

Package ‘TranScan’

September 19, 2020

Type Package
Title Translocation Detection from Hi-C data via Scan Statistics
Version 1.0
Date 2020-09-19
Author Anthony Cheng, Disheng Mao, Yuping Zhang, Joseph Glaz, Zhengqing Ouyang
Maintainer
Yuping Zhang <yuping.zhang@uconn.edu> and Zhengqing Ouyang <ouyang@schoolph.umass.edu>
Description
R-package for the paper entitled ``Translocation Detection from Hi-C data via Scan Statistics" by Anthony Cheng, Disheng Mao, Yuping Zhang, Joseph Glaz, Zhengqing Ouyang
Imports Matrix, dplyr, ggplot2, ggpubr, MASS, reshape2
Depends R (>= 3.2.3)
License GPL (>= 3)
RoxygenNote 6.1.1.9000
NeedsCompilation no

R topics documented:

chr10	1
chr10_20	2
chr20	2
TranScan	3
Index	4

chr10	<i>Chromosome 10 of T47D</i>
-------	------------------------------

Description

Intra region of chromosome 10 in cell line T47D (500kb resolution)

Usage

data(chr10)

Format

The third column refers to rounded log count

chr10_20	<i>Chromosome 10_20 of T47D</i>
----------	---------------------------------

Description

Inter region between chromosome 10 and 20 in cell line T47D (500kb resolution)

Usage

```
data(chr10)
```

Format

The third column refers to rounded log count

chr20	<i>Chromosome 20 of T47D</i>
-------	------------------------------

Description

Intra region of chromosome 20 in cell line T47D (500kb resolution)

Usage

```
data(chr20)
```

Format

The third column refers to rounded log count

TranScan

*Hi-C inter-chromosomal translocation detection via Scan clustering***Description**

This function allows you to detect the inter-chromosomal translocation from Hi-C data. It requires two intra-regions and the corresponding inter-region as inputs. We suggest rounded log counts as inputs.

Usage

```
TranScan(chr1, chr2, chr1_2, outputpath, sizes, resolution = 5e+05,
  quantitest = 256, alpha = 0.05, gammath = 0.1,
  number_of_bandwidth = 20, diag_remove = 2e+06, add_gap = 2500000)
```

Arguments

chr1	First chromosome
chr2	Second chromosome
chr1_2	Inter-chromosomal region of chr1 and chr2
outputpath	The path to store all results
sizes	Chrom sizes of chr1_2; vector of length 2
resolution	Resolution of Hi-C data
quantitest	Grid of testing, must be 2^k , default 256, meaning $256 * 256$
alpha	FDX, default 0.05
gammath	Gamma threshold, default 0.1
number_of_bandwidth	Number of bandwidths used, default 20
diag_remove	Removed short-range interactions, must be $k * \text{resolution}$, default $2e+06$
add_gap	Added gap between two chromosomes, must be $k * \text{resolution}$, default 2500000

Value

A heatmap and a rejected region plot will be generated under the output folder. The matrix form of rejected region is saved as RR.txt. All intermediate objects will also be saved.

Examples

```
data(chr10)
data(chr20)
data(chr10_20)
TranScan(chr10, chr20, chr10_20, "./test", sizes=c(133797422, 64444167))
```

Index

*Topic **datasets**

chr10, [1](#)

chr10_20, [2](#)

chr20, [2](#)

chr10, [1](#)

chr10_20, [2](#)

chr20, [2](#)

TranScan, [3](#)