Package ‘MODA’

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Type Package

Title MODA: MOdule Differential Analysis for weighted gene co-expression network

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Description MODA can be used to estimate and construct condition-specific gene co-expression networks, and identify differentially expressed subnetworks as conserved or condition specific modules which are potentially associated with relevant biological processes.

License GPL (>= 2)

Depends R (>= 3.1.0)

Imports WGCNA,dynamicTreeCut,igraph

RoxygenNote 5.0.1

biocViews GeneExpression, Microarray, DifferentialExpression, Network

Suggests BiocStyle, knitr

VignetteBuilder knitr

NeedsCompilation no

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Description

Compare the background network and a set of condition-specific network. Conserved or condition-specific modules are indicated by the plain files, based on the statistics.

Usage

```
CompareAllNets(ResultFolder, intModules, speciesName, intconditionModules, conditionNames, specificTheta, conservedTheta)
```

Arguments

- **ResultFolder**: where to store results
- **intModules**: how many modules in the background network
- **speciesName**: identifier of current profile, served as a tag in name
- **intconditionModules**: a numeric vector, each of them is the number of modules in each condition-specific network. Or just single number
- **conditionNames**: a character vector, each of them is the name of condition. Or just single name
- **specificTheta**: the threshold to define min(s)+specificTheta, less than which is considered as condition specific module. s is the sums of rows in Jaccard index matrix. See supplementary file.
- **conservedTheta**: The threshold to define max(s)-conservedTheta, greater than which is considered as condition conserved module. s is the sums of rows in Jaccard index matrix. See supplementary file.

Value

None

Author(s)

Dong Li, <dxl466@cs.bham.ac.uk>

See Also

`WeightedModulePartitionDensity`, `comparemodulestwonets`

Examples

```
data(synthetic)
ResultFolder = 'ForSynthetic' # where middle files are stored
CuttingCriterion = 'Density' # could be Density or Modularity
indicator1 = 'X' # indicator for data profile 1
indicator2 = 'Y' # indicator for data profile 2
specificTheta = 0.1 #threshold to define condition specific modules
conservedTheta = 0.1#threshold to define conserved modules
intModules1 <- WeightedModulePartitionDensity(datExpr1,ResultFolder,
```
**comparemodulestwonets**

```r
indicator1,CuttingCriterion)
intModules2 <- WeightedModulePartitionDensity(datExpr2,ResultFolder,
indicator2,CuttingCriterion)
CompareAllNets(ResultFolder,intModules1,indicator1,intModules2,indicator2,
specificTheta,conservedTheta)
```

---

**Description**

Compare the background network and a condition-specific network. A Jaccard index is used to measure the similarity of two sets, which represents the similarity of each module pairs from two networks.

**Usage**

```r
comparemodulestwonets(sourcehead, nm1, nm2, ind1, ind2)
```

**Arguments**

- `sourcehead`: prefix of where to store results
- `nm1`: how many modules in the background network
- `nm2`: how many modules in the condition-specific network
- `ind1`: indicator of the background network
- `ind2`: indicator of the condition-specific network

**Value**

A matrix where each entry is the Jaccard index of corresponding modules from two networks

**Author(s)**

Dong Li, <dxl466@cs.bham.ac.uk>

**Examples**

```r
data(synthetic)
ResultFolder = 'ForSynthetic' # where middle files are stored
CuttingCriterion = 'Density' # could be Density or Modularity
indicator1 = 'X' # indicator for data profile 1
indicator2 = 'Y' # indicator for data profile 2
intModules1 <- WeightedModulePartitionDensity(datExpr1,ResultFolder,
indicator1,CuttingCriterion)
intModules2 <- WeightedModulePartitionDensity(datExpr2,ResultFolder,
indicator2,CuttingCriterion)
JaccardMatrix <- comparemodulestwonets(ResultFolder,intModules1,intModules2,
paste('/DenseModuleGene_',indicator1,sep=''),
paste('/DenseModuleGene_',indicator2,sep=''))
```
**Description**

Synthetic gene expression profile with 20 samples and 500 genes.

**Format**

A matrix with 20 rows and 500 columns.

**Author(s)**

Dong Li, <dxl466@cs.bham.ac.uk>

**Examples**

```
data(synthetic)
## plot the heatmap of the correlation matrix ...
## Not run: heatmap(cor(as.matrix(datExpr1)))
```
**PartitionDensity**

**Illustration of partition density**

**Description**
Calculate the average density of all resulting modules from a partition. The density of each module is defined as the average adjacency of the module genes.

**Usage**

\[ \text{PartitionDensity}(ADJ, \text{PartitionSet}) \]

**Arguments**

- **ADJ**: gene similarity matrix
- **PartitionSet**: vector indicates the partition label for genes

**Value**
partition density, defined as average density of all modules

**Author(s)**
Dong Li, <dxl466@cs.bham.ac.uk>

**References**

**Examples**

```r
data(synthetic)
ADJ1=abs(cor(datExpr1,use="p"))\(^10\)
dissADJ=1-ADJ1
hierADJ=hclust(as.dist(dissADJ), method="average")
groups <- cutree(hierADJ, h = 0.8)
pDensity <- PartitionDensity(ADJ1,groups)
```

---

**PartitionModularity**

**Illustration of modularity density**

**Description**
Calculate the average modularity of a partition. The modularity of each module is defined from a natural generalization of unweighted case.

**Usage**

\[ \text{PartitionModularity}(ADJ, \text{PartitionSet}) \]

**Examples**

```r
data(synthetic)
```

---
WeightedModulePartitionDensity

Arguments

ADJ    gene similarity matrix
PartitionSet vector indicates the partition label for genes

Value

partition modularity, defined as average modularity of all modules

Author(s)

Dong Li, <dxl466@cs.bham.ac.uk>

References


Examples

data(synthetic)
ADJ1=abs(cor(datExpr1,use="p"))\*10
dissADJ=1-ADJ1
hierADJ=hclust(as.dist(dissADJ), method="average")
groups <- cutree(hierADJ, h = 0.8)
pDensity <- PartitionModularity(ADJ1,groups)

WeightedModulePartitionDensity

Illustration of Modules detection

Description

Module detection based on the optimal cutting height of dendrogram, which is selected to make the average density or modularity of resulting partition maximal. The clustering and visualization function are from WGCNA.

Usage

WeightedModulePartitionDensity(datExpr, foldername, indicatename, cutmethod = c("Density", "Modularity"), power = 10)

Arguments

datExpr    gene expression profile, rows are samples and columns genes
foldername where to store the clusters
indicatename normally a specific tag of condition
cutmethod cutting the dendrogram based on maximal average Density or Modularity
power    the power parameter of WGCNA, W_\ij=|cor(x_i,x_j)|^power
**Value**

The number of clusters

**Author(s)**

Dong Li, <dx1466@cs.bham.ac.uk>

**References**


**See Also**

PartitionDensity

PartitionModularity

**Examples**

data(synthetic)
ResultFolder = 'ForSynthetic' # where middle files are stored
CuttingCriterion = 'Density' # could be Density or Modularity
indicator1 = 'X' # indicator for data profile 1
indicator2 = 'Y' # indicator for data profile 2
specificTheta = 0.1 #threshold to define condition specific modules
conservedTheta = 0.1#threshold to define conserved modules
intModules1 <- WeightedModulePartitionDensity(datExpr1,ResultFolder,
indicator1,CuttingCriterion)
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