Package ‘SIMD’

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Type Package

Title Statistical Inferences with MeDIP-seq Data (SIMD) to infer the methylation level for each CpG site

Version 1.22.0

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Description This package provides a inferential analysis method for detecting differentially expressed CpG sites in MeDIP-seq data. It uses statistical framework and EM algorithm, to identify differentially expressed CpG sites. The methods on this package are described in the article 'Methylation-level Inferences and Detection of Differential Methylation with Medip-seq Data' by Yan Zhou, Jiadi Zhu, Mingtao Zhao, Baoxue Zhang, Chunfu Jiang and Xiyan Yang (2018, pending publication).

License GPL-3

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SIMD-package  A method to infer the methylation expression level for each CpG sites.

Description

SIMD is a package to infer the methylation expression level for each CpG sites. The main idea of SIMD is that by using statistical inference to with Medip-seq data method to infer the methylation level.

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References

Description

This data set gives 2000 CpG sites which include the chromosome of the region, the start and the stop position of the CpG sites.

Usage

all_CpGsite_bin_chr18

Format

A data.frame containing 2000 CpG sites.

Source


References


classifypvalue

calculate P-value in code EMtest.

Description

calculate P-value in code EMtest.

Usage

classifypvalue(type1, type2, type3, type4, sm1chring1, sm1chring2, sm1chring3, sm1chring4, p, typelength, sm1chringlength, pvalue = rep(0, length(sm1chring1)))
EM_H1ESB1_MeDIP_sigeCpG

Arguments

<table>
<thead>
<tr>
<th>Argument</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>type1</td>
<td>The first column of the first matrix.</td>
</tr>
<tr>
<td>type2</td>
<td>The second column of the first matrix.</td>
</tr>
<tr>
<td>type3</td>
<td>The third column of the first matrix.</td>
</tr>
<tr>
<td>type4</td>
<td>The fourth column of the first matrix.</td>
</tr>
<tr>
<td>sm1chring1</td>
<td>The first column of the second matrix.</td>
</tr>
<tr>
<td>sm1chring2</td>
<td>The second column of the second matrix.</td>
</tr>
<tr>
<td>sm1chring3</td>
<td>The third column of the second matrix.</td>
</tr>
<tr>
<td>sm1chring4</td>
<td>The fourth column of the second matrix.</td>
</tr>
<tr>
<td>p</td>
<td>P-value.</td>
</tr>
<tr>
<td>typelength</td>
<td>The number of rows of the first matrix.</td>
</tr>
<tr>
<td>sm1chringlength</td>
<td>The number of rows of the second matrix.</td>
</tr>
<tr>
<td>pvalue</td>
<td>A vector, the length equals to the number of rows of the second matrix.</td>
</tr>
</tbody>
</table>

Value

The probability.

Description

A simulation dataset of MeDIP CpG sites.

This data set gives 2000 CpG sites which include the chromosome of the region, the start and the stop position of the MeDIP CpG sites.

Usage

EM_H1ESB1_MeDIP_sigeCpG

Format

A data.frame containing 2000 MeDIP CpG sites.

Source


References

EM2_H1ESB1_MeDIP_sigleCpG

Description
This data set gives 2000 CpG sites which include the chromosome of the region, the start and the stop position of the MeDIP CpG sites.

Usage
EM2_H1ESB1_MeDIP_sigleCpG

Format
A data.frame containing 2000 MeDIP CpG sites.

Source

References

EM2_H1ESB2_MeDIP_sigleCpG

Description
This data set gives 2000 CpG sites which include the chromosome of the region, the start and the stop position of the MeDIP CpG sites.

Usage
EM2_H1ESB2_MeDIP_sigleCpG

Format
A data.frame containing 2000 MeDIP CpG sites.
EMalgorithm

Source


References


---

EMalgorithm

**EM algorithm to infer CpG sites.**

**Description**

Using EM algorithm to infer the real number of CpG sites.

**Usage**

```r
EMalgorithm(cpgsitefile, allcpgfile, category = "1", writefile = NULL, reportfile = NULL)
```

**Arguments**

- `cpgsitefile`: The path of file to store CpG site.
- `allcpgfile`: The file to store CpG sites.
- `category`: Default to "1".
- `writefile`: The path of output results. (If writefile=NULL, there will return the results back to main program.)
- `reportfile`: The path of output results.

**Value**

values or file If writefile is NULL, then return the values of results, otherwise output to write file.

**Examples**

```r
datafile <- system.file("extdata", package="methylMnM")
data(example_data)
filepath <- datafile[1]
allcpgfile <- EM_H1ESB1_MeDIP_sigleCpG
dirwrite <- paste(setwd(getwd()), "/", sep="")
readshort <- paste(filepath, "/H1ESB1_MeDIP_18.extended.txt", sep="")
writefile <- paste(dirwrite, "EM2_H1ESB1_MeDIP_sigleCpG.bed", sep="")
reportfile <- paste(dirwrite, "EM2_H1ESB1_MeDIP_sigleCpG_report.bed", sep="")
f <- EMalgorithm(cpgsitefile=readshort, allcpgfile=allcpgfile, category="1", writefile=writefile, reportfile=reportfile)
```
emalght

**Calculate the probability on condition that the sums equal to 1.**

**Description**
Calculate the probability on condition that only a single CpG contributes to a short read.

**Usage**
emalght(X)

**Arguments**
X A matrix about X, the elements in X takes values on 0,1 and satisfy the sums of each row equal to 1.

**Value**
y1 The probability when sums equal to 1.

**Examples**
```r
set.seed(123)
d <- matrix(0, nrow=200, ncol=50)
random_num <- sample(1:50, 200, replace=TRUE)
for(i in 1:nrow(d)){
  d[i,random_num[i]]<-1
}
result <- emalght(d)
head(result)
```

emalght1

**Calculate the probability on condition that the sums more than 1.**

**Description**
Calculate the probability on condition that at least a CpG contributes to a short read.

**Usage**
emalght1(X)

**Arguments**
X A matrix about X, the elements in X takes values on 0,1 and satisfy the sums of each row more than 1.
Value

\[ y_1 \] The probability when sums more than 1.

Examples

```r
set.seed(123)
d <- matrix(0, nrow=200, ncol=50)
random_num <- sample(1:10, 200, replace=TRUE)
for(i in 1:nrow(d)){
    temp <- sample(1:50, random_num[i], replace=FALSE)
d[i,temp] <- 1
}
result <- emalgth1(d)
head(result)
```

EMtest

Inferring the methylation expression level of single sites.

Description

Using statistical framework and EM algorithm to infer the methylation expression level of single sites.

Usage

```r
EMtest(datafile = NULL, chrstring = NULL, cpgfile, mrecpgfile = NULL,
writefile = NULL, reportfile = NULL, mreratio = 3/7, psd = 2,
mkadded = 1, f = 1)
```

Arguments

datafile The files of sample. (datafile should be cbind(data1, data2, data3, data4), where data1 and data2 are Medip-seq data, data3 and data4 are MRE-seq data).
chrstring The chromosome should be test.
cpgfile The file of all CpG number.
mrecpgfile The file of MRE-CpG number (If NULL, mrecpgfile will equal to cpgfile).
writefile The path of file of output result. (If writefile=NULL, there will return the results back to main program)
reportfile The path of file of output results of the number of bin, total reads before processing and total reads after processing.
mreratio The ratio of total unmethylation level with total methylation level (Defaulted mreratio is 3/7).
psd The parameters of pseudo count, which pseudo count added to Medip-seq and MRE-seq count.
mkadded Added to all CpG and MRE CpG (We set psd=2 and mkadded=1 as defaulted for robust).
f Adjustment weight, default to 1.
Value
values or file The output file "writefile" will own eleven columns, that is, "chr", "chrSt", "chrEnd", "Medip1", "Medip2", "MRE1", "MRE2", "cg", "mreCG", "pvalue" and "Ts". We also output a report file which will include parameters "s1/s2", "s3/s4", "N1", "N2", "N3", "N4", "c1", "c2", "Number of windows" and "Spend time".

Examples
data(example_data)
data1 <- EM2_H1ESB1_MeDIP_sigleCpG
data2 <- EM2_H1ESB2_MeDIP_sigleCpG
data3 <- H1ESB1_MRE_sigleCpG
data4 <- H1ESB2_MRE_sigleCpG
datafile <- cbind(data1, data2, data3, data4)
allcpg <- all_CpGsite_bin_chr18
mrecpg <- three_mre_cpg
dirwrite <- paste(setwd(getwd()), "/", sep="")
writefile <- paste(dirwrite, "pval_EM_H1ESB1_H1ESB21.bed", sep="")
reportfile <- paste(dirwrite, "report_pvalH1ESB1_H1ESB21.bed", sep="")
EMtest(datafile=datafile, chrstring=NULL, cpgfile=allcpg,
mrecpgfile=mrecpg, writefile=writefile, reportfile=reportfile,
mreratio=3/7, psd=2, mkadded=1, f=1)

H1ESB1_MRE_sigleCpG  A simulation dataset of MRE CpG sites.

Description
This data set gives 2000 CpG sites which include the chromosome of the region, the start and the stop position of the MRE CpG sites.

Usage
H1ESB1_MRE_sigleCpG

Format
A data.frame containing 2000 MRE CpG sites.

Source

References
H1ESB2_MRE_sigeCpG  *A simulation dataset of MRE CpG sites.*

**Description**

This data set gives 2000 CpG sites which include the chromosome of the region, the start and the stop position of the MRE CpG sites.

**Usage**

H1ESB2_MRE_sigeCpG

**Format**

A data.frame containing 2000 MRE CpG sites.

**Source**


**References**


---

**probBinom**

*Compute P-values for Medip-seq and MRE-seq data.*

**Description**

Compute P-values.

**Usage**

probBinom(t, size1, size2, c1, c2)

**Arguments**

- `t`: The real value for random variable according to dataset.
- `size1`: The sum of Medip-seq real reads of the each CpG site for control and treatment sample.
- `size2`: The sum of MRE-seq real reads of the each CpG site for control and treatment sample.
- `c1`: The scaling factor for MeDip-seq data.
- `c2`: The scaling factor for MRE-seq data.
three_mre_cpg

Value

p The P-values for testing the methylation expression levels for each CpG sites.

Examples

```r
set.seed(1234)
t <- 0.1
size1 <- sample(1:1000, 1, replace=TRUE)
size2 <- sample(1:1000, 1, replace=TRUE)
c1 <- 1
c2 <- 2
result <- probBinom(t, size1, size2, c1, c2)
```

three_mre_cpg  A simulation dataset of MRE CpG sites.

Description

This data set gives 2000 CpG sites which include the chromosome of the region, the start and the stop position of the MRE CpG sites.

Usage

```r
three_mre_cpg
```

Format

A data.frame containing 2000 MRE CpG sites.

Source


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