# Package: SEset (via r-universe)

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Type Package

Title Computing Statistically-Equivalent Path Models

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Author Oisín Ryan

Maintainer Oisín Ryan <o.ryan@uu.nl>

Description Tools to compute and analyze the set of statistically-equivalent (Gaussian, linear) path models which generate the input precision or (partial) correlation matrix. This procedure is useful for understanding how statistical network models such as the Gaussian Graphical Model (GGM) perform as causal discovery tools. The statistical-equivalence set of a given GGM expresses the uncertainty we have about the sign, size and direction of directed relationships based on the weights matrix of the GGM alone. The derivation of the equivalence set and its use for understanding GGMs as causal discovery tools is described by Ryan, O., Bringmann, L.F., & Schuurman, N.K. (2022) <doi:10.31234/osf.io/ryg69>.

BugReports https://github.com/ryanoisin/SEset

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```
cov_to_path
```

Path model from covariance matrix with ordering

# Description

Helper function. Takes a covariance matrix and ordering and generates a lower-triangular weights matrix.

# Usage

```
cov_to_path(sigma, ordering = NULL, digits = 2)
```

# Arguments

sigma	input matrix, with rows and columns in desired topological ordering Must be an invertible square matrix
ordering	character vector containing the dimension names of the input matrix in the de- sired ordering
digits	the number of digits used to round the output

# Value

lower triangular matrix containing regression weights of the path model. Element ij represents the effect of  $X_j$  on  $X_i$ 

# See Also

network\_to\_path

find\_parents

#### Description

Return parent indices from a (weighted) DAG for a given child

# Usage

```
find_parents(mat, child)
```

# Arguments

mat	An $p \times p$ weights or adjacency matrix
child	Index giving the position of the child node

# Value

a vector containing index numbers defining the parent nodes

# References

Ryan O, Bringmann LF, Schuurman NK (upcoming). "The challenge of generating causal hypotheses using network models." *in preperation*.

# See Also

# r2\_distribution

get_psi	Calculate residual-covariance matrix based on a path model and co-
	variance matrix

# Description

Takes an ordered path model and corresponding variance-covariance matrix and computes the appropriate residual covariance matrix (psi)

# Usage

get\_psi(B, sigma, digits = 3)

# Arguments

В	input $p \times p$ linear SEM weights matrix
sigma	variance-covariance matrix of the variables
digits	how many digits to round the result to

# Value

a  $p \times p$  residual variance-covariance matrix

network\_to\_path Path model from ordered precision matrix

# Description

Takes a precision matrix and generates a lower-triangular weights matrix.

# Usage

```
network_to_path(omega, input_type = "precision", digits = 20)
```

# Arguments

omega	input matrix, with rows and columns in desired topological ordering Must be an invertible square matrix
input_type	specifies what type of matrix 'omega' is. default is "precision", other options include a matrix of partial correlations ("parcor") or a covariance matrix "covariance"
digits	desired rounding of the output matrix

# Value

lower triangular matrix containing regression weights of the path model. Element ij represents the effect of  $X_i$  on  $X_i$ 

#### References

Ryan O, Bringmann LF, Schuurman NK (upcoming). "The challenge of generating causal hypotheses using network models." *in preperation*.

Shojaie A, Michailidis G (2010). "Penalized likelihood methods for estimation of sparse high-dimensional directed acyclic graphs." *Biometrika*, **97**(3), 519–538.

Bollen KA (1989). Structural equations with latent variables. Oxford, England, John Wiley \& Sons.

# See Also

network\_to\_SEset

# network\_to\_SEset

# Examples

```
data(riskcor)
omega <- (qgraph::EBICglasso(riskcor, n = 69, returnAllResults = TRUE))$optwi
# qgraph method estimates a non-symmetric omega matrix, but uses forceSymmetric to create
# a symmetric matrix (see qgraph:::EBICglassoCore line 65)
omega <- as.matrix(Matrix::forceSymmetric(omega)) # returns the precision matrix
B <- network_to_path(omega, digits=2)
# Path model can be plotted as a weighted DAG
pos <- matrix(c(2,0,-2,-1,-2,1,0,2,0.5,0,0,-2),6,2,byrow=TRUE)
# qgraph reads matrix elements as "from row to column"
# regression weights matrices are read "from column to row"
# path model weights matrix must be transposed for qgraph
qgraph::qgraph(t(B), labels=rownames(riskcor), layout=pos,
repulsion=.8, vsize=c(10,15), theme="colorblind", fade=FALSE)
```

network\_to\_SEset SE-set from precision matrix

# Description

Takes a precision matrix and generates the SE-set, a set of statistically equivalent path models. Unless otherwise specified, the SEset will contain one weights matrix for every possible topological ordering of the input precision matrix

#### Usage

```
network_to_SEset(
   omega,
   orderings = NULL,
   digits = 20,
   rm_duplicates = FALSE,
   input_type = "precision"
)
```

#### Arguments

omega	input $p \times p$ precision matrix
orderings	An optional matrix of $n$ orderings from which to generate the SE-set. Must be in the form of a $p \times n$ matrix with each column a vector of dimension names in the desired order. If unspecified, all $p!$ possible orderings are used
digits	desired rounding of the output weights matrices in the SE-set, in decimal places. Defaults to 20.
rm_duplicates	Logical indicating whether only unique DAGs should be returned

input_type	specifies what type of matrix 'omega' is. default is "precision", other options
	include a matrix of partial correlations ("parcor") or a model implied covariance
	or correlation matrix "MIcov"

#### Value

a  $p! \times p$  matrix containing the SE-set (or  $n \times p$  matrix if a custom set of n orderings is specified). Each row represents a lower-triangular weights matrix, stacked column-wise.

# References

Ryan O, Bringmann LF, Schuurman NK (upcoming). "The challenge of generating causal hypotheses using network models." *in preperation*.

Shojaie A, Michailidis G (2010). "Penalized likelihood methods for estimation of sparse highdimensional directed acyclic graphs." *Biometrika*, **97**(3), 519–538.

Bollen KA (1989). *Structural equations with latent variables*. Oxford, England, John Wiley \& Sons.

#### See Also

network\_to\_path, reorder\_mat, order\_gen

# Examples

```
# first estimate the precision matrix
data(riskcor)
omega <- (qgraph::EBICglasso(riskcor, n = 69, returnAllResults = TRUE))$optwi</pre>
# qgraph method estimates a non-symmetric omega matrix, but uses forceSymmetric to create
# a symmetric matrix (see qgraph:::EBICglassoCore line 65)
omega <- as.matrix(Matrix::forceSymmetric(omega)) # returns the precision matrix</pre>
SE <- network_to_SEset(omega, digits=3)</pre>
# each row of SE defines a path-model weights matrix.
# We can extract element 20 by writing it to a matrix
example <- matrix(SE[20,],6,6)</pre>
# Example path model can be plotted as a weighted DAG
pos <- matrix(c(2,0,-2,-1,-2,1,0,2,0.5,0,0,-2),6,2,byrow=TRUE)</pre>
# qgraph reads matrix elements as "from row to column"
# regression weights matrices are read "from column to row"
# path model weights matrix must be transposed for qgraph
qgraph::qgraph(t(example), labels=rownames(riskcor), layout=pos,
repulsion=.8, vsize=c(10,15), theme="colorblind", fade=FALSE)
```

order\_gen

# Description

Takes a matrix and generates a matrix containing all orderings of the rows and columns

# Usage

order\_gen(omega)

# Arguments

omega input p-dimensional square matrix

# Value

a  $p \times p!$  matrix of dimension orderings. Each column represents an ordering of dimension names as character strings.

# References

Chasalow S (2012). *combinat: combinatorics utilities*. R package version 0.0-8, https://CRAN. R-project.org/package=combinat.

#### See Also

reorder\_mat, network\_to\_SEset

# Examples

data(riskcor)
orderings <- order\_gen(riskcor)</pre>

# Each column of orderings defines an ordering of variables
print(orderings[,1])
# in the second element, the fifth and sixth variable are switched
print(orderings[,2])

path\_to\_network

#### Description

Takes a path model and generates the corresponding (standardized) precision matrix or covariance matrix. The inverse of network\_to\_path.

# Usage

path\_to\_network(B, psi = NULL, output = "precision")

# Arguments

В	input $p \times p$ weights matrix
psi	variance-covariance matrix for the residuals. If NULL (the default) will impose the constraint that the variables have variance 1 and the residuals are uncorrelated
output	Function returns the precision ("precision") or covariance ("covariance") matrix

# Value

a  $p \times p$  precision or covariance matrix

# References

Ryan O, Bringmann LF, Schuurman NK (upcoming). "The challenge of generating causal hypotheses using network models." *in preperation*.

Shojaie A, Michailidis G (2010). "Penalized likelihood methods for estimation of sparse high-dimensional directed acyclic graphs." *Biometrika*, **97**(3), 519–538.

Bollen KA (1989). *Structural equations with latent variables*. Oxford, England, John Wiley \& Sons.

# See Also

network\_to\_path, SEset\_to\_network

propcal

# Description

A function used to analyse the SEset results. Calculates the proportion of path models in a given SEset in which a particular edge is present

# Usage

```
propcal(SEmatrix, names = NULL, rm_duplicate = TRUE, directed = TRUE)
```

# Arguments

SEmatrix	An $n \times p$ matrix containing the SEset, where each row represents a $p \times p$ weights matrix stacked column-wise
names	optional character vector containing dimension names
rm_duplicate	Should duplicate weights matrices be removed from the SEset. Defaults to TRUE.
directed	If FALSE, the directionality of edges is ignored, and the output reflects in what proportion of the SEset an edge of any direction is present. If TRUE, the proportion is calculated seperately for edges of either direction. Defaults to TRUE

# Value

a  $p \times p$  matrix showing in what proportion particular edges are present. If directed=TRUE, elements ij denote the proportion of weights matrices containing a path from  $X_j$  to  $X_i$ . If directed=F, the output will be a symmetric matrix, with element ij denoting in what proprtion an edge of either direction connects  $X_i$  to  $X_j$ .

# References

Ryan O, Bringmann LF, Schuurman NK (upcoming). "The challenge of generating causal hypotheses using network models." *in preperation*.

# See Also

network\_to\_SEset

r2\_distribution

#### Description

A function used to analyse the SEset results. For each member of the SE-set, calculate the proportion of explained variance in each child node, when predicted by all of its parent nodes

#### Usage

```
r2_distribution(SEmatrix, cormat, names = NULL, indices = NULL)
```

# Arguments

SEmatrix	An $n \times p$ matrix containing the SEset, where each row represents a $p \times p$ weights matrix stacked column-wise
cormat	A $p \times p$ matrix containing the marginal covariances or correlations
names	optional character vector containing dimension names
indices	option vector of matrix indices, indicating which variables to compute the R^2 distribution for

#### Value

Returns an  $n \times p$  matrix of  $R^2$  values. For each member of the SE-set, this represents the variance explained in node  $X_i$  by it's parents in that weighted DAG.

# References

Ryan O, Bringmann LF, Schuurman NK (upcoming). "The challenge of generating causal hypotheses using network models." *in preperation*. Haslbeck JM, Waldorp LJ (2018). "How well do network models predict observations? On the importance of predictability in network models." *Behavior Research Methods*, **50**(2), 853–861.

# See Also

network\_to\_SEset, find\_parents

# Examples

```
# first estimate the precision matrix
data(riskcor)
omega <- (qgraph::EBICglasso(riskcor, n = 69, returnAllResults = TRUE))$optwi
# qgraph method estimates a non-symmetric omega matrix, but uses forceSymmetric to create
# a symmetric matrix (see qgraph:::EBICglassoCore line 65)
omega <- as.matrix(Matrix::forceSymmetric(omega)) # returns the precision matrix</pre>
```

```
SEmatrix <- network_to_SEset(omega, digits=3)</pre>
```

# reorder\_mat

```
r2set <- r2_distribution(SEmatrix, cormat = riskcor, names = NULL, indices = c(1,3,4,5,6))
# Plot results
apply(r2set,2,hist)
# For ggplot format, execute
# r2set <- tidyr::gather(r2set)</pre>
```

reorder\_mat

## Re-order rows and columns

# Description

Takes a matrix and re-orders the rows and columns to some target ordering

# Usage

```
reorder_mat(matrix, names)
```

# Arguments

matrix	input matrix to be re-arranged. Must have rows and columns named
names	character vector containing the dimension names of the input matrix in the de- sired ordering

# Value

input matrix with rows and columns sorted according to names

# See Also

order\_gen, network\_to\_SEset

# Examples

data(riskcor)

```
# first define an ordered vector of names
row_names <- rownames(riskcor)
row_names_new <- row_names[c(1,2,3,4,6,5)]</pre>
```

```
reorder_mat(riskcor,row_names_new)
```

# The fifth and sixth row and column have been switched
print(riskcor)

riskcor

#### Description

Reported sample correlation matrix from a cross-sectional study on cognitive risk and resilience factors in remitted depression patients, from Hoorelebeke, Marchetti, DE Schryver and Koster (2016). The study was conducted with 69 participants, and the correlation matrix consists of six variables. The variables are as follows:

#### Usage

data(riskcor)

#### Format

A 6 by 6 correlation matrix

#### Details

\* 'BRIEF\_WM': working memory complaints, a self-report measure of perceived cognitive control \* 'PASAT\_ACC': PASAT accuracy, performance on behavioural measure of congitive control \* 'Adapt ER': self-report adaptive emotion regulation strategies \* 'Maladapt ER': self-report maladaptive emotion regulation strategies \* 'Resilience': self-report resilience \* 'Resid Depress': self-report residual depressive symptoms

# Source

<https://ars.els-cdn.com/content/image/1-s2.0-S0165032715313252-mmc1.pdf>

#### References

Hoorelbeke K, Marchetti I, De Schryver M, Koster EH (2016). "The interplay between cognitive risk and resilience factors in remitted depression: a network analysis." *Journal of Affective Disorders*, **195**, 96–104.

#### Examples

```
data(riskcor)
print(rownames(riskcor))
print(riskcor)
```

# Description

Takes the SE-set and calculates for each weights matrix the corresponding precision matrix. Used to check the results of network\_to\_SEset to assess deviations from statistical equivalence induced due to rounding, thresholding, and numerical approximations.

# Usage

```
SEset_to_network(
   SEmatrix,
   order.ref = NULL,
   order.mat = NULL,
   output = "raw",
   omega = NULL
)
```

# Arguments

SEmatrix	a $n \times p$ matrix containing the SE-set. The output of network_to_SEset
order.ref	an optional character vector with variable names, the reference ordering of the precision matrix.
order.mat	a $n \times p$ matrix of character strings, defining the ordering of the matrix corresponding to each row of SEmatrix. If NULL it is assumed that all orderings are included and they are generated using order_gen
output	Output as "raw" or "summary". See value below
omega	Comparision precision matrix, e.g. original input precision matrix to network_to_SEset. Only necessary if output = "summary"

#### Value

If output = "raw", a  $n \times p$  matrix of precision matrices stacked column-wise in n rows. If output = "summary" returns a list containing the bias, MSE and RMSE for each re-calculated precision matrix, relative to comparison omega matrix supplied.

# References

Ryan O, Bringmann LF, Schuurman NK (upcoming). "The challenge of generating causal hypotheses using network models." *in preperation*.

Shojaie A, Michailidis G (2010). "Penalized likelihood methods for estimation of sparse highdimensional directed acyclic graphs." *Biometrika*, **97**(3), 519–538.

Bollen KA (1989). Structural equations with latent variables. Oxford, England, John Wiley \& Sons.

SEset\_to\_network

# See Also

 $network\_to\_path, path\_to\_network$ 

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