: An object of class `kmeans`.
- `clusterLabels`: A vector containing the cluster label for each gene. There is a 1-to-1 correspondence between `geneID` and `clusterLabels`.
- `conductance`: A list containing the mean, median, and individual cluster conductance index for clusters per method. The index in the `clusterConductance` field denotes the method.
- `cvGOdf`: A dataframe used for gene ontology validation. For each method, it returns the gene ontology enrichment result on the cluster with the minimum conductance index.
- `cv`: A string indicating the validation method for the number of clusters; "cvGO" means gene ontology validation was used.
- `clusterNumberPlot`: An object of class `ggplot2` for `relativeGap`, `secondOrderGap`, and `additiveGap`.
- `silhouette`: A dataframe indicating the silhouette values for genes.
- `original`: A list with matrix transformation, corresponding eigenvalues, and `n_egvec`, where the top
- `n_egvec` columns of the transformation are kept.

## See Also

[SGCP Tutorial adjacencyMatrix clustering](#)

## Examples

```
library(SGCP)
data(resClus)
summary(resClus)
resClus
```

---

`resFinalGO`*An example of the output from geneOntology function in the SGCP pipeline*

---

### Description

This is an example of the output from the `geneOntology` function, representing the final step in the SGCP pipeline. Initially, the adjacency matrix is generated using the `adjacencyMatrix` function within the SGCP framework applied to the cheng dataset. This adjacency matrix serves as input to the clustering function, resulting in the clustering outcome stored in `resClus`. The clustering result, `resClus`, is subsequently utilized in the `geneOntology` function to derive `resInitialGO`, which captures the initial gene ontology (GO) enrichment results. The `resInitialGO` output is then processed through the `semiLabeling` function to produce `resSemiLabel`, indicating the semi-labeled genes based on their clustering characteristics. This semi-labeled information is further employed in the `semiSupervised` function, yielding `resSemiSupervised`, which includes the final supervised classification outcomes for the unremarkable genes. Finally, the results from `resSemiSupervised` are fed into the `geneOntology` function once more to generate `resFinalGO`, which represents the final GO enrichment analysis.

### Usage

```
data(resFinalGO)
```

### Format

An object of class `list` containing the gene ontology information for final gene ontology.

### Details

`coderesFinalGO` is a list containing the following information:

- `GOresults`: A dataframe of significant gene ontology terms and their corresponding test statistics.
- `FinalGOTermGenes`: A list of genes belonging to significant gene ontology terms per cluster..

### See Also

[SGCP Tutorial](#) [geneOntology](#)

### Examples

```
library(SGCP)
data(resFinalGO)
summary(resFinalGO)

# dataframe of significant gene ontology terms
head(resFinalGO$GOresults)

# a list of genes belong to significant gene ontology term for cluster 1
head(resFinalGO$FinalGOTermGenes$Cluster1_GOTermGenes)

# a list of genes belong to significant gene ontology term for cluster 2
head(resFinalGO$FinalGOTermGenes$Cluster2_GOTermGenes)
```

---

resInitialGO	<i>An example of the output from the geneOntology function in the SGCP pipeline</i>
--------------	---

---

## Description

This is an example of the output from the `geneOntology` function, representing the third step in the SGCP pipeline. Initially, the adjacency matrix is generated using the `adjacencyMatrix` function within the SGCP framework applied to the cheng dataset. This adjacency matrix serves as input to the clustering function, resulting in the clustering outcome stored in `resClus`. The clustering result, `resClus`, is subsequently utilized in the `geneOntology` function to derive `resInitialGO`, which captures the initial gene ontology (GO) enrichment results.

## Usage

```
data(resInitialGO)
```

## Format

An object of class `list` containing the gene ontology information for initial gene ontology.

## Details

`resInitialGO` is a list containing the following information.

- `GOresults`: a dataframe of significant gene ontology terms and their corresponding test statistics information.
- `FinalGOTermGenes`: a list of the genes belong to significant gene ontology terms per cluster.

## See Also

[SGCP Tutorial](#) [geneOntology](#)

## Examples

```
library(SGCP)
data(resInitialGO)
summary(resInitialGO)

# dataframe of significant gene ontology terms
head(resInitialGO$GOresults)

# a list of genes belong to significant gene ontology term for cluster 1
head(resInitialGO$FinalGOTermGenes$Cluster1_GOTermGenes)

# a list of genes belong to significant gene ontology term for cluster 2
head(resInitialGO$FinalGOTermGenes$Cluster2_GOTermGenes)
```

---

resSemiLabel	<i>An example of the output from semiLabeling function in the SGCP pipeline</i>
--------------	---

---

## Description

This is an example of the output from the `semiLabeling` function, representing the semi-label step in the SGCP pipeline. Initially, the adjacency matrix is generated using the `adjacencyMatrix` function within the SGCP framework applied to the cheng dataset. This adjacency matrix serves as input to the clustering function, resulting in the clustering outcome stored in `resClus`. The clustering result, `resClus`, is subsequently utilized in the `geneOntology` function to derive `resInitialGO`, which captures the initial gene ontology (GO) enrichment results. The `resInitialGO` output is then processed through the `semiLabeling` function to produce `resSemiLabel`, indicating the semi-labeled genes based on their clustering characteristics.

## Usage

```
data(resSemiLabel)
```

## Format

An object of class `list` containing the semi-labeling information.

## Details

`resSemiLabel` is a list containing the following information.

- `cutoff`: a numeric in (0,1) that shows the base line for identifying remarkable genes.
- `geneLabel`: a dataframe of geneIDs and its corresponding label, NA labels means that corresponding genes are unremarkable.

## See Also

[SGCP Tutorial semiLabeling](#)

## Examples

```
library(SGCP)
data(resSemiLabel)
summary(resSemiLabel)

# cutoff value
head(resSemiLabel$cutoff)

# gene semi-label
head(resSemiLabel$geneLabel)
```

---

resSemiSupervised	<i>An example of the output from semiSupervised function in the SGCP pipeline</i>
-------------------	---

---

### Description

This is an example of the output from the `semiSupervised` function, representing the semi-supervised step in the SGCP pipeline. Initially, the adjacency matrix is generated using the `adjacencyMatrix` function within the SGCP framework applied to the cheng dataset. This adjacency matrix serves as input to the clustering function, resulting in the clustering outcome stored in `resClus`. The clustering result, `resClus`, is subsequently utilized in the `geneOntology` function to derive `resInitialGO`, which captures the initial gene ontology (GO) enrichment results. The `resInitialGO` output is then processed through the `semiLabeling` function to produce `resSemiLabel`, indicating the semi-labeled genes based on their clustering characteristics. This semi-labeled information is further employed in the `semiSupervised` function, yielding `resSemiSupervised`, which includes the final supervised classification outcomes for the unremarkable genes.

### Usage

```
data(resSemiSupervised)
```

### Format

An object of class `list` containing the semi-supervised information.

### Details

`resSemiSupervised` is a list containin the following information.

- `semiSupervised`: an object of `caret` for the training model.
- `prediction`: A vector of predicted labels for unremakable genes.
- `FinalLabeling`: a dataframe gene semi-label and final predicted labels.

### See Also

[SGCP Tutorial](#) [semiLabeling](#)

### Examples

```
library(SGCP)
data(resSemiSupervised)

# supervised model information
summary(resSemiSupervised$semiSupervised)

# predicted label for unremarkable genes
head(resSemiSupervised$prediction)

# gene semi and final labeling
head(resSemiSupervised$FinalLabeling)
```



---

semiLabeling	<i>Performs gene semi-labeling step in the SGCP pipeline</i>
--------------	--

---

### Description

Performs the Semi-labeling step in the SGCP pipeline to identify remarkable and unremarkable genes. This step involves collecting all gene ontology (GO) terms from all clusters and selecting terms in the top 0.1 percent. Genes associated with these terms are considered remarkable, while the remaining genes are categorized as unremarkable.

### Usage

```
semiLabeling(geneID, df_GO, GOgenes, cutoff = NULL,
             percent = 0.10, stp = 0.01)
```

### Arguments

geneID	A vector containing gene IDs, where n is the number of genes.
df_GO	The GOresults dataframe returned by the geneOntology function, containing information on GO terms in the clusters.
GOgenes	The FinalGOTermGenes list returned by the geneOntology function, listing genes associated with GO terms for each cluster.
cutoff	A numeric value in (0, 1) (default: NULL), serving as a baseline for GO term significance.
percent	A numeric value in (0, 1) (default: 0.1), indicating the percentile for selecting top GO terms.
stp	A numeric value in (0, 1) (default: 0.01), increment added to the percent parameter for stepwise selection of top GO terms.

### Details

Genes associated with GO terms more significant than the cutoff value are considered remarkable. If the cutoff value is NULL, SGCP determines the cutoff based on the significance level of the GO terms. Otherwise, SGCP selects the top percent (default: 0.1) of GO terms from all clusters combined, considering genes associated with these terms as remarkable. If all remarkable genes originate from a single cluster, SGCP incrementally increases the percent parameter by 0.01 to identify both remarkable and unremarkable genes. This process continues until remarkable genes originate from at least two clusters.

### Value

cutoff	a numeric in (0,1) which indicates the selected cutoff.
geneLabel	a dataframe containing the information of geneID and its corresponding cluster label if is remarkable otherwise NA.

### References

[Aghaieabiane, N and Koutis, I \(2022\) SGCP: A semi-supervised pipeline for gene clustering using self-training approach in gene co-expression networks](#)

**See Also**

[geneOntology](#) [SGCP Tutorial](#)

**Examples**

```
library(SGCP)
# load the output of clustering, gene ontology function

data(resClus)
data(resInitialGO)

# call the function

res <- semiLabeling(geneID = resClus$geneID, df_GO = resInitialGO$GOresults,
                   GOgenes = resInitialGO$FinalGOTermGenes)
# cutoff value
res$cutoff

# gene semi-labeling information
head(res$geneLabel)
```

---

semiSupervised

*Performs the semi-supervised step in the SGCP pipeline*

---

**Description**

Performs the semi-supervised classification step in the SGCP pipeline. It utilizes the transformed matrix from the clustering function along with gene semi-labels from the semiLabeling function. The labeled (remarkable) genes serve as the training set to train either a "k-nearest neighbor" or "logistic regression" model. The trained model then predicts labels for unlabeled (unremarkable) genes, resulting in the final modules.

**Usage**

```
semiSupervised(specExp, geneLab, model = "knn", kn = NULL)
```

**Arguments**

specExp	Matrix or dataframe where genes are in rows and features are in columns, representing the Y matrix from clustering function output.
geneLab	A dataframe returned by the semiLabeling function, containing geneIDs and their corresponding labels (remarkable or NA).
model	Classification model type: "knn" for k-nearest neighbors or "lr" for logistic regression.
kn	An integer (default: NULL) indicating the number of neighbors in k-nearest neighbors (knn) model. If kn is NULL, the default value is determined by: $kn = 20$ if $2 * k < 30$ , otherwise $kn = 20 : 30$ , where k is the number of remarkable clusters.

**Details**

Remarkable clusters are defined as clusters that contain at least one remarkable gene.

**Value**

`itemsemiSupervised` An object of the `caret` train class representing the semi-supervised classification model.

`prediction` A vector containing predicted labels for unremarkable genes.

`FinalLabeling` A dataframe containing geneIDs along with their corresponding semi-labels and final labels.

**References**

[Aghaieabiane, N and Koutis, I \(2022\) SGCP: A semi-supervised pipeline for gene clustering using self-training approach in gene co-expression networks](#)

**See Also**

[clustering semiLabeling SGCP Tutorial](#)

**Examples**

```
library(SGCP)
# load the output of clustering, gene ontology function

data(resClus)
data(resSemiLabel)

# call the function

res <- semiSupervised(specExp = resClus$Y, geneLab = resSemiLabel$geneLabel)

# model summary
summary(res$semiSupervised)

# prediction label for unremarkable genes
head(res$prediction)

# semi and final gene labels
head(res$FinalLabeling)
```

---

 sgcp

*An example of the output of ezSGCP function in the SGCP pipeline*

---

**Description**

This is an example of the output from the `ezSGCP` function, representing the entire SGCP pipeline. Initially, the adjacency matrix is generated using the `adjacencyMatrix` function within the SGCP framework applied to the `cheng` dataset. This adjacency matrix serves as input to the `clustering` function, resulting in the clustering outcome stored in `resClus`. The clustering result, `resClus`, is subsequently utilized in the `geneOntology` function to derive `resInitialGO`, which captures

the initial gene ontology (GO) enrichment results. The `resInitialGO` output is then processed through the `semiLabeling` function to produce `resSemiLabel`, indicating the semi-labeled genes based on their clustering characteristics. This semi-labeled information is further employed in the `semiSupervised` function, yielding `resSemiSupervised`, which includes the final supervised classification outcomes for the unremarkable genes. Finally, the results from `resSemiSupervised` are fed into the `geneOntology` function once more to generate `resFinalGO`, which represents the final GO enrichment analysis.

## Usage

```
data(sgcp)
```

## Format

An object of class `list` containing the `ezSGCP` function information.

## Details

`sgcp` contains a list with the following fields:

`clustering`: List of clustering

- `dropped.indices`: Dropped gene indices.
- `geneID`: Vector of geneIDs.
- `method`: Selected method for determining the number of clusters.
- `k`: Selected number of clusters.
- `Y`: Transformed matrix with  $2*k$  columns.
- `X`: Eigenvalues corresponding to the  $2*k$  columns in `Y`.
- `cluster`: Object of class `kmeans`.
- `clusterLabels`: Vector containing cluster labels for each gene.
- `conductance`: List containing mean, median, and individual cluster conductance indices. Each method's `clusterConductance` field denotes the cluster label with its corresponding conductance index.
- `cvGOdf`: `DataFrame` used for gene ontology validation. For each method, it shows gene ontology enrichment on the cluster with the smallest conductance index.
- `cv`: String indicating the validation method for the number of clusters ("`cvGO`" for gene ontology validation).
- `clusterNumberPlot`: Object of class `ggplot2` for displaying `relativeGap`, `secondOrderGap`, and `additiveGap`.
- `silhouette`: `DataFrame` indicating silhouette values for genes.
- `original`: List with matrix transformation, eigenvalues, and `n_egvec`, retaining the top columns of transformation.

`initial.GO`: List of GO term analysis results for initial clusters

- `GOresults`: `DataFrame` summarizing GO term information.
- `FinalGOTermGenes`: List containing geneIDs of each GO term per cluster.

`semiLabeling`: List of semi-labeling results

- `cutoff`: Numeric indicating selected cutoff.
- `geneLabel`: `DataFrame` with geneID and corresponding cluster label (or NA if unremarkable).

`semiSupervised`: List of semi-supervised learning results

- `semiSupervised`: Object of classification result.

- prediction: Vector of predicted labels for unremarkable genes.
- FinalLabeling: DataFrame of geneID with corresponding semi-label and final label.

final.GO

: List of GO term analysis results for final modules

- GOresults: DataFrame summarizing GO term information.
- FinalGOTermGenes: List containing geneIDs of each GO term per cluster.

### See Also

[SGCP Tutorial](#) [ezSGCP](#)

### Examples

```
library(SGCP)
data(sgcp)
summary(sgcp)

# clustering step
summary(sgcp$clustering)

# initial gene ontology step
summary(sgcp$initial.GO)

# semilabeling step
summary(sgcp$semiLabeling)

# semi-supervised step
summary(sgcp$semiSupervised)

# final gene ontology step
summary(sgcp$final.GO)
```

### Description

The plotting function for ezSGCP results visualizes various aspects. It accepts the ezSGCP output and expression data, generating the following plots: PCA of transformed and expression data, cluster conductance, gene silhouette index, method for determining the number of clusters, distribution and density of gene ontology terms, and cluster performance metrics for both initial clusters and final modules.

### Usage

```
SGCP_ezPLOT(sgcp, expreData, keep = FALSE,
            pdf.file = TRUE, pdfname = "ezSGCP.pdf",
            excel.file = TRUE, xlsxname = "ezSGCP.xlsx",
            w = 6, h = 6, sr = 2, sc = 2, ftype = "png", uni = "in",
            expressionPCA = TRUE, pointSize1 = .5,
```

```

exprePCATitle0 = "Expression Data PCA Without Labels",
exprePCATitle1 = "Expression Data PCA With Initial Labels",
exprePCATitle2 = "Expression Data PCA With Final Labels",
transformedPCA = TRUE, pointSize2 = 0.5,
transformedTitle0 = "Transformed Data PCA Without Labels",
transformedTitle1 = "Transformed Data PCA Initial Labels",
transformedTitle2 = "Transformed Data PCA Final Labels",
conduct = TRUE,
conductanceTitle = "Cluster Conductance Index",
conductx = "clusterLabel", conductivity = "conductance index",
clus_num = TRUE,
silhouette_index = FALSE,
silTitle = "Gene Silhouette Index",
silx = "genes", sily = "silhouette index",
jitt1 = TRUE,
jittTitle1 = "Initial GO p-values", jps1 = 3,
jittx1 = "cluster", jitty1 = "-log10 p-value",
jitt2 = TRUE,
jittTitle2 = "Final GO p-values", jps2 = 3,
jittx2 = "module", jitty2 = "-log10 p-value",
density1 = TRUE,
densTitle1 = "Initial GO p-values Density",
densx1 = "cluster", densy1 = "-log10 p-value",
density2 = TRUE,
densTitle2 = "Final GO p-values Density",
densx2 = "module", densy2 = "-log10 p-value",
mean1 = TRUE,
meanTitle1 = "Cluster Performance",
meanx1 = "cluster", meany1 = "mean -log10 p-value",
mean2 = TRUE,
meanTitle2 = "Module Performance",
meanx2 = "module", meany2 = "mean -log10 p-value",
pie1 = TRUE, pieTitle1 = "Initial GO Analysis",
pie1x = "cluster", pie1y = "count", posx1 = 1.8,
pie2 = TRUE, pieTitle2 = "Final GO Analysis",
pie2x = "module", pie2y = "count", posx2 = 1.8)

```

### Arguments

sgcp	Result from the SGCP pipeline, typically generated by the ezSGCP function.
expreData	Matrix containing the initial gene expression dataset.
keep	Logical, default FALSE. If TRUE, retains plotting objects.
pdf.file	Logical, default TRUE. If TRUE, saves plots in a PDF file.
pdfname	Character string, default "ezSGCP.pdf", name of the PDF file for plots.
excel.file	Logical, default TRUE. If TRUE, saves plots in an Excel file.
xlsxname	Character string, default "ezSGCP.xlsx", name of the Excel file for plots.
w	Numeric, width of plot images in Excel, default 6.
h	Numeric, height of plot images in Excel, default 6.
sr	Numeric, starting row in the Excel sheet, default 2.
sc	Numeric, starting column in the Excel sheet, default 2.

<code>ftype</code>	Character string, plot image type, default "png".
<code>uni</code>	Character string, plot image units, default "in" for inches.
<code>expressionPCA</code>	Logical, default TRUE. If TRUE, plots PCA of gene expression data.
<code>pointSize1</code>	Numeric, point size for expression PCA, default 0.5.
<code>exprePCATitle0</code>	Character string, title for expression PCA plot without labels, default "Expression Data PCA Without Labels".
<code>exprePCATitle1</code>	Character string, title for expression PCA plot with initial cluster labels, default "Expression Data PCA With Initial Labels".
<code>exprePCATitle2</code>	Character string, title for expression PCA plot with final module labels, default "Expression Data PCA With Final Labels".
<code>transformedPCA</code>	Logical, default TRUE. If TRUE, plots PCA of transformed data.
<code>pointSize2</code>	Numeric, point size for transformed PCA, default 0.5.
<code>transformedTitle0</code>	Character string, title for PCA plot without labels on transformed data, default "Transformed Data PCA Without Labels".
<code>transformedTitle1</code>	Character string, title for PCA plot with initial cluster labels on transformed data, default "Transformed Data PCA Initial Labels".
<code>transformedTitle2</code>	Character string, title for PCA plot with final labels on transformed data, default "Transformed Data PCA Final Labels".
<code>conduct</code>	Logical, default TRUE. If TRUE, plots conductance indices for clusters.
<code>conductanceTitle</code>	Character string, title for conductance indices plot, default "Cluster Conductance Index".
<code>conductx</code>	Character string, x-axis title in conductance indices plot, default "clusterLabel".
<code>conducty</code>	Character string, y-axis title in conductance indices plot, default "conductance index".
<code>clus_num</code>	Logical, default TRUE. If TRUE, plots cluster number selection methods.
<code>silhouette_index</code>	Logical, default FALSE. If TRUE, plots silhouette indices for genes.
<code>silTitle</code>	Character string, title for silhouette indices plot, default "Gene Silhouette Index".
<code>silx</code>	Character string, x-axis title in silhouette plot, default "genes".
<code>sily</code>	Character string, y-axis title in silhouette indices plot, default "silhouette index".
<code>jitt1</code>	Logical, default TRUE. If TRUE, plots jitter plot of -log10 p-values of GO terms in initial clusters.
<code>jps1</code>	Numeric, point size in jitter plot for initial clusters, default 3.
<code>jittTitle1</code>	Character string, title for jitter plot for initial clusters, default "Initial GO p-values".
<code>jittx1</code>	Character string, legend for jitter plot for initial clusters, default "cluster".
<code>jitty1</code>	Character string, y-axis title in jitter plot for initial clusters, default "-log10 p-value".
<code>jitt2</code>	Logical, default TRUE. If TRUE, plots jitter plot of -log10 p-values of GO terms in final modules.

jps2	Numeric, point size in jitter plot for final modules, default 3.
jittTitle2	Character string, title for jitter plot for final modules, default "Final GO p-values".
jittx2	Character string, legend for jitter plot for final modules, default "module".
jitty2	Character string, y-axis title in jitter plot for final modules, default "-log10 p-value".
density1	Logical, default TRUE. If TRUE, plots density plot of p-values of GO terms in initial clusters.
densTitle1	Character string, title for density plot for initial clusters, default "Initial GO p-values Density".
densx1	Character string, legend for density plot for initial clusters, default "cluster".
density1	Character string, y-axis title in density plot for initial clusters, default "-log10 p-value".
density2	Logical, default TRUE. If TRUE, plots density plot of p-values of GO terms in final modules.
densTitle2	Character string, title for density plot for final modules, default "Final GO p-values Density".
densx2	Character string, legend for density plot for final modules, default "module".
density2	Character string, y-axis title in density plot for final modules, default "-log10 p-value".
mean1	Logical, default TRUE. If TRUE, plots mean over p-values of GO terms in initial clusters.
meanTitle1	Character string, title for mean plot for initial clusters, default "Cluster Performance".
meanx1	Character string, legend for mean plot for initial clusters, default "cluster".
meany1	Character string, y-axis title in mean plot for initial clusters, default "mean -log10 p-value".
mean2	Logical, default TRUE. If TRUE, plots mean over p-values of GO terms in final modules.
meanTitle2	Character string, title for mean plot for final modules, default "Module Performance".
meanx2	Character string, legend for mean plot for final modules, default "module".
meany2	Character string, y-axis title in mean plot for final modules, default "mean -log10 p-value".
pie1	Logical, default TRUE. If TRUE, plots pie chart of direction and ontology of GO terms for initial clusters.
pieTitle1	Character string, title for pie plot for initial clusters, default "Initial GO Analysis".
piex1	Character string, x-axis title for pie plot for initial clusters, default "cluster".
piey1	Character string, y-axis title in pie plot for initial clusters, default "count".
posx1	Numeric, position of label of -log10 p-value of the most significant term, default 1.8.
pie2	Logical, default TRUE. If TRUE, plots pie chart of direction and ontology of GO terms for final modules.



pieTitle2	Character string, title for pie plot for final modules, default "Final GO Analysis".
piex2	Character string, x-axis title for pie plot for final modules, default "module".
piey2	Character string, y-axis title in pie plot for final modules, default "count".
posx2	Numeric, position of label of $-\log_{10}$ p-value of the most significant term, default 1.8.

**Value**

Returns the plotting object for each plot if keep is TRUE.

**References**

[Aghaieabiane, N and Koutis, I \(2022\) SGCP: A semi-supervised pipeline for gene clustering using self-training approach in gene co-expression networks](#)

**See Also**

[SGCP Tutorial](#)

**Examples**

```
library(SGCP)
library(SummarizedExperiment)

# load the result of ezSGCP function
data(sgcp)

# load the input expression dataset
data(cheng)
expData <- assay(cheng)

# to call the function uncomment the following
## plt <- SGCP_ezPLOT(sgcp = sgcp, expreData = cheng, keep = TRUE)

## print(plt)
```

---

SGCP_plot_bar	<i>Mean p-value bar chart for gene ontology enrichment in the SGCP pipeline</i>
---------------	---

---

**Description**

Generates a bar chart illustrating the average p-values from gene ontology enrichment across the SGCP pipeline.

**Usage**

```
SGCP_plot_bar(df, tit = "mean  $-\log_{10}$  p-values",
              xname = "module", yname = " $-\log_{10}$  p-value")
```

**Arguments**

<code>df</code>	The GOresults dataframe returned by the <code>geneOntology</code> function in the SGCP pipeline.
<code>tit</code>	Plot title (default: "Mean -log10 p-values")
<code>xname</code>	X-axis title (default: "module")
<code>yname</code>	Y-axis title (default: "-log10 p-value")

**Value**

returns the plot, an object of class `ggplot2`.

**References**

[Aghaieabiane, N and Koutis, I \(2022\) SGCP: A semi-supervised pipeline for gene clustering using self-training approach in gene co-expression networks](#)

**See Also**

[geneOntology](#) [SGCP\\_ezPLOT](#) [SGCP Tutorial](#)

**Examples**

```
library(SGCP)
# load the output of geneOntology function
data(resInitialGO)

# call the function

plt <- SGCP_plot_bar(df = resInitialGO$GOresults)
print(plt)
```

---

SGCP\_plot\_conductance *Cluster conductance index bar chart in the SGCP Pipeline*

---

**Description**

Generates a bar chart displaying the cluster conductance index in the SGCP pipeline.

**Usage**

```
SGCP_plot_conductance(conduct, tit = "Clustering Conductance Index",
                      xname = "cluster", yname = "conductance")
```

**Arguments**

<code>conduct</code>	The conductance field returned by the <code>clustering</code> function in the SGCP pipeline.
<code>tit</code>	Plot title (default: "Clustering Conductance Index")
<code>xname</code>	X-axis title (default: "cluster")
<code>yname</code>	Y-axis title (default: "conductance")

**Value**

returns the plot, an object of class ggplot2.

**References**

Aghaieabiane, N and Koutis, I (2022) SGCP: A semi-supervised pipeline for gene clustering using self-training approach in gene co-expression networks

**See Also**

[clustering SGCP\\_ezPLOT](#) [SGCP Tutorial](#)

**Examples**

```
library(SGCP)
# load the output of geneOntology function
data(resClus)

# call the function

plt <- SGCP_plot_conductance(conduct = resClus$conductance)
print(plt)
```

---

SGCP_plot_density	<i>Visualization of gene ontology term p-value distribution in the SGCP pipeline</i>
-------------------	--

---

**Description**

Generates a density chart displaying p-values of gene ontology terms in the SGCP pipeline.

**Usage**

```
SGCP_plot_density(df, tit = "p-values Density",
                  xname = "module", yname = "-log10 p-value")
```

**Arguments**

df	The GOresults dataframe returned by the geneOntology function in the SGCP pipeline.
tit	Plot title (default: "p-values Density")
xname	X-axis title (default: "module")
yname	Y-axis title (default: "-log10 p-value")

**Value**

returns the plot, an object of class ggplot2.

## References

Aghaieabiane, N and Koutis, I (2022) SGCP: A semi-supervised pipeline for gene clustering using self-training approach in gene co-expression networks

## See Also

[geneOntology](#) [SGCP\\_ezPLOT](#) [SGCP Tutorial](#)

## Examples

```
library(SGCP)
# load the output of geneOntology function
data(resInitialGO)

# call the function

plt <- SGCP_plot_density(df = resInitialGO$GOresults)
print(plt)
```

---

SGCP\_plot\_heatMap

*Adjacency matrix heatmap in the SGCP pipeline*

---

## Description

Generates a Heatmap of the the adjacency matrix (network) in the SGCP pipeline.

## Usage

```
SGCP_plot_heatMap(m, tit = "Adjacency Heatmap",
  xname = "genes", yname = "genes")
```

## Arguments

<code>m</code>	An adjacency matrix returned by the <code>adjacencyMatrix</code> function in the SGCP pipeline, or any symmetric matrix with values in (0, 1) excluding the diagonal.
<code>tit</code>	Plot title (default: "Adjacency Heatmap")
<code>xname</code>	X-axis title (default: "genes")
<code>yname</code>	Y-axis title (default: "genes")

## Value

returns the plot, an object of class `ggplot2`.

## References

Aghaieabiane, N and Koutis, I (2022) SGCP: A semi-supervised pipeline for gene clustering using self-training approach in gene co-expression networks

## See Also

[adjacencyMatrix](#) [SGCP\\_ezPLOT](#) [SGCP Tutorial](#)

## Examples

```
library(SGCP)
GeneExpression <- matrix(runif(200, 0,1), nrow = 40, ncol = 5)
diag(GeneExpression) <- 0

## call the function
adja <- adjacencyMatrix(GeneExpression)

plt <- SGCP_plot_heatMap(m = adja)
print(plt)
```

---

SGCP_plot_jitter	<i>P-value jitter chart for gene ontology enrichment in the SGCP pipeline.</i>
------------------	--

---

## Description

Generates a jitter chart illustrating the cluster gene ontology enrichment in the SGCP pipeline.

## Usage

```
SGCP_plot_jitter(df, tit = "p-values Distribution",
                 xname = "module", yname = "-log10 p-value", ps = 3)
```

## Arguments

df	The GOresults dataframe returned by the geneOntology function in the SGCP pipeline.
tit	Plot title (default: "p-values Distribution")
xname	X-axis title (default: "module")
yname	Y-axis title (default: "-log10 p-value")
ps	Numeric value for point size (default: 3)

## Value

returns the plot, an object of class ggplot2.

## References

[Aghaieabiane, N and Koutis, I \(2022\) SGCP: A semi-supervised pipeline for gene clustering using self-training approach in gene co-expression networks](#)

## See Also

[geneOntology](#) [SGCP\\_ezPLOT](#) [SGCP Tutorial](#)

## Examples

```
library(SGCP)
# load the output of geneOntology function
data(resInitialGO)

# call the function

plt <- SGCP_plot_jitter(df = resInitialGO$GOresults)
print(plt)
```

---

SGCP\_plot\_pca

*PCA visualization in the SGCP Pipeline*

---

## Description

Generates Principal Component Analysis (PCA) of the gene expression and transformed gene expression; comparison with and without labels.

## Usage

```
SGCP_plot_pca(m, clusLabs, tit = "PCA plot", ps = .5)
```

## Arguments

<code>m</code>	A numeric matrix of size $n \times m$ .
<code>clusLabs</code>	Either NULL or a vector of size $n$ indicating cluster labels. There is a 1-to-1 correspondence between the rows in <code>m</code> and <code>clusLabs</code> .
<code>tit</code>	Plot title (default: "PCA plot")
<code>ps</code>	Point size (default: 0.5)

## Value

returns the plot, an object of class `ggplot2`.

## References

[Aghaieabiane, N and Koutis, I \(2022\) SGCP: A semi-supervised pipeline for gene clustering using self-training approach in gene co-expression networks](#)

## See Also

[SGCP\\_ezPLOT](#) [SGCP Tutorial](#)

## Examples

```
library(SGCP)
GeneExpression <- matrix(runif(200, 0,1), nrow = 40, ncol = 5)
diag(GeneExpression) <- 0

## call the function
plt <- SGCP_plot_pca(m = GeneExpression, clusLabs = NULL)

print(plt)
```

---

`SGCP_plot_pie`*Gene ontology analysis pie chart in the SGCP pipeline*

---

### Description

Generate a pie chart illustrating the ontology and test direction of gene ontology terms across the SGCP pipeline

### Usage

```
SGCP_plot_pie(df, tit = "GO Analysis",  
              xname = "module", yname = "count", posx = 1.9)
```

### Arguments

<code>df</code>	The GOresults dataframe returned by the geneOntology function in the SGCP pipeline.
<code>tit</code>	Plot title (default: "GO Analysis")
<code>xname</code>	X-axis title (default: "module")
<code>yname</code>	Y-axis title (default: "count")
<code>posx</code>	Numeric value for label position in the pie chart. A higher number places labels further from the pie chart.

### Value

returns the plot, an object of class ggplot2.

### References

[Aghaieabiane, N and Koutis, I \(2022\) SGCP: A semi-supervised pipeline for gene clustering using self-training approach in gene co-expression networks](#)

### See Also

[geneOntology](#) [SGCP\\_ezPLOT](#) [SGCP Tutorial](#)

### Examples

```
library(SGCP)  
# load the output of geneOntology function  
data(resInitialGO)  
  
# call the function  
  
plt <- SGCP_plot_pie(df = resInitialGO$GOresults)  
print(plt)
```

---

SGCP\_plot\_silhouette *Cluster silhouette index chart in the SGCP Pipeline*

---

### Description

Generates a chart displaying the cluster silhouette index in the SGCP pipeline.

### Usage

```
SGCP_plot_silhouette(df, tit = "Gene Silhouette Index",  
                    xname = "genes", yname = "silhouette index")
```

### Arguments

df	The silhouette dataframe returned by the clustering function in the SGCP pipeline.
tit	Plot title (default: "Gene Silhouette Index")
xname	X-axis title (default: "genes")
yname	Y-axis title (default: "silhouette index")

### Details

In order to plot silhouette index, `sil` argument in the `clustering` function must be set to `TRUE`.

### Value

returns the plot, an object of class `ggplot2`.

### References

Aghaieabiane, N and Koutis, I (2022) SGCP: A semi-supervised pipeline for gene clustering using self-training approach in gene co-expression networks

### See Also

[clustering SGCP\\_ezPLOT](#) [SGCP Tutorial](#)

### Examples

```
library(SGCP)  
data(resClus)  
  
## call the function  
plt <- SGCP_plot_silhouette(df = resClus$silhouette)  
  
print(plt)
```



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