# Package 'HiCociety'

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<b>Description</b> Identifies chromatin interaction modules by constructing a Hi-C contact network based on statistically significant interactions, followed by network clustering. The method enables comparison of module connectivity across two Hi-C datasets and is capable of detecting cell-type-specific regulatory modules. By integrating network analysis with chromatin conformation data, this approach provides insights into the spatial organization of the genome and its functional implications in gene regulation. Author: Sora Yoon (2025) <a href="https://github.com/ysora/HiCociety">https://github.com/ysora/HiCociety</a> .			
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add\_Genes

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Add gene information

# Description

This function adds a column with a list of genes included in each locus to the ModuleSummary data frame of the hic2community function.

# Usage

```
add_Genes(df, speciesObj)
```

# **Arguments**

df The ModuleSummary data frame obtained by running hic2community function species0bj Any Txdb package name corresponding

# **Details**

Adding gene list to ModuleSummary data frame obtained from hic2community function.

### Value

A data.frame identical to the input, with an additional "Genes" column. Each entry in this column lists the gene(s) that overlap with the corresponding genomic region. If multiple genes are present, they are concatenated with commas.

#### Author(s)

Sora Yoon, PhD

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### **Examples**

```
modulefile = system.file('extdata','mouse_naiveCD4T_Vahedi_short.rds',
package = 'HiCocietyExample')
mycom = readRDS(modulefile)
mycom$ModuleSummary = add_Genes(mycom$ModuleSummary,
'TxDb.Mmusculus.UCSC.mm10.knownGene')
```

calculate\_avg\_count

Calculate Average Count within 5-pixel Padding

# **Description**

This function calculates the average count within a 25kb padding around each (x, y) coordinate pair.

# Usage

```
calculate_avg_count(x, y, counts, resol)
```

### **Arguments**

x Numeric vector of x-coordinates of contact frequency data frame.

y Numeric vector of y-coordinates of contact frequency data frame.

counts Numeric vector of contact frequency counts.

resol Integer specifying the HiC resolution.

# Value

A numeric vector of average counts.

# **Examples**

```
x <- c(1, 2, 3, 4, 5)
y <- c(1, 2, 3, 4, 5)
counts <- c(10, 20, 30, 40, 50)
resol <- 10000
calculate_avg_count(x, y, counts, resol)</pre>
```

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check\_package

Check if a package is installed

# **Description**

This function checks whether a package is installed. If the package is not installed, it informs the user that the package is missing.

# Usage

```
check_package(package)
```

# Arguments

package

The name of the package to check.

# **Details**

This function checks whether a package is installed. If the package is not installed, it informs the user that the package is missing.

### Value

A logical value: TRUE if the package is installed, FALSE otherwise.

### Author(s)

Sora Yoon, PhD

# **Examples**

```
check_package('dplyr')
```

ConnectivityDiff

Connectivity difference between two conditions

# **Description**

output table of connectivity difference of modules between cell types is generated.

# Usage

```
ConnectivityDiff(wt, ko, prefix.wt, prefix.ko, resolution = 5000)
```

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# **Arguments**

wt	hic2community result from condition 1
ko	hic2community result from condition 2

prefix.wt Prefix for wt to be presented in the column names prefix.ko Prefix for ko to be presented in the column names

resolution Resolution of Hi-C dataset

#### **Details**

Connectivity difference between two conditions

#### Value

A list of two data. frame objects, each representing the network connectivity differences of modules in condition 1 or condition 2 when compared to the counterpart cell type. Each data. frame contains the following columns: "chr", "module\_start", "module\_end", "connectivity", "transitivity", "centrality\_node", "idx" (the row index of the module in the input module object), "connectivity\_in\_(counterpart\_connectivity\_difference", and "connectivity\_foldchange".

#### Author(s)

Sora Yoon, PhD

### **Examples**

```
modulefile1 = system.file('extdata', 'mouse_naiveCD4T_Vahedi_short.rds',
package = 'HiCocietyExample')
modulefile2 = system.file('extdata', 'mouse_Th1_Vahedi_short.rds',
package = 'HiCocietyExample')
mycom1 = readRDS(modulefile1)
mycom2 = readRDS(modulefile2)
result = ConnectivityDiff(mycom1, mycom2, 'NaiveCD4T', 'Th1',
resolution = 5000)
head(print(result))
```

getContactFrequency

Get contact Frequency

# Description

Retrieve contact frequency from .hic file using strawR package.

# Usage

```
getContactFrequency(fname, chr, resol)
```

# **Arguments**

fname .hic data for any types of genome conformation capture data

chr Chromosome number of network extraction resol DNA basepair Resolution. Default=10000

#### **Details**

Get Contact Frequncy from .hic file

### Value

A data.frame containing three columns: x and y (genomic coordinate pairs), and counts (the contact frequency between them).

# Author(s)

Sora Yoon, PhD

# **Examples**

```
myhic=system.file('extdata', 'example.hic', package ='HiCocietyExample')
A = getContactFrequency(myhic,19,5000)
head(print(A))
```

getContactProbability Contact probability

### **Description**

It estimates contact probability based on the distance of a pair of a loci.

# Usage

```
getContactProbability(
  tab,
  farthest = 2000000,
  resol = 10000,
  prob,
  n_cores = NULL
)
```

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# **Arguments**

tab Output from getContactFrequency function.

farthest Maximum 1-D distance to search. Default=2Mb

resol Hi-C resolution for test. Default = 10000

prob Significance cutoff for negative binomial distribution. Default =0.975

n\_cores The number of cores used for parallel computing. If set as NULL, n\_cores is

automatically set to the number of cores in the computer if it is not exceed 30.

If it is more than 30, it is set as 30. Default = NULL

#### **Details**

Get Contact probablity

### Value

A list containing three objects: AREA, original, and len1, representing the statistical significance of each chromatin interaction pair.

#### Author(s)

Sora Yoon, PhD

### **Examples**

```
# This example might take a long time to run, so we wrap it in donttest{}

myhic = system.file('extdata','example.hic',package = 'HiCocietyExample')
mydf=getContactFrequency(myhic, 19, 5000);
myprob=getContactProbability(mydf,farthest=2000000, resol=5000,prob=0.975,
n_cores=2);
```

getElbowPoint

Estimation of elbow point from J-shaped-curve

# **Description**

It provides a point of the highest curvature from a J-shaped-plot

## Usage

```
getElbowPoint(numbers)
```

### **Arguments**

numbers Numeric vector

get\_all\_chr\_names

### **Details**

Estimation of elbow point from J-shaped curve

#### Value

A list containing two elements: index, the index of a point in a sorted vector of numbers in descending order, representing the point where the tangent is closest to one, and ConnectivityCutoff, the corresponding value at that point.

# Author(s)

Sora Yoon, PhD

# **Examples**

```
modulefile = system.file('extdata','mouse_naiveCD4T_Vahedi_short.rds',
package = 'HiCocietyExample')
mycom = readRDS(modulefile)
connec = mycom$ModuleSummary$connectivity
getElbowPoint(connec)
```

get\_all\_chr\_names

Retrieve chromosome names from .hic file

### **Description**

It retrieves all chromosome names having longer than 2.5Mbp.

# Usage

```
get_all_chr_names(fname)
```

### **Arguments**

fname

Path to .hic file

### **Details**

To extract all chromosome names from .hic file

# Value

A character vector containing the names of chromosomes whose genomic lengths exceed 2.5 Mbp.

# Author(s)

Sora Yoon, PhD

get\_txdb

# **Examples**

```
myhic=system.file('extdata', 'example.hic', package ='HiCocietyExample')
get_all_chr_names(myhic)
```

get\_txdb

All available Txdb

# Description

It finds all available Txdb packages used in add\_Genes function.

# Usage

```
get_txdb()
```

### **Details**

Check all available Txdb package

# Value

A character vector containing the names of all available TxDb packages.

# Author(s)

Sora Yoon, PhD

# **Examples**

```
get_txdb()
```

hic2community

Create module objects from the Hi-C data

# Description

It generates a list of graph of significant interactions, module table and module elements.

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### Usage

```
hic2community(
   fname,
   chr,
   resol,
   nbprob,
   farthest,
   par.noise = 1,
   network.cluster.method = "louvain",
   n_cores = NULL
)
```

### **Arguments**

fname Path to .hic file

chr chromosome numbers to run.

resol Resolution of Hi-C data

nbprob Negative binomial probability. Higher value gives smaller number of stronger

interaction.

farthest The maximum searching distance between two nodes

par.noise Parameter for noise removal. Default is 1, higher value gives more filtered in-

teractions.

network.cluster.method

Can select between 'louvain' as default and 'label\_prop' which means the label

propagation method.

n\_cores The number of cores used for parallel computing. If set as NULL, n\_cores is

automatically set to the number of cores in the computer if it is not exceed 30.

If it is more than 30, it is set as 30. Default = NULL

#### **Details**

It generates a list of graph of significant interactions, module table and module elements.

#### Value

A list containing three elements: Graphs (an igraph object representing significant chromatin interactions for each chromosome), ModuleSummary (a data.frame containing information about chromatin interaction modules), and ModuleElements (a list of nodes forming significant chromatin interactions within each module).

### Author(s)

Sora Yoon, PhD

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### **Examples**

```
# This example might take a long time to run, so we wrap it in donttest{}

myhic=system.file('extdata', 'example.hic', package ='HiCocietyExample')
mycom = hic2community(myhic, "19", 5000, 0.975, 20000000,
par.noise=1, 'louvain', n_cores=2)
```

hic2network

HiC to network data format

# Description

It converts Hi-C dataframe to network object.

### Usage

```
hic2network(ftab)
```

## **Arguments**

ftab

three-column data composed of locus1, locus2 and value

#### **Details**

Convert HiC to network data format

### Value

An igraph object representing statistically significant chromatin interactions.

# Author(s)

Sora Yoon, PhD

# **Examples**

```
# This example might take a long time to run, so we wrap it in donttest{}

myhic=system.file('extdata', 'example.hic', package ='HiCocietyExample')
ftab=getContactFrequency(myhic,19,5000);
net = hic2network(ftab[1:100,]);
plot(net)
```

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visualizeModule

Visualization of module

# Description

It draws a triangle heatmap and arcplot of a module

# Usage

```
visualizeModule(
  hicpath,
 HC.object,
 moduleNum,
 resolution,
 hic.norm,
 heatmap.color.range = NULL,
 heatmap.color = colorRampPalette(c("white", "red")),
 arc.depth = 10,
 arc.color = "gray80",
 nbnom.param = 0.99,
  txdb = "TxDb.Mmusculus.UCSC.mm10.knownGene",
  gene.strand.arrow.lwd = 3,
 gene.strand.lwd = 6,
 col.forward.gene = "purple",
  col.reverse.gene = "pink",
  highlight.centrality = FALSE,
 highlight.cent.col = FALSE,
 highlight.node = NULL,
 highlight.node.col = NULL,
  show.sig.int = TRUE,
  netinfo
)
```

# **Arguments**

hicpath	Path to the .hic file			
HC.object	The object name from hic2community result			
moduleNum	The row index of module to draw			
resolution	Resolution of HiC data			
hic.norm	Normalization method. If not, set 'NONE'			
heatmap.color.range				
	Min and max value of contact frequency, e.g., c(0,10)			
heatmap.color	$Color\ for\ heatmap.\ For\ example,\ colorRampPalette(c("white","red))$			
arc.depth	Height of arc plot			
arc.color	Arc color			

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nbnom.param Negative binomial probability cutoff. Higher cutoff gives less number of arcs.

txdb Character. One of Txdb list obtained from get\_txdb().

gene.strand.arrow.lwd

Numeric. Line width of arrowhead indicating the strands of genes. Same as arr.lwd option in Arrows function in shape package.

gene.strand.lwd

Numeric. Line width of arrow body indicating the strands of genes. Same as lwd option in Arros function in shape package.

col.forward.gene

Character. Color of arrows within gene track for forward genes.

col.reverse.gene

Character. Color of arrows within gene track for reverse genes

highlight.centrality

Boolean input to set if highlight eigenvector centrality node.

highlight.cent.col

The color of arcs stemming from the centrality node.

highlight.node The coordiante of a node of which the user will highlight the arcs stemming from this node. Default=NULL

highlight.node.col

The color of arcs stemming from the node which the user highlight.

show.sig.int Boolean. If TRUE, it marks significant contact on the triangle heatmap.

netinfo Boolean. If TRUE, it shows network information of the module as text in the plot.

#### **Details**

Visualization of module

# Value

No return value; the function generates a plot.

# Author(s)

Sora Yoon, PhD

# Examples

```
# A slow example that takes too long to run, wrapped in donttest{}

myhic = system.file('extdata','example.hic',package = 'HiCocietyExample')
HC.object = hic2community(myhic, "19", 5000, 0.975, 2000000, par.noise=1,
'louvain', n_cores=2)
mNum = 1
visualizeModule(hicpath = myhic, HC.object = HC.object, moduleNum = mNum,
resolution = 5000,
hic.norm = 'NONE', heatmap.color.range=c(0,10),
heatmap.color = colorRampPalette(c('white','red')),
```

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arc.depth=10, arc.color = "gray80", nbnom.param=0.99,
txdb = 'TxDb.Mmusculus.UCSC.mm10.knownGene',
gene.strand.arrow.lwd = 3, gene.strand.lwd = 3,
col.forward.gene = 'purple', col.reverse.gene = 'pink',
highlight.centrality=FALSE, highlight.cent.col=FALSE,
highlight.node=NULL, highlight.node.col=NULL,
show.sig.int=FALSE, netinfo=FALSE)

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