

# Package ‘corto’

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**Type** Package

**Title** Inference of Gene Regulatory Networks

**Version** 1.1.4

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**Description** We present 'corto' (Correlation Tool), a simple package to infer gene regulatory networks and visualize master regulators from gene expression data using DPI (Data Processing Inequality) and bootstrapping to recover edges. An initial step is performed to calculate all significant edges between a list of source nodes (centroids) and target genes. Then all triplets containing two centroids and one target are tested in a DPI step which removes edges. A bootstrapping process then calculates the robustness of the network, eventually re-adding edges previously removed by DPI. The algorithm has been optimized to run outside a computing cluster, using a fast correlation implementation. The package finally provides functions to calculate network enrichment analysis from RNA-Seq and ATAC-Seq signatures as described in the article by Giorgi lab (2020) <doi:10.1093/bioinformatics/btaa223>.

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corto	<i>Calculate a regulon from a data matrix</i>
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**Description**

This function applies Correlation and DPI to generate a robust regulon object based on the input data matrix and the selected centroids.

**Usage**

```
corto(
  inmat,
  centroids,
  nbootstraps = 100,
  p = 1e-30,
  nthreads = 1,
  verbose = FALSE,
  cnvmat = NULL,
  boot_threshold = 0
)
```

**Arguments**

inmat	Input matrix, with features (e.g. genes) as rows and samples as columns
centroids	A character vector indicating which features (e.g. genes) to consider as centroids (a.k.a. Master Regulators) for DPI
nbootstraps	Number of bootstraps to be performed. Default is 100
p	The p-value threshold for correlation significance (by default 1E-30)
nthreads	The number of threads to use for bootstrapping. Default is 1
verbose	Logical. Whether to print progress messages. Default is FALSE
cnvmat	An optional matrix with copy-number variation data. If specified, the program will calculate linear regression between the gene expression data in the input matrix (exp) and the cnv data, and target profiles will be transformed to the residuals of each linear model exp~cnv. Default is NULL
boot_threshold	The fraction of bootstraps in which the edge should appear to be included in the final network. It can be any number between 0.0 and 1.0. Default is 0.0.

**Value**

A list (object of class regulon), where each element is a centroid

- tfmode: a named vector containing correlation coefficients between features and the centroid
- likelihood: a numeric vector indicating the likelihood of interaction

**Examples**

```
# Load data matrix inmat (from TCGA mesothelioma project)
load(system.file("extdata", "inmat.rda", package="corto", mustWork=TRUE))
# Load centroids
load(system.file("extdata", "centroids.rda", package="corto", mustWork=TRUE))
# Run corto
regulon <- corto(inmat, centroids=centroids, nthreads=2, nbootstraps=10, verbose=TRUE)

# In a second example, a CNV matrix is provided. The analysis will be run only
# for the features (rows) and samples (columns) present in both matrices
load(system.file("extdata", "cnvmat.rda", package="corto", mustWork=TRUE))
regulon <- corto(inmat, centroids=centroids, nthreads=2, nbootstraps=6, verbose=TRUE, cnvmat=cnvmat,
p=1e-8)
```

---

fcor

*A fast correlation function*


---

**Description**

A fast correlation function

**Usage**

```
fcor(inmat, centroids, r)
```

**Arguments**

inmat	An input matrix with features as rows and samples as columns
centroids	A character vector indicating the centroids
r	A numeric correlation threshold

**Value**

A matrix describing which edges were significant in the input matrix matrix according to the r correlation threshold provided

---

fisherp	<i>Fisher integration of p-values</i>
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**Description**

This function applies the Fisher integration of p-values

**Usage**

```
fisherp(ps)
```

**Arguments**

ps	a vector of p-values
----	----------------------

**Value**

p.val an integrated p-value

**Examples**

```
ps<-c(0.01,0.05,0.03,0.2)
fisherp(ps)
```

gsea

*GSEA***Description**

This function performs Gene Set Enrichment Analysis

**Usage**

```
gsea(
  refflist,
  set,
  method = c("permutation", "pareto"),
  np = 1000,
  w = 1,
  gsea_null = NULL
)
```

**Arguments**

refflist	named vector of reference scores
set	element set
method	one of 'permutation' or 'pareto'
np	Number of permutations (Default: 1000)
w	exponent used to raise the supplied scores. Default is 1 (original scores unchanged)
gsea_null	a GSEA null distribution (Optional)

**Value**

A GSEA object. Basically a list of s components:

**ES** The enrichment score

**NES** The normalized enrichment score

**ledge** The items in the leading edge

**p.value** The permutation-based p-value

**Examples**

```
refflist<-setNames(-sort(rnorm(1000)),paste0('gene',1:1000))
set<-paste0('gene',sample(1:200,50))
obj<-gsea(refflist,set,method='pareto',np=1000)
obj$p.value
```

---

gsea2	<i>2-way GSEA GSEA Gene set enrichment analysis of two complementary gene sets using gsea</i>
-------	---

---

### Description

2-way GSEA GSEA Gene set enrichment analysis of two complementary gene sets using gsea

### Usage

```
gsea2(
  reflist,
  set1,
  set2,
  method = c("permutation", "pareto"),
  np = 1000,
  w = 1,
  gsea_null = NULL
)
```

### Arguments

reflist	named vector of reference scores
set1	element set 1
set2	element set 1
method	one of 'permutation' or 'pareto'
np	Number of permutations (Default: 1000)
w	exponent used to raise the supplied scores. Default is 1 (original scores unchanged)
gsea_null	a GSEA null distribution (Optional)

### Value

A list of 2 GSEA objects. Each of which is a list of components:

**ES** The enrichment score  
**NES** The normalized enrichment score  
**ledge** The items in the leading edge  
**p.value** The permutation-based p-value

### Examples

```
reflist<-setNames(-sort(rnorm(1000)),paste0('gene',1:1000))
set1<-paste0('gene',sample(1:200,50))
set2<-paste0('gene',sample(801:1000,50))
obj<-gsea2(reflist,set1,set2,method='pareto',np=1000)
obj$p.value
```

---

`kmgformat`*kmgformat - Nice Formatting of Numbers*

---

**Description**

This function will convert thousand numbers to K, millions to M, billions to G, trillions to T, quadrillions to P

**Usage**

```
kmgformat(input, roundParam = 1)
```

**Arguments**

<code>input</code>	A vector of values
<code>roundParam</code>	How many decimal digits you want

**Value**

A character vector of formatted numebr names

**Examples**

```
# Thousands
set.seed(1)
a<-runif(1000,0,1e4)
plot(a,yaxt='n')
kmg<-kmgformat(pretty(a))
axis(2,at=pretty(a),labels=kmg)

# Millions to Billions
set.seed(1)
a<-runif(1000,0,1e9)
plot(a,yaxt='n',pch=20,col="black")
kmg<-kmgformat(pretty(a))
axis(2,at=pretty(a),labels=kmg)
```

---

`mra`*Perform Master Regulator Analysis (mra).*

---

**Description**

The analysis is performed between two groups of samples in the form of expression matrices, with genes/features as rows and samples as columns.

**Usage**

```
mra(
  expmat1,
  expmat2 = NULL,
  regulon,
  minsize = 10,
  nperm = NULL,
  nthreads = 2,
  verbose = FALSE,
  atacseq = NULL
)
```

**Arguments**

expmat1	A numeric expression matrix, with genes/features as rows and samples as columns. If only expmat1 is provided (without expmat2), the function will perform a sample-by-sample master regulator analysis, with the mean of the dataset as a reference. If expmat2 is provided, expmat1 will be considered the "treatment" sample set. If a named vector is provided, with names as genes/features and values as signature values (e.g. T-test statistics), signature master regulator analysis is performed.
expmat2	A numeric expression matrix, with genes/features as rows and samples as columns. If provided, it will be considered as the "control" or "reference" sample set for expmat1.
regulon	A <code>_regulon_</code> object, output of the <code>_corto_</code> function.
minsize	A minimum network size for each centroid/TF to be analyzed. Default is 10.
nperm	The number of times the input data will be permuted to generate null signatures. Default is 1000 if expmat2 is provided, and 10 if expmat2 is not provided (single sample mra).
nthreads	The number of threads to use for generating null signatures. Default is 1
verbose	Boolean, whether to print full messages on progress analysis. Default is FALSE
atacseq	An optional 3 column matrix derived from an ATAC-Seq analysis, indicating 1) gene symbol, 2) $-\log_{10}(\text{FDR}) \cdot \text{sing}(\log_2\text{FC})$ of an ATAC-Seq design, 3) distance from TSS. If provided, the output will contain an <code>_atacseq_</code> field.

**Value**

A list summarizing the master regulator analysis

- nes: the normalized enrichment score: positive if the centroid/TF network is upregulated in expmat1 vs expmat2 (or in expmat1 vs the mean of the dataset), negative if downregulated. A vector in multisample mode, a matrix in sample-by-sample mode.
- pvalue: the pvalue of the enrichment.
- sig: the calculated signature (useful for plotting).
- regulon: the original regulon used in the analysis (but filtered for `_minsize_`)



- `atac`: Optionally present if atacseq data is provided. For each centroid/TF a number ranging from 0 to 1 will indicate the fraction of changes in activity due to promoter effects rather than distal effects.

---

mraplot

*Plot a master regulator analysis*


---

### Description

Plotting function for master regulator analysis performed by the `_mra_` function

### Usage

```
mraplot(
  mraobj,
  mrs = 5,
  title = "corto - Master Regulator Analysis",
  pthr = 0.01
)
```

### Arguments

<code>mraobj</code>	The input object, output of the function <code>mra</code>
<code>mrs</code>	Either a numeric value indicating how many MRs to show, sorted by significance, or a character vector specifying which TFs to show. Default is 5
<code>title</code>	Title of the plot (optional, default is "corto - Master Regulator Analysis")
<code>pthr</code>	The p-value at which the MR is considered significant. Default is 0.01

### Value

A plot is generated

---

p2r

*p2r Convert a P-value to the corresponding Correlation Coefficient*


---

### Description

p2r Convert a P-value to the corresponding Correlation Coefficient

### Usage

```
p2r(p, n)
```

**Arguments**

p                    the p-value  
 n                    the number of samples

**Value**

a correlation coefficient

**Examples**

p2r(p=0.08, n=20)

---

p2z

*p2z*

---

**Description**

This function gives a gaussian Z-score corresponding to the provided p-value Careful: sign is not provided

**Usage**

p2z(p)

**Arguments**

p                    a p-value

**Value**

z a Z score

**Examples**

p<-0.05  
 p2z(p)

---

`plot_gsea`*Plot GSEA results*

---

### Description

This function generates a GSEA plot from a gsea object

### Usage

```
plot_gsea(  
  gsea.obj,  
  twoColors = c("red", "blue"),  
  plotNames = FALSE,  
  colBarcode = "black",  
  title = "Running Enrichment Score",  
  bottomTitle = "List Values",  
  bottomYlabel = "Signature values",  
  ext_nes = NULL,  
  omit_middle = FALSE  
)
```

### Arguments

<code>gsea.obj</code>	GSEA object produced by the gsea function
<code>twoColors</code>	the two colors to use for positive[1] and negative[2] enrichment scores
<code>plotNames</code>	Logical. Should the set names be plotted?
<code>colBarcode</code>	The color of the barcode
<code>title</code>	String to be plotted above the Running Enrichment Score
<code>bottomTitle</code>	String for the title of the bottom part of the plot
<code>bottomYlabel</code>	String for the Y label of the bottom plot
<code>ext_nes</code>	Provide a NES from an external calculation
<code>omit_middle</code>	If TRUE, will not plot the running score (FALSE by default)

### Value

Nothing, a plot is generated in the default output device

### Examples

```
reflist<-setNames(-sort(rnorm(1000)),paste0('gene',1:1000))  
set<-paste0('gene',sample(1:200,50))  
obj<-gsea(reflist,set,method='pareto',np=1000)  
plot_gsea(obj)
```

---

`plot_gsea2`*Plot 2-way GSEA results*

---

### Description

This function generates a GSEA plot from a gsea object

### Usage

```
plot_gsea2(  
  gsea.obj,  
  twoColors = c("red", "blue"),  
  plotNames = FALSE,  
  title = "Running Enrichment Score",  
  bottomTitle = "List Values",  
  bottomYlabel = "Signature values"  
)
```

### Arguments

<code>gsea.obj</code>	GSEA object produced by the gsea function
<code>twoColors</code>	the two colors to use for positive[1] and negative[2] enrichment scores, and of the barcodes
<code>plotNames</code>	Logical. Should the set names be plotted?
<code>title</code>	String to be plotted above the Running Enrichment Score
<code>bottomTitle</code>	String for the title of the bottom part of the plot
<code>bottomYlabel</code>	String for the Y label of the bottom plot (FALSE by default)

### Value

Nothing, a plot is generated in the default output device

### Examples

```
reflist<-setNames(-sort(rnorm(1000)),paste0('gene',1:1000))  
set1<-paste0('gene',sample(1:200,50))  
set2<-paste0('gene',sample(801:1000,50))  
obj<-gsea2(reflist,set1,set2,method='pareto',np=1000)  
plot_gsea2(obj)
```

---

r2p	<i>r2p Convert Correlation Coefficient to P-value</i>
-----	---

---

**Description**

r2p Convert Correlation Coefficient to P-value

**Usage**

```
r2p(r, n)
```

**Arguments**

r	the correlation coefficient
n	the number of samples

**Value**

a numeric p-value

**Examples**

```
r2p(r=0.4, n=20) # 0.08
```

---

scatter	<i>scatter - XY scatter plot with extra information</i>
---------	---

---

**Description**

This function will plot two variables (based on their common names), calculate their Coefficient of Correlation (CC), plot a linear regression line and color the background if the correlation is positive (red), negative (blue) or non-significant (white)

**Usage**

```
scatter(
  x,
  y,
  method = "pearson",
  threshold = 0.01,
  showLine = TRUE,
  grid = TRUE,
  bgcol = FALSE,
  pch = 20,
  subtitle = NULL,
  extendXlim = FALSE,
  ...
)
```

**Arguments**

x	The first named vector
y	The second named vector
method	a character string indicating which correlation coefficient is to be computed. One of "pearson" (default), "kendall", or "spearman": can be abbreviated.
threshold	a numeric value indicating the significance threshold (p-value) of the correlation, in order to show a colored background. Default is 0.01.
showLine	a boolean indicating if a linear regression line should be plotted. Default is TRUE
grid	a boolean indicating whether to show a plot grid. Default is TRUE
bgcol	Boolean. Should a background coloring associated to significance and sign of correlation be used? Default is TRUE, and it will color the background in red if the correlation coefficient is positive, in blue if negative, in white if not significant (according to the <code>_threshold_</code> parameter)
pch	the <code>_pch_</code> parameter indicating the points shape. Default is 20
subtitle	NULL by default, in which case the function will print as a subtitle the correlation coefficient (CC) and its pvalue. Otherwise, a user-provided string, bypassing the predefined subtitle
extendXlim	logical. If TRUE, the x-axis limits are extended by a fraction (useful for labeling points on the margins of the plot area). Default is FALSE
...	Arguments to be passed to the core <code>_plot_</code> function

**Value**

A plot

**Examples**

```
x<-setNames(rnorm(200),paste0("var",1:200))
y<-setNames(rnorm(210),paste0("var",11:220))
scatter(x,y,xlab="Variable x",ylab="Variable y",main="Scatter plot by corto package")
```

---

slice

*Slice*

---

**Description**

This function prints a slice of a matrix

**Usage**

```
slice(matrix)
```

**Arguments**

matrix            A matrix

**Value**

A visualization of the first 5 rows and columns of the input matrix

**Examples**

```
set.seed(1)
example<-matrix(rnorm(1000),nrow=100,ncol=10)
slice(example)
```

---

ssgsea	<i>ssGSEA</i>
--------	---------------

---

**Description**

This function performs single sample GSEA

**Usage**

```
ssgsea(inmat, groups, scale = TRUE, minsize = 10)
```

**Arguments**

inmat            A numeric matrix, with rownames/rows as genes or features, and colnames/columns as sample names

groups           a named list. Names are names of the groups (e.g. pathways) and elements are character vectors indicating gene or feature names (that should match, at least partially, with the rownames of inmat)

scale            Boolean. Whether the matrix should be row-scaled.

minsize          Numeric. Include only groups with at least this many elements Default is 10

**Value**

A matrix of Normalized Enrichment Scores (NES), which can be converted to p-values using the function `_corto::z2p_`

## Examples

```
# A random matrix
set.seed(1)
inmat<-matrix(rnorm(200*50),nrow=200,ncol=50)
rownames(inmat)<-paste0("gene",1:nrow(inmat))
# A random list of groups
groups<-list()
for(i in 1:10){
  somegenes<-sample(rownames(inmat),30)
  groups[[paste0("pathway_",i)]]<-somegenes
}
# Run ssGSEA
nesmat<-ssgsea(inmat,groups)
```

---

stouffer

*Stouffer integration of Z scores*

---

## Description

This function gives a gaussian Z-score corresponding to the provided p-value Careful: sign is not provided

## Usage

```
stouffer(x)
```

## Arguments

x                    a vector of Z scores

## Value

Z an integrated Z score

## Examples

```
zs<-c(1,3,5,2,3)
stouffer(zs)
```



---

`val2col`*val2col - Convert a numeric vector into colors*

---

**Description**

`val2col` - Convert a numeric vector into colors

**Usage**

```
val2col(  
  z,  
  col1 = "navy",  
  col2 = "white",  
  col3 = "red3",  
  nbreaks = 1000,  
  center = TRUE,  
  rank = FALSE  
)
```

**Arguments**

<code>z</code>	a vector of numbers
<code>col1</code>	a color name for the min value, default 'navy'
<code>col2</code>	a color name for the middle value, default 'white'
<code>col3</code>	a color name for the max value, default 'red3'
<code>nbreaks</code>	Number of colors to be generated. Default is 30.
<code>center</code>	boolean, should the data be centered? Default is TRUE
<code>rank</code>	boolean, should the data be ranked? Default is FALSE

**Value**

a vector of colors

**Examples**

```
a<-rnorm(1000)  
cols<-val2col(a)  
plot(a,col=cols,pch=16)
```

---

`wstouffer`*Weighted Stouffer integration of Z scores*

---

**Description**

This function gives a gaussian Z-score corresponding to the provided p-value Careful: sign is not provided

**Usage**

```
wstouffer(x, w)
```

**Arguments**

`x` a vector of Z scores  
`w` weight for each Z score

**Value**

Z an integrated Z score

**Examples**

```
zs<-c(1,-3,5,2,3)  
ws<-c(1,10,1,2,1)  
wstouffer(zs,ws)
```

---

`z2p`*z2p*

---

**Description**

This function gives a gaussian p-value corresponding to the provided Z-score

**Usage**

```
z2p(z)
```

**Arguments**

`z` a Z score

**Value**

a p-value

$z2p$

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**Examples**

$z < -1.96$   
 $z2p(z)$

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