

# Package ‘wTO’

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**Type** Package

**Title** Computing Weighted Topological Overlaps (wTO) & Consensus wTO Network

**Version** 2.0.1

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**Description** Computes the Weighted Topological Overlap with positive and negative signs (wTO) networks given a data frame containing the mRNA count/ expression/ abundance per sample, and a vector containing the interested nodes of interaction (a subset of the elements of the full data frame). It also computes the cut-off threshold or p-value based on the individuals bootstrap or the values reshuffle per individual. It also allows the construction of a consensus network, based on multiple wTO networks. The package includes a visualization tool for the networks. More about the methodology can be found at <[arXiv:1711.04702](https://arxiv.org/abs/1711.04702)>.

**License** GPL-2

**LazyData** TRUE

**Imports** data.table, igraph, magrittr, plyr, parallel, som, visNetwork, reshape2, Rfast, HiClimR, methods

**Suggests** knitr, rmarkdown

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CorrelationOverlap	<i>CorrelationOverlap</i>
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### Description

This function computes the correlation between Nodes and the Overlapping Nodes of interest.

### Usage

```
CorrelationOverlap(Data, Overlap, method)
```

### Arguments

Data	data.frame containing the expression data. Nodes on the Rows, Individuals on the Columns. Don't forget to give the names to the Nodes and to the Individuals. Nodes must have the row.names() with the Node Name.
Overlap	A vector containing the names of the Nodes of interest.
method	Spearman ("s", "spearman") or Pearson ("p", "pearson") correlation

### Author(s)

Deisy Morselli Gysi <deisy at bioinf.uni-leipzig.de>

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ExampleGRF	<i>ExampleGRF</i>
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### Description

ExampleGRF data.frame containing data.frame containing names of GRFs.

### Usage

```
data(ExampleGRF)
```

### Format

data.frame 184 lines, 1 column.

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metagenomics\_abundance  
*metagenomics\_abundance*

---

**Description**

metagenomics\_abundance

**Usage**

data('metagenomics\_abundance')

**Format**

data.frame from The USC Microbial Observatory. The data is public available at <<https://www.ebi.ac.uk/metagenomics/proje>>

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Microarray\_Expression1  
*Microarray\_Expression1*

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**Description**

Microarray\_Expression1 data.frame containing expression data for 1000 genes and 18 individuals.

**Usage**

Microarray\_Expression1

**Format**

data.frame 1000 lines, 18 columns.

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Microarray\_Expression2  
*Microarray\_Expression2*

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**Description**

Microarray\_Expression2 data.frame containing expression data for 1000 genes and 18 individuals.

**Usage**

Microarray\_Expression2

**Format**

data.frame 1000 lines, 18 columns.

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 NetVis

*NetVis*


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### Description

Given a set of Nodes and the weight of the edges, a cutoff for the edges, it draws the networks. Returns a list with the nodes and edges attributes. And plots the network.

### Usage

```
NetVis(
  Node.1,
  Node.2,
  wTO,
  pval = NULL,
  MakeGroups = FALSE,
  padj = NULL,
  cutoff = list(kind = "Threshold", value = 0.5),
  layout = NULL,
  smooth.edges = T,
  path = NULL,
  Cluster = F,
  legend = T,
  shape = list(shape = "triangle", names = NULL),
  manipulation = F
)
```

### Arguments

Node.1	Names of the Nodes.1 that are connected to the Nodes.2. It's the output from wTO.Complete or Consensus.
Node.2	Names of the Nodes.2 that are connected to the Nodes.1. It's the output from wTO.Complete or Consensus.
wTO	weight of the links, the wTO output from wTO.Complete or wTO.Consensus.
pval	p-values for the wTO value. By default it is NULL.
MakeGroups	algorithm to find clusters. One of the followings: walktrap, optimal, spinglass, edge.betweenness, fast_greedy, infomap, louvain, label_prop, leading_eigen. Default to FALSE.
padj	Adjusted p-values for the wTO value. By default it is NULL.
cutoff	It's a list containing the kind of cutoff to be used (pval, Threshold or pval.adj)and it's value. Example: cutoff= list(kind = "Threshold", value = 0.5)
layout	a layout from the igraph package.
smooth.edges	If the edges should be smoothed or not.
path	If the graph should be saved specify the name of the file.

Cluster TRUE or FALSE if the nodes should be clustered (double click to uncluster).  
 legend TRUE or FALSE if the legend should appear.  
 shape a list shape=list(shape = "triangle", names = NULL), with the shape and the IDs that should have a different shape, shape can be: diamond, star, triangle, triangleDown or square.  
 manipulation TRUE or FALSE if the graph should be editable.

### Author(s)

Deisy Morselli Gysi <deisy at bioinf.uni-leipzig.de>

### Examples

```
X = wTO.Complete( k = 1, n = 5, Data = Microarray_Expression1,
  Overlap = ExampleGRF$x, method = "p", plot = FALSE)
# Plot with the default arguments.
NetVis(Node.1 = X$wTO$Node.1, Node.2 = X$wTO$Node.2,
  wTO = X$wTO$wTO_sign, cutoff = list(kind =
  "Threshold", value = 0.50))

## Not run:
# Plotting just the edges with p-value < 0.05, with straight edges, nodes clustered,
# no legend and manipulation of the graph enabled.
NetVis(Node.1 = X$wTO$Node.1, Node.2 = X$wTO$Node.2,
  wTO = X$wTO$wTO_sign, pval = X$wTO$pval_sign,
  padj = X$wTO$pval_sign,
  cutoff= list(kind = "pval", value = 0.05),
  smooth.edges = FALSE,
  Cluster = TRUE, legend = FALSE, manipulation = TRUE)
# Plotting just the edges with wTO > 0.50, no legend and the nodes:
# "ZNF738", "ZNF677" with triangle shape,
# no legend and manipulation of the graph enabled.
NetVis(Node.1 = X$wTO$Node.1, Node.2 = X$wTO$Node.2,
  wTO = X$wTO$wTO_sign, pval = X$wTO$pval_sign,
  padj = X$wTO$pval_sign, cutoff= list(kind = "Threshold", value = 0.5),legend = FALSE,
  shape = list(shape = "triangle", names = c("ZNF738", "ZNF677"))))

## End(Not run)
```

---

wTO

wTO

---

### Description

Calculates the weighted topological overlap (wTO) between a set of Nodes and the Overlapping nodes. This function implements the method from Nowick (2009).

**Usage**

```
wTO(A, sign = c("abs", "sign"))
```

**Arguments**

A	Is the weighted adjacency matrix (correlation matrix).
sign	("abs", "sign") if the user wants to use the absolute correlation or the signed correlation.

**Value**

A matrix containing the wTO values.

**Author(s)**

Deisy Morselli Gysi <deisy at bioinf.uni-leipzig.de>

**References**

Katja Nowick, Tim Gernat, Eivind Almaas and Lisa Stubbs (2009) <doi:10.1073/pnas.0911376106>

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wTO.Complete

*wTO.Complete*

---

**Description**

Compute the wTO and also the bootstraps. Proposed at: arXiv:1711.04702

**Usage**

```
wTO.Complete(  
  k = 1,  
  n = 100,  
  Data,  
  Overlap = row.names(Data),  
  method = "p",  
  method_resampling = "Bootstrap",  
  pvalmethod = "BH",  
  savecor = F,  
  expected.diff = 0.2,  
  lag = NULL,  
  ID = NULL,  
  normalize = F,  
  plot = T  
)
```

**Arguments**

k	Number of threads to be used for computing the weight Topological Overlap. Default is set to 1.
n	Number of resamplings, used to compute the empirical distributions of the links. Default is set to 100.
Data	data.frame containing the count / expression data for the correlation.
Overlap	Set of nodes of interest, where the Overlapping weights will be computed.
method	Type of the correlation that should be used. "s" / "spearman" will compute the rank spearman correlation, "p" / "pearson" will compute the linear correlation. If no value is given, the default is to use "p".
method_resampling	method of the resampling. Bootstrap, BlockBootstrap or Reshuffle. Bootstrap null hypothesis is that the wTO is random, and Reshuffle tests if the wTO is equal to zero.
pvalmethod	method to compute the multiple test correction for the pvalue. for more information check the function <a href="#">p.adjust</a> .
savecor	T/F if need to save the correlation.
expected.diff	Difference expected between the real wTO and resampled wTO By default, it is set to 0.2.
lag	time dependency, lag, if you are using the BlockedBootstrap.
ID	ID of the samples for the blocked bootstrap (for repeated measures).
normalize	T/F Should the data be normalized?
plot	T/F Should the diagnosis plot be plotted?

**Value**

a list with results.

- wTO is a data.frame containig the Nodes, the wTO computed using the signed correlations, the pvalue and the adj.pvalue.
- abs.wTO is a data.frame containig the Nodes, the wTO computed using the absolute correlations, the pvalue and the adj.pvalue.
- Correlation is a data.frame containing the correlation between all the nodes.
- Empirical.Quantile quantile values for the empirical distribution.
- Quantile quantile values for the sample distribution.

**Author(s)**

Deisy Morselli Gysi <deisy at bioinf.uni-leipzig.de>

**Examples**

```
## Not run:
# Using spearman rank correlation and bonferroni correction for the pvalues.
wTO.Complete( k =8, n = 1000, Data = Microarray_Expression1,
  Overlap = ExampleGRF$x, method = "s", pvalmethod = "bonferroni")
# Changing the resampling method to Reshuffle.
wTO.Complete( k =1, n = 1000, Data = Microarray_Expression1,
  Overlap = ExampleGRF$x, method_resampling = "Reshuffle")
# Changing the resampling method to BlockBootstrap, with a lag of 2.
row.names(metagenomics_abundance) = metagenomics_abundance$OTU
metagenomics_abundance = metagenomics_abundance[,-1]
wTO.Complete( k =1, n = 1000, Data = metagenomics_abundance, method = "s",
  Overlap = row.names(metagenomics_abundance), method_resampling = "BlockBootstrap", lag = 2)
wTO.Complete( k =2, n = 1000, Data = Microarray_Expression1, method = "s",
  Overlap = ExampleGRF$x, method_resampling = "BlockBootstrap", ID = rep(1:9,each = 2))
X = wTO.Complete( k =1, n = 1000, Data = Microarray_Expression1,
  Overlap = ExampleGRF$x, method = "p", plot = FALSE)

## End(Not run)
```

---

wTO.Consensus

*wTO.Consensus*


---

**Description**

Consensus requires a list of data.frame containing the pair of nodes, and the wTO values for all networks that need to be joined. Reference: arXiv:1711.04702

**Usage**

```
wTO.Consensus(data)
```

**Arguments**

data            list of data.frame containing the "Node.1", "Node.2" and "wTO".

**Author(s)**

Deisy Morselli Gysi <deisy at bioinf.uni-leipzig.de>

**Examples**

```
## Not run:
EXAMPLE = wTO.Complete( k =1, n = 200, Data = Microarray_Expression1,
  Overlap = ExampleGRF$x, method = "p")

# Constructing the consensus network
data = list(data.frame(Node.1 = EXAMPLE$wTO$Node.1,
  Node.2 = EXAMPLE$wTO$Node.2,
```



```

wto_sig = EXAMPLE$wTO$wTO_sign,
pvalsig = EXAMPLE$wTO$pval_sig),
data.frame(Node.1 = EXAMPLE$wTO$Node.1,
Node.2 = EXAMPLE$wTO$Node.2,
wtoabs = EXAMPLE$wTO$wTO_abs,
pvalabs = EXAMPLE$wTO$pval_abs) )
CONS = wTO.Consensus(data)

## End(Not run)

```

wTO.export

*wTO.export***Description**

Exports the significant interactions, the wTO weight and pvalues into a .txt file, tab separated. This file can be imported in other visualization tools (Cytoscape for example).

**Usage**

```
wTO.export(DATA, path, sign = TRUE, pvalue = 0.05, padj = 0.05, prop.NA = 0.5)
```

**Arguments**

DATA	Output from the function wTO.Complete or wTO.Consensus.
path	Path and file name where the .txt file should be saved.
sign	Should the network contain the results for the signed network or unsigned? Only for data coming from wTO.Complete.
pvalue	cutoff p-value for the network. Only for data coming from wTO.Complete.
padj	cutoff adjusted p-value for the network. Only for data coming from wTO.Complete.
prop.NA	cutoff proportion of NAs for the network. Only for data coming from wTO.Consensus.

**Examples**

```

## Not run:
EXAMPLE = wTO.Complete( k = 1, n = 200, Data = Microarray_Expression2,
                        Overlap = ExampleGRF$x, method = "p")
wTO.export(EXAMPLE , './EXAMPLE.txt')

#Selection of only the significant ones for the Consensus
Ex_k1_cor_p_boot_p005_sig = subset(EXAMPLE$wTO,
EXAMPLE$wTO$pval_sig < 0.05,
select = c("Node.1", "Node.2", "wTO_sign"))
Ex_k1_cor_p_boot_p005_abs = subset(EXAMPLE$wTO,
EXAMPLE$wTO$pval_abs < 0.05,
select = c("Node.1", "Node.2", "wTO_abs"))
# Constructing the consensus network

```

```

CN = wTO.Consensus(data = list(Ex_k1_cor_p_boot_p005_sig,
Ex_k1_cor_p_boot_p005_abs))
wTO.export(CN, './CN.txt')
### You can store the result on the workspace.
y = wTO.export(CN, './CN.txt')
head(y)

## End(Not run)

```

---

wTO.fast

*wTO.fast*


---

### Description

Compute the wTO and also the bootstraps. Proposed at arXiv:1711.04702. This is a quicker version of the wTO.Complete. It doesn't contain diagnose plots nor a parallel version.

### Usage

```

wTO.fast(
  Data,
  Overlap = row.names(Data),
  method = "p",
  sign = "sign",
  delta = 0.2,
  n = 10,
  method_resampling = "Bootstrap",
  lag = NULL,
  ID = NULL
)

```

### Arguments

Data	data.frame containing the count / expression data for the correlation.
Overlap	Set of nodes of interest, where the Overlapping weights will be computed.
method	Type of the correlation that should be used. "s" / "spearman" will compute the rank spearman correlation, "p" / "pearson" will compute the linear correlation. If no value is given, the default is to use "p".
sign	Should the wTO be signed?
delta	expected difference between the real wTO and the bootstrapped.
n	Number of resamplings, used to compute the empirical distributions of the links. Default is set to 100.

```

method_resampling      method of the resampling. Bootstrap or BlockBootstrap.If the second is used,
                        please give the lag (time dependency among the data).
lag                    Time dependency for the blocked bootstrap (for time series).
ID                     ID of the samples for the blocked bootstrap (for repeated measures).

```

**Author(s)**

Deisy Morselli Gysi <deisy at bioinf.uni-leipzig.de>

**Examples**

```

# wTO.fast(Data = Microarray_Expression1,
# Overlap = ExampleGRF$x,
# method = "p")

# For a time series with lag = 4
# wTO.fast(Data = Microarray_Expression1,
# Overlap = ExampleGRF$x,
# method = "p",
# method_resampling = 'BlockBootstrap',
# lag = 4)

# For a study where the individuals were measured multiple times.
# wTO.fast(Data = Microarray_Expression1,
# Overlap = ExampleGRF$x,
# method = "p",
# method_resampling = 'BlockBootstrap',
# ID = rep(1:9, each= 2))

```

---

wTO.in.line

*wTO.in.line*


---

**Description**

Transforms a correlation matrix into the line format.

**Usage**

```
wTO.in.line(d)
```

**Arguments**

d correlation matrix to be converted into the line format.

**Value**

the wTO matrix into a data.frame: Node1, Node2 and wTO.

**Author(s)**

Deisy Morselli Gysi <deisy at bioinf.uni-leipzig.de>

---

wTO.rep_measure	<i>wTO.rep_measure</i>
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---

**Description**

Compute the wTO for a repeated measures experiment and also the bootstraps. Proposed at arXiv:1711.04702. This is a quicker version of the wTO.Complete. It doesn'T contain diagnose plots nor a parallel version.

**Usage**

```
wTO.rep_measure(
  Data,
  Overlap = row.names(Data),
  ID,
  sign = "sign",
  delta = 0.2,
  n = 10
)
```

**Arguments**

Data	data.frame containing the count / expression data for the correlation.
Overlap	Set of nodes of interest, where the Overlapping weights will be computed.
ID	a vector with the individuals identification
sign	Should the wTO be signed?
delta	expected difference between the real wTO and the bootstraped.
n	Number of resamplings, used to compute the empirical distributions of the links. Default is set to 100.

**Author(s)**

Deisy Morselli Gysi <deisy at bioinf.uni-leipzig.de>

**Examples**

```
#wTO.rep_measure(Data = Microarray_Expression1, ID = rep(c(1:9),2),
#Overlap = ExampleGRF$x)
```

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