## rGMAP

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Type Package
Title Call hierarchical chromatin domains from HiC matrix by GMAP
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<b>Description</b> Call hierarchical chromatin domains from HiC contact matrix by Gaussian Mixture model And Proportion test
BugReports https://github.com/wbaopaul/rGMAP/issues
License GPL (>= 2)
LazyData TRUE
Imports data.table, ggplot2, mclust, EMD, caTools, Matrix, Rcpp (>= 0.12.5)
LinkingTo Rcpp
RoxygenNote 6.0.1
Suggests knitr, rmarkdown
VignetteBuilder knitr
R topics documented:
data_simu
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data\_simu

generate simulated hic\_mat and true tads

## **Description**

generate simulated hic\_mat and true tads

#### Usage

```
data_simu(stype = "poisson-dist", nratio = 2.5, mu0 = 200, resl = 1)
```

#### **Arguments**

Stype One of four types of simulated data in the manuscript: poission-dist, poission-

dist-hier, nb-dist, nb-dist-hier; poission- or nb- indicates poission distribution or negative bionomial distribution -hier indicated subtads are generated nestly

nratio The effect size between intra- and inter domain, larger means higher intra-tad

contacts

mu 0 The mean parameter, default 200

resl Resolution, default set to 1

#### Value

A list includes following elements:

hic\_mat n by n contact matrix
hierTads True heirarchical domains

tads\_true True TADs

hic\_rao\_IMR90\_chr15

Normalized Hi-C data for IMR90, chr15 with resolution 10kb.

## Description

Normalized Hi-C data for IMR90, chr15 with resolution 10kb.

## Usage

```
hic_rao_IMR90_chr15
```

#### **Format**

A data table with 3 variables:

**n1** bin 1

**n2** bin 2

count normalized counts

plotdom 3

#### **Source**

Rao et al., Cell 2014, A 3D map of the human genome at kilobase resolution reveals principles of chromatin looping

plotdom

visualize hierarchical domains

## **Description**

visualize hierarchical domains

## Usage

```
plotdom(hic_dat, hiertads_gmap, start_bin, end_bin, cthr = 20, resl = 10000)
```

## **Arguments**

hic\_dat hic contact matrix for a given chromosome, either a n by n matrix, or a 3 columns

data.frame <bin1> <bin2> <counts>

hiertads\_gmap

the hierarchical domains called by GMAP

start\_bin the start bin of the genome end\_bin the end bin of the genome

cthr the upper bound count threshold for color, default 20

reslution of Hi-C data, default 10000

rGMAP

Detect hierarchical choromotin domains by GMAP

## **Description**

Detect hierarchical choromotin domains by GMAP

## Usage

```
rGMAP(hic_mat, resl = 10 * 10^3, logt = T, dom_order = 2, maxDistInBin = min(200, 2 * 10^6/resl), min_d = 25, max_d = 100, min_dp = 5, max_dp = 10, hthr = 0.95, t1thr = 0.5)
```

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#### **Arguments**

Either a 3 columns Hi-C contact matrix for a given chromosome, with each row hic\_mat corrsponding to the start bin, end bin and the contact number; or a n by n matrix, n is the number of bins for a given chromosom The resolution (bin size), default 10kb resl logt Do log-transformation or not, default TRUE Maximum level of hierarchical structures, default 2 (call TADs and subTADs) dom\_order maxDistInBin Only consider contact whose distance is not greater than maxDistInBIn bins, default 200 bins (or 2Mb) The minimum d (d: window size), default 25 min\_d The maximum d (d: window size), default 100  $max_d$ The minmum dp (dp: lower bound of tad size), defalt 5 min\_dp max\_dp The maximum dp (dp: lower bound of tad size), defalt 10. min\_d, max\_d, min\_dp and max\_dp should be specified in number of bins The lower bound cutoff for posterior probability, default 0.95 hthr t1thr Lower bound for t1 for calling TAD, default 0.5 quantile of test statistics of TADs, 0.9 of subTADs

#### Value

#### A list includes following elements:

tads A data frame with columns start, end indicates the start and end coordinates of

each domain, respectively

hierTads A data frame with columns start, end, dom\_order, where dom\_order indicates

the hierarchical status of a domain, 1 indicates tads, 2 indicates subtads, and so

on

params A data frame gives the final parameters for calling TADs

#### **Examples**

```
## On simulated data:
library(rGMAP)
simu_res = data_simu('poisson-dist-hier')
true_domains = simu_res$hierTads
simu_mat = simu_res$hic_mat
predicted_domains = rGMAP(simu_mat, resl = 1)$hierTads
true_domains
predicted_domains
## On an real data example
hic_rao_IMR90_chr15
                      # normalized Hi-C data for IMR90, chr15 with resolution 10kb
res = rGMAP(hic_rao_IMR90_chr15, resl = 10 * 1000, dom_order = 2)
names (res)
#quickly visualize some hierarchical domains
pp = plotdom(hic_rao_IMR90_chr15, res$hierTads, 6000, 7000, 30, 10)
pp$p2
```

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