Package ‘CoRegNet’

January 31, 2017

Type Package

Title CoRegNet : reconstruction and integrated analysis of co-regulatory networks

Version 1.10.0

Date 2015-03-25

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Description This package provides methods to identify active transcriptional programs. Methods and classes are provided to import or infer large scale co-regulatory network from transcriptomic data. The specificity of the encoded networks is to model Transcription Factor cooperation. External regulation evidences (TFBS, ChIP,...) can be integrated to assess the inferred network and refine it if necessary. Transcriptional activity of the regulators in the network can be estimated using an measure of their influence in a given sample. Finally, an interactive UI can be used to navigate through the network of cooperative regulators and to visualize their activity in a specific sample or subgroup sample. The proposed visualization tool can be used to integrate gene expression, transcriptional activity, copy number status, sample classification and a transcriptional network including co-regulation information.

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Depends R (>= 2.14), igraph, shiny, arules, methods

Suggests RColorBrewer, gplots, BiocStyle, knitr

VignetteBuilder knitr

biocViews NetworkInference, NetworkEnrichment, GeneRegulation, GeneExpression, GraphAndNetwork, SystemsBiology, Network, Visualization, Transcription

NeedsCompilation yes

R topics documented:

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coRegnet-package

coRegnet : inference and interrogation co-regulation networks

Description

coRegnet can be used to infer and analyse regulatory networks. Although it can be used with any network, it has its own regulatory network inference routine based on a hybrid version of LICORN which combines both a discrete and a statistical model to infer regulatory network with an emphasis on regulator cooperativity.

Author(s)

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References


addEvidences

Integration of regulatory evidences.

Description

These functions can be used to integrate external data on gene regulation and co-regulation to enrich an inferred regulatory network. Additional regulatory data sets can include : ChIP-seq data, ChIP on chip, target gene promoter analysis for Transcription Factor Binding Site models, epigenetic marks or even networks inferred by other methods. Cooperative regulation data sets can include : Protein interactions, significant binding site overlap, co-localization. These additional evidence are added to enrich the network and the enrichment of the inferred interaction is tested.
addEvidences

Usage

addEvidences(object,...)
addCooperativeEvidences(object,...)

Arguments

object	A regulatory network inferred by the hLICORN function or with an externally inferred network importing through the CoRegNet class.
...
A single or several data sets of regulatory data as pairs of Transcription Factor to Target Gene regulation interactions or of pairs of transcription factors for cooperative evidences. These should be given in the form of a two column data.frame or matrix. The characters in the input dataset should correspond to the characters of the regulators and the genes in the network (accessible through regulators targets respectively).

Details

A single or several datasets of regulatory interactions can be added and enrich an inferred regulatory network. Below is an example of the format of the input data A character matrix or data.frame with two columns the first for target genes and the second for Transcription Factor.

[,1] [,2]
[1,] "TF1" "Gene1"
[2,] "TF1" "Gene2"
[3,] "TF2" "Gene1"
...

Value

A Gene Regulatory Network in a CoRegNet objects with additional informations in the core object.

Note

The variables used as the input are kept and will be used later on to refine the network. The names of target genes and of TF need to be identical to the ones in the expression dataset.

Author(s)

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See Also

refine hLICORN

Examples

#Creating a synthetic network. Upper case letters are target genes, lower case letters are regulators
acts=apply(matrix(rep(letters[1:4],4),nrow=2),2,paste,collapse=" ")
reps=apply(matrix(rep(letters[5:8],4),nrow=2),2,paste,collapse=" ")
grn=data.frame("Target"= LETTERS[1:16],"coact"=c(acts,reps),"corep"= c(reps,acts),"R2"=runif(16),stringsAsFactors=FALSE)
co=coregnet(grn)
coregnet

#Creating dummy regulatory evidence data in a data.frame
evidence1=unique(data.frame(tf=sample(letters[1:8],50, replace=TRUE),target=sample(LETTERS[1:16],50, replace=TRUE),stringsAsFactors = FALSE))
evidence2=unique(data.frame(tf=sample(letters[1:8],50, replace=TRUE),target=sample(LETTERS[1:16],50, replace=TRUE),stringsAsFactors = FALSE))

#Creating dummy cooperative evidence data in a data.frame
coregevidence1=unique(data.frame(tf1=sample(letters[1:8],50, replace=TRUE),target=sample(letters[1:8],50, replace=TRUE),stringsAsFactors = FALSE))
enrichco=addEvidences(co,evidence1,evidence2)
enrichco=addCooperativeEvidences(enrichco,coregevidence1)

---
coregnet  Initialize a co-regulatory network object.

Description

Given a previously built network encoded in a simple data.frame, initialize a co-regulatory network object coregnet. This can be used on networks inferred from gene expression data by other algorithms (ARACNe, GENIE3 ...) but also on ChIP-seq data or on network based on Transcription Factor Binding Site analysis. If the input network does not contain a description of the type of regulation (activation, inhibition) an expression data set is needed.

Usage

coregnet(GRN, expressionDATA = NULL)

Arguments

GRN  A data.frame containing the network. A two column data.frame should contain the regulator in column one and the target gene in column two. A three column data.frame should contain the target in column one, all the co-activators and co-repressors separated by a space in column two and three respectively.

expressionDATA  If the input GRN is only a two column data.frame, the expression data is needed to set each regulator of a target gene as an activator or a repressor.

Details

In the case of a two column data.frame, the pearson correlation coefficient is used to determine whether a given regulator is an activator (R-squared >= 0) or a repressor (R-squared < 0).

In order to import a network and initialize a new coregnet object the input data.frame should have three columns, the first containing the target genes, then the activators and finally the inhibitors. Target genes can be present in multiple lines. Several regulators can be present in column 2 and 3 if they are separated by a space. Below is an example of a toy network:

A:R1 R2 R3 R4
A:R5 R3 R6
B:R1 R2 R3 NA
...
Value

Returns a coregnet object.

Author(s)

Remy Nicolle <remy.c.nicolle AT gmail.com>

See Also

hLICORN

Examples

acts=apply(matrix(rep(letters[1:4],4),nrow=2),2,paste,collapse=" ")
reps=apply(matrix(rep(letters[5:8],4),nrow=2),2,paste,collapse=" ")
grn=data.frame("Target"= LETTERS[1:16] , "coact"=c(acts,reps),"corep"= c(reps,acts),"R2"=runif(16),stringsAsFactors=FALSE)
co=coregnet(grn)

---

coregnet-class

Class coregnet — Specifying the structure of the network used throughout the package.

Description

coregnet is an S4 type class which specifies the structure containing the necessary data of a Co-Regulatory network. coregnet objects are constructed by the hLICORN function which infers a regulatory network from gene expression data. A coregnet object can also be initialized by a network constructed by any other reverse engineering method as specified in the manual of the coregnet function.

Initialization

An object of type CoRegNet can be either built from a gene expression data set using the hLICORN function, using a data.frame containing a network specifying the activators and inhibitors of each gene or using an adjacency matrix (see the coregnet function).

Slots

GRN: A data.frame with three columns, Target Gene, co-activators and co-repressors. Co-regulators are separated by spaces.

adjacencyList: A description of the network through an adjacency matrix.

bytf: A named list of lists. The first entry point, the names of the list, are the Transcription Factors of the network. For each TF a list with two entries, act and rep, contains the set of activated and repressed genes respectively.

bygene: A named list of lists. The first entry point, the names of the list, are the target genes of the network. For each gene a list with two entries, act and rep, contains the set of activators and repressors of the genes respectively.

coregulators: A data.frame specifying the set of inferred co-regulators with several measures and statistics for each pairs.
evidences: A list containing all the added regulatory or co-regulatory evidences if any.
evidenceDescription: A list containing all the added regulatory or co-regulatory evidences if any.
inferenceParameters: A list of parameters of hLICORN.

Methods

Generic methods to be used with this class:

- **print**: Prints the number of genes, transcription factors and regulatory interactions.
- **summary**: Same as print
- **length**: Outputs the number of genes, transcription factors and regulatory interactions in the form of a numeric vector
- **dim**: Same as length
- **targets**: Returns the set of target genes or the targets of a given list of regulators
- **regulators**: Returns the set of regulators of the network or the set of regulators of a given list of genes.
- **activators**: Same as regulators but returns only activators of a gene
- **repressors**: Same as activators for repressor regulators
- **as.list**: Gets the network in the form of an adjacency list
- **as.data.frame**: Get the network in the form of an edge list
- **addEvidences**: Enriches the network with external regulation evidences
- **addCooperativeEvidences**: Enriches the network with external evidence of cooperative interaction between regulators
- **refine**: Refine the network based on a score which can take into account external regulation or co-regulation data.
- **regulatorInfluence**: Predicts the influence of the regulators in the network. Returns a predicted regulatory activity sample by sample
- **fitCoregnet**: Returns the fitness of the regulatory network given an expression dataset
- **display**: Starts a shiny application to display the co-regulation graph

Author(s)

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See Also

hLICORN, coregnet
**coregulators**

Extract all co-regulators of a regulatory network.

**Description**

Based on the frequency and specificity of co-regulation, this functions extracts from a coregnet network all the cooperative regulators.

**Usage**

```r
coregulators(object, maxcoreg = 2, verbose = TRUE, minCommonGenes = ifelse(maxcoreg == 2, 1, 10), adjustMethod = "fdr", alpha = 0.01)
```

**Arguments**

- `object`: The coregnet object containing the co-regulatory network
- `maxcoreg`: The maximum size of co-regulator sets to extract. The default is 2 to extract pairs of co-regulators
- `verbose`: To output information during the extraction process. By default does not output anything except warnings.
- `minCommonGenes`: The minimum number of common genes, co-regulated genes, to consider two regulators for further co-regulation analysis. Default is 1 for pairs and 10 for bigger sets.
- `adjustMethod`: The p-value adjustment method to extract significant pairs of co-regulators. Default is FDR correction. Anything that is accepted by the `p.adjust` method is fine.
- `alpha`: The threshold to consider a pair of co-regulator significant (after pvalue correction). Default is 1%

**Value**

For `maxcoreg` set to 2, a data.frame with pairs of regulators in the two first columns (`Reg1` and `Reg2`), the Support (the portion of coregulated gene networks in which this pair is found), `nGRN` for the number of times the pair was found as cooperative regulators in the net, a p-value of a FisherTest testing the specificity of the shared targets and the adjusted pvalue of this test.

For `maxcoreg` higher than 2, a two column data.frame with the set of co-regulators (collapsed in one character separated by a space) and the Support.

**Author(s)**

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**See Also**

`regulators`, `hLICORN`
**Examples**

```r
casts <- apply(rbind(rep("z", 14), matrix(rep(letters[1:4], 7), nrow = 2)), 2, paste, collapse = "")[1:13]
reps <- apply(matrix(rep(letters[5:8], 7), nrow = 2), 2, paste, collapse = "")[1:13]
grn <- data.frame("Target" = LETTERS[1:26], "coact" = c(casts, reps), "corep" = c(reps, casts), "R2" = runif(26), stringsAsFactors = FALSE)
co <- coregnet(grn)
coregulators(co)
coregulators(co, maxcoreg = 3, minCommonGenes = 3)
coregulators(co, adjustMethod = "bonferroni")
coregulators(co, alpha = 1)
```

---

**discretizeExpressionData**

Three-value discretization of gene expression data.

**Description**

Pre-process step to transform log2 numerical expression data into a three value categorical: over-expression (+1), under-expression (-1) and no change (0).

**Usage**

```r
discretizeExpressionData(numericalExpression, threshold = NULL, refSamples = NULL, standardDeviationThreshold = 1)
discretizeExpressionData(numericalExpression, threshold = NULL, refSamples = NULL, standardDeviationThreshold = 1)
```

**Arguments**

- **numericalExpression**
  
  A matrix of continuous log2 gene expression data with genes in rows and samples in column.

- **threshold**
  
  A numeric value used as a fixed fold change threshold for brut discretization. Can be NULL if a standardDeviationThreshold is not.

- **refSamples**
  
  A vector of column names used as a set of reference samples to be used to compute fold changes. Can be NULL if the input data is already centered on a reference values (tested by the presence of negative values) or if the mean if each gene should be used to center (not scale) each expression values.

- **standardDeviationThreshold**
  
  The multipicator of the whole data set standard deviation to be used as the threshold.
**discretizeExpressionData**

**Details**

Given a continuous log2 gene expression matrix this function aims at producing a matrix of discretized expression values. The numerical data must be in some form of fold change to compare each value of a gene in a sample with a reference value of the same gene. The behavior of the function will therefore depend on the form of the input data.

Given a matrix with negative values, the function will consider that the data is already in the right format and will simply apply a hard threshold to discretize the data.

Given a matrix with only positive values, which is the case for normalized RNAseq or single color microarrays, the function will center each gene based on it’s mean expression in all samples or based on the mean of expression of a set of reference sample (normal samples in a study of a particular disease for example).

In either case, the threshold will be used to transform the data in +1s if the value of a gene in a sample is above or equal to the threshold, -1s if the value is below the the negative value of the threshold and 0 otherwise.

The default is to compute a threshold based on the overall distribution of the numerical values in the dataset. This was choosen over a default fold change example (usually 1 or 2 corresponding to a two-fold or four-fold increase/decrease) after observing a large difference between technologies. However, the choice between a simple hard fold change threshold or a threshold as a multiplicator of the global standard deviation remains.

**Value**

A matrix of integers with the same number of rows and the same number of column as the input numericalExpression. Values in the output matrix are in -1,0,1. The reference samples are removed if given.

**Author(s)**

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**Examples**

```r
# Use mean of each gene as a reference
epression=matrix(2*runif(200),nrow=2,dimnames=list(paste("gene",1:2,sep=""),paste("sample",1:100,sep="")))
discExp=discretizeExpressionData(expression)
boxplot(expression~discExp,xlab="Discrete values",ylab="Continuous values")
pie(table(discExp))
discExp=discretizeExpressionData(expression,standardDeviationThreshold=2)
pie(table(discExp))
discExp=discretizeExpressionData(expression,threshold=1)
pie(table(discExp))

# Use of reference sample
epression=matrix(2*runif(200),nrow=2,dimnames=list(paste("gene",1:2,sep=""),paste("sample",1:100,sep="")))
discExp=discretizeExpressionData(expression,refSamples=1:10)
boxplot(expression~discExp,xlab="Discrete values",ylab="Continuous values")
pie(table(discExp))
discExp=discretizeExpressionData(expression,standardDeviationThreshold=2,refSamples=1:10)
pie(table(discExp))
discExp=discretizeExpressionData(expression,threshold=1,refSamples=paste("sample",1:10,sep=""))
pie(table(discExp))
```
Display

Description
Launches a shiny webpage for interactive viewing and analysis of the co-regulation network using a javascript cytoscape network.

Usage
display(coregnet, expressionData = NULL, TFA = NULL, alterationData = NULL, clinicalData = NULL, TFnotes = NULL, allTFplot = .heatplot, oneTFplot = .tfPlot)

Arguments

- **coregnet** A coregnet object
- **expressionData** A matrix or data.frame object with named columns (samples) and named rows (genes) containing gene expression data
- **TFA** A matrix or data.frame object with named columns (samples) and named rows (genes) containing transcription factor activity data such as the data obtained using the regulatorInfluence function. (unnecessary if expression data is given, makes the function run faster)
- **alterationData** optional. A matrix or data.frame object with named columns (samples) and named rows (genes) containing gene alteration data
- **clinicalData** optional. Either a list or a factor describing clinical information about samples. A list must be named and each entry should contain a vector of samples.
- **TFnotes** optional. A factor describing TFs.
- **allTFplot** A function ploting information by default. Default functions are implemented.
- **oneTFplot** A function ploting information about a TF which will be used when a single node is selected on the network. Default function are implemented.

Value
Does not return anything.

Author(s)
Remy Nicolle <remy.c.nicolle AT gmail.com>

Examples

```r
acts=apply(matrix(rep(letters[1:4],7),nrow=2),2,paste,collapse= " ")[1:13]
reps=apply(matrix(rep(letters[5:8],7),nrow=2),2,paste,collapse= " ")[1:13]
grn=data.frame("Target"= LETTERS[1:26] ,"coact"=c(acts,reps),"corep"= c(reps,acts),"R2"=runif(26),stringsAsFactors=FALSE)
co=coregnet(grn)
samples= paste("S",1:100,sep=" ")
expression=matrix(rnorm(3400),ncol=100)
dimnames(expression) = list(c(grn$Target,names(regulators(co))),samples)
TFA = regulatorInfluence(co,expression,minTarg=4)
colnames(TFA) = samples
```
if(interactive()){
  display(co,TFA=TFA,expressionData=expression)
}

CNA =matrix(sample(-2:2,800,replace=TRUE),ncol=100)
dimnames(CNA) = list(names(regulators(co)),samples)
if(interactive()){
  display(co,TFA=TFA,expressionData=expression,alteration=CNA)
}

clinicGrp = factor(paste("grp",sample(1:3,100,replace=TRUE),sep=""))
names(clinicGrp) = samples
if(interactive()){
  display(co,TFA=TFA,expressionData=expression,alteration=CNA,clinicalData=clinicGrp)
}

---

**hLICORN**

*Hybrid Learning of co-operative regulation network.*

**Description**

Parallelized inference of co-regulatory network from gene expression only.

**Usage**

```r
hLICORN(numericalExpression, discreteExpression =
  discretizeExpressionData(numericalExpression),
  TFlist, GeneList = setdiff(rownames(numericalExpression), TFlist),
  parallel = c("multicore", "no", "snow"), cluster = NULL,
  minGeneSupport=0.1,minCoregSupport = 0.1,
  maxCoreg=length(TFlist),
  searchThresh=1/3,nGRN=100,
  verbose=FALSE)
```

**Arguments**

- **numericalExpression**
  
  A numerical Matrix containing the expression of genes and of transcription factors that will be used to inferred the network. Rownames should contain the Gene names/identifiers. Samples should be in columns but Colnames are not important. The data will be gene-centered but not scaled.

- **discreteExpression**
  
  Optional. Should be in exactly the same format as numericalExpression (dimensions, colnames and rownames) and should contain value only in -1,0,1 with -1 for under-expressed, 0 for no change and 1 for over expressed. For default value see details.

- **TFlist**
  
  A character vector containing the names of the genes that are designated as Transcription Factor or Regulators. These should be contained in the rownames of the numericalExpression argument. Use data(HumanTF) for gene symbols of Human transcription factor.
optional. The list of genes for which Gene Regulatory Networks should be inferred. Should be in the rownames of the expression data. If not provided will be taken as all the genes in the rownames of the expression data that are not annotated as TF in TFlist.

parallel optional. The type of parallel method to use to run the process on several threads. Should be a value in "no" for single thread calculation, "multicore" to use several threads on the same local machine or "snow" to use the snow package in which case the cluster argument should contain a cluster object (e.g. makeCluster)

cluster A object of type cluster from the snow package which describes a cluster of computer to run the inference of each gene in parallel. Needed only when parallel is set to snow.

minGeneSupport A float between 0 and 1. Minimum number of samples in which a gene has to have non-zero discretized expression value to be considered for regulatory network inference.

minCoregSupport A float between 0 and 1. Minimum number of samples in which a set of co-regulators have the same non-zero discretized expression value to be considered as a potential set of co-regulator. Default is 0.1. Can be increased in case of limitations on memory.

maxCoreg A integer. Maximum size of co-regulator to consider. Default is set to the number of TF. Can be decreased in case of limitations on memory.

searchThresh A float between 0 and 1. Minimum proportion of sample in which a gene and a set of co-regulators must have the same non-zero discretized value in order to be considered as a potential co-regulator of the gene.

nGRN if NA, takes only the best GRN models.

verbose Sets whether information will appear in console during computation.

Details
A parallelized implementation of the h-LICORN algorithm. The inference of the network can be run in parallel using either a single machine or on a cluster using the make cluster in the parallel package (now a base R package).

The two required inputs are the continuous gene expression matrix and the list of transcription factor in the rows of the dataset. For Human, a list of TF is included as default data set using data(HumanTF) for symbols or data(HumanTF_entrezgene) for entrez gene IDs.

In case of memory overflow, a higher minCoregSupport (0.2 to 0.4) will perform more shallow search of co-regulators but will limit the quantity of memory used.

In some cases, in particular for very large datasets (several hundreds and thousands of samples), a lower minCoregSupport (down to 0.01) can be used for deep search.

Value
A object of type CoRegNet with gene regulatory network (GRN) containing several solution per gene and specifying the co-regulation interactions, the same network in the form of an adjacency list (adjacencyList) and the inferred co-regulators (coRegulators).

Note
The regulatory network inference process is done in several steps. The speed of the inference is dependant on several factors. The number of genes, the density of the discrete expression matrix (number of 1 and -1).
Author(s)

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References


See Also
discretizeExpressionData, coregnet-class

Examples

# Dummy expression data
gexp=matrix(rnorm(2600, sd=3), ncol=100)

gexp=rbind(gexp, do.call(rbind, lapply(1:26, function(i){
  tf = sample(1:26, 4)
  return((gexp[tf[1],]+gexp[tf[2],] - gexp[tf[3],]-gexp[tf[4],] + rnorm(100, sd=0.5))/2))))
dimnames(gexp)=list(c(letters, LETTERS), paste("s", 1:100, sep=""))

## Simple example of network inference
dummyNet=hLICORN(gexp, TFlist = letters)

## Infer a network only on a subset of genes
subgene = unique(dummyNet@GRN$Target)[1:2]
dummyNet=hLICORN(gexp, TFlist = letters, GeneList=subgene)

## Discretize data based on a set of reference samples (here 10 first)
disexp = discretizeExpressionData(gexp, refSamples=1:10)
dummyNet=hLICORN(gexp, TFlist = letters, discreteExpression=disexp)

## The network can be queried using the following functions
# returns the hub regulators
regulators(dummyNet)
# get the regulators of a given gene
regulators(dummyNet,"A")
activators(dummyNet,"A")
targets(dummyNet)
targets(dummyNet,"b")

# or transformed into a data.frame
coregnetToDataframe(dummyNet)
Human Data Examples

Human Transcription Factor data and Bladder cancer dataset.

Description

Internal datasets for the use of the package containing: A list of 2020 Human Transcription Factors in EntrezGene or Official Gene Symbol (HUGO) from the FANTOM consortium (Ravasi et al., 2010), A gene expression and copy number dataset of bladder cancer, Datasets of human TF protein interactions and regulatory interactions.

Usage

data(HumanTF)
data(HumanTF_entrezgene)
data(CIT_BLCA_EXP)
data(CIT_BLCA_CNV)
data(CIT_BLCA_Subgroup)
data(ENCODE_sub)
data(CHEA_sub)
data(STRING_sub)
data(HIPPIE_sub)

Format

HumanTF A character vector of Official (HUGO) gene symbols
HumanTF_entrezgene A character vector of EntrezGene
CIT_BLCA_EXP A matrix of bladder cancer gene expression from the CIT program. Only contains the 1000 genes with highest standard deviation. rownames : genes, colnames : samples. (from Rebouissou et al. 2014)
CIT_BLCA_CNV A matrix of bladder cancer DNA copy number aberrations from the CIT program. Only contains the TF and the tumor samples of CIT_BLCA_EXP. rownames : TF; colnames : samples.
CIT_BLCA_Subgroup A data.frame with sample name in first column and classification from the TCGA in the second column.
CIT_BLCA_smallGRN An example CoRegNet object.
ENCODE_sub A data.frame of regulatory interactions from the ENCODE project.
CHEA_sub A data.frame of regulatory interactions from the ChEA2 database. (http://amp.pharm.mssm.edu/ChEA2)
STRING_sub A data.frame of Protein interaction between TF from the STRING database. (http://string.embl.de)
HIPPIE_sub A data.frame of Protein interaction between TF from the HIPPIE database. (http://cbdm.mdc-berlin.de/tools/hippie)

References


Examples

data(HumanTF)
head(HumanTF)
data(HumanTF_entrezgene)
head(HumanTF_entrezgene)

---

**HumanTF**

*List of Human Transcription Factors.*

**Description**

A list of 2020 Human Transcription Factors in EntrezGene or Official Gene Symbol (HUGO) from the FANTOM consortium (Ravasi et al, 2010).

**Usage**

data(HumanTF)

**Format**

Two different objects:

- HumanTF_entrezgene: A character vector of EntrezGene
- HumanTF: A character vector of Official (HUGO) gene symbols

**References**


**Examples**

data(HumanTF)
head(HumanTF)
data(HumanTF_entrezgene)
head(HumanTF_entrezgene)
**masterRegulator**  
*Identify phenotype related Master Regulators.*

**Description**

This function implements methods to identify Master Regulators based on a regulatory network and on various type of input representing the implication of sets of gene in a phenotype of interest. These are derived from Celine Lefebvre’s algorithm MARINa (Lefebvre, 2010).

**Usage**

```r
masterRegulator(coregnet, targetGenes, method=c("set.overlap", "merge.pvalues", "list.enriched"))
```

**Arguments**

- `coregnet` A large scale co-regulatory network of type coregnet
- `targetGenes` A list of target genes. Can be either given as a character vector containing the genes or a named numeric vector containing weights (e.g. fold change) or p-values with gene as names.
- `method` The method to use to find the Master Regulators of the input target genes. The default is `set.overlap`. See details.

**Details**

Three types of input can be used depending on the objective.

To identify Master Regulators of a given set of genes (from a pathway, a list of differentially expressed genes etc ...), `MasterRegulatorInference` simply needs a CoRegNet network object and a character vector describing the Target Genes of interest. Fisher’s exact test will be used to identify TF with the set of target genes that is the most specific to these genes of interest.

To identify Master Regulators of a phenotype of interest, the p-values of the comparison with a reference phenotype (using a moderate or unmoderate t-test) or the Fold change can be used with a combined Fisher’s test or a Kolmogorov-Smirnov test to identify significant TF of these genes.

**Value**

The sorted list of TF in the input network with it’s associated p-value.

**Author(s)**

Remy Nicolle <remy.c.nicolle AT gmail.com>

**References**

A human B-cell interactome identifies MYB and FOXM1 as master regulators of proliferation in germinal centers. Molecular Systems Biology 6: 1-10
refine

### Examples

```r
# Dummy expression data and network
gexp = matrix(rnorm(2600, sd=3), ncol=100)
gexp = rbind(gexp, do.call(rbind, lapply(1:26, function(i) {
  tf = sample(1:26, 4)
  return((gexp[tf[1],] + gexp[tf[2],] - gexp[tf[3],] - gexp[tf[4],] + rnorm(100, sd=3))/2)))
})
dimnames(gexp) = list(c(letters, LETTERS), paste("s", 1:100, sep = ""))
GRN = hLICORN(gexp, TFlist = letters)

MR = masterRegulator(GRN, LETTERS[1:10])
head(MR)

eexampleWeight = rnorm(26)
names(exampleWeight) = LETTERS[1:26]
MR = masterRegulator(GRN, exampleWeight, "list")
head(MR)

eexamplePvalue = 10^(-0.1+runif(26))
names(examplePvalue) = LETTERS[1:26]
MR = masterRegulator(GRN, examplePvalue, "merg")
head(MR)
```

---

**refine**

Refine an inferred regulatory network using external evidence.

#### Description

Refine the inferred network using the integrated external evidences added by `addEvidences` or `addCooperativeEvidences`. Several strategies can be applied depending on the number and type of added evidences. These include supervised and unsupervised processes to use all the integrated data set with the inferred network and select the best Gene Regulatory Networks (GRN). Can also be used when no additional dataset has been integrated.

#### Usage

```r
refine(object, GRNselection = c("best", "maximize", "threshold"),
      integration = c("unsupervised", "supervised"),
      referenceEvidence = NULL, evidenceToMaximize = "R2", threshold = NULL, verbose = TRUE)
```

#### Arguments

- **object**
  A regulatory network inferred by the hLICORN function which can also have been enriched by external regulatory data sets using `addEvidences` or `addCooperativeEvidences`.

- **GRNselection**
  The type of Gene Regulatory Network (GRN) selection method to apply. Default is to select the best regulatory model per gene. See details for other possibilities.

- **integration**
  Defines the method to merge all the available data sets into a single score which will define the quality of a given GRN. Default is unsupervised. See details.
To be specified when using the supervised integration method. Specifies one of the integrated data set as a Gold standard to learn the best weight of each evidences to maximize the number of reference evidence in the final network.

evidenceToMaximize
To be specified when using the maximize GRN selection scheme. Instead of selecting one GRN per gene, this method will choose a threshold for the merged score that will maximize the number of interaction from a given evidence data set. When using the supervised integration method, the default is to use the referenceEvidence to be maximized.

threshold
When the automatically choosen threshold is not satisfactory, a user given threshold can be applied.

verbose
If set to TRUE (the default) sends messages at each step of the process.

Details
This function implements several strategies to select the best large scale regulatory network. Depending on the number and type of added evidences the strategies and some recommendations are detailed below.

The first step of the refinement is the integration of the different external evidences into a merged score. If no evidence data set has been added, the score given by the inference algorithm is used by its own (an adjusted R2). In the unsupervised method, the default, the merged score is simply the mean of each of the evidences, including the score given by hLICORN. For the supervised process, a weight is given to each of the evidences which is learned using a generalized linear model which will be fitted to predict the regulatory interactions of a user defined reference evidence data set.

These two (unsupervised or supervised) integration methods are derived from the network learning process used by the modENCODE consortium (Marbach et al, 2012). Once the merged score is obtained, the default is to select the best GRN per gene. However, another possibility is to select all the “good” networks by choosing a threshold on the merged score. This can be done either by a user defined threshold between 0 and 1 or by choosing automatically a threshold that will maximize the interactions originating from a user defined evidence data set.

The default behavior of the function is to integrate the data set in an unsupervised way and selecting the best GRN per gene.

It is recommanded that when no additional data has been integrated and the selection of the network only needs to be done based on the inference score, a bootstraped regression coefficient, then the simplest strategies is to use the default parameters which will select the best GRN per gene.

Value
A coRegNet object specifying a refined large scale co-regulatory network.

Author(s)
Remy Nicolle <remy.c.nicolle AT gmail.com>

References
regulatorInfluence

Regulator Influence, estimating the sample specific activity of Transcription Factors.

Description

Uses a network in the form of a coregnet object to compute regulatory influence to estimate the transcriptional activity of each regulator in each sample of the given expression data.

Usage

```r
regulatorInfluence(object, expData, minTarg = 10, withEvidences = FALSE, addCoregulators = FALSE, is.scaled = FALSE)
```

Arguments

- **object**: A network in the form of a coregnet object.
- **expData**: An expression data matrix or data.frame.
- **addCoregulators**: Compute influence for coregulators with sufficient number of targets. Default to FALSE.

See Also

addEvidences and addCooperativeEvidences

Examples

```r
# Dummy network and evidence data examples
acts = apply(rbind(rep("z", 14), matrix(rep(letters[1:4], 7), nrow = 2)), 2, paste, collapse = " ")[[1:13]]
reps = apply(matrix(rep(letters[5:8], 7), nrow = 2), 2, paste, collapse = " ")[[1:13]]
grn = data.frame("Target" = LETTERS[1:26], "coact" = c(acts, reps), "corep" = c(reps, acts), "R2" = runif(26), stringsAsFactors = FALSE)
GRN = coregnet(grn)
tfs = letters
genes = LETTERS
evidence1 = unique(data.frame(tf = sample(tfs, 100, replace = TRUE), target = sample(genes, 100, replace = TRUE), stringsAsFactors = FALSE))
evidence2 = unique(data.frame(tf = sample(tfs, 100, replace = TRUE), target = sample(genes, 100, replace = TRUE), stringsAsFactors = FALSE))
evidence3 = unique(data.frame(tf = sample(tfs, 100, replace = TRUE), target = sample(genes, 100, replace = TRUE), stringsAsFactors = FALSE))
GRNenrich = addEvidences(GRN, evidence1, evidence2, evidence3)
print(GRNenrich)

unsupervisedNet = refine(GRNenrich)
supervisedNet = refine(GRNenrich, integration = "sup", referenceEvidence = "evidence1")

# The following usually gives poor results...
supervisedNet = refine(GRNenrich, integration = "sup", referenceEvidence = "evidence1", evidenceToMaximize = "evidence1")
```
## regulators

The minimum number of targets for a regulator to be considered for activity prediction. Default set to 10.

- `withEvidences` Use only the target genes which are validated by an external validation dataset (ChIP-seq for example). This is only possible if external evidence was added using `addEvidences`. Default set to False.

- `is.scaled` Wether the input expression data is scaled, if not it will be.

**Value**

An N by R matrix with N columns the number of sample in the original expression data and R rows the number of regulators with sufficient targets to compute their influence. The expression data is centered by default but not scaled.

**Author(s)**

Remy Nicolle <remy.c.nicolle AT gmail.com>

**References**


**See Also**

- `hLICORN` and `coregnet-class` to create the network. `addEvidences` to add external evidences.

**Examples**

```r
acts=apply(matrix(rep(letters[1:4],7),nrow=2),2,paste,collapse=" ")[1:13]
reps=apply(matrix(rep(letters[5:8],7),nrow=2),2,paste,collapse=" ")[1:13]
grn=data.frame("Target"= LETTERS[1:26] ,"coact"=c(acts,reps),"corep"= c(reps,acts),"R2"=runif(26),stringsAsFactors=FALSE)
co=coregnet(grn)
samples= paste("S",1:100,sep="")
extpression=matrix(rnorm(3400),ncol=100)
dimnames(expression) = list(c(grn$Target,names(regulators(co))),samples)
#Minimum number of targets is adjusted because of the small size of the network
TFA = regulatorInfluence(co,expression,minTarg=4)
```

---

### regulators

**Interrogate a coregnet object.**

**Description**

Query the network for regulators of specific targets and targets of specific genes.

**Usage**

```r
regulators(object, target = NULL, type = c("single", "coregulators"))
activators(object, target, type=c("single", "coregulators"))
repressors(object, target, type=c("single", "coregulators"))
targets(object, regulator=NULL, type=c("regulating","activating","repressing"))
```
regulators

Arguments

object  The network in the form of a coregnet object to query.
target The target gene to query.
regulator The regulator to query.
type The type of regulation to obtain. Differs depending on the function used.

Value

For regulators if no target is given, returns integer vector with the number of targets for each regulators of the network. Given a non null vector, a vector of the union of the regulators of all the genes is returned. For activators and repressors the behavior is similar except that a target gene is needed. If type = "coregulators" then only the regulators, activators or repressors which are found to be co-regulators, co-activators or co-repressors of the target genes are given.

targets with no given regulator returns a character vector of all the target genes in the network. Specifying a vector of regulators will return a vector of the union of the targets of all these regulators. The type of regulation can be specified to return only the activated or repressed targets.

Author(s)

Remy Nicolle <remy.c.nicolle AT gmail.com>

Examples

acts=apply(matrix(rep(letters[1:4],7),nrow=2),2,paste,collapse=" ")[1:13]
reps=apply(matrix(rep(letters[5:8],7),nrow=2),2,paste,collapse=" ")[1:13]
grn=data.frame("Target"= LETTERS[1:26], "coact"=c(acts,reps),"corep"= rep(reps,acts),"R2"=runif(26),stringsAsFactors=FALSE)
c=coregnet(grn)
regulators(co)
regulators(co,"A")
regulators(co,"A","coregulators")

activators(co,"A")
activators(co,"A","coregulators")

repressors(co,"A")
repressors(co,"A","coregulators")

targets(co)
targets(co,"a")
targets(co,"a","reg")
targets(co,"a","act")
targets(co,"a","rep")
targets(co, c("a","b"),"act")
Description

Several functions to print and view info about the network enclosed in a coregnet object.

Usage

```r
summary(object,...)
## S4 method for signature 'coregnet'
show(object)
## S4 method for signature 'coregnet'
dim(x)
## S4 method for signature 'coregnet'
length(x)
## S4 method for signature 'coregnet'
print(x)
## S4 method for signature 'coregnet'
coregnetToDataFrame(network)
## S4 method for signature 'coregnet'
coregnetToList(network)
```

Arguments

- `network`: a coregnet network object.
- `object`: a coregnet network object.
- `x`: a coregnet network object.
- `...`: unused argument

Author(s)

Remy Nicolle <remy.c.nicolle AT gmail.com>

Examples

```r
regs=sample(letters,7)
grn=data.frame("Target"= LETTERS ,"activators"= sample(rep(regs,4))[1:26],"repressors"= sample(rep(regs,4))[1:26],stringsAsFactors=FALSE)
co=coregnet(grn)
print(co)
length(co)
dim(co)
co
coregnetToDataFrame(co)
coregnetToList(co)
```
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