# Package 'cooccurNet'

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Title Co-Occurrence Network

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| Description  Read and preprocess fasta format data, and construct the co-occurrence network for downstream analyses. This R package is to construct the co-occurrence network with the algorithm developed by Du (2008) <doi:10.1101 gr.6969007="">. It could be used to transform the data with high-dimension, such as DNA or protein sequence, into co-occurrence networks. Co-occurrence network could not only capture the co-variation pattern between variables, such as the positions in DNA or protein sequences, but also reflect the relationship between samples. Although it is originally used in DNA and protein sequences, it could be also used to other kinds of data, such as RNA, SNP, etc.</doi:10.1101> |
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changeLog

changeLog

## Description

Get the most recent n lines

## Usage

```
changeLog(n=20)
```

## Arguments

n

numeric, 20 by default, the number of lines will be shown up.

## **Details**

changeLog

## Value

list, the most recent n lines

## Note

Once you have installed cooccurNet, the change-log can also be viewed from the R prompt.

```
logs = changeLog(n=20)
```

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cooccurNet

cooccurNet: Co-Occurrence network computation for R

#### **Description**

Read and preprocess fasta format data, and construct the co-occurrence network for downstream analyses. This R package is to construct the co-occurrence network with the algorithm developed by Du (2008) <DOI:10.1101/gr.6969007>. It could be used to transform the data with high-dimension, such as DNA or protein sequence, into co-occurrence networks. Co-occurrence network could not only capture the co-variation pattern between variables, such as the positions in DNA or protein sequences, but also reflect the relationship between samples. Although it is originally used in DNA and protein sequences, it could be also used to other kinds of data, such as RNA, SNP, etc.

#### **Details**

Index of functions/methods (grouped in a friendly way):

- 1. readseq(dataFile="", dataType="protein", debug=FALSE)
- 2. pprocess(data=list(), conservativeFilter=0.95, memory=NULL, debug=FALSE)
- 3. gencooccur(data=list(), cooccurFilter=NULL, networkFile='cooccurNetwork', module=FALSE, moduleFile='cooccurNetworkModule', property=FALSE, propertyFile='cooccurNetworkProperty', siteCo=FALSE, siteCoFile='siteCooccurr', sampleTimes=100, debug=FALSE, parallel=FALSE)
- 4. coocnet(dataFile="", dataType="protein", conservativeFilter=0.95, memory=NULL, cooccurFilter=NULL, networkFile='cooccurNetwork', module=FALSE,moduleFile='cooccurNetworkModule', property=FALSE, propertyFile='cooccurNetworkProperty', siteCo=FALSE, siteCoFile='siteCooccurr', sampleTimes=100, debug=FALSE, parallel=FALSE)
- 5. siteco(dataFile="", dataType="protein", conservativeFilter=0.95, memory=NULL, cooccurFilter=NULL, siteCoFile='siteCooccurr', sampleTimes=100, debug=FALSE, parallel=FALSE)
- toigraph(networkFile="", networkNames=c())
- 7. getexample(dataType)
- 8. getexample\_forRCOS(dataType)
- 9. changeLog(n=20)

#### Note

Arguments:

1. dataFile

type: character

description: a FASTA data file name with full path.

2. dataType type: character

description: 'protein' by default, the type of data will be processed. It could be 'DNA', 'RNA', 'protein', 'SNP' or 'other'.

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#### 3. conservativeFilter

type: numeric

description: a number in the range of 0~1, 0.95 by default. It's used to filter the highly conservative columns which the ratio of some residue is larger than the conservationFilter.

## 4. cooccurFilter

type: numeric

description: a number in the range of  $0\sim1$ . It determines whether two columns are perfect co-occurrence. In default, for the data type of protein, it is set to be 0.9, while for the other data types, it is set to be 1.

#### 5. networkFile

type: character

description: 'cooccurNetwork' be default. It is a file name with full path for storing the cooccurrence network for each row.

#### 6. module

type: logical

description: FALSE by default, to check whether the modules in each network of the networkFile would be calculated.

#### 7. moduleFile

type: character

description: 'cooccurNetworkModule' by default. It is a file name with full path for storing the modules for co-occurrence network.

## 8. property

type: logical

description: FALSE by default, to check whether the properties for each network of the networkFile, including the network diameter, connectivity, ConnectionEffcient and so on, would be calculated.

## 9. propertyFile

type: character

description: 'cooccurNetworkProperty' by default. It is a file name with full path storing the properties for each network of the networkFile.

#### 10. siteCo

type: logical

description: FALSE by default, to check whether the residue co-occurence file would be calculated.

## 11. siteCoFile

type: character

description: 'siteCooccurr' by default. It is a file name with full path for storing the RCOS between all pairs of columns, and the related p-values.

#### 12. sampleTimes

type: numeric

description: a integer of permutations in the simulation when calculating the p-values. It should be greater than 100.

#### 13.debug

type: logical

description: FALSE by default, indicates whether the debug message will be displayed or not.

14.memory type: character

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description: the type of matrix, NULL by default. It could be 'memory' or 'sparse'. If it's set to be 'memory', all data would be manipulated in the RAM by using normal matrix and package 'bigmemory'. If it's set to be 'sparse', the package "Matrix" would be used to manipulate massive matrices in memory and initialize huge sparse matrix, which could significantly reduce the RAM consumed. In default, it is set to be NULL, so that the system would determine automatically whether all data is manipulated in the RAM or not, according to the size of data inputted and the RAM available for R.

15. parallel

type: logical

description: FALSE by default. It only supports Unix/Mac (not Windows) system.

Output files:

1. networkFile

filePath = data\$networkFile

description: A co-occurrence network (networkFile) was given for each sequence, which could be easily transformed into the format of igraph by the function toigraph().

2. moduleFile

filePath = data\$moduleFile

description: The inherent modules(moduleFile) in each co-occurrence network were given for each sequence.

3. propertyFile

filePath = data\$propertyFile

description: Some basic network attributes(propertyFile) such as connectivity and clustering coefficient for each network were given.

4. siteCoFile

filePath = data\$siteCoFile

description: The extent of co-occurrence between residues(siteCoFile), defined as residue co-occurrence score (RCOS), were given for all pairs of residues.

For more details, please see the 'Vignette: Extending cooccurNet' by using the following command. vignette("Extending-cooccurNet",package="cooccurNet")

#### Author(s)

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#### References

Du, X., Wang, Z., Wu, A., Song, L., Cao, Y., Hang, H., & Jiang, T. (2008). Networks of genomic co-occurrence capture characteristics of human influenza A (H3N2) evolution. Genome research, 18(1), 178-187. doi:10.1101/gr.6969007

## Examples

```
#example of get example data
dataFile=getexample(dataType="protein")
```

#example of get file paths for all the files available for testing RCOS.

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```
#dataFiles = getexample_forRCOS()
#example of readseq()
#read sequences from the sample fasta file
#data = readseq(dataFile=dataFile, dataType="protein")
#example of pprocess()
#preprocess the sequence dataFile
#data_process = pprocess(data=data,conservativeFilter=0.95)
#example of gencooccur()
#generate co-occurrence network and save it into the 'networkFile'
#cooccurNetwork = gencooccur(data=data_process, cooccurFilter=0.9, networkFile='cooccurNetwork')
#check the 'networkFile' path
#print(cooccurNetwork$networkFile)
#example of coocnet()
#also, you can generate the co-occurrence network by one command
#cooccurNetwork = coocnet(dataFile=getexample(dataType="protein"), dataType="protein")
#check the 'networkFile' path
#print(cooccurNetwork$networkFile)
#example of siteco()
#you can generate the co-occurrence network siteCoFile by one command
#this command will take long time to calculate the p-value.
#pairwiseCooccur = siteco(dataFile=getexample(dataType="protein"), dataType="protein")
#check the 'siteCoFile' path
#print(pairwiseCooccur$siteCoFile)
#example of toigraph()
#you can transform a network file to the igraph.data.frame
#cooccurNetwork = coocnet(dataFile=getexample(dataType="protein"),dataType="protein")
#get igraph data frame by specifying the network name
#network_igraph = toigraph(networkFile=cooccurNetwork$networkFile, networkName=c("EPI823725"))
#Plot the network (The package "igraph" must be installed and loaded firstly)
#read the names of network
#networkName = cooccurNetwork$xnames
#Transform all cooccurrence network into the igraph format
#Network_igraph = toigraph(networkFile=cooccurNetwork$networkFile, networkNames=networkName)
#example of changelog()
#logs = changeLog(n=20)
```

coocnet

coocnet

#### **Description**

Read and preprocess data, and construct the co-occurrence network in one step.

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#### Usage

```
coocnet(dataFile = "", dataType = "protein", conservativeFilter = 0.95,
  cooccurFilter = NULL, networkFile = "cooccurNetwork", module = FALSE,
  moduleFile = "cooccurNetworkModule", property = FALSE,
  propertyFile = "cooccurNetworkProperty", siteCo = FALSE,
  siteCoFile = "siteCooccurr", sampleTimes = 100, debug = FALSE,
  parallel = FALSE, memory = NULL)
```

#### **Arguments**

dataFile character, a FASTA data file name with full path.

dataType character, 'protein' by default, the type of data will be processed. It could be

'DNA', 'RNA', 'protein', 'SNP' or 'other'.

conservativeFilter

numeric, a number in the range of  $0\sim1$ , 0.95 by default. It's used to filter the highly conservative columns which the ratio of some residue is larger than the

conservationFilter.

cooccurFilter numeric, a number in the range of 0~1. It determines whether two columns are

perfect co-occurrence. In default, for the data type of protein, it is set to be 0.9,

while for the other data types, it is set to be 1.

networkFile character, 'cooccurNetwork' be default. It is a file name with full path for storing

the co-occurrence network for each row.

module logic, FALSE by default, to check whether the modules in each network of the

networkFile would be calculated.

moduleFile character, 'cooccurNetworkModule' by default. It is a file name with full path

for storing the modules for co-occurrence network.

property logic, FALSE by default, to check whether the properties for each network of the

networkFile, including the network diameter, connectivity, ConnectionEffcient

and so on, would be calculated.

propertyFile character, 'cooccurNetworkProperty' by default. It is a file name with full path

storing the properties for each network of the networkFile.

siteCo logic, FALSE by default, to check whether the residue co-occurence file would

be calculated.

siteCoFile character, 'siteCooccurr' by default. It is a file name with full path for storing

the RCOS between all pairs of columns, and the related p-values.

sampleTimes numeric, an integer of permutations in the simulation when calculating the p-

values. It should be greater than 100.

debug logic, FALSE by default, indicates whether the debug message will be displayed

or not.

parallel logic, FALSE by default. It only supports Unix/Mac (not Windows) system.

memory character, the type of matrix, NULL by default. It could be 'memory' or 'sparse'.

If it's set to be 'memory', all data would be manipulated in the RAM by using normal matrix and package 'bigmemory'. If it's set to be 'sparse', the package "Matrix" would be used to manipulate massive matrices in memory and initialize

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huge sparse matrix, which could significantly reduce the RAM consumed. In default, it is set to be NULL, so that the system would determine automatically whether all data is manipulated in the RAM or not, according to the size of data inputted and the RAM available for R.

#### Value

list, all the output file paths are attributed in it.

The attribute "networkFile" stores the co-occurrence network for each row;

The attribute "moduleFile" is optional. When the module is set to be TRUE, it would be output. It stores the modules for co-occurrence network;

The attribute "propertyFile" is optional. When the property is set to be TRUE, it would be output. It stores the properties for co-occurrence network;

The attribute "siteCoFile" is optional. When the property is set to be TRUE, it would be output. It stores all the pairwise siteCos between columns.

#### References

Du, X., Wang, Z., Wu, A., Song, L., Cao, Y., Hang, H., & Jiang, T. (2008). Networks of genomic co-occurrence capture characteristics of human influenza A (H3N2) evolution. Genome research, 18(1), 178-187. doi:10.1101/gr.6969007

#### **Examples**

```
cooccurNetwork = coocnet(dataFile=getexample(dataType="protein"), dataType="protein")
```

gencooccur

gencooccur

#### Description

Construct the co-occurrence network

#### Usage

```
gencooccur(data = list(), cooccurFilter = NULL,
  networkFile = "cooccurNetwork", module = FALSE,
  moduleFile = "cooccurNetworkModule", property = FALSE,
  propertyFile = "cooccurNetworkProperty", siteCo = FALSE,
  siteCoFile = "siteCooccurr", sampleTimes = 100, debug = FALSE,
  parallel = FALSE)
```

#### **Arguments**

data list, returns from the function "pprocess()"

cooccurFilter

numeric, a number in the range of  $0\sim1$ . It determines whether two columns are perfect co-occurrence. In default, for the data type of protein, it is set to be 0.9, while for the other data types, it is set to be 1.

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| networkFile  | character, 'cooccurNetwork' be default. It is a file name with full path for storing the co-occurrence network for each row.   |
|--------------|--|
| module       | logic, FALSE by default, to check whether the modules in each network of the networkFile would be calculated.  |
| moduleFile   | character, 'cooccurNetworkModule' by default. It is a file name with full path for storing the modules for co-occurrence network.  |
| property     | logic, FALSE by default, to check whether the properties for each network of the networkFile, including the network diameter, connectivity, ConnectionEffcient and so on, would be calculated. |
| propertyFile | character, 'cooccurNetworkProperty' by default. It is a file name with full path storing the properties for each network of the networkFile.   |
| siteCo       | logic, FALSE by default, to check whether the residue co-occurence file would be calculated.   |
| siteCoFile   | character, 'siteCooccurr' by default. It is a file name with full path for storing the RCOS between all pairs of columns, and the related p-values.  |
| sampleTimes  | numeric, an integer of permutations in the simulation when calculating the p-values. It should be greater than 100.  |
| debug        | logic, FALSE by default, indicates whether the debug message will be displayed or not.   |
| parallel     | logic, FALSE by default. It only supports Unix/Mac (not Windows) system.   |

#### Value

list, all the output file paths are attributed in it.

The attribute "networkFile" stores the co-occurrence network for each row;

The attribute "moduleFile" is optional. When the module is set to be TRUE, it would be output. It stores the modules for co-occurrence network;

The attribute "propertyFile" is optional. When the property is set to be TRUE, it would be output. It stores the properties for co-occurrence network;

The attribute "siteCoFile" is optional. When the property is set to be TRUE, it would be output. It stores all the pairwise siteCos between columns.

## References

Du, X., Wang, Z., Wu, A., Song, L., Cao, Y., Hang, H., & Jiang, T. (2008). Networks of genomic co-occurrence capture characteristics of human influenza A (H3N2) evolution. Genome research, 18(1), 178-187. doi:10.1101/gr.6969007

```
data = readseq(dataFile=getexample(dataType="protein"),dataType="protein")
data_process = pprocess(data=data)
#cooccurNetwork = gencooccur(data=data_process)
```

getexample

getexample

## Description

Get the example data

## Usage

```
getexample(dataType)
```

## **Arguments**

dataType

character, 'protein' by default. It could be 'DNA', 'RNA', 'protein', 'SNP' or 'other'.

#### **Details**

getexample

#### Value

character, the file path of the example data

#### Note

Both the dataset "DNA" and "protein" are sampled from the Hemagglutinin sequences of human influenza H3N2 viruses, while the dataset "SNP" are simulated.

## **Examples**

```
dataFile = getexample(dataType='protein')
```

getexample\_forRCOS

getexample\_forRCOS

## Description

Get the test data for testing RCOS method

## Usage

```
getexample_forRCOS()
```

#### **Details**

getexample\_forRCOS

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#### Value

character, the files available for testing RCOS.

#### Note

File paths for all the files available for testing RCOS. Currently, there is only one file "HA1protein humanH3N2", the sequences within which are derived from the database of Influenza Virus Resource.

#### **Examples**

```
dataFile = getexample_forRCOS()
```

pprocess

pprocess

#### **Description**

Filter the conservative columns (defined as the conservative score greater than the "conservative-Filter")

#### Usage

```
pprocess(data=list(), conservativeFilter=0.95, debug=FALSE, memory=NULL)
```

## **Arguments**

data

list, returns from the function "readseq()".

conservativeFilter

numeric, a number in the range of 0~1, 0.95 by default. It's used to filter the highly conservative columns which the ratio of some residue is larger than the

conservationFilter.

debug

logic, FALSE by default, indicates whether the debug message will be displayed or not.

memory

character, the type of matrix, NULL by default. It could be 'memory' or 'sparse'. If it's set to be 'memory', all data would be manipulated in the RAM by using normal matrix and package 'bigmemory'. If it's set to be 'sparse', the package "Matrix" would be used to manipulate massive matrices in memory and initialize huge sparse matrix, which could significantly reduce the RAM consumed. In default, it is set to be NULL, so that the system would determine automatically whether all data is manipulated in the RAM or not, according to the size of data inputted and the RAM available for R.

## Value

list, contains the original data matrix, frequency matrix and other informations.

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#### References

Du, X., Wang, Z., Wu, A., Song, L., Cao, Y., Hang, H., & Jiang, T. (2008). Networks of genomic co-occurrence capture characteristics of human influenza A (H3N2) evolution. Genome research, 18(1), 178-187. doi:10.1101/gr.6969007

## **Examples**

```
data = readseq(dataFile=getexample(dataType="protein"),dataType="protein")
data_process = pprocess(data=data,conservativeFilter=0.95)
```

## **Description**

Reading a sequence file in fasta format

## Usage

```
readseq(dataFile="", dataType="protein", debug=FALSE)
```

## Arguments

| dataFile | character, a FASTA data file name with full path.   |
|----------|---|
| dataType | character, 'protein' by default, the type of data will be processed. It could be 'DNA', 'RNA', 'protein', 'SNP' or 'other'. |
| debug    | logic, FALSE by default, to indicate whether the debug message will be displayed or not.                                    |

#### Value

list, contains the original data matrix and some other information.

```
data = readseq(dataFile=getexample(dataType="protein"), dataType="protein")
```

siteco 13

#### Description

Read and preprocess data, and calculate the pairwise site co-occurrence in one step

#### Usage

```
siteco(dataFile = "", dataType = "protein", conservativeFilter = 0.95,
  cooccurFilter = NULL, siteCoFile = "siteCooccurr", sampleTimes = 100,
  debug = FALSE, parallel = FALSE, memory = NULL)
```

## **Arguments**

dataFile character, a FASTA data file name with full path.

dataType character, 'protein' by default, the type of data will be processed. It could be

'DNA', 'RNA', 'protein', 'SNP' or 'other'.

conservativeFilter

numeric, a number in the range of  $0\sim1$ , 0.95 by default. It's used to filter the highly conservative columns which the ratio of some residue is larger than the

conservationFilter.

cooccurFilter numeric, a number in the range of 0~1. It determines whether two columns are

perfect co-occurrence. In default, for the data type of protein, it is set to be 0.9,

while for the other data types, it is set to be 1.

siteCoFile character, 'siteCooccurr' by default. It is a file name with full path for storing

the RCOS between all pairs of columns, and the related p-values.

sampleTimes numeric, an integer of permutations in the simulation when calculating the p-

values. It should be greater than 100.

debug logic, FALSE by default, indicates whether the debug message will be displayed

or not.

parallel logic, FALSE by default. It only supports Unix/Mac (not Windows) system.

memory character, the type of matrix, NULL by default. It could be 'memory' or 'sparse'.

If it's set to be 'memory', all data would be manipulated in the RAM by using normal matrix and package 'bigmemory'. If it's set to be 'sparse', the package "Matrix" would be used to manipulate massive matrices in memory and initialize huge sparse matrix, which could significantly reduce the RAM consumed. In default, it is set to be NULL, so that the system would determine automatically whether all data is manipulated in the RAM or not, according to the size of data

inputted and the RAM available for R.

## Value

list, the output file path of 'sitecoFile' is attributed in it. The file stores all the pairwise siteCos between columns.

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#### References

Du, X., Wang, Z., Wu, A., Song, L., Cao, Y., Hang, H., & Jiang, T. (2008). Networks of genomic co-occurrence capture characteristics of human influenza A (H3N2) evolution. Genome research, 18(1), 178-187. doi:10.1101/gr.6969007

## **Examples**

```
#pairwiseCooccur = siteco(dataFile=getexample(dataType="protein"), dataType="protein")
```

toigraph

toigraph

## Description

transform a network file to the igraph.data.frame type by specifying a network file name and a network name

## Usage

```
toigraph(networkFile="", networkNames=c())
```

#### **Arguments**

networkFile

character, a file name with full path for storing the co-occurrence network for

each row and generated from cooc() or gencooccur().

networkNames

character, a vector of network names

#### Value

a list of igraph::graph.data.frame object ordered by the input network names.

```
#cooccurNetwork = cooc(dataFile=getexample(dataType="protein"),dataType="protein")
#network_igraph = toigraph(networkFile=cooccurNetwork$networkFile, networkName=c("EPI823725")
```

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