Package 'scLink'

October 14, 2022

Title Inferring Functional Gene Co-Expression Networks from Single Cell Data
Version 1.0.1
Description Uses statistical network modeling to understand the coexpression relationships among genes and to construct sparse gene coexpression networks from single-cell gene expression data.
License GPL-3
Depends R (>= 3.5.0), parallel, glasso
Suggests knitr, rmarkdown
VignetteBuilder knitr
Encoding UTF-8
LazyData true
RoxygenNote 7.0.2
NeedsCompilation no
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Repository CRAN
Date/Publication 2020-08-26 14:20:02 UTC
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Description

Calculate scLink's correlation matrix

Usage

```
sclink_cor(expr, ncores, nthre = 20, dthre = 0.9)
```

Arguments

expr	A gene expression matrix with rows representing cells and columns representing genes. Gene names are given as column names. Can be the output of sclink_norm or user constructed gene expression matrices.
ncores	Number of cores if using parallel computation.
nthre	An integer specifying a threshold on the number of complete observations. Defaults to 20.
dthre	A number specifying the threshold on dropout probabilities. Defaults to 0.9.

Value

A correlation matrix for gene co-expression relationships.

Author(s)

```
Wei Vivian Li, <vivian.li@rutgers.edu>
```

Examples

```
count = readRDS(system.file("extdata", "example.rds", package = "scLink"))
count.norm = sclink_norm(count, scale.factor = 1e6, filter.genes = TRUE, n = 500)
corr = sclink_cor(expr = count.norm, ncores = 1)
```

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sclink_net Infer gene co-expression networks
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Description

Infer gene co-expression networks

Usage

```
sclink_net(expr, ncores, lda = seq(1, 0.1, -0.05), nthre = 20, dthre = 0.9)
```

Arguments

expr	A gene expression matrix with rows representing cells and columns representing genes. Gene names are given as column names. Can be the output of sclink_norm or user constructed gene expression matrices.
ncores	Number of cores if using parallel computation.
lda	A vector specifying a sequence of lambda values to be used in the penalized likelihood.
nthre	An integer specifying a threshold on the number of complete observations. Defaults to 20.
dthre	A number specifying the threshold on dropout probabilities. Defaults to 0.9.

Value

A list for gene co-expression relationships. The list contains a cor element for scLink's correlation matrix and a summary element for the gene networks. summary is a list with each element corresponding to the result of one lambda value. Each element of summary contains the following information:

adj: the adjacency matrix specifying the gene-gene edges;

Sigma: the estimated concentration matrix; **nedge:** number of edges in the gene network;

bic: BIC score;

lambda: value of lambda in the penalty.

Author(s)

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Examples

```
count = readRDS(system.file("extdata", "example.rds", package = "scLink"))
count.norm = sclink_norm(count, scale.factor = 1e6, filter.genes = TRUE, n = 500)
networks = sclink_net(expr = count.norm, ncores = 1, lda = seq(0.5, 0.1, -0.05))
```

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sclink_norm

Pre-process data for scLink

Description

Pre-process data for scLink

Usage

```
sclink_norm(
  count,
  scale.factor = 1e+06,
  filter.genes = FALSE,
  gene.names = NULL,
  n = 500
)
```

Arguments

count	A full gene count matrix with rows representing cells and columns representing genes. Gene names are given as column names.
scale.factor	A number specifying the sclae factor used for library size normalization. Defaults to 1e6.
filter.genes	A Boolean specifying whether scLink should select genes based on mean expression. When set to FALSE, users need to speicfy a set of genes to be used for network construction with gene.names. When set to TRUE, scLink will select genes based on their mean expression, and users need to specify the number of genes to be selected with n.
gene.names	A character vector specifying the genes used for network construction. Only needed when filter.genes = FALSE.
n	An integer specifying the number of genes to be selected by scLink (defaults to 500). Only needed when filter.genes = TRUE.

Value

A transformed and normalized gene expression matrix that can be used for correlation calculation and network construction.

Author(s)

```
Wei Vivian Li, <vivian.li@rutgers.edu>
```

Examples

```
count = readRDS(system.file("extdata", "example.rds", package = "scLink"))
count.norm = sclink_norm(count, scale.factor = 1e6, filter.genes = TRUE, n = 500)
```

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