

# rGMAP

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**Type** Package

**Title** Call hierarchical chromatin domains from HiC matrix by GMAP

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**Author** Wenbao Yu

**Maintainer** Wenbao Yu <yuw1@email.chop.edu>

**Description** Call hierarchical chromatin domains from HiC contact map 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** 6.1.0

**Suggests** knitr,  
rmarkdown

**VignetteBuilder** knitr

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data\_simu                    *generate simulated hic\_mat and true tads*

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### 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: poisson-dist, poisson-dist-hier, nb-dist, nb-dist-hier; poisson- or nb- indicates poisson distribution or negative binomial distribution -hier indicated subtads are generated nestly
nratio	The effect size between intra- and inter domain, larger means higher intra-tad contacts
mu0	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

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hic\_rao\_IMR90\_chr15

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

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### 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

**Source**

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

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plotdom	<i>visualize hierarchical domains</i>
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**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
cth	the upper bound count threshold for color, default 20
resl	reslution of Hi-C data, default 10000

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rGMAP	<i>Detect hierarchical choromotin domains by GMAP</i>
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**Description**

Detect hierarchical choromotin domains by GMAP

**Usage**

```
rGMAP(hic_mat, index_file = NULL, 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)
```

**Arguments**

<code>hic_mat</code>	<ul style="list-style-type: none"> <li>• For single chromosome, supports three types of format: <ul style="list-style-type: none"> <li>– a 3-column Hi-C contact matrix, with columns the <code>i_th</code>, <code>j_th</code> bin of a chromosome and the corresponding contact number</li> <li>– a <code>n</code> by <code>n</code> matrix, with <code>&lt;i,j&gt;</code>th element corresponding to contact number between the <code>i_th</code> and <code>j_th</code> bin of the chromosome</li> <li>– a text file of the above two types of data</li> </ul> </li> <li>• For multiple chromosomes, a <code>index_file</code> indicates genomic coordinate for each hic bin should be provided</li> </ul>
<code>index_file</code>	A 4-columns tab/space delimited text file indicates the genomic coordinates for each bin (compatible with HiC-Pro); with columns <code>bin_chr bin_start bin_end bin_id</code>
<code>resl</code>	The resolution (bin size), default 10kb
<code>logt</code>	Do log-transformation or not, default TRUE
<code>dom_order</code>	Maximum level of hierarchical structures, default 2 (call TADs and subTADs)
<code>maxDistInBin</code>	Only consider contact whose distance is not greater than <code>maxDistInBin</code> bins, default 200 bins (or 2Mb)
<code>min_d</code>	The minimum <code>d</code> ( <code>d</code> : window size), default 25
<code>max_d</code>	The maximum <code>d</code> ( <code>d</code> : window size), default 100
<code>min_dp</code>	The minimum <code>dp</code> ( <code>dp</code> : lower bound of tad size), default 5
<code>max_dp</code>	The maximum <code>dp</code> ( <code>dp</code> : lower bound of tad size), default 10. <code>min_d</code> , <code>max_d</code> , <code>min_dp</code> and <code>max_dp</code> should be specified in number of bins
<code>hthr</code>	The lower bound cutoff for posterior probability, default 0.95
<code>t1thr</code>	Lower bound for <code>t1</code> for calling TAD, default 0.5 quantile of test statistics of TADs, 0.9 of subTADs

**Value**

A list includes following elements:

<code>tads</code>	A data frame with columns <code>start</code> , <code>end</code> indicates the start and end coordinates of each domain, respectively
<code>hierTads</code>	A data frame with columns <code>start</code> , <code>end</code> , <code>dom_order</code> , where <code>dom_order</code> indicates the hierarchical status of a domain, 1 indicates tads, 2 indicates subtads, and so on
<code>params</code>	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, res1 = 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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