

# Package ‘WGCNA’

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**Title** Weighted Correlation Network Analysis

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**Depends** R (>= 2.14), stats, impute, grDevices, dynamicTreeCut (>= 1.20), utils, flashClust, Hmisc, splines, cluster, reshape, MASS, foreach, doParallel

**Suggests** GO.db, org.Hs.eg.db, org.Mm.eg.db, AnnotationDbi, infotheo, entropy, minet, survival

**ZipData** no

**License** GPL (>= 2)

**Description** Functions necessary to perform Weighted Correlation Network Analysis.

**URL** <http://www.genetics.ucla.edu/labs/horvath/CoexpressionNetwork/Rpackages/WGCNA/>

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WGCNA-package	<i>Weighted Gene Co-Expression Network Analysis</i>
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## Description

Functions necessary to perform Weighted Correlation Network Analysis. WGCNA is also known as weighted gene co-expression network analysis when dealing with gene expression data. Many functions of WGCNA can also be used for general association networks specified by a symmetric adjacency matrix.

## Details

Package:	WGCNA
Version:	1.29
Date:	2013-07-18
Depends:	R (>= 2.10), stats, impute, grDevices,
Suggests: AnnotationDbi, GO.db, org.*.eg.db, infotheo, entropy, minet ZipData:	no
License:	GPL (>= 2)
URL:	<a href="http://www.genetics.ucla.edu/labs/horv">http://www.genetics.ucla.edu/labs/horv</a>

## Index:

GTOMdist	Generalized Topological Overlap Measure
TOMdist	Topological overlap matrix dissimilarity
TOMplot	Graphical representation of the Topological Overlap Matrix
TOMsimilarity	Topological overlap matrix similarity
TOMsimilarityFromExpr	Topological overlap matrix similarity
WGCNA-package	Weighted Gene Co-Expression Network Analysis
accuracyMeasures	Accuracy measures for a 2x2 confusion matrix
addErrorBars	Add error bars to a barplot.
addGrid	Add grid lines to an existing plot.
addGuideLines	Add vertical "guide lines" to a dendrogram plot
addTraitToMEs	Add trait information to multi-set module eigengene structure
adjacency	Calculate network adjacency
adjacency.fromSimilarity	Calculate network adjacency from a similarity matrix
adjacency.polyReg	Adjacency based on polynomial regression
adjacency.splineReg	Adjacency based on natural cubic spline regression

alignExpr	Align expression data with given vector
automaticNetworkScreening	One-step automatic network gene screening
automaticNetworkScreeningGS	One-step automatic network gene screening with external gene significance
AFcorMI	Prediction of weighted mutual information adjacency matrix by correlation
bicor	Biweight Midcorrelation
bicorAndPvalue	Biweight Midcorrelation and the associated p-value
blockwiseConsensusModules	Find consensus modules across several datasets.
blockwiseIndividualTOMs	Calculate individual topological overlaps across multi-set data
blockwiseModules	Automatic network construction and module detection
BloodLists	(data) Gene sets for user enrichment analysis
blueWhiteRed	Blue-white-red color sequence
BrainLists	(data) Gene sets for user enrichment analysis
BrainRegionMarkers	(data) Gene Markers for Regions of the Human Brain
checkAdjMat	Check adjacency matrix
checkSets	Check structure and retrieve sizes of a group of datasets
checkSimilarity	Check a similarity matrix
chooseOneHubInEachModule	Choose a hub gene in each module
chooseTopHubInEachModule	Choose the top hub gene in each module
clusterCoef	Clustering coefficient calculation
coClustering	Cluster preservation based on co-clustering
coClustering.permutationTest	Permutation test for co-clustering
collapseRows	Collapse Rows in Numeric Matrix
collapseRowsUsingKME	Selects one representative row per group based on kM
collectGarbage	Iterative garbage collection
colQuantileC	Fast column-wise quantile of a matrix
conformityBasedNetworkConcepts	Calculation of conformity-based network concepts
conformityDecomposition	Conformity vector(s) and factorizability measure(s) of a network
consensusDissTOMandTree	Consensus TOM-based dissimilarity and clustering tree
consensusKME	Consensus eigengene-based connectivity
consensusMEDissimilarity	Consensus dissimilarity of module eigengenes.
consensusOrderMEs	Put close eigenvectors next to each other in several sets.
consensusProjectiveKMeans	Consensus projective K-means (pre-)clustering of expression data

cor	Faster calculation of Pearson correlations
corAndPvalue	Correlation and the associated p-value
cor1	Faster calculation of column correlations of a matrix
corFast	Faster calculation of Pearson correlations
corPredictionSuccess	~~function to do ... ~~
corPvalueFisher	Fisher's asymptotic p-value for correlation
corPvalueStudent	Student asymptotic p-value for correlation
correlationPreservation	Preservation of eigengene correlations
coxRegressionResiduals	Deviance- and martingale residuals from a Cox regression model
cutreeStatic	Constant height tree cut
cutreeStaticColor	Constant height tree cut using color labels
displayColors	Show colors used to label modules
dynamicMergeCut	Threshold for module merging
exportNetworkToVisANT	Export network data in format readable by VisANT
exportNetworkToCytoscape	Export network data in format readable by Cytoscape
fixDataStructure	Put single-set data into a form useful for multiset calculations
fundamentalNetworkConcepts	Calculation of fundamental network concepts
GOenrichmentAnalysis	Calculate enrichment p-values of clusters in GO terms
goodGenes	Filter genes with too many missing entries
goodGenesMS	Filter genes with too many missing entries across multiple sets
goodSamples	Filter samples with too many missing entries
goodSamplesGenes	Iterative filtering of samples and genes with too many missing entries
goodSamplesGenesMS	Iterative filtering of samples and genes with too many missing entries across multiple data sets
goodSamplesMS	Filter samples with too many missing entries across multiple data sets
greenBlackRed	Green-black-red color sequence
greenWhiteRed	Green-white-red color sequence
hubGeneSignificance	Hubgene significance
ImmunePathwayLists	(data) Immune Pathways with Corresponding Gene Markers
initProgInd	Inline display of progress
intramodularConnectivity	Calculation of intramodular connectivity
keepCommonProbes	Keep probes that are shared among given data sets
kMEcomparisonScatterplot	Scatterplots of eigengene-based connectivity
labeledBarplot	Barplot with text or color labels
labeledHeatmap	Produce a labeled heatmap plot
labelPoints	Attempt to intelligently label points in a scatterplot
labels2colors	Convert numerical labels to colors
lowerTri2matrix	Reconstruct a symmetric matrix from a distance (lower-triangular) representation
matchLabels	Relabel modules to best approximate a reference labeling
mergeCloseModules	Merge close modules in gene expression data
metaAnalysis	Meta-analysis significance statistics
metaZfunction	Meta-analysis Z statistic
moduleColor.getMEprefix	Get the prefix used to label module eigengenes

moduleEigengenes	Calculate module eigengenes
moduleMergeUsingKME	Merge modules and reassign genes using kME
moduleNumber	Fixed-height cut of a dendrogram
modulePreservation	Calculation of module preservation statistics
multiSetMEs	Calculate module eigengenes
multiData.eigengeneSignificance	Calculate eigengene significance for multiple data sets
mutualInfoAdjacency	Calculate weighted adjacency matrices based on mutual information
nPresent	Number of present data entries
nearestNeighborConnectivity	Connectivity to a constant number of nearest neighbors
nearestNeighborConnectivityMS	Connectivity to a constant number of nearest neighbors across multiple data sets
nearestCentroidPredictor	Nearest centroid predictor for two-class problems
networkConcepts	Calculations of network concepts
networkScreening	Network screening
networkScreeningGS	Network screening with external gene significance
normalizeLabels	Transform numerical labels into normal order
numbers2colors	Color representation for a numeric variable
orderBranchesUsingHubGenes	Optimize dendrogram using branch swaps and reflections
orderMEs	Put close eigenvectors next to each other
overlapTable	Overlap counts and Fisher exact tests for two sets of module labels
overlapTableUsingKME	Determines significant overlap between modules in two networks based on
pickHardThreshold	Analysis of scale free topology for hard-thresholding
pickHardThreshold.fromSimilarity	Analysis of scale free topology for hard-thresholding
pickSoftThreshold	Analysis of scale free topology for soft-thresholding
pickSoftThreshold.fromSimilarity	Analysis of scale free topology for soft-thresholding
plotClusterTreeSamples	Annotated clustering dendrogram of microarray samples
plotColorUnderTree	Plot color rows under a dendrogram
plotDendroAndColors	Dendrogram plot with color annotation of objects
plotEigengeneNetworks	Eigengene network plot
plotMEpairs	Pairwise scatterplots of eigengenes
plotModuleSignificance	Barplot of module significance
plotNetworkHeatmap	Network heatmap plot
pmean	Parallel mean
pmedian	Parallel median
populationMeansInAdmixture	Estimation of population-specific mean values in an admixed population
pquantile	Parallel quantile
preservationNetworkConnectivity	Network preservation calculations
projectiveKMeans	Projective K-means (pre-)clustering of expression data
propVarExplained	Proportion of variance explained by eigengenes
proportionsInAdmixture	



	Estimation of proportion of pure populations in an admixture
qvalue	q-value calculation from package qvalue
randIndex	Calculation of (adjusted) Rand index
randomGLMpredictor	Ensemble predictor based on bagging of generalized linear models
rankPvalue	Estimate the p-value for ranking consistently high (or low) on multiple
recutBlockwiseTrees	Repeat blockwise module detection from pre-calculated data
recutConsensusTrees	Repeat blockwise consensus module detection from pre-calculated data
redWhiteGreen	Red-white-green color sequence
reflectTwoBranches	Reflect branches in a dendrogram
relativeCorPredictionSuccess	Compare prediction success
removeGreyME	Removes the grey eigengene from a given collection of eigengenes.
removePrincipalComponents	Remove leading principal components from data
returnGeneSetsAsLists	Return pre-defined gene lists in several biomedical categories.
scaleFreeFitIndex	Calculation of fitting statistics for evaluating scale free topology fit.
scaleFreePlot	Visual check of scale-free topology
SCsLists	(data) Stem Cell-Related Genes with Corresponding Gene Markers
selectBranch	Find a branch in a dendrogram
setCorrelationPreservation	Summary correlation preservation measure
sigmoidAdjacencyFunction	Sigmoid-type adjacency function
signedKME	Signed eigengene-based connectivity
signumAdjacencyFunction	Hard-thresholding adjacency function
simulateDatExpr	Simulation of expression data
simulateDatExpr5Modules	
simulateEigengeneNetwork	Simulate eigengene network from a causal model
simulateModule	Simulate a gene co-expression module
simulateMultiExpr	Simulate multi-set expression data
simulateSmallLayer	Simulate small modules
sizeGrWindow	Open a graphics window of given width and height
softConnectivity	Calculation of soft (weighted) connecteivity
softConnectivity.fromSimilarity	Calculation of soft (weighted) connecteivity
spaste	Space-less paste
standardColors	Colors this library uses for labeling modules
standardScreeningBinaryTrait	Standard screening for a binary trait
standardScreeningCensoredTime	Standard screening with regard to a Censored Time Variable
stdErr	Standard error
stratifiedBarplot	Bar plots of data across two splitting parameters
swapTwoBranches	Swap branches in a dendrogram
TrueTrait	Estimate the true trait underlying a list of surrogate markers
transposeBigData	Block-by-block transpose of large matrices

unsignedAdjacency	Calculation of unsigned adjacency
userListEnrichment	Measure enrichment between inputted and user-defined lists
vectorTOM	Topological overlap for a subset of the whole set of genes
vectorizeMatrix	Turn a matrix into a vector of non-redundant components
verboseBarplot	Barplot with error bars, annotated by Kruskal-Wallis p-value
verboseBoxplot	Boxplot annotated by a Kruskal-Wallis p-value
verboseScatterplot	Scatterplot annotated by regression line and p-value
verboseIplot	Scatterplot annotated by regression line, p-value, and color for density
votingLinearPredictor	Voting linear predictor

### Author(s)

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accuracyMeasures	<i>Accuracy measures for a 2x2 confusion matrix or for vectors of predicted and observed values.</i>
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### Description

The function calculates various prediction accuracy statistics for predictions of binary or quantitative (continuous) responses. For binary classification, the function calculates the error rate, accuracy, sensitivity, specificity, positive predictive value, and other accuracy measures. For quantitative prediction, the function calculates correlation, R-squared, error measures, and the C-index.

**Usage**

```
accuracyMeasures(
  predicted,
  observed = NULL,
  type = c("auto", "binary", "quantitative"),
  levels = if (isTRUE(all.equal(dim(predicted), c(2,2)))) colnames(predicted)
              else sort(unique(c(observed, predicted))),
  negativeLevel = levels[2],
  positiveLevel = levels[1] )
```

**Arguments**

predicted	either a 2x2 confusion matrix (table) whose entries contain non-negative integers, or a vector of predicted values. Predicted values can be binary or quantitative (see type below). If a 2x2 matrix is given, it must have valid column and row names that specify the levels of the predicted and observed variables whose counts the matrix is giving (e.g., the function <code>table</code> sets the names appropriately.) If it is a 2x2 table and the table contains non-negative real (non-integer) numbers the function outputs a warning.
observed	if predicted is a vector of predicted values, this (observed) must be a vector of the same length giving the "gold standard" (or observed) values. Ignored if predicted is a 2x2 table.
type	character string specifying the type of the prediction problem (i.e., values in the predicted and observed vectors). The default "auto" decides type automatically: if predicted is a 2x2 table or if the number of unique values in the concatenation of predicted and observed is 2, the prediction problem (type) is assumed to be binary, otherwise it is assumed to be quantitative. Inconsistent specification (for example, when predicted is a 2x2 matrix and type is "quantitative") trigger errors.
levels	a 2-element vector specifying the two levels of binary variables. Only used if type is "binary" (or "auto" that results in the binary type). Defaults to either the column names of the confusion matrix (if the matrix is specified) or to the sorted unique values of observed and predicted.
negativeLevel	the binary value (level) that corresponds to the negative outcome. Note that the default is the second of the sorted levels (for example, if levels are 1,2, the default negative level is 2). Only used if type is "binary" (or "auto" that results in the binary type).
positiveLevel	the binary value (level) that corresponds to the positive outcome. Note that the default is the second of the sorted levels (for example, if levels are 1,2, the default negative level is 2). Only used if type is "binary" (or "auto" that results in the binary type).

**Details**

The rows of the 2x2 table `tab` must correspond to a test (or predicted) outcome and the columns to a true outcome ("gold standard"). A table that relates a predicted outcome to a true test outcome is also known as confusion matrix. Warning: To correctly calculate sensitivity and specificity, the positive and negative outcome must be properly specified so they can be matched to the appropriate rows and columns in the confusion table.

Interchanging the negative and positive levels swaps the estimates of the sensitivity and specificity but has no effect on the error rate or accuracy. Specifically, denote by `pos` the index of the positive

level in the confusion table, and by neg th index of the negative level in the confusion table. The function then defines number of true positives=TP=tab[pos, pos], no.false positives =FP=tab[pos, neg], no.false negatives=FN=tab[neg, pos], no.true negatives=TN=tab[neg, neg]. Then Specificity=  $TN/(FP+TN)$  Sensitivity=  $TP/(TP+FN)$  NegativePredictiveValue=  $TN/(FN + TN)$  PositivePredictiveValue=  $TP/(TP + FP)$  FalsePositiveRate =  $1-Specificity$  FalseNegativeRate =  $1-Sensitivity$  Power = Sensitivity LikelihoodRatioPositive =  $Sensitivity / (1-Specificity)$  LikelihoodRatioNegative =  $(1-Sensitivity)/Specificity$ . The naive error rate is the error rate of a constant (naive) predictor that assigns the same outcome to all samples. The prediction of the naive predictor equals the most frequently observed outcome. Example: Assume you want to predict disease status and 70 percent of the observed samples have the disease. Then the naive predictor has an error rate of 30 percent (since it only misclassifies 30 percent of the healthy individuals).

### Value

Data frame with two columns:

Measure	this column contains character strings that specify name of the accuracy measure.
Value	this column contains the numeric estimates of the corresponding accuracy measures.

### Author(s)

Steve Horvath and Peter Langfelder

### References

[http://en.wikipedia.org/wiki/Sensitivity\\_and\\_specificity](http://en.wikipedia.org/wiki/Sensitivity_and_specificity)

### Examples

```
m=100
trueOutcome=sample( c(1,2),m,replace=TRUE)
predictedOutcome=trueOutcome
# now we noise half of the entries of the predicted outcome
predictedOutcome[ 1:(m/2)] =sample(predictedOutcome[ 1:(m/2)] )
tab=table(predictedOutcome, trueOutcome)
accuracyMeasures(tab)

# Same result:
accuracyMeasures(predictedOutcome, trueOutcome)
```

---

addErrorBars	<i>Add error bars to a barplot.</i>
--------------	-------------------------------------

---

### Description

This function adds error bars to an existing barplot.

### Usage

```
addErrorBars(means, errors, two.side = FALSE)
```

**Arguments**

means	vector of means plotted in the barplot
errors	vector of standard errors (single positive values) to be plotted.
two.side	should the error bars be two-sided?

**Value**

None.

**Author(s)**

Steve Horvath and Peter Langfelder

---

addGrid *Add grid lines to an existing plot.*

---

**Description**

This function adds horizontal and/or vertical grid lines to an existing plot. The grid lines are aligned with tick marks.

**Usage**

```
addGrid(linesPerTick = NULL, horiz = TRUE, vert = FALSE, col = "grey30", lty = 3)
```

**Arguments**

linesPerTick	Number of lines between successive tick marks (including the line on the tick-marks themselves)
horiz	Draw horizontal grid lines?
vert	Draw vertical tick lines?
col	Specifies color of the grid lines
lty	Specifies line type of grid lines. See <a href="#">par</a> .

**Details**

If `linesPerTick` is not specified, it is set to 5 if number of tick s is 5 or less, and it is set to 2 if number of ticks is greater than 5.

**Note**

The function does not work whenever logarithmic scales are in use.

**Author(s)**

Peter Langfelder

**Examples**

```
plot(c(1:10), c(1:10))
addGrid();
```

---

addGuideLines                      *Add vertical “guide lines” to a dendrogram plot*

---

### Description

Adds vertical “guide lines” to a dendrogram plot.

### Usage

```
addGuideLines(dendro,
              all = FALSE,
              count = 50,
              positions = NULL,
              col = "grey30",
              lty = 3,
              hang = 0)
```

### Arguments

dendro	The dendrogram (see <a href="#">hclust</a> ) to which the guide lines are to be added.
all	Add a guide line to every object on the dendrogram? Useful if the number of objects is relatively low.
count	Number of guide lines to be plotted. The lines will be equidistantly spaced.
positions	Horizontal positions of the added guide lines. If given, overrides count.
col	Color of the guide lines
lty	Line type of the guide lines. See <a href="#">par</a> .
hang	Fraction of the figure height that will separate top ends of guide lines and the merge heights of the corresponding objects.

### Author(s)

Peter Langfelder

---

addTraitToMEs                      *Add trait information to multi-set module eigengene structure*

---

### Description

Adds trait information to multi-set module eigengene structure.

### Usage

```
addTraitToMEs(multiME, multiTraits)
```

**Arguments**

<code>multiME</code>	Module eigengenes in multi-set format. A vector of lists, one list per set. Each list must contain an element named <code>data</code> that is a data frame with module eigengenes.
<code>multiTraits</code>	Microarray sample trait(s) in multi-set format. A vector of lists, one list per set. Each list must contain an element named <code>data</code> that is a data frame in which each column corresponds to a trait, and each row to an individual sample.

**Details**

The function simply `cbind`'s the module eigengenes and traits for each set. The number of sets and numbers of samples in each set must be consistent between `multiMEs` and `multiTraits`.

**Value**

A multi-set structure analogous to the input: a vector of lists, one list per set. Each list will contain a component `data` with the merged eigengenes and traits for the corresponding set.

**Author(s)**

Peter Langfelder

**See Also**

[checkSets](#), [moduleEigengenes](#)

---

adjacency

*Calculate network adjacency*


---

**Description**

Calculates (correlation or distance) network adjacency from given expression data or from a similarity.

**Usage**

```
adjacency(datExpr,
          selectCols = NULL,
          type = "unsigned",
          power = if (type=="distance") 1 else 6,
          corFnc = "cor", corOptions = "use = 'p'",
          distFnc = "dist", distOptions = "method = 'euclidean'")

adjacency.fromSimilarity(similarity,
                        type = "unsigned",
                        power = if (type=="distance") 1 else 6)
```

**Arguments**

<code>datExpr</code>	data frame containing expression data. Columns correspond to genes and rows to samples.
<code>similarity</code>	a (signed) similarity matrix: square, symmetric matrix with entries between -1 and 1.
<code>selectCols</code>	for correlation networks only (see below); can be used to select genes whose adjacencies will be calculated. Should be either a numeric vector giving the indices of the genes to be used, or a boolean vector indicating which genes are to be used.
<code>type</code>	network type. Allowed values are (unique abbreviations of) "unsigned", "signed", "signed hybrid", "distance".
<code>power</code>	soft thresholding power.
<code>corFnc</code>	character string specifying the function to be used to calculate co-expression similarity for correlation networks. Defaults to Pearson correlation. Any function returning values between -1 and 1 can be used.
<code>corOptions</code>	character string specifying additional arguments to be passed to the function given by <code>corFnc</code> . Use "use = 'p', method = 'spearman'" to obtain Spearman correlation.
<code>distFnc</code>	character string specifying the function to be used to calculate co-expression similarity for distance networks. Defaults to the function <code>dist</code> . Any function returning non-negative values can be used.
<code>distOptions</code>	character string specifying additional arguments to be passed to the function given by <code>distFnc</code> . For example, when the function <code>dist</code> is used, the argument <code>method</code> can be used to specify various ways of computing the distance.

**Details**

The argument `type` determines whether a correlation (type one of "unsigned", "signed", "signed hybrid"), or a distance network (type equal "distance") will be calculated. In correlation networks the adjacency is constructed from correlations (values between -1 and 1, with high numbers meaning high similarity). In distance networks, the adjacency is constructed from distances (non-negative values, high values mean low similarity).

The function calculates the similarity of columns (genes) in `datExpr` by calling the function given in `corFnc` (for correlation networks) or `distFnc` (for distance networks), transforms the similarity according to `type` and raises it to `power`, resulting in a weighted network adjacency matrix. If `selectCols` is given, the `corFnc` function will be given arguments `(datExpr, datExpr[selectCols], ...)`; hence the returned adjacency will have rows corresponding to all genes and columns corresponding to genes selected by `selectCols`.

Correlation and distance are transformed as follows: for `type = "unsigned"`,  $\text{adjacency} = |\text{cor}|^{\text{power}}$ ; for `type = "signed"`,  $\text{adjacency} = (0.5 * (1 + \text{cor}))^{\text{power}}$ ; for `type = "signed hybrid"`,  $\text{adjacency} = \text{cor}^{\text{power}}$  if  $\text{cor} > 0$  and 0 otherwise; and for `type = "distance"`,  $\text{adjacency} = (1 - (\text{dist}/\max(\text{dist}))^2)^{\text{power}}$ .

The function `adjacency.fromSimilarity` inputs a similarity matrix, that is it skips the correlation calculation step but is otherwise identical.

**Value**

Adjacency matrix of dimensions `ncol(datExpr)` times `ncol(datExpr)` (or the same dimensions as `similarity`). If `selectCols` was given, the number of columns will be the length (if numeric) or sum (if boolean) of `selectCols`.



**Note**

When calculated from the `datExpr`, the network is always calculated among the columns of `datExpr` irrespective of whether a correlation or a distance network is requested.

**Author(s)**

Peter Langfelder and Steve Horvath

**References**

Bin Zhang and Steve Horvath (2005) A General Framework for Weighted Gene Co-Expression Network Analysis, *Statistical Applications in Genetics and Molecular Biology*, Vol. 4 No. 1, Article 17

Langfelder P, Horvath S (2007) Eigengene networks for studying the relationships between co-expression modules. *BMC Systems Biology* 2007, 1:54

---

adjacency.polyReg      *Adjacency matrix based on polynomial regression*

---

**Description**

`adjacency.polyReg` calculates a network adjacency matrix by fitting polynomial regression models to pairs of variables (i.e. pairs of columns from `datExpr`). Each polynomial fit results in a model fitting index `R.squared`. Thus, the `n` columns of `datExpr` result in an `n x n` dimensional matrix whose entries contain `R.squared` measures. This matrix is typically non-symmetric. To arrive at a (symmetric) adjacency matrix, one can specify different symmetrization methods with `symmetrizationMethod`.

**Usage**

```
adjacency.polyReg(datExpr, degree=3, symmetrizationMethod = "mean")
```

**Arguments**

<code>datExpr</code>	data frame containing numeric variables. Example: Columns may correspond to genes and rows to observations (samples).
<code>degree</code>	the degree of the polynomial. Must be less than the number of unique points.
<code>symmetrizationMethod</code>	character string (eg "none", "min", "max", "mean") that specifies the method used to symmetrize the pairwise model fitting index matrix (see details).

**Details**

A network adjacency matrix is a symmetric matrix whose entries lie between 0 and 1. It is a special case of a similarity matrix. Each variable (column of `datExpr`) is regressed on every other variable, with each model fitting index recorded in a square matrix. Note that the model fitting index of regressing variable `x` and variable `y` is usually different from that of regressing `y` on `x`. From the polynomial regression model `glm(y ~ poly(x,degree))` one can calculate the model fitting index `R.squared(y,x)`. `R.squared(y,x)` is a number between 0 and 1. The closer it is to 1, the better the polynomial describes the relationship between `x` and `y` and the more significant is the pairwise relationship between the 2 variables. One can also reverse the roles of `x` and `y` to arrive at a model fitting

index  $R.squared(x,y)$ . If  $degree > 1$  then  $R.squared(x,y)$  is typically different from  $R.squared(y,x)$ . Assume a set of  $n$  variables  $x_1, \dots, x_n$  (corresponding to the columns of `datExpr` then one can define  $R.squared(x_i, x_j)$ . The model fitting indices for the elements of an  $n \times n$  dimensional matrix ( $R.squared(i,j)$ ). `symmetrizationMethod` implements the following symmetrization methods:  $A.min(i,j) = \min(R.squared(i,j), R.squared(j,i))$ ,  $A.ave(i,j) = (R.squared(i,j) + R.squared(j,i)) / 2$ ,  $A.max(i,j) = \max(R.squared(i,j), R.squared(j,i))$ .

### Value

An adjacency matrix of dimensions `ncol(datExpr)` times `ncol(datExpr)`.

### Author(s)

Lin Song, Steve Horvath

### References

Song L, Langfelder P, Horvath S Avoiding mutual information based co-expression measures (to appear).

Horvath S (2011) Weighted Network Analysis. Applications in Genomics and Systems Biology. Springer Book. ISBN: 978-1-4419-8818-8

### See Also

For more information about polynomial regression, please refer to functions [poly](#) and [glm](#)

### Examples

```
#Simulate a data frame datE which contains 5 columns and 50 observations
m=50
x1=rnorm(m)
r=.5; x2=r*x1+sqrt(1-r^2)*rnorm(m)
r=.3; x3=r*(x1-.5)^2+sqrt(1-r^2)*rnorm(m)
x4=rnorm(m)
r=.3; x5=r*x4+sqrt(1-r^2)*rnorm(m)
datE=data.frame(x1,x2,x3,x4,x5)
#calculate adjacency by symmetrizing using max
A.max=adjacency.polyReg(datE, symmetrizationMethod="max")
A.max
#calculate adjacency by symmetrizing using mean
A.mean=adjacency.polyReg(datE, symmetrizationMethod="mean")
A.mean
# output the unsymmetrized pairwise model fitting indices R.squared
R.squared=adjacency.polyReg(datE, symmetrizationMethod="none")
R.squared
```

## Description

adjacency.splineReg calculates a network adjacency matrix by fitting spline regression models to pairs of variables (i.e. pairs of columns from datExpr). Each spline regression model results in a fitting index R.squared. Thus, the n columns of datExpr result in an n x n dimensional matrix whose entries contain R.squared measures. This matrix is typically non-symmetric. To arrive at a (symmetric) adjacency matrix, one can specify different symmetrization methods with symmetrizationMethod.

## Usage

```
adjacency.splineReg(
  datExpr,
  df = 6-(nrow(datExpr)<100)-(nrow(datExpr)<30),
  symmetrizationMethod = "mean",
  ...)
```

## Arguments

datExpr	data frame containing numeric variables. Example: Columns may correspond to genes and rows to observations (samples).
df	degrees of freedom in generating natural cubic spline. The default is as follows: if nrow(datExpr)>100 use 6, if nrow(datExpr)>30 use 4, otherwise use 5.
symmetrizationMethod	character string (eg "none", "min", "max", "mean") that specifies the method used to symmetrize the pairwise model fitting index matrix (see details).
...	other arguments from function <a href="#">ns</a>

## Details

A network adjacency matrix is a symmetric matrix whose entries lie between 0 and 1. It is a special case of a similarity matrix. Each variable (column of datExpr) is regressed on every other variable, with each model fitting index recorded in a square matrix. Note that the model fitting index of regressing variable x and variable y is usually different from that of regressing y on x. From the spline regression model  $\text{glm}(y \sim \text{ns}(x, \text{df}))$  one can calculate the model fitting index  $R.\text{squared}(y,x)$ .  $R.\text{squared}(y,x)$  is a number between 0 and 1. The closer it is to 1, the better the spline regression model describes the relationship between x and y and the more significant is the pairwise relationship between the 2 variables. One can also reverse the roles of x and y to arrive at a model fitting index  $R.\text{squared}(x,y)$ .  $R.\text{squared}(x,y)$  is typically different from  $R.\text{squared}(y,x)$ . Assume a set of n variables  $x_1, \dots, x_n$  (corresponding to the columns of datExpr) then one can define  $R.\text{squared}(x_i, x_j)$ . The model fitting indices for the elements of an n x n dimensional matrix ( $R.\text{squared}(ij)$ ). symmetrizationMethod implements the following symmetrization methods:  $A.\text{min}(ij) = \min(R.\text{squared}(ij), R.\text{squared}(ji))$ ,  $A.\text{ave}(ij) = (R.\text{squared}(ij) + R.\text{squared}(ji)) / 2$ ,  $A.\text{max}(ij) = \max(R.\text{squared}(ij), R.\text{squared}(ji))$ . For more information about natural cubic spline regression, please refer to functions "ns" and "glm".

## Value

An adjacency matrix of dimensions  $\text{ncol}(\text{datExpr})$  times  $\text{ncol}(\text{datExpr})$ .

## Author(s)

Lin Song, Steve Horvath

## References

Song L, Langfelder P, Horvath S Avoiding mutual information based co-expression measures (to appear).

Horvath S (2011) Weighted Network Analysis. Applications in Genomics and Systems Biology. Springer Book. ISBN: 978-1-4419-8818-8

## See Also

[ns](#), [glm](#)

## Examples

```
#Simulate a data frame datE which contains 5 columns and 50 observations
m=50
x1=rnorm(m)
r=.5; x2=r*x1+sqrt(1-r^2)*rnorm(m)
r=.3; x3=r*(x1-.5)^2+sqrt(1-r^2)*rnorm(m)
x4=rnorm(m)
r=.3; x5=r*x4+sqrt(1-r^2)*rnorm(m)
datE=data.frame(x1,x2,x3,x4,x5)
#calculate adjacency by symmetrizing using max
A.max=adjacency.splineReg(datE, symmetrizationMethod="max")
A.max
#calculate adjacency by symmetrizing using mean
A.mean=adjacency.splineReg(datE, symmetrizationMethod="mean")
A.mean
# output the unsymmetrized pairwise model fitting indices R.squared
R.squared=adjacency.splineReg(datE, symmetrizationMethod="none")
R.squared
```

---

AFcorMI

*Prediction of Weighted Mutual Information Adjacency Matrix by Correlation*

---

## Description

AFcorMI computes a predicted weighted mutual information adjacency matrix from a given correlation matrix.

## Usage

```
AFcorMI(r, m)
```

## Arguments

**r** symmetric correlation matrix with values from -1 to 1.  
**m** number of observations from which the correlation was calculated.

## Details

This function is a one-to-one prediction when we consider correlation as unsigned. The prediction corresponds to the `AdjacencyUniversalVersion2` discussed in the help file for the function [mutualInfoAdjacency](#). For more information about the generation and features of the predicted mutual information adjacency, please refer to the function [mutualInfoAdjacency](#).

**Value**

A matrix with the same size as the input correlation matrix, containing the predicted mutual information of type AdjacencyUniversalVersion2.

**Author(s)**

Steve Horvath, Lin Song, Peter Langfelder

**See Also**

[mutualInfoAdjacency](#)

**Examples**

```
#Simulate a data frame datE which contains 5 columns and 50 observations
m=50
x1=rnorm(m)
r=.5; x2=r*x1+sqrt(1-r^2)*rnorm(m)
r=.3; x3=r*(x1-.5)^2+sqrt(1-r^2)*rnorm(m)
x4=rnorm(m)
r=.3; x5=r*x4+sqrt(1-r^2)*rnorm(m)
datE=data.frame(x1,x2,x3,x4,x5)
#calculate predicted AUV2
cor.data=cor(datE, use="p")
AUV2=AFcorMI(r=cor.data, m=nrow(datE))
```

---

alignExpr

*Align expression data with given vector*


---

**Description**

Multiplies genes (columns) in given expression data such that their correlation with given reference vector is non-negative.

**Usage**

```
alignExpr(datExpr, y = NULL)
```

**Arguments**

datExpr	expression data to be aligned. A data frame with columns corresponding to genes and rows to samples.
y	reference vector of length equal the number of samples (rows) in datExpr

**Details**

The function basically multiplies each column in datExpr by the sign of its correlation with y. If y is not given, the first column in datExpr will be used as the reference vector.

**Value**

A data frame containing the aligned expression data, of the same dimensions as the input data frame.

**Author(s)**

Steve Horvath and Peter Langfelder

---

allocateJobs

*Divide tasks among workers*

---

**Description**

This function calculates an even splitting of a given number of tasks among a given number of workers (threads).

**Usage**

```
allocateJobs(nTasks, nWorkers)
```

**Arguments**

nTasks	number of tasks to be divided
nWorkers	number of workers

**Details**

Tasks are labeled consecutively 1,2,..., nTasks. The tasks are split in contiguous blocks as evenly as possible.

**Value**

A list with one component per worker giving the task indices to be worked on by each worker. If there are more workers than tasks, the tasks for the extra workers are 0-length numeric vectors.

**Author(s)**

Peter Langfelder

**Examples**

```
allocateJobs(10, 3);  
allocateJobs(2,4);
```

---

allowWGCNAThreads	<i>Allow and disable multi-threading for certain WGCNA calculations</i>
-------------------	---

---

### Description

These functions allow and disable multi-threading for WGCNA calculations that can optionally be multi-threaded, which includes all functions using `cor` or `bicor` functions.

### Usage

```
allowWGCNAThreads(nThreads = NULL)
```

```
enableWGCNAThreads(nThreads = NULL)
```

```
disableWGCNAThreads()
```

```
WGCNANThreads()
```

### Arguments

nThreads	Number of threads to allow. If not given, the number of processors online (as reported by system configuration) will be used. There appear to be some cases where the automatically-determined number is wrong; please check the output to see that the number of threads makes sense. Except for testing and/or torturing your system, the number of threads should be no more than the number of actual processors/cores.
----------	---

### Details

`allowWGCNAThreads` enables parallel calculation within the compiled code in WGCNA, principally for calculation of correlations in the presence of missing data. This function is now deprecated; use `enableWGCNAThreads` instead.

`enableWGCNAThreads` enables parallel calculations within user-level R functions as well as within the compiled code, and registers an appropriate parallel calculation back-end for the operating system/platform.

`disableWGCNAThreads` disables parallel processing.

`WGCNANThreads` returns the number of threads (parallel processes) that WGCNA is currently configured to run with.

### Value

`allowWGCNAThreads`, `enableWGCNAThreads`, and `disableWGCNAThreads` return the maximum number of threads WGCNA calculations will be allowed to use.

### Note

Multi-threading within compiled code is not available on Windows; R code parallelization works on all platforms.

**Author(s)**

Peter Langfelder

automaticNetworkScreening

*One-step automatic network gene screening***Description**

This function performs gene screening based on a given trait and gene network properties

**Usage**

```
automaticNetworkScreening(
  datExpr,
  y,
  power = 6,
  networkType = "unsigned",
  detectCutHeight = 0.995,
  minModuleSize = min(20, ncol(as.matrix(datExpr))/2),
  datME = NULL,
  getQValues = TRUE,
  ...)
```

**Arguments**

datExpr	data frame containing the expression data, columns corresponding to genes and rows to samples
y	vector containing trait values for all samples in datExpr
power	soft thresholding power used in network construction
networkType	character string specifying network type. Allowed values are (unique abbreviations of) "unsigned", "signed", "hybrid".
detectCutHeight	cut height of the gene hierarchical clustering dendrogram. See <code>cutreeDynamic</code> for details.
minModuleSize	minimum module size to be used in module detection procedure.
datME	optional specification of module eigengenes. A data frame whose columns are the module eigengenes. If given, module analysis will not be performed.
getQValues	logical: should q-values (local FDR) be calculated?
...	other arguments to the module identification function <a href="#">blockwiseModules</a>

**Details**

Network screening is a method for identifying genes that have a high gene significance and are members of important modules at the same time. If `datME` is given, the function calls [networkScreening](#) with the default parameters. If `datME` is not given, module eigengenes are first calculated using network analysis based on supplied parameters.



**Value**

A list with the following components:

networkScreening	a data frame containing results of the network screening procedure. See <a href="#">networkScreening</a> for more details.
datME	calculated module eigengenes (or a copy of the input datME, if given).
hubGeneSignificance	hub gene significance for all calculated modules. See <a href="#">hubGeneSignificance</a> .

**Author(s)**

Steve Horvath

**See Also**

[networkScreening](#), [hubGeneSignificance](#), [networkScreening](#), [cutreeDynamic](#)

---

automaticNetworkScreeningGS

*One-step automatic network gene screening with external gene significance*

---

**Description**

This function performs gene screening based on external gene significance and their network properties.

**Usage**

```
automaticNetworkScreeningGS(
  datExpr, GS,
  power = 6, networkType = "unsigned",
  detectCutHeight = 0.995, minModuleSize = min(20, ncol(as.matrix(datExpr))/2),
  datME = NULL)
```

**Arguments**

datExpr	data frame containing the expression data, columns corresponding to genes and rows to samples
GS	vector containing gene significance for all genes given in datExpr
power	soft thresholding power used in network construction
networkType	character string specifying network type. Allowed values are (unique abbreviations of) "unsigned", "signed", "hybrid".
detectCutHeight	cut height of the gene hierarchical clustering dendrogram. See <a href="#">cutreeDynamic</a> for details.
minModuleSize	minimum module size to be used in module detection procedure.
datME	optional specification of module eigengenes. A data frame whose columns are the module eigengenes. If given, module analysis will not be performed.

## Details

Network screening is a method for identifying genes that have a high gene significance and are members of important modules at the same time. If `datME` is given, the function calls [networkScreeningGS](#) with the default parameters. If `datME` is not given, module eigengenes are first calculated using network analysis based on supplied parameters.

## Value

A list with the following components:

`networkScreening`  
a data frame containing results of the network screening procedure. See [networkScreeningGS](#) for more details.

`datME`  
calculated module eigengenes (or a copy of the input `datME`, if given).

`hubGeneSignificance`  
hub gene significance for all calculated modules. See [hubGeneSignificance](#).

## Author(s)

Steve Horvath

## See Also

[networkScreening](#), [hubGeneSignificance](#), [networkScreening](#), [cutreeDynamic](#)

---

`bicor`

*Biweight Midcorrelation*

---

## Description

Calculate biweight midcorrelation efficiently for matrices.

## Usage

```
bicor(x, y = NULL,  
      robustX = TRUE, robustY = TRUE,  
      use = "all.obs",  
      maxPOutliers = 1,  
      quick = 0,  
      pearsonFallback = "individual",  
      cosine = FALSE,  
      cosineX = cosine,  
      cosineY = cosine,  
      nThreads = 0,  
      verbose = 0, indent = 0)
```

**Arguments**

x	a vector or matrix-like numeric object
y	a vector or matrix-like numeric object
robustX	use robust calculation for x?
robustY	use robust calculation for y?
use	specifies handling of NAs. One of (unique abbreviations of) "all.obs", "pairwise.complete.obs".
maxPOutliers	specifies the maximum percentile of data that can be considered outliers on either side of the median separately. For each side of the median, if higher percentile than maxPOutliers is considered an outlier by the weight function based on $9 \cdot \text{mad}(x)$ , the width of the weight function is increased such that the percentile of outliers on that side of the median equals maxPOutliers. Using maxPOutliers=1 will effectively disable all weight function broadening; using maxPOutliers=0 will give results that are quite similar (but not equal to) Pearson correlation.
quick	real number between 0 and 1 that controls the handling of missing data in the calculation of correlations. See details.
pearsonFallback	Specifies whether the bicor calculation should revert to Pearson when median absolute deviation (mad) is zero. Recognized values are (abbreviations of) "none", "individual", "all". If set to "none", zero mad will result in NA for the corresponding correlation. If set to "individual", Pearson calculation will be used only for columns that have zero mad. If set to "all", the presence of a single zero mad will cause the whole variable to be treated in Pearson correlation manner (as if the corresponding robust option was set to FALSE).
cosine	logical: calculate cosine biweight midcorrelation? Cosine bicorrelation is similar to standard bicorrelation but the median subtraction is not performed.
cosineX	logical: use the cosine calculation for x? This setting does not affect y and can be used to give a hybrid cosine-standard bicorrelation.
cosineY	logical: use the cosine calculation for y? This setting does not affect x and can be used to give a hybrid cosine-standard bicorrelation.
nThreads	non-negative integer specifying the number of parallel threads to be used by certain parts of correlation calculations. This option only has an effect on systems on which a POSIX thread library is available (which currently includes Linux and Mac OSX, but excludes Windows). If zero, the number of online processors will be used if it can be determined dynamically, otherwise correlation calculations will use 2 threads.
verbose	if non-zero, the underlying C function will print some diagnostics.
indent	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

**Details**

This function implements biweight midcorrelation calculation (see references). If y is not supplied, midcorrelation of columns of x will be calculated; otherwise, the midcorrelation between columns of x and y will be calculated. Thus, `bicor(x)` is equivalent to `bicor(x, x)` but is more efficient.

The options `robustX`, `robustY` allow the user to revert the calculation to standard correlation calculation. This is important, for example, if any of the variables is binary (or, more generally, discrete)

as in such cases the robust methods produce meaningless results. If both `robustX`, `robustY` are set to `FALSE`, the function calculates the standard Pearson correlation (but is slower than the function `cor`).

The argument `quick` specifies the precision of handling of missing data in the correlation calculations. Value `quick = 0` will cause all calculations to be executed accurately, which may be significantly slower than calculations without missing data. Progressively higher values will speed up the calculations but introduce progressively larger errors. Without missing data, all column medians and median absolute deviations (MADs) can be pre-calculated before the covariances are calculated. When missing data are present, exact calculations require the column medians and MADs to be calculated for each covariance. The approximate calculation uses the pre-calculated median and MAD and simply ignores missing data in the covariance calculation. If the number of missing data is high, the pre-calculated medians and MADs may be very different from the actual ones, thus potentially introducing large errors. The `quick` value times the number of rows specifies the maximum difference in the number of missing entries for median and MAD calculations on the one hand and covariance on the other hand that will be tolerated before a recalculation is triggered. The hope is that if only a few missing data are treated approximately, the error introduced will be small but the potential speedup can be significant.

The choice `"all"` for `pearsonFallback` is not fully implemented in the sense that there are rare but possible cases in which the calculation is equivalent to `"individual"`. This may happen if the `use` option is set to `"pairwise.complete.obs"` and the missing data are arranged such that each individual mad is non-zero, but when two columns are analyzed together, the missing data from both columns may make a mad zero. In such a case, the calculation is treated as Pearson, but other columns will be treated as bicor.

### Value

A matrix of biweight midcorrelations. Dimnames on the result are set appropriately.

### Author(s)

Peter Langfelder

### References

- Peter Langfelder, Steve Horvath (2012) Fast R Functions for Robust Correlations and Hierarchical Clustering. *Journal of Statistical Software*, 46(11), 1-17. <http://www.jstatsoft.org/v46/i11/>
- "Dealing with Outliers in Bivariate Data: Robust Correlation", Rich Herrington, <http://www.unt.edu/benchmarks/archives>
- "Introduction to Robust Estimation and Hypothesis Testing", Rand Wilcox, Academic Press, 1997.
- "Data Analysis and Regression: A Second Course in Statistics", Mosteller and Tukey, Addison-Wesley, 1977, pp. 203-209.

---

bicorAndPvalue

*Calculation of biweight midcorrelations and associated p-values*

---

### Description

A faster, one-step calculation of Student correlation p-values for multiple biweight midcorrelations, properly taking into account the actual number of observations.

**Usage**

```
bicorAndPvalue(x, y = NULL,
               use = "pairwise.complete.obs",
               alternative = c("two.sided", "less", "greater"),
               ...)
```

**Arguments**

x	a vector or a matrix
y	a vector or a matrix. If NULL, the correlation of columns of x will be calculated.
use	determines handling of missing data. See <a href="#">bicor</a> for details.
alternative	specifies the alternative hypothesis and must be (a unique abbreviation of) one of "two.sided", "greater" or "less". the initial letter. "greater" corresponds to positive association, "less" to negative association.
...	other arguments to the function <a href="#">bicor</a> .

**Details**

The function calculates the biweight midcorrelations of a matrix or of two matrices and the corresponding Student p-values. The output is not as full-featured as [cor.test](#), but can work with matrices as input.

**Value**

A list with the following components, each a matrix:

bicor	the calculated correlations
p	the Student p-values corresponding to the calculated correlations
Z	Fisher transform of the calculated correlations
t	Student t statistics of the calculated correlations
nObs	Numbers of observations for the correlation, p-values etc.

**Author(s)**

Peter Langfelder and Steve Horvath

**References**

Peter Langfelder, Steve Horvath (2012) Fast R Functions for Robust Correlations and Hierarchical Clustering. *Journal of Statistical Software*, 46(11), 1-17. <http://www.jstatsoft.org/v46/i11/>

**See Also**

[bicor](#) for calculation of correlations only;

[cor.test](#) for another function for significance test of correlations

**Examples**

```
# generate random data with non-zero correlation
set.seed(1);
a = rnorm(100);
b = rnorm(100) + a;
x = cbind(a, b);
# Call the function and display all results
bicorAndPvalue(x)
# Set some components to NA
x[c(1:4), 1] = NA
corAndPvalue(x)
# Note that changed number of observations.
```

---

blockSize	<i>Attempt to calculate an appropriate block size to maximize efficiency of block-wise calculations.</i>
-----------	--

---

**Description**

The function uses a rather primitive way to estimate available memory and use it to suggest a block size appropriate for the many block-by-block calculations in this package.

**Usage**

```
blockSize(
  matrixSize,
  rectangularBlocks = TRUE,
  maxMemoryAllocation = NULL,
  overheadFactor = 3)
```

**Arguments**

matrixSize	the relevant dimension (usually the number of columns) of the matrix that is to be operated on block-by-block.
rectangularBlocks	logical indicating whether the blocks of data are rectangular (of size blockSize times matrixSize) or square (of size blockSize times blockSize).
maxMemoryAllocation	maximum desired memory allocation, in bytes. Should not exceed 2GB or total installed RAM (whichever is greater) on 32-bit systems, while on 64-bit systems it should not exceed the total installed RAM. If not supplied, the available memory will be estimated internally.
overheadFactor	overhead factor for the memory use by R. Recommended values are between 2 (for simple calculations) and 4 or more for complicated calculations where intermediate results (for which R must also allocate memory) take up a lot of space.

## Details

Multiple functions within the WGCNA package use a divide-and-conquer (also known as block-by-block, or block-wise) approach to handling large data sets. This function is meant to assist in choosing a suitable block size, given the size of the data and the available memory.

If the entire expected result fits into the allowed memory (after taking into account the expected overhead), the returned block size will equal the input `matrixSize`.

The internal estimation of available memory works by returning the size of largest successfully allocated block of memory. It is hoped that this will lead to reasonable results but some operating systems may actually allocate more than is available. It is therefore preferable that the user specifies the available memory by hand.

## Value

A single integer giving the suggested block size, or `matrixSize` if the entire calculation is expected to fit into memory in one piece.

## Author(s)

Peter Langfelder

## Examples

```
# Suitable blocks for handling 30,000 genes within 2GB (=2^31 bytes) of memory
blockSize(30000, rectangularBlocks = TRUE, maxMemoryAllocation = 2^31)
```

---

`blockwiseConsensusModules`

*Find consensus modules across several datasets.*

---

## Description

Perform network construction and consensus module detection across several datasets.

## Usage

```
blockwiseConsensusModules(
  multiExpr,

  # Data checking options

  checkMissingData = TRUE,

  # Blocking options

  blocks = NULL,
  maxBlockSize = 5000,
  randomSeed = 12345,

  # TOM precalculation arguments, if available

  individualTOMInfo = NULL,
```

```
useIndivTOMSubset = NULL,

# Network construction arguments: correlation options

corType = "pearson",
maxPOutliers = 1,
quickCor = 0,
pearsonFallback = "individual",
cosineCorrelation = FALSE,

# Adjacency function options

power = 6,
networkType = "unsigned",
checkPower = TRUE,

# Topological overlap options

TOMType = "unsigned",
TOMDenom = "min",

# Save individual TOMs?

saveIndividualTOMs = TRUE,
individualTOMFileNames = "individualTOM-Set%s-Block%b.RData",

# Consensus calculation options

consensusQuantile = 0,
scaleTOMs = TRUE, scaleQuantile = 0.95,

# Sampling for scaling (speeds up calculation)

sampleForScaling = TRUE, sampleForScalingFactor = 1000,
getTOMScalingSamples = FALSE,

# Returning the consensus TOM

saveTOMs = FALSE,
consensusTOMFileNames = "consensusTOM-block.%b.RData",

# Internal handling of TOMs

useDiskCache = TRUE, chunkSize = NULL,
cacheBase = ".blockConsModsCache",

# Basic tree cut options

deepSplit = 2,
detectCutHeight = 0.995, minModuleSize = 20,
checkMinModuleSize = TRUE,

# Advanced tree cut options
```



```

maxCoreScatter = NULL, minGap = NULL,
maxAbsCoreScatter = NULL, minAbsGap = NULL,
pamStage = TRUE, pamRespectsDendro = TRUE,

# Gene reassignment and trimming from a module, and module "significance" criteria

reassignThresholdPS = 1e-4,
trimmingConsensusQuantile = consensusQuantile,
minCoreKME = 0.5, minCoreKMESize = minModuleSize/3,
minKMEtoStay = 0.2,

# Module eigengene calculation options

impute = TRUE,
trapErrors = FALSE,

#Module merging options

mergeCutHeight = 0.15,
mergeConsensusQuantile = consensusQuantile,

# Output options

numericLabels = FALSE,

# General options

nThreads = 0,
verbose = 2, indent = 0)

```

### Arguments

<code>multiExpr</code>	expression data in the multi-set format (see <a href="#">checkSets</a> ). A vector of lists, one per set. Each set must contain a component data that contains the expression data, with rows corresponding to samples and columns to genes or probes.
<code>checkMissingData</code>	logical: should data be checked for excessive numbers of missing entries in genes and samples, and for genes with zero variance? See details.
<code>blocks</code>	optional specification of blocks in which hierarchical clustering and module detection should be performed. If given, must be a numeric vector with one entry per gene of <code>multiExpr</code> giving the number of the block to which the corresponding gene belongs.
<code>maxBlockSize</code>	integer giving maximum block size for module detection. Ignored if <code>blocks</code> above is non-NULL. Otherwise, if the number of genes in <code>datExpr</code> exceeds <code>maxBlockSize</code> , genes will be pre-clustered into blocks whose size should not exceed <code>maxBlockSize</code> .
<code>randomSeed</code>	integer to be used as seed for the random number generator before the function starts. If a current seed exists, it is saved and restored upon exit. If NULL is given, the function will not save and restore the seed.
<code>individualTOMInfo</code>	Optional data for TOM matrices in individual data sets. This object is returned

by the function `blockwiseIndividualTOMs`. If not given, appropriate topological overlaps will be calculated using the network construction options below.

<code>useIndivTOMSubset</code>	If <code>individualTOMInfo</code> is given, this argument allows to only select a subset of the individual set networks contained in <code>individualTOMInfo</code> . It should be a numeric vector giving the indices of the individual sets to be used. Note that this argument is NOT applied to <code>multiExpr</code> .
<code>corType</code>	character string specifying the correlation to be used. Allowed values are (unique abbreviations of) "pearson" and "bicor", corresponding to Pearson and bid-weight midcorrelation, respectively. Missing values are handled using the <code>parwise.complete.obs</code> option.
<code>maxPOutliers</code>	only used for <code>corType=="bicor"</code> . Specifies the maximum percentile of data that can be considered outliers on either side of the median separately. For each side of the median, if higher percentile than <code>maxPOutliers</code> is considered an outlier by the weight function based on $9 \times \text{mad}(x)$ , the width of the weight function is increased such that the percentile of outliers on that side of the median equals <code>maxPOutliers</code> . Using <code>maxPOutliers=1</code> will effectively disable all weight function broadening; using <code>maxPOutliers=0</code> will give results that are quite similar (but not equal to) Pearson correlation.
<code>quickCor</code>	real number between 0 and 1 that controls the handling of missing data in the calculation of correlations. See details.
<code>pearsonFallback</code>	Specifies whether the bicor calculation, if used, should revert to Pearson when median absolute deviation ( <code>mad</code> ) is zero. Recognized values are (abbreviations of) "none", "individual", "all". If set to "none", zero <code>mad</code> will result in NA for the corresponding correlation. If set to "individual", Pearson calculation will be used only for columns that have zero <code>mad</code> . If set to "all", the presence of a single zero <code>mad</code> will cause the whole variable to be treated in Pearson correlation manner (as if the corresponding robust option was set to FALSE). Has no effect for Pearson correlation. See <code>bicor</code> .
<code>cosineCorrelation</code>	logical: should the cosine version of the correlation calculation be used? The cosine calculation differs from the standard one in that it does not subtract the mean.
<code>power</code>	soft-thresholding power for network construction.
<code>networkType</code>	network type. Allowed values are (unique abbreviations of) "unsigned", "signed", "signed hybrid". See <code>adjacency</code> .
<code>checkPower</code>	logical: should basic sanity check be performed on the supplied power? If you would like to experiment with unusual powers, set the argument to FALSE and proceed with caution.
<code>TOMType</code>	one of "none", "unsigned", "signed". If "none", adjacency will be used for clustering. If "unsigned", the standard TOM will be used (more generally, TOM function will receive the adjacency as input). If "signed", TOM will keep track of the sign of correlations between neighbors.
<code>TOMDenom</code>	a character string specifying the TOM variant to be used. Recognized values are "min" giving the standard TOM described in Zhang and Horvath (2005), and "mean" in which the <code>min</code> function in the denominator is replaced by <code>mean</code> . The "mean" may produce better results but at this time should be considered experimental.

saveIndividualTOMs	logical: should individual TOMs be saved to disk for later use?
individualTOMFileNames	character string giving the file names to save individual TOMs into. The following tags should be used to make the file names unique for each set and block: %s will be replaced by the set number; %N will be replaced by the set name (taken from names(multiExpr)) if it exists, otherwise by set number; %b will be replaced by the block number. If the file names turn out to be non-unique, an error will be generated.
consensusQuantile	quantile at which consensus is to be defined. See details.
scaleTOMs	should set-specific TOM matrices be scaled to the same scale?
scaleQuantile	if scaleTOMs is TRUE, topological overlaps (or adjacencies if TOMs are not computed) will be scaled such that their scaleQuantile quantiles will agree.
sampleForScaling	if TRUE, scale quantiles will be determined from a sample of network similarities. Note that using all data can double the memory footprint of the function and the function may fail.
sampleForScalingFactor	determines the number of samples for scaling: the number is $1/\text{scaleQuantile} * \text{sampleForScaling}$ . Should be set well above 1 to ensure accuracy of the sampled quantile.
getTOMScalingSamples	logical: should samples used for TOM scaling be saved for future analysis? This option is only available when sampleForScaling is TRUE.
saveTOMs	logical: should the consensus topological overlap matrices for each block be saved and returned?
consensusTOMFileNames	character string containing the file names containing the consensus topological overlaps. The tag %b will be replaced by the block number. If the resulting file names are non-unique (for example, because the user gives a file name without a %b tag), an error will be generated. These files are standard R data files and can be loaded using the <a href="#">load</a> function.
useDiskCache	should calculated network similarities in individual sets be temporarily saved to disk? Saving to disk is somewhat slower than keeping all data in memory, but for large blocks and/or many sets the memory footprint may be too big.
chunkSize	network similarities are saved in smaller chunks of size chunkSize.
cacheBase	character string containing the desired name for the cache files. The actual file names will consist of cacheBase and a suffix to make the file names unique.
deepSplit	integer value between 0 and 4. Provides a simplified control over how sensitive module detection should be to module splitting, with 0 least and 4 most sensitive. See <a href="#">cutreeDynamic</a> for more details.
detectCutHeight	dendrogram cut height for module detection. See <a href="#">cutreeDynamic</a> for more details.
minModuleSize	minimum module size for module detection. See <a href="#">cutreeDynamic</a> for more details.
checkMinModuleSize	logical: should sanity checks be performed on minModuleSize?

maxCoreScatter	maximum scatter of the core for a branch to be a cluster, given as the fraction of cutHeight relative to the 5th percentile of joining heights. See <a href="#">cutreeDynamic</a> for more details.
minGap	minimum cluster gap given as the fraction of the difference between cutHeight and the 5th percentile of joining heights. See <a href="#">cutreeDynamic</a> for more details.
maxAbsCoreScatter	maximum scatter of the core for a branch to be a cluster given as absolute heights. If given, overrides maxCoreScatter. See <a href="#">cutreeDynamic</a> for more details.
minAbsGap	minimum cluster gap given as absolute height difference. If given, overrides minGap. See <a href="#">cutreeDynamic</a> for more details.
pamStage	logical. If TRUE, the second (PAM-like) stage of module detection will be performed. See <a href="#">cutreeDynamic</a> for more details.
pamRespectsDendro	Logical, only used when pamStage is TRUE. If TRUE, the PAM stage will respect the dendrogram in the sense an object can be PAM-assigned only to clusters that lie below it on the branch that the object is merged into. See <a href="#">cutreeDynamic</a> for more details.
reassignThresholdPS	per-set p-value ratio threshold for reassigning genes between modules. See Details.
trimmingConsensusQuantile	a number between 0 and 1 specifying the consensus quantile used for kME calculation that determines module trimming according to the arguments below.
minCoreKME	a number between 0 and 1. If a detected module does not have at least minModuleKMESize genes with eigengene connectivity at least minCoreKME, the module is disbanded (its genes are unlabeled and returned to the pool of genes waiting for module detection).
minCoreKMESize	see minCoreKME above.
minKMEtoStay	genes whose eigengene connectivity to their module eigengene is lower than minKMEtoStay are removed from the module.
impute	logical: should imputation be used for module eigengene calculation? See <a href="#">moduleEigengenes</a> for more details.
trapErrors	logical: should errors in calculations be trapped?
mergeCutHeight	dendrogram cut height for module merging.
mergeConsensusQuantile	consensus quantile for module merging. See <a href="#">mergeCloseModules</a> for details.
numericLabels	logical: should the returned modules be labeled by colors (FALSE), or by numbers (TRUE)?
nThreads	non-negative integer specifying the number of parallel threads to be used by certain parts of correlation calculations. This option only has an effect on systems on which a POSIX thread library is available (which currently includes Linux and Mac OSX, but excludes Windows). If zero, the number of online processors will be used if it can be determined dynamically, otherwise correlation calculations will use 2 threads.
verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.
indent	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

## Details

The function starts by optionally filtering out samples that have too many missing entries and genes that have either too many missing entries or zero variance in at least one set. Genes that are filtered out are left unassigned by the module detection. Returned eigengenes will contain NA in entries corresponding to filtered-out samples.

If `blocks` is not given and the number of genes exceeds `maxBlockSize`, genes are pre-clustered into blocks using the function `consensusProjectiveKMeans`; otherwise all genes are treated in a single block.

For each block of genes, the network is constructed and (if requested) topological overlap is calculated in each set. To minimize memory usage, calculated topological overlaps are optionally saved to disk in chunks until they are needed again for the calculation of the consensus network topological overlap. If requested, the consensus topological overlaps are saved to disk for later use. Genes are then clustered using average linkage hierarchical clustering and modules are identified in the resulting dendrogram by the Dynamic Hybrid tree cut. Found modules are trimmed of genes whose consensus module membership `kME` (that is, correlation with module eigengene) is less than `minKMEtoStay`. Modules in which fewer than `minCoreKMESize` genes have consensus `KME` higher than `minCoreKME` are disbanded, i.e., their constituent genes are pronounced unassigned.

After all blocks have been processed, the function checks whether there are genes whose `KME` in the module they assigned is lower than `KME` to another module. If `p-values` of the higher correlations are smaller than those of the native module by the factor `reassignThresholdPS` (in every set), the gene is re-assigned to the closer module.

In the last step, modules whose eigengenes are highly correlated are merged. This is achieved by clustering module eigengenes using the dissimilarity given by one minus their correlation, cutting the dendrogram at the height `mergeCutHeight` and merging all modules on each branch. The process is iterated until no modules are merged. See `mergeCloseModules` for more details on module merging.

The argument `quick` specifies the precision of handling of missing data in the correlation calculations. Zero will cause all calculations to be executed precisely, which may be significantly slower than calculations without missing data. Progressively higher values will speed up the calculations but introduce progressively larger errors. Without missing data, all column means and variances can be pre-calculated before the covariances are calculated. When missing data are present, exact calculations require the column means and variances to be calculated for each covariance. The approximate calculation uses the pre-calculated mean and variance and simply ignores missing data in the covariance calculation. If the number of missing data is high, the pre-calculated means and variances may be very different from the actual ones, thus potentially introducing large errors. The `quick` value times the number of rows specifies the maximum difference in the number of missing entries for mean and variance calculations on the one hand and covariance on the other hand that will be tolerated before a recalculation is triggered. The hope is that if only a few missing data are treated approximately, the error introduced will be small but the potential speedup can be significant.

## Value

A list with the following components:

<code>colors</code>	module assignment of all input genes. A vector containing either character strings with module colors (if input <code>numericLabels</code> was unset) or numeric module labels (if <code>numericLabels</code> was set to <code>TRUE</code> ). The color "grey" and the numeric label 0 are reserved for unassigned genes.
<code>unmergedColors</code>	module colors or numeric labels before the module merging step.

multiMEs	module eigengenes corresponding to the modules returned in colors, in multi-set format. A vector of lists, one per set, containing eigengenes, proportion of variance explained and other information. See <a href="#">multiSetMEs</a> for a detailed description.
goodSamples	a list, with one component per input set. Each component is a logical vector with one entry per sample from the corresponding set. The entry indicates whether the sample in the set passed basic quality control criteria.
goodGenes	a logical vector with one entry per input gene indicating whether the gene passed basic quality control criteria in all sets.
dendrograms	a list with one component for each block of genes. Each component is the hierarchical clustering dendrogram obtained by clustering the consensus gene dissimilarity in the corresponding block.
TOMfiles	if saveTOMs==TRUE, a vector of character strings, one string per block, giving the file names of files (relative to current directory) in which blockwise topological overlaps were saved.
blockGenes	a list with one component for each block of genes. Each component is a vector giving the indices (relative to the input multiExpr) of genes in the corresponding block.
blocks	if input blocks was given, its copy; otherwise a vector of length equal number of genes giving the block label for each gene. Note that block labels are not necessarily sorted in the order in which the blocks were processed (since we do not require this for the input blocks). See blockOrder below.
blockOrder	a vector giving the order in which blocks were processed and in which blockGenes above is returned. For example, blockOrder[1] contains the label of the first-processed block.
originCount	if the input consensusQuantile==0, this vector will contain counts of how many times each set contributed the consensus gene similarity value. If the counts are highly unbalanced, the consensus may be biased.
TOMScalingSamples	if the input getTOMScalingSamples is TRUE, this component is a list with one component per block. Each component is again a list with two components: sampleIndex contains indices of the distance structure in which TOM is stored that were sampled, and TOMSamples is a matrix whose rows correspond to TOM samples and columns to individual set. Hence, TOMScalingSamples[[blockNo]]\$TOMSamples[ind] contains the TOM entry that corresponds to element TOMScalingSamples[[blockNo]]\$sampleIndex of the TOM distance structure in block blockNo and set setNo. (For details on the distance structure, see <a href="#">dist</a> .)

### Note

If the input datasets have large numbers of genes, consider carefully the maxBlockSize as it significantly affects the memory footprint (and whether the function will fail with a memory allocation error). From a theoretical point of view it is advantageous to use blocks as large as possible; on the other hand, using smaller blocks is substantially faster and often the only way to work with large numbers of genes. As a rough guide, it is unlikely a standard desktop computer with 4GB memory or less will be able to work with blocks larger than 7000 genes.

### Author(s)

Peter Langfelder

## References

Langfelder P, Horvath S (2007) Eigengene networks for studying the relationships between co-expression modules. *BMC Systems Biology* 2007, 1:54

## See Also

[goodSamplesGenesMS](#) for basic quality control and filtering;  
[adjacency](#), [TOMsimilarity](#) for network construction;  
[hclust](#) for hierarchical clustering;  
[cutreeDynamic](#) for adaptive branch cutting in hierarchical clustering dendrograms;  
[mergeCloseModules](#) for merging of close modules.

---

blockwiseIndividualTOMs

*Calculation of block-wise topological overlaps*

---

## Description

Calculates topological overlaps in the given (expression) data. If the number of variables (columns) in the input data is too large, the data is first split using pre-clustering, then topological overlaps are calculated in each block.

## Usage

```
blockwiseIndividualTOMs(  
  multiExpr,  
  
  # Data checking options  
  
  checkMissingData = TRUE,  
  
  # Blocking options  
  
  blocks = NULL,  
  maxBlockSize = 5000,  
  randomSeed = 12345,  
  
  # Network construction arguments: correlation options  
  
  corType = "pearson",  
  maxPOutliers = 1,  
  quickCor = 0,  
  pearsonFallback = "individual",  
  cosineCorrelation = FALSE,  
  
  # Adjacency function options  
  
  power = 6,  
  networkType = "unsigned",
```

```

checkPower = TRUE,

# Topological overlap options

TOMType = "unsigned",
TOMDenom = "min",

# Save individual TOMs? If not, they will be returned in the session.

saveTOMs = TRUE,
individualTOMFileNames = "individualTOM-Set%s-Block%b.RData",

# General options

nThreads = 0,
verbose = 2, indent = 0)

```

### Arguments

<code>multiExpr</code>	expression data in the multi-set format (see <a href="#">checkSets</a> ). A vector of lists, one per set. Each set must contain a component data that contains the expression data, with rows corresponding to samples and columns to genes or probes.
<code>checkMissingData</code>	logical: should data be checked for excessive numbers of missing entries in genes and samples, and for genes with zero variance? See details.
<code>blocks</code>	optional specification of blocks in which hierarchical clustering and module detection should be performed. If given, must be a numeric vector with one entry per gene of <code>multiExpr</code> giving the number of the block to which the corresponding gene belongs.
<code>maxBlockSize</code>	integer giving maximum block size for module detection. Ignored if <code>blocks</code> above is non-NULL. Otherwise, if the number of genes in <code>datExpr</code> exceeds <code>maxBlockSize</code> , genes will be pre-clustered into blocks whose size should not exceed <code>maxBlockSize</code> .
<code>randomSeed</code>	integer to be used as seed for the random number generator before the function starts. If a current seed exists, it is saved and restored upon exit. If NULL is given, the function will not save and restore the seed.
<code>corType</code>	character string specifying the correlation to be used. Allowed values are (unique abbreviations of) "pearson" and "bicor", corresponding to Pearson and bid-weight midcorrelation, respectively. Missing values are handled using the <code>pariwise.complete.obs</code> option.
<code>maxPOutliers</code>	only used for <code>corType=="bicor"</code> . Specifies the maximum percentile of data that can be considered outliers on either side of the median separately. For each side of the median, if higher percentile than <code>maxPOutliers</code> is considered an outlier by the weight function based on $9 \cdot \text{mad}(x)$ , the width of the weight function is increased such that the percentile of outliers on that side of the median equals <code>maxPOutliers</code> . Using <code>maxPOutliers=1</code> will effectively disable all weight function broadening; using <code>maxPOutliers=0</code> will give results that are quite similar (but not equal to) Pearson correlation.
<code>quickCor</code>	real number between 0 and 1 that controls the handling of missing data in the calculation of correlations. See details.



pearsonFallback	Specifies whether the bicor calculation, if used, should revert to Pearson when median absolute deviation (mad) is zero. Recognized values are (abbreviations of) "none", "individual", "all". If set to "none", zero mad will result in NA for the corresponding correlation. If set to "individual", Pearson calculation will be used only for columns that have zero mad. If set to "all", the presence of a single zero mad will cause the whole variable to be treated in Pearson correlation manner (as if the corresponding robust option was set to FALSE). Has no effect for Pearson correlation. See <a href="#">bicor</a> .
cosineCorrelation	logical: should the cosine version of the correlation calculation be used? The cosine calculation differs from the standard one in that it does not subtract the mean.
power	soft-thresholding power for network construction.
networkType	network type. Allowed values are (unique abbreviations of) "unsigned", "signed", "signed hybrid". See <a href="#">adjacency</a> .
checkPower	logical: should basic sanity check be performed on the supplied power? If you would like to experiment with unusual powers, set the argument to FALSE and proceed with caution.
TOMType	one of "none", "unsigned", "signed". If "none", adjacency will be used for clustering. If "unsigned", the standard TOM will be used (more generally, TOM function will receive the adjacency as input). If "signed", TOM will keep track of the sign of correlations between neighbors. Note that the "unsigned" vs. "signed" distinction is only relevant when networkType is "unsigned". When networkType is "signed" or "signed hybrid", there is no difference between TOMType="signed" and TOMType="unsigned".
TOMDenom	a character string specifying the TOM variant to be used. Recognized values are "min" giving the standard TOM described in Zhang and Horvath (2005), and "mean" in which the min function in the denominator is replaced by mean. The "mean" may produce better results in certain special situations but at this time should be considered experimental.
saveTOMs	logical: should calculated TOMs be saved to disk (TRUE) or returned in the return value (FALSE)? Returning calculated TOMs via the return value may be more convenient but not always feasible if the matrices are too big to fit all in memory at the same time.
individualTOMFileNames	character string giving the file names to save individual TOMs into. The following tags should be used to make the file names unique for each set and block: %s will be replaced by the set number; %N will be replaced by the set name (taken from names(multiExpr)) if it exists, otherwise by set number; %b will be replaced by the block number. If the file names turn out to be non-unique, an error will be generated.
nThreads	non-negative integer specifying the number of parallel threads to be used by certain parts of correlation calculations. This option only has an effect on systems on which a POSIX thread library is available (which currently includes Linux and Mac OSX, but excludes Windows). If zero, the number of online processors will be used if it can be determined dynamically, otherwise correlation calculations will use 2 threads.
verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.

`indent` indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

### Details

The function starts by optionally filtering out samples that have too many missing entries and genes that have either too many missing entries or zero variance in at least one set. Genes that are filtered out are excluded from the TOM calculations.

If `blocks` is not given and the number of genes exceeds `maxBlockSize`, genes are pre-clustered into blocks using the function [consensusProjectiveKMeans](#); otherwise all genes are treated in a single block.

For each block of genes, the network is constructed and (if requested) topological overlap is calculated in each set. The topological overlaps can be saved to disk as RData files, or returned directly within the return value (see below). Note that the matrices can be big and returning them within the return value can quickly exhaust the system's memory. In particular, if the block-wise calculation is necessary, it is nearly certain that returning all matrices via the return value will be impossible.

### Value

A list with the following components:

`actualTOMFileNames`

Only returned if input `saveTOMs` is TRUE. A matrix of character strings giving the file names in which each block TOM is saved. Rows correspond to data sets and columns to blocks.

`TOMSimilarities`

Only returned if input `saveTOMs` is FALSE. A list in which each component corresponds to one block. Each component is a matrix of dimensions (number of sets) times N, where N is the length of a distance structure corresponding to the block. That is, if the block contains  $n$  genes,  $N = n*(n-1)/2$ . Each row of the matrix contains the topological overlap of variables in the corresponding set (and the corresponding block), arranged as a distance structure. Do note however that the topological overlap is a similarity (not a distance).

`blocks`

if input `blocks` was given, its copy; otherwise a vector of length equal number of genes giving the block label for each gene. Note that block labels are not necessarily sorted in the order in which the blocks were processed (since we do not require this for the input `blocks`). See `blockOrder` below.

`blockGenes`

a list with one component for each block of genes. Each component is a vector giving the indices (relative to the input `multiExpr`) of genes in the corresponding block.

`goodSamplesAndGenes`

if input `checkMissingData` is TRUE, the output of the function [goodSamplesGenesMS](#). A list with components `goodGenes` (logical vector indicating which genes passed the missing data filters), `goodSamples` (a list of logical vectors indicating which samples passed the missing data filters in each set), and `allOK` (a logical indicating whether all genes and all samples passed the filters). See [goodSamplesGenesMS](#) for more details. If `checkMissingData` is FALSE, `goodSamplesAndGenes` contains a list of the same type but indicating that all genes and all samples passed the missing data filters.

The following components are present mostly to streamline the interaction of this function with [blockwiseConsensusModules](#).

nGenes	Number of genes that passed missing data filters (if input checkMissingData is TRUE), or the number of all genes (if checkMissingData is FALSE).
gBlocks	the vector blocks (above), restricted to good genes only.
nThreads	number of threads used to calculate correlation and TOM matrices.
TOMSavedInFiles	logical: were calculated matrices saved in files (TRUE) or returned in the return value (FALSE)?
intNetworkType, intCorType	integer codes for network and correlation type.

**Author(s)**

Peter Langfelder

**References**

For a general discussion of the weighted network formalism, see

Bin Zhang and Steve Horvath (2005) "A General Framework for Weighted Gene Co-Expression Network Analysis", *Statistical Applications in Genetics and Molecular Biology*: Vol. 4: No. 1, Article 17

The blockwise approach is briefly described in the article describing this package,

Langfelder P, Horvath S (2008) "WGCNA: an R package for weighted correlation network analysis". *BMC Bioinformatics* 2008, 9:559

**See Also**

[blockwiseConsensusModules](#)

---

blockwiseModules	<i>Automatic network construction and module detection</i>
------------------	--

---

**Description**

This function performs automatic network construction and module detection on large expression datasets in a block-wise manner.

**Usage**

```
blockwiseModules(
  datExpr,
  blocks = NULL,
  maxBlockSize = 5000,
  randomSeed = 12345,
  corType = "pearson",
  power = 6,
  networkType = "unsigned",
  TOMType = "signed",
  TOMDenom = "min",
  deepSplit = 2,
  detectCutHeight = 0.995, minModuleSize = min(20, ncol(datExpr)/2 ),
```

```

maxCoreScatter = NULL, minGap = NULL,
maxAbsCoreScatter = NULL, minAbsGap = NULL,
pamStage = TRUE, pamRespectsDendro = TRUE,
minCoreKME = 0.5, minCoreKMESize = minModuleSize/3,
minKMEtoStay = 0.3,
reassignThreshold = 1e-6,
mergeCutHeight = 0.15, impute = TRUE,
getTOMs = NULL,
saveTOMs = FALSE,
saveTOMFileBase = "blockwiseTOM",
trapErrors = FALSE, numericLabels = FALSE,
checkMissingData = TRUE,
maxPOutliers = 1,
quickCor = 0,
pearsonFallback = "individual",
cosineCorrelation = FALSE,
nThreads = 0,
verbose = 0, indent = 0)

```

### Arguments

datExpr	expression data. A data frame in which columns are genes and rows are samples. NAs are allowed, but not too many.
blocks	optional specification of blocks in which hierarchical clustering and module detection should be performed. If given, must be a numeric vector with one entry per column (gene) of exprData giving the number of the block to which the corresponding gene belongs.
maxBlockSize	integer giving maximum block size for module detection. Ignored if blocks above is non-NULL. Otherwise, if the number of genes in datExpr exceeds maxBlockSize, genes will be pre-clustered into blocks whose size should not exceed maxBlockSize.
randomSeed	integer to be used as seed for the random number generator before the function starts. If a current seed exists, it is saved and restored upon exit. If NULL is given, the function will not save and restore the seed.
corType	character string specifying the correlation to be used. Allowed values are (unique abbreviations of) "pearson" and "bicor", corresponding to Pearson and bid-weight midcorrelation, respectively. Missing values are handled using the pairwise.complete.obs option.
power	soft-thresholding power for network construction.
networkType	network type. Allowed values are (unique abbreviations of) "unsigned", "signed", "signed hybrid". See <a href="#">adjacency</a> .
TOMType	one of "none", "unsigned", "signed". If "none", adjacency will be used for clustering. If "unsigned", the standard TOM will be used (more generally, TOM function will receive the adjacency as input). If "signed", TOM will keep track of the sign of correlations between neighbors.
TOMDenom	a character string specifying the TOM variant to be used. Recognized values are "min" giving the standard TOM described in Zhang and Horvath (2005), and "mean" in which the min function in the denominator is replaced by mean. The "mean" may produce better results but at this time should be considered experimental.

deepSplit	integer value between 0 and 4. Provides a simplified control over how sensitive module detection should be to module splitting, with 0 least and 4 most sensitive. See <a href="#">cutreeDynamic</a> for more details.
detectCutHeight	dendrogram cut height for module detection. See <a href="#">cutreeDynamic</a> for more details.
minModuleSize	minimum module size for module detection. See <a href="#">cutreeDynamic</a> for more details.
maxCoreScatter	maximum scatter of the core for a branch to be a cluster, given as the fraction of cutHeight relative to the 5th percentile of joining heights. See <a href="#">cutreeDynamic</a> for more details.
minGap	minimum cluster gap given as the fraction of the difference between cutHeight and the 5th percentile of joining heights. See <a href="#">cutreeDynamic</a> for more details.
maxAbsCoreScatter	maximum scatter of the core for a branch to be a cluster given as absolute heights. If given, overrides maxCoreScatter. See <a href="#">cutreeDynamic</a> for more details.
minAbsGap	minimum cluster gap given as absolute height difference. If given, overrides minGap. See <a href="#">cutreeDynamic</a> for more details.
pamStage	logical. If TRUE, the second (PAM-like) stage of module detection will be performed. See <a href="#">cutreeDynamic</a> for more details.
pamRespectsDendro	Logical, only used when pamStage is TRUE. If TRUE, the PAM stage will respect the dendrogram in the sense an object can be PAM-assigned only to clusters that lie below it on the branch that the object is merged into. See <a href="#">cutreeDynamic</a> for more details.
minCoreKME	a number between 0 and 1. If a detected module does not have at least minModuleKMESize genes with eigengene connectivity at least minCoreKME, the module is disbanded (its genes are unlabeled and returned to the pool of genes waiting for module detection).
minCoreKMESize	see minCoreKME above.
minKMEtoStay	genes whose eigengene connectivity to their module eigengene is lower than minKMEtoStay are removed from the module.
reassignThreshold	p-value ratio threshold for reassigning genes between modules. See Details.
mergeCutHeight	dendrogram cut height for module merging.
impute	logical: should imputation be used for module eigengene calculation? See <a href="#">moduleEigengenes</a> for more details.
getTOMs	deprecated, please use saveTOMs below.
saveTOMs	logical: should the consensus topological overlap matrices for each block be saved and returned?
saveTOMfileBase	character string containing the file name base for files containing the consensus topological overlaps. The full file names have "block.1.RData", "block.2.RData" etc. appended. These files are standard R data files and can be loaded using the <a href="#">load</a> function.
trapErrors	logical: should errors in calculations be trapped?

numericLabels	logical: should the returned modules be labeled by colors (FALSE), or by numbers (TRUE)?
checkMissingData	logical: should data be checked for excessive numbers of missing entries in genes and samples, and for genes with zero variance? See details.
maxPOutliers	only used for corType=="bicor". Specifies the maximum percentile of data that can be considered outliers on either side of the median separately. For each side of the median, if higher percentile than maxPOutliers is considered an outlier by the weight function based on $9 * \text{mad}(x)$ , the width of the weight function is increased such that the percentile of outliers on that side of the median equals maxPOutliers. Using maxPOutliers=1 will effectively disable all weight function broadening; using maxPOutliers=0 will give results that are quite similar (but not equal to) Pearson correlation.
quickCor	real number between 0 and 1 that controls the handling of missing data in the calculation of correlations. See details.
pearsonFallback	Specifies whether the bicor calculation, if used, should revert to Pearson when median absolute deviation (mad) is zero. Recognized values are (abbreviations of) "none", "individual", "all". If set to "none", zero mad will result in NA for the corresponding correlation. If set to "individual", Pearson calculation will be used only for columns that have zero mad. If set to "all", the presence of a single zero mad will cause the whole variable to be treated in Pearson correlation manner (as if the corresponding robust option was set to FALSE). Has no effect for Pearson correlation. See <a href="#">bicor</a> .
cosineCorrelation	logical: should the cosine version of the correlation calculation be used? The cosine calculation differs from the standard one in that it does not subtract the mean.
nThreads	non-negative integer specifying the number of parallel threads to be used by certain parts of correlation calculations. This option only has an effect on systems on which a POSIX thread library is available (which currently includes Linux and Mac OSX, but excludes Windows). If zero, the number of online processors will be used if it can be determined dynamically, otherwise correlation calculations will use 2 threads.
verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.
indent	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

## Details

Before module detection starts, genes and samples are optionally checked for the presence of NAs. Genes and/or samples that have too many NAs are flagged as bad and removed from the analysis; bad genes will be automatically labeled as unassigned, while the returned eigengenes will have NA entries for all bad samples.

If blocks is not given and the number of genes exceeds maxBlockSize, genes are pre-clustered into blocks using the function [projectiveKMeans](#); otherwise all genes are treated in a single block.

For each block of genes, the network is constructed and (if requested) topological overlap is calculated. If requested, the topological overlaps are returned as part of the return value list. Genes are then clustered using average linkage hierarchical clustering and modules are identified in the

resulting dendrogram by the Dynamic Hybrid tree cut. Found modules are trimmed of genes whose correlation with module eigengene (KME) is less than `minKMEtoStay`. Modules in which fewer than `minCoreKMESize` genes have KME higher than `minCoreKME` are disbanded, i.e., their constituent genes are pronounced unassigned.

After all blocks have been processed, the function checks whether there are genes whose KME in the module they assigned is lower than KME to another module. If p-values of the higher correlations are smaller than those of the native module by the factor `reassignThresholdPS`, the gene is re-assigned to the closer module.

In the last step, modules whose eigengenes are highly correlated are merged. This is achieved by clustering module eigengenes using the dissimilarity given by one minus their correlation, cutting the dendrogram at the height `mergeCutHeight` and merging all modules on each branch. The process is iterated until no modules are merged. See [mergeCloseModules](#) for more details on module merging.

The argument `quick` specifies the precision of handling of missing data in the correlation calculations. Zero will cause all calculations to be executed precisely, which may be significantly slower than calculations without missing data. Progressively higher values will speed up the calculations but introduce progressively larger errors. Without missing data, all column means and variances can be pre-calculated before the covariances are calculated. When missing data are present, exact calculations require the column means and variances to be calculated for each covariance. The approximate calculation uses the pre-calculated mean and variance and simply ignores missing data in the covariance calculation. If the number of missing data is high, the pre-calculated means and variances may be very different from the actual ones, thus potentially introducing large errors. The `quick` value times the number of rows specifies the maximum difference in the number of missing entries for mean and variance calculations on the one hand and covariance on the other hand that will be tolerated before a recalculation is triggered. The hope is that if only a few missing data are treated approximately, the error introduced will be small but the potential speedup can be significant.

## Value

A list with the following components:

<code>colors</code>	a vector of color or numeric module labels for all genes.
<code>unmergedColors</code>	a vector of color or numeric module labels for all genes before module merging.
<code>MEs</code>	a data frame containing module eigengenes of the found modules (given by <code>colors</code> ).
<code>goodSamples</code>	numeric vector giving indices of good samples, that is samples that do not have too many missing entries.
<code>goodGenes</code>	numeric vector giving indices of good genes, that is genes that do not have too many missing entries.
<code>dendrograms</code>	a list whose components contain hierarchical clustering dendrograms of genes in each block.
<code>TOMfiles</code>	if <code>saveTOMs==TRUE</code> , a vector of character strings, one string per block, giving the file names of files (relative to current directory) in which blockwise topological overlaps were saved.
<code>blockGenes</code>	a list whose components give the indices of genes in each block.
<code>blocks</code>	if <code>input blocks</code> was given, its copy; otherwise a vector of length equal number of genes giving the block label for each gene. Note that block labels are not necessarily sorted in the order in which the blocks were processed (since we do not require this for the input blocks). See <code>blockOrder</code> below.

blockOrder	a vector giving the order in which blocks were processed and in which blockGenes above is returned. For example, blockOrder[1] contains the label of the first-processed block.
MEsOK	logical indicating whether the module eigengenes were calculated without errors.

**Note**

If the input dataset has a large number of genes, consider carefully the `maxBlockSize` as it significantly affects the memory footprint (and whether the function will fail with a memory allocation error). From a theoretical point of view it is advantageous to use blocks as large as possible; on the other hand, using smaller blocks is substantially faster and often the only way to work with large numbers of genes. As a rough guide, it is unlikely a standard desktop computer with 4GB memory or less will be able to work with blocks larger than 8000 genes.

**Author(s)**

Peter Langfelder

**References**

Bin Zhang and Steve Horvath (2005) "A General Framework for Weighted Gene Co-Expression Network Analysis", *Statistical Applications in Genetics and Molecular Biology*: Vol. 4: No. 1, Article 17

**See Also**

[goodSamplesGenes](#) for basic quality control and filtering;  
[adjacency](#), [TOMsimilarity](#) for network construction;  
[hclust](#) for hierarchical clustering;  
[cutreeDynamic](#) for adaptive branch cutting in hierarchical clustering dendrograms;  
[mergeCloseModules](#) for merging of close modules.

---

BloodLists

*Blood Cell Types with Corresponding Gene Markers*

---

**Description**

This matrix gives a predefined set of marker genes for many blood cell types, as reported in several previously-published studies. It is used with `userListEnrichment` to search user-defined gene lists for enrichment.

**Usage**

```
data(BloodLists)
```



**Format**

A 2048 x 2 matrix of characters containing Gene / Category pairs. The first column (Gene) lists genes corresponding to a given category (second column). Each Category entry is of the form <Blood cell type>\_\_<reference>, where the references can be found at [userListEnrichment](#). Note that the matrix is sorted first by Category and then by Gene, such that all genes related to the same category are listed sequentially.

**Source**

For references used in this variable, please see [userListEnrichment](#)

**Examples**

```
data(BloodLists)
head(BloodLists)
```

---

blueWhiteRed	<i>Blue-white-red color sequence</i>
--------------	--------------------------------------

---

**Description**

Generate a blue-white-red color sequence of a given length.

**Usage**

```
blueWhiteRed(n, gamma = 1)
```

**Arguments**

n	number of colors to be returned
gamma	color change power

**Details**

The function returns a color vector that starts with blue, gradually turns into white and then to red. The power gamma can be used to control the behaviour of the quarter- and three quarter-values (between blue and white, and white and red, respectively). Higher powers will make the mid-colors more white, while lower powers will make the colors more saturated, respectively.

**Value**

A vector of colors of length n.

**Author(s)**

Peter Langfelder

**See Also**

[numbers2colors](#) for a function that produces a color representation for continuous numbers.

**Examples**

```
par(mfrow = c(3, 1))
displayColors(blueWhiteRed(50));
title("gamma = 1")
displayColors(blueWhiteRed(50, 3));
title("gamma = 3")
displayColors(blueWhiteRed(50, 0.5));
title("gamma = 0.5")
```

BrainLists

*Brain-Related Categories with Corresponding Gene Markers***Description**

This matrix gives a predefined set of marker genes for many brain-related categories (ie., cell type, organelle, changes with disease, etc.), as reported in several previously-published studies. It is used with `userListEnrichment` to search user-defined gene lists for enrichment.

**Usage**

```
data(BrainLists)
```

**Format**

A 48319 x 2 matrix of characters containing Gene / Category pairs. The first column (Gene) lists genes corresponding to a given category (second column). Each Category entry is of the form `<Brain descriptor>__<reference>`, where the references can be found at [userListEnrichment](#). Note that the matrix is sorted first by Category and then by Gene, such that all genes related to the same category are listed sequentially.

**Source**

For references used in this variable, please see [userListEnrichment](#)

**Examples**

```
data(BrainLists)
head(BrainLists)
```

BrainRegionMarkers

*Gene Markers for Regions of the Human Brain***Description**

This matrix gives a predefined set of marker genes for many regions of the human brain, using data from the Allen Human Brain Atlas (<http://human.brain-map.org/>) as reported in: Hawrylycz MJ, Lein ES, Guillozet-Bongaarts AL, Shen EH, Ng L, Miller JA, et al. (2012) An Anatomically Comprehensive Atlas of the Adult Human Brain Transcriptome. Nature (in press). It is used with `userListEnrichment` to search user-defined gene lists for enrichment.

**Usage**

```
data(BrainRegionMarkers)
```

**Format**

A 28477 x 2 matrix of characters containing Gene / Category pairs. The first column (Gene) lists genes corresponding to a given category (second column). Each Category entry is of the form <Brain Region>\_<Marker Type>\_HBA. Note that the matrix is sorted first by Category and then by Gene, such that all genes related to the same category are listed sequentially.

**Source**

For references used in this variable, or other information, please see [userListEnrichment](#)

**Examples**

```
data(BrainRegionMarkers)
head(BrainRegionMarkers)
```

---

checkAdjMat	<i>Check adjacency matrix</i>
-------------	-------------------------------

---

**Description**

Checks a given matrix for properties that an adjacency matrix must satisfy.

**Usage**

```
checkAdjMat(adjMat, min = 0, max = 1)
checkSimilarity(similarity, min = -1, max = 1)
```

**Arguments**

adjMat	matrix to be checked
similarity	matrix to be checked
min	minimum allowed value for entries of the input
max	maximum allowed value for entries of the input

**Details**

The function checks whether the given matrix really is a 2-dimensional numeric matrix, whether it is square, symmetric, and all finite entries are between min and max. If any of the conditions is not met, the function issues an error.

**Value**

None. The function returns normally if all conditions are met.

**Author(s)**

Peter Langfelder

**See Also**

[adjacency](#)

---

checkSets	<i>Check structure and retrieve sizes of a group of datasets.</i>
-----------	---

---

**Description**

Checks whether given sets have the correct format and retrieves dimensions.

**Usage**

```
checkSets(data, checkStructure = FALSE, useSets = NULL)
```

**Arguments**

data	A vector of lists; in each list there must be a component named data whose content is a matrix or dataframe or array of dimension 2.
checkStructure	If FALSE, incorrect structure of data will trigger an error. If TRUE, an appropriate flag (see output) will be set to indicate whether data has correct structure.
useSets	Optional specification of entries of the vector data that are to be checked. Defaults to all components. This may be useful when data only contains information for some of the sets.

**Details**

For multiset calculations, many quantities (such as expression data, traits, module eigengenes etc) are presented by a common structure, a vector of lists (one list for each set) where each list has a component data that contains the actual (expression, trait, eigengene) data for the corresponding set in the form of a dataframe. This function checks whether data conforms to this convention and retrieves some basic dimension information (see output).

**Value**

A list with components

nSets	Number of sets (length of the vector data).
nGenes	Number of columns in the data components in the lists. This number must be the same for all sets.
nSamples	A vector of length nSets giving the number of rows in the data components.
structureOK	Only set if the argument checkStructure equals TRUE. The value is TRUE if the parameter data passes a few tests of its structure, and FALSE otherwise. The tests are not exhaustive and are meant to catch obvious user errors rather than be bulletproof.

**Author(s)**

Peter Langfelder, <Peter.Langfelder@gmail.com>

---

`chooseOneHubInEachModule`*Chooses a single hub gene in each module*

---

**Description**

`chooseOneHubInEachModule` returns one gene in each module with high connectivity, given a number of randomly selected genes to test.

**Usage**

```
chooseOneHubInEachModule(  
  datExpr,  
  colorh,  
  numGenes = 100,  
  omitColors = "grey",  
  power = 2,  
  type = "signed",  
  ...)
```

**Arguments**

<code>datExpr</code>	Gene expression data with rows as samples and columns as genes.
<code>colorh</code>	The module assignments (color vectors) corresponding to the rows in <code>datExpr</code> .
<code>numGenes</code>	The number of random genes to select per module. Higher number of genes increases the accuracy of hub selection but slows down the function.
<code>omitColors</code>	All colors in this character vector (default is "grey") are ignored by this function.
<code>power</code>	Power to use for the adjacency network (default = 2).
<code>type</code>	What type of network is being entered. Common choices are "signed" (default) and "unsigned". With "signed" negative correlations count against, whereas with "unsigned" negative correlations are treated identically as positive correlations.
<code>...</code>	Any other parameters accepted by the <code>*adjacency*</code> function

**Value**

Both functions output a character vector of genes, where the genes are the hub gene picked for each module, and the names correspond to the module in which each gene is a hub.

**Author(s)**

Jeremy Miller

**Examples**

```
## Example: first simulate some data.  
  
MEturquoise = sample(1:100,50)  
MEblue      = sample(1:100,50)  
MEbrown     = sample(1:100,50)  
MEyellow    = sample(1:100,50)
```

```

MEgreen      = c(MEyellow[1:30], sample(1:100,20))
MERed       = c(MEbrown [1:20], sample(1:100,30))
MEblack     = c(MEblue  [1:25], sample(1:100,25))
ME          = data.frame(MEturquoise, MEblue, MEbrown, MEyellow, MEgreen, MERed, MEblack)
dat1        = simulateDatExpr(ME,300,c(0.2,0.1,0.08,0.051,0.05,0.042,0.041,0.3),
                             signed=TRUE)
TOM1        = TOMsimilarityFromExpr(dat1$datExpr, networkType="signed")
colnames(TOM1) <- rownames(TOM1) <- colnames(dat1$datExpr)
tree1 <- tree2 <- flashClust(as.dist(1-TOM1),method="average")
colorh      = labels2colors(dat1$allLabels)
hubs        = chooseOneHubInEachModule(dat1$datExpr, colorh)
hubs

```

---

chooseTopHubInEachModule

*Chooses the top hub gene in each module*

---

### Description

chooseTopHubInEachModule returns the gene in each module with the highest connectivity, looking at all genes in the expression file.

### Usage

```

chooseTopHubInEachModule(
  datExpr,
  colorh,
  omitColors = "grey",
  power = 2,
  type = "signed",
  ...)

```

### Arguments

datExpr	Gene expression data with rows as samples and columns as genes.
colorh	The module assignments (color vectors) corresponding to the rows in datExpr.
omitColors	All colors in this character vector (default is "grey") are ignored by this function.
power	Power to use for the adjacency network (default = 2).
type	What type of network is being entered. Common choices are "signed" (default) and "unsigned". With "signed" negative correlations count against, whereas with "unsigned" negative correlations are treated identically as positive correlations.
...	Any other parameters accepted by the *adjacency* function

### Value

Both functions output a character vector of genes, where the genes are the hub gene picked for each module, and the names correspond to the module in which each gene is a hub.

### Author(s)

Jeremy Miller

**Examples**

```
## Example: first simulate some data.

MEturquoise = sample(1:100,50)
MEblue      = sample(1:100,50)
MEbrown     = sample(1:100,50)
MEyellow    = sample(1:100,50)
MEgreen     = c(MEyellow[1:30], sample(1:100,20))
MERed      = c(MEbrown [1:20], sample(1:100,30))
MEblack     = c(MEblue  [1:25], sample(1:100,25))
ME         = data.frame(MEturquoise, MEblue, MEbrown, MEyellow, MEgreen, MERed, MEblack)
dat1       = simulateDatExpr(ME,300,c(0.2,0.1,0.08,0.051,0.05,0.042,0.041,0.3), signed=TRUE)
TOM1      = TOMsimilarityFromExpr(dat1$datExpr, networkType="signed")
colnames(TOM1) <- rownames(TOM1) <- colnames(dat1$datExpr)
tree1 <- tree2 <- flashClust(as.dist(1-TOM1),method="average")
colorh = labels2colors(dat1$allLabels)
hubs   = chooseTopHubInEachModule(dat1$datExpr, colorh)
hubs
```

---

clusterCoef

*Clustering coefficient calculation*

---

**Description**

This function calculates the clustering coefficients for all nodes in the network given by the input adjacency matrix.

**Usage**

```
clusterCoef(adjMat)
```

**Arguments**

adjMat            adjacency matrix

**Value**

A vector of clustering coefficients for each node.

**Author(s)**

Steve Horvath

---

coClustering	<i>Co-clustering measure of cluster preservation between two clusterings</i>
--------------	--

---

**Description**

The function calculates the co-clustering statistics for each module in the reference clustering.

**Usage**

```
coClustering(clusters.ref, clusters.test, tupleSize = 2, unassignedLabel = 0)
```

**Arguments**

<code>clusters.ref</code>	Reference input clustering. A vector in which each element gives the cluster label of an object.
<code>clusters.test</code>	Test input clustering. Must be a vector of the same size as <code>clusters.ref</code> .
<code>tupleSize</code>	Co-clustering tuple size.
<code>unassignedLabel</code>	Optional specification of a clustering label that denotes unassigned objects. Objects with this label are excluded from the calculation.

**Details**

Co-clustering of cluster  $q$  in the reference clustering and cluster  $q'$  in the test clustering measures the overlap of clusters  $q$  and  $q'$  by the number of tuples that can be chosen from the overlap of clusters  $q$  and  $q'$  relative to the number of tuples in cluster  $q$ . To arrive at a co-clustering measure for cluster  $q$ , we sum the co-clustering of  $q$  and  $q'$  over all clusters  $q'$  in the test clustering. A value close to 1 indicates high preservation of the reference cluster in the test clustering, while a value close to zero indicates a low preservation.

**Value**

A vector in which each component corresponds to a cluster in the reference clustering. Entries give the co-clustering measure of cluster preservation.

**Author(s)**

Peter Langfelder

**References**

For example, see Langfelder P, Luo R, Oldham MC, Horvath S (2011) Is My Network Module Preserved and Reproducible? PLoS Comput Biol 7(1): e1001057. Co-clustering is discussed in the Methods Supplement (Supplementary text 1) of that article.

**See Also**

[modulePreservation](#) for a large suite of module preservation statistics [coClustering.permutationTest](#) for a permutation test for co-clustering significance



**Examples**

```
# An example with random (unrelated) clusters:

set.seed(1);
nModules = 10;
nGenes = 1000;
c11 = sample(c(1:nModules), nGenes, replace = TRUE);
c12 = sample(c(1:nModules), nGenes, replace = TRUE);
coClustering(c11, c12)

# For the same reference and test clustering:

coClustering(c11, c11)
```

---

```
coClustering.permutationTest
```

*Permutation test for co-clustering*

---

**Description**

This function calculates permutation  $Z$  statistics that measure how different the co-clustering of modules in a reference and test clusterings is from random.

**Usage**

```
coClustering.permutationTest(
  clusters.ref, clusters.test,
  tupleSize = 2,
  nPermutations = 100,
  unassignedLabel = 0,
  randomSeed = 12345, verbose = 0, indent = 0)
```

**Arguments**

<code>clusters.ref</code>	Reference input clustering. A vector in which each element gives the cluster label of an object.
<code>clusters.test</code>	Test input clustering. Must be a vector of the same size as <code>clusters.ref</code> .
<code>tupleSize</code>	Co-clustering tuple size.
<code>nPermutations</code>	Number of permutations to execute. Since the function calculates parametric p-values, a relatively small number of permutations (at least 50) should be sufficient.
<code>unassignedLabel</code>	Optional specification of a clustering label that denotes unassigned objects. Objects with this label are excluded from the calculation.
<code>randomSeed</code>	Random seed for initializing the random number generator. If NULL, the generator is not initialized (useful for calling the function sequentially). The default assures reproducibility.
<code>verbose</code>	If non-zero, function will print out progress messages.
<code>indent</code>	Indentation for progress messages. Each unit adds two spaces.

**Details**

This function performs a permutation test to determine whether observed co-clustering statistics are significantly different from those expected by chance. It returns the observed co-clustering as well as the permutation Z statistic, calculated as  $(\text{observed} - \text{mean})/\text{sd}$ , where mean and sd are the mean and standard deviation of the co-clustering when the test clustering is repeatedly randomly permuted.

**Value**

observed	the observed co-clustering measures for clusters in <code>clusters.ref</code>
Z	permutation Z statics
permuted.mean	means of the co-clustering measures when the test clustering is permuted
permuted.sd	standard deviations of the co-clustering measures when the test clustering is permuted
permuted.cc	values of the co-clustering measure for each permutation of the test clustering. A matrix of dimensions (number of permutations)x(number of clusters in reference clustering).

**Author(s)**

Peter Langfelder

**References**

For example, see Langfelder P, Luo R, Oldham MC, Horvath S (2011) Is My Network Module Preserved and Reproducible? PLoS Comput Biol 7(1): e1001057. Co-clustering is discussed in the Methods Supplement (Supplementary text 1) of that article.

**See Also**

[coClustering](#) for calculation of the "observed" co-clustering measure [modulePreservation](#) for a large suite of module preservation statistics

**Examples**

```
set.seed(1);
nModules = 5;
nGenes = 100;
c11 = sample(c(1:nModules), nGenes, replace = TRUE);
c12 = sample(c(1:nModules), nGenes, replace = TRUE);

cc = coClustering(c11, c12)

# Choose a low number of permutations to make the example fast
ccPerm = coClustering.permutationTest(c11, c12, nPermutations = 20, verbose = 1);

ccPerm$observed
ccPerm$Z

# Combine c11 and c12 to obtain clustering that is somewhat similar to c11:

c13 = c12;
from1 = sample(c(TRUE, FALSE), nGenes, replace = TRUE);
```

```

c13[from1] = c11[from1];

ccPerm = coClustering.permutationTest(c11, c13, nPermutations = 20, verbose = 1);

# observed co-clustering is higher than before:
ccPerm$observed

# Note the high preservation Z statistics:
ccPerm$Z

```

collapseRows

*Select one representative row per group***Description**

Abstractly speaking, the function allows one to collapse the rows of a numeric matrix, e.g. by forming an average or selecting one representative row for each group of rows specified by a grouping variable (referred to as `rowGroup`). The word "collapse" reflects the fact that the method yields a new matrix whose rows correspond to other rows of the original input data. The function implements several network-based and biostatistical methods for finding a representative row for each group specified in `rowGroup`. Optionally, the function identifies the representative row according to the least number of missing data, the highest sample mean, the highest sample variance, the highest connectivity. One of the advantages of this function is that it implements default settings which have worked well in numerous applications. Below, we describe these default settings in more detail.

**Usage**

```

collapseRows(datET, rowGroup, rowID,
             method="MaxMean", connectivityBasedCollapsing=FALSE,
             methodFunction=NULL, connectivityPower=1,
             selectFewestMissing=TRUE, thresholdCombine=NA)

```

**Arguments**

<code>datET</code>	matrix or data frame containing numeric values where rows correspond to variables (e.g. microarray probes) and columns correspond to observations (e.g. microarrays). Each row of <code>datET</code> must have a unique row identifier (specified in the vector <code>rowID</code> ). The group label of each row is encoded in the vector <code>rowGroup</code> . While <code>rowID</code> should have non-missing, unique values (identifiers), the values of the vector <code>rowGroup</code> will typically not be unique since the function aims to pick a representative row for each group.
<code>rowGroup</code>	character vector whose components contain the group label (e.g. a character string) for each row of <code>datET</code> . This vector needs to have the same length as the vector <code>rowID</code> . In gene expression applications, this vector could contain the gene symbol (or a co-expression module label).
<code>rowID</code>	character vector of row identifiers. This should include all the rows from <code>rownames(datET)</code> , but can include other rows. Its entries should be unique (no duplicates) and no missing values are permitted. If the row identifier is missing for a given row, we suggest you remove this row from <code>datET</code> before applying the function.

**method** character string for determining which method is used to choose a probe among exactly 2 corresponding rows or when `connectivityBasedCollapsing=FALSE`. These are the options: "MaxMean" (default) or "MinMean" = choose the row with the highest or lowest mean value, respectively. "maxRowVariance" = choose the row with the highest variance (across the columns of `datET`). "absMaxMean" or "absMinMean" = choose the row with the highest or lowest mean absolute value. "ME" = choose the eigenrow (first principal component of the rows in each group). Note that with this method option, `connectivityBasedCollapsing` is automatically set to `FALSE`. "Average" = for each column, take the average value of the rows in each group "function" = use this method for a user-input function (see the description of the argument "methodFunction"). Note: if `method="ME"`, "Average" or "function", the output parameters "group2row" and "selectedRow" are not informative.

**connectivityBasedCollapsing**

logical value. If `TRUE`, groups with 3 or more corresponding rows will be represented by the row with the highest connectivity according to a signed weighted correlation network adjacency matrix among the corresponding rows. Recall that the connectivity is defined as the rows sum of the adjacency matrix. The signed weighted adjacency matrix is defined as  $A=(0.5+0.5*\text{COR})^{\text{power}}$  where power is determined by the argument `connectivityPower` and `COR` denotes the matrix of pairwise Pearson correlation coefficients among the corresponding rows.

**methodFunction** character string. It only needs to be specified if `method="function"` otherwise its input is ignored. Must be a function that takes a  $N_r \times N_c$  matrix of numbers as input and outputs a vector with the length  $N_c$  (e.g., `colMeans`). This will then be the method used for collapsing values for multiple rows into a single value for the row.

**connectivityPower**

Positive number (typically integer) for specifying the threshold (power) used to construct the signed weighted adjacency matrix, see the description of `connectivityBasedCollapsing`. This option is only used if `connectivityBasedCollapsing=TRUE`.

**selectFewestMissing**

logical values. If `TRUE` (default), the input expression matrix is trimmed such that for each group only the rows with the fewest number of missing values are retained. In situations where an equal number of values are missing (or where there is no missing data), all rows for a given group are retained. Whether this value is set to `TRUE` or `FALSE`, all rows with  $>90\%$  missing data are omitted from the analysis.

**thresholdCombine**

Number between -1 and 1, or `NA`. If `NA` (default), this input is ignored. If a number between -1 and 1 is input, this value is taken as a threshold value, and `collapseRows` proceeds following the "maxMean" method, but ONLY for ids with correlations of  $R > \text{thresholdCombine}$ . Specifically: ...1) If there is one id/group, keep the id ...2) If there are 2 ids/group, take the maximum mean expression if their correlation is  $> \text{thresholdCombine}$  ...3) If there are 3+ ids/group, iteratively repeat (2) for the 2 ids with the highest correlation until all ids remaining have correlation  $< \text{thresholdCombine}$  for each group Note that this option usually results in more than one id per group; therefore, one must use care when implementing this option for use in comparisons between multiple matrices / data frames.

## Details

The function is robust to missing data. Also, if rowIDs are missing, they are inferred according to the rownames of `datET` when possible. When a group corresponds to only 1 row then it is represented by this row since there is no other choice. Having said this, the row may be removed if it contains an excessive amount of missing data (90 percent or more missing values), see the description of the argument `selectFewestMissing` for more details.

A group is represented by a corresponding row with the fewest number of missing data if `selectFewestMissing` has been set to `TRUE`. Often several rows have the same minimum number of missing values (or no missing values) and a representative must be chosen among those rows. In this case we distinguish 2 situations: (1) If a group corresponds to exactly 2 rows then the corresponding row with the highest average is selected if `method="maxMean"`. Alternative methods can be chosen as described in `method`. (2) If a group corresponds to more than 2 rows, then the function calculates a signed weighted correlation network (with power specified in `connectivityPower`) among the corresponding rows if `connectivityBasedCollapsing=TRUE`. Next the function calculates the network connectivity of each row (closely related to the sum or correlations with the other matching rows). Next it chooses the most highly connected row as representative. If `connectivityBasedCollapsing=FALSE`, then `method` is used. For both situations, if more than one row has the same value, the first such row is chosen.

Setting `thresholdCombine` is a special case of this function, as not all ids for a single group are necessarily collapsed—only those with similar expression patterns are collapsed. We suggest using this option when the goal is to decrease the number of ids for computational reasons, but when ALL ids for a single group should not be combined (for example, if two probes could represent different splice variants for the same gene for many genes on a microarray).

Example application: when dealing with microarray gene expression data then the rows of `datET` may correspond to unique probe identifiers and `rowGroup` may contain corresponding gene symbols. Recall that multiple probes (specified using `rowID=ProbeID`) may correspond to the same gene symbol (specified using `rowGroup=GeneSymbol`). In this case, `datET` contains the input expression data with rows as rowIDs and output expression data with rows as gene symbols, collapsing all probes for a given gene symbol into one representative.

## Value

The output is a list with the following components.

- `datETcollapsed` is a numeric matrix with the same columns as the input matrix `datET`, but with rows corresponding to the different row groups rather than individual row identifiers. (If `thresholdCombine` is set, then rows still correspond to individual row identifiers.)
- `group2row` is a matrix whose rows correspond to the unique group labels and whose 2 columns report which group label (first column called `group`) is represented by what row label (second column called `selectedRowID`). Set to `NULL` if `method="ME"` or `"function"`.
- .
- `selectedRow` is a logical vector whose components are `TRUE` for probes selected as representatives and `FALSE` otherwise. It has the same length as the vector `probeID`. Set to `NULL` if `method="ME"` or `"function"`.

## Author(s)

Jeremy A. Miller, Steve Horvath, Peter Langfelder, Chaochao Cai

## References

Miller JA, Langfelder P, Cai C, Horvath S (2010) Strategies for optimally aggregating gene expression data: The collapseRows R function. Technical Report.

## Examples

```
#####
# EXAMPLE 1:
# The code simulates a data frame (called dat1) of correlated rows.
# You can skip this part and start at the line called Typical Input Data
# The first column of the data frame will contain row identifiers
# number of columns (e.g. observations or microarrays)
m=60
# number of rows (e.g. variables or probes on a microarray)
n=500
# seed module eigenvector for the simulateModule function
MEtrue=rnorm(m)
# numeric data frame of n rows and m columns
datNumeric=data.frame(t(simulateModule(MEtrue,n)))
RowIdentifier=paste("Probe", 1:n, sep="")
ColumnName=paste("Sample",1:m, sep="")
dimnames(datNumeric)[[2]]=ColumnName
# Let us now generate a data frame whose first column contains the rowID
dat1=data.frame(RowIdentifier, datNumeric)
#we simulate a vector with n/5 group labels, i.e. each row group corresponds to 5 rows
rowGroup=rep( paste("Group",1:(n/5), sep=""), 5 )

# Typical Input Data
# Since the first column of dat1 contains the RowIdentifier, we use the following code
datET=dat1[,-1]
rowID=dat1[,1]

# assign row names according to the RowIdentifier
dimnames(datET)[[1]]=rowID
# run the function and save it in an object

collapse.object=collapseRows(datET=datET, rowGroup=rowGroup, rowID=rowID)

# this creates the collapsed data where
# the first column contains the group name
# the second column reports the corresponding selected row name (the representative)
# and the remaining columns report the values of the representative row
dat1Collapsed=data.frame( collapse.object$group2row, collapse.object$datETcollapsed)
dat1Collapsed[1:5,1:5]

#####
# EXAMPLE 2:
# Using the same data frame as above, run collapseRows with a user-inputted function.
# In this case we will use the mean. Note that since we are choosing some combination
# of the probe values for each gene, the group2row and selectedRow output
# parameters are not meaningful.

collapse.object.mean=collapseRows(datET=datET, rowGroup=rowGroup, rowID=rowID,
                                method="function", methodFunction=colMeans)[[1]]

# Note that in this situation, running the following code produces the identical results:
```

```
collapse.object.mean.2=collapseRows(datET=datET, rowGroup=rowGroup, rowID=rowID,
  method="Average")[[1]]

#####
# EXAMPLE 3:
# Using collapseRows to calculate the module eigengene.
# First we create some sample data as in example 1 (or use your own!)
m=60
n=500
MEtrue=rnorm(m)
datNumeric=data.frame(t(simulateModule(MEtrue,n)))

# In this example, rows are genes, and groups are modules.
RowIdentifier=paste("Gene", 1:n, sep="")
ColumnName=paste("Sample",1:m, sep="")
dimnames(datNumeric)[[2]]=ColumnName
dat1=data.frame(RowIdentifier, datNumeric)
# We simulate a vector with n/100 modules, i.e. each row group corresponds to 100 rows
rowGroup=rep( paste("Module",1:(n/100), sep=""), 100 )
datET=dat1[,-1]
rowID=dat1[,1]
dimnames(datET)[[1]]=rowID

# run the function and save it in an object
collapse.object.ME=collapseRows(datET=datET, rowGroup=rowGroup, rowID=rowID, method="ME")[[1]]

# Note that in this situation, running the following code produces the identical results:
collapse.object.ME.2 = t(moduleEigengenes(expr=t(datET),colors=rowGroup)$eigengene)
colnames(collapse.object.ME.2) = ColumnName
rownames(collapse.object.ME.2) = sort(unique(rowGroup))
```

---

collapseRowsUsingKME    *Selects one representative row per group based on kME*

---

## Description

This function selects only the most informative probe for each gene in a kME table, only keeping the probe which has the highest kME with respect to any module in the module membership matrix. This function is a special case of the function collapseRows.

## Usage

```
collapseRowsUsingKME(MM, Gin, Pin = NULL, kMEcols = 1:dim(MM)[2])
```

## Arguments

MM	A module membership (kME) table with at least a subset of the columns corresponding to kME values.
Gin	Genes labels in a 1 to 1 correspondence with the rows of MM.
Pin	If NULL (default), rownames of MM are assumed to be probe IDs. If entered, Pin must be the same length as Gin and correspond to probe IDs for MM.
kMEcols	A numeric vector showing which columns in MM correspond to kME values. The default is all of them.

**Value**

datETcollapsed	A numeric matrix with the same columns as the input matrix MM, but with rows corresponding to the genes rather than the probes.
group2row	A matrix whose rows correspond to the unique gene labels and whose 2 columns report which gene label (first column called group) is represented by what probe (second column called selectedRowID)
selectedRow	A logical vector whose components are TRUE for probes selected as representatives and FALSE otherwise. It has the same length as the vector Pin.

**Author(s)**

Jeremy Miller

**See Also**

[collapseRows](#)

**Examples**

```
# Example: first simulate some data
set.seed(100)
ME.A = sample(1:100,50); ME.B = sample(1:100,50)
ME.C = sample(1:100,50); ME.D = sample(1:100,50)
ME1 = data.frame(ME.A, ME.B, ME.C, ME.D)
simDatA = simulateDatExpr(ME1,1000,c(0.2,0.1,0.08,0.05,0.3), signed=TRUE)
simDatB = simulateDatExpr(ME1,1000,c(0.2,0.1,0.08,0.05,0.3), signed=TRUE)
Gin = c(colnames(simDatA$datExpr),colnames(simDatB$datExpr))
Pin = paste("Probe",1:length(Gin),sep=".")
datExpr = cbind(simDatA$datExpr, simDatB$datExpr)
MM = corAndPvalue(datExpr,ME1)$cor

# Now run the function and see some example output
results = collapseRowsUsingKME(MM, Gin, Pin)
head(results$MMcollapsed)
head(results$group2Row)
head(results$selectedRow)
```

---

collectGarbage

*Iterative garbage collection.*

---

**Description**

Performs garbage collection until free memory indicators show no change.

**Usage**

```
collectGarbage()
```

**Value**

None.



**Author(s)**

Steve Horvath

---

`colQuantileC`*Fast column-wise quantile of a matrix.*

---

**Description**

Fast calculation of column-wise quantiles of a matrix at a single probability. Implemented via compiled code, it is much faster than the equivalent `apply(data, 2, quantile, prob = p)`.

**Usage**

```
colQuantileC(data, p)
```

**Arguments**

<code>data</code>	a numerical matrix column-wise quantiles are desired. Missing values are currently not allowed.
<code>p</code>	a single probability at which the quantile is to be calculated.

**Value**

A vector of length equal the number of columns in `data` containing the column-wise quantiles.

**Author(s)**

Peter Langfelder

**See Also**[quantile](#)

---

`conformityBasedNetworkConcepts`*Calculation of conformity-based network concepts.*

---

**Description**

This function computes 3 types of network concepts (also known as network indices or statistics) based on an adjacency matrix and optionally a node significance measure.

**Usage**

```
conformityBasedNetworkConcepts(adj, GS = NULL)
```

**Arguments**

adj	adjacency matrix. A symmetric matrix with components between 0 and 1.
GS	optional node significance measure. A vector with length equal the dimension of adj.

**Details**

This function computes 3 types of network concepts (also known as network indices or statistics) based on an adjacency matrix and optionally a node significance measure. Specifically, it computes I) fundamental network concepts, II) conformity based network concepts, and III) approximate conformity based network concepts. These network concepts are defined for any symmetric adjacency matrix (weighted and unweighted). The network concepts are described in Dong and Horvath (2007) and Horvath and Dong (2008). In the following, we use the term gene and node interchangeably since these methods were originally developed for gene networks. In the following, we briefly describe the 3 types of network concepts:

Type I: fundamental network concepts are defined as a function of the off-diagonal elements of an adjacency matrix  $A$  and/or a node significance measure  $GS$ . Type II: conformity-based network concepts are functions of the off-diagonal elements of the conformity based adjacency matrix  $A.CF=CF*t(CF)$  and/or the node significance measure. These network concepts are defined for any network for which a conformity vector can be defined. Details: For any adjacency matrix  $A$ , the conformity vector  $CF$  is calculated by requiring that  $A[i,j]$  is approximately equal to  $CF[i]*CF[j]$ . Using the conformity one can define the matrix  $A.CF=CF*t(CF)$  which is the outer product of the conformity vector with itself. In general,  $A.CF$  is not an adjacency matrix since its diagonal elements are different from 1. If the off-diagonal elements of  $A.CF$  are similar to those of  $A$  according to the Frobenius matrix norm, then  $A$  is approximately factorizable. To measure the factorizability of a network, one can calculate the Factorizability, which is a number between 0 and 1 (Dong and Horvath 2007). The conformity is defined using a monotonic, iterative algorithm that maximizes the factorizability measure. Type III: approximate conformity based network concepts are functions of all elements of the conformity based adjacency matrix  $A.CF$  (including the diagonal) and/or the node significance measure  $GS$ . These network concepts are very useful for deriving relationships between network concepts in networks that are approximately factorizable.

**Value**

A list with the following components:

Factorizability	number between 0 and 1 giving the factorizability of the matrix. The closer to 1 the higher the evidence of factorizability, that is, $A-I$ is close to $outer(CF,CF)-diag(CF^2)$ .
fundamentalNCS	fundamental network concepts, that is network concepts calculated directly from the given adjacency matrix $adj$ . A list with components <i>ScaledConnectivity</i> (giving the scaled connectivity of each node), <i>Connectivity</i> (connectivity of each node), <i>ClusterCoef</i> (the clustering coefficient of each node), <i>MAR</i> (maximum adjacency ratio of each node), <i>Density</i> (the mean density of the network), <i>Centralization</i> (the centralization of the network), <i>Heterogeneity</i> (the heterogeneity of the network). If the input node significance $GS$ is specified, the following additional components are included: <i>NetworkSignificance</i> (network significance, the mean node significance), and <i>HubNodeSignificance</i> (hub node significance given by the linear regression of node significance on connectivity).

## conformityBasedNCs

network concepts based on an approximate adjacency matrix given by the outer product of the conformity vector but with unit diagonal. A list with components Conformity (the conformity vector) and Connectivity.CF, ClusterCoef.CF, MAR.CF, Density.CF giving the conformity-based analogs of the above network concepts.

## approximateConformityBasedNCs

network concepts based on an approximate adjacency matrix given by the outer product of the conformity vector. A list with components Conformity (the conformity vector) and Connectivity.CF.App, ClusterCoef.CF.App, MAR.CF.App, Density.CF.App giving the conformity-based analogs of the above network concepts.

**Author(s)**

Steve Horvath

**References**

Dong J, Horvath S (2007) Understanding Network Concepts in Modules, BMC Systems Biology 2007, 1:24 Horvath S, Dong J (2008) Geometric Interpretation of Gene Coexpression Network Analysis. PLoS Comput Biol 4(8): e1000117

**See Also**

[networkConcepts](#) for calculation of eigennode based network concepts for a correlation network;  
[fundamentalNetworkConcepts](#) for calculation of fundamental network concepts only.

## conformityDecomposition

*Conformity and module based decomposition of a network adjacency matrix.*

**Description**

The function calculates the conformity based approximation A.CF of an adjacency matrix and a factorizability measure codeFactorizability. If a module assignment C1 is provided, it also estimates a corresponding intermodular adjacency matrix. In this case, function automatically carries out the module- and conformity based decomposition of the adjacency matrix described in chapter 2 of (Horvath 2011).

**Usage**

```
conformityDecomposition(adj, C1 = NULL)
```

**Arguments**

adj a symmetric numeric matrix (or data frame) whose entries lie between 0 and 1.  
 C1 a vector (or factor variable) of length equal to the number of rows of adj. The variable assigns each network node (row of adj) to a module. The entries of C1 could be integers or character strings.

## Details

We distinguish two situations depending on whether or not `C1` equals `NULL`. 1) Let us start out assuming that `C1 = NULL`. In this case, the function calculates the conformity vector for a general, possibly non-factorizable network `adj` by minimizing a quadratic (sums of squares) loss function. The conformity and factorizability for an adjacency matrix is defined in (Dong and Horvath 2007, Horvath and Dong 2008) but we briefly describe it in the following. A network is called exactly factorizable if the pairwise connection strength (adjacency) between 2 network nodes can be factored into node specific contributions, named node 'conformity', i.e. if  $adj[i, j] = Conformity[i] * Conformity[j]$ . The conformity turns out to be highly related to the network connectivity (aka degree). If `adj` is not exactly factorizable, then the function `conformityDecomposition` calculates a conformity vector of the exactly factorizable network that best approximates `adj`. The factorizability measure `Factorizability` is a number between 0 and 1. The higher `Factorizability`, the more factorizable is `adj`. Warning: the algorithm may only converge to a local optimum and it may not converge at all. Also see the notes below.

2) Let us now assume that `C1` is not `NULL`, i.e. it specifies the module assignment of each node. Then the function calculates a module- and CF-based approximation of `adj` (explained in chapter 2 in Horvath 2011). In this case, the function calculates a conformity vector `Conformity` and a matrix `IntermodularAdjacency` such that  $adj[i, j]$  is approximately equal to  $Conformity[i] * Conformity[j] * IntermodularAdjacency[module.index[i], module.index[j]]$  where `module.index[i]` is the row of the matrix `IntermodularAdjacency` that corresponds to the module assigned to node `i`. To estimate `Conformity` and a matrix `IntermodularAdjacency`, the function attempts to minimize a quadratic loss function (sums of squares). Currently, the function only implements a heuristic algorithm for optimizing the objective function (chapter 2 of Horvath 2011). Another, more accurate Majorization Minorization (MM) algorithm for the decomposition is implemented in the function `propensityDecomposition` by Ranola et al (2011).

## Value

<code>A.CF</code>	a symmetric matrix that approximates the input matrix <code>adj</code> . Roughly speaking, the $i,j$ -th element of the matrix equals $Conformity[i] * Conformity[j] * IntermodularAdjacency[module.index[i], module.index[j]]$ where <code>module.index[i]</code> is the row of the matrix <code>IntermodularAdjacency</code> that corresponds to the module assigned to node <code>i</code> .
<code>Conformity</code>	a numeric vector whose entries correspond to the rows of <code>codeadj</code> . If <code>C1=NULL</code> then <code>Conformity[i]</code> is the conformity. If <code>C1</code> is not <code>NULL</code> then <code>Conformity[i]</code> is the intramodular conformity with respect to the module that node <code>i</code> belongs to.
<code>IntermodularAdjacency</code>	a symmetric matrix (data frame) whose rows and columns correspond to the number of modules specified in <code>C1</code> . Interpretation: it measures the similarity (adjacency) between the modules. In this case, the rows (and columns) of <code>IntermodularAdjacency</code> correspond to the entries of <code>C1.level</code> .
<code>Factorizability</code>	is a number between 0 and 1. If <code>C1=NULL</code> then it equals 1, if (and only if) <code>adj</code> is exactly factorizable. If <code>C1</code> is a vector, then it measures how well the module- and CF based decomposition approximates <code>adj</code> .
<code>C1.level</code>	is a vector of character strings which correspond to the factor levels of the module assignment <code>C1</code> . Incidentally, the function automatically turns <code>C1</code> into a factor variable. The components of <code>Conformity</code> and <code>IntramodularFactorizability</code> correspond to the entries of <code>C1.level</code> .
<code>IntramodularFactorizability</code>	is a numeric vector of length equal to the number of modules specified by <code>C1</code> . Its entries report the factorizability measure for each module. The components

correspond to the entries of `Cl.level1`.

`listConformity`

### Note

Regarding the situation when `Cl=NULL`. One can easily show that the conformity vector is not unique if `adj` contains only 2 nodes. However, for more than 2 nodes the conformity is uniquely defined when dealing with an exactly factorizable weighted network whose entries `adj[i, j]` are larger than 0. In this case, one can get explicit formulas for the conformity (Dong and Horvath 2007).

### Author(s)

Steve Horvath

### References

Dong J, Horvath S (2007) Understanding Network Concepts in Modules. *BMC Systems Biology* 2007, June 1:24  
 Horvath S, Dong J (2008) Geometric Interpretation of Gene Co-Expression Network Analysis. *PloS Computational Biology*. 4(8): e1000117. PMID: 18704157  
 Horvath S (2011) Weighted Network Analysis. Applications in Genomics and Systems Biology. Springer Book. ISBN: 978-1-4419-8818-8  
 Ranola JMO, Langfelder P, Song L, Horvath S, Lange K (2011) An MM algorithm for the module- and propensity based decomposition of a network. Currently a draft.

### See Also

[conformityBasedNetworkConcepts](#)

### Examples

```
# assume the number of nodes can be divided by 2 and by 3
n=6
# here is a perfectly factorizable matrix
A=matrix(1,nrow=n,ncol=n)
# this provides the conformity vector and factorizability measure
conformityDecomposition(adj=A)
# now assume we have a class assignment
Cl=rep(c(1,2),c(n/2,n/2))
conformityDecomposition(adj=A,Cl=Cl)
# here is a block diagonal matrix
blockdiag.A=A
blockdiag.A[1:(n/3),(n/3+1):n]=0
blockdiag.A[(n/3+1):n, 1:(n/3)]=0
block.Cl=rep(c(1,2),c(n/3,2*n/3))
conformityDecomposition(adj= blockdiag.A,Cl=block.Cl)

# another block diagonal matrix
blockdiag.A=A
blockdiag.A[1:(n/3),(n/3+1):n]=0.3
blockdiag.A[(n/3+1):n, 1:(n/3)]=0.3
block.Cl=rep(c(1,2),c(n/3,2*n/3))
conformityDecomposition(adj= blockdiag.A,Cl=block.Cl)
```

---

consensusDissTOMandTree

*Consensus clustering based on topological overlap and hierarchical clustering*

---

### Description

This function makes a consensus network using all of the default values in the WGCNA library. Details regarding how consensus modules are formed can be found here: <http://www.genetics.ucla.edu/labs/horvath/CoexpressionNetworkConstruction-man.pdf>

### Usage

```
consensusDissTOMandTree(multiExpr, softPower, TOM = NULL)
```

### Arguments

multiExpr	Expression data in the multi-set format (see checkSets). A vector of lists, one per set. Each set must contain a component data that contains the expression data. Rows correspond to samples and columns to genes or probes. Two or more sets of data must be included and adjacencies cannot be used.
softPower	Soft thresholding power used to make each of the networks in multiExpr.
TOM	A LIST of matrices holding the topological overlap corresponding to the sets in multiExpr, if they have already been calculated. Otherwise, keep TOM set as NULL (default), and TOM similarities will be calculated using the WGCNA defaults. If inputted, this variable must be a list with each entree a TOM corresponding to the same entries in multiExpr.

### Value

consensusTOM	The TOM difference matrix (1-TOM similarity) corresponding to the consensus network.
consTree	Returned value is the same as that of hclust: An object of class hclust which describes the tree produced by the clustering process. This tree corresponds to the dissimilarity matrix consensusTOM.

### Author(s)

Peter Langfelder, Steve Horvath, Jeremy Miller

### References

Langfelder P, Horvath S (2007) Eigengene networks for studying the relationships between co-expression modules. BMC Systems Biology 2007, 1:54

### See Also

[blockwiseConsensusModules](#)

**Examples**

```
# Example consensus network using two simulated data sets

set.seed      = 100
MEturquoise  = sample(1:100,50)
MEblue       = sample(1:100,50)
MEbrown      = sample(1:100,50)
MEyellow     = sample(1:100,50)
MEgreen      = sample(1:100,50)

ME = data.frame(MEturquoise, MEblue, MEbrown, MEyellow, MEGreen)
dat1 = simulateDatExpr(ME,300,c(0.2, 0.10, 0.10, 0.10, 0.10, 0.2), signed=TRUE)
dat2 = simulateDatExpr(ME,300,c(0.18, 0.11, 0.11, 0.09, 0.11, 0.23),signed=TRUE)
multiExpr = list(S1=list(data=dat1$datExpr),S2=list(data=dat2$datExpr))
softPower=8

consensusNetwork = consensusDissTOMandTree(multiExpr, softPower)
plotDendroAndColors(consensusNetwork$consTree, cbind(labels2colors(dat1$allLabels),
  labels2colors(dat2$allLabels)),c("S1","S2"), dendroLabels=FALSE)
```

---

consensusKME

*Calculate consensus kME (eigengene-based connectivities) across multiple data sets.*

---

**Description**

Calculate consensus kME (eigengene-based connectivities) across multiple data sets, typically following a consensus module analysis.

**Usage**

```
consensusKME(
  multiExpr,
  moduleLabels,
  multiEigengenes = NULL,
  consensusQuantile = 0,
  signed = TRUE,
  useModules = NULL,
  metaAnalysisWeights = NULL,
  corAndPvalueFnc = corAndPvalue, corOptions = list(), corComponent = "cor",
  getQvalues = FALSE,
  useRankPvalue = TRUE,
  rankPvalueOptions = list(calculateQvalue = getQvalues, pValueMethod = "scale"),
  setNames = NULL,
  excludeGrey = TRUE, greyLabel = ifelse(is.numeric(moduleLabels), 0, "grey"))
```

**Arguments**

**multiExpr** Expression (or other numeric) data in a multi-set format. A vector of lists; in each list there must be a component named 'data' whose content is a matrix or dataframe or array of dimension 2.

moduleLabels	Module labels: one label for each gene in multiExpr.
multiEigengenes	Optional eigengenes of modules specified in moduleLabels. If not given, will be calculated from multiExpr.
signed	logical: should the network be considered signed? In signed networks (TRUE), negative kME values are not considered significant and the corresponding p-values will be one-sided. In unsigned networks (FALSE), negative kME values are considered significant and the corresponding p-values will be two-sided.
useModules	Optional specification of module labels to which the analysis should be restricted. This could be useful if there are many modules, most of which are not interesting. Note that the "grey" module cannot be used with useModules.
consensusQuantile	Quantile for the consensus calculation. Should be a number between 0 (minimum) and 1.
metaAnalysisWeights	Optional specification of meta-analysis weights for each input set. If given, must be a numeric vector of length equal the number of input data sets (i.e., length(multiExpr)). These weights will be used in addition to constant weights and weights proportional to number of samples (observations) in each set.
corAndPvalueFnc	Function that calculates associations between expression profiles and eigengenes. See details.
corOptions	List giving additional arguments to function corAndPvalueFnc. See details.
corComponent	Name of the component of output of corAndPvalueFnc that contains the actual correlation.
getQvalues	logical: should q-values (estimates of FDR) be calculated?
useRankPvalue	Logical: should the rankPvalue function be used to obtain alternative meta-analysis statistics?
rankPvalueOptions	Additional options for function rankPvalue. These include na.last (default "keep"), ties.method (default "average"), calculateQvalue (default copied from input getQvalues), and pvalueMethod (default "scale"). See the help file for rankPvalue for full details.
setNames	names for the input sets. If not given, will be taken from names(multiExpr). If those are NULL as well, the names will be "Set_1", "Set_2", ....
excludeGrey	logical: should the grey module be excluded from the kME tables? Since the grey module is typically not a real module, it makes little sense to report kME values for it.
greyLabel	label that labels the grey module.

## Details

The function corAndPvalueFnc is currently is expected to accept arguments x (gene expression profiles), y (eigengene expression profiles), and alternative with possibilities at least "greater", "two.sided". Any additional arguments can be passed via corOptions.

The function corAndPvalueFnc should return a list which at the least contains (1) a matrix of associations of genes and eigengenes (this component should have the name given by corComponent), and (2) a matrix of the corresponding p-values, named "p" or "p.value". Other components are optional but for full functionality should include (3) nObs giving the number of observations for



each association (which is the number of samples less number of missing data - this can in principle vary from association to association), and (4) Z giving a Z static for each observation. If these are missing, nObs is calculated in the main function, and calculations using the Z statistic are skipped.

## Value

Data frame with the following components (for easier readability the order here is not the same as in the actual output):

ID	Gene ID, taken from the column names of the first input data set
consensus.kME.1, consensus.kME.2, ...	Consensus kME (that is, the requested quantile of the kMEs in the individual data sets) in each module for each gene across the input data sets. The module labels (here 1, 2, etc.) correspond to those in <code>moduleLabels</code> .
weightedAverage.equalWeights.kME1, weightedAverage.equalWeights.kME2, ...	Average kME in each module for each gene across the input data sets.
weightedAverage.RootDoFWeights.kME1, weightedAverage.RootDoFWeights.kME2, ...	Weighted average kME in each module for each gene across the input data sets. The weight of each data set is proportional to the square root of the number of samples in the set.
weightedAverage.DoFWeights.kME1, weightedAverage.DoFWeights.kME2, ...	Weighted average kME in each module for each gene across the input data sets. The weight of each data set is proportional to number of samples in the set.
weightedAverage.userWeights.kME1, weightedAverage.userWeights.kME2, ...	(Only present if input <code>metaAnalysisWeights</code> is non-NULL.) Weighted average kME in each module for each gene across the input data sets. The weight of each data set is given in <code>metaAnalysisWeights</code> .
meta.Z.equalWeights.kME1, meta.Z.equalWeights.kME2, ...	Meta-analysis Z statistic for kME in each module, obtained by weighing the Z scores in each set equally. Only returned if the function <code>corAndPvalueFnc</code> returns the Z statistics corresponding to the correlations.
meta.Z.RootDoFWeights.kME1, meta.Z.RootDoFWeights.kME2, ...	Meta-analysis Z statistic for kME in each module, obtained by weighing the Z scores in each set by the square root of the number of samples. Only returned if the function <code>corAndPvalueFnc</code> returns the Z statistics corresponding to the correlations.
meta.Z.DoFWeights.kME1, meta.Z.DoFWeights.kME2, ...	Meta-analysis Z statistic for kME in each module, obtained by weighing the Z scores in each set by the number of samples. Only returned if the function <code>corAndPvalueFnc</code> returns the Z statistics corresponding to the correlations.
meta.Z.userWeights.kME1, meta.Z.userWeights.kME2, ...	Meta-analysis Z statistic for kME in each module, obtained by weighing the Z scores in each set by <code>metaAnalysisWeights</code> . Only returned if <code>metaAnalysisWeights</code> is non-NULL and the function <code>corAndPvalueFnc</code> returns the Z statistics corresponding to the correlations.
meta.p.equalWeights.kME1, meta.p.equalWeights.kME2, ...	p-values obtained from the equal-weight meta-analysis Z statistics. Only returned if the function <code>corAndPvalueFnc</code> returns the Z statistics corresponding to the correlations.
meta.p.RootDoFWeights.kME1, meta.p.RootDoFWeights.kME2, ...	p-values obtained from the meta-analysis Z statistics with weights proportional to the square root of the number of samples. Only returned if the function <code>corAndPvalueFnc</code> returns the Z statistics corresponding to the correlations.

meta.p.DoFWeights.kME1, meta.p.DoFWeights.kME2, ...  
 p-values obtained from the degree-of-freedom weight meta-analysis Z statistics. Only returned if the function corAndPvalueFnc returns the Z statistics corresponding to the correlations.

meta.p.userWeights.kME1, meta.p.userWeights.kME2, ...  
 p-values obtained from the user-supplied weight meta-analysis Z statistics. Only returned if metaAnalysisWeights is non-NULL and the function corAndPvalueFnc returns the Z statistics corresponding to the correlations.

meta.q.equalWeights.kME1, meta.q.equalWeights.kME2, ...  
 q-values obtained from the equal-weight meta-analysis p-values. Only present if getQvalues is TRUE and the function corAndPvalueFnc returns the Z statistics corresponding to the kME values.

meta.q.RootDoFWeights.kME1, meta.q.RootDoFWeights.kME2, ...  
 q-values obtained from the meta-analysis p-values with weights proportional to the square root of the number of samples. Only present if getQvalues is TRUE and the function corAndPvalueFnc returns the Z statistics corresponding to the kME values.

meta.q.DoFWeights.kME1, meta.q.DoFWeights.kME2, ...  
 q-values obtained from the degree-of-freedom weight meta-analysis p-values. Only present if getQvalues is TRUE and the function corAndPvalueFnc returns the Z statistics corresponding to the kME values.

meta.q.userWeights.kME1, meta.q.userWeights.kME2, ...  
 q-values obtained from the user-specified weight meta-analysis p-values. Only present if metaAnalysisWeights is non-NULL, getQvalues is TRUE and the function corAndPvalueFnc returns the Z statistics corresponding to the kME values.

The next set of columns contain the results of function `rankPvalue` and are only present if input `useRankPvalue` is TRUE. Some columns may be missing depending on the options specified in `rankPvalueOptions`. We explicitly list columns that are based on weighing each set equally; names of these columns carry the suffix `.equalWeights`

pValueExtremeRank.ME1.equalWeights, pValueExtremeRank.ME2.equalWeights, ...  
 This is the minimum between pValueLowRank and pValueHighRank, i.e.  $\min(\text{pValueLow}, \text{pValueHigh})$

pValueLowRank.ME1.equalWeights, pValueLowRank.ME2.equalWeights, ...  
 Asymptotic p-value for observing a consistently low value across the columns of datS based on the rank method.

pValueHighRank.ME1.equalWeights, pValueHighRank.ME2.equalWeights, ...  
 Asymptotic p-value for observing a consistently low value across the columns of datS based on the rank method.

pValueExtremeScale.ME1.equalWeights, pValueExtremeScale.ME2.equalWeights, ...  
 This is the minimum between pValueLowScale and pValueHighScale, i.e.  $\min(\text{pValueLow}, \text{pValueHigh})$

pValueLowScale.ME1.equalWeights, pValueLowScale.ME2.equalWeights, ...  
 Asymptotic p-value for observing a consistently low value across the columns of datS based on the Scale method.

pValueHighScale.ME1.equalWeights, pValueHighScale.ME2.equalWeights, ...  
 Asymptotic p-value for observing a consistently low value across the columns of datS based on the Scale method.

qValueExtremeRank.ME1.equalWeights, qValueExtremeRank.ME2.equalWeights, ...  
 local false discovery rate (q-value) corresponding to the p-value pValueExtremeRank

`qValueLowRank.ME1.equalWeights`, `qValueLowRank.ME2.equalWeights`, ...  
 local false discovery rate (q-value) corresponding to the p-value `pValueLowRank`

`qValueHighRank.ME1.equalWeights`, `qValueHighRank.ME2.equalWeights`, ...  
 local false discovery rate (q-value) corresponding to the p-value `pValueHighRank`

`qValueExtremeScale.ME1.equalWeights`, `qValueExtremeScale.ME2.equalWeights`, ...  
 local false discovery rate (q-value) corresponding to the p-value `pValueExtremeScale`

`qValueLowScale.ME1.equalWeights`, `qValueLowScale.ME2.equalWeights`, ...  
 local false discovery rate (q-value) corresponding to the p-value `pValueLowScale`

`qValueHighScale.ME1.equalWeights`, `qValueHighScale.ME2.equalWeights`, ...  
 local false discovery rate (q-value) corresponding to the p-value `pValueHighScale`

... Analogous columns corresponding to weighing individual sets by the square root of the number of samples, by number of samples, and by user weights (if given). The corresponding column name suffixes are `.RootDoFWWeights`, `.DoFWWeights`, and `.userWeights`.

The following set of columns summarize kME in individual input data sets.

`kME1.Set_1`, `kME1.Set_2`, ..., `kME2.Set_1`, `kME2.Set_2`, ...  
 kME values for each gene in each module in each given data set.

`p.kME1.Set_1`, `p.kME1.Set_2`, ..., `p.kME2.Set_1`, `p.kME2.Set_2`, ...  
 p-values corresponding to kME values for each gene in each module in each given data set.

`q.kME1.Set_1`, `q.kME1.Set_2`, ..., `q.kME2.Set_1`, `q.kME2.Set_2`, ...  
 q-values corresponding to kME values for each gene in each module in each given data set. Only returned if `getQvalues` is TRUE.

`Z.kME1.Set_1`, `Z.kME1.Set_2`, ..., `Z.kME2.Set_1`, `Z.kME2.Set_2`, ...  
 Z statistics corresponding to kME values for each gene in each module in each given data set. Only present if the function `corAndPvalueFnc` returns the Z statistics corresponding to the kME values.

### Author(s)

Peter Langfelder

### References

Langfelder P, Horvath S., WGCNA: an R package for weighted correlation network analysis. *BMC Bioinformatics*. 2008 Dec 29; 9:559.

### See Also

[signedKME](#) for eigengene based connectivity in a single data set. [corAndPvalue](#), [bicorAndPvalue](#) for two alternatives for calculating correlations and the corresponding p-values and Z scores. Both can be used with this function.

---

`consensusMEDissimilarity`*Consensus dissimilarity of module eigengenes.*

---

**Description**

Calculates consensus dissimilarity (1-cor) of given module eigengenes realized in several sets.

**Usage**

```
consensusMEDissimilarity(MEs, useAbs = FALSE, useSets = NULL, method = "consensus")
```

**Arguments**

<code>MEs</code>	Module eigengenes of the same modules in several sets.
<code>useAbs</code>	Controls whether absolute value of correlation should be used instead of correlation in the calculation of dissimilarity.
<code>useSets</code>	If the consensus is to include only a selection of the given sets, this vector (or scalar in the case of a single set) can be used to specify the selection. If NULL, all sets will be used.
<code>method</code>	A character string giving the method to use. Allowed values are (abbreviations of) "consensus" and "majority". The consensus dissimilarity is calculated as the minimum of given set dissimilarities for "consensus" and as the average for "majority".

**Details**

This function calculates the individual set dissimilarities of the given eigengenes in each set, then takes the (parallel) maximum or average over all sets. For details on the structure of input data, see [checkSets](#).

**Value**

A dataframe containing the matrix of dissimilarities, with names and rownames set appropriately.

**Author(s)**

Peter Langfelder, <Peter.Langfelder@gmail.com>

**See Also**

[checkSets](#)

---

consensusOrderMEs      *Put close eigenvectors next to each other in several sets.*

---

## Description

Reorder given (eigen-)vectors such that similar ones (as measured by correlation) are next to each other. This is a multi-set version of [orderMEs](#); the dissimilarity used can be of consensus type (for each pair of eigenvectors the consensus dissimilarity is the maximum of individual set dissimilarities over all sets) or of majority type (for each pair of eigenvectors the consensus dissimilarity is the average of individual set dissimilarities over all sets).

## Usage

```
consensusOrderMEs(MEs, useAbs = FALSE, useSets = NULL,
  greyLast = TRUE,
  greyName = paste(moduleColor.getMEprefix(), "grey", sep=""),
  method = "consensus")
```

## Arguments

MEs	Module eigengenes of several sets in a multi-set format (see <a href="#">checkSets</a> ). A vector of lists, with each list corresponding to one dataset and the module eigengenes in the component data, that is <code>MEs[[set]]\$data[sample, module]</code> is the expression of the eigengene of module <code>module</code> in sample <code>sample</code> in dataset <code>set</code> . The number of samples can be different between the sets, but the modules must be the same.
useAbs	Controls whether vector similarity should be given by absolute value of correlation or plain correlation.
useSets	Allows the user to specify for which sets the eigengene ordering is to be performed.
greyLast	Normally the color grey is reserved for unassigned genes; hence the grey module is not a proper module and it is conventional to put it last. If this is not desired, set the parameter to FALSE.
greyName	Name of the grey module eigengene.
method	A character string giving the method to be used calculating the consensus dissimilarity. Allowed values are (abbreviations of) "consensus" and "majority". The consensus dissimilarity is calculated as the maximum of given set dissimilarities for "consensus" and as the average for "majority".

## Details

Ordering module eigengenes is useful for plotting purposes. This function calculates the consensus or majority dissimilarity of given eigengenes over the sets specified by `useSets` (defaults to all sets). A hierarchical dendrogram is calculated using the dissimilarity and the order given by the dendrogram is used for the eigengenes in all other sets.

## Value

A vector of lists of the same type as `MEs` containing the re-ordered eigengenes.

**Author(s)**

Peter Langfelder, <Peter.Langfelder@gmail.com>

**See Also**

[moduleEigengenes](#), [multiSetMEs](#), [orderMEs](#)

---

consensusProjectiveKMeans

*Consensus projective K-means (pre-)clustering of expression data*

---

**Description**

Implementation of a consensus variant of K-means clustering for expression data across multiple data sets.

**Usage**

```
consensusProjectiveKMeans(
  multiExpr,
  preferredSize = 5000,
  nCenters = NULL,
  sizePenaltyPower = 4,
  networkType = "unsigned",
  randomSeed = 54321,
  checkData = TRUE,
  useMean = (length(multiExpr) > 3),
  maxIterations = 1000,
  verbose = 0, indent = 0)
```

**Arguments**

multiExpr	expression data in the multi-set format (see <a href="#">checkSets</a> ). A vector of lists, one per set. Each set must contain a component data that contains the expression data, with rows corresponding to samples and columns to genes or probes.
preferredSize	preferred maximum size of clusters.
nCenters	number of initial clusters. Empirical evidence suggests that more centers will give a better preclustering; the default is <code>as.integer(min(nGenes/20, preferredSize^2/nGenes))</code> and is an attempt to arrive at a reasonable number given the resources available.
sizePenaltyPower	parameter specifying how severe is the penalty for clusters that exceed preferredSize.
networkType	network type. Allowed values are (unique abbreviations of) "unsigned", "signed", "signed hybrid". See <a href="#">adjacency</a> .
randomSeed	integer to be used as seed for the random number generator before the function starts. If a current seed exists, it is saved and restored upon exit.
checkData	logical: should data be checked for genes with zero variance and genes and samples with excessive numbers of missing samples? Bad samples are ignored; returned cluster assignment for bad genes will be NA.

useMean	logical: should mean distance across sets be used instead of maximum? See details.
maxIterations	maximum iterations to be attempted.
verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.
indent	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

## Details

The principal aim of this function within WGCNA is to pre-cluster a large number of genes into smaller blocks that can be handled using standard WGCNA techniques.

This function implements a variant of K-means clustering that is suitable for co-expression analysis. Cluster centers are defined by the first principal component, and distances by correlation. Consensus distance across several sets is defined as the maximum of the corresponding distances in individual sets; however, if useMean is set, the mean distance will be used instead of the maximum. The distance between a gene and a center of a cluster is multiplied by a factor of  $\max(\text{clusterSize}/\text{preferredSize}, 1)^{\text{sizePenal}}$  thus penalizing clusters whose size exceeds preferredSize. The function starts with randomly generated cluster assignment (hence the need to set the random seed for repeatability) and executes iterations of calculating new centers and reassigning genes to nearest (in the consensus sense) center until the clustering becomes stable. Before returning, nearby clusters are iteratively combined if their combined size is below preferredSize.

Consensus distance defined as maximum of distances in all sets is consistent with the approach taken in [blockwiseConsensusModules](#), but the procedure may not converge. Hence it is advisable to use the mean as consensus in cases where there are multiple data sets (4 or more, say) and/or if the input data sets are very different.

The standard principal component calculation via the function svd fails from time to time (likely a convergence problem of the underlying lapack functions). Such errors are trapped and the principal component is approximated by a weighted average of expression profiles in the cluster. If verbose is set above 2, an informational message is printed whenever this approximation is used.

## Value

A list with the following components:

clusters	a numerical vector with one component per input gene, giving the cluster number in which the gene is assigned.
centers	a vector of lists, one list per set. Each list contains a component data that contains a matrix whose columns are the cluster centers in the corresponding set.
unmergedClusters	a numerical vector with one component per input gene, giving the cluster number in which the gene was assigned before the final merging step.
unmergedCenters	a vector of lists, one list per set. Each list contains a component data that contains a matrix whose columns are the cluster centers before merging in the corresponding set.

## Author(s)

Peter Langfelder

**See Also**[projectiveKMeans](#)

---

`cor`*Fast calculations of Pearson correlation.*

---

**Description**

These functions implements a faster calculation of Pearson correlation.

The speedup against the R's standard `cor` function will be substantial particularly if the input matrix only contains a small number of missing data. If there are no missing data, or the missing data are numerous, the speedup will be smaller but still present.

**Usage**

```
cor(x, y = NULL,
    use = "all.obs",
    method = c("pearson", "kendall", "spearman"),
    quick = 0,
    cosine = FALSE,
    cosineX = cosine,
    cosineY = cosine,
    nThreads = 0,
    verbose = 0, indent = 0)
```

```
corFast(x, y = NULL,
        use = "all.obs",
        quick = 0, nThreads = 0,
        verbose = 0, indent = 0)
```

```
cor1(x, use = "all.obs", verbose = 0, indent = 0)
```

**Arguments**

<code>x</code>	a numeric vector or a matrix. If <code>y</code> is null, <code>x</code> must be a matrix.
<code>y</code>	a numeric vector or a matrix. If not given, correlations of columns of <code>x</code> will be calculated.
<code>use</code>	a character string specifying the handling of missing data. The fast calculations currently support <code>"all.obs"</code> and <code>"pairwise.complete.obs"</code> ; for other options, see R's standard correlation function <code>cor</code> . Abbreviations are allowed.
<code>method</code>	a character string specifying the method to be used. Fast calculations are currently available only for <code>"pearson"</code> .
<code>quick</code>	real number between 0 and 1 that controls the precision of handling of missing data in the calculation of correlations. See details.
<code>cosine</code>	logical: calculate cosine correlation? Only valid for <code>method="pearson"</code> . Cosine correlation is similar to Pearson correlation but the mean subtraction is not performed. The result is the cosine of the angle(s) between (the columns of) <code>x</code> and <code>y</code> .



cosineX	logical: use the cosine calculation for x? This setting does not affect y and can be used to give a hybrid cosine-standard correlation.
cosineY	logical: use the cosine calculation for y? This setting does not affect x and can be used to give a hybrid cosine-standard correlation.
nThreads	non-negative integer specifying the number of parallel threads to be used by certain parts of correlation calculations. This option only has an effect on systems on which a POSIX thread library is available (which currently includes Linux and Mac OSX, but excludes Windows). If zero, the number of online processors will be used if it can be determined dynamically, otherwise correlation calculations will use 2 threads.
verbose	Controls the level of verbosity. Values above zero will cause a small amount of diagnostic messages to be printed.
indent	Indentation of printed diagnostic messages. Each unit above zero adds two spaces.

### Details

The fast calculations are currently implemented only for `method="pearson"` and use either `"all.obs"` or `"pairwise.complete.obs"`. The `corFast` function is a wrapper that calls the function `cor`. If the combination of `method` and `use` is implemented by the fast calculations, the fast code is executed; otherwise, R's own correlation `cor` is executed.

The argument `quick` specifies the precision of handling of missing data. Zero will cause all calculations to be executed precisely, which may be significantly slower than calculations without missing data. Progressively higher values will speed up the calculations but introduce progressively larger errors. Without missing data, all column means and variances can be pre-calculated before the covariances are calculated. When missing data are present, exact calculations require the column means and variances to be calculated for each covariance. The approximate calculation uses the pre-calculated mean and variance and simply ignores missing data in the covariance calculation. If the number of missing data is high, the pre-calculated means and variances may be very different from the actual ones, thus potentially introducing large errors. The `quick` value times the number of rows specifies the maximum difference in the number of missing entries for mean and variance calculations on the one hand and covariance on the other hand that will be tolerated before a recalculation is triggered. The hope is that if only a few missing data are treated approximately, the error introduced will be small but the potential speedup can be significant.

### Value

The matrix of the Pearson correlations of the columns of `x` with columns of `y` if `y` is given, and the correlations of the columns of `x` if `y` is not given.

### Note

The implementation uses the BLAS library matrix multiplication function for the most expensive step of the calculation. Using a tuned, architecture-specific BLAS may significantly improve the performance of this function.

The values returned by the `corFast` function may differ from the values returned by R's function `cor` by rounding errors on the order of  $1e-15$ .

### Author(s)

Peter Langfelder

## References

Peter Langfelder, Steve Horvath (2012) Fast R Functions for Robust Correlations and Hierarchical Clustering. *Journal of Statistical Software*, 46(11), 1-17. <http://www.jstatsoft.org/v46/i11/>

## See Also

R's standard Pearson correlation function `cor`.

## Examples

```
## Test the speedup compared to standard function cor

# Generate a random matrix with 200 rows and 1000 columns

set.seed(10)
nrow = 100;
ncol = 500;
data = matrix(rnorm(nrow*ncol), nrow, ncol);

## First test: no missing data

system.time( {corStd = stats::cor(data)} );

system.time( {corFast = cor(data)} );

all.equal(corStd, corFast)

# Here R's standard correlation performs very well.

# We now add a few missing entries.

data[sample(nrow, 10), 1] = NA;

# And test the correlations again...

system.time( {corStd = stats::cor(data, use = 'p')} );

system.time( {corFast = cor(data, use = 'p')} );

all.equal(corStd, corFast)

# Here the R's standard correlation slows down considerably
# while corFast still retains it speed. Choosing
# higher ncol above will make the difference more pronounced.
```

---

corAndPvalue

*Calculation of correlations and associated p-values*

---

## Description

A faster, one-step calculation of Student correlation p-values for multiple correlations, properly taking into account the actual number of observations.

## Usage

```
corAndPvalue(x, y = NULL,  
             use = "pairwise.complete.obs",  
             alternative = c("two.sided", "less", "greater"),  
             ...)
```

## Arguments

x	a vector or a matrix
y	a vector or a matrix. If NULL, the correlation of columns of x will be calculated.
use	determines handling of missing data. See <code>cor</code> for details.
alternative	specifies the alternative hypothesis and must be (a unique abbreviation of) one of "two.sided", "greater" or "less". the initial letter. "greater" corresponds to positive association, "less" to negative association.
...	other arguments to the function <code>cor</code> .

## Details

The function calculates correlations of a matrix or of two matrices and the corresponding Student p-values. The output is not as full-featured as `cor.test`, but can work with matrices as input.

## Value

A list with the following components, each a matrix:

cor	the calculated correlations
p	the Student p-values corresponding to the calculated correlations
Z	Fisher transforms of the calculated correlations
t	Student t statistics of the calculated correlations
nObs	Numbers of observations for the correlation, p-values etc.

## Author(s)

Peter Langfelder and Steve Horvath

## References

Peter Langfelder, Steve Horvath (2012) Fast R Functions for Robust Correlations and Hierarchical Clustering. Journal of Statistical Software, 46(11), 1-17. <http://www.jstatsoft.org/v46/i11/>

## See Also

`cor` for calculation of correlations only;

`cor.test` for another function for significance test of correlations



**See Also**

[relativeCorPredictionSuccess](#)

---

corPvalueFisher      *Fisher's asymptotic p-value for correlation*

---

**Description**

Calculates Fisher's asymptotic p-value for given correlations.

**Usage**

```
corPvalueFisher(cor, nSamples, twoSided = TRUE)
```

**Arguments**

cor	A vector of correlation values whose corresponding p-values are to be calculated
nSamples	Number of samples from which the correlations were calculated
twoSided	logical: should the calculated p-values be two sided?

**Value**

A vector of p-values of the same length as the input correlations.

**Author(s)**

Steve Horvath and Peter Langfelder

---

corPvalueStudent      *Student asymptotic p-value for correlation*

---

**Description**

Calculates Student asymptotic p-value for given correlations.

**Usage**

```
corPvalueStudent(cor, nSamples)
```

**Arguments**

cor	A vector of correlation values whose corresponding p-values are to be calculated
nSamples	Number of samples from which the correlations were calculated

**Value**

A vector of p-values of the same length as the input correlations.

**Author(s)**

Steve Horvath and Peter Langfelder

---

`correlationPreservation`*Preservation of eigengene correlations*

---

**Description**

Calculates a summary measure of preservation of eigengene correlations across data sets

**Usage**

```
correlationPreservation(multiME, setLabels, excludeGrey = TRUE, greyLabel = "grey")
```

**Arguments**

<code>multiME</code>	consensus module eigengenes in a multi-set format. A vector of lists with one list corresponding to each set. Each list must contain a component data that is a data frame whose columns are consensus module eigengenes.
<code>setLabels</code>	names to be used for the sets represented in <code>multiME</code> .
<code>excludeGrey</code>	logical: exclude the 'grey' eigengene from preservation measure?
<code>greyLabel</code>	module label corresponding to the 'grey' module. Usually this will be the character string "grey" if the labels are colors, and the number 0 if the labels are numeric.

**Details**

The function calculates the preservation of correlation of each eigengene with all other eigengenes (optionally except the 'grey' eigengene) in all pairs of sets.

**Value**

A data frame whose rows correspond to consensus module eigengenes given in the input `multiME`, and columns correspond to all possible set comparisons. The two sets compared in each column are indicated in the column name.

**Author(s)**

Peter Langfelder

**References**

Langfelder P, Horvath S (2007) Eigengene networks for studying the relationships between co-expression modules. *BMC Systems Biology* 2007, 1:54

**See Also**

[multiSetMEs](#) and [modulecheckSets](#) in package `moduleColor` for more on eigengenes and the multi-set format

---

 coxRegressionResiduals

*Deviance- and martingale residuals from a Cox regression model*


---

### Description

The function inputs a censored time variable which is specified by two input variables `time` and `event`. It outputs i) the martingale residual and ii) deviance residual corresponding to a Cox regression model. By default, the Cox regression model is an intercept only Cox regression model. But optionally, the user can input covariates using the argument `datCovariates`. The function makes use of the `coxph` function in the survival library. See `help(residuals.coxph)` to learn more.

### Usage

```
coxRegressionResiduals(time, event, datCovariates = NULL)
```

### Arguments

<code>time</code>	is a numeric variable that contains follow up time or time to event.
<code>event</code>	is a binary variable that takes on values 1 and 0. 1 means that the event took place (e.g. person died, or tumor recurred). 0 means censored, i.e. event has not yet been observed or loss to follow up.
<code>datCovariates</code>	a data frame whose columns correspond to covariates that should be used in the Cox regression model. By default, the only covariate the intercept term 1.

### Details

Residuals are often used to investigate the lack of fit of a model. For Cox regression, there is no easy analog to the usual "observed minus predicted" residual of linear regression. Instead, several specialized residuals have been proposed for Cox regression analysis. The function calculates residuals that are well defined for an intercept only Cox regression model: the martingale and deviance residuals (Therneau et al 1990). The martingale residual of a subject (person) specifies excess failures beyond the expected baseline hazard. For example, a subject who was censored at 3 years, and whose predicted cumulative hazard at 3 years was 30. Another subject who had an event at 10 years, and whose predicted cumulative hazard at 10 years was 60. Since martingale residuals are not symmetrically distributed, even when the fitted model is correct, it is often advantageous to transform them into more symmetrically distributed residuals: deviance residuals. Thus, deviance residuals are defined as transformations of the martingale residual and the event variable. Deviance residuals are often symmetrically distributed around zero. Deviance Residuals are similar to residuals from ordinary linear regression in that they are symmetrically distributed around 0 and have standard deviation of 1.0. A subject with a large deviance residual is poorly predicted by the model, i.e. is different from the baseline cumulative hazard. A negative value indicates a longer than expected survival time. When covariates are specified in `datCovariates`, then one can plot deviance (or martingale) residuals against the covariates. Unusual patterns may indicate poor fit of the Cox model. Cryptic comments: Deviance (or martingale) residuals can sometimes be used as (uncensored) quantitative variables instead of the original time censored variable. For example, they could be used as outcome in a regression tree or regression forest predictor.

### Value

It outputs a data frame with 2 columns. The first and second column correspond to martingale and deviance residuals respectively.

**Note**

This function can be considered a wrapper of the `coxph` function.

**Author(s)**

Steve Horvath

**References**

Therneau TM, Grambsch PM, Fleming TR (1990) Martingale-based residuals for survival models. *Biometrika* (1990), 77, 1, pp. 147-60

**Examples**

```
library(survival)
# simulate time and event data
time1=sample(1:100)
event1=sample(c(1,0), 100,replace=TRUE)

event1[1:5]=NA
time1[1:5]=NA
# no covariates
datResiduals= coxRegressionResiduals(time=time1,event=event1)

# now we simulate a covariate
z= rnorm(100)
cor(datResiduals,use="p")
datResiduals=coxRegressionResiduals(time=time1,event=event1,datCovariates=data.frame(z))
cor(datResiduals,use="p")
```

---

cutreeStatic

*Constant-height tree cut*

---

**Description**

Module detection in hierarchical dendrograms using a constant-height tree cut. Only branches whose size is at least `minSize` are retained.

**Usage**

```
cutreeStatic(dendro, cutHeight = 0.9, minSize = 50)
```

**Arguments**

<code>dendro</code>	a hierarchical clustering dendrogram such as returned by <a href="#">hclust</a> .
<code>cutHeight</code>	height at which branches are to be cut.
<code>minSize</code>	minimum number of object on a branch to be considered a cluster.



**Details**

This function performs a straightforward constant-height cut as implemented by [cutree](#), then calculates the number of objects on each branch and only keeps branches that have at least `minSize` objects on them.

**Value**

A numeric vector giving labels of objects, with 0 meaning unassigned. The largest cluster is conventionally labeled 1, the next largest 2, etc.

**Author(s)**

Peter Langfelder

**See Also**

[hclust](#) for hierarchical clustering, [cutree](#) and [cutreeStatic](#) for other constant-height branch cuts, [standardColors](#) to convert the returned numerical labels into colors for easier visualization.

---

cutreeStaticColor	<i>Constant height tree cut using color labels</i>
-------------------	--

---

**Description**

Cluster detection by a constant height cut of a hierarchical clustering dendrogram.

**Usage**

```
cutreeStaticColor(dendro, cutHeight = 0.9, minSize = 50)
```

**Arguments**

<code>dendro</code>	a hierarchical clustering dendrogram such as returned by <a href="#">hclust</a> .
<code>cutHeight</code>	height at which branches are to be cut.
<code>minSize</code>	minimum number of object on a branch to be considered a cluster.

**Details**

This function performs a straightforward constant-height cut as implemented by [cutree](#), then calculates the number of objects on each branch and only keeps branches that have at least `minSize` objects on them.

**Value**

A character vector giving color labels of objects, with "grey" meaning unassigned. The largest cluster is conventionally labeled "turquoise", next "blue" etc. Run `standardColors()` to see the sequence of standard color labels.

**Author(s)**

Peter Langfelder

**See Also**

[hclust](#) for hierarchical clustering, [cutree](#) and [cutreeStatic](#) for other constant-height branch cuts, [standardColors](#) to see the sequence of color labels that can be assigned.

---

displayColors	<i>Show colors used to label modules</i>
---------------	--

---

**Description**

The function plots a barplot using colors that label modules.

**Usage**

```
displayColors(colors = NULL)
```

**Arguments**

`colors` colors to be displayed. Defaults to all colors available for module labeling.

**Details**

To see the first `n` colors, use argument `colors = standardColors(n)`.

**Value**

None.

**Author(s)**

Peter Langfelder

**See Also**

[standardColors](#)

**Examples**

```
displayColors(standardColors(10))
```

---

dynamicMergeCut	<i>Threshold for module merging</i>
-----------------	-------------------------------------

---

**Description**

Calculate a suitable threshold for module merging based on the number of samples and a desired Z quantile.

**Usage**

```
dynamicMergeCut(n, mergeCor = 0.9, Zquantile = 2.35)
```

**Arguments**

n	number of samples
mergeCor	theoretical correlation threshold for module merging
Zquantile	Z quantile for module merging

**Details**

This function calculates the threshold for module merging. The threshold is calculated as the lower boundary of the interval around the theoretical correlation mergeCor whose width is given by the Z value Zquantile.

**Value**

The correlation threshold for module merging; a single number.

**Author(s)**

Steve Horvath

**See Also**

[moduleEigengenes](#), [mergeCloseModules](#)

**Examples**

```
dynamicMergeCut(20)  
dynamicMergeCut(50)  
dynamicMergeCut(100)
```

---

 exportNetworkToCytoscape

*Export network to Cytoscape*


---

### Description

This function exports a network in edge and node list files in a format suitable for importing to Cytoscape.

### Usage

```
exportNetworkToCytoscape(
  adjMat,
  edgeFile = NULL,
  nodeFile = NULL,
  weighted = TRUE,
  threshold = 0.5,
  nodeNames = NULL,
  altNodeNames = NULL,
  nodeAttr = NULL,
  includeColNames = TRUE)
```

### Arguments

adjMat	adjacency matrix giving connection strengths among the nodes in the network.
edgeFile	file name of the file to contain the edge information.
nodeFile	file name of the file to contain the node information.
weighted	logical: should the exported network be weighted?
threshold	adjacency threshold for including edges in the output.
nodeNames	names of the nodes. If not given, dimnames of adjMat will be used.
altNodeNames	optional alternate names for the nodes, for example gene names if nodes are labeled by probe IDs.
nodeAttr	optional node attribute, for example module color. Can be a vector or a data frame.
includeColNames	logical: should column names be included in the output files? Note that Cytoscape can read files both with and without column names.

### Details

If the corresponding file names are supplied, the edge and node data is written to the appropriate files. The edge and node data is also returned as return value (see below).

### Value

A list with the following components:

edgeData	a data frame containing the edge data, with one row per edge
nodeData	a data frame containing the node data, with one row per node

**Author(s)**

Peter Langfelder

**See Also**

[exportNetworkToVisANT](#)

---

exportNetworkToVisANT *Export network data in format readable by VisANT*

---

**Description**

Exports network data in a format readable and displayable by the VisANT software.

**Usage**

```
exportNetworkToVisANT(  
  adjMat,  
  file = NULL,  
  weighted = TRUE,  
  threshold = 0.5,  
  probeToGene = NULL)
```

**Arguments**

adjMat	adjacency matrix of the network to be exported.
file	character string specifying the file name of the file in which the data should be written. If not given, no file will be created. The file is in a plain text format.
weighted	logical: should the exported network be weighted?
threshold	adjacency threshold for including edges in the output.
probeToGene	optional specification of a conversion between probe names (that label columns and rows of adjacency) and gene names (that should label nodes in the output).

**Details**

The adjacency matrix is checked for validity. The entries can be negative, however. The adjacency matrix is expected to also have valid names or `dimnames[[2]]` that represent the probe names of the corresponding edges.

Whether the output is a weighted network or not, only edges whose (absolute value of) adjacency are above `threshold` will be included in the output.

If `probeToGene` is given, it is expected to have two columns, the first one corresponding to the probe names, the second to their corresponding gene names that will be used in the output.

**Value**

A data frame containing the network information suitable as input to VisANT. The same data frame is also written into a file specified by `file`, if given.

**Author(s)**

Peter Langfelder

**References**

VisANT software is available from <http://visant.bu.edu/>.

---

fixDataStructure      *Put single-set data into a form useful for multiset calculations.*

---

**Description**

Encapsulates single-set data in a wrapper that makes the data suitable for functions working on multiset data collections.

**Usage**

```
fixDataStructure(data, verbose = 0, indent = 0)
```

**Arguments**

data	A dataframe, matrix or array with two dimensions to be encapsulated.
verbose	Controls verbosity. 0 is silent.
indent	Controls indentation of printed progress messages. 0 means no indentation, every unit adds two spaces.

**Details**

For multiset calculations, many quantities (such as expression data, traits, module eigengenes etc) are presented by a common structure, a vector of lists (one list for each set) where each list has a component data that contains the actual (expression, trait, eigengene) data for the corresponding set in the form of a dataframe. This function creates a vector of lists of length 1 and fills the component data with the content of parameter data.

**Value**

As described above, input data in a format suitable for functions operating on multiset data collections.

**Author(s)**

Peter Langfelder, <[Peter.Langfelder@gmail.com](mailto:Peter.Langfelder@gmail.com)>

**See Also**

[checkSets](#)

**Examples**

```

singleSetData = matrix(rnorm(100), 10,10);
encapsData = fixDataStructure(singleSetData);
length(encapsData)
names(encapsData[[1]])
dim(encapsData[[1]]$data)
all.equal(encapsData[[1]]$data, singleSetData);

```

---

fundamentalNetworkConcepts

*Calculation of fundamental network concepts from an adjacency matrix.*

---

**Description**

This function computes fundamental network concepts (also known as network indices or statistics) based on an adjacency matrix and optionally a node significance measure. These network concepts are defined for any symmetric adjacency matrix (weighted and unweighted). The network concepts are described in Dong and Horvath (2007) and Horvath and Dong (2008). Fundamental network concepts are defined as a function of the off-diagonal elements of an adjacency matrix `adj` and/or a node significance measure `GS`.

**Usage**

```
fundamentalNetworkConcepts(adj, GS = NULL)
```

**Arguments**

<code>adj</code>	an adjacency matrix, that is a square, symmetric matrix with entries between 0 and 1
<code>GS</code>	a node significance measure: a vector of the same length as the number of rows (and columns) of the adjacency matrix.

**Value**

A list with the following components:

<code>Connectivity</code>	a numerical vector that reports the connectivity (also known as degree) of each node. This fundamental network concept is also known as whole network connectivity. One can also define the scaled connectivity $K = \text{Connectivity} / \max(\text{Connectivity})$ which is used for computing the hub gene significance.
<code>ScaledConnectivity</code>	the <code>Connectivity</code> vector scaled by the highest connectivity in the network, i.e., $\text{Connectivity} / \max(\text{Connectivity})$ .
<code>ClusterCoef</code>	a numerical vector that reports the cluster coefficient for each node. This fundamental network concept measures the cliquishness of each node.

MAR	a numerical vector that reports the maximum adjacency ratio for each node. MAR[i] equals 1 if all non-zero adjacencies between node i and the remaining network nodes equal 1. This fundamental network concept is always 1 for nodes of an unweighted network. This is a useful measure for weighted networks since it allows one to determine whether a node has high connectivity because of many weak connections (small MAR) or because of strong (but few) connections (high MAR), see Horvath and Dong 2008.
Density	the density of the network.
Centralization	the centralization of the network.
Heterogeneity	the heterogeneity of the network.

**Author(s)**

Steve Horvath

**References**

- Dong J, Horvath S (2007) Understanding Network Concepts in Modules, BMC Systems Biology 2007, 1:24
- Horvath S, Dong J (2008) Geometric Interpretation of Gene Coexpression Network Analysis. PLoS Comput Biol 4(8): e1000117

**See Also**

[conformityBasedNetworkConcepts](#) for calculation of conformity based network concepts for a network adjacency matrix;

[networkConcepts](#), for calculation of conformity based and eigennode based network concepts for a correlation network.

---

GOenrichmentAnalysis    *Calculation of GO enrichment (experimental)*

---

**Description**

**WARNING:** This function should be considered experimental. The arguments and resulting values (in particular, the enrichment p-values) are not yet finalized and may change in the future. The function should only be used to get a quick and rough overview of GO enrichment in the modules in a data set; for a publication-quality analysis, please use an established tool.

Using Bioconductor's annotation packages, this function calculates enrichments and returns terms with best enrichment values.

**Usage**

```
GOenrichmentAnalysis(labels,
                      entrezCodes,
                      yeastORFs = NULL,
                      organism = "human",
                      ontologies = c("BP", "CC", "MF"),
                      evidence = "all",
                      includeOffspring = TRUE,
```



```

backgroundType = "givenInGO",
removeDuplicates = TRUE,
leaveOutLabel = NULL,
nBestP = 10, pCut = NULL,
nBiggest = 0,
getTermDetails = TRUE,
verbose = 2, indent = 0)

```

## Arguments

labels	cluster (module, group) labels of genes to be analyzed. Either a single vector, or a matrix. In the matrix case, each column will be analyzed separately; analyzing a collection of module assignments in one function call will be faster than calling the function several times. For each row, the labels in all columns must correspond to the same gene specified in entrezCodes.
entrezCodes	Entrez (a.k.a. LocusLink) codes of the genes whose labels are given in labels. A single vector; the i-th entry corresponds to row i of the matrix labels (or to the i-th entry if labels is a vector).
yeastORFs	if organism=="yeast" (below), this argument can be used to input yeast open reading frame (ORF) identifiers instead of Entrez codes. Since the GO mappings for yeast are provided in terms of ORF identifiers, this may lead to a more accurate GO enrichment analysis. If given, the argument entrezCodes is ignored.
organism	character string specifying the organism for which to perform the analysis. Recognized values are (unique abbreviations of) "human", "mouse", "rat", "malaria", "yeast",
ontologies	vector of character strings specifying GO ontologies to be included in the analysis. Can be any subset of "BP", "CC", "MF". The result will contain the terms with highest enrichment in each specified category, plus a separate list of terms with best enrichment in all ontologies combined.
evidence	vector of character strings specifying admissible evidence for each gene in its specific term, or "all" for all evidence codes. See Details or <a href="http://www.geneontology.org/GO.evidence">http://www.geneontology.org/GO.evidence</a> for available evidence codes and their meaning.
includeOffspring	logical: should genes belonging to the offspring of each term be included in the term? As a default, only genes belonging directly to each term are associated with the term. Note that the calculation of enrichments with offspring included can be quite slow for large data sets.
backgroundType	specification of the background to use. Recognized values are (unique abbreviations of) "allGiven", "allInGO", "givenInGO", meaning that the functions will take all genes given in labels as background ("allGiven"), all genes present in any of the GO categories ("allInGO"), or the intersection of given genes and genes present in GO ("givenInGO"). The default is recommended for genome-wide enrichment studies.
removeDuplicates	logical: should duplicate entries in entrezCodes be removed? If TRUE, only the first occurrence of each unique Entrez code will be kept. The cluster labels will be adjusted accordingly.
leaveOutLabel	optional specifications of module labels for which enrichment calculation is not desired. Can be a single label or a vector of labels to be ignored. However, if in any of the sets no labels are left to calculate enrichment of, the function will stop with an error.

nBestP	specifies the number of terms with highest enrichment whose detailed information will be returned.
pCut	alternative specification of terms to be returned: all terms whose enrichment p-value is more significant than pCut will be returned. If pCut is given, nBestP is ignored.
nBiggest	in addition to returning terms with highest enrichment, terms that contain most of the genes in each cluster can be returned by specifying the number of biggest terms per cluster to be returned. This may be useful for development and testing purposes.
getTermDetails	logical indicating whether detailed information on the most enriched terms should be returned.
verbose	integer specifying the verbosity of the function. Zero means silent, positive values will cause the function to print progress reports.
indent	integer specifying indentation of the diagnostic messages. Zero means no indentation, each unit adds two spaces.

## Details

This function is basically a wrapper for the annotation packages available from Bioconductor. It requires the packages GO.db, AnnotationDbi, and org.xx.eg.db, where xx is the code corresponding to the organism that the user wishes to analyze (e.g., Hs for human Homo Sapiens, Mm for mouse Mus Musculus etc). For each cluster specified in the input, the function calculates all enrichments in the specified ontologies, and collects information about the terms with highest enrichment. The enrichment p-value is calculated using Fisher exact test. As background we use all of the supplied genes that are present in at least one term in GO (in any of the ontologies).

For best results, the newest annotation libraries should be used. Because of the way Bioconductor is set up, to get the newest annotation libraries you may have to use the current version of R.

According to <http://www.geneontology.org/GO.evidence.shtml>, the following codes are used by GO:

### Experimental Evidence Codes

EXP: Inferred from Experiment  
 IDA: Inferred from Direct Assay  
 IPI: Inferred from Physical Interaction  
 IMP: Inferred from Mutant Phenotype  
 IGI: Inferred from Genetic Interaction  
 IEP: Inferred from Expression Pattern

### Computational Analysis Evidence Codes

ISS: Inferred from Sequence or Structural Similarity  
 ISO: Inferred from Sequence Orthology  
 ISA: Inferred from Sequence Alignment  
 ISM: Inferred from Sequence Model  
 IGC: Inferred from Genomic Context  
 IBA: Inferred from Biological aspect of Ancestor  
 IBD: Inferred from Biological aspect of Descendant  
 IKR: Inferred from Key Residues  
 IRD: Inferred from Rapid Divergence  
 RCA: inferred from Reviewed Computational Analysis

### Author Statement Evidence Codes

TAS: Traceable Author Statement  
 NAS: Non-traceable Author Statement

Curator Statement Evidence Codes  
 IC: Inferred by Curator  
 ND: No biological Data available

Automatically-assigned Evidence Codes  
 IEA: Inferred from Electronic Annotation

Obsolete Evidence Codes  
 NR: Not Recorded

## Value

A list with the following components:

keptForAnalysis

logical vector with one entry per given gene. TRUE if the entry was used for enrichment analysis. Depending on the setting of `removeDuplications` above, only a single entry per gene may be used.

inGO

logical vector with one entry per given gene. TRUE if the gene belongs to any GO term, FALSE otherwise. Also FALSE for genes not used for the analysis because of duplication.

If input `labels` contained only one vector of labels, the following components:

countsInTerms

a matrix whose rows correspond to given cluster, and whose columns correspond to GO terms, containing number of genes in the intersection of the corresponding module and GO term. Row and column names are set appropriately.

enrichmentP

a matrix whose rows correspond to given cluster, and whose columns correspond to GO terms, containing enrichment p-values of each term in each cluster. Row and column names are set appropriately.

bestPTerms

a list of lists with each inner list corresponding to an ontology given in `ontologies` in input, plus one component corresponding to all given ontologies combined. The name of each component is set appropriately. Each inner list contains two components: `enrichment` is a data frame containing the highest enriched terms for each module; and `forModule` is a list of lists with one inner list per module, appropriately named. Each inner list contains one component per term. If input `getTermDetails` is TRUE, this component is yet another list and contains components `termName` (term name), `enrichmentP` (enrichment P value), `termDefinition` (GO term definition), `termOntology` (GO term ontology), `geneCodes` (Entrez codes of module genes in this term), `genePositions` (indices of the genes listed in `geneCodes` within the given labels). Thus, to obtain information on say the second term of the 5th module in ontology BP, one can look at the appropriate row of `bestPTerms$BP$enrichment`, or one can reference `bestPTerms$BP$forModule[[5]][[2]]`. The author of the function apologizes for any confusion this structure of the output may cause.

biggestTerms

a list of the same format as `bestPTerms`, containing information about the terms with most genes in the module for each supplied ontology.

If input `labels` contained more than one vector, instead of the above components the return value contains a list named `setResults` that has one component per given set; each component is a list containing the above components for the corresponding set.

**Author(s)**

Peter Langfelder

**See Also**

Bioconductor's annotation packages such as GO.db and organism-specific annotation packages such as org.Hs.eg.db.

---

goodGenes

*Filter genes with too many missing entries*

---

**Description**

This function checks data for missing entries and returns a list of genes that have non-zero variance and pass two criteria on maximum number of missing values: the fraction of missing values must be below a given threshold and the total number of missing samples must be below a given threshold.

**Usage**

```
goodGenes(datExpr,
          useSamples = NULL,
          useGenes = NULL,
          minFraction = 1/2,
          minNSamples = ..minNSamples,
          minNGenes = ..minNGenes,
          verbose = 1, indent = 0)
```

**Arguments**

datExpr	expression data. A data frame in which columns are genes and rows are samples.
useSamples	optional specifications of which samples to use for the check. Should be a logical vector; samples whose entries are FALSE will be ignored for the missing value counts. Defaults to using all samples.
useGenes	optional specifications of genes for which to perform the check. Should be a logical vector; genes whose entries are FALSE will be ignored. Defaults to using all genes.
minFraction	minimum fraction of non-missing samples for a gene to be considered good.
minNSamples	minimum number of non-missing samples for a gene to be considered good.
minNGenes	minimum number of good genes for the data set to be considered fit for analysis. If the actual number of good genes falls below this threshold, an error will be issued.
verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.
indent	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

**Details**

The constants `..minNSamples` and `..minNGenes` are both set to the value 4. For most data sets, the fraction of missing samples criterion will be much more stringent than the absolute number of missing samples criterion.

**Value**

A logical vector with one entry per gene that is TRUE if the gene is considered good and FALSE otherwise. Note that all genes excluded by `useGenes` are automatically assigned FALSE.

**Author(s)**

Peter Langfelder and Steve Horvath

**See Also**

[goodSamples](#), [goodSamplesGenes](#)

---

goodGenesMS

*Filter genes with too many missing entries across multiple sets*

---

**Description**

This function checks data for missing entries and returns a list of genes that have non-zero variance in all sets and pass two criteria on maximum number of missing values in each given set: the fraction of missing values must be below a given threshold and the total number of missing samples must be below a given threshold

**Usage**

```
goodGenesMS(multiExpr,
            useSamples = NULL,
            useGenes = NULL,
            minFraction = 1/2,
            minNSamples = ..minNSamples,
            minNGenes = ..minNGenes,
            verbose = 1, indent = 0)
```

**Arguments**

<code>multiExpr</code>	expression data in the multi-set format (see <a href="#">checkSets</a> ). A vector of lists, one per set. Each set must contain a component data that contains the expression data, with rows corresponding to samples and columns to genes or probes.
<code>useSamples</code>	optional specifications of which samples to use for the check. Should be a logical vector; samples whose entries are FALSE will be ignored for the missing value counts. Defaults to using all samples.
<code>useGenes</code>	optional specifications of genes for which to perform the check. Should be a logical vector; genes whose entries are FALSE will be ignored. Defaults to using all genes.

minFraction	minimum fraction of non-missing samples for a gene to be considered good.
minNSamples	minimum number of non-missing samples for a gene to be considered good.
minNGenes	minimum number of good genes for the data set to be considered fit for analysis. If the actual number of good genes falls below this threshold, an error will be issued.
verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.
indent	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

### Details

The constants `..minNSamples` and `..minNGenes` are both set to the value 4. For most data sets, the fraction of missing samples criterion will be much more stringent than the absolute number of missing samples criterion.

### Value

A logical vector with one entry per gene that is TRUE if the gene is considered good and FALSE otherwise. Note that all genes excluded by `useGenes` are automatically assigned FALSE.

### Author(s)

Peter Langfelder

### See Also

[goodGenes](#), [goodSamples](#), [goodSamplesGenes](#) for cleaning individual sets separately;  
[goodSamplesMS](#), [goodSamplesGenesMS](#) for additional cleaning of multiple data sets together.

---

goodSamples	<i>Filter samples with too many missing entries</i>
-------------	---

---

### Description

This function checks data for missing entries and returns a list of samples that pass two criteria on maximum number of missing values: the fraction of missing values must be below a given threshold and the total number of missing genes must be below a given threshold.

### Usage

```
goodSamples(datExpr,
            useSamples = NULL,
            useGenes = NULL,
            minFraction = 1/2,
            minNSamples = ..minNSamples,
            minNGenes = ..minNGenes,
            verbose = 1, indent = 0)
```

**Arguments**

datExpr	expression data. A data frame in which columns are genes and rows are samples.
useSamples	optional specifications of which samples to use for the check. Should be a logical vector; samples whose entries are FALSE will be ignored for the missing value counts. Defaults to using all samples.
useGenes	optional specifications of genes for which to perform the check. Should be a logical vector; genes whose entries are FALSE will be ignored. Defaults to using all genes.
minFraction	minimum fraction of non-missing samples for a gene to be considered good.
minNSamples	minimum number of good samples for the data set to be considered fit for analysis. If the actual number of good samples falls below this threshold, an error will be issued.
minNGenes	minimum number of non-missing samples for a sample to be considered good.
verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.
indent	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

**Details**

The constants `..minNSamples` and `..minNGenes` are both set to the value 4. For most data sets, the fraction of missing samples criterion will be much more stringent than the absolute number of missing samples criterion.

**Value**

A logical vector with one entry per sample that is TRUE if the sample is considered good and FALSE otherwise. Note that all samples excluded by `useSamples` are automatically assigned FALSE.

**Author(s)**

Peter Langfelder and Steve Horvath

**See Also**

[goodSamples](#), [goodSamplesGenes](#)

---

goodSamplesGenes

*Iterative filtering of samples and genes with too many missing entries*

---

**Description**

This function checks data for missing entries and zero-variance genes, and returns a list of samples and genes that pass criteria maximum number of missing values. If necessary, the filtering is iterated.

**Usage**

```
goodSamplesGenes(  
  datExpr,  
  minFraction = 1/2,  
  minNSamples = ..minNSamples,  
  minNGenes = ..minNGenes,  
  verbose = 1, indent = 0)
```

**Arguments**

<code>datExpr</code>	expression data. A data frame in which columns are genes and rows are samples.
<code>minFraction</code>	minimum fraction of non-missing samples for a gene to be considered good.
<code>minNSamples</code>	minimum number of non-missing samples for a gene to be considered good.
<code>minNGenes</code>	minimum number of good genes for the data set to be considered fit for analysis. If the actual number of good genes falls below this threshold, an error will be issued.
<code>verbose</code>	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.
<code>indent</code>	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

**Details**

This function iteratively identifies samples and genes with too many missing entries and genes with zero variance. Iterations may be required since excluding samples effectively changes criteria on genes and vice versa. The process is repeated until the lists of good samples and genes are stable. The constants `..minNSamples` and `..minNGenes` are both set to the value 4.

**Value**

A list with the following components:

<code>goodSamples</code>	A logical vector with one entry per sample that is TRUE if the sample is considered good and FALSE otherwise.
<code>goodGenes</code>	A logical vector with one entry per gene that is TRUE if the gene is considered good and FALSE otherwise.

**Author(s)**

Peter Langfelder

**See Also**

[goodSamples](#), [goodGenes](#)



---

goodSamplesGenesMS	<i>Iterative filtering of samples and genes with too many missing entries across multiple data sets</i>
--------------------	---

---

### Description

This function checks data for missing entries and zero variance across multiple data sets and returns a list of samples and genes that pass criteria maximum number of missing values. If necessary, the filtering is iterated.

### Usage

```
goodSamplesGenesMS(
  multiExpr,
  minFraction = 1/2,
  minNSamples = ..minNSamples,
  minNGenes = ..minNGenes,
  verbose = 2, indent = 0)
```

### Arguments

multiExpr	expression data in the multi-set format (see <a href="#">checkSets</a> ). A vector of lists, one per set. Each set must contain a component data that contains the expression data, with rows corresponding to samples and columns to genes or probes.
minFraction	minimum fraction of non-missing samples for a gene to be considered good.
minNSamples	minimum number of non-missing samples for a gene to be considered good.
minNGenes	minimum number of good genes for the data set to be considered fit for analysis. If the actual number of good genes falls below this threshold, an error will be issued.
verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.
indent	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

### Details

This function iteratively identifies samples and genes with too many missing entries, and genes with zero variance. Iterations may be required since excluding samples effectively changes criteria on genes and vice versa. The process is repeated until the lists of good samples and genes are stable. The constants `..minNSamples` and `..minNGenes` are both set to the value 4.

### Value

A list with the following components:

goodSamples	A list with one component per given set. Each component is a logical vector with one entry per sample in the corresponding set that is TRUE if the sample is considered good and FALSE otherwise.
goodGenes	A logical vector with one entry per gene that is TRUE if the gene is considered good and FALSE otherwise.

**Author(s)**

Peter Langfelder

**See Also**

[goodGenes](#), [goodSamples](#), [goodSamplesGenes](#) for cleaning individual sets separately;  
[goodSamplesMS](#), [goodGenesMS](#) for additional cleaning of multiple data sets together.

goodSamplesMS

*Filter samples with too many missing entries across multiple data sets***Description**

This function checks data for missing entries and returns a list of samples that pass two criteria on maximum number of missing values: the fraction of missing values must be below a given threshold and the total number of missing genes must be below a given threshold.

**Usage**

```
goodSamplesMS(multiExpr,
              useSamples = NULL,
              useGenes = NULL,
              minFraction = 1/2,
              minNSamples = ..minNSamples,
              minNGenes = ..minNGenes,
              verbose = 1, indent = 0)
```

**Arguments**

multiExpr	expression data in the multi-set format (see <a href="#">checkSets</a> ). A vector of lists, one per set. Each set must contain a component data that contains the expression data, with rows corresponding to samples and columns to genes or probes.
useSamples	optional specifications of which samples to use for the check. Should be a logical vector; samples whose entries are FALSE will be ignored for the missing value counts. Defaults to using all samples.
useGenes	optional specifications of genes for which to perform the check. Should be a logical vector; genes whose entries are FALSE will be ignored. Defaults to using all genes.
minFraction	minimum fraction of non-missing samples for a gene to be considered good.
minNSamples	minimum number of good samples for the data set to be considered fit for analysis. If the actual number of good samples falls below this threshold, an error will be issued.
minNGenes	minimum number of non-missing samples for a sample to be considered good.
verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.
indent	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

**Details**

The constants `..minNSamples` and `..minNGenes` are both set to the value 4. For most data sets, the fraction of missing samples criterion will be much more stringent than the absolute number of missing samples criterion.

**Value**

A list with one component per input set. Each component is a logical vector with one entry per sample in the corresponding set, indicating whether the sample passed the missing value criteria.

**Author(s)**

Peter Langfelder and Steve Horvath

**See Also**

[goodGenes](#), [goodSamples](#), [goodSamplesGenes](#) for cleaning individual sets separately;  
[goodGenesMS](#), [goodSamplesGenesMS](#) for additional cleaning of multiple data sets together.

---

`greenBlackRed`*Green-black-red color sequence*

---

**Description**

Generate a green-black-red color sequence of a given length.

**Usage**

```
greenBlackRed(n, gamma = 1)
```

**Arguments**

<code>n</code>	number of colors to be returned
<code>gamma</code>	color correction power

**Details**

The function returns a color vector that starts with pure green, gradually turns into black and then to red. The power `gamma` can be used to control the behaviour of the quarter- and three quarter-values (between green and black, and black and red, respectively). Higher powers will make the mid-colors more green and red, respectively.

**Value**

A vector of colors of length `n`.

**Author(s)**

Peter Langfelder

## Examples

```
par(mfrow = c(3, 1))
displayColors(greenBlackRed(50));
displayColors(greenBlackRed(50, 2));
displayColors(greenBlackRed(50, 0.5));
```

---

greenWhiteRed

*Green-white-red color sequence*

---

## Description

Generate a green-white-red color sequence of a given length.

## Usage

```
greenWhiteRed(n, gamma = 1, warn = TRUE)
```

## Arguments

n	number of colors to be returned
gamma	color change power
warn	logical: should the user be warned that this function produces a palette unsuitable for people with most common color blindness?

## Details

The function returns a color vector that starts with green, gradually turns into white and then to red. The power gamma can be used to control the behaviour of the quarter- and three quarter-values (between green and white, and white and red, respectively). Higher powers will make the mid-colors more white, while lower powers will make the colors more saturated, respectively.

Typical use of this function is to produce (via function [numbers2colors](#)) a color representation of numbers within a symmetric interval around 0, for example, the interval [-1, 1]. Note though that since green and red are not distinguishable by people with the most common type of color blindness, we recommend using the analogous palette returned by the function [blueWhiteRed](#).

## Value

A vector of colors of length n.

## Author(s)

Peter Langfelder

## See Also

[blueWhiteRed](#) for a color sequence more friendly to people with the most common type of color blindness;

[numbers2colors](#) for a function that produces a color representation for continuous numbers.

**Examples**

```
par(mfrow = c(3, 1))
displayColors(greenWhiteRed(50));
title("gamma = 1")
displayColors(greenWhiteRed(50, 3));
title("gamma = 3")
displayColors(greenWhiteRed(50, 0.5));
title("gamma = 0.5")
```

---

**GTOMdist***Generalized Topological Overlap Measure*

---

**Description**

Generalized Topological Overlap Measure, taking into account interactions of higher degree.

**Usage**

```
GTOMdist(adjMat, degree = 1)
```

**Arguments**

adjMat            adjacency matrix. See details below.  
degree            integer specifying the maximum degree to be calculated.

**Value**

Matrix of the same dimension as the input adjMat.

**Author(s)**

Steve Horvath and Andy Yip

**References**

Yip A, Horvath S (2007) Gene network interconnectedness and the generalized topological overlap measure. *BMC Bioinformatics* 2007, 8:22

---

**hubGeneSignificance***Hubgene significance*

---

**Description**

Calculate approximate hub gene significance for all modules in network.

**Usage**

```
hubGeneSignificance(datKME, GS)
```

**Arguments**

- datKME a data frame (or a matrix-like object) containing eigengene-based connectivities of all genes in the network.
- GS a vector with one entry for every gene containing its gene significance.

**Details**

In datKME rows correspond to genes and columns to modules.

**Value**

A vector whose entries are the hub gene significances for each module.

**Author(s)**

Steve Horvath

**References**

Dong J, Horvath S (2007) Understanding Network Concepts in Modules, BMC Systems Biology 2007, 1:24

---

ImmunePathwayLists *Immune Pathways with Corresponding Gene Markers*

---

**Description**

This matrix gives a predefined set of marker genes for many immune response pathways, as assembled by Brian Modena (a member of Daniel R Salomon's lab at Scripps Research Institute), and colleagues. It is used with `userListEnrichment` to search user-defined gene lists for enrichment.

**Usage**

```
data(ImmunePathwayLists)
```

**Format**

A 3597 x 2 matrix of characters containing Gene / Category pairs. The first column (Gene) lists genes corresponding to a given category (second column). Each Category entry is of the form <Immune Pathway>\_\_ImmunePathway. Note that the matrix is sorted first by Category and then by Gene, such that all genes related to the same category are listed sequentially.

**Source**

For more information about this list, please see [userListEnrichment](#)

**Examples**

```
data(ImmunePathwayLists)
head(ImmunePathwayLists)
```

---

Inline display of progress

*Inline display of progress*

---

## Description

These functions provide an inline display of progress.

## Usage

```
initProgInd(leadStr = "..", trailStr = "", quiet = !interactive())  
updateProgInd(newFrac, progInd, quiet = !interactive())
```

## Arguments

leadStr	character string that will be printed before the actual progress number.
trailStr	character string that will be printed after the actual progress number.
quiet	can be used to silence the indicator for non-interactive sessions whose output is typically redirected to a file.
newFrac	new fraction of progress to be displayed.
progInd	an object of class <code>progressIndicator</code> that encodes previously printed message.

## Details

A progress indicator is a simple inline display of progress intended to satisfy impatient users during lengthy operations. The function `initProgInd` initializes a progress indicator (at zero); `updateProgInd` updates it to a specified fraction.

Note that excessive use of `updateProgInd` may lead to a performance penalty (see examples).

## Value

Both functions return an object of class `progressIndicator` that holds information on the last printed value and should be used for subsequent updates of the indicator.

## Author(s)

Peter Langfelder

## Examples

```
max = 10;  
prog = initProgInd("Counting: ", "done");  
for (c in 1:max)  
{  
  Sys.sleep(0.10);  
  prog = updateProgInd(c/max, prog);  
}  
printFlush("");  
  
printFlush("Example 2:");  
prog = initProgInd();
```

```

for (c in 1:max)
{
  Sys.sleep(0.10);
  prog = updateProgInd(c/max, prog);
}
printFlush("");

## Example of a significant slowdown:

## Without progress indicator:

system.time( {a = 0; for (i in 1:10000) a = a+i; } )

## With progress indicator, some 50 times slower:

system.time(
{
  prog = initProgInd("Counting: ", "done");
  a = 0;
  for (i in 1:10000)
  {
    a = a+i;
    prog = updateProgInd(i/10000, prog);
  }
}
)

```

---

intramodularConnectivity

*Calculation of intramodular connectivity*


---

### Description

Calculates intramodular connectivity, i.e., connectivity of nodes to other nodes within the same module.

### Usage

```
intramodularConnectivity(adjMat, colors, scaleByMax = FALSE)
```

### Arguments

adjMat	adjacency matrix, a square, symmetric matrix with entries between 0 and 1.
colors	module labels. A vector of length ncol(adjMat) giving a module label for each gene (node) of the network.
scaleByMax	logical: should intramodular connectivities be scaled by the maximum IM connectivity in each module?

### Details

The module labels can be numeric or character. For each node (gene), the function sums adjacency entries (excluding the diagonal) to other nodes within the same module. Optionally, the connectivities can be scaled by the maximum connectivity in each module.



**Value**

A data frame with 4 columns giving the total connectivity, intramodular connectivity, extra-modular connectivity, and the difference of the intra- and extra-modular connectivities for all genes.

**Author(s)**

Steve Horvath and Peter Langfelder

**References**

Dong J, Horvath S (2007) Understanding Network Concepts in Modules, BMC Systems Biology 2007, 1:24

**See Also**

[adjacency](#)

---

keepCommonProbes	<i>Keep probes that are shared among given data sets</i>
------------------	--

---

**Description**

This function strips out probes that are not shared by all given data sets, and orders the remaining common probes using the same order in all sets.

**Usage**

```
keepCommonProbes(multiExpr, orderBy = 1)
```

**Arguments**

multiExpr	expression data in the multi-set format (see <a href="#">checkSets</a> ). A vector of lists, one per set. Each set must contain a component data that contains the expression data, with rows corresponding to samples and columns to genes or probes.
orderBy	index of the set by which probes are to be ordered.

**Value**

Expression data in the same format as the input data, containing only common probes.

**Author(s)**

Peter Langfelder

**See Also**

[checkSets](#)

---

kMEcomparisonScatterplot

*Function to plot kME values between two comparable data sets.*


---

### Description

Plots the kME values of genes in two groups of expression data for each module in an inputted color vector.

### Usage

```
kMEcomparisonScatterplot(
  datExpr1, datExpr2, colorh,
  inA = NULL, inB = NULL, MEsA = NULL, MEsB = NULL,
  nameA = "A", nameB = "B",
  plotAll = FALSE, noGrey = TRUE, maxPlot = 1000, pch = 19,
  fileName = if (plotAll) paste("kME_correlations_between_", nameA, "_and_",
                                nameB, "_all.pdf", sep="") else
                                paste("kME_correlations_between_", nameA, "_and_",
                                      nameB, "_inMod.pdf", sep=""), ...)
```

### Arguments

datExpr1	The first expression matrix (samples=rows, genes=columns). This can either include only the data for group A (in which case dataExpr2 must be entered), or can contain all of the data for groups A and B (in which case inA and inB must be entered).
datExpr2	The second expression matrix, or set to NULL if all data is from same expression matrix. If entered, datExpr2 must contain the same genes as datExpr1 in the same order.
colorh	The common color vector (module labels) corresponding to both sets of expression data.
inA, inB	Vectors of TRUE/FALSE indicating whether a sample is in group A/B, or a vector of numeric indices indicating which samples are in group A/B. If datExpr2 is entered, these inputs are ignored (thus default = NULL). For these and all other A/B inputs, "A" corresponds to datExpr1 and "B" corresponds to datExpr2 if datExpr2 is entered; otherwise "A" corresponds to datExpr1[inA,] while "B" corresponds to datExpr1[inB,].
MEsA, MEsB	Either the module eigengenes or NULL (default) in which case the module eigengenes will be calculated. In inputted, MEs MUST be calculated using "moduleEigengenes(<parameters>)\$eigengenes" for function to work properly.
nameA, nameB	The names of these groups (defaults = "A" and "B"). The resulting file name (see below) and x and y axis labels for each scatter plot depend on these names.
plotAll	If TRUE, plot gene-ME correlations for all genes. If FALSE, plot correlations for only genes in the plotted module (default). Note that the output file name will be different depending on this parameter, so both can be run without overwriting results.
noGrey	If TRUE (default), the grey module genes are ignored. This parameter is only used if MEsA and MEsB are calculated.

maxPlot	The maximum number of random genes to include (default=1000). Smaller values lead to smaller and less cluttered plots, usually without significantly affecting the resulting correlations. This parameter is only used if plotAll=TRUE.
pch	See help file for "points". Setting pch=19 (default) produces solid circles.
fileName	Name of the file to hold the plots. Since the output format is pdf, the extension should be .pdf .
...	Other plotting parameters that are allowable inputs to verboseScatterplot.

**Value**

The default output is a file called "kME\_correlations\_between\_[nameA]\_and\_[nameB]\_[all/inMod].pdf", where [nameA] and [nameB] correspond to the nameA and nameB input parameters, and [all/inMod] depends on whether plotAll=TRUE or FALSE. This output file contains all of the plots as separate pdf images, and will be located in the current working directory.

**Note**

The function "pdf", which can be found in the grDevices library, is required to run this function.

**Author(s)**

Jeremy Miller

**Examples**

```
# Example output file ("kME_correlations_between_A_and_B_inMod.pdf") using simulated data.

set.seed = 100
ME=matrix(0,50,5)
for (i in 1:5) ME[,i]=sample(1:100,50)
simData1 = simulateDatExpr5Modules(MEturquoise=ME[,1],MEblue=ME[,2],
                                   MEbrown=ME[,3],MEyellow=ME[,4], MEgreen=ME[,5])
simData2 = simulateDatExpr5Modules(MEturquoise=ME[,1],MEblue=ME[,2],
                                   MEbrown=ME[,3],MEyellow=ME[,4], MEgreen=ME[,5])
kMEcomparisonScatterplot(simData1$datExpr, simData2$datExpr, simData1$trueModule)
```

---

labeledBarplot

*Barplot with text or color labels.*


---

**Description**

Produce a barplot with extra annotation.

**Usage**

```
labeledBarplot(
  Matrix, labels,
  colorLabels = FALSE,
  colored = TRUE,
  setStdMargins = TRUE,
  stdErrors = NULL,
```

```

    cex.lab = NULL,
    xLabelsAngle = 45,
    ...)

```

### Arguments

<code>Matrix</code>	vector or a matrix to be plotted.
<code>labels</code>	labels to annotate the bars underneath the barplot.
<code>colorLabels</code>	logical: should the labels be interpreted as colors? If TRUE, the bars will be labeled by colored squares instead of text. See details.
<code>colored</code>	logical: should the bars be divided into segments and colored? If TRUE, assumes the labels can be interpreted as colors, and the input <code>Matrix</code> is square and the rows have the same labels as the columns. See details.
<code>setStdMargins</code>	if TRUE, the function will set margins <code>c(3, 3, 2, 2)+0.2</code> .
<code>stdErrors</code>	if given, error bars corresponding to $1.96 \times \text{stdErrors}$ will be plotted on top of the bars.
<code>cex.lab</code>	character expansion factor for axis labels, including the text labels underneath the barplot.
<code>xLabelsAngle</code>	angle at which text labels under the barplot will be printed.
<code>...</code>	other parameters for the function <code>barplot</code> .

### Details

Individual bars in the barplot can be identified either by printing the text of the corresponding entry in `labels` underneath the bar at the angle specified by `xLabelsAngle`, or by interpreting the `labels` entry as a color (see below) and drawing a correspondingly colored square underneath the bar.

For reasons of compatibility with other functions, `labels` are interpreted as colors after stripping the first two characters from each label. For example, the label "MEturquoise" is interpreted as the color turquoise.

If `colored` is set, the code assumes that `labels` can be interpreted as colors, and the input `Matrix` is square and the rows have the same labels as the columns. Each bar in the barplot is then sectioned into contributions from each row entry in `Matrix` and is colored by the color given by the entry in `labels` that corresponds to the row.

### Value

None.

### Author(s)

Peter Langfelder

---

labeledHeatmap	<i>Produce a labeled heatmap plot</i>
----------------	---------------------------------------

---

### Description

Plots a heatmap plot with color legend, row and column annotation, and optional text within the heatmap.

### Usage

```
labeledHeatmap(
  Matrix,
  xLabels, yLabels = NULL,
  xSymbols = NULL, ySymbols = NULL,
  colorLabels = NULL,
  xColorLabels = FALSE, yColorLabels = FALSE,
  checkColorsValid = TRUE,
  invertColors = FALSE,
  setStdMargins = TRUE,
  xLabelsPosition = "bottom",
  xLabelsAngle = 45,
  xLabelsAdj = 1,
  xColorWidth = 0.05,
  yColorWidth = 0.05,
  colors = NULL,
  naColor = "grey",
  textMatrix = NULL,
  cex.text = NULL, cex.lab = NULL,
  cex.lab.x = cex.lab,
  cex.lab.y = cex.lab,
  colors.lab.x = 1,
  colors.lab.y = 1,
  plotLegend = TRUE, ...)
```

### Arguments

Matrix	numerical matrix to be plotted in the heatmap.
xLabels	labels for the columns. See Details.
yLabels	labels for the rows. See Details.
xSymbols	additional labels used when xLabels are interpreted as colors. See Details.
ySymbols	additional labels used when yLabels are interpreted as colors. See Details.
colorLabels	logical: should xLabels and yLabels be interpreted as colors? If given, overrides xColorLabels and yColorLabels below.
xColorLabels	logical: should xLabels be interpreted as colors?
yColorLabels	logical: should yLabels be interpreted as colors?
checkColorsValid	logical: should given colors be checked for validity against the output of colors()? If this argument is FALSE, invalid color specification will trigger an error.

<code>invertColors</code>	logical: should the color order be inverted?
<code>setStdMargins</code>	logical: should standard margins be set before calling the plot function? Standard margins depend on <code>colorLabels</code> : they are wider for text labels and narrower for color labels. The defaults are static, that is the function does not attempt to guess the optimal margins.
<code>xLabelsPosition</code>	a character string specifying the position of labels for the columns. Recognized values are (unique abbreviations of) "top", "bottom".
<code>xLabelsAngle</code>	angle by which the column labels should be rotated.
<code>xLabelsAdj</code>	justification parameter for column labels. See <code>par</code> and the description of parameter "adj".
<code>xColorWidth</code>	width of the color labels for the x axis expressed as a fraction of the smaller of the range of the x and y axis.
<code>yColorWidth</code>	width of the color labels for the y axis expressed as a fraction of the smaller of the range of the x and y axis.
<code>colors</code>	color palette to be used in the heatmap. Defaults to <code>heat.colors</code> .
<code>naColor</code>	color to be used for encoding missing data.
<code>textMatrix</code>	optional matrix of text entries of the same dimensions as <code>Matrix</code> .
<code>cex.text</code>	character expansion factor for <code>textMatrix</code> .
<code>cex.lab</code>	character expansion factor for text labels labeling the axes.
<code>cex.lab.x</code>	character expansion factor for text labels labeling the x axis. Overrides <code>cex.lab</code> above.
<code>cex.lab.y</code>	character expansion factor for text labels labeling the y axis. Overrides <code>cex.lab</code> above.
<code>colors.lab.x</code>	colors for character labels or symbols along x axis.
<code>colors.lab.y</code>	colors for character labels or symbols along y axis.
<code>plotLegend</code>	logical: should a color legend be plotted?
<code>...</code>	other arguments to function <code>heatmap</code> .

## Details

The function basically plots a standard heatmap plot of the given `Matrix` and embellishes it with row and column labels and/or with text within the heatmap entries. Row and column labels can be either character strings or color squares, or both.

To get simple text labels, use `colorLabels=FALSE` and pass the desired row and column labels in `yLabels` and `xLabels`, respectively.

To label rows and columns by color squares, use `colorLabels=TRUE`; `yLabels` and `xLabels` are then expected to represent valid colors. For reasons of compatibility with other functions, each entry in `yLabels` and `xLabels` is expected to consist of a color designation preceded by 2 characters: an example would be `MEturquoise`. The first two characters can be arbitrary, they are stripped. Any labels that do not represent valid colors will be considered text labels and printed in full, allowing the user to mix text and color labels.

It is also possible to label rows and columns by both color squares and additional text annotation. To achieve this, use the above technique to get color labels and, additionally, pass the desired text annotation in the `xSymbols` and `ySymbols` arguments.

**Value**

None.

**Author(s)**

Peter Langfelder

**See Also**

[heatmap](#), [colors](#)

**Examples**

```
# This example illustrates 4 main ways of annotating columns and rows of a heatmap.
# Copy and paste the whole example into an R session with an interactive plot window;
# alternatively, you may replace the command sizeGrWindow below by opening
# another graphical device such as pdf.
```

```
# Generate a matrix to be plotted
```

```
nCol = 8; nRow = 7;
mat = matrix(runif(nCol*nRow, min = -1, max = 1), nRow, nCol);
```

```
rowColors = standardColors(nRow);
colColors = standardColors(nRow + nCol)[(nRow+1):(nRow + nCol)];
```

```
rowColors;
colColors;
```

```
sizeGrWindow(9,7)
par(mfrow = c(2,2))
par(mar = c(4, 5, 4, 6));
```

```
# Label rows and columns by text:
```

```
labeledHeatmap(mat, xLabels = colColors, yLabels = rowColors,
               colors = greenWhiteRed(50),
               setStdMargins = FALSE,
               textMatrix = signif(mat, 2),
               main = "Text-labeled heatmap");
```

```
# Label rows and columns by colors:
```

```
rowLabels = paste("ME", rowColors, sep="");
colLabels = paste("ME", colColors, sep="");
```

```
labeledHeatmap(mat, xLabels = colLabels, yLabels = rowLabels,
               colorLabels = TRUE,
               colors = greenWhiteRed(50),
               setStdMargins = FALSE,
               textMatrix = signif(mat, 2),
               main = "Color-labeled heatmap");
```

```
# Mix text and color labels:
```

```

rowLabels[3] = "Row 3";
colLabels[1] = "Column 1";

labeledHeatmap(mat, xLabels = colLabels, yLabels = rowLabels,
               colorLabels = TRUE,
               colors = greenWhiteRed(50),
               setStdMargins = FALSE,
               textMatrix = signif(mat, 2),
               main = "Mix-labeled heatmap");

# Color labels and additional text labels

rowLabels = paste("ME", rowColors, sep="");
colLabels = paste("ME", colColors, sep="");

extraRowLabels = paste("Row", c(1:nRow));
extraColLabels = paste("Column", c(1:nCol));

# Extend margins to fit all labels
par(mar = c(6, 6, 4, 6));
labeledHeatmap(mat, xLabels = colLabels, yLabels = rowLabels,
               xSymbols = extraColLabels,
               ySymbols = extraRowLabels,
               colorLabels = TRUE,
               colors = greenWhiteRed(50),
               setStdMargins = FALSE,
               textMatrix = signif(mat, 2),
               main = "Text- + color-labeled heatmap");

```

---

labelPoints

*Label scatterplot points*


---

## Description

Given scatterplot point coordinates, the function tries to place labels near the points such that the labels overlap as little as possible. User beware: the algorithm implemented here is quite primitive and while it will help in many cases, it is by no means perfect. Consider this function experimental. We hope to improve the algorithm in the future to make it useful in a broader range of situations.

## Usage

```

labelPoints(
  x, y, labels,
  cex = 0.7, offs = 0.01, xpd = TRUE,
  jiggle = 0, protectEdges = TRUE, ...)

```

## Arguments

x	a vector of x coordinates of the points
y	a vector of y coordinates of the points
labels	labels to be placed next to the points
cex	character expansion factor for the labels



offs	offset of the labels from the plotted coordinates in inches
xpd	logical: controls truncating labels to fit within the plotting region. See <a href="#">par</a> .
jiggle	amount of random noise to be added to the coordinates. This may be useful if the scatterplot is too regular (such as all points on one straight line).
protectEdges	logical: should labels be shifted inside the (actual or virtual) frame of the plot?
...	other arguments to function <a href="#">text</a> .

### Details

The algorithm basically works by finding the direction of most surrounding points, and attempting to place the label in the opposite direction. There are (not uncommon) situations in which this placement is suboptimal; the author promises to further develop the function sometime in the future.

Note that this function does not plot the actual scatterplot; only the labels are plotted. Plotting the scatterplot is the responsibility of the user.

The argument `offs` needs to be carefully tuned to the size of the plotted symbols. Sorry, no automation here yet.

The argument `protectEdges` can be used to shift labels that would otherwise extend beyond the plot to within the plot. Sometimes this may cause some overlapping with other points or labels; use with care.

### Value

None.

### Author(s)

Peter Langfelder

### See Also

[plot.default](#), [text](#)

### Examples

```
# generate some random points
set.seed(11);
n = 20;
x = runif(n);
y = runif(n);

# Create a basic scatterplot
col = standardColors(n);
plot(x,y, pch = 21, col = col, bg = col, cex = 2.6,
      xlim = c(-0.1, 1.1), ylim = c(-0.1, 1.0));
labelPoints(x, y, paste("Pt", c(1:n), sep=""), offs = 0.10, cex = 1);

# label points using longer text labels. Note the positioning is not perfect, but close enough.

plot(x,y, pch = 21, col = col, bg = col, cex = 2.6,
      xlim = c(-0.1, 1.1), ylim = c(-0.1, 1.0));
labelPoints(x, y, col, offs = 0.10, cex = 0.8);
```

---

labels2colors	<i>Convert numerical labels to colors.</i>
---------------	--

---

### Description

Converts a vector or array of numerical labels into a corresponding vector or array of colors corresponding to the labels.

### Usage

```
labels2colors(labels, zeroIsGrey = TRUE, colorSeq = NULL, naColor = "grey",  
              commonColorCode = TRUE)
```

### Arguments

labels	Vector or matrix of non-negative integer or other (such as character) labels. See details.
zeroIsGrey	If TRUE, labels 0 will be assigned color grey. Otherwise, labels below 1 will trigger an error.
colorSeq	Color sequence corresponding to labels. If not given, a standard sequence will be used.
naColor	Color that will encode missing values.
commonColorCode	logical: if labels is a matrix, should each column have its own colors?

### Details

If labels is numeric, it is used directly as index to the standard color sequence. If 0 is present among the labels and zeroIsGrey=TRUE, labels 0 are given grey color.

If labels is not numeric, its columns are turned into factors and the numeric representation of each factor is used to assign the corresponding colors. In this case commonColorCode governs whether each column gets its own color code, or whether the color code will be universal.

The standard sequence start with well-distinguishable colors, and after about 40 turns into a quasi-random sampling of all colors available in R with the exception of all shades of grey (and gray).

If the input labels have a dimension attribute, it is copied into the output, meaning the dimensions of the returned value are the same as those of the input labels.

### Value

A vector or array of character strings of the same length or dimensions as labels.

### Author(s)

Peter Langfelder, <Peter.Langfelder@gmail.com>

**Examples**

```

labels = c(0:20);
labels2colors(labels);
labels = matrix(letters[1:9], 3,3);
labels2colors(labels)
# Note the difference when commonColorCode = FALSE
labels2colors(labels, commonColorCode = FALSE)

```

---

lowerTri2matrix	<i>Reconstruct a symmetric matrix from a distance (lower-triangular) representation</i>
-----------------	---

---

**Description**

Assuming the input vector contains a vectorized form of the distance representation of a symmetric matrix, this function creates the corresponding matrix. This is useful when re-forming symmetric matrices that have been vectorized to save storage space.

**Usage**

```
lowerTri2matrix(x, diag = 1)
```

**Arguments**

x	a numeric vector
diag	value to be put on the diagonal. Recycled if necessary.

**Details**

The function assumes that x contains the vectorized form of the distance representation of a symmetric matrix. In particular, x must have a length that can be expressed as  $n*(n-1)/2$ , with n an integer. The result of the function is then an n times n matrix.

**Value**

A symmetric matrix whose lower triangle is given by x.

**Author(s)**

Peter Langfelder

**Examples**

```

# Create a symmetric matrix
m = matrix(c(1:16), 4,4)
mat = (m + t(m));
diag(mat) = 0;

# Print the matrix
mat

# Take the lower triangle and vectorize it (in two ways)

```

```
x1 = mat[lower.tri(mat)]
x2 = as.vector(as.dist(mat))

all.equal(x1, x2) # The vectors are equal

# Turn the vectors back into matrices
new.mat = lowerTri2matrix(x1, diag = 0);

# Did we get back the same matrix?

all.equal(mat, new.mat)
```

---

matchLabels

*Relabel module labels to best match the given reference labels*

---

### Description

Given a source and reference vectors of module labels, the function produces a module labeling that is equivalent to source, but individual modules are re-labeled so that modules with significant overlap in source and reference have the same labels.

### Usage

```
matchLabels(source, reference, pThreshold = 5e-2)
```

### Arguments

source	a vector or a matrix of reference labels. The labels may be numeric or character.
reference	a vector of reference labels.
pThreshold	threshold of Fisher's exact test for considering modules to have a significant overlap.

### Details

Each column of source is treated separately. Source and reference labels are assumed to be of the same type, that is noth should be either numeric or character. If the labels are character, they are assumed to be color labels such as the ones returned by [standardColors](#).

The function calculates the overlap of the source and reference modules using Fisher's exact test. It then attempts to relabel source modules such that modules with the highest overlap with the reference modules have the same color. Where this is not possible (for example because one reference module has the highest overlap with two source modules), the source modules will be relabeled using labels that are not present among the reference labels.

### Value

A vector (if the input source labels are a vector) or a matrix (if the input source labels are a matrix) of the new labels.

### Author(s)

Peter Langfelder

**See Also**

[standardColors](#)

---

matrixToNetwork	<i>Construct a network from a matrix</i>
-----------------	--

---

**Description**

Constructs a network

**Usage**

```
matrixToNetwork(
  mat,
  symmetrizeMethod = c("average", "min", "max"),
  signed = TRUE,
  min = NULL, max = NULL,
  power = 12,
  diagEntry = 1)
```

**Arguments**

mat	matrix to be turned into a network. Must be square.
symmetrizeMethod	method for symmetrizing the matrix. The method will be applied to each component of mat and its transpose.
signed	logical: should the resulting network be signed? Unsigned networks are constructed from <code>abs(mat)</code> .
min	minimum allowed value for mat. If NULL, the actual attained minimum of mat will be used. Missing data are ignored. Values below min are truncated to min.
max	maximum allowed value for mat. If NULL, the actual attained maximum of mat will be used. Missing data are ignored. Values below max are truncated to max.
power	the soft-thresholding power.
diagEntry	the value of the entries on the diagonal in the result. This is usually 1 but some applications may require a zero (or even NA) diagonal.

**Details**

If `signed` is FALSE, the matrix `mat` is first converted to its absolute value.

This function then symmetrizes the matrix using the `symmetrizeMethod` component-wise on `mat` and `t(mat)` (i.e., the transpose of `mat`).

In the next step, the symmetrized matrix is linearly scaled to the interval [0,1] using either `min` and `max` (each either supplied or determined from the matrix). Values outside of the [min, max] range are truncated to min or max.

Lastly, the adjacency is calculated by raising the matrix to `power`. The diagonal of the result is set to `diagEntry`. Note that most WGCNA functions expect the diagonal of an adjacency matrix to be 1.

**Value**

The adjacency matrix that encodes the network.

**Author(s)**

Peter Langfelder

**See Also**

adjacency for calculation of a correlation network (adjacency) from a numeric matrix such as expression data

adjacency.fromSimilarity for simpler calculation of a network from a symmetric similarity matrix.

---

mergeCloseModules      *Merge close modules in gene expression data*

---

**Description**

Merges modules in gene expression networks that are too close as measured by the correlation of their eigengenes.

**Usage**

```
mergeCloseModules(exprData, colors,
                  consensusQuantile = 0,
                  cutHeight = 0.2,
                  MEs = NULL,
                  impute = TRUE,
                  useAbs = FALSE,
                  corFnc = cor, corOptions = list(use = 'p'),
                  iterate = TRUE,
                  relabel = FALSE,
                  colorSeq = NULL,
                  getNewMEs = TRUE,
                  getNewUnassdME = TRUE,
                  useSets = NULL,
                  checkDataFormat = TRUE,
                  unassdColor = ifelse(is.numeric(colors), 0, "grey"),
                  trapErrors = FALSE,
                  verbose = 1, indent = 0)
```

**Arguments**

**exprData**      Expression data, either a single data frame with rows corresponding to samples and columns to genes, or in a multi-set format (see [checkSets](#)). See [checkDataStructure](#) below.

**colors**      A vector (numeric, character or a factor) giving module colors for genes. The method only makes sense when genes have the same color label in all sets, hence a single vector.

consensusQuantile	A number giving the desired quantile to use in the consensus similarity calculation (see details).
cutHeight	Maximum dissimilarity (i.e., 1-correlation) that qualifies modules for merging.
MEs	If module eigengenes have been calculated before, the user can save some computational time by inputting them. MEs should have the same format as <code>exprData</code> . If they are not given, they will be calculated.
impute	Should missing values be imputed in eigengene calculation? If imputation is disabled, the presence of NA entries will cause the eigengene calculation to fail and eigengenes will be replaced by their hubgene approximation. See <a href="#">moduleEigengenes</a> for more details.
useAbs	Specifies whether absolute value of correlation or plain correlation (of module eigengenes) should be used in calculating module dissimilarity.
corFnc	Correlation function to be used to calculate correlation of module eigengenes.
corOptions	Can be used to specify options to the correlation function, in addition to argument <code>x</code> which is used to pass the actual data to calculate the correlation of.
iterate	Controls whether the merging procedure should be repeated until there is no change. If FALSE, only one iteration will be executed.
relabel	Controls whether, after merging, color labels should be ordered by module size.
colorSeq	Color labels to be used for relabeling. Defaults to the standard color order used in this package if <code>colors</code> are not numeric, and to integers starting from 1 if <code>colors</code> is numeric.
getNewMEs	Controls whether module eigengenes of merged modules should be calculated and returned.
getNewUnassdME	When doing module eigengene manipulations, the function does not normally calculate the eigengene of the 'module' of unassigned ('grey') genes. Setting this option to TRUE will force the calculation of the unassigned eigengene in the returned <code>newMEs</code> , but not in the returned <code>oldMEs</code> .
useSets	A vector of scalar allowing the user to specify which sets will be used to calculate the consensus dissimilarity of module eigengenes. Defaults to all given sets.
checkDataFormat	If TRUE, the function will check <code>exprData</code> and MEs for correct multi-set structure. If single set data is given, it will be converted into a format usable for the function. If FALSE, incorrect structure of input data will trigger an error.
unassdColor	Specifies the string that labels unassigned genes. Module of this color will not enter the module eigengene clustering and will not be merged with other modules.
trapErrors	Controls whether computational errors in calculating module eigengenes, their dissimilarity, and merging trees should be trapped. If TRUE, errors will be trapped and the function will return the input colors. If FALSE, errors will cause the function to stop.
verbose	Controls verbosity of printed progress messages. 0 means silent, up to (about) 5 the verbosity gradually increases.
indent	A single non-negative integer controlling indentation of printed messages. 0 means no indentation, each unit above that adds two spaces.

**Details**

This function returns the color labels for modules that are obtained from the input modules by merging ones that are closely related. The relationships are quantified by correlations of module eigengenes; a “consensus” measure is defined as the “consensus quantile” over the corresponding relationship in each set. Once the (dis-)similarity is calculated, average linkage hierarchical clustering of the module eigengenes is performed, the dendrogram is cut at the height `cutHeight` and modules on each branch are merged. The process is (optionally) repeated until no more modules are merged.

If, for a particular module, the module eigene calculation fails, a hubgene approximation will be used.

The user should be aware that if a computational error occurs and `trapErrors==TRUE`, the returned list (see below) will not contain all of the components returned upon normal execution.

**Value**

If no errors occurred, a list with components

<code>colors</code>	Color labels for the genes corresponding to merged modules. The function attempts to mimic the mode of the input <code>colors</code> : if the input <code>colors</code> is numeric, character and factor, respectively, so is the output. Note, however, that if the function performs relabeling, a standard sequence of labels will be used: integers starting at 1 if the input <code>colors</code> is numeric, and a sequence of color labels otherwise (see <code>colorSeq</code> above).
<code>dendro</code>	Hierarchical clustering dendrogram (average linkage) of the eigengenes of the most recently computed tree. If <code>iterate</code> was set <code>TRUE</code> , this will be the dendrogram of the merged modules, otherwise it will be the dendrogram of the original modules.
<code>oldDendro</code>	Hierarchical clustering dendrogram (average linkage) of the eigengenes of the original modules.
<code>cutHeight</code>	The input <code>cutHeight</code> .
<code>oldMEs</code>	Module eigengenes of the original modules in the sets given by <code>useSets</code> .
<code>newMEs</code>	Module eigengenes of the merged modules in the sets given by <code>useSets</code> .
<code>allOK</code>	A boolean set to <code>TRUE</code> .

If an error occurred and `trapErrors==TRUE`, the list only contains these components:

<code>colors</code>	A copy of the input <code>colors</code> .
<code>allOK</code>	a boolean set to <code>FALSE</code> .

**Author(s)**

Peter Langfelder, <Peter.Langfelder@gmail.com>



metaAnalysis

*Meta-analysis of binary and continuous variables***Description**

This is a meta-analysis complement to functions [standardScreeningBinaryTrait](#) and [standardScreeningNumericTrait](#). Given expression (or other) data from multiple independent data sets, and the corresponding clinical traits or outcomes, the function calculates multiple screening statistics in each data set, then calculates meta-analysis Z scores, p-values, and optionally q-values (False Discovery Rates). Three different ways of calculating the meta-analysis Z scores are provided: the Stouffer method, weighted Stouffer method, and using user-specified weights.

**Usage**

```
metaAnalysis(multiExpr, multiTrait,
             binary = NULL,
             metaAnalysisWeights = NULL,
             corFnc = cor, corOptions = list(use = "p"),
             getQvalues = FALSE,
             getAreaUnderROC = FALSE,
             useRankPvalue = TRUE,
             rankPvalueOptions = list(),
             setNames = NULL,
             kruskalTest = FALSE, var.equal = FALSE,
             metaKruskal = kruskalTest, na.action = "na.exclude")
```

**Arguments**

multiExpr	Expression data (or other data) in multi-set format (see <a href="#">checkSets</a> ). A vector of lists; in each list there must be a component named data whose content is a matrix or dataframe or array of dimension 2.
multiTrait	Trait or outcome data in multi-set format. Only one trait is allowed; consequently, the data component of each component list can be either a vector or a dataframe (matrix, array of dimension 2).
binary	Logical: is the trait binary (TRUE) or continuous (FALSE)? If not given, the decision will be made based on the content of multiTrait.
metaAnalysisWeights	Optional specification of set weights for meta-analysis. If given, must be a vector of non-negative weights, one entry for each set contained in multiExpr.
corFnc	Correlation function to be used for screening. Should be either the default <a href="#">cor</a> or its robust alternative, <a href="#">bicor</a> .
corOptions	A named list giving extra arguments to be passed to the correlation function.
getQvalues	Logical: should q-values (FDRs) be calculated?
getAreaUnderROC	Logical: should area under the ROC be calculated? Caution, enabling the calculation will slow the function down considerably for large data sets.
useRankPvalue	Logical: should the <a href="#">rankPvalue</a> function be used to obtain alternative meta-analysis statistics?

rankPvalueOptions	Additional options for function <code>rankPvalue</code> . These include <code>na.last</code> (default "keep"), <code>ties.method</code> (default "average"), <code>calculateQvalue</code> (default copied from input <code>getQvalues</code> ), and <code>pvalueMethod</code> (default "all"). See the help file for <code>rankPvalue</code> for full details.
setNames	Optional specification of set names (labels). These are used to label the corresponding components of the output. If not given, will be taken from the <code>names</code> attribute of <code>multiExpr</code> . If <code>names(multiExpr)</code> is NULL, generic names of the form <code>Set_1</code> , <code>Set2</code> , ... will be used.
kruskalTest	Logical: should the Kruskal test be performed in addition to t-test? Only applies to binary traits.
var.equal	Logical: should the t-test assume equal variance in both groups? If TRUE, the function will warn the user that the returned test statistics will be different from the results of the standard <code>t.test</code> function.
metaKruskal	Logical: should the meta-analysis be based on the results of Kruskal test (TRUE) or Student t-test (FALSE)?
na.action	Specification of what should happen to missing values in <code>t.test</code> .

## Details

The Stouffer method of combines Z statistics by simply taking a mean of input Z statistics and multiplying it by  $\sqrt{n}$ , where n is the number of input data sets. We refer to this method as `Stouffer.equalWeights`. In general, a better (i.e., more powerful) method of combining Z statistics is to weigh them by the number of degrees of freedom (which approximately equals n). We refer to this method as `weightedStouffer`. Finally, the user can also specify custom weights, for example if a data set needs to be downweighted due to technical concerns; however, specifying own weights by hand should be done carefully to avoid possible selection biases.

## Value

Data frame with the following components:

ID	Identifier of the input genes (or other variables)
Z.equalWeights	Meta-analysis Z statistics obtained using Stouffer's method with equal weights
p.equalWeights	p-values corresponding to Z.Stouffer.equalWeights
q.equalWeights	q-values corresponding to p.Stouffer.equalWeights, only present if <code>getQvalues</code> is TRUE.
Z.RootDoFWeights	Meta-analysis Z statistics obtained using Stouffer's method with weights given by the square root of the number of (non-missing) samples in each data set
p.RootDoFWeights	p-values corresponding to Z.DoFWeights
q.RootDoFWeights	q-values corresponding to p.DoFWeights, only present if <code>getQvalues</code> is TRUE.
Z.DoFWeights	Meta-analysis Z statistics obtained using Stouffer's method with weights given by the number of (non-missing) samples in each data set
p.DoFWeights	p-values corresponding to Z.DoFWeights
q.DoFWeights	q-values corresponding to p.DoFWeights, only present if <code>getQvalues</code> is TRUE.
Z.userWeights	Meta-analysis Z statistics obtained using Stouffer's method with user-defined weights. Only present if input <code>metaAnalysisWeights</code> are present.

p.userWeights p-values corresponding to Z.userWeights  
 q.userWeights q-values corresponding to p.userWeights, only present if getQvalues is TRUE.

The next set of columns is present only if input useRankPvalue is TRUE and contain the output of the function [rankPvalue](#) with the same column weights as the above meta-analysis. Depending on the input options calculateQvalue and pvalueMethod in rankPvalueOptions, some columns may be missing. The following columns are calculated using equal weights for each data set.

pValueExtremeRank.equalWeights  
 This is the minimum between pValueLowRank and pValueHighRank, i.e.  $\min(\text{pValueLow}, \text{pValueHigh})$

pValueLowRank.equalWeights  
 Asymptotic p-value for observing a consistently low value across the columns of datS based on the rank method.

pValueHighRank.equalWeights  
 Asymptotic p-value for observing a consistently low value across the columns of datS based on the rank method.

pValueExtremeScale.equalWeights  
 This is the minimum between pValueLowScale and pValueHighScale, i.e.  $\min(\text{pValueLow}, \text{pValueHigh})$

pValueLowScale.equalWeights  
 Asymptotic p-value for observing a consistently low value across the columns of datS based on the Scale method.

pValueHighScale.equalWeights  
 Asymptotic p-value for observing a consistently low value across the columns of datS based on the Scale method.

qValueExtremeRank.equalWeights  
 local false discovery rate (q-value) corresponding to the p-value pValueExtremeRank

qValueLowRank.equalWeights  
 local false discovery rate (q-value) corresponding to the p-value pValueLowRank

qValueHighRank.equalWeights  
 local false discovery rate (q-value) corresponding to the p-value pValueHighRank

qValueExtremeScale.equalWeights  
 local false discovery rate (q-value) corresponding to the p-value pValueExtremeScale

qValueLowScale.equalWeights  
 local false discovery rate (q-value) corresponding to the p-value pValueLowScale

qValueHighScale.equalWeights  
 local false discovery rate (q-value) corresponding to the p-value pValueHighScale

... Analogous columns calculated by weighting each input set using the square root of the number of samples, number of samples, and user weights (if given). The corresponding column names carry the suffixes RootDofWeights, DofWeights, userWeights.

The following columns contain results returned by [standardScreeningBinaryTrait](#) or [standardScreeningNumericTrait](#) (depending on whether the input trait is binary or continuous).

For binary traits, the following information is returned for each set:

`corPearson.Set_1`, `corPearson.Set_2`, ...  
 Pearson correlation with a binary numeric version of the input variable. The numeric variable equals 1 for level 1 and 2 for level 2. The levels are given by `levels(factor(y))`.

`t.Student.Set_1`, `t.Student.Set_2`, ...  
 Student t-test statistic

`pvalueStudent.Set_1`, `pvalueStudent.Set_2`, ...  
 two-sided Student t-test p-value.

`qvalueStudent.Set_1`, `qvalueStudent.Set_2`, ...  
 (if input `qValues==TRUE`) q-value (local false discovery rate) based on the Student T-test p-value (Storey et al 2004).

`foldChange.Set_1`, `foldChange.Set_2`, ...  
 a (signed) ratio of mean values. If the mean in the first group (corresponding to level 1) is larger than that of the second group, it equals `meanFirstGroup/meanSecondGroup`. But if the mean of the second group is larger than that of the first group it equals `-meanSecondGroup/meanFirstGroup` (notice the minus sign).

`meanFirstGroup.Set_1`, `meanSecondGroup.Set_2`, ...  
 means of columns in input `datExpr` across samples in the second group.

`SE.FirstGroup.Set_1`, `SE.FirstGroup.Set_2`, ...  
 standard errors of columns in input `datExpr` across samples in the first group. Recall that  $SE(x)=\sqrt{\text{var}(x)/n}$  where `n` is the number of non-missing values of `x`.

`SE.SecondGroup.Set_1`, `SE.SecondGroup.Set_2`, ...  
 standard errors of columns in input `datExpr` across samples in the second group.

`areaUnderROC.Set_1`, `areaUnderROC.Set_2`, ...  
 the area under the ROC, also known as the concordance index or C.index. This is a measure of discriminatory power. The measure lies between 0 and 1 where 0.5 indicates no discriminatory power. 0 indicates that the "opposite" predictor has perfect discriminatory power. To compute it we use the function `rcorr.cens` with `outx=TRUE` (from Frank Harrel's package `Hmisc`).

`nPresentSamples.Set_1`, `nPresentSamples.Set_2`, ...  
 number of samples with finite measurements for each gene.

If input `kruskalTest` is `TRUE`, the following columns further summarize results of Kruskal-Wallis test:

`stat.Kruskal.Set_1`, `stat.Kruskal.Set_2`, ...  
 Kruskal-Wallis test statistic.

`stat.Kruskal.signed.Set_1`, `stat.Kruskal.signed.Set_2`, ...  
 (Warning: experimental) Kruskal-Wallis test statistic including a sign that indicates whether the average rank is higher in second group (positive) or first group (negative).

`pvaluekruskal.Set_1`, `pvaluekruskal.Set_2`, ...  
 Kruskal-Wallis test p-value.

`qkruskal.Set_1`, `qkruskal.Set_2`, ...  
 q-values corresponding to the Kruskal-Wallis test p-value (if input `qValues==TRUE`).

`Z.Set1`, `Z.Set2`, ...  
 Z statistics obtained from `pvalueStudent.Set1`, `pvalueStudent.Set2`, ... or from `pvaluekruskal.Set1`, `pvaluekruskal.Set2`, ..., depending on input `metaKruskal`.

For numeric traits, the following columns are returned:

cor.Set\_1, cor.Set\_2, ...  
     correlations of all genes with the trait

Z.Set1, Z.Set2, ...  
     Fisher Z statistics corresponding to the correlations

pvalueStudent.Set\_1, pvalueStudent.Set\_2, ...  
     Student p-values of the correlations

qvalueStudent.Set\_1, qvalueStudent.Set\_1, ...  
     (if input qValues==TRUE) q-values of the correlations calculated from the p-values

AreaUnderROC.Set\_1, AreaUnderROC.Set\_2, ...  
     area under the ROC

nPresentSamples.Set\_1, nPresentSamples.Set\_2, ...  
     number of samples present for the calculation of each association.

### Author(s)

Peter Langfelder

### References

For Stouffer's method, see

Stouffer, S.A., Suchman, E.A., DeVinney, L.C., Star, S.A. & Williams, R.M. Jr. 1949. The American Soldier, Vol. 1: Adjustment during Army Life. Princeton University Press, Princeton.

A discussion of weighted Stouffer's method can be found in

Whitlock, M. C., Combining probability from independent tests: the weighted Z-method is superior to Fisher's approach, *Journal of Evolutionary Biology* 18:5 1368 (2005)

### See Also

[standardScreeningBinaryTrait](#), [standardScreeningNumericTrait](#) for screening functions for individual data sets

---

metaZfunction	<i>Meta-analysis Z statistic</i>
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---

### Description

The function calculates a meta analysis Z statistic based on an input data frame of Z statistics.

### Usage

```
metaZfunction(datZ, columnweights = NULL)
```

### Arguments

**datZ**                    Matrix or data frame of Z statistics (assuming standard normal distribution under the null hypothesis). Rows correspond to genes, columns to independent data sets.

**columnweights**       optional vector of non-negative numbers for weighing the columns of datZ.

**Details**

For example, if datZ has 3 columns whose columns are labelled Z1,Z2,Z3 then ZMeta=(Z1+Z2+Z3)/sqrt(3). Under the null hypothesis (where all Z statistics follow a standard normal distribution and the Z statistics are independent), ZMeta also follows a standard normal distribution. To calculate a 2 sided p-value, one can use the following code `pvalue=2*pnorm(-abs(ZMeta) )`

**Value**

Vector of meta analysis Z statistic. Under the null hypothesis this should follow a standard normal distribution.

**Author(s)**

Steve Horvath

---

```
moduleColor.getMEprefix
```

*Get the prefix used to label module eigengenes.*

---

**Description**

Returns the currently used prefix used to label module eigengenes. When returning module eigengenes in a dataframe, names of the corresponding columns will start with the given prefix.

**Usage**

```
moduleColor.getMEprefix()
```

**Details**

Returns the prefix used to label module eigengenes. When returning module eigengenes in a dataframe, names of the corresponding columns will consist of the corresponding color label preceded by the given prefix. For example, if the prefix is "PC" and the module is turquoise, the corresponding module eigengene will be labeled "PCturquoise". Most of old code assumes "PC", but "ME" is more instructive and used in some newer analyses.

**Value**

A character string.

**Note**

Currently the standard prefix is "ME" and there is no way to change it.

**Author(s)**

Peter Langfelder, <Peter.Langfelder@gmail.com>

**See Also**

[moduleEigengenes](#)

---

moduleEigengenes	<i>Calculate module eigengenes.</i>
------------------	-------------------------------------

---

### Description

Calculates module eigengenes (1st principal component) of modules in a given single dataset.

### Usage

```
moduleEigengenes(expr,
                 colors,
                 impute = TRUE,
                 nPC = 1,
                 align = "along average",
                 excludeGrey = FALSE,
                 grey = ifelse(is.numeric(colors), 0, "grey"),
                 subHubs = TRUE,
                 trapErrors = FALSE,
                 returnValidOnly = trapErrors,
                 softPower = 6,
                 scale = TRUE,
                 verbose = 0, indent = 0)
```

### Arguments

expr	Expression data for a single set in the form of a data frame where rows are samples and columns are genes (probes).
colors	A vector of the same length as the number of probes in expr, giving module color for all probes (genes). Color "grey" is reserved for unassigned genes.
impute	If TRUE, expression data will be checked for the presence of NA entries and if the latter are present, numerical data will be imputed, using function <code>impute.knn</code> and probes from the same module as the missing datum. The function <code>impute.knn</code> uses a fixed random seed giving repeatable results.
nPC	Number of principal components and variance explained entries to be calculated. Note that only the first principal component is returned; the rest are used only for the calculation of proportion of variance explained. The number of returned variance explained entries is currently <code>min(nPC, 10)</code> . If given nPC is greater than 10, a warning is issued.
align	Controls whether eigengenes, whose orientation is undetermined, should be aligned with average expression ( <code>align = "along average"</code> , the default) or left as they are ( <code>align = ""</code> ). Any other value will trigger an error.
excludeGrey	Should the improper module consisting of 'grey' genes be excluded from the eigengenes?
grey	Value of colors designating the improper module. Note that if colors is a factor of numbers, the default value will be incorrect.
subHubs	Controls whether hub genes should be substituted for missing eigengenes. If TRUE, each missing eigengene (i.e., eigengene whose calculation failed and the error was trapped) will be replaced by a weighted average of the most connected hub genes in the corresponding module. If this calculation fails, or if

	subHubs==FALSE, the value of trapErrors will determine whether the offending module will be removed or whether the function will issue an error and stop.
trapErrors	Controls handling of errors from that may arise when there are too many NA entries in expression data. If TRUE, errors from calling these functions will be trapped without abnormal exit. If FALSE, errors will cause the function to stop. Note, however, that subHubs takes precedence in the sense that if subHubs==TRUE and trapErrors==FALSE, an error will be issued only if both the principal component and the hubgene calculations have failed.
returnValidOnly	logical; controls whether the returned data frame of module eigengenes contains columns corresponding only to modules whose eigengenes or hub genes could be calculated correctly (TRUE), or whether the data frame should have columns for each of the input color labels (FALSE).
softPower	The power used in soft-thresholding the adjacency matrix. Only used when the hubgene approximation is necessary because the principal component calculation failed. It must be non-negative. The default value should only be changed if there is a clear indication that it leads to incorrect results.
scale	logical; can be used to turn off scaling of the expression data before calculating the singular value decomposition. The scaling should only be turned off if the data has been scaled previously, in which case the function can run a bit faster. Note however that the function first imputes, then scales the expression data in each module. If the expression contain missing data, scaling outside of the function and letting the function impute missing data may lead to slightly different results than if the data is scaled within the function.
verbose	Controls verbosity of printed progress messages. 0 means silent, up to (about) 5 the verbosity gradually increases.
indent	A single non-negative integer controlling indentation of printed messages. 0 means no indentation, each unit above that adds two spaces.

### Details

Module eigengene is defined as the first principal component of the expression matrix of the corresponding module. The calculation may fail if the expression data has too many missing entries. Handling of such errors is controlled by the arguments subHubs and trapErrors. If subHubs==TRUE, errors in principal component calculation will be trapped and a substitute calculation of hubgenes will be attempted. If this fails as well, behaviour depends on trapErrors: if TRUE, the offending module will be ignored and the return value will allow the user to remove the module from further analysis; if FALSE, the function will stop.

From the user's point of view, setting trapErrors=FALSE ensures that if the function returns normally, there will be a valid eigengene (principal component or hubgene) for each of the input colors. If the user sets trapErrors=TRUE, all calculational (but not input) errors will be trapped, but the user should check the output (see below) to make sure all modules have a valid returned eigengene.

While the principal component calculation can fail even on relatively sound data (it does not take all that many "well-placed" NA to torpedo the calculation), it takes many more irregularities in the data for the hubgene calculation to fail. In fact such a failure signals there likely is something seriously wrong with the data.

### Value

A list with the following components:



eigengenes	Module eigengenes in a dataframe, with each column corresponding to one eigengene. The columns are named by the corresponding color with an "ME" prepended, e.g., MEturquoise etc. If <code>returnValidOnly==FALSE</code> , module eigengenes whose calculation failed have all components set to NA.
averageExpr	If <code>align == "along average"</code> , a dataframe containing average normalized expression in each module. The columns are named by the corresponding color with an "AE" prepended, e.g., AEturquoise etc.
varExplained	A dataframe in which each column corresponds to a module, with the component <code>varExplained[PC, module]</code> giving the variance of module explained by the principal component no. PC. The calculation is exact irrespective of the number of computed principal components. At most 10 variance explained values are recorded in this dataframe.
nPC	A copy of the input nPC.
validMEs	A boolean vector. Each component (corresponding to the columns in data) is TRUE if the corresponding eigengene is valid, and FALSE if it is invalid. Valid eigengenes include both principal components and their hubgene approximations. When <code>returnValidOnly==FALSE</code> , by definition all returned eigengenes are valid and the entries of <code>validMEs</code> are all TRUE.
validColors	A copy of the input colors with entries corresponding to invalid modules set to grey if given, otherwise 0 if colors is numeric and "grey" otherwise.
allOK	Boolean flag signalling whether all eigengenes have been calculated correctly, either as principal components or as the hubgene average approximation.
allPC	Boolean flag signalling whether all returned eigengenes are principal components.
isPC	Boolean vector. Each component (corresponding to the columns in eigengenes) is TRUE if the corresponding eigengene is the first principal component and FALSE if it is the hubgene approximation or is invalid.
isHub	Boolean vector. Each component (corresponding to the columns in eigengenes) is TRUE if the corresponding eigengene is the hubgene approximation and FALSE if it is the first principal component or is invalid.
validAEs	Boolean vector. Each component (corresponding to the columns in eigengenes) is TRUE if the corresponding module average expression is valid.
allAEOK	Boolean flag signalling whether all returned module average expressions contain valid data. Note that <code>returnValidOnly==TRUE</code> does not imply <code>allAEOK==TRUE</code> : some invalid average expressions may be returned if their corresponding eigengenes have been calculated correctly.

**Author(s)**

Steve Horvath <SHorvath@mednet.ucla.edu>, Peter Langfelder <Peter.Langfelder@gmail.com>

**References**

Zhang, B. and Horvath, S. (2005), "A General Framework for Weighted Gene Co-Expression Network Analysis", *Statistical Applications in Genetics and Molecular Biology*: Vol. 4: No. 1, Article 17

**See Also**

[svd](#), [impute.knn](#)

---

moduleMergeUsingKME     *Merge modules and reassign genes using kME.*

---

### Description

This function takes an expression data matrix (and other user-defined parameters), calculates the module membership (kME) values, and adjusts the module assignments, merging modules that are not sufficiently distinct and reassigning modules that were originally assigned suboptimally.

### Usage

```
moduleMergeUsingKME(
  datExpr, colorh, ME = NULL,
  threshPercent = 50, mergePercent = 25,
  reassignModules = TRUE,
  convertGrey = TRUE,
  omitColors = "grey",
  reassignScale = 1,
  threshNumber = NULL)
```

### Arguments

datExpr	An expression data matrix, with samples as rows, genes (or probes) as column.
colorh	The color vector (module assignments) corresponding to the columns of datExpr.
ME	Either NULL (default), at which point the module eigengenes will be calculated, or pre-calculated module eigengenes for each of the modules, with samples as rows (corresponding to datExpr), and modules corresponding to columns (column names MUST be module colors or module colors prefixed by "ME" or "PC").
threshPercent	Threshold percent of the number of genes in the module that should be included for the various analyses. For example, in a module with 200 genes, if threshPercent=50 (default), then 50 genes will be checked for reassignment and used to test whether two modules should be merged. See also threshNumber.
mergePercent	If greater than this percent of the assigned genes are above the threshold are in a module other than the assigned module, then these two modules will be merged. For example, if mergePercent=25 (default), and the 70 out of 200 genes in the blue module were more highly correlated with the black module eigengene, then all genes in the blue module would be reassigned to the black module.
reassignModules	If TRUE (default), genes are reassigned to the module with which they have the highest module membership (kME), but only if their kME is above the threshPercent (or threshNumber) threshold of that module.
convertGrey	If TRUE (default), unassigned (grey) genes are assigned as in "reassignModules"
omitColors	These are all of the module assignments which indicate genes that are not assigned to modules (default="grey"). These genes will all be assigned as "grey" by this function.

- reassignScale A value between 0 and 1 (default) which determines how the threshPercent gets scaled for reassigning genes. Smaller values reassign more genes, but does not affect the merging process.
- threshNumber Either NULL (default) or, if entered, every module is counted as having exactly threshNumber genes, and threshPercent is ignored. This parameter should have the effect of

**Value**

- moduleColors The NEW color vector (module assignments) corresponding to the columns of datExpr, after module merging and reassignments.
- mergeLog A log of the order in which modules were merged, for reference.

**Note**

Note that this function should be considered "experimental" as it has only been beta tested. Please e-mail [jeremyinla@gmail.com](mailto:jeremyinla@gmail.com) if you have any issues with the function.

**Author(s)**

Jeremy Miller

**Examples**

```
## First simulate some data and the resulting network dendrogram
set.seed(100)
MEturquoise = sample(1:100,50)
MEblue      = sample(1:100,50)
MEbrown     = sample(1:100,50)
MEyellow    = sample(1:100,50)
MEgreen     = c(MEyellow[1:30], sample(1:100,20))
MERed      = c(MEbrown [1:20], sample(1:100,30))
#MEblack    = c(MEblue  [1:25], sample(1:100,25))
ME = data.frame(MEturquoise, MEblue, MEbrown, MEyellow, MEGreen, MERed)#, MEblack)
dat1 = simulateDatExpr(ME, 400, c(0.15,0.13,0.12,0.10,0.09,0.09,0.1), signed=TRUE)
TOM1 = TOMsimilarityFromExpr(dat1$datExpr, networkType="signed")
tree1 = flashClust(as.dist(1-TOM1),method="average")

## Here is an example using different mergePercentages,
# setting an inclusive threshPercent (91)
colorh1 <- colorPlot <- labels2colors(dat1$allLabels)
merges = c(65,40,20,5)
for (m in merges)
  colorPlot = cbind(colorPlot,
                    moduleMergeUsingKME(dat1$datExpr,colorh1,
                                         threshPercent=91, mergePercent=m)$moduleColors)
plotDendroAndColors(tree1, colorPlot, c("ORIG",merges), dendroLabels=FALSE)

## Here is an example using a lower reassignScale (so that more genes get reassigned)
colorh1 <- colorPlot <- labels2colors(dat1$allLabels)
merges = c(65,40,20,5)
for (m in merges) colorPlot = cbind(colorPlot,
                                     moduleMergeUsingKME(dat1$datExpr,colorh1,threshPercent=91,
                                                         reassignScale=0.7, mergePercent=m)$moduleColors)
```

```

plotDendroAndColors(tree1, colorPlot, c("ORIG",merges), dendroLabels=FALSE)

## Here is an example using a less-inclusive threshPercent (75),
# little if anything is merged.

colorh1 <- colorPlot <- labels2colors(dat1$allLabels)
merges = c(65,40,20,5)
for (m in merges) colorPlot = cbind(colorPlot,
  moduleMergeUsingKME(dat1$datExpr,colorh1,
    threshPercent=75, mergePercent=m)$moduleColors)
plotDendroAndColors(tree1, colorPlot, c("ORIG",merges), dendroLabels=FALSE)
# (Note that with real data, the default threshPercent=50 usually results
# in some modules being merged)

```

---

moduleNumber	<i>Fixed-height cut of a dendrogram.</i>
--------------	--

---

### Description

Detects branches of on the input dendrogram by performing a fixed-height cut.

### Usage

```
moduleNumber(dendro, cutHeight = 0.9, minSize = 50)
```

### Arguments

dendro	a hierarchical clustering dendrogram such as one returned by hclust.
cutHeight	Maximum joining heights that will be considered.
minSize	Minimum cluster size.

### Details

All contiguous branches below the height cutHeight that contain at least minSize objects are assigned unique positive numerical labels; all unassigned objects are assigned label 0.

### Value

A vector of numerical labels giving the assignment of each object.

### Note

The numerical labels may not be sequential. See [normalizeLabels](#) for a way to put the labels into a standard order.

### Author(s)

Peter Langfelder, <Peter.Langfelder@gmail.com>

### See Also

[hclust](#), [cutree](#), [normalizeLabels](#)

---

modulePreservation	<i>Calculation of module preservation statistics</i>
--------------------	--

---

## Description

Calculations of module preservation statistics between independent data sets.

## Usage

```
modulePreservation(
  multiData,
  multiColor,
  dataIsExpr = TRUE,
  networkType = "unsigned",
  corFnc = "cor",
  corOptions = "use = 'p'",
  referenceNetworks = 1,
  nPermutations = 100,
  includekMEallInSummary = FALSE,
  restrictSummaryForGeneralNetworks = TRUE,
  calculateQvalue = FALSE,
  randomSeed = 12345,
  maxGoldModuleSize = 1000,
  maxModuleSize = 1000,
  quickCor = 1,
  ccTupletSize = 2,
  calculateCor.kIMall = FALSE,
  calculateClusterCoeff = FALSE,
  useInterpolation = FALSE,
  checkData = TRUE,
  greyName = NULL,
  savePermutedStatistics = TRUE,
  loadPermutedStatistics = FALSE,
  permutedStatisticsFile = if (useInterpolation) "permutedStats-intrModules.RData"
                           else "permutedStats-actualModules.RData",

  plotInterpolation = TRUE,
  interpolationPlotFile = "modulePreservationInterpolationPlots.pdf",
  discardInvalidOutput = TRUE,
  verbose = 1, indent = 0)
```

## Arguments

- |            |  |
|------------|--|
| multiData  | expression data or adjacency data in the multi-set format (see <a href="#">checkSets</a> ). A vector of lists, one per set. Each set must contain a component data that contains the expression or adjacency data. If expression data are used, rows correspond to samples and columns to genes or probes. In case of adjacencies, each data matrix should be a symmetric matrix with entries between 0 and 1 and unit diagonal. Each component of the outermost list should be named. |
| multiColor | a list in which every component is a vector giving the module labels of genes in multiExpr. The components must be named using the same names that are   |

	used in <code>multiExpr</code> ; these names are used to match labels to expression data sets. See details.
<code>dataIsExpr</code>	logical: if TRUE, <code>multiData</code> will be interpreted as expression data; if FALSE, <code>multiData</code> will be interpreted as adjacencies.
<code>networkType</code>	network type. Allowed values are (unique abbreviations of) "unsigned", "signed", "signed hybrid". See <a href="#">adjacency</a> .
<code>corFnc</code>	character string specifying the function to be used to calculate co-expression similarity. Defaults to Pearson correlation. Another useful choice is <a href="#">bicor</a> . More generally, any function returning values between -1 and 1 can be used.
<code>corOptions</code>	character string specifying additional arguments to be passed to the function given by <code>corFnc</code> . Use "use = 'p', method = 'spearman'" to obtain Spearman correlation.
<code>referenceNetworks</code>	a vector giving the indices of expression data to be used as reference networks. Reference networks must have their module labels given in <code>multiColor</code> .
<code>nPermutations</code>	specifies the number of permutations that will be calculated in the permutation test.
<code>includekMEallInSummary</code>	logical: should <code>cor.kMEall</code> be included in the calculated summary statistics? Because <code>kMEall</code> takes into account all genes in the network, this statistic measures preservation of the full network with respect to the eigengene of the module. This may be undesirable, hence the default is FALSE.
<code>restrictSummaryForGeneralNetworks</code>	logical: should the summary statistics for general (not correlation) networks be restricted (density to <code>meanAdj</code> , connectivity to <code>cor.kIM</code> and <code>cor.Adj</code> )? The default TRUE corresponds to published work.
<code>calculateQvalue</code>	logical: should q-values (local FDR estimates) be calculated? Package <code>qvalue</code> must be installed for this calculation. Note that q-values may not be meaningful when the number of modules is small and/or most modules are preserved.
<code>randomSeed</code>	seed for the random number generator. If NULL, the seed will not be set. If non-NULL and the random generator has been initialized prior to the function call, the latter's state is saved and restored upon exit
<code>maxGoldModuleSize</code>	maximum size of the "gold" module, i.e., the random sample of all network genes.
<code>maxModuleSize</code>	maximum module size used for calculations. Modules larger than <code>maxModuleSize</code> will be reduced by randomly sampling <code>maxModuleSize</code> genes.
<code>quickCor</code>	number between 0 and 1 specifying the handling of missing data in calculation of correlation. Zero means exact but potentially slower calculations; one means potentially faster calculations, but with potentially inaccurate results if the proportion of missing data is large. See <a href="#">cor</a> for more details.
<code>ccTupletSize</code>	tuplet size for co-clustering calculations.
<code>calculateCor.kIMall</code>	logical: should <code>cor.kMEall</code> be calculated? This option is only valid for adjacency input. If FALSE, <code>cor.kIMall</code> will not be calculated, potentially saving significant amount of time if the input adjacencies are large and contain many modules.

calculateClusterCoeff	logical: should statistics based on the clustering coefficient be calculated? While these statistics may be interesting, the calculations are also computationally expensive.
checkData	logical: should data be checked for excessive number of missing entries? See <a href="#">goodSamplesGenesMS</a> for details.
greyName	label used for unassigned genes. Traditionally such genes are labeled by grey color or numeric label 0. These values are the default when multiColor contains character or numeric vectors, respectively.
savePermutedStatistics	logical: should calculated permutation statistics be saved? Saved statistics may be re-used if the calculation needs to be repeated.
permutedStatisticsFile	file name to save the permutation statistics into.
loadPermutedStatistics	logical: should permutation statistics be loaded? If a previously executed calculation needs to be repeated, loading permutation study results can cut the calculation time many-fold.
useInterpolation	logical: should permutation statistics be calculated by interpolating an artificial set of evenly spaced modules? This option may potentially speed up the calculations, but it restricts calculations to density measures.
plotInterpolation	logical: should interpolation plots be saved? If interpolation is used (see useInterpolation above), the function can optionally generate diagnostic plots that can be used to assess whether the interpolation makes sense.
interpolationPlotFile	file name to save the interpolation plots into.
discardInvalidOutput	logical: should output columns containing no valid data be discarded? This option may be useful when input dataIsExpr is FALSE and some of the output statistics cannot be calculated. This option causes such statistics to be dropped from output.
verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.
indent	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

## Details

This function calculates module preservation statistics pair-wise between given reference sets and all other sets in multiExpr. Reference sets must have their corresponding module assignment specified in multiColor; module assignment is optional for test sets. Individual expression sets and their module labels are matched using names of the corresponding components in multiExpr and multiColor.

For each reference-test pair, the function calculates module preservation statistics that measure how well the modules of the reference set are preserved in the test set. If the multiColor also contains module assignment for the test set, the calculated statistics also include cross-tabulation statistics that make use of the test module assignment.

For each reference-test pair, the function only uses genes (columns of the data component of each component of `multiExpr`) that are in common between the reference and test set. Columns are matched by column names, so column names must be valid.

In addition to preservation statistics, the function also calculates several statistics of module quality, that is measures of how well-defined modules are in the reference set. The quality statistics are calculated with respect to genes in common with with a test set; thus the function calculates a set of quality statistics for each reference-test pair. This may be somewhat counter-intuitive, but it allows a direct comparison of corresponding quality and preservation statistics.

The calculated p-values are determined from the Z scores of individual measures under assumption of normality. No p-value is calculated for the Zsummary measures. Bonferoni correction to the number of tested modules. Because the p-values for strongly preserved modules are often extremely low, the function reports natural logarithms (base e) of the p-values. However, q-values are reported untransformed since they are calculated that way in package `qvalue`.

Missing data are removed (but see `quickCor` above).

## Value

The function returns a nested list of preservation statistics. At the top level, the list components are:

<code>quality</code>	observed values, Z scores, log p-values, Bonferoni-corrected log p-values, and (optionally) q-values of quality statistics. All logarithms are in base 10.
<code>preservation</code>	observed values, Z scores, log p-values, Bonferoni-corrected log p-values, and (optionally) q-values of density and connectivity preservation statistics. All logarithms are in base 10.
<code>accuracy</code>	observed values, Z scores, log p-values, Bonferoni-corrected log p-values, and (optionally) q-values of cross-tabulation statistics. All logarithms are in base 10.
<code>referenceSeparability</code>	observed values, Z scores, log p-values, Bonferoni-corrected log p-values, and (optionally) q-values of module separability in the reference network. All logarithms are in base 10.
<code>testSeparability</code>	observed values, Z scores, p-values, Bonferoni-corrected p-values, and (optionally) q-values of module separability in the test network. All logarithms are in base 10.
<code>permutationDetails</code>	results of individual permutations, useful for diagnostics

All of the above are lists. The lists `quality`, `preservation`, `referenceSeparability`, and `testSeparability` each contain 4 or 5 components: `observed` contains observed values, `Z` contains the corresponding Z scores, `log.p` contains base 10 logarithms of the p-values, `log.pBonf` contains base 10 logarithms of the Bonferoni corrected p-values, and optionally `q` contains the associated q-values. The list `accuracy` contains `observed`, `Z`, `log.p`, `log.pBonf`, optionally `q`, and additional components `observedOverlapCounts` and `observedFisherPvalues` that contain the observed matrices of overlap counts and Fisher test p-values.

Each of the lists `observed`, `Z`, `log.p`, `log.pBonf`, optionally `q`, `observedOverlapCounts` and `observedFisherPvalues` is structured as a 2-level list where the outer components correspond to reference sets and the inner components to tests sets. As an example, `preservation$observed[[1]][[2]]` contains the density and connectivity preservation statistics for the preservation of set 1 modules in set 2, that is set 1 is the reference set and set 2 is the test set. `preservation$observed[[1]][[2]]` is a data frame in which each row corresponds to a module in the reference network 1 plus one row for the unassigned objects, and one row for a "module" that contains randomly sampled objects and



that represents a whole-network average. Each column corresponds to a statistic as indicated by the column name.

### Note

For large data sets, the permutation study may take a while (typically on the order of several hours). Use `verbose = 3` to get detailed progress report as the calculations advance.

### Author(s)

Rui Luo and Peter Langfelder

### References

Peter Langfelder, Rui Luo, Michael C. Oldham, and Steve Horvath, to appear

### See Also

Network construction and module detection functions in the WGCNA package such as [adjacency](#), [blockwiseModules](#); rudimentary cleaning in [goodSamplesGenesMS](#); the WGCNA implementation of correlation in [cor](#).

---

multiData.eigengeneSignificance

*Eigengene significance across multiple sets*

---

### Description

This function calculates eigengene significance and the associated significance statistics (p-values, q-values etc) across several data sets.

### Usage

```
multiData.eigengeneSignificance(
  multiData, multiTrait,
  moduleLabels, multiEigengenes = NULL,
  useModules = NULL,
  corAndPvalueFnc = corAndPvalue, corOptions = list(),
  corComponent = "cor",
  getQvalues = FALSE, setNames = NULL,
  excludeGrey = TRUE, greyLabel = ifelse(is.numeric(moduleLabels), 0, "grey"))
```

### Arguments

multiData	Expression data (or other data) in multi-set format (see <a href="#">checkSets</a> ). A vector of lists; in each list there must be a component named <code>data</code> whose content is a matrix or dataframe or array of dimension 2.
multiTrait	Trait or outcome data in multi-set format. Only one trait is allowed; consequently, the <code>data</code> component of each component list can be either a vector or a dataframe (matrix, array of dimension 2).
moduleLabels	Module labels: one label for each gene in <code>multiExpr</code> .

multiEigengenes	Optional eigengenes of modules specified in moduleLabels. If not given, will be calculated from multiExpr.
useModules	Optional specification of module labels to which the analysis should be restricted. This could be useful if there are many modules, most of which are not interesting. Note that the "grey" module cannot be used with useModules.
corAndPvalueFnc	Function that calculates associations between expression profiles and eigengenes. See details.
corOptions	List giving additional arguments to function corAndPvalueFnc. See details.
corComponent	Name of the component of output of corAndPvalueFnc that contains the actual correlation.
getQvalues	logical: should q-values (estimates of FDR) be calculated?
setNames	names for the input sets. If not given, will be taken from names(multiExpr). If those are NULL as well, the names will be "Set_1", "Set_2", ....
excludeGrey	logical: should the grey module be excluded from the kME tables? Since the grey module is typically not a real module, it makes little sense to report kME values for it.
greyLabel	label that labels the grey module.

### Details

This is a convenience function that calculates module eigengene significances (i.e., correlations of module eigengenes with a given trait) across all sets in a multi-set analysis. Also returned are p-values, Z scores, numbers of present (i.e., non-missing) observations for each significance, and optionally the q-values (false discovery rates) corresponding to the p-values.

The function corAndPvalueFnc is currently is expected to accept arguments *x* (gene expression profiles) and *y* (eigengene expression profiles). Any additional arguments can be passed via corOptions.

The function corAndPvalueFnc should return a list which at the least contains (1) a matrix of associations of genes and eigengenes (this component should have the name given by corComponent), and (2) a matrix of the corresponding p-values, named "p" or "p.value". Other components are optional but for full functionality should include (3) nObs giving the number of observations for each association (which is the number of samples less number of missing data - this can in principle vary from association to association), and (4) Z giving a Z static for each observation. If these are missing, nObs is calculated in the main function, and calculations using the Z statistic are skipped.

### Value

A list containing the following components. Each component is a matrix in which the rows correspond to module eigengenes and columns to data sets. Row and column names are set appropriately.

eigengeneSignificance	Module eigengene significance.
p.value	p-values (returned by corAndPvalueFnc).
q.value	q-values corresponding to the p-values above. Only returned in input getWvalues is TRUE.
Z	Z statistics (if returned by corAndPvalueFnc).
nObservations	Number of non-missing observations in each correlation/p-value.

**Author(s)**

Peter Langfelder

multiSetMEs

*Calculate module eigengenes.***Description**

Calculates module eigengenes for several sets.

**Usage**

```
multiSetMEs(exprData,
            colors,
            universalColors = NULL,
            useSets = NULL,
            useGenes = NULL,
            impute = TRUE,
            nPC = 1,
            align = "along average",
            excludeGrey = FALSE,
            grey = ifelse(is.null(universalColors), ifelse(is.numeric(colors), 0, "grey"),
                          ifelse(is.numeric(universalColors), 0, "grey")),
            subHubs = TRUE,
            trapErrors = FALSE,
            returnValidOnly = trapErrors,
            softPower = 6,
            verbose = 1, indent = 0)
```

**Arguments**

exprData	Expression data in a multi-set format (see <a href="#">checkSets</a> ). A vector of lists, with each list corresponding to one microarray dataset and expression data in the component data, that is <code>expr[[set]]\$data[sample, probe]</code> is the expression of probe <code>probe</code> in sample <code>sample</code> in dataset <code>set</code> . The number of samples can be different between the sets, but the probes must be the same.
colors	A matrix of dimensions (number of probes, number of sets) giving the module assignment of each gene in each set. The color "grey" is interpreted as unassigned.
universalColors	Alternative specification of module assignment. A single vector of length (number of probes) giving the module assignment of each gene in all sets (that is the modules are common to all sets). If given, takes precedence over <code>color</code> .
useSets	If calculations are requested in (a) selected set(s) only, the set(s) can be specified here. Defaults to all sets.
useGenes	Can be used to restrict calculation to a subset of genes (the same subset in all sets). If given, <code>validColors</code> in the returned list will only contain colors for the genes specified in <code>useGenes</code> .

impute	Logical. If TRUE, expression data will be checked for the presence of NA entries and if the latter are present, numerical data will be imputed, using function <code>impute.knn</code> and probes from the same module as the missing datum. The function <code>impute.knn</code> uses a fixed random seed giving repeatable results.
nPC	Number of principal components to be calculated. If only eigengenes are needed, it is best to set it to 1 (default). If variance explained is needed as well, use value NULL. This will cause all principal components to be computed, which is slower.
align	Controls whether eigengenes, whose orientation is undetermined, should be aligned with average expression ( <code>align = "along average"</code> , the default) or left as they are ( <code>align = ""</code> ). Any other value will trigger an error.
excludeGrey	Should the improper module consisting of 'grey' genes be excluded from the eigengenes?
grey	Value of <code>colors</code> or <code>universalColors</code> (whichever applies) designating the improper module. Note that if the appropriate <code>colors</code> argument is a factor of numbers, the default value will be incorrect.
subHubs	Controls whether hub genes should be substituted for missing eigengenes. If TRUE, each missing eigengene (i.e., eigengene whose calculation failed and the error was trapped) will be replaced by a weighted average of the most connected hub genes in the corresponding module. If this calculation fails, or if <code>subHubs==FALSE</code> , the value of <code>trapErrors</code> will determine whether the offending module will be removed or whether the function will issue an error and stop.
trapErrors	Controls handling of errors from that may arise when there are too many NA entries in expression data. If TRUE, errors from calling these functions will be trapped without abnormal exit. If FALSE, errors will cause the function to stop. Note, however, that <code>subHubs</code> takes precedence in the sense that if <code>subHubs==TRUE</code> and <code>trapErrors==FALSE</code> , an error will be issued only if both the principal component and the hubgene calculations have failed.
returnValidOnly	Boolean. Controls whether the returned data frames of module eigengenes contain columns corresponding only to modules whose eigengenes or hub genes could be calculated correctly in every set (TRUE), or whether the data frame should have columns for each of the input color labels (FALSE).
softPower	The power used in soft-thresholding the adjacency matrix. Only used when the hubgene approximation is necessary because the principal component calculation failed. It must be non-negative. The default value should only be changed if there is a clear indication that it leads to incorrect results.
verbose	Controls verbosity of printed progress messages. 0 means silent, up to (about) 5 the verbosity gradually increases.
indent	A single non-negative integer controlling indentation of printed messages. 0 means no indentation, each unit above that adds two spaces.

## Details

This function calls `moduleEigengenes` for each set in `exprData`.

Module eigengene is defined as the first principal component of the expression matrix of the corresponding module. The calculation may fail if the expression data has too many missing entries. Handling of such errors is controlled by the arguments `subHubs` and `trapErrors`. If `subHubs==TRUE`, errors in principal component calculation will be trapped and a substitute calculation of hubgenes will be attempted. If this fails as well, behaviour depends on `trapErrors`: if TRUE, the offending

module will be ignored and the return value will allow the user to remove the module from further analysis; if FALSE, the function will stop. If `universalColors` is given, any offending module will be removed from all sets (see `validMEs` in return value below).

From the user's point of view, setting `trapErrors=FALSE` ensures that if the function returns normally, there will be a valid eigengene (principal component or hubgene) for each of the input colors. If the user sets `trapErrors=TRUE`, all calculational (but not input) errors will be trapped, but the user should check the output (see below) to make sure all modules have a valid returned eigengene.

While the principal component calculation can fail even on relatively sound data (it does not take all that many "well-placed" NA to torpedo the calculation), it takes many more irregularities in the data for the hubgene calculation to fail. In fact such a failure signals there likely is something seriously wrong with the data.

## Value

A vector of lists similar in spirit to the input `exprData`. For each set there is a list with the following components:

<code>data</code>	Module eigengenes in a data frame, with each column corresponding to one eigengene. The columns are named by the corresponding color with an "ME" prepended, e.g., MEturquoise etc. Note that, when <code>trapErrors == TRUE</code> and <code>returnValidOnly==FALSE</code> , this data frame also contains entries corresponding to removed modules, if any. ( <code>validMEs</code> below indicates which eigengenes are valid and <code>allOK</code> whether all module eigengens were successfully calculated.)
<code>averageExpr</code>	If <code>align == "along average"</code> , a dataframe containing average normalized expression in each module. The columns are named by the corresponding color with an "AE" prepended, e.g., AEturquoise etc.
<code>varExplained</code>	A dataframe in which each column corresponds to a module, with the component <code>varExplained[PC, module]</code> giving the variance of module explained by the principal component no. PC. This is only accurate if all principal components have been computed (input <code>nPC = NULL</code> ). At most 5 principal components are recorded in this dataframe.
<code>nPC</code>	A copy of the input <code>nPC</code> .
<code>validMEs</code>	A boolean vector. Each component (corresponding to the columns in <code>data</code> ) is TRUE if the corresponding eigengene is valid, and FALSE if it is invalid. Valid eigengenes include both principal components and their hubgene approximations. When <code>returnValidOnly==FALSE</code> , by definition all returned eigengenes are valid and the entries of <code>validMEs</code> are all TRUE.
<code>validColors</code>	A copy of the input colors ( <code>universalColors</code> if set, otherwise <code>colors[, set]</code> ) with entries corresponding to invalid modules set to grey if given, otherwise 0 if the appropriate input colors are numeric and "grey" otherwise.
<code>allOK</code>	Boolean flag signalling whether all eigengenes have been calculated correctly, either as principal components or as the hubgene approximation. If <code>universalColors</code> is set, this flag signals whether all eigengenes are valid in all sets.
<code>allPC</code>	Boolean flag signalling whether all returned eigengenes are principal components. This flag (as well as the subsequent ones) is set independently for each set.
<code>isPC</code>	Boolean vector. Each component (corresponding to the columns in <code>eigengenes</code> ) is TRUE if the corresponding eigengene is the first principal component and FALSE if it is the hubgene approximation or is invalid.

isHub	Boolean vector. Each component (corresponding to the columns in eigengenes) is TRUE if the corresponding eigengene is the hubgene approximation and FALSE if it is the first principal component or is invalid.
validAEs	Boolean vector. Each component (corresponding to the columns in eigengenes) is TRUE if the corresponding module average expression is valid.
allAEOK	Boolean flag signalling whether all returned module average expressions contain valid data. Note that returnValidOnly==TRUE does not imply allAEOK==TRUE: some invalid average expressions may be returned if their corresponding eigengenes have been calculated correctly.

**Author(s)**

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**See Also**

[moduleEigengenes](#)

---

mutualInfoAdjacency     *Calculate weighted adjacency matrices based on mutual information*

---

**Description**

The function calculates different types of weighted adjacency matrices based on the mutual information between vectors (corresponding to the columns of the input data frame datE). The mutual information between pairs of vectors is divided by an upper bound so that the resulting normalized measure lies between 0 and 1.

**Usage**

```
mutualInfoAdjacency(
  datE,
  discretizeColumns = TRUE,
  entropyEstimationMethod = "MM",
  numberBins = NULL)
```

**Arguments**

**datE**                    datE is a data frame or matrix whose columns correspond to variables and whose rows correspond to measurements. For example, the columns may correspond to genes while the rows correspond to microarrays. The number of nodes in the mutual information network equals the number of columns of datE.

**discretizeColumns**     is a logical variable. If it is set to TRUE then the columns of datE will be discretized into a user-defined number of bins (see numberBins).

**entropyEstimationMethod**     takes a text string for specifying the entropy and mutual information estimation method. If entropyEstimationMethod="MM" then the Miller-Madow asymptotic bias corrected empirical estimator is used. If entropyEstimationMethod="ML" the maximum likelihood estimator (also known as plug-in or empirical estimator) is used. If entropyEstimationMethod="shrink", the shrinkage estimator

of a Dirichlet probability distribution is used. If `entropyEstimationMethod="SG"`, the Schurmann-Grassberger estimator of the entropy of a Dirichlet probability distribution is used.

`numberBins` is an integer larger than 0 which specifies how many bins are used for the discretization step. This argument is only relevant if `discretizeColumns` has been set to `TRUE`. By default `numberBins` is set to  $\sqrt{m}$  where  $m$  is the number of samples, i.e. the number of rows of `datE`. Thus the default is `numberBins=sqrt(nrow(datE))`.

## Details

The function inputs a data frame `datE` and outputs a list whose components correspond to different weighted network adjacency measures defined between the columns of `datE`. Make sure to install the following R packages `entropy`, `minet`, `infotheo` since the function `mutualInfoAdjacency` makes use of the `entropy` function from the R package `entropy` (Hausser and Strimmer 2008) and functions from the `minet` and `infotheo` package (Meyer et al 2008). A weighted network adjacency matrix is a symmetric matrix whose entries take on values between 0 and 1. Each weighted adjacency matrix contains scaled versions of the mutual information between the columns of the input data frame `datE`. We assume that `datE` contains numeric values which will be discretized unless the user chooses the option `discretizeColumns=FALSE`. The raw (unscaled) mutual information and entropy measures have units "nat", i.e. natural logarithms are used in their definition (base  $e=2.71..$ ). Several mutual information estimation methods have been proposed in the literature (reviewed in Hausser and Strimmer 2008, Meyer et al 2008). While mutual information networks allows one to detect non-linear relationships between the columns of `datE`, they may overfit the data if relatively few observations are available. Thus, if the number of rows of `datE` is smaller than say 200, it may be better to fit a correlation using the function `adjacency`.

## Value

The function outputs a list with the following components:

`Entropy` is a vector whose components report entropy estimates of each column of `datE`. The natural logarithm (base  $e$ ) is used in the definition. Using the notation from the Wikipedia entry ([http://en.wikipedia.org/wiki/Mutual\\_information](http://en.wikipedia.org/wiki/Mutual_information)), this vector contains the values  $H_x$  where  $x$  corresponds to a column in `datE`.

`MutualInformation` is a symmetric matrix whose entries contain the pairwise mutual information measures between the columns of `datE`. The diagonal of the matrix `MutualInformation` equals `Entropy`. In general, the entries of this matrix can be larger than 1, i.e. this is not an adjacency matrix. Using the notation from the Wikipedia entry, this matrix contains the mutual information estimates  $I(X;Y)$

`AdjacencySymmetricUncertainty` is a weighted adjacency matrix whose entries are based on the mutual information. Using the notation from the Wikipedia entry, this matrix contains the mutual information estimates  $AdjacencySymmetricUncertainty=2*I(X;Y)/(H(X)+H(Y))$ . Since  $I(X;X)=H(X)$ , the diagonal elements of `AdjacencySymmetricUncertainty` equal 1. In general the entries of this symmetric matrix `AdjacencySymmetricUncertainty` lie between 0 and 1.

`AdjacencyUniversalVersion1` is a weighted adjacency matrix that is a simple function of the `AdjacencySymmetricUncertainty`. Specifically,  $AdjacencyUniversalVersion1= AdjacencySymmetricUncertainty/(2- AdjacencySymmetricUncertainty)$ . Note that  $f(x)= x/(2-x)$  is a monotonically increasing function on the unit interval  $[0,1]$  whose values lie between 0 and 1. The reason why we call it the universal

adjacency is that  $dissUA=1-AdjacencyUniversalVersion1$  turns out to be a universal distance function, i.e. it satisfies the properties of a distance (including the triangle inequality) and it takes on a small value if any other distance measure takes on a small value (Kraskov et al 2003).

`AdjacencyUniversalVersion2`

is a weighted adjacency matrix for which  $dissUAVersion2=1-AdjacencyUniversalVersion2$  is also a universal distance measure. Using the notation from Wikipedia, the entries of the symmetric matrix `AdjacencyUniversalVersion2` are defined as follows  $AdjacencyUniversalVersion2=I(X;Y)/\max(H(X),H(Y))$ .

### Author(s)

Steve Horvath, Lin Song, Peter Langfelder

### References

Hausser J, Strimmer K (2008) Entropy inference and the James-Stein estimator, with application to nonlinear gene association networks. See <http://arxiv.org/abs/0811.3579>

Patrick E. Meyer, Frederic Lafitte, and Gianluca Bontempi. `minet`: A R/Bioconductor Package for Inferring Large Transcriptional Networks Using Mutual Information. *BMC Bioinformatics*, Vol 9, 2008

Kraskov A, Stoegbauer H, Andrzejak RG, Grassberger P (2003) Hierarchical Clustering Based on Mutual Information. *ArXiv q-bio/0311039*

### See Also

[adjacency](#)

### Examples

```
# Load requisite packages. These packages are considered "optional",
# so WGCNA does not load them automatically.

if (require(infotheo, quietly = TRUE) &&
    require(minet, quietly = TRUE) &&
    require(entropy, quietly = TRUE))
{
  # Example can be executed.
  #Simulate a data frame datE which contains 5 columns and 50 observations
  m=50
  x1=rnorm(m)
  r=.5; x2=r*x1+sqrt(1-r^2)*rnorm(m)
  r=.3; x3=r*(x1-.5)^2+sqrt(1-r^2)*rnorm(m)
  x4=rnorm(m)
  r=.3; x5=r*x4+sqrt(1-r^2)*rnorm(m)
  datE=data.frame(x1,x2,x3,x4,x5)

  #calculate entropy, mutual information matrix and weighted adjacency
  # matrices based on mutual information.
  MIadj=mutualInfoAdjacency(datE=datE)
} else
  printFlush(paste("Please install packages infotheo, minet and entropy",
                  "before running this example."));
```



**Description**

Basic statistical functions for handling missing values or NA.

In `log.na`, `sum.na`, `mean.na` and `var.na`, `quantile.na`, `length.na`, missing values are omitted from the calculation.

The function `cor.na` calls `cor` with the argument `use="pairwise.complete.obs"`.

The function `order.na` only handles vector arguments and not lists. However, it gives the option of omitting the NAs (`na.last=NA`), of placing the NAs at the start of the ordered vector (`na.last=F`) or at the end (`na.last=T`).

The function `scale.na` is a modified version of [scale](#) which allows NAs in the variance calculation.

If `scale = T`, the function `f` in `scale.na` uses `var.na` to perform the variance calculation. The function `prod.na` is similar to the [prod](#) function with `na.rm=TRUE`. This function returns the product of all the values present in its arguments, omitting any missing values.

**Author(s)**

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**See Also**

[log](#), [sum](#), [mean](#), [var](#), [cor](#), [order](#), [scale](#), [link{prod}](#).

---

nearestCentroidPredictor

*Nearest centroid predictor*

---

**Description**

Nearest centroid predictor for binary (i.e., two-outcome) data. Implements a whole host of options and improvements such as accounting for within-class heterogeneity using sample networks, various ways of feature selection and weighing etc.

**Usage**

```
nearestCentroidPredictor(
  # Input training and test data
  x, y,
  xtest = NULL,

  # Feature weights and selection criteria
  featureSignificance = NULL,
  assocFnc = "cor", assocOptions = "use = 'p'",
  assocCut.hi = NULL, assocCut.lo = NULL,
  nFeatures.hi = 10, nFeatures.lo = 10,
```

```

weighFeaturesByAssociation = 0,
scaleFeatureMean = TRUE, scaleFeatureVar = TRUE,

# Predictor options
centroidMethod = c("mean", "eigensample"),
simFnc = "cor", simOptions = "use = 'p'",
useQuantile = NULL,
sampleWeights = NULL,
weighSimByPrediction = 0,

# What should be returned
CVfold = 0, returnFactor = FALSE,

# General options
randomSeed = 12345,
verbose = 2, indent = 0)

```

### Arguments

x	Training features (predictive variables). Each column corresponds to a feature and each row to an observation.
y	The response variable. Can be a single vector or a matrix with arbitrary many columns. Number of rows (observations) must equal to the number of rows (observations) in x.
xtest	Optional test set data. A matrix of the same number of columns (i.e., features) as x. If test set data are not given, only the prediction on training data will be returned.
featureSignificance	Optional vector of feature significance for the response variable. If given, it is used for feature selection (see details). Should preferably be signed, that is features can have high negative significance.
assocFnc	Character string specifying the association function. The association function should behave roughly as <code>link{cor}</code> in that it takes two arguments (a matrix and a vector) plus options and returns the vector of associations between the columns of the matrix and the vector. The associations may be signed (i.e., negative or positive).
assocOptions	Character string specifying options to the association function.
assocCut.hi	Association (or featureSignificance) threshold for including features in the predictor. Features with association higher than <code>assocCut.hi</code> will be included. If not given, the threshold method will not be used; instead, a fixed number of features will be included as specified by <code>nFeatures.hi</code> and <code>nFeatures.lo</code> .
assocCut.lo	Association (or featureSignificance) threshold for including features in the predictor. Features with association lower than <code>assocCut.lo</code> will be included. If not given, defaults to <code>-assocCut.hi</code> . If <code>assocCut.hi</code> is NULL, the threshold method will not be used; instead, a fixed number of features will be included as specified by <code>nFeatures.hi</code> and <code>nFeatures.lo</code> .
nFeatures.hi	Number of highest-associated features (or features with highest featureSignificance) to include in the predictor. Only used if <code>assocCut.hi</code> is NULL.
nFeatures.lo	Number of lowest-associated features (or features with highest featureSignificance) to include in the predictor. Only used if <code>assocCut.hi</code> is NULL.

weighFeaturesByAssociation	(Optional) power to downweigh features that are less associated with the response. See details.
scaleFeatureMean	Logical: should the training features be scaled to mean zero? Unless there are good reasons not to scale, the features should be scaled.
scaleFeatureVar	Logical: should the training features be scaled to unit variance? Again, unless there are good reasons not to scale, the features should be scaled.
centroidMethod	One of "mean" and "eigensample", specifies how the centroid should be calculated. "mean" takes the mean across all samples (or all samples within a sample module, if sample networks are used), whereas "eigensample" calculates the first principal component of the feature matrix and uses that as the centroid.
simFnc	Character string giving the similarity function for measuring the similarity between test samples and centroids. This function should behave roughly like the function <code>cor</code> in that it takes two arguments (x, y) and calculates the pair-wise similarities between columns of x and y. For convenience, the value "dist" is treated specially: the Euclidean distance between the columns of x and y is calculated and its negative is returned (so that smallest distance corresponds to highest similarity). Since values of this function are only used for ranking centroids, its values are not restricted to be positive or within certain bounds.
simOptions	Character string specifying the options to the similarity function.
useQuantile	If non-NULL, the "nearest quantiloid" will be used instead of the nearest centroid. See details.
sampleWeights	Optional specification of sample weights. Useful for example if one wants to explore boosting.
weighSimByPrediction	(Optional) power to downweigh features that are not well predicted between training and test sets. See details.
CVfold	Non-negative integer specifying cross-validation. Zero means no cross-validation will be performed. values above zero specify the number of samples to be considered test data for each step of cross-validation.
returnFactor	Logical: should a factor be returned?
randomSeed	Integere specifying the seed for the random number generator. If NULL, the seed will not be set. See <a href="#">set.seed</a> .
verbose	Integer controlling how verbose the diagnostic messages should be. Zero means silent.
indent	Indentation for the diagnostic messages. Zero means no indentation, each unit adds two spaces.

## Details

Nearest centroid predictor works by forming a representative profile (centroid) across features for each class from the training data, then assigning each test sample to the class of the nearest representative profile. The representative profile can be formed either as mean or as the first principal component ("eigensample"; this choice is governed by the option `centroidMethod`).

When the number of features is large and only a small fraction is likely to be associated with the outcome, feature selection can be used to restrict the features that actually enter the centroid. Feature selection can be based either on their association with the outcome calculated from the training

data using `assocFnc`, or on user-supplied feature significance (e.g., derived from literature, argument `featureSignificance`). In either case, features can be selected by high and low association thresholds or by taking a fixed number of highest- and lowest-associated features.

As an alternative to centroids, the predictor can also assign test samples based on a given quantile of the distances from the training samples in each class (argument `useQuantile`). This may be advantageous if the samples in each class form irregular clusters. Note that setting `useQuantile=0` (i.e., using minimum distance in each class) essentially gives a nearest neighbor predictor: each test sample will be assigned to the class of its nearest training neighbor.

If features exhibit non-trivial correlations among themselves (such as, for example, in gene expression data), one can attempt to down-weight features that do not exhibit the same correlation in the test set. This is done by using essentially the same predictor to predict `_features_` from all other features in the test data (using the training data to train the feature predictor). Because test features are known, the prediction accuracy can be evaluated. If a feature is predicted badly (meaning the error in the test set is much larger than the error in the cross-validation prediction in training data), it may mean that its quality in the training or test data is low (for example, due to excessive noise or outliers). Such features can be downweighted using the argument `weighByPrediction`. The extra factor is  $\min(1, (\text{root mean square prediction error in test set})/(\text{root mean square cross-validation prediction error in the training data})^{\text{weighByPrediction}})$ , that is it is never bigger than 1.

Unless the features' mean and variance can be ascribed clear meaning, the (training) features should be scaled to mean 0 and variance 1 before the centroids are formed.

The function implements a basic option for removal of spurious effects in the training and test data, by removing a fixed number of leading principal components from the features. This sometimes leads to better prediction accuracy but should be used with caution.

If samples within each class are heterogenous, a single centroid may not represent each class well. This function can deal with within-class heterogeneity by clustering samples (separately in each class), then using a one representative (mean, eigensample) or quantile for each cluster in each class to assign test samples. Various similarity measures, specified by `adjFnc`, can be used to construct the sample network adjacency. Similarly, the user can specify a clustering function using `clusteringFnc`. The requirements on the clustering function are described in a separate section below.

## Value

A list with the following components:

`predicted`            The back-substitution prediction in the training set.

`predictedTest`      Prediction in the test set.

`featureSignificance`

A vector of feature significance calculated by `assocFnc` or a copy of the input `featureSignificance` if the latter is non-NULL.

`selectedFeatures`

A vector giving the indices of the features that were selected for the predictor.

`centroidProfile`

The representative profiles of each class (or cluster). Only returned in `useQuantile` is NULL.

`testSample2centroidSimilarities`

A matrix of calculated similarities between the test samples and class/cluster centroids.

`featureValidationWeights`

A vector of validation weights (see Details) for the selected features. If `weighFeaturesByValidation` is 0, a unit vector is used and returned.

CVpredicted	Cross-validation prediction on the training data. Present only if CVfold is non-zero.
sampleClusterLabels	A list with two components (one per class). Each component is a vector of sample cluster labels for samples in the class.

**Author(s)**

Peter Langfelder

**See Also**

[votingLinearPredictor](#)

---

nearestNeighborConnectivity

*Connectivity to a constant number of nearest neighbors*

---

**Description**

Given expression data and basic network parameters, the function calculates connectivity of each gene to a given number of nearest neighbors.

**Usage**

```
nearestNeighborConnectivity(datExpr,
  nNeighbors = 50, power = 6, type = "unsigned",
  corFnc = "cor", corOptions = "use = 'p'",
  blockSize = 1000,
  sampleLinks = NULL, nLinks = 5000, setSeed = 38457,
  verbose = 1, indent = 0)
```

**Arguments**

datExpr	a data frame containing expression data, with rows corresponding to samples and columns to genes. Missing values are allowed and will be ignored.
nNeighbors	number of nearest neighbors to use.
power	soft thresholding power for network construction. Should be a number greater than 1.
type	a character string encoding network type. Recognized values are (unique abbreviations of) "unsigned", "signed", and "signed hybrid".
corFnc	character string containing the name of the function to calculate correlation. Suggested functions include "cor" and "bicor".
corOptions	further argument to the correlation function.
blockSize	correlation calculations will be split into square blocks of this size, to prevent running out of memory for large gene sets.
sampleLinks	logical: should network connections be sampled (TRUE) or should all connections be used systematically (FALSE)?

nLinks	number of links to be sampled. Should be set such that nLinks * nNeighbors be several times larger than the number of genes.
setSeed	seed to be used for sampling, for repeatability. If a seed already exists, it is saved before the sampling starts and restored upon exit.
verbose	integer controlling the level of verbosity. 0 means silent.
indent	integer controlling indentation of output. Each unit above 0 adds two spaces.

### Details

Connectivity of gene *i* is the sum of adjacency strengths between gene *i* and other genes; in this case we take the nNeighbors nodes with the highest connection strength to gene *i*. The adjacency strengths are calculated by correlating the given expression data using the function supplied in corFNC and transforming them into adjacency according to the given network type and power.

### Value

A vector with one component for each gene containing the nearest neighbor connectivity.

### Author(s)

Peter Langfelder

### See Also

[adjacency](#), [softConnectivity](#)

---

nearestNeighborConnectivityMS

*Connectivity to a constant number of nearest neighbors across multiple data sets*

---

### Description

Given expression data from several sets and basic network parameters, the function calculates connectivity of each gene to a given number of nearest neighbors in each set.

### Usage

```
nearestNeighborConnectivityMS(multiExpr, nNeighbors = 50, power = 6,
  type = "unsigned", corFnc = "cor", corOptions = "use = 'p'",
  blockSize = 1000,
  sampleLinks = NULL, nLinks = 5000, setSeed = 36492,
  verbose = 1, indent = 0)
```

**Arguments**

multiExpr	expression data in multi-set format. A vector of lists, one list per set. In each list there must be a component named <code>data</code> whose content is a matrix or dataframe or array of dimension 2 containing the expression data. Rows correspond to samples and columns to genes (probes).
nNeighbors	number of nearest neighbors to use.
power	soft thresholding power for network construction. Should be a number greater than 1.
type	a character string encoding network type. Recognized values are (unique abbreviations of) "unsigned", "signed", and "signed hybrid".
corFnc	character string containing the name of the function to calculate correlation. Suggested functions include "cor" and "bicor".
corOptions	further argument to the correlation function.
blockSize	correlation calculations will be split into square blocks of this size, to prevent running out of memory for large gene sets.
sampleLinks	logical: should network connections be sampled (TRUE) or should all connections be used systematically (FALSE)?
nLinks	number of links to be sampled. Should be set such that $nLinks * nNeighbors$ be several times larger than the number of genes.
setSeed	seed to be used for sampling, for repeatability. If a seed already exists, it is saved before the sampling starts and restored after.
verbose	integer controlling the level of verbosity. 0 means silent.
indent	integer controlling indentation of output. Each unit above 0 adds two spaces.

**Details**

Connectivity of gene *i* is the sum of adjacency strengths between gene *i* and other genes; in this case we take the `nNeighbors` nodes with the highest connection strength to gene *i*. The adjacency strengths are calculated by correlating the given expression data using the function supplied in `corFNC` and transforming them into adjacency according to the given network type and power.

**Value**

A matrix in which columns correspond to sets and rows to genes; each entry contains the nearest neighbor connectivity of the corresponding gene.

**Author(s)**

Peter Langfelder

**See Also**

[adjacency](#), [softConnectivity](#), [nearestNeighborConnectivity](#)

networkConcepts

*Calculations of network concepts***Description**

This functions calculates various network concepts (topological properties, network indices) of a network calculated from expression data. See details for a detailed description.

**Usage**

```
networkConcepts(datExpr, power = 1, trait = NULL, networkType = "unsigned")
```

**Arguments**

datExpr	a data frame containing the expression data, with rows corresponding to samples and columns to genes (nodes).
power	soft thresholding power.
trait	optional specification of a sample trait. A vector of length equal the number of samples in datExpr.
networkType	network type. Recognized values are (unique abbreviations of) "unsigned", "signed", and "signed hybrid".

**Details**

This function computes various network concepts (also known as network statistics, topological properties, or network indices) for a weighted correlation network. The nodes of the weighted correlation network will be constructed between the columns (interpreted as nodes) of the input datExpr. If the option networkType="unsigned" then the adjacency between nodes  $i$  and  $j$  is defined as  $A[i, j] = \text{abs}(\text{cor}(\text{datExpr}[, i], \text{datExpr}[, j]))^{\text{power}}$ . In the following, we use the term gene and node interchangeably since these methods were originally developed for gene networks. The function computes the following 4 types of network concepts (introduced in Horvath and Dong 2008):

Type I: fundamental network concepts are defined as a function of the off-diagonal elements of an adjacency matrix  $A$  and/or a node significance measure  $GS$ . These network concepts can be defined for any network (not just correlation networks). The adjacency matrix of an unsigned weighted correlation network is given by  $A = \text{abs}(\text{cor}(\text{datExpr}, \text{use} = "p"))^{\text{power}}$  and the trait based gene significance measure is given by  $GS = \text{abs}(\text{cor}(\text{datExpr}, \text{trait}, \text{use} = "p"))^{\text{power}}$  where datExpr, trait, power are input parameters.

Type II: conformity-based network concepts are functions of the off-diagonal elements of the conformity based adjacency matrix  $A.CF = CF * t(CF)$  and/or the node significance measure. These network concepts are defined for any network for which a conformity vector can be defined. Details: For any adjacency matrix  $A$ , the conformity vector  $CF$  is calculated by requiring that  $A[i, j]$  is approximately equal to  $CF[i] * CF[j]$ . Using the conformity one can define the matrix  $A.CF = CF * t(CF)$  which is the outer product of the conformity vector with itself. In general,  $A.CF$  is not an adjacency matrix since its diagonal elements are different from 1. If the off-diagonal elements of  $A.CF$  are similar to those of  $A$  according to the Frobenius matrix norm, then  $A$  is approximately factorizable. To measure the factorizability of a network, one can calculate the Factorizability, which is a number between 0 and 1 (Dong and Horvath 2007). The conformity is defined using a monotonic, iterative algorithm that maximizes the factorizability measure.



Type III: approximate conformity based network concepts are functions of all elements of the conformity based adjacency matrix  $A$ .  $CF$  (including the diagonal) and/or the node significance measure  $GS$ . These network concepts are very useful for deriving relationships between network concepts in networks that are approximately factorizable.

Type IV: eigengene-based (also known as eigennode-based) network concepts are functions of the eigengene-based adjacency matrix  $A.E=ConformityE*t(ConformityE)$  (diagonal included) and/or the corresponding eigengene-based gene significance measure  $GSE$ . These network concepts can only be defined for correlation networks. Details: The columns (nodes) of  $datExpr$  can be summarized with the first principal component, which is referred to as *Eigengene* in coexpression network analysis. In general correlation networks, it is called *eigennode*. The eigengene-based conformity  $ConformityE[i]$  is defined as  $abs(cor(datE[, i], Eigengene))^power$  where the power corresponds to the power used for defining the weighted adjacency matrix  $A$ . The eigengene-based conformity can also be used to define an eigengene-based adjacency matrix  $A.E=ConformityE*t(ConformityE)$ . The eigengene based factorizability  $EF(datE)$  is a number between 0 and 1 that measures how well  $A.E$  approximates  $A$  when the power parameter equals 1.  $EF(datE)$  is defined with respect to the singular values of  $datExpr$ . For a trait based node significance measure  $GS=abs(cor(datE, trait))^power$ , one can also define an eigengene-based node significance measure  $GSE[i]=ConformityE[i]*EigengeneSignificance$  where the eigengene significance  $abs(cor(Eigengene, trait))^power$  is defined as power of the absolute value of the correlation between *eigengene* and *trait*. *Eigengene*-based network concepts are very useful for providing a geometric interpretation of network concepts and for deriving relationships between network concepts. For example, the hub gene significance measure and its eigengene-based analog have been used to characterize networks where highly connected hub genes are important with regard to a trait based gene significance measure (Horvath and Dong 2008).

## Value

A list with the following components:

Summary	a data frame whose rows report network concepts that only depend on the adjacency matrix. Density (mean adjacency), Centralization, Heterogeneity (coefficient of variation of the connectivity), Mean ClusterCoef, Mean Connectivity. The columns of the data frame report the 4 types of network concepts mentioned in the description: Fundamental concepts, eigengene-based concepts, conformity-based concepts, and approximate conformity-based concepts.
Size	reports the network size, i.e. the number of nodes, which equals the number of columns of the input data frame $datExpr$ .
Factorizability	a number between 0 and 1. The closer it is to 1, the better the off-diagonal elements of the conformity based network $A.CF$ approximate those of $A$ (according to the Frobenius norm).
Eigengene	the first principal component of the standardized columns of $datExpr$ . The number of components of this vector equals the number of rows of $datExpr$ .
VarExplained	the proportion of variance explained by the first principal component (the <i>Eigengene</i> ). It is numerically different from the eigengene based factorizability. While <i>VarExplained</i> is based on the squares of the singular values of $datExpr$ , the eigengene-based factorizability is based on fourth powers of the singular values.
Conformity	numerical vector giving the conformity. The number of components of the conformity vector equals the number of columns in $datExpr$ . The conformity is often highly correlated with the vector of node connectivities. The conformity is computed using an iterative algorithm for maximizing the factorizability measure. The algorithm and related network concepts are described in Dong and Horvath 2007.

ClusterCoef	a numerical vector that reports the cluster coefficient for each node. This fundamental network concept measures the cliquishness of each node.
Connectivity	a numerical vector that reports the connectivity (also known as degree) of each node. This fundamental network concept is also known as whole network connectivity. One can also define the scaled connectivity $K = \text{Connectivity} / \max(\text{Connectivity})$ which is used for computing the hub gene significance.
MAR	a numerical vector that reports the maximum adjacency ratio for each node. $\text{MAR}[i]$ equals 1 if all non-zero adjacencies between node $i$ and the remaining network nodes equal 1. This fundamental network concept is always 1 for nodes of an unweighted network. This is a useful measure for weighted networks since it allows one to determine whether a node has high connectivity because of many weak connections (small MAR) or because of strong (but few) connections (high MAR), see Horvath and Dong 2008.
ConformityE	a numerical vector that reports the eigengene based (aka eigenenode based) conformity for the correlation network. The number of components equals the number of columns of <code>datExpr</code> .
GS	a numerical vector that encodes the node (gene) significance. The $i$ -th component equals the node significance of the $i$ -th column of <code>datExpr</code> if a sample trait was supplied to the function (input trait). $\text{GS}[i] = \text{abs}(\text{cor}(\text{datE}[, i], \text{trait}, \text{use} = "p"))^{\text{power}}$
GSE	a numerical vector that reports the eigengene based gene significance measure. Its $i$ -th component is given by $\text{GSE}[i] = \text{ConformityE}[i] * \text{EigengeneSignificance}$ where the eigengene significance $\text{abs}(\text{cor}(\text{Eigengene}, \text{trait}))^{\text{power}}$ is defined as power of the absolute value of the correlation between eigengene and trait.
Significance	a data frame whose rows report network concepts that also depend on the trait based node significance measure. The rows correspond to network concepts and the columns correspond to the type of network concept (fundamental versus eigengene based). The first row of the data frame reports the network significance. The fundamental version of this network concepts is the average gene significance = $\text{mean}(\text{GS})$ . The eigengene based analog of this concept is defined as $\text{mean}(\text{GSE})$ . The second row reports the hub gene significance which is defined as slope of the intercept only regression model that regresses the gene significance on the scaled network connectivity $K$ . The third row reports the eigengene significance $\text{abs}(\text{cor}(\text{Eigengene}, \text{trait}))^{\text{power}}$ . More details can be found in Horvath and Dong (2008).

**Author(s)**

Jun Dong, Steve Horvath, Peter Langfelder

**References**

- Bin Zhang and Steve Horvath (2005) "A General Framework for Weighted Gene Co-Expression Network Analysis", *Statistical Applications in Genetics and Molecular Biology*: Vol. 4: No. 1, Article 17
- Dong J, Horvath S (2007) Understanding Network Concepts in Modules, *BMC Systems Biology* 2007, 1:24
- Horvath S, Dong J (2008) Geometric Interpretation of Gene Coexpression Network Analysis. *PLoS Comput Biol* 4(8): e1000117

**See Also**

[conformityBasedNetworkConcepts](#) for approximate conformity-based network concepts  
[fundamentalNetworkConcepts](#) for calculation of fundamental network concepts only.

---

networkScreening      *Identification of genes related to a trait*

---

**Description**

This function blends standard and network approaches to selecting genes (or variables in general) highly related to a given trait.

**Usage**

```
networkScreening(y, datME, datExpr,
                 corFnc = "cor", corOptions = "use = 'p'",
                 oddPower = 3,
                 blockSize = 1000,
                 minimumSampleSize = ..minNSamples,
                 addMEy = TRUE, removeDiag = FALSE,
                 weightESy = 0.5, getQValues = TRUE)
```

**Arguments**

y	clinical trait given as a numeric vector (one value per sample)
datME	data frame of module eigengenes
datExpr	data frame of expression data
corFnc	character string specifying the function to be used to calculate co-expression similarity. Defaults to Pearson correlation. Any function returning values between -1 and 1 can be used.
corOptions	character string specifying additional arguments to be passed to the function given by corFnc. Use "use = 'p', method = 'spearman'" to obtain Spearman correlation.
oddPower	odd integer used as a power to raise module memberships and significances
blockSize	block size to use for calculations with large data sets
minimumSampleSize	minimum acceptable number of samples. Defaults to the default minimum number of samples used throughout the WGCNA package, currently 4.
addMEy	logical: should the trait be used as an additional "module eigengene"?
removeDiag	logical: remove the diagonal?
weightESy	weight to use for the trait as an additional eigengene; should be between 0 and 1
getQValues	logical: should q-values be calculated?

**Details**

This function should be considered experimental. It takes into account both the "standard" and the network measures of gene importance for the trait.

**Value**

datout = data.frame(p.Weighted, q.Weighted, Cor.Weighted, Z.Weighted, p.Standard, q.Standard, Cor.Standard, Z.Standard) Data frame reporting the following quantities for each given gene:

p.Weighted	weighted p-value of association with the trait
q.Weighted	q-value (local FDR) calculated from p.Weighted
cor.Weighted	correlation of trait with gene expression weighted by a network term
Z.Weighted	Fisher Z score of the weighted correlation
p.Standard	standard Student p-value of association of the gene with the trait
q.Standard	q-value (local FDR) calculated from p.Standard
cor.Standard	correlation of gene with the trait
Z.Standard	Fisher Z score of the standard correlation

**Author(s)**

Steve Horvath

---

networkScreeningGS	<i>Network gene screening with an external gene significance measure</i>
--------------------	--

---

**Description**

This function blends standard and network approaches to selecting genes (or variables in general) with high gene significance

**Usage**

```
networkScreeningGS(
  datExpr,
  datME,
  GS,
  oddPower = 3,
  blockSize = 1000,
  minimumSampleSize = ..minNSamples,
  addGS = TRUE)
```

**Arguments**

datExpr	data frame of expression data
datME	data frame of module eigengenes
GS	numeric vector of gene significances
oddPower	odd integer used as a power to raise module memberships and significances
blockSize	block size to use for calculations with large data sets
minimumSampleSize	minimum acceptable number of samples. Defaults to the default minimum number of samples used throughout the WGCNA package, currently 4.
addGS	logical: should gene significances be added to the screening statistics?

**Details**

This function should be considered experimental. It takes into account both the "standard" and the network measures of gene importance for the trait.

**Value**

GS.Weighted	weighted gene significance
GS	copy of the input gene significances (only if addGS=TRUE)

**Author(s)**

Steve Horvath

**See Also**

[networkScreening](#), [automaticNetworkScreeningGS](#)

---

normalizeLabels	<i>Transform numerical labels into normal order.</i>
-----------------	--

---

**Description**

Transforms numerical labels into normal order, that is the largest group will be labeled 1, next largest 2 etc. Label 0 is optionally preserved.

**Usage**

```
normalizeLabels(labels, keepZero = TRUE)
```

**Arguments**

labels	Numerical labels.
keepZero	If TRUE (the default), labels 0 are preserved.

**Value**

A vector of the same length as input, containing the normalized labels.

**Author(s)**

Peter Langfelder, <Peter.Langfelder@gmail.com>

---

nPresent	<i>Number of present data entries.</i>
----------	--

---

**Description**

A simple sum of present entries in the argument.

**Usage**

```
nPresent(x)
```

**Arguments**

x                      data in which to count number of present entries.

**Value**

A single number giving the number of present entries in x.

**Author(s)**

Steve Horvath

---

nSets	<i>Number of sets in a multi-set variable</i>
-------	---

---

**Description**

A convenience function that returns the number of sets in a multi-set variable.

**Usage**

```
nSets(multiData, ...)
```

**Arguments**

multiData            vector of lists; in each list there must be a component named data whose content is a matrix or dataframe or array of dimension 2.

...                    Other arguments to function [checkSets](#).

**Value**

A single integer that equals the number of sets given in the input multiData.

**Author(s)**

Peter Langfelder

**See Also**

[checkSets](#)

---

numbers2colors	<i>Color representation for a numeric variable</i>
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---

## Description

The function creates a color representation for the given numeric input.

## Usage

```
numbers2colors(  
  x,  
  signed = NULL,  
  centered = signed,  
  lim = NULL,  
  commonLim = FALSE,  
  colors = if (signed) blueWhiteRed(100) else blueWhiteRed(100)[51:100],  
  naColor = "grey")
```

## Arguments

x	a vector or matrix of numbers. Missing values are allowed and will be assigned the color given in naColor. If a matrix, each column of the matrix is processed separately and the return value will be a matrix of colors.
signed	logical: should x be considered signed? If TRUE, the default setting is to use to use a palette that starts with green for the most negative values, continues with white for values around zero and turns red for positive values. If FALSE, the default palette ranges from white for minimum values to red for maximum values. If not given, the behaviour is controlled by values in x: if there are both positive and negative values, signed will be considered TRUE, otherwise FALSE.
centered	logical. If TRUE and signed==TRUE, numeric value zero will correspond to the middle of the color palette. If FALSE or signed==FALSE, the middle of the color palette will correspond to the average of the minimum and maximum value. If neither signed nor centered are given, centered will follow signed (see above).
lim	optional specification of limits, that is numeric values that should correspond to the first and last entry of colors.
commonLim	logical: should limits be calculated separately for each column of x, or should the limits be the same for all columns? Only applies if lim is NULL.
colors	color palette to represent the given numbers.
naColor	color to represent missing values in x.

## Details

Each column of x is processed individually, meaning that the color palette is adjusted individually for each column of x.

## Value

A vector or matrix (of the same dimensions as x) of colors.

**Author(s)**

Peter Langfelder

**See Also**[labels2colors](#) for color coding of ordinal labels.

---

`orderBranchesUsingHubGenes`*Optimize dendrogram using branch swaps and reflections.*

---

**Description**

This function takes as input the hierarchical clustering tree as well as a subset of genes in the network (generally corresponding to branches in the tree), then returns a semi-optimally ordered tree. The idea is to maximize the correlations between adjacent branches in the dendrogram, in as much as that is possible by adjusting the arbitrary positionings of the branches by swapping and reflecting branches.

**Usage**

```
orderBranchesUsingHubGenes(
  hierTOM,
  datExpr = NULL, colorh = NULL,
  type = "signed", adj = NULL, iter = NULL,
  useReflections = FALSE, allowNonoptimalSwaps = FALSE)
```

**Arguments**

<code>hierTOM</code>	A hierarchical clustering object (or gene tree) that is used to plot the dendrogram. For example, the output object from the function <code>hclust</code> or <code>flashClust</code> . Note that elements of <code>hierTOM\$order</code> MUST be named (for example, with the corresponding gene name).
<code>datExpr</code>	Gene expression data with rows as samples and columns as genes, or NULL if a pre-made adjacency is entered. Column names of <code>datExpr</code> must be a subset of gene names of <code>hierTOM\$order</code> .
<code>colorh</code>	The module assignments (color vectors) corresponding to the rows in <code>datExpr</code> , or NULL if a pre-made adjacency is entered.
<code>type</code>	What type of network is being entered. Common choices are "signed" (default) and "unsigned". With "signed" negative correlations count against, whereas with "unsigned" negative correlations are treated identically as positive correlations.
<code>adj</code>	Either NULL (default) or an adjacency (or any other square) matrix with rows and columns corresponding to a subset of the genes in <code>hierTOM\$order</code> . If entered, <code>datExpr</code> , <code>colorh</code> , and <code>type</code> are all ignored. Typically, this would be left blank but could include correlations between module eigengenes, with rows and columns renamed as genes in the corresponding modules, for example.
<code>iter</code>	The number of iterations to run the function in search of optimal branch ordering. The default is the square of the number of modules (or the square of the number of genes in the adjacency matrix).



useReflections If TRUE, both reflections and branch swapping will be used to optimize dendrogram. If FALSE (default) only branch swapping will be used.

allowNonoptimalSwaps

If TRUE, there is chance (that decreases with each iteration) of swapping / reflecting branches whether or not the new correlation between expression of genes in adjacent branches is better or worse. The idea (which has not been sufficiently tested), is that this would prevent the function from getting stuck at a local maxima of correlation. If FALSE (default), the swapping / reflection of branches only occurs if it results in a higher correlation between adjacent branches.

### Value

hierTOM A hierarchical clustering object with the hierTOM\$order variable properly adjusted, but all other variables identical as the heirTOM input.

changeLog A log of all of the changes that were made to the dendrogram, including what change was made, on what iteration, and the Old and New scores based on correlation. These scores have arbitrary units, but higher is better.

### Note

This function is very slow and is still in an *\*experimental\** function. We have not had problems with ~10 modules across ~5000 genes, although theoretically it should work for many more genes and modules, depending upon the speed of the computer running R. Please address any problems or suggestions to [jeremyinla@gmail.com](mailto:jeremyinla@gmail.com).

### Author(s)

Jeremy Miller

### Examples

```
## Example: first simulate some data.

MEturquoise = sample(1:100,50)
MEblue      = c(MEturquoise[1:25], sample(1:100,25))
MEbrown     = sample(1:100,50)
MEyellow    = sample(1:100,50)
MEgreen     = c(MEyellow[1:30], sample(1:100,20))
MERed       = c(MEbrown [1:20], sample(1:100,30))
ME          = data.frame(MEturquoise, MEblue, MEbrown, MEyellow, MEgreen, MERed)
dat1        = simulateDatExpr(ME,400,c(0.16,0.12,0.11,0.10,0.10,0.10,0.1), signed=TRUE)
TOM1        = TOMsimilarityFromExpr(dat1$datExpr, networkType="signed")
colnames(TOM1) <- rownames(TOM1) <- colnames(dat1$datExpr)
tree1       = flashClust(as.dist(1-TOM1),method="average")
colorh      = labels2colors(dat1$allLabels)

plotDendroAndColors(tree1,colorh,dendroLabels=FALSE)

## Reassign modules using the selectBranch and chooseOneHubInEachModule functions

datExpr = dat1$datExpr
hubs    = chooseOneHubInEachModule(datExpr, colorh)
colorh2 = rep("grey", length(colorh))
colorh2 [selectBranch(tree1,hubs["blue"],hubs["turquoise"])] = "blue"
```

```

colorh2 [selectBranch(tree1,hubs["turquoise"],hubs["blue"])] = "turquoise"
colorh2 [selectBranch(tree1,hubs["green"],hubs["yellow"])]   = "green"
colorh2 [selectBranch(tree1,hubs["yellow"],hubs["green"])]   = "yellow"
colorh2 [selectBranch(tree1,hubs["red"],hubs["brown"])]      = "red"
colorh2 [selectBranch(tree1,hubs["brown"],hubs["red"])]      = "brown"
plotDendroAndColors(tree1,cbind(colorh,colorh2),c("Old","New"),dendroLabels=FALSE)

## Now swap and reflect some branches, then optimize the order of the branches
# and output pdf with resulting images

## Not run:
pdf("DENDROGRAM_PLOTS.pdf",width=10,height=5)
plotDendroAndColors(tree1,colorh2,dendroLabels=FALSE,main="Starting Dendrogram")

tree1 = swapTwoBranches(tree1,hubs["red"],hubs["turquoise"])
plotDendroAndColors(tree1,colorh2,dendroLabels=FALSE,main="Swap blue/turquoise and red/brown")

tree1 = reflectBranch(tree1,hubs["blue"],hubs["green"])
plotDendroAndColors(tree1,colorh2,dendroLabels=FALSE,main="Reflect turquoise/blue")

# (This function will take a few minutes)
out = orderBranchesUsingHubGenes(tree1,datExpr,colorh2,useReflections=TRUE,iter=100)
tree1 = out$geneTree
plotDendroAndColors(tree1,colorh2,dendroLabels=FALSE,main="Semi-optimal branch order")

out$changeLog

dev.off()

## End(Not run)

```

---

orderMEs

*Put close eigenvectors next to each other*


---

## Description

Reorder given (eigen-)vectors such that similar ones (as measured by correlation) are next to each other.

## Usage

```

orderMEs(MEs, greyLast = TRUE,
         greyName = paste(moduleColor.getMEprefix(), "grey", sep=""),
         orderBy = 1, order = NULL,
         useSets = NULL, verbose = 0, indent = 0)

```

## Arguments

MEs            Module eigengenes in a multi-set format (see [checkSets](#)). A vector of lists, with each list corresponding to one dataset and the module eigengenes in the component data, that is `MEs[[set]]$data[sample, module]` is the expression of the eigengene of module `module` in sample `sample` in dataset `set`. The number of samples can be different between the sets, but the modules must be the same.

greyLast	Normally the color grey is reserved for unassigned genes; hence the grey module is not a proper module and it is conventional to put it last. If this is not desired, set the parameter to FALSE.
greyName	Name of the grey module eigengene.
orderBy	Specifies the set by which the eigengenes are to be ordered (in all other sets as well). Defaults to the first set in useSets (or the first set, if useSets is not given).
order	Allows the user to specify a custom ordering.
useSets	Allows the user to specify for which sets the eigengene ordering is to be performed.
verbose	Controls verbosity of printed progress messages. 0 means silent, nonzero verbose.
indent	A single non-negative integer controlling indentation of printed messages. 0 means no indentation, each unit above zero adds two spaces.

### Details

Ordering module eigengenes is useful for plotting purposes. For this function the order can be specified explicitly, or a set can be given in which the correlations of the eigengenes will determine the order. For the latter, a hierarchical dendrogram is calculated and the order given by the dendrogram is used for the eigengenes in all other sets.

### Value

A vector of lists of the same type as MEs containing the re-ordered eigengenes.

### Author(s)

Peter Langfelder, <Peter.Langfelder@gmail.com>

### See Also

[moduleEigengenes](#), [multiSetMEs](#), [consensusOrderMEs](#)

---

overlapTable	<i>Calculate overlap of modules</i>
--------------	-------------------------------------

---

### Description

The function calculates overlap counts and Fisher exact test p-values for the given two sets of module assignments.

### Usage

```
overlapTable(labels1, labels2)
```

### Arguments

labels1	a vector containing module labels.
labels2	a vector containing module labels to be compared to labels1.

**Value**

A list with the following components:

countTable	a matrix whose rows correspond to modules (unique labels) in labels1 and whose columns correspond to modules (unique labels) in labels2, giving the number of objects in the intersection of the two respective modules.
pTable	a matrix whose rows correspond to modules (unique labels) in labels1 and whose columns correspond to modules (unique labels) in labels2, giving Fisher's exact test significance p-values for the overlap of the two respective modules.

**Author(s)**

Peter Langfelder

**See Also**

[fisher.test](#), [matchLabels](#)

---

overlapTableUsingKME     *Determines significant overlap between modules in two networks based on kME tables.*

---

**Description**

Takes two sets of expression data (or kME tables) as input and returns a table listing the significant overlap between each module in each data set, as well as the actual genes in common for every module pair. Modules can be defined in several ways (generally involving kME) based on user input.

**Usage**

```
overlapTableUsingKME(
  dat1, dat2,
  colorh1, colorh2,
  MEs1 = NULL, MEs2 = NULL,
  name1 = "MM1", name2 = "MM2",
  cutoffMethod = "assigned", cutoff = 0.5,
  omitGrey = TRUE, datIsExpression = TRUE)
```

**Arguments**

dat1, dat2	Either expression data sets (with samples as rows and genes as columns) or module membership (kME) tables (with genes as rows and modules as columns). Function reads these inputs based on whether datIsExpression=TRUE or FALSE. <b>***Be sure that these inputs include relevant row and column names, or else the function will not work properly.***</b>
colorh1, colorh2	Color vector (module assignments) corresponding to the genes from dat1/2. This vector must be the same length as the Gene dimension from dat1/2.

MEs1, MEs2	If entered (default=NULL), these are the module eigengenes that will be used to form the kME tables. Rows are samples and columns are module assignments. Note that if datIsExpression=FALSE, these inputs are ignored.
name1, name2	The names of the two data sets being compared. These names affect the output parameters.
cutoffMethod	This variable is used to determine how modules are defined in each data set. Must be one of four options: (1) "assigned" -> use the module assignments in colorh (default); (2) "kME" -> any gene with kME > cutoff is in the module; (3) "numGenes" -> the top cutoff number of genes based on kME is in the module; and (4) "pvalue" -> any gene with correlation pvalue < cutoff is in the module (this includes both positively and negatively-correlated genes).
cutoff	For all cutoffMethods other than "assigned", this parameter is used as the described cutoff value.
omitGrey	If TRUE the grey modules (non-module genes) for both networks are not returned.
datIsExpression	If TRUE (default), dat1/2 is assumed to be expression data. If FALSE, dat1/2 is assumed to be a table of kME values.

**Value**

## PvaluesHypergeo

A table of p-values showing significance of module overlap based on the hypergeometric test. Note that these p-values are not corrected for multiple comparisons.

AllCommonGenes A character vector of all genes in common between the two data sets.

Genes<name1/2> A list of character vectors of all genes in each module in both data sets. All genes in the MOD module in data set MM1 could be found using "<outputVariableName>\$GenesMM1\$MM1\_MOD"

## OverlappingGenes

A list of character vectors of all genes for each between-set comparison from PvaluesHypergeo. All genes in MOD.A from MM1 that are also in MOD.B from MM2 could be found using "<outputVariableName>\$OverlappingGenes\$MM1\_MOD.A\_MM2"

**Author(s)**

Jeremy Miller

**See Also**

[overlapTable](#)

**Examples**

```
# Example: first generate simulated data.

set.seed(100)
ME.A = sample(1:100,50); ME.B = sample(1:100,50)
ME.C = sample(1:100,50); ME.D = sample(1:100,50)
ME.E = sample(1:100,50); ME.F = sample(1:100,50)
ME.G = sample(1:100,50); ME.H = sample(1:100,50)
ME1   = data.frame(ME.A, ME.B, ME.C, ME.D, ME.E)
```

```

ME2      = data.frame(ME.A, ME.C, ME.D, ME.E, ME.F, ME.G, ME.H)
simDat1 = simulateDatExpr(ME1,1000,c(0.2,0.1,0.08,0.05,0.04,0.3), signed=TRUE)
simDat2 = simulateDatExpr(ME2,1000,c(0.2,0.1,0.08,0.05,0.04,0.03,0.02,0.3),
                          signed=TRUE)

# Now run the function using assigned genes
results = overlapTableUsingKME(simDat1$datExpr, simDat2$datExpr,
                              labels2colors(simDat1$allLabels), labels2colors(simDat2$allLabels),
                              cutoffMethod="assigned")
results$PvaluesHypergeo

# Now run the function using a p-value cutoff, and inputting the original MEs
colnames(ME1) = standardColors(5); colnames(ME2) = standardColors(7)
results = overlapTableUsingKME(simDat1$datExpr, simDat2$datExpr,
                              labels2colors(simDat1$allLabels),
                              labels2colors(simDat2$allLabels),
                              ME1, ME2, cutoffMethod="pvalue", cutoff=0.05)
results$PvaluesHypergeo

# Check which genes are in common between the black modules from set 1 and
# the green module from set 2
results$OverlappingGenes$MM1_green_MM2_black

```

---

`pickHardThreshold`

*Analysis of scale free topology for hard-thresholding.*

---

## Description

Analysis of scale free topology for multiple hard thresholds. The aim is to help the user pick an appropriate threshold for network construction.

## Usage

```

pickHardThreshold(
  data,
  dataIsExpr,
  RsquaredCut = 0.85,
  cutVector = seq(0.1, 0.9, by = 0.05),
  moreNetworkConcepts = FALSE,
  removeFirst = FALSE, nBreaks = 10,
  corFnc = "cor", corOptions = "use = 'p'")

```

```

pickHardThreshold.fromSimilarity(
  similarity,
  RsquaredCut = 0.85,
  cutVector = seq(0.1, 0.9, by = 0.05),
  moreNetworkConcepts=FALSE,
  removeFirst = FALSE, nBreaks = 10)

```

## Arguments

`data` expression data in a matrix or data frame. Rows correspond to samples and columns to genes.

dataIsExpr	logical: should the data be interpreted as expression (or other numeric) data, or as a similarity matrix of network nodes?
similarity	similarity matrix: a symmetric matrix with entries between -1 and 1 and unit diagonal.
RsquaredCut	desired minimum scale free topology fitting index $R^2$ .
cutVector	a vector of hard threshold cuts for which the scale free topology fit indices are to be calculated.
moreNetworkConcepts	logical: should additional network concepts be calculated? If TRUE, the function will calculate how the network density, the network heterogeneity, and the network centralization depend on the power. For the definition of these additional network concepts, see Horvath and Dong (2008). PLoS Comp Biol.
removeFirst	should the first bin be removed from the connectivity histogram?
nBreaks	number of bins in connectivity histograms
corFnc	a character string giving the correlation function to be used in adjacency calculation.
corOptions	further options to the correlation function specified in corFnc.

### Details

The function calculates unsigned networks by thresholding the correlation matrix using thresholds given in cutVector. For each power the scale free topology fit index is calculated and returned along with other information on connectivity.

### Value

A list with the following components:

cutEstimate	estimate of an appropriate hard-thresholding cut: the lowest cut for which the scale free topology fit $R^2$ exceeds RsquaredCut. If $R^2$ is below RsquaredCut for all cuts, NA is returned.
fitIndices	a data frame containing the fit indices for scale free topology. The columns contain the hard threshold, adjusted $R^2$ for the linear fit, the linear coefficient, adjusted $R^2$ for a more complicated fit models, mean connectivity, median connectivity and maximum connectivity. If input moreNetworkConcepts is TRUE, 3 additional columns containing network density, centralization, and heterogeneity.

### Author(s)

Steve Horvath

### References

- Bin Zhang and Steve Horvath (2005) "A General Framework for Weighted Gene Co-Expression Network Analysis", Statistical Applications in Genetics and Molecular Biology: Vol. 4: No. 1, Article 17
- Horvath S, Dong J (2008) Geometric Interpretation of Gene Coexpression Network Analysis. PLoS Comput Biol 4(8): e1000117

**See Also**

[signumAdjacencyFunction](#)

---

pickSoftThreshold      *Analysis of scale free topology for soft-thresholding*

---

**Description**

Analysis of scale free topology for multiple soft thresholding powers. The aim is to help the user pick an appropriate soft-thresholding power for network construction.

**Usage**

```
pickSoftThreshold(
  data,
  dataIsExpr = TRUE,
  RsquaredCut = 0.85,
  powerVector = c(seq(1, 10, by = 1), seq(12, 20, by = 2)),
  removeFirst = FALSE, nBreaks = 10, blockSize = NULL,
  corFnc = cor, corOptions = list(use = 'p'),
  networkType = "unsigned",
  moreNetworkConcepts = FALSE,
  verbose = 0, indent = 0)

pickSoftThreshold.fromSimilarity(
  similarity,
  RsquaredCut = 0.85,
  powerVector = c(seq(1, 10, by = 1), seq(12, 20, by = 2)),
  removeFirst = FALSE, nBreaks = 10, blockSize = 1000,
  networkType = "unsigned",
  moreNetworkConcepts=FALSE,
  verbose = 0, indent = 0)
```

**Arguments**

data	expression data in a matrix or data frame. Rows correspond to samples and columns to genes.
dataIsExpr	logical: should the data be interpreted as expression (or other numeric) data, or as a similarity matrix of network nodes?
similarity	similarity matrix: a symmetric matrix with entries between -1 and 1 and unit diagonal.
RsquaredCut	desired minimum scale free topology fitting index $R^2$ .
powerVector	a vector of soft thresholding powers for which the scale free topology fit indices are to be calculated.
removeFirst	should the first bin be removed from the connectivity histogram?
nBreaks	number of bins in connectivity histograms



blockSize	block size into which the calculation of connectivity should be broken up. If not given, a suitable value will be calculated using function blockSize and printed if verbose>0. If R runs into memory problems, decrease this value.
corFnc	the correlation function to be used in adjacency calculation.
corOptions	a list giving further options to the correlation function specified in corFnc.
networkType	network type. Allowed values are (unique abbreviations of) "unsigned", "signed", "signed hybrid". See <a href="#">adjacency</a> .
moreNetworkConcepts	logical: should additional network concepts be calculated? If TRUE, the function will calculate how the network density, the network heterogeneity, and the network centralization depend on the power. For the definition of these additional network concepts, see Horvath and Dong (2008). <i>PLoS Comp Biol</i> .
verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.
indent	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

### Details

The function calculates weighted networks either by interpreting data directly as similarity, or first transforming it to similarity of the type specified by networkType. The weighted networks are obtained by raising the similarity to the powers given in powerVector. For each power the scale free topology fit index is calculated and returned along with other information on connectivity.

On systems with multiple cores or processors, the function pickSoftThreshold takes advantage of parallel processing if the function [enableWGCNAThreads](#) has been called to allow parallel processing and set up the parallel calculation back-end.

### Value

A list with the following components:

powerEstimate	estimate of an appropriate soft-thresholding power: the lowest power for which the scale free topology fit $R^2$ exceeds RsquaredCut. If $R^2$ is below RsquaredCut for all powers, NA is returned.
fitIndices	a data frame containing the fit indices for scale free topology. The columns contain the soft-thresholding power, adjusted $R^2$ for the linear fit, the linear coefficient, adjusted $R^2$ for a more complicated fit models, mean connectivity, median connectivity and maximum connectivity. If input moreNetworkConcepts is TRUE, 3 additional columns containing network density, centralization, and heterogeneity.

### Author(s)

Steve Horvath and Peter Langfelder

### References

- Bin Zhang and Steve Horvath (2005) "A General Framework for Weighted Gene Co-Expression Network Analysis", *Statistical Applications in Genetics and Molecular Biology*: Vol. 4: No. 1, Article 17
- Horvath S, Dong J (2008) Geometric Interpretation of Gene Coexpression Network Analysis. *PLoS Comput Biol* 4(8): e1000117

**See Also**

[adjacency](#), [softConnectivity](#)

---

plotClusterTreeSamples

*Annotated clustering dendrogram of microarray samples*

---

**Description**

This function plots an annotated clustering dendrogram of microarray samples.

**Usage**

```
plotClusterTreeSamples(
  datExpr,
  y = NULL,
  traitLabels = NULL,
  yLabels = NULL,
  main = if (is.null(y)) "Sample dendrogram" else
           "Sample dendrogram and trait indicator",
  setLayout = TRUE, autoColorHeight = TRUE, colorHeight = 0.3,
  dendroLabels = NULL,
  addGuide = FALSE, guideAll = TRUE,
  guideCount = NULL, guideHang = 0.2,
  cex.traitLabels = 0.8,
  cex.dendroLabels = 0.9,
  marAll = c(1, 5, 3, 1),
  saveMar = TRUE,
  abHeight = NULL, abCol = "red",
  ...)
```

**Arguments**

datExpr	a data frame containing expression data, with rows corresponding to samples and columns to genes. Missing values are allowed and will be ignored.
y	microarray sample trait. Either a vector with one entry per sample, or a matrix in which each column corresponds to a (different) trait and each row to a sample.
traitLabels	labels to be printed next to the color rows depicting sample traits. Defaults to column names of y.
yLabels	Optional labels to identify colors in the row identifying the sample classes. If given, must be of the same dimensions as y. Each label that occurs will be displayed once.
main	title for the plot.
setLayout	logical: should the plotting device be partitioned into a standard layout? If FALSE, the user is responsible for partitioning. The function expects two regions of the same width, the first one immediately above the second one.
autoColorHeight	logical: should the height of the color area below the dendrogram be automatically adjusted for the number of traits? Only effective if setLayout is TRUE.

colorHeight	Specifies the height of the color area under dendrogram as a fraction of the height of the dendrogram area. Only effective when autoColorHeight above is FALSE.
dendroLabels	dendrogram labels. Set to FALSE to disable dendrogram labels altogether; set to NULL to use row labels of datExpr.
addGuide	logical: should vertical "guide lines" be added to the dendrogram plot? The lines make it easier to identify color codes with individual samples.
guideAll	logical: add a guide line for every sample? Only effective for addGuide set TRUE.
guideCount	number of guide lines to be plotted. Only effective when addGuide is TRUE and guideAll is FALSE.
guideHang	fraction of the dendrogram height to leave between the top end of the guide line and the dendrogram merge height. If the guide lines overlap with dendrogram labels, increase guideHang to leave more space for the labels.
cex.traitLabels	character expansion factor for trait labels.
cex.dendroLabels	character expansion factor for dendrogram (sample) labels.
marAll	a 4-element vector giving the bottom, left, top and right margins around the combined plot. Note that this is not the same as setting the margins via a call to <code>par</code> , because the bottom margin of the dendrogram and the top margin of the color underneath are always zero.
saveMar	logical: save margins setting before starting the plot and restore on exit?
abHeight	optional specification of the height for a horizontal line in the dendrogram, see <a href="#">abline</a> .
abCol	color for plotting the horizontal line.
...	other graphical parameters to <a href="#">plot.hclust</a> .

### Details

The function generates an average linkage hierarchical clustering dendrogram (see [hclust](#)) of samples from the given expression data, using Eclidean distance of samples. The dendrogram is plotted together with color annotation for the samples.

The trait `y` must be numeric. If `y` is integer, the colors will correspond to values. If `y` is continuous, it will be dichotomized to two classes, below and above median.

### Value

None.

### Author(s)

Steve Horvath and Peter Langfelder

### See Also

[dist](#), [hclust](#), [plotDendroAndColors](#)

---

plotColorUnderTree      *Plot color rows under a dendrogram*

---

### Description

Plot color rows encoding information about objects in a clustering dendrogram, usually below the dendrogram.

### Usage

```
plotColorUnderTree(
  dendro,
  colors,
  rowLabels = NULL,
  rowWidths = NULL,
  rowText = NULL,
  rowTextAlignment = c("left", "center", "right"),
  rowTextIgnore = NULL,
  textPositions = NULL,
  addTextGuide = TRUE,
  cex.rowLabels = 1,
  cex.rowText = 0.8,
  ...)
```

### Arguments

dendro	A dendrogram such as returned by <a href="#">hclust</a> .
colors	Coloring of objects on the dendrogram. Either a vector (one color per object) or a matrix (can also be an array or a data frame) with each column giving one color per object. Each column will be plotted as a horizontal row of colors under the dendrogram.
rowLabels	Labels for the colorings given in colors. The labels will be printed to the left of the color rows in the plot. If the argument is given, it must be a vector of length equal to the number of columns in colors. If not given, names(colors) will be used if available. If not, sequential numbers starting from 1 will be used.
rowWidths	Optional specification of relative row widths for the color and text (if given) rows. Need not sum to 1.
rowText	Optional labels to identify colors in the color rows. If given, must be of the same dimensions as colors. Each label that occurs will be displayed once.
rowTextAlignment	Character string specifying whether the labels should be left-justified to the start of the largest block of each label, centered in the middle, or right-justified to the end of the largest block.
rowTextIgnore	Optional specifications of labels that should be ignored when displaying them using rowText above.
textPositions	optional numeric vector of the same length as the number of columns in rowText giving the color rows under which the text rows should appear.
addTextGuide	logical: should guide lines be added for the text rows (if given)?

`cex.rowLabels` Font size scale factor for the row labels. See [par](#).  
`cex.rowText` character expansion factor for text rows (if given).  
... Other parameters to be passed on to the plotting method (such as `main` for the main title etc).

### Details

It is often useful to plot dendrograms of objects together with additional information about the objects, for example module assignment (by color) that was obtained by cutting a hierarchical dendrogram or external color-coded measures such as gene significance. This function provides a way to do so. The calling code should section the screen into two (or more) parts, plot the dendrogram (via `plot(hclust)`) in the upper section and use this function to plot color annotation in the order corresponding to the dendrogram in the lower section.

### Value

None.

### Note

This function is identical to the function `plotHclustColors` in package `moduleColor`.

### Author(s)

Steve Horvath <[SHorvath@mednet.ucla.edu](mailto:SHorvath@mednet.ucla.edu)> and Peter Langfelder <[Peter.Langfelder@gmail.com](mailto:Peter.Langfelder@gmail.com)>

### See Also

[cutreeDynamic](#) for module detection in a dendrogram;

[plotDendroAndColors](#) for automated plotting of dendrograms and colors in one step.

---

plotCor

*Red and Green Color Image of Correlation Matrix*

---

### Description

This function produces a red and green color image of a correlation matrix using an RGB color specification. Increasingly positive correlations are represented with reds of increasing intensity, and increasingly negative correlations are represented with greens of increasing intensity.

### Usage

```
plotCor(x, new=FALSE, nrgcols=50, labels=FALSE, labcols=1, title="", ...)
```

**Arguments**

<code>x</code>	a matrix of numerical values.
<code>new</code>	If <code>new=F</code> , <code>x</code> must already be a correlation matrix. If <code>new=T</code> , the correlation matrix for the columns of <code>x</code> is computed and displayed in the image.
<code>nrgcols</code>	the number of colors ( $\geq 1$ ) to be used in the red and green palette.
<code>labels</code>	vector of character strings to be placed at the tickpoints, labels for the columns of <code>x</code> .
<code>labcols</code>	colors to be used for the labels of the columns of <code>x</code> . <code>labcols</code> can have either length 1, in which case all the labels are displayed using the same color, or the same length as <code>labels</code> , in which case a color is specified for the label of each column of <code>x</code> .
<code>title</code>	character string, overall title for the plot.
<code>...</code>	graphical parameters may also be supplied as arguments to the function (see <a href="#">par</a> ). For comparison purposes, it is good to set <code>zlim=c(-1,1)</code> .

**Author(s)**

Sandrine Dudoit, <sandrine@stat.berkeley.edu>

**See Also**

[plotMat,rgcolors.func](#), [cor.na](#), [cor](#), [image](#), [rgb](#).

---

`plotDendroAndColors`     *Dendrogram plot with color annotation of objects*

---

**Description**

This function plots a hierarchical clustering dendrogram and color annotation(s) of objects in the dendrogram underneath.

**Usage**

```
plotDendroAndColors(
  dendro,
  colors,
  groupLabels = NULL,
  rowText = NULL,
  rowTextAlignment = c("left", "center", "right"),
  rowTextIgnore = NULL,
  textPositions = NULL,
  setLayout = TRUE,
  autoColorHeight = TRUE,
  colorHeight = 0.2,
  rowWidths = NULL,
  dendroLabels = NULL,
  addGuide = FALSE, guideAll = FALSE,
  guideCount = 50, guideHang = 0.2,
  addTextGuide = FALSE,
```

```

cex.colorLabels = 0.8, cex.dendroLabels = 0.9,
cex.rowText = 0.8,
marAll = c(1, 5, 3, 1), saveMar = TRUE,
abHeight = NULL, abCol = "red", ...)

```

## Arguments

dendro	a hierarchical clustering dendrogram such as one produced by <a href="#">hclust</a> .
colors	Coloring of objects on the dendrogram. Either a vector (one color per object) or a matrix (can also be an array or a data frame) with each column giving one color per object. Each column will be plotted as a horizontal row of colors under the dendrogram.
groupLabels	Labels for the colorings given in colors. The labels will be printed to the left of the color rows in the plot. If the argument is given, it must be a vector of length equal to the number of columns in colors. If not given, names(colors) will be used if available. If not, sequential numbers starting from 1 will be used.
rowText	Optional labels to identify colors in the color rows. If given, must be either the same dimensions as colors or must have the same number of rows and textPositions must be used to specify which columns of colors each column of rowText corresponds to. Each label that occurs will be displayed once, under the largest continuous block of the corresponding colors.
rowTextAlignment	Character string specifying whether the labels should be left-justified to the start of the largest block of each label, centered in the middle, or right-justified to the end of the largest block.
rowTextIgnore	Optional specifications of labels that should be ignored when displaying them using rowText above.
textPositions	optional numeric vector of the same length as the number of columns in rowText giving the color rows under which the text rows should appear.
setLayout	logical: should the plotting device be partitioned into a standard layout? If FALSE, the user is responsible for partitioning. The function expects two regions of the same width, the first one immediately above the second one.
autoColorHeight	logical: should the height of the color area below the dendrogram be automatically adjusted for the number of traits? Only effective if setLayout is TRUE.
colorHeight	specifies the height of the color area under dendrogram as a fraction of the height of the dendrogram area. Only effective when autoColorHeight above is FALSE.
rowWidths	optional specification of relative row widths for the color and text (if given) rows. Need not sum to 1.
dendroLabels	dendrogram labels. Set to FALSE to disable dendrogram labels altogether; set to NULL to use row labels of datExpr.
addGuide	logical: should vertical "guide lines" be added to the dendrogram plot? The lines make it easier to identify color codes with individual samples.
guideAll	logical: add a guide line for every sample? Only effective for addGuide set TRUE.
guideCount	number of guide lines to be plotted. Only effective when addGuide is TRUE and guideAll is FALSE.

guideHang	fraction of the dendrogram height to leave between the top end of the guide line and the dendrogram merge height. If the guide lines overlap with dendrogram labels, increase guideHang to leave more space for the labels.
addTextGuide	logical: should guide lines be added for the text rows (if given)?
cex.colorLabels	character expansion factor for trait labels.
cex.dendroLabels	character expansion factor for dendrogram (sample) labels.
cex.rowText	character expansion factor for text rows (if given).
marAll	a vector of length 4 giving the bottom, left, top and right margins of the combined plot. There is no margin between the dendrogram and the color plot underneath.
saveMar	logical: save margins setting before starting the plot and restore on exit?
abHeight	optional specification of the height for a horizontal line in the dendrogram, see <a href="#">abline</a> .
abCol	color for plotting the horizontal line.
...	other graphical parameters to <a href="#">plot.hclust</a> .

### Details

The function splits the plotting device into two regions, plots the given dendrogram in the upper region, then plots color rows in the region below the dendrogram.

### Value

None.

### Author(s)

Peter Langfelder

### See Also

[plotColorUnderTree](#)

---

plotEigengeneNetworks *Eigengene network plot*

---

### Description

This function plots dendrogram and eigengene representations of (consensus) eigengenes networks. In the case of consensus eigengene networks the function also plots pairwise preservation measures between consensus networks in different sets.



**Usage**

```
plotEigengeneNetworks(
  multiME,
  setLabels,
  letterSubPlots = FALSE, Letters = NULL,
  excludeGrey = TRUE, greyLabel = "grey",
  plotDendrograms = TRUE, plotHeatmaps = TRUE,
  setMargins = TRUE, marDendro = NULL, marHeatmap = NULL,
  colorLabels = TRUE, signed = TRUE,
  heatmapColors = NULL,
  plotAdjacency = TRUE,
  printAdjacency = FALSE, cex.adjacency = 0.9,
  coloredBarplot = TRUE, barplotMeans = TRUE, barplotErrors = FALSE,
  plotPreservation = "standard",
  zlimPreservation = c(0, 1),
  printPreservation = FALSE, cex.preservation = 0.9,
  ...)
```

**Arguments**

multiME	either a single data frame containing the module eigengenes, or module eigengenes in the multi-set format (see <a href="#">checkSets</a> ). The multi-set format is a vector of lists, one per set. Each set must contain a component data whose rows correspond to samples and columns to eigengenes.
setLabels	A vector of character strings that label sets in multiME.
letterSubPlots	logical: should subplots be lettered?
Letters	optional specification of a sequence of letters for lettering. Defaults to "ABCD"...
excludeGrey	logical: should the grey module eigengene be excluded from the plots?
greyLabel	label for the grey module. Usually either "grey" or the number 0.
plotDendrograms	logical: should eigengene dendrograms be plotted?
plotHeatmaps	logical: should eigengene network heatmaps be plotted?
setMargins	logical: should margins be set? See <a href="#">par</a> .
marDendro	a vector of length 4 giving the margin setting for dendrogram plots. See <a href="#">par</a> . If setMargins is TRUE and marDendro is not given, the function will provide reasonable default values.
marHeatmap	a vector of length 4 giving the margin setting for heatmap plots. See <a href="#">par</a> . If setMargins is TRUE and marDendro is not given, the function will provide reasonable default values.
colorLabels	logical: should module eigengene names be interpreted as color names and the colors used to label heatmap plots and barplots?
signed	logical: should eigengene networks be constructed as signed?
heatmapColors	color palette for heatmaps. Defaults to <a href="#">heat.colors</a> when signed is FALSE, and to <a href="#">redWhiteGreen</a> when signed is TRUE.
plotAdjacency	logical: should module eigengene heatmaps plot adjacency (ranging from 0 to 1), or correlation (ranging from -1 to 1)?
printAdjacency	logical: should the numerical values be printed into the adjacency or correlation heatmap?

<code>cex.adjacency</code>	character expansion factor for printing of numerical values into the adjacency or correlation heatmap
<code>coloredBarplot</code>	logical: should the barplot of eigengene adjacency preservation distinguish individual contributions by color? This is possible only if <code>colorLabels</code> is TRUE and module eigengene names encode valid colors.
<code>barplotMeans</code>	logical: plot mean preservation in the barplot? This option effectively rescales the preservation by the number of eigengenes in the network. If means are plotted, the barplot is not colored.
<code>barplotErrors</code>	logical: should standard errors of the mean preservation be plotted?
<code>plotPreservation</code>	a character string specifying which type of preservation measure to plot. Allowed values are (unique abbreviations of) "standard", "hyperbolic", "both".
<code>zlimPreservation</code>	a vector of length 2 giving the value limits for the preservation heatmaps.
<code>printPreservation</code>	logical: should preservation values be printed within the heatmap?
<code>cex.preservation</code>	character expansion factor for preservation display.
<code>...</code>	other graphical arguments to function <a href="#">labeledHeatmap</a> .

## Details

Consensus eigengene networks consist of a fixed set of eigengenes "expressed" in several different sets. Network connection strengths are given by eigengene correlations. This function aims to visualize the networks as well as their similarities and differences across sets.

The function partitions the screen appropriately and plots eigengene dendrograms in the top row, then a square matrix of plots: heatmap plots of eigengene networks in each set on the diagonal, heatmap plots of pairwise preservation networks below the diagonal, and barplots of aggregate network preservation of individual eigengenes above the diagonal. A preservation plot or barplot in the row *i* and column *j* of the square matrix represents the preservation between sets *i* and *j*.

Individual eigengenes are labeled by their name in the dendrograms; in the heatmaps and barplots they can optionally be labeled by color squares. For compatibility with other functions, the color labels are encoded in the eigengene names by prefixing the color with two letters, such as "MEturquoise".

Two types of network preservation can be plotted: the "standard" is simply the difference between adjacencies in the two compared sets. The "hyperbolic" difference de-emphasizes the preservation of low adjacencies. When "both" is specified, standard preservation is plotted in the lower triangle and hyperbolic in the upper triangle of each preservation heatmap.

If the eigengenes are labeled by color, the bars in the barplot can be split into segments representing the contribution of each eigengene and labeled by the contribution. For example, a yellow segment in a bar labeled by a turquoise square represents the preservation of the adjacency between the yellow and turquoise eigengenes in the two networks compared by the barplot.

For large numbers of eigengenes and/or sets, it may be difficult to get a meaningful plot fit a standard computer screen. In such cases we recommend using a device such as [postscript](#) or [pdf](#) where the user can specify large dimensions; such plots can be conveniently viewed in standard pdf or postscript viewers.

## Value

None.

**Author(s)**

Peter Langfelder

**References**

For theory and applications of consensus eigengene networks, see

Langfelder P, Horvath S (2007) Eigengene networks for studying the relationships between co-expression modules. *BMC Systems Biology* 2007, 1:54

**See Also**

[labeledHeatmap](#), [labeledBarplot](#) for annotated heatmaps and barplots;

[hclust](#) for hierarchical clustering and dendrogram plots

---

 plotMat

*Red and Green Color Image of Data Matrix*


---

**Description**

This function produces a red and green color image of a data matrix using an RGB color specification. Larger entries are represented with reds of increasing intensity, and smaller entries are represented with greens of increasing intensity.

**Usage**

```
plotMat(x, nrgcols=50, rlabels=FALSE, clabels=FALSE, rcols=1, ccols=1, title="",...)
```

**Arguments**

<code>x</code>	a matrix of numbers.
<code>nrgcols</code>	the number of colors ( $\geq 1$ ) to be used in the red and green palette.
<code>rlabels</code>	vector of character strings to be placed at the row tickpoints, labels for the rows of <code>x</code> .
<code>clabels</code>	vector of character strings to be placed at the column tickpoints, labels for the columns of <code>x</code> .
<code>rcols</code>	colors to be used for the labels of the rows of <code>x</code> . <code>rcols</code> can have either length 1, in which case all the labels are displayed using the same color, or the same length as <code>rlabels</code> , in which case a color is specified for the label of each row of <code>x</code> .
<code>ccols</code>	colors to be used for the labels of the columns of <code>x</code> . <code>ccols</code> can have either length 1, in which case all the labels are displayed using the same color, or the same length as <code>clabels</code> , in which case a color is specified for the label of each column of <code>x</code> .
<code>title</code>	character string, overall title for the plot.
<code>...</code>	graphical parameters may also be supplied as arguments to the function (see <a href="#">par</a> ). E.g. <code>zlim=c(-3,3)</code>

**Author(s)**

Sandrine Dudoit, <sandrine@stat.berkeley.edu>

**See Also**

[plotCor](#), [rgcolors.func](#), [cor.na](#), [cor](#), [image](#), [rgb](#).

---

plotMEpairs

*Pairwise scatterplots of eigengenes*

---

**Description**

The function produces a matrix of plots containing pairwise scatterplots of given eigengenes, the distribution of their values and their pairwise correlations.

**Usage**

```
plotMEpairs(  
  datME,  
  y = NULL,  
  main = "Relationship between module eigengenes",  
  clusterMEs = TRUE,  
  ...)
```

**Arguments**

datME	a data frame containing expression data, with rows corresponding to samples and columns to genes. Missing values are allowed and will be ignored.
y	optional microarray sample trait vector. Will be treated as an additional eigengene.
main	main title for the plot.
clusterMEs	logical: should the module eigengenes be ordered by their dendrogram?
...	additional graphical parameters to the function <a href="#">pairs</a>

**Details**

The function produces an NxN matrix of plots, where N is the number of eigengenes. In the upper triangle it plots pairwise scatterplots of module eigengenes (plus the trait y, if given). On the diagonal it plots histograms of sample values for each eigengene. Below the diagonal, it displays the pairwise correlations of the eigengenes.

**Value**

None.

**Author(s)**

Steve Horvath

**See Also**

[pairs](#)

---

`plotModuleSignificance`*Barplot of module significance*

---

**Description**

Plot a barplot of gene significance.

**Usage**

```
plotModuleSignificance(  
  geneSignificance,  
  colors,  
  boxplot = FALSE,  
  main = "Gene significance across modules",  
  ylab = "Gene Significance", ...)
```

**Arguments**

<code>geneSignificance</code>	a numeric vector giving gene significances.
<code>colors</code>	a character vector specifying module assignment for the genes whose significance is given in <code>geneSignificance</code> . The modules should be labeled by colors.
<code>boxplot</code>	logical: should a boxplot be produced instead of a barplot?
<code>main</code>	main title for the plot.
<code>ylab</code>	y axis label for the plot.
<code>...</code>	other graphical parameters to <a href="#">plot</a> .

**Details**

Given individual gene significances and their module assignment, the function calculates the module significance for each module as the average gene significance of the genes within the module. The result is plotted in a barplot or boxplot form. Each bar or box is labeled by the corresponding module color.

**Value**

None.

**Author(s)**

Steve Horvath

**References**

Bin Zhang and Steve Horvath (2005) "A General Framework for Weighted Gene Co-Expression Network Analysis", *Statistical Applications in Genetics and Molecular Biology*: Vol. 4: No. 1, Article 17

Dong J, Horvath S (2007) Understanding Network Concepts in Modules, *BMC Systems Biology* 2007, 1:24

**See Also**[barplot](#), [boxplot](#)

---

plotNetworkHeatmap	<i>Network heatmap plot</i>
--------------------	-----------------------------

---

**Description**

Network heatmap plot.

**Usage**

```
plotNetworkHeatmap(  
  datExpr,  
  plotGenes,  
  useTOM = TRUE,  
  power = 6,  
  networkType = "unsigned",  
  main = "Heatmap of the network")
```

**Arguments**

datExpr	a data frame containing expression data, with rows corresponding to samples and columns to genes. Missing values are allowed and will be ignored.
plotGenes	a character vector giving the names of genes to be included in the plot. The names will be matched against names(datExpr).
useTOM	logical: should TOM be plotted (TRUE), or correlation-based adjacency (FALSE)?
power	soft-thresholding power for network construction.
networkType	a character string giving the network type. Recognized values are (unique abbreviations of) "unsigned", "signed", and "signed hybrid".
main	main title for the plot.

**Details**

The function constructs a network from the given expression data (selected by plotGenes) using the soft-thresholding procedure, optionally calculates Topological Overlap (TOM) and plots a heatmap of the network.

Note that all network calculations are done in one block and may fail due to memory allocation issues for large numbers of genes.

**Value**

None.

**Author(s)**

Steve Horvath

## References

Bin Zhang and Steve Horvath (2005) "A General Framework for Weighted Gene Co-Expression Network Analysis", *Statistical Applications in Genetics and Molecular Biology*: Vol. 4: No. 1, Article 17

## See Also

[adjacency](#), [TOMsimilarity](#)

---

populationMeansInAdmixture

*Estimate the population-specific mean values in an admixed population.*

---

## Description

Uses the expression values from an admixed population and estimates of the proportions of sub-populations to estimate the population specific mean values. For example, this function can be used to estimate the cell type specific mean gene expression values based on expression values from a mixture of cells. The method is described in Shen-Orr et al (2010) where it was used to estimate cell type specific gene expression levels based on a mixture sample.

## Usage

```
populationMeansInAdmixture(
  datProportions, datE.Admixture,
  scaleProportionsTo1 = TRUE,
  scaleProportionsInCelltype = TRUE,
  setMissingProportionsToZero = FALSE)
```

## Arguments

- datProportions** a matrix of non-negative numbers (ideally proportions) where the rows correspond to the samples (rows of `datE.Admixture`) and the columns correspond to the sub-populations of the mixture. The function calculates a mean expression value for each column of `datProportions`. Negative entries in `datProportions` lead to an error message. But the rows of `datProportions` do not have to sum to 1, see the argument `scaleProportionsTo1`.
- datE.Admixture** a matrix of numbers. The rows correspond to samples (mixtures of populations). The columns contain the variables (e.g. genes) for which the means should be estimated.
- scaleProportionsTo1** logical. If set to `TRUE` (default) then the proportions in each row of `datProportions` are scaled so that they sum to 1, i.e. `datProportions[i,]=datProportions[i,]/max(datProportions[i,])`. In general, we recommend to set it to `TRUE`.
- scaleProportionsInCelltype** logical. If set to `TRUE` (default) then the proportions in each cell types are recalc and make the mean to 0.
- setMissingProportionsToZero** logical. Default is `FALSE`. If set to `TRUE` then it sets missing values in `datProportions` to zero.

**Details**

The function outputs a matrix of coefficients resulting from fitting a regression model. If the proportions sum to 1, then  $i$ -th row of the output matrix reports the coefficients of the following model  $\text{lm}(\text{datE.Admixture}[,i] \sim -1, \text{data}=\text{datProportions})$ . Aside, the minus 1 in the formula indicates that no intercept term will be fit. Under certain assumptions, the coefficients can be interpreted as the mean expression values in the sub-populations (Shen-Orr 2010).

**Value**

a numeric matrix whose rows correspond to the columns of `datE.Admixture` (e.g. to genes) and whose columns correspond to the columns of `datProportions` (e.g. sub populations or cell types).

**Note**

This can be considered a wrapper of the `lm` function.

**Author(s)**

Steve Horvath, Chaochao Cai

**References**

Shen-Orr SS, Tibshirani R, Khatri P, Bodian DL, Staedtler F, Perry NM, Hastie T, Sarwal MM, Davis MM, Butte AJ (2010) Cell type-specific gene expression differences in complex tissues. *Nature Methods*, vol 7 no.4

**Examples**

```
set.seed(1)
# this is the number of complex (mixed) tissue samples, e.g. arrays
m=10
# true count data (e.g. pure cells in the mixed sample)
datTrueCounts=as.matrix(data.frame(TrueCount1=rpois(m,lambda=16),
TrueCount2=rpois(m,lambda=8),TrueCount3=rpois(m,lambda=4),
TrueCount4=rpois(m,lambda=2)))
no.pure=dim(datTrueCounts)[[2]]

# now we transform the counts into proportions
divideBySum=function(x) t(x)/sum(x)
datProportions= t(apply(datTrueCounts,1,divideBySum))
dimnames(datProportions)[[2]]=paste("TrueProp",1:dim(datTrueCounts)[[2]],sep=".")

# number of genes that are highly expressed in each pure population
no.genesPerPure=rep(5, no.pure)
no.genes= sum(no.genesPerPure)
GeneIndicator=rep(1:no.pure, no.genesPerPure)
# true mean values of the genes in the pure populations
# in the end we hope to estimate them from the mixed samples
datTrueMeans0=matrix( rnorm(no.genes*no.pure,sd=.3), nrow= no.genes,ncol=no.pure)
for (i in 1:no.pure ){
datTrueMeans0[GeneIndicator==i,i]= datTrueMeans0[GeneIndicator==i,i]+1
}
dimnames(datTrueMeans0)[[1]]=paste("Gene",1:dim(datTrueMeans0)[[1]],sep=".")
dimnames(datTrueMeans0)[[2]]=paste("MeanPureCellType",1:dim(datTrueMeans0)[[2]],
sep=".")
```



```

# plot.mat(datTrueMeans0)
# simulate the (expression) values of the admixed population samples

noise=matrix(rnorm(m*no.genes,sd=.1),nrow=m,ncol= no.genes)
datE.Admixture= as.matrix(datProportions) %*% t(datTrueMeans0) + noise
dimnames(datE.Admixture)[[1]]=paste("MixedTissue",1:m,sep=".")

datPredictedMeans=populationMeansInAdmixture(datProportions,datE.Admixture)

par(mfrow=c(2,2))
for (i in 1:4 ){
  verboseScatterplot(datPredictedMeans[,i],datTrueMeans0[,i],
  xlab="predicted mean",ylab="true mean",main="all populations")
  abline(0,1)
}

#assume we only study 2 populations (ie we ignore the others)
selectPopulations=c(1,2)
datPredictedMeansTooFew=populationMeansInAdmixture(datProportions[,selectPopulations],
  datE.Admixture)

par(mfrow=c(2,2))
for (i in 1:length(selectPopulations) ){
  verboseScatterplot(datPredictedMeansTooFew[,i],datTrueMeans0[,i],
  xlab="predicted mean",ylab="true mean",main="too few populations")
  abline(0,1)
}

#assume we erroneously add a population
datProportionsTooMany=data.frame(datProportions,WrongProp=sample(datProportions[,1]))
datPredictedMeansTooMany=populationMeansInAdmixture(datProportionsTooMany,
  datE.Admixture)

par(mfrow=c(2,2))
for (i in 1:4 ){
  verboseScatterplot(datPredictedMeansTooMany[,i],datTrueMeans0[,i],
  xlab="predicted mean",ylab="true mean",main="too many populations")
  abline(0,1)
}

```

---

pquantile

*Parallel quantile, median, mean*


---

## Description

Calculation of “parallel” quantiles, medians, and means, across given arguments.

## Usage

```

pquantile(prob, ...)
pmedian(...)
pmean(...)

```

**Arguments**

`prob` A number or vector of probabilities at which to calculate the quantile. See [quantile](#).

... Numeric arguments. All arguments must have the same dimensions. See details.

**Details**

Given the arguments, say `x,y,z`, of equal dimensions, the `pquantile` calculates and returns the quantile of the first components of `x,y,z`, then the second components, etc. Similarly, `pmedian` and `pmean` calculate the median and mean, respectively.

**Value**

A vector or array containing the quantiles, medians, or means. The dimensions are determined by the dimensions of the input arguments and whether the `prob` input is scalar or a vector. If any of the input variables have `dimnames`, the first non-NULL `dimnames` are copied into the output.

**Author(s)**

Peter Langfelder and Steve Horvath

**See Also**

[pmin](#) and [pmax](#) for analogous functions for minimum and maximum, [quantile](#), [median](#), [mean](#) for the underlying statistics.

**Examples**

```
# Generate 2 simple matrices
a = matrix(c(1:12), 3, 4);
b = a+ 1;
c = a + 2;

# Set the colnames on matrix a
colnames(a) = spaste("col_", c(1:4));

# Example use

pquantile(prob = 0.5, a, b, c)
pquantile(prob = c(0, 0.5, 1), a,b, c);

pmean(a,b,c)
pmedian(a,b,c)
```

---

`prepComma`*Prepend a comma to a non-empty string*

---

**Description**

Utility function that prepends a comma before the input string if the string is non-empty.

**Usage**

```
prepComma(s)
```

**Arguments**

`s` Character string.

**Value**

If `s` is non-empty, returns `paste(", ", s)`, otherwise returns `s`.

**Author(s)**

Peter Langfelder

**Examples**

```
prepComma("abc");  
prepComma("");
```

---

`preservationNetworkConnectivity`*Network preservation calculations*

---

**Description**

This function calculates several measures of gene network preservation. Given gene expression data in several individual data sets, it calculates the individual adjacency matrices, forms the preservation network and finally forms several summary measures of adjacency preservation for each node (gene) in the network.

**Usage**

```
preservationNetworkConnectivity(  
  multiExpr,  
  useSets = NULL, useGenes = NULL,  
  corFnc = "cor", corOptions = "use='p'",  
  networkType = "unsigned",  
  power = 6,  
  sampleLinks = NULL, nLinks = 5000,  
  blockSize = 1000,  
  setSeed = 12345,
```

```
weightPower = 2,
verbose = 2, indent = 0)
```

### Arguments

multiExpr	expression data in the multi-set format (see <a href="#">checkSets</a> ). A vector of lists, one per set. Each set must contain a component data that contains the expression data, with rows corresponding to samples and columns to genes or probes.
useSets	optional specification of sets to be used for the preservation calculation. Defaults to using all sets.
useGenes	optional specification of genes to be used for the preservation calculation. Defaults to all genes.
corFnc	character string containing the name of the function to calculate correlation. Suggested functions include "cor" and "bicor".
corOptions	further argument to the correlation function.
networkType	a character string encoding network type. Recognized values are (unique abbreviations of) "unsigned", "signed", and "signed hybrid".
power	soft thresholding power for network construction. Should be a number greater than 1.
sampleLinks	logical: should network connections be sampled (TRUE) or should all connections be used systematically (FALSE)?
nLinks	number of links to be sampled. Should be set such that nLinks * nNeighbors be several times larger than the number of genes.
blockSize	correlation calculations will be split into square blocks of this size, to prevent running out of memory for large gene sets.
setSeed	seed to be used for sampling, for repeatability. If a seed already exists, it is saved before the sampling starts and restored upon exit.
weightPower	power with which higher adjacencies will be weighted in weighted means
verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.
indent	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

### Details

The preservation network is formed from adjacencies of compared sets. For 'complete' preservations, all given sets are compared at once; for 'pairwise' preservations, the sets are compared in pairs. Unweighted preservations are simple mean preservations for each node; their weighted counterparts are weighted averages in which a preservation of adjacencies  $A_{ij}^{(1)}$  and  $A_{ij}^{(2)}$  of nodes  $i, j$  between sets 1 and 2 is weighted by  $[(A_{ij}^{(1)} + A_{ij}^{(2)})/2]^{weightPower}$ . The hyperbolic preservation is based on  $\tanh[(max - min)/(max + min)^2]$ , where  $max$  and  $min$  are the componentwise maximum and minimum of the compared adjacencies, respectively.

### Value

A list with the following components:

pairwise	a matrix with rows corresponding to genes and columns to unique pairs of given sets, giving the pairwise preservation of the adjacencies connecting the gene to all other genes.
complete	a vector with one entry for each input gene containing the complete mean preservation of the adjacencies connecting the gene to all other genes.
pairwiseWeighted	a matrix with rows corresponding to genes and columns to unique pairs of given sets, giving the pairwise weighted preservation of the adjacencies connecting the gene to all other genes.
completeWeighted	a vector with one entry for each input gene containing the complete weighted mean preservation of the adjacencies connecting the gene to all other genes.
pairwiseHyperbolic	a matrix with rows corresponding to genes and columns to unique pairs of given sets, giving the pairwise hyperbolic preservation of the adjacencies connecting the gene to all other genes.
completeHyperbolic	a vector with one entry for each input gene containing the complete mean hyperbolic preservation of the adjacencies connecting the gene to all other genes.
pairwiseWeightedHyperbolic	a matrix with rows corresponding to genes and columns to unique pairs of given sets, giving the pairwise weighted hyperbolic preservation of the adjacencies connecting the gene to all other genes.
completeWeightedHyperbolic	a vector with one entry for each input gene containing the complete weighted hyperbolic mean preservation of the adjacencies connecting the gene to all other genes.

**Author(s)**

Peter Langfelder

**References**

Langfelder P, Horvath S (2007) Eigengene networks for studying the relationships between co-expression modules. *BMC Systems Biology* 2007, 1:54

**See Also**

[adjacency](#) for calculation of adjacency;

---

projectiveKMeans

*Projective K-means (pre-)clustering of expression data*

---

**Description**

Implementation of a variant of K-means clustering for expression data.

**Usage**

```
projectiveKMeans(
  datExpr,
  preferredSize = 5000,
  nCenters = as.integer(min(ncol(datExpr)/20, preferredSize^2/ncol(datExpr))),
  sizePenaltyPower = 4,
  networkType = "unsigned",
  randomSeed = 54321,
  checkData = TRUE,
  maxIterations = 1000,
  verbose = 0, indent = 0)
```

**Arguments**

datExpr	expression data. A data frame in which columns are genes and rows are samples. NAs are allowed, but not too many.
preferredSize	preferred maximum size of clusters.
nCenters	number of initial clusters. Empirical evidence suggests that more centers will give a better preclustering; the default is an attempt to arrive at a reasonable number.
sizePenaltyPower	parameter specifying how severe is the penalty for clusters that exceed preferredSize.
networkType	network type. Allowed values are (unique abbreviations of) "unsigned", "signed", "signed hybrid". See <a href="#">adjacency</a> .
randomSeed	integer to be used as seed for the random number generator before the function starts. If a current seed exists, it is saved and restored upon exit.
checkData	logical: should data be checked for genes with zero variance and genes and samples with excessive numbers of missing samples? Bad samples are ignored; returned cluster assignment for bad genes will be NA.
maxIterations	maximum iterations to be attempted.
verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.
indent	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

**Details**

The principal aim of this function within WGCNA is to pre-cluster a large number of genes into smaller blocks that can be handled using standard WGCNA techniques.

This function implements a variant of K-means clustering that is suitable for co-expression analysis. Cluster centers are defined by the first principal component, and distances by correlation (more precisely, 1-correlation). The distance between a gene and a cluster is multiplied by a factor of  $\max(\text{clusterSize}/\text{preferredSize}, 1)^{\text{sizePenaltyPower}}$ , thus penalizing clusters whose size exceeds preferredSize. The function starts with randomly generated cluster assignment (hence the need to set the random seed for repeatability) and executes iterations of calculating new centers and reassigning genes to nearest center until the clustering becomes stable. Before returning, nearby clusters are iteratively combined if their combined size is below preferredSize.

The standard principal component calculation via the function `svd` fails from time to time (likely a convergence problem of the underlying lapack functions). Such errors are trapped and the principal

component is approximated by a weighted average of expression profiles in the cluster. If verbose is set above 2, an informational message is printed whenever this approximation is used.

### Value

A list with the following components:

clusters	a numerical vector with one component per input gene, giving the cluster number in which the gene is assigned.
centers	cluster centers, that is their first principal components.

### Author(s)

Peter Langfelder

---

proportionsInAdmixture

*Estimate the proportion of pure populations in an admixed population based on marker expression values.*

---

### Description

Assume that `datE.Admixture` provides the expression values from a mixture of cell types (admixed population) and you want to estimate the proportion of each pure cell type in the mixed samples (rows of `datE.Admixture`). The function allows you to do this as long as you provide a data frame `MarkerMeansPure` that reports the mean expression values of markers in each of the pure cell types.

### Usage

```
proportionsInAdmixture(
  MarkerMeansPure,
  datE.Admixture,
  calculateConditionNumber = FALSE,
  coefToProportion = TRUE)
```

### Arguments

`MarkerMeansPure`

is a data frame whose first column reports the name of the marker and the remaining columns report the mean values of the markers in each of the pure populations. The function will estimate the proportion of pure cells which correspond to columns 2 through of `dim(MarkerMeansPure)[[2]]` of `MarkerMeansPure`. Rows that contain missing values (NA) will be removed.

`datE.Admixture`

is a data frame of expression data, e.g. the columns of `datE.Admixture` could correspond to thousands of genes. The rows of `datE.Admixture` correspond to the admixed samples for which the function estimates the proportions of pure populations. Some of the markers specified in the first column of `MarkerMeansPure` should correspond to column names of `datE.Admixture`.

**calculateConditionNumber**

logical. Default is FALSE. If set to TRUE then it uses the kappa function to calculate the condition number of the matrix `MarkerMeansPure[, -1]`. This allows one to determine whether the linear model for estimating the proportions is well specified. Type `help(kappa)` to learn more. `kappa()` computes by default (an estimate of) the 2-norm condition number of a matrix or of the R matrix of a QR decomposition, perhaps of a linear fit.

**coefToProportion**

logical. By default, it is set to TRUE. When estimating the proportions the function fits a multivariate linear model. Ideally, the coefficients of the linear model correspond to the proportions in the admixed samples. But sometimes the coefficients take on negative values or do not sum to 1. If `coefToProportion=TRUE` then negative coefficients will be set to 0 and the remaining coefficients will be scaled so that they sum to 1.

**Details**

The methods implemented in this function were motivated by the gene expression deconvolution approach described by Abbas et al (2009), Lu et al (2003), Wang et al (2006). This approach can be used to predict the proportions of (pure) cells in a complex tissue, e.g. the proportion of blood cell types in whole blood. To define the markers, you may need to have expression data from pure populations. Then you can define markers based on a significant t-test or ANOVA across the pure populations. Next use the pure population data to estimate corresponding mean expression values. Hopefully, the array platforms and normalization methods for `datE.MarkersAdmixtureTranspose` and `MarkerMeansPure` are comparable. When dealing with Affymetrix data: we have successfully used it on untransformed MAS5 data. For statisticians: To estimate the proportions, we use the coefficients of a linear model. Specifically: `datCoef= t(lm(datE.MarkersAdmixtureTranspose ~MarkerMeansPure[, -1])` where `datCoef` is a matrix whose rows correspond to the mixed samples (rows of `datE.Admixture`) and the columns correspond to pure populations (e.g. cell types), i.e. the columns of `MarkerMeansPure[, -1]`. More details can be found in Abbas et al (2009).

**Value**

A list with the following components

**PredictedProportions**

data frame that contains the predicted proportions. The rows of `PredictedProportions` correspond to the admixed samples, i.e. the rows of `datE.Admixture`. The columns of `PredictedProportions` correspond to the pure populations, i.e. the columns of `MarkerMeansPure[, -1]`.

**datCoef=datCoef**

data frame of numbers that is analogous to `PredictedProportions`. In general, `datCoef` will only be different from `PredictedProportions` if `coefToProportion=TRUE`. See the description of `coefToProportion`

**conditionNumber**

This is the condition number resulting from the kappa function. See the description of `calculateConditionNumber`.

**markersUsed**

vector of character strings that contains the subset of marker names (specified in the first column of `MarkerMeansPure`) that match column names of `datE.Admixture` and that contain non-missing pure mean values.

**Note**

This function can be considered a wrapper of the `lm` function.



**Author(s)**

Steve Horvath, Chaochao Cai

**References**

Abbas AR, Wolslegel K, Seshasayee D, Modrusan Z, Clark HF (2009) Deconvolution of Blood Microarray Data Identifies Cellular Activation Patterns in Systemic Lupus Erythematosus. PLoS ONE 4(7): e6098. doi:10.1371/journal.pone.0006098

Lu P, Nakorchevskiy A, Marcotte EM (2003) Expression deconvolution: a reinterpretation of DNA microarray data reveals dynamic changes in cell populations. Proc Natl Acad Sci U S A 100: 10370-10375.

Wang M, Master SR, Chodosh LA (2006) Computational expression deconvolution in a complex mammalian organ. BMC Bioinformatics 7: 328.

**See Also**

[lm, kappa](#)

---

propVarExplained

*Proportion of variance explained by eigengenes.*

---

**Description**

This function calculates the proportion of variance of genes in each module explained by the respective module eigengene.

**Usage**

```
propVarExplained(datExpr, colors, MEs, corFnc = "cor", corOptions = "use = 'p'")
```

**Arguments**

datExpr	expression data. A data frame in which columns are genes and rows are samples. NAs are allowed and will be ignored.
colors	a vector giving module assignment for genes given in datExpr. Unique values should correspond to the names of the eigengenes in MEs.
MEs	a data frame of module eigengenes in which each column is an eigengene and each row corresponds to a sample.
corFnc	character string containing the name of the function to calculate correlation. Suggested functions include "cor" and "bicor".
corOptions	further argument to the correlation function.

**Details**

For compatibility with other functions, entries in color are matched to a substring of names(MEs) starting at position 3. For example, the entry "turquoise" in colors will be matched to the eigengene named "MEturquoise". The first two characters of the eigengene name are ignored and can be arbitrary.

**Value**

A vector with one entry per eigengene containing the proportion of variance of the module explained by the eigengene.

**Author(s)**

Peter Langfelder

**See Also**

[moduleEigengenes](#)

---

PWLists

*Pathways with Corresponding Gene Markers - Compiled by Mike Palazzolo and Jim Wang from CHDI*

---

**Description**

This matrix gives a predefined set of marker genes for many immune response pathways, as assembled by Mike Palazzolo and Jim Wang from CHDI, and colleagues. It is used with `userListEnrichment` to search user-defined gene lists for enrichment.

**Usage**

```
data(PWLists)
```

**Format**

A 124350 x 2 matrix of characters containing 2724 Gene / Category pairs. The first column (Gene) lists genes corresponding to a given category (second column). Each Category entry is of the form `<gene set>__<reference>`.

**Source**

For more information about this list, please see [userListEnrichment](#)

**Examples**

```
data(PWLists)
head(PWLists)
```

---

qvalue	<i>Estimate the q-values for a given set of p-values</i>
--------	--

---

### Description

Estimate the q-values for a given set of p-values. The q-value of a test measures the proportion of false positives incurred (called the false discovery rate) when that particular test is called significant.

### Usage

```
qvalue(p, lambda=seq(0,0.90,0.05), pi0.method="smoother", fdr.level=NULL, robust=FALSE,
       smooth.df=3, smooth.log.pi0=FALSE)
```

### Arguments

p	A vector of p-values (only necessary input)
lambda	The value of the tuning parameter to estimate $\pi_0$ . Must be in [0,1). Optional, see Storey (2002).
pi0.method	Either "smoother" or "bootstrap"; the method for automatically choosing tuning parameter in the estimation of $\pi_0$ , the proportion of true null hypotheses
fdr.level	A level at which to control the FDR. Must be in (0,1]. Optional; if this is selected, a vector of TRUE and FALSE is returned that specifies whether each q-value is less than fdr.level or not.
robust	An indicator of whether it is desired to make the estimate more robust for small p-values and a direct finite sample estimate of pFDR. Optional.
smooth.df	Number of degrees-of-freedom to use when estimating $\pi_0$ with a smoother. Optional.
smooth.log.pi0	If TRUE and pi0.method = "smoother", $\pi_0$ will be estimated by applying a smoother to a scatterplot of $\log \pi_0$ estimates against the tuning parameter $\lambda$ . Optional.

### Details

If no options are selected, then the method used to estimate  $\pi_0$  is the smoother method described in Storey and Tibshirani (2003). The bootstrap method is described in Storey, Taylor & Siegmund (2004).

### Value

A list containing:

call	function call
pi0	an estimate of the proportion of null p-values
qvalues	a vector of the estimated q-values (the main quantity of interest)
pvalues	a vector of the original p-values
significant	if fdr.level is specified, and indicator of whether the q-value fell below fdr.level (taking all such q-values to be significant controls FDR at level fdr.level)

**Note**

This function is adapted from package `qvalue`. The reason we provide our own copy is that package `qvalue` contains additional functionality that relies on Tcl/Tk which has led to multiple problems. Our copy does not require Tcl/Tk.

**Author(s)**

John D. Storey <jstorey@u.washington.edu>, adapted for WGCNA by Peter Langfelder

**References**

Storey JD. (2002) A direct approach to false discovery rates. *Journal of the Royal Statistical Society, Series B*, 64: 479-498.

Storey JD and Tibshirani R. (2003) Statistical significance for genome-wide experiments. *Proceedings of the National Academy of Sciences*, 100: 9440-9445.

Storey JD. (2003) The positive false discovery rate: A Bayesian interpretation and the q-value. *Annals of Statistics*, 31: 2013-2035.

Storey JD, Taylor JE, and Siegmund D. (2004) Strong control, conservative point estimation, and simultaneous conservative consistency of false discovery rates: A unified approach. *Journal of the Royal Statistical Society, Series B*, 66: 187-205.

QVALUE Manual <http://faculty.washington.edu/~jstorey/qvalue/manual.pdf>

---

qvalue.restricted      *qvalue convenience wrapper*

---

**Description**

This function calls `qvalue` on finite input p-values, optionally traps errors from the q-value calculation, and returns just the q values.

**Usage**

```
qvalue.restricted(p, trapErrors = TRUE, ...)
```

**Arguments**

<code>p</code>	a vector of p-values. Missing data are allowed and will be removed.
<code>trapErrors</code>	logical: should errors generated by function <code>qvalue</code> be trapped? If TRUE, the errors will be silently ignored and the returned q-values will all be NA.
<code>...</code>	other arguments to function <code>qvalue</code> .

**Value**

A vector of q-values. Entries whose corresponding p-values were not finite will be NA.

**Author(s)**

Peter Langfelder

**See Also**[qvalue](#)

---

randIndex	<i>Rand index of two partitions</i>
-----------	-------------------------------------

---

**Description**

Computes the Rand index, a measure of the similarity between two clusterings.

**Usage**

```
randIndex(tab, adjust = TRUE)
```

**Arguments**

tab	a matrix giving the cross-tabulation table of two clusterings.
adjust	logical: should the "adjusted" version be computed?

**Value**

the Rand index of the input table.

**Author(s)**

Steve Horvath

**References**

W. M. Rand (1971). "Objective criteria for the evaluation of clustering methods". *Journal of the American Statistical Association* 66: 846-850

---

randomGLMpredictor	<i>Random generalized linear model predictor</i>
--------------------	--

---

**Description**

An ensemble predictor based on bootstrap aggregation (bagging) of generalized linear models whose covariates are selected using forward stepwise regression according to AIC criteria.

**Usage**

```
randomGLMpredictor(
  x, y, xtest = NULL,
  classify = TRUE,
  nBags = 100,
  replace = TRUE,
  nObsInBag = if (replace) nrow(x) else as.integer(0.632 * nrow(x)),
  nFeaturesInBag = ceiling(ifelse(ncol(x) <= 10, ncol(x), ifelse(ncol(x) <= 300,
    (1.0276 - 0.00276 * ncol(x)) * ncol(x), ncol(x)/5))),
  nCandidateCovariates=50,
  candidateCorFnc= cor,
  candidateCorOptions = list(method = "pearson", use="p"),
  mandatoryCovariates = NULL,
  randomSeed = 12345,
  verbose =1)
```

**Arguments**

x	a matrix with rows correspond to observations and columns corresponding to features (covariates).
y	class outcome (factor variable) or quantitative outcome (numeric variable).
xtest	an optional matrix (whose columns correspond to those in x) which contain test (validation) data. The number of rows will typically be different from those in x.
classify	logical: should the response be treated as a binary variable (TRUE) or as a continuous variable (FALSE)?
nBags	number of bags in the ensemble predictor.
replace	logical which determines whether the observations for the bag (bootstrap data) are sampled with or without replacement. The function randomly select bagging observations with or without replacement.
nObsInBag	number of observations selected for each bag. Typically, a bootstrap sample (bag) has the same number of observations as in the original observed data (i.e. the rows of x).
nFeaturesInBag	number of features selected into each bag. Features are randomly selected without replacement.
nCandidateCovariates	top number of features selected with highest absolute correlation with the outcome in individual bag. These features/covariates become the candidates for forward stepwise regression.
candidateCorFnc	the correlation function used to select candidate covariates. Either cor or bicor.
candidateCorOptions	list of arguments to correlation function. If bicor is chosen for class outcome, make sure to include "robustY=F".
mandatoryCovariates	indices of features that forced into all regression models across bags. As default, no feature is mandatory.
randomSeed	NULL or integer. The seed for the random number generator. If NULL, the seed will not be set. If non-NULL and the random generator has been initialized prior to the function call, the latter's state is saved and restored upon exit.

`verbose` integer which determines the level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.

### Details

The `randomGLMpredictor` function requires the R package MASS since it makes use of the function `stepAIC`. Basically, `randomGLMpredictor` first selects bootstrapping samples and features randomly for each bag, and then restricts the analysis to features that are highly correlated with the outcome. Prediction in each bag is made based on forward stepwise regression (logistic for binary outcomes, linear for quantitative outcomes). An overall prediction is obtained by averaging results from all bags. Generally, `nCandidateCovariates > 100` is not recommended, because the forward selection process is time-consuming. If `"nBags=1, replace=F, nObsInBag=nrow(x)"` is used, the function becomes a stepwise generalized linear model predictor without bagging.

### Value

The function returns a list with the following components:

- `predictedOOB` the predicted classification of the input data based on out-of-bag samples. Only for binary outcomes.
- `predictedOOB.cont`  
In case of a binary outcome, this is the predicted probability of each outcome specified by `y` based on out-of-bag samples. In case of a continuous outcome, this is the predicted value based on out-of-bag samples.
- `predictedTest` if test set is given, the predicted classification for test data. Only for binary outcomes.
- `predictedTest.cont`  
if test set is given, the predicted probability of each outcome specified by `y` for test data for binary outcomes. In case of a continuous outcome, this is the test set predicted value.
- `bagObsIndx` a matrix with `nBags` rows and `nObsInBag` columns, giving the indices of observations selected for each bag.
- `datSelectedAsCandidates`  
a (0,1) matrix with `nBags` rows and columns corresponding to features, indicating which features are selected as candidate regression covariates in each bag.
- `datSelectedByForwardRegression`  
a (0,1) matrix with `nBags` rows and columns corresponding to features, indicating which features/covariates are selected into the final regression model in each bag.
- `datCoefOfForwardRegression`  
a matrix with `nBags` rows and columns corresponding to features, giving the final generalized linear model coefficients for features in each bag.
- `timesSelectedByForwardRegression`  
a variable importance measure, giving the times each feature is selected into final models in all bags.

### Author(s)

Lin Song

## References

Lin Song, Peter Langfelder, Steve Horvath: Random generalized linear model: a superior ensemble predictor involving few features. *BMC Bioinformatics*, future.

## Examples

```
## binary outcome prediction
# data generation
data(iris)
iris=iris[1:100,]
iris$Species = as.factor(as.character(iris$Species))
set.seed(1)
indx=sample(100, 67, replace=FALSE)
alldat1=iris[indx, ]
alldat2=iris[-indx,]
dat1=alldat1[,-5]
y1=alldat1[,5]
dat2=alldat2[,-5]
y2=alldat2[,5]

# predict with a small number of bags - normally nBags should be at least 100.
RGLM = randomGLMpredictor(dat1, y1, dat2, nCandidateCovariates=ncol(dat1), nBags=30)
y2predict = RGLM$predictedTest
table(y2predict, y2)
```

---

rankPvalue	<i>Estimate the p-value for ranking consistently high (or low) on multiple lists</i>
------------	--

---

## Description

The function rankPvalue calculates the p-value for observing that an object (corresponding to a row of the input data frame datS) has a consistently high ranking (or low ranking) according to multiple ordinal scores (corresponding to the columns of the input data frame datS).

## Usage

```
rankPvalue(datS, columnweights = NULL,
           na.last = "keep", ties.method = "average",
           calculateQvalue = TRUE, pValueMethod = "all")
```

## Arguments

datS	a data frame whose rows represent objects that will be ranked. Each column of datS represents an ordinal variable (which can take on negative values). The columns correspond to (possibly signed) object significance measures, e.g., statistics (such as Z statistics), ranks, or correlations.
columnweights	allows the user to input a vector of non-negative numbers reflecting weights for the different columns of datZ. If it is set to NULL then all weights are equal.



<code>na.last</code>	controls the treatment of missing values (NAs) in the rank function. If TRUE, missing values in the data are put last (i.e. they get the highest rank values). If FALSE, they are put first; if NA, they are removed; if "keep" they are kept with rank NA. See <a href="#">rank</a> for more details.
<code>ties.method</code>	represents the ties method used in the rank function for the percentile rank method. See <a href="#">rank</a> for more details.
<code>calculateQvalue</code>	logical: should q-values be calculated? If set to TRUE then the function calculates corresponding q-values (local false discovery rates) using the qvalue package, see Storey JD and Tibshirani R. (2003). This option assumes that qvalue package has been installed.
<code>pValueMethod</code>	determines which method is used for calculating p-values. By default it is set to "all", i.e. both methods are used. If it is set to "rank" then only the percentile rank method is used. If it set to "scale" then only the scale method will be used.

## Details

The function calculates asymptotic p-values (and optionally q-values) for testing the null hypothesis that the values in the columns of `datS` are independent. This allows us to find objects (rows) with consistently high (or low) values across the columns.

Example: Imagine you have 5 vectors of Z statistics corresponding to the columns of `datS`. Further assume that a gene has ranks 1,1,1,1,20 in the 5 lists. It seems very significant that the gene ranks number 1 in 4 out of the 5 lists. The function `rankPvalue` can be used to calculate a p-value for this occurrence.

The function uses the central limit theorem to calculate asymptotic p-values for two types of test statistics that measure consistently high or low ordinal values. The first method (referred to as percentile rank method) leads to accurate estimates of p-values if `datS` has at least 4 columns but it can be overly conservative. The percentile rank method replaces each column `datS` by the ranked version `rank(datS[,i])` (referred to as low ranking) and by `rank(-datS[,i])` (referred to as high ranking). Low ranking and high ranking allow one to find consistently small values or consistently large values of `datS`, respectively. All ranks are divided by the maximum rank so that the result lies in the unit interval  $[0,1]$ . In the following, we refer to `rank/max(rank)` as percentile rank. For a given object (corresponding to a row of `datS`) the observed percentile rank follows approximately a uniform distribution under the null hypothesis. The test statistic is defined as the sum of the percentile ranks (across the columns of `datS`). Under the null hypothesis that there is no relationship between the rankings of the columns of `datS`, this (row sum) test statistic follows a distribution that is given by the convolution of random uniform distributions. Under the null hypothesis, the individual percentile ranks are independent and one can invoke the central limit theorem to argue that the row sum test statistic follows asymptotically a normal distribution. It is well-known that the speed of convergence to the normal distribution is extremely fast in case of identically distributed uniform distributions. Even when `datS` has only 4 columns, the difference between the normal approximation and the exact distribution is negligible in practice (Killmann et al 2001). In summary, we use the central limit theorem to argue that the sum of the percentile ranks follows a normal distribution whose mean and variance can be calculated using the fact that the mean value of a uniform random variable (on the unit interval) equals 0.5 and its variance equals 1/12.

The second method for calculating p-values is referred to as scale method. It is often more powerful but its asymptotic p-value can only be trusted if either `datS` has a lot of columns or if the ordinal scores (columns of `datS`) follow an approximate normal distribution. The scale method scales (or standardizes) each ordinal variable (column of `datS`) so that it has mean 0 and variance 1. Under the null hypothesis of independence, the row sum follows approximately a normal distribution if the assumptions of the central limit theorem are met. In practice, we find that the second approach is often more powerful but it makes more distributional assumptions (if `datS` has few columns).

**Value**

A list whose actual content depends on which p-value methods is selected, and whether q0values are calculated. The following inner components are calculated, organized in outer components datoutrank and datoutscale,:

pValueExtremeRank	This is the minimum between pValueLowRank and pValueHighRank, i.e. $\min(\text{pValueLow}, \text{pValueHigh})$
pValueLowRank	Asymptotic p-value for observing a consistently low value across the columns of datS based on the rank method.
pValueHighRank	Asymptotic p-value for observing a consistently low value across the columns of datS based on the rank method.
pValueExtremeScale	This is the minimum between pValueLowScale and pValueHighScale, i.e. $\min(\text{pValueLow}, \text{pValueHigh})$
pValueLowScale	Asymptotic p-value for observing a consistently low value across the columns of datS based on the Scale method.
pValueHighScale	Asymptotic p-value for observing a consistently low value across the columns of datS based on the Scale method.
qValueExtremeRank	local false discovery rate (q-value) corresponding to the p-value pValueExtremeRank
qValueLowRank	local false discovery rate (q-value) corresponding to the p-value pValueLowRank
qValueHighRank	local false discovery rate (q-value) corresponding to the p-value pValueHighRank
qValueExtremeScale	local false discovery rate (q-value) corresponding to the p-value pValueExtremeScale
qValueLowScale	local false discovery rate (q-value) corresponding to the p-value pValueLowScale
qValueHighScale	local false discovery rate (q-value) corresponding to the p-value pValueHighScale

**Author(s)**

Steve Horvath

**References**

- Killmann F, VonCollani E (2001) A Note on the Convolution of the Uniform and Related Distributions and Their Use in Quality Control. *Economic Quality Control* Vol 16 (2001), No. 1, 17-41. ISSN 0940-5151
- Storey JD and Tibshirani R. (2003) Statistical significance for genome-wide experiments. *Proceedings of the National Academy of Sciences*, 100: 9440-9445.

**See Also**

[rank](#), [qvalue](#)

---

recutBlockwiseTrees      *Repeat blockwise module detection from pre-calculated data*

---

### Description

Given consensus networks constructed for example using `blockwiseModules`, this function (re-)detects modules in them by branch cutting of the corresponding dendrograms. If repeated branch cuts of the same gene network dendrograms are desired, this function can save substantial time by re-using already calculated networks and dendrograms.

### Usage

```
recutBlockwiseTrees(
  datExpr,
  goodSamples, goodGenes,
  blocks,
  TOMfiles,
  dendrograms,
  corType = "pearson",
  networkType = "unsigned",
  deepSplit = 2,
  detectCutHeight = 0.995, minModuleSize = min(20, ncol(datExpr)/2 ),
  maxCoreScatter = NULL, minGap = NULL,
  maxAbsCoreScatter = NULL, minAbsGap = NULL,
  pamStage = TRUE, pamRespectsDendro = TRUE,
  minCoreKME = 0.5, minCoreKMESize = minModuleSize/3,
  minKMEtoStay = 0.3,
  reassignThreshold = 1e-6,
  mergeCutHeight = 0.15, impute = TRUE,
  trapErrors = FALSE, numericLabels = FALSE,
  verbose = 0, indent = 0)
```

### Arguments

<code>datExpr</code>	expression data. A data frame in which columns are genes and rows are samples. NAs are allowed, but not too many.
<code>goodSamples</code>	a logical vector specifying which samples are considered "good" for the analysis. See <code>goodSamplesGenes</code> .
<code>goodGenes</code>	a logical vector with length equal number of genes in <code>multiExpr</code> that specifies which genes are considered "good" for the analysis. See <code>goodSamplesGenes</code> .
<code>blocks</code>	specification of blocks in which hierarchical clustering and module detection should be performed. A numeric vector with one entry per gene of <code>multiExpr</code> giving the number of the block to which the corresponding gene belongs.
<code>TOMfiles</code>	a vector of character strings specifying file names in which the block-wise topological overlaps are saved.
<code>dendrograms</code>	a list of length equal the number of blocks, in which each component is a hierarchical clustering dendrograms of the genes that belong to the block.

corType	character string specifying the correlation to be used. Allowed values are (unique abbreviations of) "pearson" and "bicor", corresponding to Pearson and bid-weight midcorrelation, respectively. Missing values are handled using the <code>parwise.complete.obs</code> option.
networkType	network type. Allowed values are (unique abbreviations of) "unsigned", "signed", "signed hybrid". See <a href="#">adjacency</a> .
deepSplit	integer value between 0 and 4. Provides a simplified control over how sensitive module detection should be to module splitting, with 0 least and 4 most sensitive. See <a href="#">cutreeDynamic</a> for more details.
detectCutHeight	dendrogram cut height for module detection. See <a href="#">cutreeDynamic</a> for more details.
minModuleSize	minimum module size for module detection. See <a href="#">cutreeDynamic</a> for more details.
maxCoreScatter	maximum scatter of the core for a branch to be a cluster, given as the fraction of cutHeight relative to the 5th percentile of joining heights. See <a href="#">cutreeDynamic</a> for more details.
minGap	minimum cluster gap given as the fraction of the difference between cutHeight and the 5th percentile of joining heights. See <a href="#">cutreeDynamic</a> for more details.
maxAbsCoreScatter	maximum scatter of the core for a branch to be a cluster given as absolute heights. If given, overrides maxCoreScatter. See <a href="#">cutreeDynamic</a> for more details.
minAbsGap	minimum cluster gap given as absolute height difference. If given, overrides minGap. See <a href="#">cutreeDynamic</a> for more details.
pamStage	logical. If TRUE, the second (PAM-like) stage of module detection will be performed. See <a href="#">cutreeDynamic</a> for more details.
pamRespectsDendro	Logical, only used when pamStage is TRUE. If TRUE, the PAM stage will respect the dendrogram in the sense an object can be PAM-assigned only to clusters that lie below it on the branch that the object is merged into. See <a href="#">cutreeDynamic</a> for more details.
minCoreKME	a number between 0 and 1. If a detected module does not have at least minModuleKMESize genes with eigengene connectivity at least minCoreKME, the module is disbanded (its genes are unlabeled and returned to the pool of genes waiting for module detection).
minCoreKMESize	see minCoreKME above.
minKMEtoStay	genes whose eigengene connectivity to their module eigengene is lower than minKMEtoStay are removed from the module.
reassignThreshold	p-value ratio threshold for reassigning genes between modules. See <a href="#">Details</a> .
mergeCutHeight	dendrogram cut height for module merging.
impute	logical: should imputation be used for module eigengene calculation? See <a href="#">moduleEigengenes</a> for more details.
trapErrors	logical: should errors in calculations be trapped?
numericLabels	logical: should the returned modules be labeled by colors (FALSE), or by numbers (TRUE)?

verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.
indent	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

## Details

For details on blockwise module detection, see [blockwiseModules](#). This function implements the module detection subset of the functionality of [blockwiseModules](#); network construction and clustering must be performed in advance. The primary use of this function is to experiment with module detection settings without having to re-execute long network and clustering calculations whose results are not affected by the cutting parameters.

This function takes as input the networks and dendrograms that are produced by [blockwiseModules](#). Working block by block, modules are identified in the dendrogram by the Dynamic Hybrid Tree Cut algorithm. Found modules are trimmed of genes whose correlation with module eigengene (KME) is less than `minKMEtoStay`. Modules in which fewer than `minCoreKMESize` genes have KME higher than `minCoreKME` are disbanded, i.e., their constituent genes are pronounced unassigned.

After all blocks have been processed, the function checks whether there are genes whose KME in the module they assigned is lower than KME to another module. If p-values of the higher correlations are smaller than those of the native module by the factor `reassignThresholdPS`, the gene is re-assigned to the closer module.

In the last step, modules whose eigengenes are highly correlated are merged. This is achieved by clustering module eigengenes using the dissimilarity given by one minus their correlation, cutting the dendrogram at the height `mergeCutHeight` and merging all modules on each branch. The process is iterated until no modules are merged. See [mergeCloseModules](#) for more details on module merging.

## Value

A list with the following components:

colors	a vector of color or numeric module labels for all genes.
unmergedColors	a vector of color or numeric module labels for all genes before module merging.
MEs	a data frame containing module eigengenes of the found modules (given by colors).
MEsOK	logical indicating whether the module eigengenes were calculated without errors.

## Author(s)

Peter Langfelder

## References

Bin Zhang and Steve Horvath (2005) "A General Framework for Weighted Gene Co-Expression Network Analysis", *Statistical Applications in Genetics and Molecular Biology*: Vol. 4: No. 1, Article 17

**See Also**

[blockwiseModules](#) for full module calculation;  
[cutreeDynamic](#) for adaptive branch cutting in hierarchical clustering dendrograms;  
[mergeCloseModules](#) for merging of close modules.

---

recutConsensusTrees     *Repeat blockwise consensus module detection from pre-calculated data*

---

**Description**

Given consensus networks constructed for example using [blockwiseConsensusModules](#), this function (re-)detects modules in them by branch cutting of the corresponding dendrograms. If repeated branch cuts of the same gene network dendrograms are desired, this function can save substantial time by re-using already calculated networks and dendrograms.

**Usage**

```
recutConsensusTrees(
  multiExpr,
  goodSamples, goodGenes,
  blocks,
  TOMfiles,
  dendrograms,
  corType = "pearson",
  networkType = "unsigned",
  deepSplit = 2,
  detectCutHeight = 0.995, minModuleSize = 20,
  checkMinModuleSize = TRUE,
  maxCoreScatter = NULL, minGap = NULL,
  maxAbsCoreScatter = NULL, minAbsGap = NULL,
  pamStage = TRUE, pamRespectsDendro = TRUE,
  trimmingConsensusQuantile = 0,
  minCoreKME = 0.5, minCoreKMESize = minModuleSize/3,
  minKMEtoStay = 0.2,
  reassignThresholdPS = 1e-4,
  mergeCutHeight = 0.15,
  mergeConsensusQuantile = trimmingConsensusQuantile,
  impute = TRUE,
  trapErrors = FALSE,
  numericLabels = FALSE,
  verbose = 2, indent = 0)
```

**Arguments**

`multiExpr`     expression data in the multi-set format (see [checkSets](#)). A vector of lists, one per set. Each set must contain a component data that contains the expression data, with rows corresponding to samples and columns to genes or probes.

goodSamples	a list with one component per set. Each component is a logical vector specifying which samples are considered "good" for the analysis. See <a href="#">goodSamplesGenesMS</a> .
goodGenes	a logical vector with length equal number of genes in <code>multiExpr</code> that specifies which genes are considered "good" for the analysis. See <a href="#">goodSamplesGenesMS</a> .
blocks	specification of blocks in which hierarchical clustering and module detection should be performed. A numeric vector with one entry per gene of <code>multiExpr</code> giving the number of the block to which the corresponding gene belongs.
TOMFiles	a vector of character strings specifying file names in which the block-wise topological overlaps are saved.
dendrograms	a list of length equal the number of blocks, in which each component is a hierarchical clustering dendrograms of the genes that belong to the block.
corType	character string specifying the correlation to be used. Allowed values are (unique abbreviations of) "pearson" and "bicor", corresponding to Pearson and bid-weight midcorrelation, respectively. Missing values are handled using the <code>parwise.complete.obs</code> option.
networkType	network type. Allowed values are (unique abbreviations of) "unsigned", "signed", "signed hybrid". See <a href="#">adjacency</a> . Note that while no new networks are computed in this function, this parameter affects the interpretation of correlations in this function.
deepSplit	integer value between 0 and 4. Provides a simplified control over how sensitive module detection should be to module splitting, with 0 least and 4 most sensitive. See <a href="#">cutreeDynamic</a> for more details.
detectCutHeight	dendrogram cut height for module detection. See <a href="#">cutreeDynamic</a> for more details.
minModuleSize	minimum module size for module detection. See <a href="#">cutreeDynamic</a> for more details.
checkMinModuleSize	logical: should sanity checks be performed on <code>minModuleSize</code> ?
maxCoreScatter	maximum scatter of the core for a branch to be a cluster, given as the fraction of <code>cutHeight</code> relative to the 5th percentile of joining heights. See <a href="#">cutreeDynamic</a> for more details.
minGap	minimum cluster gap given as the fraction of the difference between <code>cutHeight</code> and the 5th percentile of joining heights. See <a href="#">cutreeDynamic</a> for more details.
maxAbsCoreScatter	maximum scatter of the core for a branch to be a cluster given as absolute heights. If given, overrides <code>maxCoreScatter</code> . See <a href="#">cutreeDynamic</a> for more details.
minAbsGap	minimum cluster gap given as absolute height difference. If given, overrides <code>minGap</code> . See <a href="#">cutreeDynamic</a> for more details.
pamStage	logical. If TRUE, the second (PAM-like) stage of module detection will be performed. See <a href="#">cutreeDynamic</a> for more details.
pamRespectsDendro	Logical, only used when <code>pamStage</code> is TRUE. If TRUE, the PAM stage will respect the dendrogram in the sense an object can be PAM-assigned only to clusters that lie below it on the branch that the object is merged into. See <a href="#">cutreeDynamic</a> for more details.

trimmingConsensusQuantile	a number between 0 and 1 specifying the consensus quantile used for kME calculation that determines module trimming according to the arguments below.
minCoreKME	a number between 0 and 1. If a detected module does not have at least minModuleKMESize genes with eigengene connectivity at least minCoreKME, the module is disbanded (its genes are unlabeled and returned to the pool of genes waiting for module detection).
minCoreKMESize	see minCoreKME above.
minKMEtoStay	genes whose eigengene connectivity to their module eigengene is lower than minKMEtoStay are removed from the module.
reassignThresholdPS	per-set p-value ratio threshold for reassigning genes between modules. See Details.
mergeCutHeight	dendrogram cut height for module merging.
mergeConsensusQuantile	consensus quantile for module merging. See mergeCloseModules for details.
impute	logical: should imputation be used for module eigengene calculation? See <a href="#">moduleEigengenes</a> for more details.
trapErrors	logical: should errors in calculations be trapped?
numericLabels	logical: should the returned modules be labeled by colors (FALSE), or by numbers (TRUE)?
verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.
indent	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

## Details

For details on blockwise consensus module detection, see [blockwiseConsensusModules](#). This function implements the module detection subset of the functionality of [blockwiseConsensusModules](#); network construction and clustering must be performed in advance. The primary use of this function is to experiment with module detection settings without having to re-execute long network and clustering calculations whose results are not affected by the cutting parameters.

This function takes as input the networks and dendrograms that are produced by [blockwiseConsensusModules](#). Working block by block, modules are identified in the dendrograms by the Dynamic Hybrid tree cut. Found modules are trimmed of genes whose consensus module membership kME (that is, correlation with module eigengene) is less than minKMEtoStay. Modules in which fewer than minCoreKMESize genes have consensus KME higher than minCoreKME are disbanded, i.e., their constituent genes are pronounced unassigned.

After all blocks have been processed, the function checks whether there are genes whose KME in the module they assigned is lower than KME to another module. If p-values of the higher correlations are smaller than those of the native module by the factor reassignThresholdPS (in every set), the gene is re-assigned to the closer module.

In the last step, modules whose eigengenes are highly correlated are merged. This is achieved by clustering module eigengenes using the dissimilarity given by one minus their correlation, cutting the dendrogram at the height mergeCutHeight and merging all modules on each branch. The process is iterated until no modules are merged. See [mergeCloseModules](#) for more details on module merging.



**Value**

A list with the following components:

colors	module assignment of all input genes. A vector containing either character strings with module colors (if input <code>numericLabels</code> was unset) or numeric module labels (if <code>numericLabels</code> was set to TRUE). The color "grey" and the numeric label 0 are reserved for unassigned genes.
unmergedColors	module colors or numeric labels before the module merging step.
multiMEs	module eigengenes corresponding to the modules returned in <code>colors</code> , in multi-set format. A vector of lists, one per set, containing eigengenes, proportion of variance explained and other information. See <a href="#">multiSetMEs</a> for a detailed description.

**Note**

Basic sanity checks are performed on given arguments, but it is left to the user's responsibility to provide valid input.

**Author(s)**

Peter Langfelder

**References**

Langfelder P, Horvath S (2007) Eigengene networks for studying the relationships between co-expression modules. *BMC Systems Biology* 2007, 1:54

**See Also**

[blockwiseConsensusModules](#) for the full blockwise modules calculation. Parts of its output are natural input for this function.

[cutreeDynamic](#) for adaptive branch cutting in hierarchical clustering dendrograms;

[mergeCloseModules](#) for merging of close modules.

---

redWhiteGreen

*Red-white-green color sequence*

---

**Description**

Generate a red-white-green color sequence of a given length.

**Usage**

```
redWhiteGreen(n, gamma = 1)
```

**Arguments**

n	number of colors to be returned
gamma	color correction power

**Details**

The function returns a color vector that starts with pure green, gradually turns into white and then to red. The power gamma can be used to control the behaviour of the quarter- and three quarter-values (between red and white, and white and green, respectively). Higher powers will make the mid-colors more white, while lower powers will make the colors more saturated, respectively.

**Value**

A vector of colors of length n.

**Author(s)**

Peter Langfelder

**Examples**

```
par(mfrow = c(3, 1))
displayColors(redWhiteGreen(50));
displayColors(redWhiteGreen(50, 3));
displayColors(redWhiteGreen(50, 0.5));
```

---

relativeCorPredictionSuccess

*Compare prediction success*

---

**Description**

Compare prediction success of several gene screening methods.

**Usage**

```
relativeCorPredictionSuccess(
  corPredictionNew,
  corPredictionStandard,
  corTestSet,
  topNumber = 100)
```

**Arguments**

corPredictionNew	Matrix of predictor statistics
corPredictionStandard	Reference predictor statistics
corTestSet	Correlations of predictor variables with trait in test set
topNumber	A vector giving the numbers of top genes to consider

**Value**

Data frame with components

topNumber	copy of the input topNumber
kruskalp	Kruskal-Wallis p-values

**Author(s)**

Steve Horvath

**See Also**[corPredictionSuccess](#)

---

`removeGreyME`*Removes the grey eigengene from a given collection of eigengenes.*

---

**Description**

Given module eigengenes either in a single data frame or in a multi-set format, removes the grey eigengenes from each set. If the grey eigengenes are not found, a warning is issued.

**Usage**

```
removeGreyME(MEs, greyMENAME = paste(moduleColor.getMEprefix(), "grey", sep=""))
```

**Arguments**

MEs	Module eigengenes, either in a single data frame (typically for a single set), or in a multi-set format. See <a href="#">checkSets</a> for a description of the multi-set format.
greyMENAME	Name of the module eigengene (in each corresponding data frame) that corresponds to the grey color. This will typically be "PCgrey" or "MEgrey". If the module eigengenes were calculated using standard functions in this library, the default should work.

**Value**

Module eigengenes in the same format as input (either a single data frame or a vector of lists) with the grey eigengene removed.

**Author(s)**

Peter Langfelder, &lt;Peter.Langfelder@gmail.com&gt;

---

`removePrincipalComponents`*Remove leading principal components from data*

---

**Description**

This function calculates a fixed number of the first principal components of the given data and returns the residuals of a linear regression of each column on the principal components.

**Usage**

```
removePrincipalComponents(x, n)
```

**Arguments**

- x                    Input data, a numeric matrix. All entries must be non-missing and finite.
- n                    Number of principal components to remove. This must be smaller than the smaller of the number of rows and columns in x.

**Value**

A matrix of residuals of the same dimensions as x.

**Author(s)**

Peter Langfelder

**See Also**

[svd](#) for singular value decomposition, [lm](#) for linear regression

---

returnGeneSetsAsList    *Return pre-defined gene lists in several biomedical categories.*

---

**Description**

This function returns gene sets for use with other R functions. These gene sets can include inputted lists of genes and files containing user-defined lists of genes, as well as a pre-made collection of brain, blood, and other biological lists. The function returns gene lists associated with each category for use with other enrichment strategies (i.e., GSEA).

**Usage**

```
returnGeneSetsAsList(
  fnIn = NULL, catNmIn = fnIn,
  useBrainLists = FALSE, useBloodAtlases = FALSE,
  useStemCellLists = FALSE, useBrainRegionMarkers = FALSE,
  useImmunePathwayLists = FALSE, geneSubset=NULL)
```

**Arguments**

- fnIn                    A vector of file names containing user-defined lists. These files must be in one of three specific formats (see details section). The default (NULL) may only be used if one of the "use\_\_\_\_\_" parameters is TRUE.
- catNmIn                A vector of category names corresponding to each fnIn. This name will be appended to each overlap corresponding to that filename. The default sets the category names as the corresponding file names.
- useBrainLists         If TRUE, a pre-made set of brain-derived enrichment lists will be added to any user-defined lists for enrichment comparison. The default is FALSE. See references section for related references.
- useBloodAtlases        If TRUE, a pre-made set of blood-derived enrichment lists will be added to any user-defined lists for enrichment comparison. The default is FALSE. See references section for related references.

**useStemCellLists**

If TRUE, a pre-made set of stem cell (SC)-derived enrichment lists will be added to any user-defined lists for enrichment comparison. The default is FALSE. See references section for related references.

**useBrainRegionMarkers**

If TRUE, a pre-made set of enrichment lists for human brain regions will be added to any user-defined lists for enrichment comparison. The default is FALSE. These lists are derived from data from the Allen Human Brain Atlas (<http://human.brain-map.org/>). See references section for more details.

**useImmunePathwayLists**

If TRUE, a pre-made set of enrichment lists for immune system pathways will be added to any user-defined lists for enrichment comparison. The default is FALSE. These lists are derived from the lab of Daniel R Saloman. See references section for more details.

**geneSubset**

A vector of gene (or other) identifiers. If entered, only genes in this list will be returned in the output, otherwise all genes in each category will be returned (default, geneSubset=NULL).

**Details**

User-inputted files for `fnIn` can be in one of three formats:

1) Text files (must end in ".txt") with one list per file, where the first line is the list descriptor and the remaining lines are gene names corresponding to that list, with one gene per line. For example Ribosome RPS4 RPS8 ...

2) Gene / category files (must be csv files), where the first line is the column headers corresponding to Genes and Lists, and the remaining lines correspond to the genes in each list, for any number of genes and lists. For example: Gene, Category RPS4, Ribosome RPS8, Ribosome ... NDUF1, Mitochondria NDUF3, Mitochondria ... MAPT, AlzheimersDisease PSEN1, AlzheimersDisease PSEN2, AlzheimersDisease ...

3) Module membership (kME) table in csv format. Currently, the module assignment is the only thing that is used, so as long as the Gene column is 2nd and the Module column is 3rd, it doesn't matter what is in the other columns. For example, PSID, Gene, Module, <other columns> <psid>, RPS4, blue, <other columns> <psid>, NDUF1, red, <other columns> <psid>, RPS8, blue, <other columns> <psid>, NDUF3, red, <other columns> <psid>, MAPT, green, <other columns> ...

**Value****geneSets**

A list of categories in alphabetical order, where each component of the list is a character vector of all genes corresponding to the named category. For example:  
`geneSets = list(category1=c("gene1", "gene2"), category2=c("gene3", "gene4", "gene5"))`

**Author(s)**

Jeremy Miller

**References**

Please see the help file for `userListEnrichment` in the WGCNA library for references for the pre-defined lists.

## Examples

```
# Example: Return a list of genes for various immune pathways
geneSets = returnGeneSetsAsList(useImmunePathwayLists=TRUE)
geneSets[7:8]
```

---

rgcolors.func

*Red and Green Color Specification*

---

## Description

This function creates a vector of  $n$  “contiguous” colors, corresponding to  $n$  intensities (between 0 and 1) of the red, green and blue primaries, with the blue intensities set to zero. The values returned by `rgcolors.func` can be used with a `col=` specification in graphics functions or in [par](#).

## Usage

```
rgcolors.func(n=50)
```

## Arguments

`n` the number of colors ( $\geq 1$ ) to be used in the red and green palette.

## Value

a character vector of color names. Colors are specified directly in terms of their RGB components with a string of the form “\#RRGGBB”, where each of the pairs RR, GG, BB consist of two hexadecimal digits giving a value in the range 00 to FF.

## Author(s)

Sandrine Dudoit, <sandrine@stat.berkeley.edu>  
Jane Fridlyand, <janef@stat.berkeley.edu>

## See Also

[plotCor](#), [plotMat](#), [colors](#), [rgb](#), [image](#).

## Examples

```
rgcolors.func(n=5)
## The following vector is returned:
## "#00FF00" "#40BF00" "#808000" "#BF4000" "#FF0000"
```

---

scaleFreeFitIndex      *Calculation of fitting statistics for evaluating scale free topology fit.*

---

### Description

The function `scaleFreeFitIndex` calculates several indices (fitting statistics) for evaluating scale free topology fit. The input is a vector (of connectivities) `k`. Next `k` is discretized into `nBreaks` number of equal-width bins. Let's denote the resulting vector `dk`. The relative frequency for each bin is denoted `p.dk`.

### Usage

```
scaleFreeFitIndex(k, nBreaks = 10, removeFirst = FALSE)
```

### Arguments

`k`                      numeric vector whose components contain non-negative values  
`nBreaks`                positive integer. This determines the number of equal width bins.  
`removeFirst`           logical. If TRUE then the first bin will be removed.

### Value

Data frame with columns

`Rsquared.SFT`        the model fitting index (R.squared) from the following model `lm(log.p.dk ~ log.dk)`  
`slope.SFT`            the slope estimate from model `lm(log(p(k))~log(k))`  
`truncatedExponentialAdjRsquared`  
                           the adjusted R.squared measure from the truncated exponential model given by  
`lm2 = lm(log.p.dk ~ log.dk + dk)`.

### Author(s)

Steve Horvath

---

scaleFreePlot            *Visual check of scale-free topology*

---

### Description

A simple visula check of scale-free network ropology.

### Usage

```
scaleFreePlot(  
  connectivity,  
  nBreaks = 10,  
  truncated = FALSE,  
  removeFirst = FALSE,  
  main = "", ...)
```

**Arguments**

<code>connectivity</code>	vector containing network connectivities.
<code>nBreaks</code>	number of breaks in the connectivity dendrogram.
<code>truncated</code>	logical: should a truncated exponential fit be calculated and plotted in addition to the linear one?
<code>removeFirst</code>	logical: should the first bin be removed from the fit?
<code>main</code>	main title for the plot.
<code>...</code>	other graphical parameter to the <code>plot</code> function.

**Details**

The function plots a log-log plot of a histogram of the given connectivities, and fits a linear model plus optionally a truncated exponential model. The  $R^2$  of the fit can be considered an index of the scale freedom of the network topology.

**Value**

None.

**Author(s)**

Steve Horvath

**References**

Bin Zhang and Steve Horvath (2005) "A General Framework for Weighted Gene Co-Expression Network Analysis", *Statistical Applications in Genetics and Molecular Biology*: Vol. 4: No. 1, Article 17

**See Also**

[softConnectivity](#) for connectivity calculation in weighted networks.

---

SCsLists

*Stem Cell-Related Genes with Corresponding Gene Markers*

---

**Description**

This matrix gives a predefined set of genes related to several stem cell (SC) types, as reported in two previously-published studies. It is used with `userListEnrichment` to search user-defined gene lists for enrichment.

**Usage**

```
data(SCsLists)
```



**Format**

A 14003 x 2 matrix of characters containing Gene / Category pairs. The first column (Gene) lists genes corresponding to a given category (second column). Each Category entry is of the form <Stem cell-related category>\_\_<reference>, where the references can be found at [userListEnrichment](#). Note that the matrix is sorted first by Category and then by Gene, such that all genes related to the same category are listed sequentially.

**Source**

For references used in this variable, please see [userListEnrichment](#)

**Examples**

```
data(SCsLists)
head(SCsLists)
```

---

```
setCorrelationPreservation
```

*Summary correlation preservation measure*

---

**Description**

Given consensus eigengenes, the function calculates the average correlation preservation pair-wise for all pairs of sets.

**Usage**

```
setCorrelationPreservation(
  multiME,
  setLabels,
  excludeGrey = TRUE, greyLabel = "grey",
  method = "absolute")
```

**Arguments**

multiME	consensus module eigengenes in a multi-set format. A vector of lists with one list corresponding to each set. Each list must contain a component data that is a data frame whose columns are consensus module eigengenes.
setLabels	names to be used for the sets represented in multiME.
excludeGrey	logical: exclude the 'grey' eigengene from preservation measure?
greyLabel	module label corresponding to the 'grey' module. Usually this will be the character string "grey" if the labels are colors, and the number 0 if the labels are numeric.
method	character string giving the correlation preservation measure to use. Recognized values are (unique abbreviations of) "absolute", "hyperbolic".

**Details**

For each pair of sets, the function calculates the average preservation of correlation among the eigengenes. Two preservation measures are available, the absolute preservation (high if the two correlations are similar and low if they are different), and the hyperbolically scaled preservation, which de-emphasizes preservation of low correlation values.

**Value**

A data frame with each row and column corresponding to a set given in multiME, containing the pairwise average correlation preservation values. Names and rownames are set to entries of setLabels.

**Author(s)**

Peter Langfelder

**References**

Langfelder P, Horvath S (2007) Eigengene networks for studying the relationships between co-expression modules. BMC Systems Biology 2007, 1:54

**See Also**

[multiSetMEs](#) for module eigengene calculation;  
[plotEigengeneNetworks](#) for eigengene network visualization.

---

sigmoidAdjacencyFunction

*Sigmoid-type adjacency function.*

---

**Description**

Sigmoid-type function that converts a similarity to a weighted network adjacency.

**Usage**

```
sigmoidAdjacencyFunction(ss, mu = 0.8, alpha = 20)
```

**Arguments**

ss	similarity, a number between 0 and 1. Can be given as a scalar, vector or a matrix.
mu	shift parameter.
alpha	slope parameter.

**Details**

The sigmoid adjacency function is defined as  $1/(1 + \exp[-\alpha(ss - \mu)])$ .

**Value**

Adjacencies returned in the same form as the input ss

**Author(s)**

Steve Horvath

## References

Bin Zhang and Steve Horvath (2005) "A General Framework for Weighted Gene Co-Expression Network Analysis", *Statistical Applications in Genetics and Molecular Biology*: Vol. 4: No. 1, Article 17

---

signedKME

*Signed eigengene-based connectivity*

---

## Description

Calculation of (signed) eigengene-based connectivity, also known as module membership.

## Usage

```
signedKME(  
  datExpr, datME,  
  outputColumnName = "kME",  
  corFnc = "cor", corOptions = "use = 'p'")
```

## Arguments

datExpr	a data frame containing the gene expression data. Rows correspond to samples and columns to genes. Missing values are allowed and will be ignored.
datME	a data frame containing module eigengenes. Rows correspond to samples and columns to module eigengenes.
outputColumnName	a character string specifying the prefix of column names of the output.
corFnc	character string specifying the function to be used to calculate co-expression similarity. Defaults to Pearson correlation. Any function returning values between -1 and 1 can be used.
corOptions	character string specifying additional arguments to be passed to the function given by corFnc. Use "use = 'p', method = 'spearman'" to obtain Spearman correlation.

## Details

Signed eigengene-based connectivity of a gene in a module is defined as the correlation of the gene with the corresponding module eigengene. The samples in datExpr and datME must be the same.

## Value

A data frame in which rows correspond to input genes and columns to module eigengenes, giving the signed eigengene-based connectivity of each gene with respect to each eigengene.

## Author(s)

Steve Horvath

## References

Dong J, Horvath S (2007) Understanding Network Concepts in Modules, BMC Systems Biology 2007, 1:24

Horvath S, Dong J (2008) Geometric Interpretation of Gene Coexpression Network Analysis. PLoS Comput Biol 4(8): e1000117

---

signumAdjacencyFunction

*Hard-thresholding adjacency function*

---

## Description

This function transforms correlations or other measures of similarity into an unweighted network adjacency.

## Usage

```
signumAdjacencyFunction(corMat, threshold)
```

## Arguments

corMat	a matrix of correlations or other measures of similarity.
threshold	threshold for connecting nodes: all nodes whose corMat is above the threshold will be connected in the resulting network.

## Value

An unweighted adjacency matrix of the same dimensions as the input corMat.

## Author(s)

Steve Horvath

## References

Bin Zhang and Steve Horvath (2005) "A General Framework for Weighted Gene Co-Expression Network Analysis", Statistical Applications in Genetics and Molecular Biology: Vol. 4: No. 1, Article 17

## See Also

[adjacency](#) for soft-thresholding and creating weighted networks.

---

simulateDatExpr                      *Simulation of expression data*

---

### Description

Simulation of expression data with a customizable modular structure and several different types of noise.

### Usage

```
simulateDatExpr(
  eigengenes,
  nGenes,
  modProportions,
  minCor = 0.3,
  maxCor = 1,
  corPower = 1,
  signed = FALSE,
  propNegativeCor = 0.3,
  geneMeans = NULL,
  backgroundNoise = 0.1,
  leaveOut = NULL,
  nSubmoduleLayers = 0,
  nScatteredModuleLayers = 0,
  averageNGenesInSubmodule = 10,
  averageExprInSubmodule = 0.2,
  submoduleSpacing = 2,
  verbose = 1, indent = 0)
```

### Arguments

eigengenes	a data frame containing the seed eigengenes for the simulated modules. Rows correspond to samples and columns to modules.
nGenes	total number of genes to be simulated.
modProportions	a numeric vector with length equal the number of eigengenes in eigengenes plus one, containing fractions of the total number of genes to be put into each of the modules and into the "grey module", which means genes not related to any of the modules. See details.
minCor	minimum correlation of module genes with the corresponding eigengene. See details.
maxCor	maximum correlation of module genes with the corresponding eigengene. See details.
corPower	controls the dropoff of gene-eigengene correlation. See details.
signed	logical: should the genes be simulated as belonging to a signed network? If TRUE, all genes will be simulated to have positive correlation with the eigengene. If FALSE, a proportion given by propNegativeCor will be simulated with negative correlations of the same absolute values.
propNegativeCor	proportion of genes to be simulated with negative gene-eigengene correlations. Only effective if signed is FALSE.

geneMeans	optional vector of length nGenes giving desired mean expression for each gene. If not given, the returned expression profiles will have mean zero.
backgroundNoise	amount of background noise to be added to the simulated expression data.
leaveOut	optional specification of modules that should be left out of the simulation, that is their genes will be simulated as unrelated ("grey"). This can be useful when simulating several sets, in some which a module is present while in others it is absent.
nSubmoduleLayers	number of layers of ordered submodules to be added. See details.
nScatteredModuleLayers	number of layers of scattered submodules to be added. See details.
averageNGenesInSubmodule	average number of genes in a submodule. See details.
averageExprInSubmodule	average strength of submodule expression vectors.
submoduleSpacing	a number giving submodule spacing: this multiple of the submodule size will lie between the submodule and the next one.
verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.
indent	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

## Details

Given eigengenes can be unrelated or they can exhibit non-trivial correlations. Each module is simulated separately from others. The expression profiles are chosen such that their correlations with the eigengene run from just below `maxCor` to `minCor` (hence `minCor` must be between 0 and 1, not including the bounds). The parameter `corPower` can be chosen to control the behaviour of the simulated correlation with the gene index; values higher than 1 will result in the correlation approaching `minCor` faster and lower than 1 slower.

Numbers of genes in each module are specified (as fractions of the total number of genes `nGenes`) by `modProportions`. The last entry in `modProportions` corresponds to the genes that will be simulated as unrelated to anything else ("grey" genes). The proportion must add up to 1 or less. If the sum is less than one, the remaining genes will be partitioned into groups and simulated to be "close" to the proper modules, that is with small but non-zero correlations (between `minCor` and 0) with the module eigengene.

If `signed` is set `FALSE`, the correlation for some of the module genes is chosen negative (but the absolute values remain the same as they would be for positively correlated genes). To ensure consistency for simulations of multiple sets, the indices of the negatively correlated genes are fixed and distributed evenly.

In addition to the primary module structure, a secondary structure can be optionally simulated. Modules in the secondary structure have sizes chosen from an exponential distribution with mean equal `averageNGenesInSubmodule`. Expression vectors simulated in the secondary structure are simulated with expected standard deviation chosen from an exponential distribution with mean equal `averageExprInSubmodule`; the higher this coefficient, the more pronounced will the submodules be in the main modules. The secondary structure can be simulated in several layers; their number is given by `SubmoduleLayers`. Genes in these submodules are ordered in the same order as in the main modules.

In addition to the ordered submodule structure, a scattered submodule structure can be simulated as well. This structure can be viewed as noise that tends to correlate random groups of genes. The size and effect parameters are the same as for the ordered submodules, and the number of layers added is controlled by `nScatteredModuleLayers`.

### Value

A list with the following components:

<code>datExpr</code>	simulated expression data in a data frame whose columns correspond genes and rows to samples.
<code>setLabels</code>	simulated module assignment. Module labels are numeric, starting from 1. Genes simulated to be outside of proper modules have label 0. Modules that are left out (specified in <code>leaveOut</code> ) are indicated as 0 here.
<code>allLabels</code>	simulated module assignment. Genes that belong to leftout modules (specified in <code>leaveOut</code> ) are indicated by their would-be assignment here.
<code>labelOrder</code>	a vector specifying the order in which labels correspond to the given eigengenes, that is <code>labelOrder[1]</code> is the label assigned to module whose seed is <code>eigengenes[, 1]</code> etc.

### Author(s)

Peter Langfelder

### References

A short description of the simulation method can also be found in the Supplementary Material to the article

Langfelder P, Horvath S (2007) Eigengene networks for studying the relationships between co-expression modules. *BMC Systems Biology* 2007, 1:54.

The material is posted at <http://www.genetics.ucla.edu/labs/horvath/CoexpressionNetwork/EigengeneNetwork/Supplemental>

### See Also

[simulateEigengeneNetwork](#) for a simulation of eigengenes with a given causal structure;

[simulateModule](#) for simulations of individual modules;

[simulateDatExpr5Modules](#) for a simplified interface to expression simulations;

[simulateMultiExpr](#) for a simulation of several related data sets.

---

`simulateDatExpr5Modules`

*Simplified simulation of expression data*

---

### Description

This function provides a simplified interface to the expression data simulation, at the cost of considerably less flexibility.

**Usage**

```
simulateDatExpr5Modules(
  nGenes = 2000,
  colorLabels = c("turquoise", "blue", "brown", "yellow", "green"),
  simulateProportions = c(0.1, 0.08, 0.06, 0.04, 0.02),
  METurquoise, MEblue, MEbrown, MEyellow, MEgreen,
  SDnoise = 1, backgroundCor = 0.3)
```

**Arguments**

nGenes	total number of genes to be simulated.
colorLabels	labels for simulated modules.
simulateProportions	a vector of length 5 giving proportions of the total number of genes to be placed in each individual module. The entries must be positive and sum to at most 1. If the sum is less than 1, the leftover genes will be simulated outside of modules.
MEturquoise	seed module eigengene for the first module.
MEblue	seed module eigengene for the second module.
MEbrown	seed module eigengene for the third module.
MEyellow	seed module eigengene for the fourth module.
MEgreen	seed module eigengene for the fifth module.
SDnoise	level of noise to be added to the simulated expressions.
backgroundCor	background correlation. If non-zero, a component will be added to all genes such that the average correlation of otherwise unrelated genes will be backgroundCor.

**Details**

Roughly one-third of the genes are simulated with a negative correlation to their seed eigengene. See the functions [simulateModule](#) and [simulateDatExpr](#) for more details.

**Value**

A list with the following components:

datExpr	the simulated expression data in a data frame, with rows corresponding to samples and columns to genes.
truemodule	a vector with one entry per gene containing the simulated module membership.
datME	a data frame containing a copy of the input module eigengenes.

**Author(s)**

Steve Horvath and Peter Langfelder

**See Also**

[simulateModule](#) for simulation of individual modules;  
[simulateDatExpr](#) for a more comprehensive data simulation interface.



---

`simulateEigengeneNetwork`*Simulate eigengene network from a causal model*

---

**Description**

Simulates a set of eigengenes (vectors) from a given set of causal anchors and a causal matrix.

**Usage**

```
simulateEigengeneNetwork(  
  causeMat,  
  anchorIndex, anchorVectors,  
  noise = 1,  
  verbose = 0, indent = 0)
```

**Arguments**

<code>causeMat</code>	causal matrix. The entry $[i, j]$ is the influence (path coefficient) of vector $j$ on vector $i$ .
<code>anchorIndex</code>	specifies the indices of the anchor vectors.
<code>anchorVectors</code>	a matrix giving the actual anchor vectors as columns. Their number must equal the length of <code>anchorIndex</code> .
<code>noise</code>	standard deviation of the noise added to each simulated vector.
<code>verbose</code>	level of verbosity. 0 means silent.
<code>indent</code>	indentation for diagnostic messages. Zero means no indentation; each unit adds two spaces.

**Details**

The algorithm starts with the anchor vectors and iteratively generates the rest from the path coefficients given in the matrix `causeMat`.

**Value**

A list with the following components:

<code>eigengenes</code>	generated eigengenes.
<code>causeMat</code>	a copy of the input causal matrix
<code>levels</code>	useful for debugging. A vector with one entry for each eigengene giving the number of generations of parents of the eigengene. Anchors have level 0, their direct causal children have level 1 etc.
<code>anchorIndex</code>	a copy of the input <code>anchorIndex</code> .

**Author(s)**

Peter Langfelder

---

simulateModule                      *Simulate a gene co-expression module*

---

### Description

Simulation of a single gene co-expression module.

### Usage

```
simulateModule(
  ME,
  nGenes,
  nNearGenes = 0,
  minCor = 0.3, maxCor = 1, corPower = 1,
  signed = FALSE, propNegativeCor = 0.3,
  geneMeans = NULL,
  verbose = 0, indent = 0)
```

### Arguments

ME	seed module eigengene.
nGenes	number of genes in the module to be simulated. Must be non-zero.
nNearGenes	number of genes to be simulated with low correlation with the seed eigengene.
minCor	minimum correlation of module genes with the eigengene. See details.
maxCor	maximum correlation of module genes with the eigengene. See details.
corPower	controls the dropoff of gene-eigengene correlation. See details.
signed	logical: should the genes be simulated as belonging to a signed network? If TRUE, all genes will be simulated to have positive correlation with the eigengene. If FALSE, a proportion given by propNegativeCor will be simulated with negative correlations of the same absolute values.
propNegativeCor	proportion of genes to be simulated with negative gene-eigengene correlations. Only effective if signed is FALSE.
geneMeans	optional vector of length nGenes giving desired mean expression for each gene. If not given, the returned expression profiles will have mean zero.
verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.
indent	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

### Details

Module genes are simulated around the eigengene by choosing them such that their (expected) correlations with the seed eigengene decrease progressively from (just below) maxCor to minCor. The genes are otherwise independent from one another. The variable corPower determines how fast the correlation drops towards minCor. Higher powers lead to a faster drop-off; corPower must be above zero but need not be integer.

If signed is FALSE, the genes are simulated so as to be part of an unsigned network module, that is some genes will be simulated with a negative correlation with the seed eigengene (but of the same absolute value that a positively correlated gene would be simulated with). The proportion of genes with negative correlation is controlled by propNegativeCor.

Optionally, the function can also simulate genes that are "near" the module, meaning they are simulated with a low but non-zero correlation with the seed eigengene. The correlations run between minCor and zero.

### Value

A matrix containing the expression data with rows corresponding to samples and columns to genes.

### Author(s)

Peter Langfelder

### References

A short description of the simulation method can also be found in the Supplementary Material to the article

Langfelder P, Horvath S (2007) Eigengene networks for studying the relationships between co-expression modules. BMC Systems Biology 2007, 1:54.

The material is posted at <http://www.genetics.ucla.edu/labs/horvath/CoexpressionNetwork/EigengeneNetwork/Supplemental>

### See Also

[simulateEigengeneNetwork](#) for a simulation of eigengenes with a given causal structure;

[simulateDatExpr](#) for simulations of whole datasets consisting of multiple modules;

[simulateDatExpr5Modules](#) for a simplified interface to expression simulations;

[simulateMultiExpr](#) for a simulation of several related data sets.

---

simulateMultiExpr

*Simulate multi-set expression data*

---

### Description

Simulation of expression data in several sets with relate module structure.

### Usage

```
simulateMultiExpr(eigengenes,  
                  nGenes,  
                  modProportions,  
                  minCor = 0.5, maxCor = 1,  
                  corPower = 1,  
                  backgroundNoise = 0.1,  
                  leaveOut = NULL,  
                  signed = FALSE,  
                  propNegativeCor = 0.3,  
                  geneMeans = NULL,
```

```

nSubmoduleLayers = 0,
nScatteredModuleLayers = 0,
averageNGenesInSubmodule = 10,
averageExprInSubmodule = 0.2,
submoduleSpacing = 2,
verbose = 1, indent = 0)

```

## Arguments

eigengenes	the seed eigengenes for the simulated modules in a multi-set format. A list with one component per set. Each component is again a list that must contain a component data. This is a data frame of seed eigengenes for the corresponding data set. Columns correspond to modules, rows to samples. Number of samples in the simulated data is determined from the number of samples of the eigengenes.
nGenes	integer specifying the number of simulated genes.
modProportions	a numeric vector with length equal the number of eigengenes in eigengenes plus one, containing fractions of the total number of genes to be put into each of the modules and into the "grey module", which means genes not related to any of the modules. See details.
minCor	minimum correlation of module genes with the corresponding eigengene. See details.
maxCor	maximum correlation of module genes with the corresponding eigengene. See details.
corPower	controls the dropoff of gene-eigengene correlation. See details.
backgroundNoise	amount of background noise to be added to the simulated expression data.
leaveOut	optional specification of modules that should be left out of the simulation, that is their genes will be simulated as unrelated ("grey"). A logical matrix in which columns correspond to sets and rows to modules. Wherever TRUE, the corresponding module in the corresponding data set will not be simulated, that is its genes will be simulated independently of the eigengene.
signed	logical: should the genes be simulated as belonging to a signed network? If TRUE, all genes will be simulated to have positive correlation with the eigengene. If FALSE, a proportion given by propNegativeCor will be simulated with negative correlations of the same absolute values.
propNegativeCor	proportion of genes to be simulated with negative gene-eigengene correlations. Only effective if signed is FALSE.
geneMeans	optional vector of length nGenes giving desired mean expression for each gene. If not given, the returned expression profiles will have mean zero.
nSubmoduleLayers	number of layers of ordered submodules to be added. See details.
nScatteredModuleLayers	number of layers of scattered submodules to be added. See details.
averageNGenesInSubmodule	average number of genes in a submodule. See details.
averageExprInSubmodule	average strength of submodule expression vectors.

submoduleSpacing	a number giving submodule spacing: this multiple of the submodule size will lie between the submodule and the next one.
verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.
indent	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

## Details

For details of simulation of individual data sets and the meaning of individual set simulation arguments, see [simulateDatExpr](#). This function simulates several data sets at a time and puts the result in a multi-set format. The number of genes is the same for all data sets. Module memberships are also the same, but modules can optionally be “dissolved”, that is their genes will be simulated as unassigned. Such “dissolved”, or left out, modules can be specified in the matrix `leaveOut`.

## Value

A list with the following components:

<code>multiExpr</code>	simulated expression data in multi-set format analogous to that of the input <code>eigengenes</code> . A list with one component per set. Each component is again a list that must contain a component data. This is a data frame of expression data for the corresponding data set. Columns correspond to genes, rows to samples.
<code>setLabels</code>	a matrix of dimensions (number of genes) times (number of sets) that contains module labels for each genes in each simulated data set.
<code>allLabels</code>	a matrix of dimensions (number of genes) times (number of sets) that contains the module labels that would be simulated if no module were left out using <code>leaveOut</code> . This means that all columns of the matrix are equal; the columns are repeated for convenience so <code>allLabels</code> has the same dimensions as <code>setLabels</code> .
<code>labelOrder</code>	a matrix of dimensions (number of modules) times (number of sets) that contains the order in which module labels were assigned to genes in each set. The first label is assigned to genes 1...(module size of module labeled by first label), the second label to the following batch of genes etc.

## Author(s)

Peter Langfelder

## References

A short description of the simulation method can also be found in the Supplementary Material to the article

Langfelder P, Horvath S (2007) Eigengene networks for studying the relationships between co-expression modules. *BMC Systems Biology* 2007, 1:54.

The material is posted at <http://www.genetics.ucla.edu/labs/horvath/CoexpressionNetwork/EigengeneNetwork/Supplemental>

**See Also**

[simulateEigengeneNetwork](#) for a simulation of eigengenes with a given causal structure;  
[simulateDatExpr](#) for simulation of individual data sets;  
[simulateDatExpr5Modules](#) for a simple simulation of a data set consisting of 5 modules;  
[simulateModule](#) for simulations of individual modules;

---

simulateSmallLayer      *Simulate small modules*

---

**Description**

This function simulates a set of small modules. The primary purpose is to add a submodule structure to the main module structure simulated by [simulateDatExpr](#).

**Usage**

```
simulateSmallLayer(  
  order,  
  nSamples,  
  minCor = 0.3, maxCor = 0.5, corPower = 1,  
  averageModuleSize,  
  averageExpr,  
  moduleSpacing,  
  verbose = 4, indent = 0)
```

**Arguments**

order	a vector giving the simulation order for vectors. See details.
nSamples	integer giving the number of samples to be simulated.
minCor	a multiple of maxCor (see below) giving the minimum correlation of module genes with the corresponding eigengene. See details.
maxCor	maximum correlation of module genes with the corresponding eigengene. See details.
corPower	controls the dropoff of gene-eigengene correlation. See details.
averageModuleSize	average number of genes in a module. See details.
averageExpr	average strength of module expression vectors.
moduleSpacing	a number giving module spacing: this multiple of the module size will lie between the module and the next one.
verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.
indent	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

**Details**

Module eigenvectors are chosen randomly and independently. Module sizes are chosen randomly from an exponential distribution with mean equal averageModuleSize. Two thirds of genes in each module are simulated as proper module genes and one third as near-module genes (see [simulateModule](#) for details). Between each successive pairs of modules a number of genes given by moduleSpacing will be left unsimulated (zero expression). Module expression, that is the expected standard deviation of the module expression vectors, is chosen randomly from an exponential distribution with mean equal averageExpr. The expression profiles are chosen such that their correlations with the eigengene run from just below maxCor to minCor \* maxCor (hence minCor must be between 0 and 1, not including the bounds). The parameter corPower can be chosen to control the behaviour of the simulated correlation with the gene index; values higher than 1 will result in the correlation approaching minCor \* maxCor faster and lower than 1 slower.

The simulated genes will be returned in the order given in order.

**Value**

A matrix of simulated gene expressions, with dimension (nSamples, length(order)).

**Author(s)**

Peter Langfelder

**See Also**

[simulateModule](#) for simulation of individual modules;

[simulateDatExpr](#) for the main gene expression simulation function.

---

sizeGrWindow

*Opens a graphics window with specified dimensions*

---

**Description**

If a graphic device window is already open, it is closed and re-opened with specified dimensions (in inches); otherwise a new window is opened.

**Usage**

```
sizeGrWindow(width, height)
```

**Arguments**

width                   desired width of the window, in inches.

height                   desired heigh of the window, in inches.

**Value**

None.

**Author(s)**

Peter Langfelder

---

softConnectivity      *Calculates connectivity of a weighted network.*

---

### Description

Given expression data or a similarity, the function constructs the adjacency matrix and for each node calculates its connectivity, that is the sum of the adjacency to the other nodes.

### Usage

```
softConnectivity(
  datExpr,
  corFnc = "cor", corOptions = "use = 'p'",
  type = "unsigned",
  power = if (type == "signed") 15 else 6,
  blockSize = 1500,
  minNSamples = NULL,
  verbose = 2, indent = 0)

softConnectivity.fromSimilarity(
  similarity,
  type = "unsigned",
  power = if (type == "signed") 15 else 6,
  blockSize = 1500,
  verbose = 2, indent = 0)
```

### Arguments

datExpr	a data frame containing the expression data, with rows corresponding to samples and columns to genes.
similarity	a similarity matrix: a square symmetric matrix with entries between -1 and 1.
corFnc	character string giving the correlation function to be used for the adjacency calculation. Recommended choices are "cor" and "bicor", but other functions can be used as well.
corOptions	character string giving further options to be passed to the correlation function.
type	network type. Allowed values are (unique abbreviations of) "unsigned", "signed", "signed hybrid".
power	soft thresholding power.
blockSize	block size in which adjacency is to be calculated. Too low (say below 100) may make the calculation inefficient, while too high may cause R to run out of physical memory and slow down the computer. Should be chosen such that an array of doubles of size (number of genes) * (block size) fits into available physical memory.
minNSamples	minimum number of samples available for the calculation of adjacency for the adjacency to be considered valid. If not given, defaults to the greater of <code>..minNSamples</code> (currently 4) and number of samples divided by 3. If the number of samples falls below this threshold, the connectivity of the corresponding gene will be returned as NA.



verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.
indent	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

**Value**

A vector with one entry per gene giving the connectivity of each gene in the weighted network.

**Author(s)**

Steve Horvath

**References**

Bin Zhang and Steve Horvath (2005) "A General Framework for Weighted Gene Co-Expression Network Analysis", *Statistical Applications in Genetics and Molecular Biology*: Vol. 4: No. 1, Article 17

**See Also**

[adjacency](#)

---

spaste	<i>Space-less paste</i>
--------	-------------------------

---

**Description**

A convenient wrapper for the [paste](#) function with `sep=""`.

**Usage**

```
spaste(...)
```

**Arguments**

... standard arguments to function [paste](#) except `sep`.

**Value**

The result of the corresponding [paste](#).

**Note**

Do not use the `sep` argument. Using will lead to an error.

**Author(s)**

Peter Langfelder

**See Also**

[paste](#)

**Examples**

```
a = 1;
paste("a=", a);
spaste("a=", a);
```

---

standardColors	<i>Colors this library uses for labeling modules.</i>
----------------	---

---

**Description**

Returns the vector of color names in the order they are assigned by other functions in this library.

**Usage**

```
standardColors(n = NULL)
```

**Arguments**

**n** Number of colors requested. If NULL, all (approx. 450) colors will be returned. Any other invalid argument such as less than one or more than maximum (`length(standardColors())`) will trigger an error.

**Value**

A vector of character color names of the requested length.

**Author(s)**

Peter Langfelder, <Peter.Langfelder@gmail.com>

**Examples**

```
standardColors(10);
```

---

standardScreeningBinaryTrait	<i>Standard screening for binary traits</i>
------------------------------	---

---

**Description**

The function `standardScreeningBinaryTrait` computes widely used statistics for relating the columns of the input data frame (argument `datE`) to a binary sample trait (argument `y`). The statistics include Student t-test p-value and the corresponding local false discovery rate (known as q-value, Storey et al 2004), the fold change, the area under the ROC curve (also known as C-index), mean values etc. If the input option `KruskalTest` is set to `TRUE`, it also computes the Kruskal Wallist test p-value and corresponding q-value. The Kruskal Wallis test is a non-parametric, rank-based group comparison test.

**Usage**

```
standardScreeningBinaryTrait(
  datExpr, y,
  corFnc = cor, corOptions = list(use = 'p'),
  kruskalTest = FALSE, qValues = FALSE,
  var.equal=FALSE, na.action="na.exclude",
  getAreaUnderROC = TRUE)
```

**Arguments**

datExpr	a data frame or matrix whose columns will be related to the binary trait
y	a binary vector whose length (number of components) equals the number of rows of datE
corFnc	correlation function. Defaults to Pearson correlation.
corOptions	a list specifying options to corFnc. An empty list must be specified as list() (supplying NULL instead will trigger an error).
kruskalTest	logical: should the Kruskal test be performed?
qValues	logical: should the q-values be calculated?
var.equal	logical input parameter for the Student t-test. It indicates whether to treat the two variances (corresponding to the binary grouping) are being equal. If TRUE then the pooled variance is used to estimate the variance otherwise the Welch (or Satterthwaite) approximation to the degrees of freedom is used. Warning: here the default value is TRUE which is different from the default value of t.test. Type help(t.test) for more details.
na.action	character string for the Student t-test: indicates what should happen when the data contain missing values NAs.
getAreaUnderROC	logical: should area under the ROC curve be calculated? The calculation slows the function down somewhat.

**Value**

A data frame whose rows correspond to the columns of datE and whose columns report

ID	column names of the input datExpr.
corPearson	pearson correlation with a binary numeric version of the input variable. The numeric variable equals 1 for level 1 and 2 for level 2. The levels are given by levels(factor(y)).
t.Student	Student's t-test statistic
pvalueStudent	two-sided Student t-test p-value.
qvalueStudent	(if input qValues==TRUE) q-value (local false discovery rate) based on the Student T-test p-value (Storey et al 2004).
foldChange	a (signed) ratio of mean values. If the mean in the first group (corresponding to level 1) is larger than that of the second group, it equals meanFirstGroup/meanSecondGroup. But if the mean of the second group is larger than that of the first group it equals -meanSecondGroup/meanFirstGroup (notice the minus sign).
meanFirstGroup	means of columns in input datExpr across samples in the first group.

meanSecondGroup means of columns in input datExpr across samples in the second group.

SE.FirstGroup standard errors of columns in input datExpr across samples in the first group. Recall that  $SE(x)=\sqrt{\text{var}(x)/n}$  where n is the number of non-missing values of x.

SE.SecondGroup standard errors of columns in input datExpr across samples in the second group.

areaUnderROC the area under the ROC, also known as the concordance index or C.index. This is a measure of discriminatory power. The measure lies between 0 and 1 where 0.5 indicates no discriminatory power. 0 indicates that the "opposite" predictor has perfect discriminatory power. To compute it we use the function [rcorr.cens](#) with outx=TRUE (from Frank Harrel's package Hmisc). Only present if input getAreUnderROC is TRUE.

nPresentSamples number of samples with finite measurements for each gene.

If input kruskalTest is TRUE, the following columns further summarize results of Kruskal-Wallis test:

stat.Kruskal Kruskal-Wallis test statistic.

stat.Kruskal.signed (Warning: experimental) Kruskal-Wallis test statistic including a sign that indicates whether the average rank is higher in second group (positive) or first group (negative).

pvaluekruskal Kruskal-Wallis test p-values.

qkruskal q-values corresponding to the Kruskal-Wallis test p-value (if input qValues==TRUE).

**Author(s)**

Steve Horvath

**References**

Storey JD, Taylor JE, and Siegmund D. (2004) Strong control, conservative point estimation, and simultaneous conservative consistency of false discovery rates: A unified approach. *Journal of the Royal Statistical Society, Series B*, 66: 187-205.

**Examples**

```
m=50
y=sample(c(1,2),m,replace=TRUE)
datExprSignal=simulateModule(scale(y),30)
datExprNoise=simulateModule(rnorm(m),150)
datExpr=data.frame(datExprSignal,datExprNoise)

Result1=standardScreeningBinaryTrait(datExpr,y)
Result1[1:5,]

# use unequal variances and calculate q-values
Result2=standardScreeningBinaryTrait(datExpr,y, var.equal=FALSE,qValue=TRUE)
Result2[1:5,]

# calculate Kruskal Wallis test and q-values
```

```
Result3=standardScreeningBinaryTrait(datExpr,y,kruskalTest=TRUE,qValue=TRUE)
Result3[1:5,]
```

---

standardScreeningCensoredTime

*Standard Screening with regard to a Censored Time Variable*

---

## Description

The function `standardScreeningCensoredTime` computes association measures between the columns of the input data `datE` and a censored time variable (e.g. survival time). The censored time is specified using two input variables "time" and "event". The event variable is binary where 1 indicates that the event took place (e.g. the person died) and 0 indicates censored (i.e. lost to follow up). The function fits univariate Cox regression models (one for each column of `datE`) and outputs a Wald test p-value, a logrank p-value, corresponding local false discovery rates (known as q-values, Storey et al 2004), hazard ratios. Further it reports the concordance index (also known as area under the ROC curve) and optionally results from dichotomizing the columns of `datE`.

## Usage

```
standardScreeningCensoredTime(
  time,
  event,
  datExpr,
  percentiles = seq(from = 0.1, to = 0.9, by = 0.2),
  dichotomizationResults = FALSE,
  qValues = TRUE,
  fastCalculation = TRUE)
```

## Arguments

<code>time</code>	numeric variable showing time to event or time to last follow up.
<code>event</code>	Input variable <code>time</code> specifies the time to event or time to last follow up. Input variable <code>event</code> indicates whether the event happened (=1) or whether there was censoring (=0).
<code>datExpr</code>	a data frame or matrix whose columns will be related to the censored time.
<code>percentiles</code>	numeric vector which is only used when <code>dichotomizationResults=T</code> . Each value should lie between 0 and 1. For each value specified in the vector <code>percentiles</code> , a binary vector will be defined by dichotomizing the column value according to the corresponding quantile. Next a corresponding p-value will be calculated.
<code>dichotomizationResults</code>	logical. If this option is set to <code>TRUE</code> then the values of the columns of <code>datE</code> will be dichotomized and corresponding Cox regression p-values will be calculated.
<code>qValues</code>	logical. If this option is set to <code>TRUE</code> (default) then q-values will be calculated for the Cox regression p-values.
<code>fastCalculation</code>	logical. If set to <code>TRUE</code> , the function outputs correlation test p-values (and q-values) for correlating the columns of <code>datE</code> with the expected hazard (if no covariate is fit). Specifically, the expected hazard is defined as the deviance

residual of an intercept only Cox regression model. The results are very similar to those resulting from a univariate Cox model where the censored time is regressed on the columns of `dat`. Specifically, this computational speed up is facilitated by the insight that the p-values resulting from a univariate Cox regression `coxph(Surv(time,event)~datE[,i])` are very similar to those from `corPvalueFisher(cor(devianceResidual,datE[,i]), nSamples)`.

## Details

If input option `fastCalculation=TRUE`, then the function outputs correlation test p-values (and q-values) for correlating the columns of `datE` with the expected hazard (if no covariate is fit). Specifically, the expected hazard is defined as the deviance residual of an intercept only Cox regression model. The results are very similar to those resulting from a univariate Cox model where the censored time is regressed on the columns of `dat`. Specifically, this computational speed up is facilitated by the insight that the p-values resulting from a univariate Cox regression `coxph(Surv(time,event)~datE[,i])` are very similar to those from `corPvalueFisher(cor(devianceResidual,datE[,i]), nSamples)`

## Value

If `fastCalculation` is `FALSE`, the function outputs a data frame whose rows correspond to the columns of `datE` and whose columns report

<code>ID</code>	column names of the input data <code>datExpr</code> .
<code>pvalueWald</code>	Wald test p-value from fitting a univariate Cox regression model where the censored time is regressed on each column of <code>datExpr</code> .
<code>qValueWald</code>	local false discovery rate (q-value) corresponding to the Wald test p-value.
<code>pvalueLogrank</code>	Logrank p-value resulting from the Cox regression model. Also known as score test p-value. For large sample sizes this should be similar to the Wald test p-value.
<code>qValueLogrank</code>	local false discovery rate (q-value) corresponding to the Logrank test p-value.
<code>HazardRatio</code>	hazard ratio resulting from the Cox model. If the value is larger than 1, then high values of the column are associated with shorter time, e.g. increased hazard of death. A hazard ratio equal to 1 means no relationship between the column and time. $HR < 1$ means that high values are associated with longer time, i.e. lower hazard.
<code>CI.LowerLimitHR</code>	Lower bound of the 95 percent confidence interval of the hazard ratio.
<code>CI.UpperLimitHR</code>	Upper bound of the 95 percent confidence interval of the hazard ratio.
<code>C.index</code>	concordance index, also known as C-index or area under the ROC curve. Calculated with the <code>rcorr.cens</code> option <code>outx=TRUE</code> (ties are ignored).
<code>MinimumDichotPvalue</code>	This is the smallest p-value from the dichotomization results. To see which dichotomized variable (and percentile) corresponds to the minimum, study the following columns.
<code>pValueDichot0.1</code>	This columns report the p-value when the column is dichotomized according to the specified percentile (here 0.1). The percentiles are specified in the input option <code>percentiles</code> .

- `pvalueDeviance` The p-value resulting from using a correlation test to relate the expected hazard (deviance residual) with each (undichotomized) column of `datE`. Specifically, the Fisher transformation is used to calculate the p-value for the Pearson correlation. The resulting p-value should be very similar to that of a univariate Cox regression model.
- `qvalueDeviance` Local false discovery rate (q-value) corresponding to `pvalueDeviance`.
- `corDeviance` Pearson correlation between the expected hazard (deviance residual) with each (undichotomized) column of `datExpr`.

**Author(s)**

Steve Horvath

---

`standardScreeningNumericTrait`
*Standard screening for numeric traits***Description**

Standard screening for numeric traits based on Pearson correlation.

**Usage**

```
standardScreeningNumericTrait(datExpr, yNumeric, corFnc = cor,
                              corOptions = list(use = 'p'),
                              alternative = c("two.sided", "less", "greater"),
                              qValues = TRUE,
                              areaUnderROC = TRUE)
```

**Arguments**

- `datExpr` data frame containing expression data (or more generally variables to be screened), with rows corresponding to samples and columns to genes (variables)
- `yNumeric` a numeric vector giving the trait measurements for each sample
- `corFnc` correlation function. Defaults to Pearson correlation but can also be [bicor](#).
- `corOptions` list specifying additional arguments to be passed to the correlation function given by `corFnc`.
- `alternative` alternative hypothesis for the correlation test
- `qValues` logical: should q-values be calculated?
- `areaUnderROC` logical: should area under the receiver-operating curve be calculated?

**Details**

The function calculates the correlations, associated p-values, area under the ROC, and q-values

**Value**

Data frame with the following components:

ID	Gene (or variable) identifiers copied from <code>colnames(datExpr)</code>
cor	correlations of all genes with the trait
Z	Fisher Z statistics corresponding to the correlations
pvalueStudent	Student p-values of the correlations
qvalueStudent	(if input <code>qValues==TRUE</code> ) q-values of the correlations calculated from the p-values
AreaUnderROC	(if input <code>areaUnderROC==TRUE</code> ) area under the ROC
nPresentSamples	number of samples present for the calculation of each association.

**Author(s)**

Steve Horvath

**See Also**

[standardScreeningBinaryTrait](#), [standardScreeningCensoredTime](#)

---

stat.bwss

*Between and Within Group Sum of Squares Calculation*

---

**Description**

This function computes the between and within group sum of squares for each row of a matrix which may have NAs.

**Usage**

```
stat.bwss(x, cl)
```

**Arguments**

x	a matrix, NAs allowed. In the microarray context, the rows of X could correspond to genes and the columns to different hybridizations.
cl	column labels, must be consecutive integers.

**Value**

List containing the following components

wn	matrix with class sizes for each row of X;
BW	vector of BSS/WSS for each row of X;
BSS	between group sum of squares for each row of X;
WSS	within group sum of squares for each row of X;
TSS	total sum of squares for each row of X;
tvar	variance for each row of X.



**Author(s)**

Sandrine Dudoit, <sandrine@stat.berkeley.edu>  
Jane Fridlyand, <janef@stat.berkeley.edu>

**References**

S. Dudoit, J. Fridlyand, and T. P. Speed. Comparison of Discrimination Methods for the Classification of Tumors Using Gene Expression Data. June 2000. (Statistics, UC Berkeley, Tech Report # 576).

**See Also**

[var.na](#), [var](#), [apply](#).

---

stat.diag.da

*Diagonal Discriminant Analysis*

---

**Description**

This function implements a simple Gaussian maximum likelihood discriminant rule, for diagonal class covariance matrices.

**Usage**

```
stat.diag.da(ls, cll, ts, pool=1)
```

**Arguments**

ls	learning set data matrix, with rows corresponding to cases (i.e., mRNA samples) and columns to predictor variables (i.e., genes).
cll	class labels for learning set, must be consecutive integers.
ts	test set data matrix, with rows corresponding to cases and columns to predictor variables.
pool	logical flag. If pool=1, the covariance matrices are assumed to be constant across classes and the discriminant rule is linear in the data. If pool=0, the covariance matrices may vary across classes and the discriminant rule is quadratic in the data.

**Value**

List containing the following components

pred	vector of class predictions for the test set.
------	---

**Author(s)**

Sandrine Dudoit, <sandrine@stat.berkeley.edu>  
Jane Fridlyand, <janef@stat.berkeley.edu>

**References**

S. Dudoit, J. Fridlyand, and T. P. Speed. Comparison of Discrimination Methods for the Classification of Tumors Using Gene Expression Data. June 2000. (Statistics, UC Berkeley, Tech Report #576).

---

stdErr	<i>Standard error of the mean of a given vector.</i>
--------	--

---

**Description**

Returns the standard error of the mean of a given vector. Missing values are ignored.

**Usage**

```
stdErr(x)
```

**Arguments**

x                    a numeric vector

**Value**

Standard error of the mean of x.

**Author(s)**

Steve Horvath

---

stratifiedBarplot	<i>Bar plots of data across two splitting parameters</i>
-------------------	--

---

**Description**

This function takes an expression matrix which can be split using two separate splitting parameters (ie, control vs AD with multiple brain regions), and plots the results as a barplot. Group average, standard deviations, and relevant Kruskal-Wallis p-values are returned.

**Usage**

```
stratifiedBarplot(  
  expAll,  
  groups, split, subset,  
  genes = NA,  
  scale = "N", graph = TRUE,  
  las1 = 2, cex1 = 1.5, ...)
```

**Arguments**

expAll	An expression matrix, with rows as samples and genes/probes as columns. If genes=NA, then column names must be included.
groups	A character vector corresponding to the samples in expAll, with each element the group name of the relevant sample or NA for samples not in any group. For, example: NA, NA, NA, Con, Con, Con, Con, AD, AD, AD, AD, NA, NA. This trait will be plotted as adjacent bars for each split.
split	A character vector corresponding to the samples in expAll, with each element the group splitting name of the relevant sample or NA for samples not in any group. For, example: NA, NA, NA, Hip, Hip, EC, EC, Hip, Hip, EC, EC, NA, NA. This trait will be plotted as the same color across each split of the barplot. For the function to work properly, the same split values should be inputted for each group.
subset	A list of one or more genes to compare the expression with. If the list contains more than one gene, the first element contains the group name. For example, Ribosomes, RPL3, RPL4, RPS3.
genes	If entered, this parameter is a list of gene/probe identifiers corresponding to the columns in expAll.
scale	For subsets of genes that include more than one gene, this parameter determines how the genes are combined into a single value. Currently, there are five options: 1) ("N")o scaling (default); 2) first divide each gene by the ("A")verage across samples; 3) first scale genes to ("Z")-score across samples; 4) only take the top ("H")ub gene (ignore all but the highest-connected gene); and 5) take the ("M")odule eigengene. Note that these scaling methods have not been sufficiently tested, and should be considered experimental.
graph	If TRUE (default), bar plot is made. If FALSE, only the results are returned, and no plot is made.
cex1	Sets the graphing parameters of cex.axis and cex.names (default=1.5)
las1	Sets the graphing parameter las (default=2).
...	Other graphing parameters allowed in the barplot function. Note that the parameters for cex.axis, cex.names, and las are superseded by cex1 and las1 and will therefore be ignored.

**Value**

splitGroupMeans	The group/split averaged expression across each group and split combination. This is the height of the bars in the graph.
splitGroupSDs	The standard deviation of group/split expression across each group and split combination. This is the height of the error bars in the graph.
splitPvals	Kruskal-Wallis p-values for each splitting parameter across groups.
groupPvals	Kruskal-Wallis p-values for each group parameter across splits.

**Author(s)**

Jeremy Miller

**See Also**

[barplot](#), [verboseBarplot](#)

**Examples**

```
# Example: first simulate some data
set.seed(100)
ME.A = sample(1:100,50); ME.B = sample(1:100,50)
ME.C = sample(1:100,50); ME.D = sample(1:100,50)
ME1 = data.frame(ME.A, ME.B, ME.C, ME.D)
simDatA = simulateDatExpr(ME1,1000,c(0.2,0.1,0.08,0.05,0.3), signed=TRUE)
datExpr = simDatA$datExpr+5
datExpr[1:10,] = datExpr[1:10,]+2
datExpr[41:50,] = datExpr[41:50,]-1

# Now split up the data and plot it!
subset = c("Random Genes", "Gene.1", "Gene.234", "Gene.56", "Gene.789")
groups = rep(c("A","A","A","B","B","B","C","C","C","C"),5)
split = c(rep("ZZ",10), rep("YY",10), rep("XX",10), rep("WW",10), rep("VV",10))
par(mfrow = c(1,1))
results = stratifiedBarplot(datExpr, groups, split, subset)
results

# Now plot it the other way
results = stratifiedBarplot(datExpr, split, groups, subset)
```

subsetTOM

*Topological overlap for a subset of a whole set of genes***Description**

This function calculates topological overlap of a subset of vectors with respect to a whole data set.

**Usage**

```
subsetTOM(
  datExpr,
  subset,
  corFnc = "cor", corOptions = "use = 'p'",
  networkType = "unsigned",
  power = 6,
  verbose = 1, indent = 0)
```

**Arguments**

datExpr	a data frame containing the expression data of the whole set, with rows corresponding to samples and columns to genes.
subset	a single logical or numeric vector giving the indices of the nodes for which the TOM is to be calculated.
corFnc	character string giving the correlation function to be used for the adjacency calculation. Recommended choices are "cor" and "bicor", but other functions can be used as well.
corOptions	character string giving further options to be passed to the correlation function.

networkType	character string giving network type. Allowed values are (unique abbreviations of) "unsigned", "signed", "signed hybrid". See <a href="#">adjacency</a> .
power	soft-thresholding power for network construction.
verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.
indent	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

### Details

This function is designed to calculate topological overlaps of small subsets of large expression data sets, for example in individual modules.

### Value

A matrix of dimensions  $n \times n$ , where  $n$  is the number of entries selected by block.

### Author(s)

Peter Langfelder

### References

Bin Zhang and Steve Horvath (2005) "A General Framework for Weighted Gene Co-Expression Network Analysis", *Statistical Applications in Genetics and Molecular Biology*: Vol. 4: No. 1, Article 17

### See Also

[TOMsimilarity](#) for standard calculation of topological overlap.

---

swapTwoBranches	<i>Select, swap, or reflect branches in a dendrogram.</i>
-----------------	---

---

### Description

swapTwoBranches takes a gene tree object and two genes as input, and swaps the branches containing these two genes at the nearest branch point of the dendrogram.

reflectBranch takes a gene tree object and two genes as input, and reflects the branch containing the first gene at the nearest branch point of the dendrogram.

selectBranch takes a gene tree object and two genes as input, and outputs indices for all genes in the branch containing the first gene, up to the nearest branch point of the dendrogram.

### Usage

```
swapTwoBranches(hierTOM, g1, g2)
reflectBranch(hierTOM, g1, g2, both = FALSE)
selectBranch(hierTOM, g1, g2)
```

**Arguments**

hierTOM	A hierarchical clustering object (or gene tree) that is used to plot the dendrogram. For example, the output object from the function <code>hclust</code> or <code>flashClust</code> . Note that elements of <code>hierTOM\$order</code> MUST be named (for example, with the corresponding gene name).
g1	Any gene in the branch of interest.
g2	Any gene in a branch directly adjacent to the branch of interest.
both	Logical: should the selection include the branch gene <code>g2</code> ?

**Value**

`swapTwoBranches` and `reflectBranch` return a hierarchical clustering object with the `hierTOM$order` variable properly adjusted, but all other variables identical as the `hierTOM` input.

`selectBranch` returns a numeric vector corresponding to all genes in the requested branch.

**Author(s)**

Jeremy Miller

**Examples**

```
## Example: first simulate some data.

MEturquoise = sample(1:100,50)
MEblue      = c(MEturquoise[1:25], sample(1:100,25))
MEbrown     = sample(1:100,50)
MEyellow    = sample(1:100,50)
MEgreen     = c(MEyellow[1:30], sample(1:100,20))
MERed      = c(MEbrown [1:20], sample(1:100,30))
ME         = data.frame(MEturquoise, MEblue, MEbrown, MEyellow, MEgreen, MERed)
dat1       = simulateDatExpr(ME,400 ,c(0.16,0.12,0.11,0.10,0.10,0.09,0.15),
                             signed=TRUE)

TOM1      = TOMsimilarityFromExpr(dat1$datExpr, networkType="signed")
colnames(TOM1) <- rownames(TOM1) <- colnames(dat1$datExpr)
tree1     = flashClust(as.dist(1-TOM1),method="average")
colorh    = labels2colors(dat1$allLabels)

plotDendroAndColors(tree1,colorh,dendroLabels=FALSE)

## Reassign modules using the selectBranch and chooseOneHubInEachModule functions

datExpr = dat1$datExpr
hubs    = chooseOneHubInEachModule(datExpr, colorh)
colorh2 = rep("grey", length(colorh))
colorh2 [selectBranch(tree1,hubs["blue"],hubs["turquoise"])] = "blue"
colorh2 [selectBranch(tree1,hubs["turquoise"],hubs["blue"])] = "turquoise"
colorh2 [selectBranch(tree1,hubs["green"],hubs["yellow"])]   = "green"
colorh2 [selectBranch(tree1,hubs["yellow"],hubs["green"])]   = "yellow"
colorh2 [selectBranch(tree1,hubs["red"],hubs["brown"])]      = "red"
colorh2 [selectBranch(tree1,hubs["brown"],hubs["red"])]       = "brown"
plotDendroAndColors(tree1,cbind(colorh,colorh2),c("Old","New"),dendroLabels=FALSE)

## Now swap and reflect some branches, then optimize the order of the branches
```

```

# Open a suitably sized graphics window

sizeGrWindow(12,9);

# partition the screen for 3 dendrogram + module color plots

layout(matrix(c(1:6), 6, 1), heights = c(0.8, 0.2, 0.8, 0.2, 0.8, 0.2));

plotDendroAndColors(tree1,colorh2,dendroLabels=FALSE,main="Starting Dendrogram",
                    setLayout = FALSE)

tree1 = swapTwoBranches(tree1,hubs["red"],hubs["turquoise"])
plotDendroAndColors(tree1,colorh2,dendroLabels=FALSE,main="Swap blue/turquoise and red/brown",
                    setLayout = FALSE)

tree1 = reflectBranch(tree1,hubs["blue"],hubs["green"])
plotDendroAndColors(tree1,colorh2,dendroLabels=FALSE,main="Reflect turquoise/blue",
                    setLayout = FALSE)

```

---

TOMplot

*Graphical representation of the Topological Overlap Matrix*


---

### Description

Graphical representation of the Topological Overlap Matrix using a heatmap plot combined with the corresponding hierarchical clustering dendrogram and module colors.

### Usage

```

TOMplot(
  dissim,
  dendro,
  Colors = NULL,
  ColorsLeft = Colors,
  terrainColors = FALSE,
  setLayout = TRUE,
  ...)

```

### Arguments

dissim	a matrix containing the topological overlap-based dissimilarity
dendro	the corresponding hierarchical clustering dendrogram
Colors	optional specification of module colors to be plotted on top
ColorsLeft	optional specification of module colors on the left side. If NULL, Colors will be used.
terrainColors	logical: should terrain colors be used?
setLayout	logical: should layout be set? If TRUE, standard layout for one plot will be used. Note that this precludes multiple plots on one page. If FALSE, the user is responsible for setting the correct layout.
...	other graphical parameters to <a href="#">heatmap</a> .

**Details**

The standard heatmap function uses the [layout](#) function to set the following layout (when Colors is given):

```
0 0 5
0 0 2
4 1 3
```

To get a meaningful heatmap plot, user-set layout must respect this geometry.

**Value**

None.

**Author(s)**

Steve Horvath and Peter Langfelder

**See Also**

[heatmap](#), the workhorse function doing the plotting.

---

TOMsimilarity

*Topological overlap matrix similarity and dissimilarity*

---

**Description**

Calculation of the topological overlap matrix from a given adjacency matrix.

**Usage**

```
TOMsimilarity(adjMat, TOMType = "unsigned", TOMDenom = "min", verbose = 1, indent = 0)
TOMdist(adjMat, TOMType = "unsigned", TOMDenom = "min", verbose = 1, indent = 0)
```

**Arguments**

adjMat	adjacency matrix, that is a square, symmetric matrix with entries between 0 and 1 (negative values are allowed if TOMType=="signed").
TOMType	a character string specifying TOM type to be calculated. One of "unsigned", "signed". If "unsigned", the standard TOM will be used (more generally, TOM function will receive the adjacency as input). If "signed", TOM will keep track of the sign of the adjacency between neighbors.
TOMDenom	a character string specifying the TOM variant to be used. Recognized values are "min" giving the standard TOM described in Zhang and Horvath (2005), and "mean" in which the min function in the denominator is replaced by mean. The "mean" may produce better results but at this time should be considered experimental.
verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.
indent	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.



## Details

The functions perform basically the same calculations of topological overlap. TOMdist turns the overlap (which is a measure of similarity) into a measure of dissimilarity by subtracting it from 1.

Basic checks on the adjacency matrix are performed and missing entries are replaced by zeros. If TOMType = "unsigned", entries of the adjacency matrix are required to lie between 0 and 1; for TOMType = "signed" they can be between -1 and 1. In both cases the resulting TOM entries, as well as the corresponding dissimilarities, lie between 0 and 1.

The underlying C code assumes that the diagonal of the adjacency matrix equals 1. If this is not the case, the diagonal of the input is set to 1 before the calculation begins.

## Value

A matrix holding the topological overlap.

## Author(s)

Peter Langfelder

## References

Bin Zhang and Steve Horvath (2005) "A General Framework for Weighted Gene Co-Expression Network Analysis", *Statistical Applications in Genetics and Molecular Biology*: Vol. 4: No. 1, Article 17

## See Also

[TOMsimilarityFromExpr](#)

---

TOMsimilarityFromExpr *Topological overlap matrix*

---

## Description

Calculation of the topological overlap matrix from given expression data.

## Usage

```
TOMsimilarityFromExpr(  
  datExpr,  
  corType = "pearson",  
  networkType = "unsigned",  
  power = 6,  
  TOMType = "signed",  
  TOMDenom = "min",  
  maxPOutliers = 1,  
  quickCor = 0,  
  pearsonFallback = "individual",  
  cosineCorrelation = FALSE,  
  nThreads = 0,  
  verbose = 1, indent = 0)
```

**Arguments**

datExpr	expression data. A data frame in which columns are genes and rows are samples. NAs are allowed, but not too many.
corType	character string specifying the correlation to be used. Allowed values are (unique abbreviations of) "pearson" and "bicor", corresponding to Pearson and bid-weight midcorrelation, respectively. Missing values are handled using the <code>pairwise.complete.obs</code> option.
networkType	network type. Allowed values are (unique abbreviations of) "unsigned", "signed", "signed hybrid". See <a href="#">adjacency</a> .
power	soft-thresholding power for network construction.
TOMType	one of "none", "unsigned", "signed". If "none", adjacency will be used for clustering. If "unsigned", the standard TOM will be used (more generally, TOM function will receive the adjacency as input). If "signed", TOM will keep track of the sign of correlations between neighbors.
TOMDenom	a character string specifying the TOM variant to be used. Recognized values are "min" giving the standard TOM described in Zhang and Horvath (2005), and "mean" in which the min function in the denominator is replaced by mean. The "mean" may produce better results but at this time should be considered experimental.
maxPOutliers	only used for <code>corType=="bicor"</code> . Specifies the maximum percentile of data that can be considered outliers on either side of the median separately. For each side of the median, if higher percentile than <code>maxPOutliers</code> is considered an outlier by the weight function based on $9 * \text{mad}(x)$ , the width of the weight function is increased such that the percentile of outliers on that side of the median equals <code>maxPOutliers</code> . Using <code>maxPOutliers=1</code> will effectively disable all weight function broadening; using <code>maxPOutliers=0</code> will give results that are quite similar (but not equal to) Pearson correlation.
quickCor	real number between 0 and 1 that controls the handling of missing data in the calculation of correlations. See details.
pearsonFallback	Specifies whether the bicor calculation, if used, should revert to Pearson when median absolute deviation ( <code>mad</code> ) is zero. Recognized values are (abbreviations of) "none", "individual", "all". If set to "none", zero <code>mad</code> will result in NA for the corresponding correlation. If set to "individual", Pearson calculation will be used only for columns that have zero <code>mad</code> . If set to "all", the presence of a single zero <code>mad</code> will cause the whole variable to be treated in Pearson correlation manner (as if the corresponding robust option was set to FALSE). Has no effect for Pearson correlation. See <a href="#">bicor</a> .
cosineCorrelation	logical: should the cosine version of the correlation calculation be used? The cosine calculation differs from the standard one in that it does not subtract the mean.
nThreads	non-negative integer specifying the number of parallel threads to be used by certain parts of correlation calculations. This option only has an effect on systems on which a POSIX thread library is available (which currently includes Linux and Mac OSX, but excludes Windows). If zero, the number of online processors will be used if it can be determined dynamically, otherwise correlation calculations will use 2 threads.
verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.

indent            indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

**Value**

A matrix holding the topological overlap.

**Author(s)**

Peter Langfelder

**References**

Bin Zhang and Steve Horvath (2005) "A General Framework for Weighted Gene Co-Expression Network Analysis", *Statistical Applications in Genetics and Molecular Biology*: Vol. 4: No. 1, Article 17

**See Also**

[TOMsimilarity](#)

---

transposeBigData            *Transpose a big matrix or data frame*

---

**Description**

This transpose command partitions a big matrix (or data frame) into blocks and applies the `t()` function to each block separately.

**Usage**

```
transposeBigData(x, blocksize = 20000)
```

**Arguments**

`x`                    a matrix or data frame  
`blocksize`           a positive integer larger than 1, which determines the block size. Default is 20k.

**Details**

Assume you have a very large matrix with say 500k columns. In this case, the standard transpose function of R `t()` can take a long time. Solution: Split the original matrix into sub-matrices by dividing the columns into blocks. Next apply `t()` to each sub-matrix. The same holds if the large matrix contains a large number of rows. The function `transposeBigData` automatically checks whether the large matrix contains more rows or more columns. If the number of columns is larger than or equal to the number of rows then the block wise splitting will be applied to columns otherwise to the rows.

**Value**

A matrix or data frame (depending on the input `x`) which is the transpose of `x`.

**Note**

This function can be considered a wrapper of `t()`

**Author(s)**

Steve Horvath, UCLA

**References**

Any linear algebra book will explain the transpose.

**See Also**

The standard function `t`.

**Examples**

```
x=data.frame(matrix(1:10000,nrow=4,ncol=2500))
dimnames(x)[[2]]=paste("Y",1:2500,sep="")
xTranspose=transposeBigData(x)
x[1:4,1:4]
xTranspose[1:4,1:4]
```

---

TrueTrait

*Estimate the true trait underlying a list of surrogate markers.*

---

**Description**

Assume an imprecisely measured trait  $y$  that is related to the true, unobserved trait  $y_{TRUE}$  as follows  $y_{TRUE}=y+\text{noise}$  where noise is assumed to have mean zero and a constant variance. Assume you have 1 or more surrogate markers for  $y_{TRUE}$  corresponding to the columns of `datX`. The function implements several approaches for estimating  $y_{TRUE}$  based on the inputs  $y$  and/or `datX`.

**Usage**

```
TrueTrait(datX, y, datXtest=NULL,
          corFnc = "cor", corOptions = "use = 'pairwise.complete.obs'",
          LeaveOneOut.CV=FALSE, skipMissingVariables=TRUE,
          addLinearModel=FALSE, Strata=NULL)
```

**Arguments**

<code>datX</code>	is a vector or data frame whose columns correspond to the surrogate markers (variables) for the true underlying trait. The number of rows of <code>datX</code> equals the number of observations, i.e. it should equal the length of $y$
$y$	is a numeric vector which specifies the observed trait.
<code>datXtest</code>	can be set as a matrix or data frame of a second, independent test data set. Its columns should correspond to those of <code>datX</code> , i.e. the two data sets should have the same number of columns but the number or rows (test set observations) can be different.

<code>corFnc</code>	Character string specifying the correlation function to be used in the calculations. Recommended values are the default Pearson correlation "cor" or biweight mid-correlation "bicor". Additional arguments to the correlation function can be specified using <code>corOptions</code> .
<code>corOptions</code>	Character string giving additional arguments to the function specified in <code>corFnc</code> .
<code>LeaveOneOut.CV</code>	logical. If TRUE then leave one out cross validation estimates will be calculated for <code>y.true1</code> and <code>y.true2</code> based on <code>datX</code> .
<code>skipMissingVariables</code>	logical. If TRUE then variables whose values are missing for a given observation will be skipped when estimating the true trait of that particular observation. Thus, the estimate of a particular observation are determined by all the variables whose values are non-missing.
<code>addLinearModel</code>	logical. If TRUE then the function also estimates the true trait based on the predictions of the linear model <code>lm(y~., data=datX)</code>
<code>Strata</code>	vector whose length show equal the number of rows of <code>datX</code> . This vector allows one to specify subsets of observations (called strata or batches or independent data sets) to which the function should be applied. For example, if the observations were measured in different batches then one can apply the function to each individual batch of data. No estimates will be calculated for <code>Strata</code> components that equal NA or "exclude".

## Details

This R function implements formulas described in Klemra and Doubal (2006). The assumptions underlying these formulas are described in Klemra et al. But briefly, the function provides several estimates of the true underlying trait under the following assumptions: 1) There is a true underlying trait that affects  $y$  and a list of surrogate markers corresponding to the columns of `datX`. 2) There is a linear relationship between the true underlying trait and  $y$  and the surrogate markers. 3)  $y_{TRUE} = y + \text{Noise}$  where the Noise term has a mean of zero and a fixed variance. 4) Weighted least squares estimation is used to relate the surrogate markers to the underlying trait where the weights are proportional to  $1/\text{ssq}_j$  where  $\text{ssq}_j$  is the noise variance of the  $j$ -th marker.

Specifically, output `y.true1` corresponds to formula 31, `y.true2` corresponds to formula 25, and `y.true3` corresponds to formula 34.

Although the true underlying trait  $y_{TRUE}$  is not known, one can estimate the standard deviation between the estimate `y.true2` and  $y_{TRUE}$  using formula 33. Similarly, one can estimate the SD for the estimate `y.true3` using formula 42. These estimated SDs correspond to output components 2 and 3, respectively. These SDs are valuable since they provide a sense of how accurate the measure is.

To estimate the correlations between  $y$  and the surrogate markers, one can specify different correlation measures. The default method is based on the Person correlation but one can also specify the biweight midcorrelation by choosing "bicor", see `help(bicor)` to learn more.

When the `datX` is comprised of observations measured in different strata (e.g. different batches or independent data sets) then one can obtain stratum specific estimates by specifying the strata using the argument `Strata`. In this case, the estimation focuses on one stratum at a time.

## Value

A list with the following components.

<code>datEstimates</code>	is a data frame whose columns corresponds to estimates of the true underlying trait. The number of rows equals the number of observations, i.e. the length of $y$ .
---------------------------	---

The first column `y.true1` is the average value of standardized columns of `datX` where standardization subtracts out the intercept term and divides by the slope of the linear regression model `lm(marker~y)`. Since this estimate ignores the fact that the surrogate markers have different correlations with `y`, it is typically inferior to `y.true2`. The second column `y.true2` equals the weighted average value of standardized columns of `datX`. The standardization is described in section 2.4 of Klemra et al. The weights are proportional to  $r^2/(1+r^2)$  where  $r$  denotes the correlation between the surrogate marker and `y`. Since this estimate does not include `y` as additional surrogate marker, it may be slightly inferior to `y.true3`. Having said this, the difference between `y.true2` and `y.true3` is often negligible. An additional column called `y.lm` is added if `codeaddLinearModel=TRUE`. In this case, `y.lm` reports the linear model predictions. Finally, the column `y.true3` is very similar to `y.true2` but it includes `y` as additional surrogate marker. It is expected to be the best estimate of the underlying true trait (see Klemra et al 2006).

`datEstimatestest`

is output only if a test data set has been specified in the argument `datXtest`. In this case, it contains a data frame with columns `ytrue1` and `ytrue2`. The number of rows equals the number of test set observations, i.e the number of rows of `datXtest`. Since the value of `y` is not known in case of a test data set, one cannot calculate `y.true3`. An additional column with linear model predictions `y.lm` is added if `codeaddLinearModel=TRUE`.

`datEstimates.LeaveOneOut.CV`

is output only if the argument `LeaveOneOut.CV` has been set to `TRUE`. In this case, it contains a data frame with leave-one-out cross validation estimates of `ytrue1` and `ytrue2`. The number of rows equals the length of `y`. Since the value of `y` is not known in case of a test data set, one cannot calculate `y.true3`.

`SD.ytrue2`

is a scalar. This is an estimate of the standard deviation between the estimate `y.true2` and the true (unobserved) `yTRUE`. It corresponds to formula 33.

`SD.ytrue3`

is a scalar. This is an estimate of the standard deviation between `y.true3` and the true (unobserved) `yTRUE`. It corresponds to formula 42.

`datVariableInfo`

is a data frame that reports information for each variable (column of `datX`) when it comes to the definition of `y.true2`. The rows correspond to the number of variables. Columns report the variable name, the center (intercept that is subtracted to scale each variable), the scale (i.e. the slope that is used in the denominator), and finally the weights used in the weighted sum of the scaled variables.

`datEstimatesByStratum`

a data frame that will only be output if `Strata` is different from `NULL`. In this case, it has the same dimensions as `datEstimates` but the estimates were calculated separately for each level of `Strata`.

`SD.ytrue2ByStratum`

a vector of length equal to the different levels of `Strata`. Each component reports the estimate of `SD.ytrue2` for observations in the stratum specified by `unique(Strata)`.

`datVariableInfoByStratum`

a list whose components are matrices with variable information. Each list component reports the variable information in the stratum specified by `unique(Strata)`.

### Author(s)

Steve Horvath

## References

Klemera P, Doubal S (2006) A new approach to the concept and computation of biological age. *Mechanisms of Ageing and Development* 127 (2006) 240-248

Choa IH, Parka KS, Limb CJ (2010) An Empirical Comparative Study on Validation of Biological Age Estimation Algorithms with an Application of Work Ability Index. *Mechanisms of Ageing and Development* Volume 131, Issue 2, February 2010, Pages 69-78

## Examples

```
# observed trait
y=rnorm(1000,mean=50,sd=20)
# unobserved, true trait
yTRUE =y +rnorm(100,sd=10)
# now we simulate surrogate markers around the true trait
datX=simulateModule(yTRUE,nGenes=20, minCor=.4,maxCor=.9, geneMeans=rnorm(20,50,30) )
True1=TrueTrait(datX=datX,y=y)
datTrue=True1$datEstimates
par(mfrow=c(2,2))
for (i in 1:dim(datTrue)[[2]]) {
  meanAbsDev= mean(abs(yTRUE-datTrue[,i]))
  verboseScatterplot(datTrue[,i],yTRUE,xlab=names(datTrue)[i],
                    main=paste(i, "MeanAbsDev=", signif(meanAbsDev,3)));
  abline(0,1)
}
#compare the estimated standard deviation of y.true2
True1[[2]]
# with the true SD
sqrt(var(yTRUE-datTrue$y.true2))
#compare the estimated standard deviation of y.true3
True1[[3]]
# with the true SD
sqrt(var(yTRUE-datTrue$y.true3))
```

---

unsignedAdjacency

*Calculation of unsigned adjacency*

---

## Description

Calculation of the unsigned network adjacency from expression data. The restricted set of parameters for this function should allow a faster and less memory-hungry calculation.

## Usage

```
unsignedAdjacency(
  datExpr,
  datExpr2 = NULL,
  power = 6,
  corFnc = "cor", corOptions = "use = 'p'")
```

### Arguments

datExpr	expression data. A data frame in which columns are genes and rows are samples. Missing values are ignored.
datExpr2	optional specification of a second set of expression data. See details.
power	soft-thresholding power for network construction.
corFnc	character string giving the correlation function to be used for the adjacency calculation. Recommended choices are "cor" and "bicor", but other functions can be used as well.
corOptions	character string giving further options to be passed to the correlation function

### Details

The correlation function will be called with arguments `datExpr`, `datExpr2` plus any extra arguments given in `corOptions`. If `datExpr2` is `NULL`, the standard correlation functions will calculate the correlation of columns in `datExpr`.

### Value

Adjacency matrix of dimensions  $n \times n$ , where  $n$  is the number of genes in `datExpr`.

### Author(s)

Steve Horvath and Peter Langfelder

### References

Bin Zhang and Steve Horvath (2005) "A General Framework for Weighted Gene Co-Expression Network Analysis", *Statistical Applications in Genetics and Molecular Biology*: Vol. 4: No. 1, Article 17

### See Also

[adjacency](#)

---

userListEnrichment      *Measure enrichment between inputted and user-defined lists*

---

### Description

This function measures list enrichment between inputted lists of genes and files containing user-defined lists of genes. Significant enrichment is measured using a hypergeometric test. A pre-made collection of brain-related lists can also be loaded. The function writes the significant enrichments to a file, but also returns all overlapping genes across all comparisons.



**Usage**

```

userListEnrichment(
  geneR, labelR,
  fnIn = NULL, catNmIn = fnIn,
  nameOut = "enrichment.csv",
  useBrainLists = FALSE, useBloodAtlases = FALSE, omitCategories = "grey",
  outputCorrectedPvalues = TRUE, useStemCellLists = FALSE,
  outputGenes = FALSE,
  minGenesInCategory = 1,
  useBrainRegionMarkers = FALSE, useImmunePathwayLists = FALSE,
  usePalazzoloWang = FALSE)

```

**Arguments**

geneR	A vector of gene (or other) identifiers. This vector should include ALL genes in your analysis (i.e., the genes corresponding to your labeled lists AND the remaining background reference genes).
labelR	A vector of labels (for example, module assignments) corresponding to the geneR list. NOTE: For all background reference genes that have no corresponding label, use the label "background" (or any label included in the omitCategories parameter).
fnIn	A vector of file names containing user-defined lists. These files must be in one of three specific formats (see details section). The default (NULL) may only be used if one of the "use_____" parameters is TRUE.
catNmIn	A vector of category names corresponding to each fnIn. This name will be appended to each overlap corresponding to that filename. The default sets the category names as the corresponding file names.
nameOut	Name of the file where the output enrichment information will be written. (Note that this file includes only a subset of what is returned by the function.)
useBrainLists	If TRUE, a pre-made set of brain-derived enrichment lists will be added to any user-defined lists for enrichment comparison. The default is FALSE. See references section for related references.
useBloodAtlases	If TRUE, a pre-made set of blood-derived enrichment lists will be added to any user-defined lists for enrichment comparison. The default is FALSE. See references section for related references.
omitCategories	Any labelR entries corresponding to these categories will be ignored. The default ("grey") will ignore unassigned genes in a standard WGCNA network.
outputCorrectedPvalues	If TRUE (default) only pvalues that are significant after correcting for multiple comparisons (using Bonferroni method) will be outputted to nameOut. Otherwise the uncorrected p-values will be outputted to the file. Note that both sets of p-values for all comparisons are reported in the returned "pValues" parameter.
useStemCellLists	If TRUE, a pre-made set of stem cell (SC)-derived enrichment lists will be added to any user-defined lists for enrichment comparison. The default is FALSE. See references section for related references.
outputGenes	If TRUE, will output a list of all genes in each returned category, as well as a count of the number of genes in each category. The default is FALSE.

minGenesInCategory	Will omit all significant categories with fewer than minGenesInCategory genes (default is 1).
useBrainRegionMarkers	If TRUE, a pre-made set of enrichment lists for human brain regions will be added to any user-defined lists for enrichment comparison. The default is FALSE. These lists are derived from data from the Allen Human Brain Atlas ( <a href="http://human.brain-map.org/">http://human.brain-map.org/</a> ). See references section for more details.
useImmunePathwayLists	If TRUE, a pre-made set of enrichment lists for immune system pathways will be added to any user-defined lists for enrichment comparison. The default is FALSE. These lists are derived from the lab of Daniel R Saloman. See references section for more details.
usePalazzoloWang	If TRUE, a pre-made set of enrichment lists compiled by Mike Palazzolo and Jim Wang from CHDI will be added to any user-defined lists for enrichment comparison. The default is FALSE. See references section for more details.

## Details

User-inputted files for `fnIn` can be in one of three formats:

- 1) Text files (must end in ".txt") with one list per file, where the first line is the list descriptor and the remaining lines are gene names corresponding to that list, with one gene per line. For example  
Ribosome RPS4 RPS8 ...
- 2) Gene / category files (must be csv files), where the first line is the column headers corresponding to Genes and Lists, and the remaining lines correspond to the genes in each list, for any number of genes and lists. For example: Gene, Category RPS4, Ribosome RPS8, Ribosome ... NDUF1, Mitochondria NDUF3, Mitochondria ... MAPT, AlzheimersDisease PSEN1, AlzheimersDisease PSEN2, AlzheimersDisease ...
- 3) Module membership (kME) table in csv format. Currently, the module assignment is the only thing that is used, so as long as the Gene column is 2nd and the Module column is 3rd, it doesn't matter what is in the other columns. For example, PSID, Gene, Module, <other columns> <psid>, RPS4, blue, <other columns> <psid>, NDUF1, red, <other columns> <psid>, RPS8, blue, <other columns> <psid>, NDUF3, red, <other columns> <psid>, MAPT, green, <other columns> ...

## Value

pValues	A data frame showing, for each comparison, the input category, user defined category, type, the number of overlapping genes and both the uncorrected and Bonferroni corrected p-values for every pair of list overlaps tested.
ovGenes	A list of character vectors corresponding to the overlapping genes for every pair of list overlaps tested. Specific overlaps can be found by typing <variable-Name>\$ovGenes\$'<labelR> - <comparisonCategory>'. See example below.
sigOverlaps	Identical information that is written to nameOut. A data frame with columns giving the input category, user defined category, type, and P-values (corrected or uncorrected, depending on outputCorrectedPvalues) corresponding to all significant enrichments.

## Author(s)

Jeremy Miller

## References

The primary reference for this function is: Miller JA, Cai C, Langfelder P, Geschwind DH, Kurian SM, Salomon DR, Horvath S. (2011) Strategies for aggregating gene expression data: the collapseRows R function. *BMC Bioinformatics* 12:322.

If you have any suggestions for lists to add to this function, please e-mail Jeremy Miller at [jere-myinla@gmail.com](mailto:jere-myinla@gmail.com)

---

References for the pre-defined brain lists (useBrainLists=TRUE, in alphabetical order by category descriptor) are as follows:

ABA ==> Cell type markers from: Lein ES, et al. (2007) Genome-wide atlas of gene expression in the adult mouse brain. *Nature* 445:168-176.

ADvsCT\_inCA1 ==> Lists of genes found to be increasing or decreasing with Alzheimer's disease in 3 studies: 1. Blalock => Blalock E, Geddes J, Chen K, Porter N, Markesbery W, Landfield P (2004) Incipient Alzheimer's disease: microarray correlation analyses reveal major transcriptional and tumor suppressor responses. *PNAS* 101:2173-2178. 2. Colangelo => Colangelo V, Schurr J, Ball M, Pelaez R, Bazan N, Lukiw W (2002) Gene expression profiling of 12633 genes in Alzheimer hippocampal CA1: transcription and neurotrophic factor down-regulation and up-regulation of apoptotic and pro-inflammatory signaling. *J Neurosci Res* 70:462-473. 3. Liang => Liang WS, et al (2008) Altered neuronal gene expression in brain regions differentially affected by Alzheimer's disease: a reference data set. *Physiological genomics* 33:240-56.

Bayes ==> Postsynaptic Density Proteins from: Bayes A, et al. (2011) Characterization of the proteome, diseases and evolution of the human postsynaptic density. *Nat Neurosci*. 14(1):19-21.

Blalock\_AD ==> Modules from a network using the data from: Blalock E, Geddes J, Chen K, Porter N, Markesbery W, Landfield P (2004) Incipient Alzheimer's disease: microarray correlation analyses reveal major transcriptional and tumor suppressor responses. *PNAS* 101:2173-2178.

CA1vsCA3 ==> Lists of genes enriched in CA1 and CA3 relative to other each and to other areas of the brain, from several studies: 1. Ginsberg => Ginsberg SD, Che S (2005) Expression profile analysis within the human hippocampus: comparison of CA1 and CA3 pyramidal neurons. *J Comp Neurol* 487:107-118. 2. Lein => Lein E, Zhao X, Gage F (2004) Defining a molecular atlas of the hippocampus using DNA microarrays and high-throughput in situ hybridization. *J Neurosci* 24:3879-3889. 3. Newrzella => Newrzella D, et al (2007) The functional genome of CA1 and CA3 neurons under native conditions and in response to ischemia. *BMC Genomics* 8:370. 4. Torres => Torres-Munoz JE, Van Waveren C, Keegan MG, Bookman RJ, Petit CK (2004) Gene expression profiles in microdissected neurons from human hippocampal subregions. *Brain Res Mol Brain Res* 127:105-114. 5. GorLorT => In either Ginsberg or Lein or Torres list.

Cahoy ==> Definite (10+ fold) and probable (1.5+ fold) enrichment from: Cahoy JD, et al. (2008) A transcriptome database for astrocytes, neurons, and oligodendrocytes: A new resource for understanding brain development and function. *J Neurosci* 28:264-278.

CTX ==> Modules from the CTX (cortex) network from: Oldham MC, et al. (2008) Functional organization of the transcriptome in human brain. *Nat Neurosci* 11:1271-1282.

DiseaseGenes ==> Probable (C or better rating as of 16 Mar 2011) and possible (all genes in database as of ~2008) genetics-based disease genes from: <http://www.alzforum.org/>

EarlyAD ==> Genes whose expression is related to cognitive markers of early Alzheimer's disease vs. non-demented controls with AD pathology, from: Parachikova, A., et al (2007) Inflammatory changes parallel the early stages of Alzheimer disease. *Neurobiology of Aging* 28:1821-1833.

HumanChimp ==> Modules showing region-specificity in both human and chimp from: Oldham MC, Horvath S, Geschwind DH (2006) Conservation and evolution of gene coexpression networks in human and chimpanzee brains. *Proc Natl Acad Sci USA* 103: 17973-17978.

HumanMeta ==> Modules from the human network from: Miller J, Horvath S, Geschwind D (2010) Divergence of human and mouse brain transcriptome highlights Alzheimer disease pathways. Proc Natl Acad Sci 107:12698-12703.

JAXdiseaseGene ==> Genes where mutations in mouse and/or human are known to cause any disease. WARNING: this list represents an oversimplification of data! This list was created from the Jackson Laboratory: Bult CJ, Eppig JT, Kadin JA, Richardson JE, Blake JA; Mouse Genome Database Group (2008) The Mouse Genome Database (MGD): Mouse biology and model systems. Nucleic Acids Res 36 (database issue):D724-D728.

Lu\_Aging ==> Modules from a network using the data from: Lu T, Pan Y, Kao S-Y, Li C, Kohane I, Chan J, Yankner B (2004) Gene regulation and DNA damage in the ageing human brain. Nature 429:883-891.

MicroglialMarkers ==> Markers for microglia and macrophages from several studies: 1. GSE772 => Gan L, et al. (2004) Identification of cathepsin B as a mediator of neuronal death induced by Abeta-activated microglial cells using a functional genomics approach. J Biol Chem 279:5565-5572. 2. GSE1910 => Albright AV, Gonzalez-Scarano F (2004) Microarray analysis of activated mixed glial (microglia) and monocyte-derived macrophage gene expression. J Neuroimmunol 157:27-38. 3. AitGhezala => Ait-Ghezala G, Mathura VS, Laporte V, Quadros A, Paris D, Patel N, et al. Genomic regulation after CD40 stimulation in microglia: relevance to Alzheimer's disease. Brain Res Mol Brain Res 2005;140(1-2):73-85. 4. 3treatments\_Thomas => Thomas, DM, Francescutti-Verbeem, DM, Kuhn, DM (2006) Gene expression profile of activated microglia under conditions associated with dopamine neuronal damage. The FASEB Journal 20:515-517.

MitochondrialType ==> Mitochondrial genes from the somatic vs. synaptic fraction of mouse cells from: Winden KD, et al. (2009) The organization of the transcriptional network in specific neuronal classes. Mol Syst Biol 5:291.

MO ==> Markers for many different things provided to my by Mike Oldham. These were originally from several sources: 1. 2+\_26Mar08 => Genetics-based disease genes in two or more studies from <http://www.alzforum.org/> (compiled by Mike Oldham). 2. Bachoo => Bachoo, R.M. et al. (2004) Molecular diversity of astrocytes with implications for neurological disorders. PNAS 101, 8384-8389. 3. Foster => Foster, LJ, de Hoog, CL, Zhang, Y, Zhang, Y, Xie, X, Mootha, VK, Mann, M. (2006) A Mammalian Organelle Map by Protein Correlation Profiling. Cell 125(1): 187-199. 4. Morciano => Morciano, M. et al. Immunolocalization of two synaptic vesicle pools from synaptosomes: a proteomics analysis. J. Neurochem. 95, 1732-1745 (2005). 5. Sugino => Sugino, K. et al. Molecular taxonomy of major neuronal classes in the adult mouse forebrain. Nat. Neurosci. 9, 99-107 (2006).

MouseMeta ==> Modules from the mouse network from: Miller J, Horvath S, Geschwind D (2010) Divergence of human and mouse brain transcriptome highlights Alzheimer disease pathways. Proc Natl Acad Sci 107:12698-12703.

Sugino/Winden ==> Conservative list of genes in modules from the network from: Winden K, Oldham M, Mirnics K, Ebert P, Swan C, Levitt P, Rubenstein J, Horvath S, Geschwind D (2009). The organization of the transcriptional network in specific neuronal classes. Molecular systems biology 5. NOTE: Original data came from this neuronal-cell-type-selection experiment in mouse: Sugino K, Hempel C, Miller M, Hattox A, Shapiro P, Wu C, Huang J, Nelson S (2006). Molecular taxonomy of major neuronal classes in the adult mouse forebrain. Nat Neurosci 9:99-107

Voineagu ==> Several Autism-related gene categories from: Voineagu I, Wang X, Johnston P, Lowe JK, Tian Y, Horvath S, Mill J, Cantor RM, Blencowe BJ, Geschwind DH. (2011). Transcriptomic analysis of autistic brain reveals convergent molecular pathology. Nature 474(7351):380-4

—————References for the pre-defined blood atlases (useBloodAtlases=TRUE, in alphabetical order by category descriptor) are as follows:

Blood(composite) ==> Lists for blood cell types with this label are made from combining marker genes from the following three publications: 1. Abbas AB, Baldwin D, Ma Y, Ouyang W, Gur-

ney A, et al. (2005). Immune response in silico (IRIS): immune-specific genes identified from a compendium of microarray expression data. *Genes Immun.* 6(4):319-31. 2. Grigoryev YA, Kurian SM, Avnur Z, Borie D, Deng J, et al. (2010). Deconvoluting post-transplant immunity: cell subset-specific mapping reveals pathways for activation and expansion of memory T, monocytes and B cells. *PLoS One.* 5(10):e13358. 3. Watkins NA, Gusnanto A, de Bono B, De S, Miranda-Saavedra D, et al. (2009). A HaemAtlas: characterizing gene expression in differentiated human blood cells. *Blood.* 113(19):e1-9.

Gnatenko ==> Top 50 marker genes for platelets from: Gnatenko DV, et al. (2009) Transcript profiling of human platelets using microarray and serial analysis of gene expression (SAGE). *Methods Mol Biol.* 496:245-72.

Gnatenko2 ==> Platelet-specific genes on a custom microarray from: Gnatenko DV, et al. (2010) Class prediction models of thrombocytosis using genetic biomarkers. *Blood.* 115(1):7-14.

Kabanova ==> Red blood cell markers from: Kabanova S, et al. (2009) Gene expression analysis of human red blood cells. *Int J Med Sci.* 6(4):156-9.

Whitney ==> Genes corresponding to individual variation in blood from: Whitney AR, et al. (2003) Individuality and variation in gene expression patterns in human blood. *PNAS.* 100(4):1896-1901.

————— References for the pre-defined stem cell (SC) lists (useStemCellLists=TRUE, in alphabetical order by category descriptor) are as follows:

Cui ==> genes differentiating erythrocyte precursors (CD36+ cells) from multipotent human primary hematopoietic stem cells/progenitor cells (CD133+ cells), from: Cui K, Zang C, Roh TY, Schones DE, Childs RW, Peng W, Zhao K. (2009). Chromatin signatures in multipotent human hematopoietic stem cells indicate the fate of bivalent genes during differentiation. *Cell Stem Cell* 4:80-93

Lee ==> gene lists related to Polycomb proteins in human embryonic SCs, from (a highly-cited paper!): Lee TI, Jenner RG, Boyer LA, Guenther MG, Levine SS, Kumar RM, Chevalier B, Johnstone SE, Cole MF, Isono K, et al. (2006) Control of developmental regulators by polycomb in human embryonic stem cells. *Cell* 125:301-313

————— References and more information for the pre-defined human brain region lists (useBrainRegionMarkers=TRUE):

HBA ==> Hawrylycz MJ, Lein ES, Guillozet-Bongaarts AL, Shen EH, Ng L, Miller JA, et al. (2012) An Anatomically Comprehensive Atlas of the Adult Human Brain Transcriptome. *Nature* (in press) Three categories of marker genes are presented: 1. globalMarker(top200) = top 200 global marker genes for 22 large brain structures. Genes are ranked based on fold change enrichment (expression in region vs. expression in rest of brain) and the ranks are averaged between brains 2001 and 2002 (human.brain-map.org). 2. localMarker(top200) = top 200 local marker genes for 90 large brain structures. Same as 1, except fold change is defined as expression in region vs. expression in larger region (format: <region>\_IN\_<largerRegion>). For example, enrichment in CA1 is relative to other subcompartments of the hippocampus. 3. localMarker(FC>2) = same as #2, but only local marker genes with fold change > 2 in both brains are included. Regions with <10 marker genes are omitted.

————— More information for the pre-defined immune pathways lists (useImmunePathwayLists=TRUE):

ImmunePathway ==> These lists were created by Brian Modena (a member of Daniel R Salomon's lab at Scripps Research Institute), with input from Sunil M Kurian and Dr. Salomon, using Ingenuity, WikiPathways and literature search to assemble them. They reflect knowledge-based immune pathways and were in part informed by Dr. Salomon and colleague's work in expression profiling of biopsies and peripheral blood but not in some highly organized process. These lists are not from any particular publication, but are culled to include only genes of reasonably high confidence.

References for the pre-defined lists from CHDI (usePalazzoloWang=TRUE, in alphabetical order by category descriptor) are as follows:

Biocyc NCBI Biosystems ==> Several gene sets from the "Biocyc" component of NCBI Biosystems: Geer LY, Marchler-Bauer A, Geer RC, Han L, He J, He S, Liu C, Shi W, Bryant SH (2010) The NCBI BioSystems database. *Nucleic Acids Res.* 38(Database issue):D492-6.

Kegg NCBI Biosystems ==> Several gene sets from the "Kegg" component of NCBI Biosystems: Geer LY et al 2010 (full citation above).

Palazzolo and Wang ==> These gene sets were compiled from a variety of sources by Mike Palazzolo and Jim Wang at CHDI.

Pathway Interaction Database NCBI Biosystems ==> Several gene sets from the "Pathway Interaction Database" component of NCBI Biosystems: Geer LY et al 2010 (full citation above).

PMID 17500595 Kaltenbach 2007 ==> Several gene sets from: Kaltenbach LS, Romero E, Becklin RR, Chettier R, Bell R, Phansalkar A, et al. (2007) Huntingtin interacting proteins are genetic modifiers of neurodegeneration. *PLoS Genet.* 3(5):e82

PMID 22348130 Schaefer 2012 ==> Several gene sets from: Schaefer MH, Fontaine JF, Vinayagam A, Porras P, Wanker EE, Andrade-Navarro MA (2012) HIPPIE: Integrating protein interaction networks with experiment based quality scores. *PLoS One.* 7(2):e31826

PMID 22556411 Culver 2012 ==> Several gene sets from: Culver BP, Savas JN, Park SK, Choi JH, Zheng S, Zeitlin SO, Yates JR 3rd, Tanese N. (2012) Proteomic analysis of wild-type and mutant huntingtin-associated proteins in mouse brains identifies unique interactions and involvement in protein synthesis. *J Biol Chem.* 287(26):21599-614

PMID 22578497 Cajigas 2012 ==> Several gene sets from: Cajigas IJ, Tushev G, Will TJ, tom Dieck S, Fuerst N, Schuman EM. (2012) The local transcriptome in the synaptic neuropil revealed by deep sequencing and high-resolution imaging. *Neuron.* 74(3):453-66

Reactome NCBI Biosystems ==> Several gene sets from the "Reactome" component of NCBI Biosystems: Geer LY et al 2010 (full citation above).

Wiki Pathways NCBI Biosystems ==> Several gene sets from the "Wiki Pathways" component of NCBI Biosystems: Geer LY et al 2010 (full citation above).

Yang ==> These gene sets were compiled from a variety of sources by Mike Palazzolo and Jim Wang at CHDI.

## Examples

```
# Example: first, read in some gene names and split them into categories
data(BrainLists);
listGenes = unique(as.character(BrainLists[,1]))
set.seed(100)
geneR = sort(sample(listGenes,2000))
categories = sort(rep(standardColors(10),200))
categories[sample(1:2000,200)] = "grey"
write(c("TESTLIST1",geneR[300:400], sep="\n"),"TESTLIST1.txt")
write(c("TESTLIST2",geneR[800:1000],sep="\n"),"TESTLIST2.txt")

# Now run the function!
testResults = userListEnrichment(geneR, labelR=categories,
                                fnIn=c("TESTLIST1.txt","TESTLIST2.txt"),
                                catNmIn=c("TEST1","TEST2"),
                                nameOut = "testEnrichment.csv",useBrainLists=TRUE, omitCategories ="grey")

# To see a list of all significant enrichments, either open
# the file "testEnrichments.csv" in the current directory, or type:
```

```

testResults$sigOverlaps

# To see all of the overlapping genes between two categories
#(whether or not the p-value is significant), type
#restResults$ovGenes$'<labelR> -- <comparisonCategory>'. For example:

testResults$ovGenes$"black -- TESTLIST1__TEST1"
testResults$ovGenes$"red -- salmon_M12_Ribosome__HumanMeta"

# More detailed overlap information is in the pValue output. For example:
head(testResults$pValue)

```

---

vectorizeMatrix	<i>Turn a matrix into a vector of non-redundant components</i>
-----------------	--

---

### Description

A convenient function to turn a matrix into a vector of non-redundant components. If the matrix is non-symmetric, returns a vector containing all entries of the matrix. If the matrix is symmetric, only returns the upper triangle and optionally the diagonal.

### Usage

```
vectorizeMatrix(M, diag = FALSE)
```

### Arguments

M	the matrix or data frame to be vectorized.
diag	logical: should the diagonal be included in the output?

### Value

A vector containing the non-redundant entries of the input matrix.

### Author(s)

Steve Horvath

---

vectorTOM	<i>Topological overlap for a subset of the whole set of genes</i>
-----------	---

---

### Description

This function calculates topological overlap of a small set of vectors with respect to a whole data set.

**Usage**

```
vectorTOM(
  datExpr,
  vect,
  subtract1 = FALSE,
  blockSize = 2000,
  corFnc = "cor", corOptions = "use = 'p'",
  networkType = "unsigned",
  power = 6,
  verbose = 1, indent = 0)
```

**Arguments**

datExpr	a data frame containing the expression data of the whole set, with rows corresponding to samples and columns to genes.
vect	a single vector or a matrix-like object containing vectors whose topological overlap is to be calculated.
subtract1	logical: should calculation be corrected for self-correlation? Set this to TRUE if vect contains a subset of datExpr.
blockSize	maximum block size for correlation calculations. Only important if vect contains a large number of columns.
corFnc	character string giving the correlation function to be used for the adjacency calculation. Recommended choices are "cor" and "bcor", but other functions can be used as well.
corOptions	character string giving further options to be passed to the correlation function.
networkType	character string giving network type. Allowed values are (unique abbreviations of) "unsigned", "signed", "signed hybrid". See <a href="#">adjacency</a> .
power	soft-thresholding power for network construction.
verbose	integer level of verbosity. Zero means silent, higher values make the output progressively more and more verbose.
indent	indentation for diagnostic messages. Zero means no indentation, each unit adds two spaces.

**Details**

Topological overlap can be viewed as the normalized count of shared neighbors encoded in an adjacency matrix. In this case, the adjacency matrix is calculated between the columns of vect and datExpr and the topological overlap of vectors in vect measures the number of shared neighbors in datExpr that vectors of vect share.

**Value**

A matrix of dimensions  $n \times n$ , where  $n$  is the number of columns in vect.

**Author(s)**

Peter Langfelder



## References

Bin Zhang and Steve Horvath (2005) "A General Framework for Weighted Gene Co-Expression Network Analysis", *Statistical Applications in Genetics and Molecular Biology*: Vol. 4: No. 1, Article 17

## See Also

[TOMsimilarity](#) for standard calculation of topological overlap.

---

verboseBarplot	<i>Barplot with error bars, annotated by Kruskal-Wallis or ANOVA p-value</i>
----------------	--

---

## Description

Produce a barplot with error bars, annotated by Kruskal-Wallis or ANOVA p-value.

## Usage

```
verboseBarplot(x, g,
               main = "", xlab = NA, ylab = NA,
               cex = 1, cex.axis = 1.5, cex.lab = 1.5, cex.main = 1.5,
               color = "grey", numberStandardErrors = 1,
               KruskalTest = TRUE, AnovaTest = FALSE, two.sided = TRUE,
               horiz = FALSE, ...)
```

## Arguments

x	numerical or binary vector of data whose group means are to be plotted
g	a factor or a an object coercible to a factor giving the groups whose means are to be calculated.
main	main title for the plot.
xlab	label for the x-axis.
ylab	label for the y-axis.
cex	character expansion factor for plot annotations.
cex.axis	character expansion factor for axis annotations.
cex.lab	character expansion factor for axis labels.
cex.main	character expansion factor for the main title.
color	a vector giving the colors of the bars in the barplot.
numberStandardErrors	size of the error bars in terms of standard errors. See details.
KruskalTest	logical: should Kruskal-Wallis test be performed? See details.
AnovaTest	logical: should ANOVA be performed? See details.
two.sided	logical: should the printed p-value be two-sided? See details.
horiz	logical: should the bars be drawn horizontally?
...	other parameters to function <a href="#">barplot</a>

## Details

This function creates a barplot of a numeric variable (input  $x$ ) across the levels of a grouping variable (input  $g$ ). The height of the bars equals the mean value of  $x$  across the observations with a given level of  $g$ . By default, the barplot also shows plus/minus one standard error. If you want only plus one standard error (not minus) choose `two.sided=TRUE`. But the number of standard errors can be determined with the input `numberStandardErrors`. For example, if you want a 95% confidence interval around the mean, choose `numberStandardErrors=2`. If you don't want any standard errors set `numberStandardErrors=-1`. The function also outputs the p-value of a Kruskal Wallis test (Fisher test for binary input data), which is a non-parametric multi group comparison test. Alternatively, one can use Analysis of Variance (Anova) to compute a p-value by setting `AnovaTest=TRUE`. Anova is a generalization of the Student t-test to multiple groups. In case of two groups, the Anova p-value equals the Student t-test p-value. Anova should only be used if  $x$  follows a normal distribution. Anova also assumes homoscedasticity (equal variances). The Kruskal Wallis test is often advantageous since it makes no distributional assumptions. Since the Kruskal Wallis test is based on the ranks of  $x$ , it is more robust with regard to outliers. All p-values are two-sided.

## Value

None.

## Author(s)

Steve Horvath

## See Also

[barplot](#)

## Examples

```
group=sample(c(1,2),100,replace=TRUE)

height=rnorm(100,mean=group)

par(mfrow=c(2,2))
verboseBarplot(height,group, main="1 SE, Kruskal Test")

verboseBarplot(height,group,numberStandardErrors=2,
               main="2 SE, Kruskal Test")

verboseBarplot(height,group,numberStandardErrors=2,AnovaTest=TRUE,
               main="2 SE, Anova")

verboseBarplot(height,group,numberStandardErrors=2,AnovaTest=TRUE,
               main="2 SE, Anova, only plus SE", two.sided=FALSE)
```

---

verboseBoxplot	<i>Boxplot annotated by a Kruskal-Wallis p-value</i>
----------------	--

---

### Description

Plot a boxplot annotated by the Kruskal-Wallis p-value. Uses the function [boxplot](#) for the actual drawing.

### Usage

```
verboseBoxplot(x, g, main = "", xlab = NA, ylab = NA,  
              cex = 1, cex.axis = 1.5, cex.lab = 1.5, cex.main = 1.5,  
              notch = TRUE, varwidth = TRUE, ...)
```

### Arguments

x	numerical vector of data whose group means are to be plotted
g	a factor or a an object coercible to a factor giving the groups that will go into each box.
main	main title for the plot.
xlab	label for the x-axis.
ylab	label for the y-axis.
cex	character expansion factor for plot annotations.
cex.axis	character expansion factor for axis annotations.
cex.lab	character expansion factor for axis labels.
cex.main	character expansion factor for the main title.
notch	logical: should the notches be drawn? See <a href="#">boxplot</a> and <a href="#">boxplot.stats</a> for details.
varwidth	logical: if TRUE, the boxes are drawn with widths proportional to the square-roots of the number of observations in the groups.
...	other arguments to the function <a href="#">boxplot</a> . Of note is the argument <code>las</code> that specifies label orientation. Value <code>las=1</code> will result in horizontal labels (the default), while <code>las=2</code> will result in vertical labels, useful when the labels are long.

### Value

Returns the value returned by the function [boxplot](#).

### Author(s)

Steve Horvath

### See Also

[boxplot](#)

---

 verboseIplot

*Scatterplot with density*


---

### Description

Produce a scatterplot that shows density with color and is annotated by the correlation, MSE, and regression line.

### Usage

```
verboseIplot(
  x, y,
  xlim = NA, ylim = NA,
  nBinsX = 150, nBinsY = 150,
  ztransf = function(x) {x}, gamma = 1,
  sample = NULL, corFnc = "cor", corOptions = "use = 'p'",
  main = "", xlab = NA, ylab = NA, cex = 1,
  cex.axis = 1.5, cex.lab = 1.5, cex.main = 1.5,
  abline = FALSE, abline.color = 1, abline.lty = 1,
  corLabel = corFnc, ...)
```

### Arguments

x	numerical vector to be plotted along the x axis.
y	numerical vector to be plotted along the y axis.
xlim	define the range in x axis
ylim	define the range in y axis
nBinsX	number of bins along the x axis
nBinsY	number of bins along the y axis
ztransf	Function to transform the number of counts per pixel, which will be mapped by the function in colramp to well defined colors. The user has to make sure that the transformed density lies in the range [0,zmax], where zmax is any positive number ( $\geq 2$ ).
gamma	color correction power
sample	either a number of points to be sampled or a vector of indices input x and y for points to be plotted. Useful when the input vectors are large and plotting all points is not practical.
corFnc	character string giving the correlation function to annotate the plot.
corOptions	character string giving further options to the correlation function.
main	main title for the plot.
xlab	label for the x-axis.
ylab	label for the y-axis.
cex	character expansion factor for plot annotations.
cex.axis	character expansion factor for axis annotations.
cex.lab	character expansion factor for axis labels.
cex.main	character expansion factor for the main title.



**Arguments**

<code>x</code>	numerical vector to be plotted along the x axis.
<code>y</code>	numerical vector to be plotted along the y axis.
<code>sample</code>	determines whether x and y should be sampled for plotting, useful to keep the plot manageable when x and y are large vectors. The default NULL value implies no sampling. A single numeric value will be interpreted as the number of points to sample randomly. If a vector is given, it will be interpreted as the indices of the entries in x and y that should be plotted. In either case, the correlation and p value will be determined from the full vectors x and y.
<code>corFnc</code>	character string giving the correlation function to annotate the plot.
<code>corOptions</code>	character string giving further options to the correlation function.
<code>main</code>	main title for the plot.
<code>xlab</code>	label for the x-axis.
<code>ylab</code>	label for the y-axis.
<code>cex</code>	character expansion factor for plot annotations.
<code>cex.axis</code>	character expansion factor for axis annotations.
<code>cex.lab</code>	character expansion factor for axis labels.
<code>cex.main</code>	character expansion factor for the main title.
<code>abline</code>	logical: should the linear regression fit line be plotted?
<code>abline.color</code>	color specification for the fit line.
<code>abline.lty</code>	line type for the fit line.
<code>corLabel</code>	character string to be used as the label for the correlation value printed in the main title.
<code>displayAsZero</code>	Correlations whose absolute value is smaller than this number will be displayed as zero. This can result in a more intuitive display (for example, <code>cor=0</code> instead of <code>cor=2.6e-17</code> ).
<code>col</code>	color of the plotted symbols. Recycled as necessary.
<code>bg</code>	fill color of the plotted symbols (used for certain symbols). Recycled as necessary.
<code>lmFnc</code>	linear model fit function. Used to calculate the linear model fit line if 'abline' is TRUE. For example, robust linear models are implemented in the function <code>rlm</code> .
<code>...</code>	other arguments to the function <code>plot</code> .

**Details**

Irrespective of the specified correlation function, the p-value is always calculated for pearson correlation.

**Value**

If `sample` above is given, the indices of the plotted points are returned invisibly.

**Author(s)**

Steve Horvath and Peter Langfelder

**See Also**

`plot.default` for standard scatterplots

---

votingLinearPredictor *Voting linear predictor*


---

### Description

Predictor based on univariate regression on all or selected given features that pools all predictions using weights derived from the univariate linear models.

### Usage

```
votingLinearPredictor(
  x, y, xtest = NULL,
  classify = FALSE,
  CVfold = 0,
  randomSeed = 12345,
  assocFnc = "cor", assocOptions = "use = 'p'",
  featureWeightPowers = NULL, priorWeights = NULL,
  weighByPrediction = 0,
  nFeatures.hi = NULL, nFeatures.lo = NULL,
  dropUnusedDimensions = TRUE,
  verbose = 2, indent = 0)
```

### Arguments

x	Training features (predictive variables). Each column corresponds to a feature and each row to an observation.
y	The response variable. Can be a single vector or a matrix with arbitrary many columns. Number of rows (observations) must equal to the number of rows (observations) in x.
xtest	Optional test set data. A matrix of the same number of columns (i.e., features) as x. If test set data are not given, only the prediction on training data will be returned.
classify	Should the response be treated as a categorical variable? Classification really only works with two classes. (The function will run for multiclass problems as well, but the results will be sub-optimal.)
CVfold	Optional specification of cross-validation fold. If 0 (the default), no cross-validation is performed.
randomSeed	Random seed, used for observation selection for cross-validation. If NULL, the random generator is not reset.
assocFnc	Function to measure association. Usually a measure of correlation, for example Pearson correlation or <a href="#">bicor</a> .
assocOptions	Character string specifying the options to be passed to the association function.
featureWeightPowers	Powers to which to raise the result of assocFnc to obtain weights. Can be a single number or a vector of arbitrary length; the returned value will contain one prediction per power.

priorWeights	Prior weights for the features. If given, must be either (1) a vector of the same length as the number of features (columns in $x$ ); (2) a matrix of dimensions $\text{length}(\text{featureWeightPowers}) \times (\text{number of features})$ ; or (3) array of dimensions $(\text{number of response variables}) \times \text{length}(\text{featureWeightPowers}) \times (\text{number of features})$ .
weighByPrediction	(Optional) power to downweigh features that are not well predicted between training and test sets. See details.
nFeatures.hi	Optional restriction of the number of features to use. If given, this many features with the highest association and lowest association (if nFeatures.lo is not given) will be used for prediction.
nFeatures.lo	Optional restriction of the number of lowest (i.e., most negatively) associated features to use. Only used if nFeatures.hi is also non-NULL.
dropUnusedDimensions	Logical: should unused dimensions be dropped from the result?
verbose	Integer controlling how verbose the diagnostic messages should be. Zero means silent.
indent	Indentation for the diagnostic messages. Zero means no indentation, each unit adds two spaces.

### Details

The predictor calculates the association of each (selected) feature with the response and uses the association to calculate the weight of the feature as  $\text{sign}(\text{association}) * (\text{association})^{\text{featureWeightPower}}$ . Optionally, this weight is multiplied by `priorWeights`. Further, a feature prediction weight can be used to downweigh features that are not well predicted by other features (see below).

For classification, the (continuous) result of the above calculation is turned into ordinal values essentially by rounding.

If features exhibit non-trivial correlations among themselves (such as, for example, in gene expression data), one can attempt to down-weight features that do not exhibit the same correlation in the test set. This is done by using essentially the same predictor to predict `_features_` from all other features in the test data (using the training data to train the feature predictor). Because test features are known, the prediction accuracy can be evaluated. If a feature is predicted badly (meaning the error in the test set is much larger than the error in the cross-validation prediction in training data), it may mean that its quality in the training or test data is low (for example, due to excessive noise or outliers). Such features can be downweighed using the argument `weighByPrediction`. The extra factor is  $\min(1, (\text{root mean square prediction error in test set}) / (\text{root mean square cross-validation prediction error in the training data})^{\text{weighByPrediction}})$ , that is it is never bigger than 1.

### Value

A list with the following components:

predicted	The back-substitution prediction on the training data. Normally an array of dimensions $(\text{number of observations}) \times (\text{number of response variables}) \times \text{length}(\text{featureWeightPowers})$ , but unused are dropped unless <code>dropUnusedDimensions = FALSE</code> .
weightBase	Absolute value of the associations of each feature with each response.
variableImportance	The weight of each feature in the prediction (including the sign).
predictedTest	If input <code>xtest</code> is non-NULL, the predicted test response, in format analogous to predicted above.



CVpredicted     If input CVfold is non-zero, cross-validation prediction on the training data.

**Note**

It makes little practical sense to supply neither xtest nor CVfold since the prediction accuracy on training data will be highly biased.

**Author(s)**

Peter Langfelder

**See Also**

[bicor](#) for robust correlation that can be used as an association measure

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