

# Package ‘netresponse’

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**Type** Package

**Title** Functional Network Analysis

**Version** 1.32.2

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**Description** Algorithms for functional network analysis. Includes an implementation of a variational Dirichlet process Gaussian mixture model for nonparametric mixture modeling.

**License** GPL (>=2)

**Depends** R (>= 2.15.1), Rgraphviz, methods, minet, mclust, reshape2

**Imports** dmt, ggplot2, graph, igraph, parallel, plyr, qvalue, RColorBrewer

**URL** <https://github.com/antagomir/netresponse>

**BugReports** <https://github.com/antagomir/netresponse/issues>

**biocViews** CellBiology, Clustering, GeneExpression, Genetics, Network, GraphAndNetwork, DifferentialExpression, Microarray, Transcription

**RoxygenNote** 5.0.1

**NeedsCompilation** yes

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---

netresponse-package    *NetResponse: Global modeling of transcriptional responses in interaction networks*

---

## Description

Global modeling of transcriptional responses in interaction networks.

Package:    netresponse  
Type:        Package  
Version:    See sessionInfo() or DESCRIPTION file  
Date:        2011-02-03  
License:    GNU GPL >=2  
LazyLoad:   yes

## Author(s)

Leo Lahti, Olli-Pekka Huovilainen, Antonio Gusmao and Juuso Parkkinen. Maintainer: Leo Lahti <leo.lahti@iki.fi>

## References

Leo Lahti et al.: Global modeling of transcriptional responses in interaction networks. *Bioinformatics* (2010). See citation("netresponse") for details.

## Examples

```
# Define parameters for toy data
Ns <- 200 # number of samples (conditions)
Nf <- 10  # number of features (nodes)
feature.names <- paste("feat", seq(Nf), sep="")
sample.names <- paste("sample", seq(Ns), sep="")
# random seed
set.seed( 123 )
# Random network
netw <- pmax(array(sign(rnorm(Nf^2))), dim = c(Nf, Nf)), 0)
# in pathway analysis nodes correspond to genes
rownames(netw) <- colnames(netw) <- feature.names
# Random responses of the nodes across conditions
D <- array(rnorm(Ns*Nf), dim = c(Ns,Nf), dimnames = list(sample.names, feature.names))
D[1:100, 4:6] <- t(sapply(1:(Ns/2),function(x){rnorm(3, mean = 1:3)}))
D[101:Ns, 4:6] <- t(sapply(1:(Ns/2),function(x){rnorm(3, mean = 7:9)}))
# Calculate the model
model <- detect.responses(D, netw)
# Subnets (each is a list of nodes)
get.subnets( model )
```

```
# Retrieve model for one subnetwork
# means, standard deviations and weights for the components
inds <- which(sapply(model@last.grouping, length) > 2)
subnet.id <- names(model@subnets)[[1]]
m <- get.model.parameters(model, subnet.id)
print(m)
```

---

add.ellipse

*Add ellipse to an existing plot*

---

### Description

Calculates and plots ellipse corresponding to specified confidence interval in 2-dimensional plot

### Usage

```
add.ellipse(centroid, covmat, confidence = 0.95, npoints = 100, col =
"black", ...)
```

### Arguments

centroid	Vector with two elements defining the ellipse centroid.
covmat	Covariance matrix for the investigated data. Only diagonal covariances supported.
confidence	Confidence level determining the ellipse borders based on the covariance matrix.
npoints	Number of plotting points.
col	Color.
...	Other arguments to be passed.

### Value

Used for plotting side effects.

### Author(s)

Leo Lahti <leo.lahti@iki.fi>

### Examples

```
#add.ellipse(centroid = c(0, 0), covmat = diag(c(1,2)))
```

---

bic.mixture	<i>BIC mixture</i>
-------------	--------------------

---

**Description**

Latent class analysis based on (infinite) Gaussian mixture model. If the input is data matrix, a multivariate model is fitted; if the input is a vector, a univariate model is fitted

**Usage**

```
bic.mixture(x, max.modes, bic.threshold = 0, min.modes = 1, ...)
```

**Arguments**

x	samples x features matrix for multivariate analysis, or a vector for univariate analysis
max.modes	Maximum number of modes to be checked for mixture model selection
bic.threshold	BIC threshold which needs to be exceeded before a new mode is added to the mixture.
min.modes	minimum number of modes
...	Further optional arguments to be passed

**Value**

Fitted latent class model (parameters and free energy)

**Author(s)**

Contact: Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse")

---

bic.mixture.multivariate	<i>Multivariate BIC mixture</i>
--------------------------	---------------------------------

---

**Description**

Latent class analysis based on (infinite) Gaussian mixture model. If the input (dat) is data matrix, a multivariate model is fitted.

**Usage**

```
bic.mixture.multivariate(x, max.modes, bic.threshold = 0, min.modes = 1,
  ...)
```

**Arguments**

x	matrix (for multivariate analysis)
max.modes	Maximum number of modes to be checked for mixture model selection
bic.threshold	BIC threshold which needs to be exceeded before a new mode is added to the mixture.
min.modes	Minimum number of modes to be checked for mixture model selection
...	Further optional arguments to be passed

**Value**

Fitted latent class model (parameters and free energy)

**Author(s)**

Contact: Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse")

---

bic.mixture.univariate

*Univariate BIC mixture*

---

**Description**

Latent class analysis based on (infinite) Gaussian mixture model. If the input (dat) is data matrix, a multivariate model is fitted. If the input is a vector or a 1-dimensional matrix, a univariate model is fitted.

**Usage**

```
bic.mixture.univariate(x, max.modes, bic.threshold = 0, min.modes = 1, ...)
```

**Arguments**

x	dat vector (for univariate analysis) or a matrix (for multivariate analysis)
max.modes	Maximum number of modes to be checked for mixture model selection
bic.threshold	BIC threshold which needs to be exceeded before a new mode is added to the mixture.
min.modes	minimum number of modes
...	Further optional arguments to be passed

**Value**

Fitted latent class model (parameters and free energy)

**Author(s)**

Contact: Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse")

---

`bic.select.best.mode` *Select best mode with BIC*

---

**Description**

Select optimal number of mixture components by adding components until the increase in objective function is below threshold.

**Usage**

```
bic.select.best.mode(x, max.modes, bic.threshold, min.modes = 1)
```

**Arguments**

<code>x</code>	dat vector (for univariate analysis) or a matrix (for multivariate analysis)
<code>max.modes</code>	Maximum number of modes to be checked for mixture model selection
<code>bic.threshold</code>	BIC threshold which needs to be exceeded before a new mode is added to the mixture.
<code>min.modes</code>	Optional. Minimum number of modes.

**Value**

Fitted latent class model (parameters and free energy)

**Author(s)**

Contact: Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse")

---

centerData	<i>Center data matrix.</i>
------------	----------------------------

---

**Description**

Center data matrix to 0 for each variable by removing the means.

**Usage**

```
centerData(X, rm.na = TRUE, meanvalue = NULL)
```

**Arguments**

X	The data set: samples x features. Each feature will be centered.
rm.na	Ignore NAs.
meanvalue	Can be used to set a desired center value. The default is 0.

**Value**

Centered data matrix.

**Note**

Note that the model assumes samples x features matrix, and centers each feature.

**Author(s)**

Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse").

**Examples**

```
#centerData(X)
```



---

continuous.responses    *Continuous responses*

---

### Description

Quantify association between modes and continuous variable

### Usage

```
continuous.responses(annotation.vector, model, method = "t-test",
  min.size = 2, data = NULL)
```

### Arguments

annotation.vector	annotation vector with discrete factor levels, and named by the samples
model	NetResponse model object
method	method for enrichment calculation
min.size	minimum sample size for a response
data	data matrix (samples x features)

### Value

List with each element corresponding to one variable and listing the responses according to association strength

### Author(s)

Contact: Leo Lahti <leo.lahti@iki.fi>

### References

See citation("netresponse")

---

detect.responses    *detect.responses*

---

### Description

Main function of the NetResponse algorithm. Detect condition-specific network responses, given network and a set of measurements of node activity in a set of conditions. Returns a set of subnetworks and their estimated context-specific responses.

**Usage**

```
detect.responses(datamatrix, network = NULL, initial.responses = 1,
  max.responses = 10, max.subnet.size = 10, verbose = TRUE,
  prior.alpha = 1, prior.alphaKsi = 0.01, prior.betaKsi = 0.01,
  update.hyperparams = 0, implicit.noise = 0, vdp.threshold = 1e-05,
  merging.threshold = 0, ite = Inf, information.criterion = "BIC",
  speedup = TRUE, speedup.max.edges = 10, positive.edges = FALSE,
  mc.cores = 1, mixture.method = "vdp", bic.threshold = 0,
  pca.basis = FALSE, ...)
```

**Arguments**

<code>datamatrix</code>	Matrix of samples x features. For example, gene expression matrix with conditions on the rows, and genes on the columns. The matrix contains same features than the 'network' object, characterizing the network states across the different samples.
<code>network</code>	Binary network describing undirected pairwise interactions between features of 'datamatrix'. The following formats are supported: binary matrix, graphNEL, igraph, graphAM, Matrix, dgCMatrix, dgeMatrix
<code>initial.responses</code>	Initial number of components for each subnetwork model. Used to initialize calculations.
<code>max.responses</code>	Maximum number of responses for each subnetwork. Can be used to limit the potential number of network states.
<code>max.subnet.size</code>	Numeric. Maximum allowed subnetwork size.
<code>verbose</code>	Logical. Verbose parameter.
<code>prior.alpha</code> , <code>prior.alphaKsi</code> , <code>prior.betaKsi</code>	Prior parameters for Gaussian mixture model that is calculated for each subnetwork (normal-inverse-Gamma prior). <code>alpha</code> tunes the mean; <code>alphaKsi</code> and <code>betaKsi</code> are the shape and scale parameters of the inverse Gamma function, respectively.
<code>update.hyperparams</code>	Logical. Indicate whether to update hyperparameters during modeling.
<code>implicit.noise</code>	Implicit noise parameter. Add implicit noise to vdp mixture model. Can help to avoid overfitting to local optima, if this appears to be a problem.
<code>vdp.threshold</code>	Minimal free energy improvement after which the variational Gaussian mixture algorithm is deemed converged.
<code>merging.threshold</code>	Minimal cost value improvement required for merging two subnetworks.
<code>ite</code>	Defines maximum number of iterations on posterior update ( <code>updatePosterior</code> ). Increasing this can potentially lead to more accurate results, but computation may take longer.
<code>information.criterion</code>	Information criterion for model selection. Default is BIC (Bayesian Information Criterion); other options include AIC and AICc.

speedup	Takes advantage of approximations to PCA, mutual information etc in various places to speed up calculations. Particularly useful with large and densely connected networks and/or large sample size.
speedup.max.edges	Used if speedup = TRUE. Applies prefiltering of edges for calculating new joint models between subnetwork pairs when potential cost changes (delta) are updated for a newly merged subnetwork and its neighborghs. Empirical mutual information between each such subnetwork pair is calculated based on their first principal components, and joint models will be calculated only for the top candidates up to the number specified by speedup.max.edges. It is expected that the subnetwork pair that will benefit most from joint modeling will be among the top mutual infomation candidates. This way it is possible to avoid calculating exhaustive many models on the network hubs.
positive.edges	Consider only the edges with positive association. Currently measured with Spearman correlation.
mc.cores	Number of cores to be used in parallelization. See help(mclapply) for details.
mixture.method	Specify the approach to use in mixture modeling. Options. vdp (nonparametric Variational Dirichlet process mixture model); bic (based on Gaussian mixture modeling with EM, using BIC to select the optimal number of components)
bic.threshold	BIC threshold which needs to be exceeded before a new mode is added to the mixture with mixture.method = "bic"
pca.basis	Transform data first onto PCA basis to try to avoid problems with non-diagonal covariances.
...	Further optional arguments to be passed.

**Value**

NetResponseModel object.

**Author(s)**

Maintainer: Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse").

**Examples**

```
data(toydata)          # Load toy data set
D <- toydata$emat      # Response matrix (for example, gene expression)
netw <- toydata$netw   # Network

# Run NetReponse algorithm
## Not run: model <- detect.responses(D, netw, verbose = FALSE)
```

---

dna

*Dna damage data set (PPI and expression)*

---

### Description

A combined yeast data set with protein-protein interactions and gene expression (dna damage). Gene expression profiles are transformed into links by computing a Pearson correlation for all pairs of genes and treating all correlations above 0.85 as additional links. Number of genes: 1823, number of interactions: 12382, number of gene expression observations: 52, number of total links with PPI and expression links: 15547.

### Usage

data(dna)

### Format

List of following objects:

**ppi** PPI data matrix

**exp** gene expression profiles data matrix

**gids** Vector of gene ids corresponding to indices used in data matrices

**obs** Gene expression observation details

**combined.links** pooled matrix of PPI and expression links

### Source

PPI data pooled from yeast data sets of [1] and [2]. Dna damage expression set of [3].

### References

Ulitsky, I. and Shamir, R. *Identification of functional modules using network topology and high-throughput data*. BMC Systems Biology 2007, 1:8.

Nariai, N., Kolaczyk, E. D. and Kasif, S. *Probabilistic Protein Function Prediction from Heterogeneous Genome-Wide Data*. PLoS ONE 2007, 2(3):e337.

Gasch, A., Huang, M., Metzner, S., Botstein, D. and Elledge, S. *Genomic expression responses to DNA-damaging agents and the regulatory role of the yeast ATR homolog Mex1p*. Molecular Biology of the Cell 2001, 12:2987-3003.

### Examples

data(dna)

---

`enrichment.list.factor`*enrichment.list.factor*

---

**Description**

Orders the responses by association strength (enrichment score) to a given sample set. For instance, if the samples correspond to a particular experimental factor, this function can be used to prioritize the responses according to their association strength to this factor.

**Usage**

```
enrichment.list.factor(models, level.samples, method, verbose = FALSE)
```

**Arguments**

<code>models</code>	List of models. Each model should have a sample-cluster assignment matrix <code>qofz</code> .
<code>level.samples</code>	Measure enrichment of this sample (set) across the observed responses.
<code>method</code>	'hypergeometric' measures enrichment of factor levels in this response; 'precision' measures response purity for each factor level; 'dependency' measures log-arithm of the joint density between response and factor level vs. their marginal densities: $\log(P(r,s)/(P(r)P(s)))$
<code>verbose</code>	Follow progress by intermediate messages.

**Value**

A data frame which gives a data frame of responses ordered by enrichment score for the investigated sample. The model, response id and enrichment score are shown. The method field indicates the enrichment calculation method. The sample field lists the samples `et` for which the enrichments were calculated. The info field lists additional information on enrichment statistics.

**Author(s)**

Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse") for citation details.

**Examples**

```
#
```

---

```
enrichment.list.factor.minimal
      enrichment.list.factor
```

---

**Description**

Orders the responses by association strength (enrichment score) to a given sample set. For instance, if the samples correspond to a particular experimental factor, this function can be used to prioritize the responses according to their association strength to this factor.

**Usage**

```
enrichment.list.factor.minimal(groupings, method, verbose = FALSE,
  annotation.vector, level)
```

**Arguments**

groupings	List of groupings. Each model should have a sample-cluster assignment matrix qofz.
method	'hypergeometric' measures enrichment of factor levels in this response; 'precision' measures response purity for each factor level; 'dependency' measures logarithm of the joint density between response and factor level vs. their marginal densities: $\log(P(r,s)/(P(r)P(s)))$
verbose	Follow progress by intermediate messages.
annotation.vector	annotation vector
level	level

**Value**

A data frame which gives a data frame of responses ordered by enrichment score for the investigated sample. The model, response id and enrichment score are shown. The method field indicates the enrichment calculation method. The sample field lists the samples et for which the enrichments were calculated. The info field lists additional information on enrichment statistics.

**Author(s)**

Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse") for citation details.

**Examples**

```
#
```

---

factor.responses	<i>Factor responses</i>
------------------	-------------------------

---

**Description**

List responses for each level of the given factor

**Usage**

```
factor.responses(annotation.vector, groupings, method = "hypergeometric",  
  min.size = 2, data = NULL)
```

**Arguments**

annotation.vector	annotation vector with discrete factor levels, and named by the samples
groupings	List of groupings. Each model should have a sample-cluster assignment matrix qofz, or a vector of cluster indices named by the samples.
method	method for enrichment calculation
min.size	minimum sample size for a response
data	data (samples x features; or a vector in univariate case)

**Value**

List with each element corresponding to one factor level and listing the responses according to association strength

**Author(s)**

Contact: Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse")

---

factor.responses.minimal  
*Factor responses (minimal)*

---

### Description

List responses for each level of the given factor

### Usage

```
factor.responses.minimal(annotation.vector, groupings,  
  method = "hypergeometric", min.size = 2, data = NULL)
```

### Arguments

annotation.vector	annotation vector with discrete factor levels, and named by the samples
groupings	List of groupings. Each model should have a sample-cluster assignment matrix qofz, or a vector of cluster indices named by the samples.
method	method for enrichment calculation
min.size	minimum sample size for a response
data	data (samples x features; or a vector in univariate case)

### Value

List with each element corresponding to one factor level and listing the responses according to association strength

### Author(s)

Contact: Leo Lahti <leo.lahti@iki.fi>

### References

See citation("netresponse")



---

find.similar.features *Find similar features with a given subnetwork.*

---

### Description

Given subnetwork, orders the remaining features (genes) in the input data based on similarity with the subnetwork. Allows the identification of similar features that are not directly connected in the input network.

### Usage

```
find.similar.features(model, subnet.id, datamatrix = NULL, verbose =  
FALSE, information.criterion = NULL)
```

### Arguments

model	NetResponseModel object.
subnet.id	Investigated subnetwork.
datamatrix	Optional. Can be used to compare subnetwork similarity with new data which was not used for learning the subnetworks.
verbose	Logical indicating whether progress of the algorithm should be indicated on the screen.
information.criterion	Information criterion for model selection. By default uses the same than in the 'model' object.

### Details

The same similarity measure is used as when agglomerating the subnetworks: the features are ordered by delta (change) in the cost function, assuming that the feature would be merged in the subnetwork. The smaller the change, the more similar the feature is (change would minimize the new cost function value). Negative values of delta mean that the cost function would be improved by merging the new feature in the subnetwork, indicating features having coordinated response.

### Value

A data frame with elements feature.names (e.g. gene IDs) and delta, which indicates similarity level. See details for details. The smaller, the more similar. The data frame is ordered such that the features are listed by decreasing similarity.

### Author(s)

Leo Lahti <leo.lahti@iki.fi>

### References

See citation("netresponse") for reference details.

## Examples

```
data(toydata)
model <- toydata$model
subnet.id <- "Subnet-1"
g <- find.similar.features(model, subnet.id)
# List features that are similar to this subnetwork (delta < 0)
# (ordered by decreasing similarity)
subset(g, delta < 0)
```

---

get.dat,NetResponseModel-method

*Get subnetwork data*

---

## Description

Get subnetwork data

## Usage

```
## S4 method for signature 'NetResponseModel'
get.dat(model, subnet.id, sample = NULL)
```

## Arguments

model	Result from NetResponse (detect.responses function).
subnet.id	Subnet identifier. A natural number which specifies one of the subnetworks within the 'model' object.
sample	Define the retrieved samples

## Value

Subnet data matrix

## Author(s)

Leo Lahti <leo.lahti@iki.fi>

## References

See citation("netresponse")

## Examples

```
## Load a pre-calculated netresponse model obtained with
# model <- detect.responses(toydata$emat, toydata$netw, verbose = FALSE)
# data( toydata ); get.dat(toydata$model)
```

---

get.model.parameters    *get.model.parameters*

---

## Description

Retrieve the mixture model parameters of the NetResponse algorithm for a given subnetwork.

## Usage

```
get.model.parameters(model, subnet.id = NULL)
```

## Arguments

model	Result from NetResponse (detect.responses function).
subnet.id	Subnet identifier. A natural number which specifies one of the subnetworks within the 'model' object.

## Details

Only the non-empty components are returned. Note: the original data matrix needs to be provided for function call separately.

## Value

A list with the following elements:

mu	Centroids for the mixture components. Components x nodes.
sd	Standard deviations for the mixture components. A vector over the nodes for each component, implying the diagonal covariance matrix of the model (i.e. $\text{diag}(\text{std}^2)$ ). Components x nodes
w	Vector of component weights.
nodes	List of nodes in the subnetwork.
K	Number of mixture components.

## Author(s)

Leo Lahti <leo.lahti@iki.fi>

## References

Leo Lahti et al.: Global modeling of transcriptional responses in interaction networks. Bioinformatics (2010). See citation("netresponse") for details.

**Examples**

```
# Load toy data
data( toydata )           # Load toy data set
D     <- toydata$emat     # Response matrix (for example, gene expression)
model <- toydata$model    # Pre-calculated model

# Get model parameters for a given subnet
# (Gaussian mixture: mean, covariance diagonal, mixture proportions)
get.model.parameters(model, subnet.id = 1)
```

---

```
get.subnets,NetResponseModel-method
      get.subnets
```

---

**Description**

List the detected subnetworks (each is a list of nodes in the corresponding subnetwork).

**Usage**

```
## S4 method for signature 'NetResponseModel'
get.subnets(model, get.names = TRUE,
             min.size = 2, max.size = Inf, min.responses = 2)
```

**Arguments**

<code>model</code>	Output from the <code>detect.responses</code> function. An object of <code>NetResponseModel</code> class.
<code>get.names</code>	Logical. Indicate whether to return subnetwork nodes using node names (TRUE) or node indices (FALSE).
<code>min.size, max.size</code>	Numeric. Filter out subnetworks whose size is not within the limits specified here.
<code>min.responses</code>	Numeric. Filter out subnetworks with less responses (mixture components) than specified here.

**Value**

A list of subnetworks.

**Author(s)**

Leo Lahti <leo.lahti@iki.fi>

## References

Leo Lahti et al.: Global modeling of transcriptional responses in interaction networks. *Bioinformatics* (2010). See citation("netresponse") for details.

## Examples

```
## Load a pre-calculated netresponse model obtained with
# model <- detect.responses(toydata$emat, toydata$netw, verbose = FALSE)
# data( toydata ); get.subnets(toydata$model)
```

---

ICMg.combined.sampler *ICMg.combined.sampler*

---

## Description

Main function of the ICMg algorithm. ICMg.combined.sampler computes samples from the posterior of the assignments of datapoints (interactions and expression profiles) to latent components. From these we can then obtain component membership distributions and clusterings for genes.

## Usage

```
ICMg.combined.sampler(L, X, C, alpha = 10, beta = 0.01, pm0 = 0, V0 = 1,
  V = 0.1, B.num = 8, B.size = 100, S.num = 20, S.size = 10,
  C.boost = 1)
```

## Arguments

L	N x 2 matrix of link endpoints (N = number of links).
X	M x D matrix of gene expression profiles (M = number of nodes, D = number of observations).
C	Number of components.
alpha	Hyperparameter describing the global distribution over components, larger alpha gives a more uniform distribution.
beta	Hyperparameter describing the component-wise distributions over nodes, larger beta gives a more uniform distribution.
pm0	Hyperparameter describing the prior mean of the expression profiles, should be zero.
V0	Hyperparameter describing the variation of the component-wise expression profiles means around pm0.
V	Hyperparameter describing the variation of gene-specific expression profiles around the component-wise means.
B.num	Number of burnin rounds.*
B.size	Size of one burnin round.*
S.num	Number of sample rounds.*
S.size	Size of one sample round.*
C.boost	Set to 1 to use faster iteration with C, set to 0 to use slower R functions.

**Details**

One run consists of two parts, during burnin the sampler is expected to mix, after which the samples are taken. Information about convergence (convN and convL are estimates of convergence for link and node sampling, respectively) and component sizes are printed after each burnin/sample round. For example: B.num=8, B.size=100, S.num=20, S.size=10, runs 800 burnin iterations in 8 rounds and then takes 20 samples with an interval of 10 iterations.

**Value**

Returns samples as a list:

z	S.num x N matrix of samples of component assignments for links.
w	S.num x M matrix of samples of component assignments for gene expression profiles.
convl	Vector of length (B.num + S.num) with convergence estimator values for link sampling.
convn	Vector of length (B.num + S.num) with convergence estimator values for node sampling.
countsl	(B.num + S.num) x C matrix of link component sizes.
countsn	(B.num + S.num) x C matrix of node component sizes.

additionally all parameters of the run are included in the list.

**Author(s)**

Juuso Parkkinen

**References**

Parkkinen, J. and Kaski, S. Searching for functional gene modules with interaction component models. BMC Systems Biology 4 (2010), 4.

**See Also**

[ICMg.links.sampler](#)

**Examples**

```
data(osmo) # Load data set
res <- ICMg.combined.sampler(osmo$ppi, osmo$exp, C=10)
```

---

ICMg.get.comp.memberships  
*ICMg.get.comp.memberships*

---

**Description**

Function for computing the component memberships for each data point from the MCMC samples.

**Usage**

```
ICMg.get.comp.memberships(links, samples)
```

**Arguments**

links	N x 2 matrix of link endpoints (N = number of links).
samples	Posterior samples, as given by either ICMg.combined.sampler or ICMg.links.sampler.

**Value**

A matrix containing the component memberships for each data point (node).

**Author(s)**

Juuso Parkkinen

**References**

Parkkinen, J. and Kaski, S. Searching for functional gene modules with interaction component models. BMC Systems Biology 4 (2010), 4.

**See Also**

[ICMg.combined.sampler](#), [ICMg.links.sampler](#)

---

ICMg.links.sampler     *ICMg.links.sampler*

---

**Description**

ICMg.links.sampler computes samples from the posterior of the assignments of datapoints (interactions) to latent components. From these we can then obtain component membership distributions and clusterings for genes.

**Usage**

```
ICMg.links.sampler(L, C, alpha=10, beta=0.01, B.num=8, B.size=100,  
S.num=20, S.size=10, C.boost=1)
```

**Arguments**

L	N x 2 matrix of link endpoints (N = number of links).
C	Number of components.
alpha	Hyperparameter describing the global distribution over components, larger alpha gives a more uniform distribution.
beta	Hyperparameter describing the component-wise distributions over nodes, larger beta gives a more uniform distribution.
B.num	Number of burnin rounds.*
B.size	Size of one burnin round.*
S.num	Number of sample rounds.*
S.size	Size of one sample round.*
C.boost	Set to 1 to use faster iteration with C, set to 0 to use slower R functions.

**Details**

One run consists of two parts, during burnin the sampler is expected to mix, after which the samples are taken. Information about convergence (convN and convL are estimates of convergence for link and node sampling, respectively) and component sizes are printed after each burnin/sample round. For example: B.num=8, B.size=100, S.num=20, S.size=10, runs 800 burnin iterations in 8 rounds and then takes 20 samples with an interval of 10 iterations.

**Value**

Returns samples as a list:

z	S.num x N matrix of samples of component assignments for links.
conv	Vector of length (B.num + S.num) with convergence estimator values for link sampling.
counts	(B.num + S.num) x C matrix of link component sizes.

additionally all parameters of the run are included in the list.

**Author(s)**

Juuso Parkkinen

**References**

Parkkinen, J. and Kaski, S. Searching for functional gene modules with interaction component models. BMC Systems Biology 4 (2010), 4.

**See Also**

[ICMg.combined.sampler](#)



**Examples**

```
data(osmo) # Load data

## Run ICMg links sampler
res <- ICMg.links.sampler(osmo$ppi, C=10)
```

---

```
list.responses.continuous.multi
```

*Investigate association of a continuous variable and the modes*

---

**Description**

Investigate association of a continuous variable and the modes given a list of groupings

**Usage**

```
list.responses.continuous.multi(annotation.df, groupings, method = "t-test",
  pth = Inf, verbose = TRUE, rounding = NULL)
```

**Arguments**

annotation.df	annotation data.frame with discrete factor levels, rows named by the samples
groupings	Sample mode information. Each element corresponds to one grouping; each grouping lists samples for the modes within that grouping.
method	method for quantifying the association
pth	p-value threshold applied to adjusted p-values
verbose	verbose
rounding	rounding digits

**Value**

Table listing all associations between the factor levels and responses

**Author(s)**

Contact: Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse")

```
list.responses.continuous.single
```

*Investigate association of a continuous variable and the modes*

---

### Description

Investigate association of a continuous variable and the modes.

### Usage

```
list.responses.continuous.single(annotation.df, groupings, method = "t-test",  
  pth = Inf, verbose = TRUE, rounding = NULL, adjust.p = TRUE)
```

### Arguments

annotation.df	annotation data.frame with discrete factor levels, rows named by the samples
groupings	Sample mode information. Each element corresponds to one of the modes and lists the samples assignment matrix qofz. Alternatively, a vector of mode indices named by the samples can be given.
method	method for quantifying the association
pth	p-value threshold (for adjusted p-values)
verbose	verbose
rounding	rounding digits
adjust.p	Adjust p-values (this will add p.adj column and remove pvalue column in the output table)

### Value

Table listing all associations between the factor levels and responses

### Author(s)

Contact: Leo Lahti <leo.lahti@iki.fi>

### References

See citation("netresponse")

---

list.responses.factor *List significant responses*

---

## Description

List significantly associated responses for all factors and levels in the given annotation matrix

## Usage

```
list.responses.factor(annotation.df, models, method = "hypergeometric",  
  min.size = 2, qth = Inf, verbose = TRUE, data = NULL,  
  rounding = NULL)
```

## Arguments

annotation.df	annotation data.frame with discrete factor levels, rows named by the samples
models	List of models. Each model should have a sample-cluster assignment matrix qofz, or a vector of cluster indices named by the samples.
method	method for enrichment calculation
min.size	minimum sample size for a response
qth	q-value threshold
verbose	verbose
data	data (samples x features; or a vector in univariate case)
rounding	rounding digits

## Value

Table listing all associations between the factor levels and responses

## Author(s)

Contact: Leo Lahti <leo.lahti@iki.fi>

## References

See citation("netresponse")

---

`list.responses.factor.minimal`*List factor responses (minimal)*

---

**Description**

List significantly associated responses for all factors and levels in the given annotation matrix

**Usage**

```
list.responses.factor.minimal(annotation.df, groupings,  
  method = "hypergeometric", min.size = 2, pth = Inf, verbose = TRUE,  
  data = NULL, rounding = NULL)
```

**Arguments**

<code>annotation.df</code>	annotation data.frame with discrete factor levels, rows named by the samples
<code>groupings</code>	List of groupings. Each model should have a sample-cluster assignment matrix qofz, or a vector of cluster indices named by the samples.
<code>method</code>	method for enrichment calculation
<code>min.size</code>	minimum sample size for a response
<code>pth</code>	p-value threshold; applied to adjusted p-value
<code>verbose</code>	verbose
<code>data</code>	data (samples x features; or a vector in univariate case)
<code>rounding</code>	rounding digits

**Value**

A list with two elements: Table listing all associations between the factor levels and responses; multiple p-value adjustment method

**Author(s)**

Contact: Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse")

---

`list.significant.responses`  
*Listing significant responses*

---

### **Description**

List responses with significant associations to a given sample group.

### **Usage**

```
list.significant.responses(model, sample, qth = 1,  
  method = "hypergeometric")
```

### **Arguments**

<code>model</code>	NetResponseModel object.
<code>sample</code>	User-specified samples group for which the enrichments are calculated. For instance, an annotation category.
<code>qth</code>	q-value threshold for enrichments
<code>method</code>	Enrichment method.

### **Value**

Statistics of the significantly associated responses.

### **Author(s)**

Leo Lahti <leo.lahti@iki.fi>

### **References**

See citation("netresponse")

### **See Also**

`response.enrichment`

### **Examples**

```
#
```

---

listify.groupings	<i>Convert grouping info into a list; each element corresponds to a group and lists samples in that group.</i>
-------------------	--

---

**Description**

Convert grouping info into a list; each element corresponds to a group and lists samples in that group.

**Usage**

```
listify.groupings(groupings, verbose = FALSE)
```

**Arguments**

groupings	a list, a vector, or a samplesxmodes assignment matrix
verbose	verbose

**Value**

Group list

**Author(s)**

Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse")

**Examples**

```
#
```

---

mixture.model	<i>Mixture model</i>
---------------	----------------------

---

**Description**

Fit Gaussian mixture model

**Usage**

```
mixture.model(x, mixture.method = "vdp", max.responses = 10,
  implicit.noise = 0, prior.alpha = 1, prior.alphaKsi = 0.01,
  prior.betaKsi = 0.01, vdp.threshold = 1e-05, initial.responses = 1,
  ite = Inf, speedup = TRUE, bic.threshold = 0, pca.basis = FALSE,
  min.responses = 1, ...)
```

**Arguments**

<code>x</code>	data matrix (samples x features, for multivariate analysis) or a vector (for univariate analysis)
<code>mixture.method</code>	Specify the approach to use in mixture modeling. Options. <code>vdp</code> (nonparametric Variational Dirichlet process mixture model); <code>bic</code> (based on Gaussian mixture modeling with EM, using BIC to select the optimal number of components)
<code>max.responses</code>	Maximum number of responses for each subnetwork. Can be used to limit the potential number of network states.
<code>implicit.noise</code>	Implicit noise parameter. Add implicit noise to <code>vdp</code> mixture model. Can help to avoid overfitting to local optima, if this appears to be a problem.
<code>prior.alpha</code> , <code>prior.alphaKsi</code> , <code>prior.betaKsi</code>	Prior parameters for Gaussian mixture model that is calculated for each subnetwork (normal-inverse-Gamma prior). <code>alpha</code> tunes the mean; <code>alphaKsi</code> and <code>betaKsi</code> are the shape and scale parameters of the inverse Gamma function, respectively.
<code>vdp.threshold</code>	Minimal free energy improvement after which the variational Gaussian mixture algorithm is deemed converged.
<code>initial.responses</code>	Initial number of components for each subnetwork model. Used to initialize calculations.
<code>ite</code>	Maximum number of iterations on posterior update ( <code>updatePosterior</code> ). Increasing this can potentially lead to more accurate results, but computation may take longer.
<code>speedup</code>	Takes advantage of approximations to PCA, mutual information etc in various places to speed up calculations. Particularly useful with large and densely connected networks and/or large sample size.
<code>bic.threshold</code>	BIC threshold which needs to be exceeded before a new mode is added to the mixture with <code>mixture.method = "bic"</code>
<code>pca.basis</code>	<code>pca.basis</code>
<code>min.responses</code>	minimum number of responses
<code>...</code>	Further optional arguments to be passed.

**Value**

List with two elements: `model`: fitted mixture model (parameters and free energy); `model.params`: model parameters

**Author(s)**

Contact: Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse")

---

`model.stats`*model.stats*

---

**Description**

Subnetwork statistics: size and number of distinct responses for each subnet.

**Usage**

```
model.stats(models)
```

**Arguments**

`models`            NetResponse object or list of models

**Value**

A 'subnetworks x properties' data frame containing the following elements.

`subnet.size`:     Vector of subnetwork sizes.

`subnet.responses`:

Vector giving the number of responses in each subnetwork.

**Author(s)**

Leo Lahti <leo.lahti@iki.fi>

**References**

Leo Lahti et al.: Global modeling of transcriptional responses in interaction networks. *Bioinformatics* (2010). See citation("netresponse") for reference details.

**Examples**

```
# Load a pre-calculated netresponse model obtained with
# model <- detect.responses(toydata$emat, toydata$netw, verbose = FALSE)
data(toydata)
# Calculate summary statistics for the model
stat <- model.stats(toydata$model)
```



---

 NetResponseModel-class

*Class "NetResponseModel"*


---

### Description

A NetResponse model.

### Objects from the Class

Returned by `detect.responses` function.

### Author(s)

Leo Lahti <leo.lahti@iki.fi>

### Examples

```
showClass("NetResponseModel")
```

---

`order.responses`

*order.responses*

---

### Description

Orders the responses by association strength (enrichment score) to a given sample set. For instance, if the samples correspond to a particular experimental factor, this function can be used to prioritize the responses according to their association strength to this factor.

### Usage

```
order.responses(models, sample, method = "hypergeometric", min.size = 2,
  max.size = Inf, min.responses = 2, subnet.ids = NULL, verbose = FALSE,
  data = NULL)
```

### Arguments

<code>models</code>	List of models. Each model should have a sample-cluster assignment matrix <code>qofz</code> .
<code>sample</code>	Measure enrichment of this sample (set) across the observed responses.
<code>method</code>	'hypergeometric' measures enrichment of factor levels in this response; 'precision' measures response purity for each factor level; 'dependency' measures logarithm of the joint density between response and factor level vs. their marginal densities: $\log(P(r,s)/(P(r)P(s)))$

<code>min.size</code> , <code>max.size</code> , <code>min.responses</code>	Optional parameters to filter the results based on subnet size and number of responses.
<code>subnet.ids</code>	Specify subnets for which the responses shall be ordered. By default, use all subnets.
<code>verbose</code>	Follow progress by intermediate messages.
<code>data</code>	data (samples x features; or a vector in univariate case)

**Value**

A data frame with elements 'ordered.responses' which gives a data frame of responses ordered by enrichment score for the investigated sample. The subnetwork, response id and enrichment score are shown. The method field indicates the enrichment calculation method. The sample field lists the samples et for which the enrichments were calculated. The info field lists additional information on enrichment statistics.

**Note**

Tools for analyzing end results of the model.

**Author(s)**

Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse") for citation details.

**Examples**

```
#
# - for given sample/s (factor level),
#   order responses (across all subnets) by association strength
#   (enrichment score); overrepresentation
# order.responses(model, sample, method = "hypergeometric")
```

---

osmo

*Osmoshock data set (PPI and expression)*

---

**Description**

A combined yeast data set with protein-protein interactions and gene expression (osmotick shock response). Gene expression profiles are transformed into links by computing a Pearson correlation for all pairs of genes and treating all correlations above 0.85 as additional links. Number of genes: 1711, number of interactions: 10250, number of gene expression observations: 133, number of total links with PPI and expression links: 14256.

**Usage**

```
data(osmo)
```

**Format**

List of following objects:

**ppi** PPI data matrix

**exp** gene expression profiles data matrix

**gids** Vector of gene ids corresponding to indices used in data matrices

**obs** Gene expression observation details

**combined.links** pooled matrix of PPI and expression links

**Source**

PPI data pooled from yeast data sets of [1] and [2]. Dna damage expression set of [3].

**References**

Ulitsky, I. and Shamir, R. *Identification of functional modules using network topology and high-throughput data*. BMC Systems Biology 2007, 1:8.

Nariai, N., Kolaczyk, E. D. and Kasif, S. *Probabilistic Protein Function Prediction from Heterogeneous Genome-Wide Data*. PLoS ONE 2007, 2(3):e337.

O'Rourke, S. and Herskowitz, I. *Unique and redundant roles for Hog MAPK pathway components as revealed by whole-genome expression analysis*. Molecular Biology of the Cell 2004, 15:532-42.

**Examples**

```
data(osmo)
```

---

PlotMixture

*Plot mixtures*

---

**Description**

Plot mixtures.

**Usage**

```
PlotMixture(x, qofz, binwidth = 0.05, xlab.text = NULL, ylab.text = NULL,  
title.text = NULL)
```

**Arguments**

x	data vector
qofz	Mode assignment probabilities for each sample. Samples x modes.
binwidth	binwidth for histogram
xlab.text	xlab.text
ylab.text	ylab.text
title.text	title.text

**Value**

Used for its side-effects

**Author(s)**

Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse") for citation details.

**Examples**

```
# PlotMixture(x, qofz)
```

---

PlotMixtureBivariate *PlotMixtureBivariate*

---

**Description**

Visualize data, centroids and response confidence intervals for a given Gaussian mixture model in two-dimensional (bivariate) case. Optionally, color the samples according to annotations labels.

**Usage**

```
PlotMixtureBivariate(x, means, sds, ws, labels = NULL, confidence = 0.95,
  main = "", ...)
```

**Arguments**

x	data matrix (samples x features)
means	mode centroids (modes x features)
sds	mode standard deviations, assuming diagonal covariance matrices (modes x features, each row giving the sqrt of covariance diagonal for the corresponding mode)
ws	weight for each mode

labels	Optional: sample class labels to be indicated in colors.
confidence	Confidence interval for the responses based on the covariances of each response. If NULL, no plotting.
main	title text
...	Further arguments for plot function.

**Value**

Used for its side-effects.

**Author(s)**

Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse") for citation details.

**Examples**

```
#plotMixture(dat, means, sds, ws)
```

---

PlotMixtureMultivariate  
*PlotMixtureMultivariate*

---

**Description**

Visualize data, centroids and response confidence intervals for a given Gaussian mixture model with PCA. Optionally, color the samples according to annotations labels.

**Usage**

```
PlotMixtureMultivariate(x, means, sds, ws, labels = NULL, title = NULL,
  modes = NULL, pca = FALSE, qofz = NULL, ...)
```

**Arguments**

x	data matrix (samples x features)
means	mode centroids (modes x features)
sds	mode standard deviations, assuming diagonal covariance matrices (modes x features, each row giving the sqrt of covariance diagonal for the corresponding mode)
ws	weight for each mode
labels	Optional: sample class labels to be indicated in colors.
title	title

modes	Optional: provide sample modes for visualization already in the input
pca	The data is projected on PCA plane by default ( <code>pca = TRUE</code> ). By setting this off ( <code>pca = FALSE</code> ) it is possible to visualize two-dimensional data in the original domain.
qofz	Sample-response probabilistic assignments matrix (samples x responses)
...	Further arguments for plot function.

**Value**

Used for its side-effects.

**Author(s)**

Leo Lahti <leo.lahti@iki.fi>

**References**

See `citation("netresponse")` for citation details.

**Examples**

```
#plotMixture(dat, means, sds, ws)
```

---

`PlotMixtureMultivariate.deprecated`

*PlotMixtureMultivariate.deprecated*

---

**Description**

Visualize data, centroids and response confidence intervals for a given Gaussian mixture model with PCA. Optionally, color the samples according to annotations labels.

**Usage**

```
PlotMixtureMultivariate.deprecated(x, means, sds, ws, labels = NULL,
  confidence = 0.95, ...)
```

**Arguments**

x	data matrix (samples x features)
means	mode centroids (modes x features)
sds	mode standard deviations, assuming diagonal covariance matrices (modes x features, each row giving the sqrt of covariance diagonal for the corresponding mode)
ws	weight for each mode
labels	Optional: sample class labels to be indicated in colors.

confidence	Confidence interval for the responses based on the covariances of each response. If NULL, no plotting.
...	Further arguments for plot function.

**Value**

Used for its side-effects.

**Author(s)**

Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse") for citation details.

**Examples**

```
#plotMixture(dat, means, sds, ws)
```

---

PlotMixtureUnivariate *Plot univariate mixtures*

---

**Description**

Visualize data, centroids and stds for a given univariate Gaussian mixture model with PCA.

**Usage**

```
PlotMixtureUnivariate(x, means = NULL, sds = NULL, ws = NULL,
  title.text = NULL, xlab.text = NULL, ylab.text = NULL,
  binwidth = 0.05, qofz = NULL, density.color = "darkgray",
  cluster.assignments = NULL, ...)
```

**Arguments**

x	data vector
means	mode centroids
sds	mode standard deviations
ws	weight for each mode
title.text	Plot title
xlab.text	xlab.text
ylab.text	ylab.text
binwidth	binwidth for histogram
qofz	Mode assignment probabilities for each sample. Samples x modes.

`density.color` Color for density lines  
`cluster.assignments` Vector of cluster indices, indicating cluster for each data point  
`...` Further arguments for plot function.

**Value**

Used for its side-effects

**Author(s)**

Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse") for citation details.

**Examples**

```
# plotMixtureUnivariate(dat, means, sds, ws)
```

---

plotPCA

*plotPCA*

---

**Description**

Visualize data, centroids and response confidence intervals for a given subnetwork with PCA. Optionally, color the samples according to annotations labels.

**Usage**

```
plotPCA(x, subnet.id, labels = NULL, confidence = 0.95, ...)
```

**Arguments**

`x` NetResponseModel object. Output from the detect.responses function.  
`subnet.id` Subnetwork id. Either character as 'Subnetwork-2' or numeric as 2, which is then converted to character.  
`labels` Optional: sample class labels to be indicated in colors.  
`confidence` Confidence interval for the responses based on the covariances of each response. If NULL, no plotting.  
`...` Further arguments for plot function.

**Value**

Used for its side-effects.



**Author(s)**

Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse") for citation details.

**Examples**

```
#plotPCA(x, subnet.id)
```

---

plot\_associations      *Association strength between category labels and responses.*

---

**Description**

Plot association strength between user-defined category labels and responses in a selected subnetwork. Associations are shown in terms  $-\log_{10}(p)$  enrichment values for the annotation categories for the responses within the specified subnetwork. No correction for multiple testing.

**Usage**

```
plot_associations(x, subnet.id, labels, method = "hypergeometric",  
  mode = "group.by.classes", ...)
```

**Arguments**

x	NetResponseModel object
subnet.id	Subnetwork.
labels	Factor. Labels for the data samples. Name by samples, or provide in the same order as in the original data.
method	Method to calculate association strength.
mode	group.by.responses or group.by.classes: indicate barplot grouping type.
...	Other arguments to be passed for plot_

**Value**

Used for side effect (plotting).

**Author(s)**

Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse").

**See Also**

plot\_responses

**Examples**

```
#
```

---

plot\_data

*Plot observed data.*

---

**Description**

Plotting tool for measurement data. Produces boxplot for each feature in each annotation category for the selected subnetwork.

**Usage**

```
plot_data(x, subnet.id, labels, ...)
```

**Arguments**

x	NetResponseModel object.
subnet.id	Specify the subnetwork.
labels	Annotation categories.
...	Further arguments for plot function.

**Value**

ggplot2 plot object

**Author(s)**

Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse")

**See Also**

plot\_responses

**Examples**

```
#
```

---

plot\_expression      *plot\_expression*

---

### Description

Plot expression matrix in color scale. For one-channel data; plot expression of each gene relative to its mean expression level over all samples. Blue indicates decreased expression and red indicates increased expression. Brightness of the color indicates magnitude of the change. Black denotes no change.

### Usage

```
plot_expression(x, maintext, ...)
```

### Arguments

x	samples x features matrix
maintext	main title
...	optional arguments

### Value

Used for its side effects.

### Author(s)

Leo Lahti <leo.lahti@iki.fi>

### References

See citation("netresponse").

### See Also

[plot\\_scale](#)

### Examples

```
#plot_expression(x)
```

---

plot\_matrix

*Visualize a matrix with one or two-way color scale.*


---

### Description

Fast investigation of matrix objects; standard visualization choices are made automatically; fast and easy-to-use but does not necessarily provide optimal visualization.

### Usage

```
plot_matrix(mat, type = "twoway", midpoint = 0, palette = NULL,
  colors = NULL, col.breaks = NULL, interval = 0.1, plot_axes = "both",
  row.tick = 1, col.tick = 1, cex.xlab = 0.9, cex.ylab = 0.9,
  xlab = NULL, ylab = NULL, limit.trunc = 0, mar = c(5, 4, 4, 2), ...)
```

### Arguments

mat	matrix
type	String. Specifies visualization type. Options: 'oneway' (color scale ranges from white to dark red; the color can be changed if needed); 'twoway' (color scale ranges from dark blue through white to dark red; colors can be changed if needed)
midpoint	middle point for the color plot: smaller values are shown with blue, larger are shown with red in type = 'twoway'
palette	Optional. Color palette.
colors	Optional. Colors.
col.breaks	breakpoints for the color palette
interval	interval for palette color switches
plot_axes	String. Indicates whether to plot x-axis ('x'), y-axis ('y'), or both ('both').
row.tick	interval for plotting row axis texts
col.tick	interval for plotting column axis texts
cex.xlab	use this to specify distinct font size for the x axis
cex.ylab	use this to specify distinct font size for the y axis
xlab	optional x axis labels
ylab	optional y axis labels
limit.trunc	color scale limit breakpoint
mar	image margins
...	optional parameters to be passed to function 'image', see help(image) for further details

### Value

A list with the color palette (colors), color breakpoints (breaks), and palette function (palette.function)

**Author(s)**

Leo Lahti <microbiome-admin@googlegroups.com>

**References**

See citation('microbiome')

**Examples**

```
mat <- rbind(c(1,2,3,4,5), c(1, 3, 1), c(4,2,2))
plot_matrix(mat, 'tway', midpoint = 3)
```

---

plot_response	<i>plot_response</i>
---------------	----------------------

---

**Description**

Plot a specific transcriptional response for a given subnetwork. TRUE, colors = TRUE, plot\_type = "twopi", ...)

**Usage**

```
plot_response(x, mynet, mybreaks, mypalette, plot_names = TRUE,
             colors = TRUE, plot_type = "twopi", ...)
```

**Arguments**

x	A numerical vector, or NULL.
mynet	Binary matrix specifying the interactions between nodes.
mybreaks	Specify breakpoints for color plot_
mypalette	Specify palette for color plot_
plot_names	Plot node names (TRUE) or indices (FALSE).
colors	Plot colors. Logical.
plot_type	Network plot mode. For instance, 'neato' or 'twopi'.
...	Further arguments for plot function.

**Value**

Used for its side-effects.

**Author(s)**

Leo Lahti, Olli-Pekka Huovilainen and Antonio Gusmao. Maintainer: Leo Lahti <leo.lahti@iki.fi>

## References

L. Lahti et al.: Global modeling of transcriptional responses in interaction networks. Submitted.

## Examples

```
#tmp <- plot_response(model, mynet,
# maintext = paste("Subnetwork", subnet.id))
```

---

plot_responses	<i>plot_responses</i>
----------------	-----------------------

---

## Description

Plot the detected transcriptional responses for a given subnetwork. `plot_mode = "network"`, `xaxis = TRUE`, `yaxis = TRUE`, `plot_type = "twopi"`, `mar = c(5, 4, 4, 2)`, `horiz = TRUE`, `datamatrix = NULL`, `scale = FALSE`, ...)

## Usage

```
plot_responses(x, subnet.id, nc = 3, plot_names = TRUE,
plot_mode = "network", xaxis = TRUE, yaxis = TRUE,
plot_type = "twopi", mar = c(5, 4, 4, 2), horiz = TRUE,
datamatrix = NULL, scale = FALSE, ...)
```

## Arguments

<code>x</code>	Result from <code>NetResponse</code> ( <code>detect.responses</code> function).
<code>subnet.id</code>	Subnet id.
<code>nc</code>	Number of columns for an array of images.
<code>plot_names</code>	Plot node names (TRUE) or indices (FALSE).
<code>plot_mode</code>	<code>network</code> : plot responses as a subnetwork graph; <code>matrix</code> , <code>heatmap</code> : plot subnetwork expression matrix. For both, expression of each gene is shown relative to the mean expression level of the gene; <code>boxplot_data</code> : feature-wise boxplots for hard sample-to-response assignments; <code>response.barplot</code> : estimated response centroids as barplot including 95 confidence intervals for the means; <code>pca</code> : PCA projection with estimated centroids and 95 two-dimensional case the original coordinates are used.
<code>xaxis</code> , <code>yaxis</code>	Logical. Plot row/column names.
<code>plot_type</code>	Network plot mode. For instance, 'neato' or 'twopi'.
<code>mar</code>	Figure margins.
<code>horiz</code>	Logical. Horizontal barplot_
<code>datamatrix</code>	<code>datamatrix</code>
<code>scale</code>	scale the phylotypes to unit length (only implemented for <code>plot_mode = "matrix"</code> )
...	Further arguments for plot function.

**Value**

Used for its side-effects.

**Author(s)**

Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse")

**See Also**

[plot\\_scale](#)

**Examples**

```
#
#res <- detect.responses(D, netw)
#vis <- plot_responses(res, subnet.id)
```

---

plot\_scale

*plot\_scale*

---

**Description**

Plot the color scale used in visualization.

**Usage**

```
plot_scale(x, y, m = NULL, cex.axis = 1.5, label.step = 2,
  interval = 0.1, two.sided = TRUE, label.start = NULL, Nlab = 3, ...)
```

**Arguments**

x	Breakpoints for the plot_
y	Color palette.
m	Breakpoints' upper limit.
cex.axis	Axis scale.
label.step	Density of the labels.
interval	Interval.
two.sided	Plot two-sided (TRUE) or one-sided (FALSE) visualization.
label.start	Label starting point.
Nlab	Number of labels to plot_
...	Further arguments for plot function.

**Value**

Used for its side-effects.

**Author(s)**

Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse")

**Examples**

```
#
#res <- detect.responses(D, netw, verbose = FALSE)
#vis <- plot_responses(res, subnet.idx)
#plot_scale(vis$breaks, vis$palette)
```

---

plot\_subnet

*plot\_subnet*

---

**Description**

Plot the given subnetwork.

**Usage**

```
plot_subnet(x, subnet.id, network, plot_names = TRUE, ...)
```

**Arguments**

x	Result from NetResponse (detect.responses function).
subnet.id	Subnet id.
network	Original network used in the modelling.
plot_names	Plot node names (TRUE) or indices (FALSE).
...	Further arguments for plot function.

**Value**

Used for its side-effects. Returns a matrix that describes the investigated subnetwork.

**Author(s)**

Leo Lahti, Olli-Pekka Huovilainen and Antonio Gusmao. Maintainer: Leo Lahti <leo.lahti@iki.fi>



## References

L. Lahti et al.: Global modeling of transcriptional responses in interaction networks. Submitted.

## Examples

```
#  
# res <- detect.responses(D, netw, verbose = FALSE)  
# net <- plot_subnet(res, subnet.idx = 1)
```

---

read.sif	<i>Reading network files</i>
----------	------------------------------

---

## Description

Function to read network files.

## Usage

```
read.sif(sif.file, format = "graphNEL", directed = FALSE, header =  
TRUE, sep = "\t", ...)
```

## Arguments

sif.file	Name of network file in SIF format.
format	Output format: igraph or graphNEL
directed	Logical. Directed/undirected graph. Not used in the current model.
header	Logical. Indicate whether the SIF file has header or not.
sep	Field separator.
...	Further optional arguments to be passed for file reading.

## Details

Read in SIF network file, return R graph object in igraph or graphNEL format.

## Value

R graph object in igraph or graphNEL format.

## Author(s)

Leo Lahti <leo.lahti@iki.fi>

## Examples

```
#net <- read.sif("network.sif")
```

---

response.enrichment    *Enrichment for a specified sample group in the given response.*

---

**Description**

Calculate enrichment values for a specified sample group in the given response.

**Usage**

```
response.enrichment(total.samples, response.samples, annotated.samples,  
method = "hypergeometric")
```

**Arguments**

total.samples	All samples in the data
response.samples	Samples in the investigated subset
annotated.samples	Samples at the investigated annotation level for enrichment calculation
method	Enrichment method.

**Value**

List with enrichment statistics, depending on enrichment method.

**Author(s)**

Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse")

**See Also**

order.responses

**Examples**

```
#enr <- response.enrichment(subnet.id, models, sample, response, method)
```

---

response2sample	<i>response2sample</i>
-----------------	------------------------

---

### Description

List the most strongly associated response of a given subnetwork for each sample.

### Usage

```
response2sample(model, subnet.id = NULL, component.list = TRUE,  
  verbose = FALSE, data = NULL)
```

### Arguments

model	A NetResponseModel object or list.
subnet.id	Subnet id. A natural number which specifies one of the subnetworks within the 'model' object.
component.list	List samples separately for each mixture component (TRUE). Else list the most strongly associated component for each sample (FALSE).
verbose	Follow progress by intermediate messages.
data	Data (features x samples; or a vector for univariate case) to predict response for given data points (currently implemented only for mixture.model output) Return:

### Value

A list. Each element corresponds to one subnetwork response, and contains a list of samples that are associated with the response (samples for which this response has the highest probability  $P(\text{response} | \text{sample})$ ).

### Author(s)

Leo Lahti <leo.lahti@iki.fi>

### References

Leo Lahti et al.: Global modeling of transcriptional responses in interaction networks. *Bioinformatics* (2010). See citation("netresponse") for citation details.

### Examples

```
# Load example data  
data( toydata )      # Load toy data set  
D   <- toydata$emat  # Response matrix (for example, gene expression)  
model <- toydata$model # Pre-calculated model  
  
# Find the samples for each response (for a given subnetwork)
```

```
response2sample(model, subnet.id = 1)
```

---

```
sample2response      sample2response
```

---

## Description

Probabilistic sample-response assignments for given subnet.

## Usage

```
sample2response(model, subnet.id, mode = "soft")
```

## Arguments

<code>model</code>	Result from <code>NetResponse</code> ( <code>detect.responses</code> function).
<code>subnet.id</code>	Subnet identifier. A natural number which specifies one of the subnetworks within the 'model' object.
<code>mode</code>	soft: gives samples x responses probabilistic assignment matrix; hard: gives the most likely response for each sample

## Value

A matrix of probabilities. Sample-response assignments for given subnet, listing the probability of each response, given a sample.

## Author(s)

Leo Lahti <leo.lahti@iki.fi>

## References

Leo Lahti et al.: Global modeling of transcriptional responses in interaction networks. *Bioinformatics* (2010). See `citation("netresponse")` for citation details.

## Examples

```
#data( toydata )      # Load toy data set
#D   <- toydata$emat  # Response matrix (for example, gene expression)
#netw <- toydata$netw # Network

# Detect network responses
#model <- detect.responses(D, netw, verbose = FALSE)

# Assign samples to responses (soft, probabilistic assignments sum to 1)
#response.probabilities <- sample2response(model, subnet.id = "Subnet-1")
```

---

set.breaks	<i>Set breaks</i>
------------	-------------------

---

**Description**

Set breakpoints for two-way color palette.

**Usage**

```
set.breaks(mat, interval = 0.1)
```

**Arguments**

mat	Matrix to visualize.
interval	Density of color breakpoints.

**Value**

A vector listing the color breakpoints.

**Author(s)**

Leo Lahti, Olli-Pekka Huovilainen and Antonio Gusmao. Maintainer: Leo Lahti <leo.lahti@iki.fi>

**References**

L. Lahti et al.: Global modeling of transcriptional responses in interaction networks. Submitted.

**Examples**

```
set.breaks(array(rnorm(100), dim = c(10, 10)), interval = .1)
```

---

toydata	<i>toydata</i>
---------	----------------

---

**Description**

Toy data for NetResponse examples.

**Usage**

```
data(toydata)
```

**Format**

Toy data: a list with three elements:

emat: Data matrix (samples x features). This contains the same features that are provided in the network (toydata\$netw). The matrix characterizes measurements of network states across different conditions.

netw: Binary matrix that describes pairwise interactions between features. This defines an undirected network over the features. A link between two nodes is denoted by 1.

model: A pre-calculated model. Object of NetResponseModel class, resulting from applying the netresponse algorithm on the toydata with `model <- detect.responses(D, netw)`.

**References**

Leo Lahti et al.: Global modeling of transcriptional responses in interaction networks. *Bioinformatics* (2010).

**Examples**

```
data(toydata)
D <- toydata$emat # Response matrix (samples x features)
netw <- toydata$netw # Network between the features
model <- toydata$model # Pre-calculated NetResponseModel obtained with
# model <- detect.responses(D, netw)
```

---

vdp.mixt

*vdp.mixt*


---

**Description**

Accelerated variational Dirichlet process Gaussian mixture.

**Usage**

```
vdp.mixt(dat, prior.alpha = 1, prior.alphaKsi = 0.01,
  prior.betaKsi = 0.01, do.sort = TRUE, threshold = 1e-05,
  initial.K = 1, ite = Inf, implicit.noise = 0, c.max = 10,
  speedup = TRUE, min.size = 5)
```

**Arguments**

**dat** Data matrix (samples x features).

**prior.alpha, prior.alphaKsi, prior.betaKsi** Prior parameters for Gaussian mixture model (normal-inverse-Gamma prior). alpha tunes the mean; alphaKsi and betaKsi are the shape and scale parameters of the inverse Gamma function, respectively.

**do.sort** When true, qOFz will be sorted in decreasing fashion by component size, based on colSums(qOFz). The qOFz matrix describes the sample-component assignments in the mixture model.

threshold	Defines the minimal free energy improvement that stops the algorithm: used to define convergence limit.
initial.K	Initial number of mixture components.
ite	Defines maximum number of iterations on posterior update (updatePosterior). Increasing this can potentially lead to more accurate results, but computation may take longer.
implicit.noise	Adds implicit noise; used by vdp.mk.log.lambda.so and vdp.mk.hp.posterior.so. By adding noise (positive values), one can avoid overfitting to local optima in some cases, if this happens to be a problem.
c.max	Maximum number of candidates to consider in find.best.splitting. During mixture model calculations new mixture components can be created until this upper limit has been reached. Defines the level of truncation for a truncated stick-breaking process.
speedup	When learning the number of components, each component is splitted based on its first PCA component. To speed up, approximate by using only subset of data to calculate PCA.
min.size	Minimum size for a component required for potential splitting during mixture estimation.

## Details

Implementation of the Accelerated variational Dirichlet process Gaussian mixture model algorithm by Kenichi Kurihara et al., 2007.

**ALGORITHM SUMMARY** This code implements Gaussian mixture models with diagonal covariance matrices. The following greedy iterative approach is taken in order to obtain the number of mixture models and their corresponding parameters:

1. Start from one cluster,  $T = 1$ .
2. Select a number of candidate clusters according to their values of  $Nc = \sum_{n=1}^N q_{z_n}(z_n = c)$  (larger is better).
3. For each of the candidate clusters,  $c$ :
  - 3a. Split  $c$  into two clusters,  $c_1$  and  $c_2$ , through the bisector of its principal component. Initialise the responsibilities  $q_{z_n}(z_n = c_1)$  and  $q_{z_n}(z_n = c_2)$ .
  - 3b. Update only the parameters of  $c_1$  and  $c_2$  using the observations that belonged to  $c$ , and determine the new value for the free energy,  $FT+1$ .
  - 3c. Reassign cluster labels so that cluster 1 corresponds to the largest cluster, cluster 2 to the second largest, and so on.
4. Select the split that lead to the maximal reduction of free energy,  $FT+1$ .
5. Update the posterior using the newly split data.
6. If  $FT - FT+1 < \epsilon$  then halt, else set  $T := T + 1$  and go to step 2.

The loop is implemented in the function greedy(...)

## Value

prior	Prior parameters of the vdp-gm model (qofz: priors on observation lables; Mu: centroids; S2: variance).
posterior	Posterior estimates for the model parameters and statistics.
weights	Mixture proportions, or weights, for the Gaussian mixture components.
centroids	Centroids of the mixture components.

sds	Standard deviations for the mixture model components (posterior modes of the covariance diagonals square root). Calculated as $\sqrt{(\text{invgam.scale}/(\text{invgam.shape} + 1))}$ .
qOfz	Sample-to-cluster assignments (soft probabilistic associations).
Nc	Component sizes
invgam.shape	Shape parameter (alpha) of the inverse Gamma distribution
invgam.scale	Scale parameter (beta) of the inverse Gamma distribution
Nparams	Number of model parameters
K	Number of components in the mixture model
opts	Model parameters that were used.
free.energy	Free energy of the model.

### Note

This implementation is based on the Variational Dirichlet Process Gaussian Mixture Model implementation, Copyright (C) 2007 Kenichi Kurihara (all rights reserved) and the Agglomerative Independent Variable Group Analysis package (in Matlab): Copyright (C) 2001-2007 Esa Alhoniemi, Antti Honkela, Krista Lagus, Jeremias Seppa, Harri Valpola, and Paul Wagner.

### Author(s)

Maintainer: Leo Lahti <leo.lahti@iki.fi>

### References

Kenichi Kurihara, Max Welling and Nikos Vlassis: Accelerated Variational Dirichlet Process Mixtures. In B. Schölkopf and J. Platt and T. Hoffman (eds.), Advances in Neural Information Processing Systems 19, 761–768. MIT Press, Cambridge, MA 2007.

### Examples

```
set.seed(123)

# Generate toy data with two Gaussian components
dat <- rbind(array(rnorm(400), dim = c(200,2)) + 5,
            array(rnorm(400), dim = c(200,2)))

# Infinite Gaussian mixture model with
# Variational Dirichlet Process approximation
mixt <- vdp.mixt( dat )

# Centroids of the detected Gaussian components
mixt$posterior$centroids

# Hard mixture component assignments for the samples
apply(mixt$posterior$qOfz, 1, which.max)
```



---

vectorize.groupings    *Convert grouping info into a vector; each element corresponds to a group and lists samples in that group.*

---

**Description**

Convert grouping info into a vector; each element corresponds to a group and lists samples in that group.

**Usage**

```
vectorize.groupings(groupings, verbose = FALSE)
```

**Arguments**

groupings	a list, a vector, or a samplesxmodes assignment matrix
verbose	verbose

**Value**

Indicator vector

**Author(s)**

Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse")

**Examples**

```
#
```

---

```
write.netresponse.results
```

*Write NetResponse results summary into a file.*

---

**Description**

Write NetResponse results summary into a file.

**Usage**

```
write.netresponse.results(x, subnet.ids = NULL, filename)
```

**Arguments**

x	NetResponseModel
subnet.ids	List of subnet ids to consider. By default, all subnets.
filename	Output file name.

**Details**

Experimental version.

**Value**

Used for side effects.

**Author(s)**

Leo Lahti <leo.lahti@iki.fi>

**References**

See citation("netresponse")

**Examples**

#

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