Package ‘MEDME’

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Type Package

Title Modelling Experimental Data from MeDIP Enrichment

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Description Description: MEDME allows the prediction of absolute and relative methylation levels based on measures obtained by MeDIP-microarray experiments

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Depends R (>= 2.15), grDevices, graphics, methods, stats, utils

Imports Biostrings, MASS, drc

Suggests BSgenome.Hsapiens.UCSC.hg18, BSgenome.Mmusculus.UCSC.mm9

License GPL (>= 2)

LazyLoad yes

biocViews Microarray, CpGIsland, DNAMethylation

NeedsCompilation yes

R topics documented:

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Determining the count of CpG dinucleotides for a set of genomic locations

Description

The count of CpGs is determined in each window of size wsize, with or without weighting, for each probe according to its position, chromosome and genome release.

Usage

CGcount(data, wsize = 1000, wFunction = "linear")

Arguments

data : An object of class MEDMEset
wsize : number; the size of the smoothing window, in bp
wFunction : string; the type of weighting function, to choose among linear, exp, log or none

Details

Only human and mouse are currently supported. The respective genomic sequence metadata library needs to be downloaded from the Bioconductor website, installed and loaded (around 800Mb). Please note that only the last genome release should be used. LiftOver UCSC tool could be used for batch conversion of old genomic position to the last genome release.

Value

An object of class MEDMEset is returned where the count of CpGs for each probe has been saved on the CGcount slot.

See Also

smooth

Examples

data(testMEDMEset)
## just an example with the first 1000 probes
testMEDMEset = smooth(data = testMEDMEset[1:1000,])
library(BSgenome.Hsapiens.UCSC.hg18)
testMEDMEset = CGcount(data = testMEDMEset)
MEDME

Determining the logistic model of MeDIP enrichment in respect to the expected DNA methylation level

Description

Probe-level MeDIP weighted enrichment is compared to the expected DNA methylation level. The former is determined applying MeDIP protocol to a fully methylated DNA. The latter is determined as the count of CpGs for each probe. This is assumed to be the methylation level of each probe in a fully methylated sample.

Usage

MEDME(data, sample, CGcountThr = 1, figName = NULL)

Arguments

data: An object of class MEDMEset
sample: Integer; the number of the sample to be used to fit the model, based on the order of samples in the smoothed slot
CGcountThr: number; the threshold to avoid modelling probes with really low methylation level, i.e. CpG count
figName: string; the name of the file reporting the model fitting

Details

The model should be applied on calibration data containing MeDIP enrichment of fully methylated DNA, most likely artificially generated (see references). Nevertheless, in case chromosome or genome-wide human tiling arrays are used a regular sample could be used too. In fact, human genomic DNA is known to be hyper-methylated but in the promoter regions. Of course the performance of the method is expected to be somehow affected by this approximation.

Value

The logistic model as returned from the multdrc function from the drc R library

References

http://genome.cshlp.org/cgi/content/abstract/gr.080721.108v1

See Also

smooth, CGcount

Examples

data(testMEDMEset)
## just an example with the first 1000 probes
testMEDMEset = smooth(data = testMEDMEset[1:1000, ])
library(BSgenome.Hsapiens.UCSC.hg18)
testMEDMEset = CGcount(data = testMEDMEset)
MEDMEmodel = MEDME(data = testMEDMEset, sample = 1, CGcountThr = 1, figName = NULL)
MEDME.predict

Applying the logistic model on MeDIP enrichment data

Description
This allows the probe-level determination of MeDIP smoothed data, as well as absolute and relative methylation levels (AMS and RMS respectively)

Usage
MEDME.predict(data, MEDMEfit, MEDMEextremes = c(1,32), wsize = 1000, wFunction='linear')

Arguments
data
MEDMEfit
MEDMEextremes
wsize
wFunction

Value
An object of class MEDMEset. The resulting smoothed data, the absolute and relative methylation score (AMS and RMS) are saved in the smoothed, AMS and RMS slots, respectively.

References
http://genome.cshlp.org/cgi/content/abstract/gr.080721.108v1

See Also
smooth, CGcount, MEDME

Examples
data(testMEDMEset)
# just an example with the first 1000 probes
testMEDMEset = smooth(data = testMEDMEset[1:1000, ])
library(BSgenome.Hsapiens.UCSC.hg18)
testMEDMEset = CGcount(data = testMEDMEset)
MEDMEmodel = MEDME(data = testMEDMEset, sample = 1, CGcountThr = 1, figName = NULL)
testMEDMEset = MEDME.predict(data = testMEDMEset, MEDMEfit = MEDMEmodel, MEDMEextremes = c(1,32), wsize = 100)
**MEDME.readFiles**

*reading sgr or gff files for MEDME*

**Description**

allows to read sgr or gff files before submitting the data to MEDME analysis

**Usage**

MEDME.readFiles(path = getwd(), files = NULL, format, organism)

**Arguments**

- **path**: string; the path where the files are stored; the current working directory is the default
- **files**: vector; optional vector of file names
- **format**: string; either sgr or gff to indicate the respective file formats
- **organism**: string; either hsa or mmu for homo sapiens and mus musculus respectively

**Details**

In case of GFF files (recommendend), tab-delimited files with header are expected with following fields: chromosome, probe ids, start and stop chromosomal positions, and score are expected in columns 1, 3, 4, 5 and 6 respectively. Multiple files are also expected to be in the same order of rows.

In case of sgr files (GFF is the preferred format), tab-delimited files with no header and chr, chr positions and score are expected in columns 1, 2 and 3 respectively. Multiple files are also expected to be in the same order of rows.

**Value**

An object of class MEDMEmset. The column headers in the logR slot are determined from the file names. in case of SGR files the are not probe names and progressive numbers are used in place of them. In case of GFF files the probe names are determined from the 3rd column.

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**MEDME.writeFiles**

*writeFiles sgr or gff files from MEDME output*

**Description**

allows to write sgr or gff files after MEDME analysis

**Usage**

MEDME.writeFiles(data, output, path = getwd(), format, featureLength = NULL)
MEDMEset-class

Arguments

data: An object of class MEDMEset
output: string; the name of the data slot to be written on the disk, either logR, smoothed, AMS or RMS
path: string; the path where the files are stored; the current working directory is the default
format: string; either sgr or gff to indicate the respective file formats
featureLength: integer; in case of GFF file format the length of the features has to be provided to determine start and end positions

Details

One GFF or SGR file is provided for each sample of the data MEDMEset object.
In case of GFF files, tab-delimited files with header are provided with following fields for each probe: chromosome, empty field, probe ids, start and stop chromosomal positions, and score and empty fields.
In case of sgr files, tab-delimited files with no header and chr, chr positions and score are provided.

MEDMEset-class

Description

This class is used in MEDME library to store MeDIP derived DNA-methylation estimates and to save further elaboration of these, in association with chromosomal and positional probe information

Objects from the Class

Objects can be created by calls of the form new("MEDMEset", ...). This object could initially host the MeDIP normalized logRatio data, as returned by the MEDME.readFiles function. Afterwards, the same object is returned by most of the MEDME library function. Each time, a new slot is filled with additional data, as smoothed logR or Absolute/Relative Methylation Scores (AMS and RMS respectively). At the end of the analysis, usually after a call to the MEDME.predict function, the MEDME.writeFiles function can be used to generate SGR or GFF files from this object.

Slots

chr: Object of class "character": the probe-level chromosome asignments
pos: Object of class "numeric": the probe-level genomic position
logR: Object of class "matrix": the probe-level un-trasformed normalized MeDIP logRatios for each sample
smoothed: Object of class "matrix": the probe-level smoothed MeDIP logRatios for each sample
AMS: Object of class "matrix": the probe-level Absolute Methylation Score for each sample
RMS: Object of class "matrix": the probe-level Relative Methylation Score for each sample
CGcounts: Object of class "numeric": the probe-level count of CpGs
organism: Object of class "character": the organism that the probe genomic positions are referring to, either hsa or mmu for homo sapiens or mus musculus respectively
Methods

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<th>Signature</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>$\text{AMS}$ signature(object = &quot;MEDMEset&quot;)</td>
<td>extracts the Absolute Methylation Score from the AMS slot</td>
</tr>
<tr>
<td>$\text{CG}$ signature(object = &quot;MEDMEset&quot;)</td>
<td>extracts the probe CpG count from the CGcounts slot</td>
</tr>
<tr>
<td>$\text{chr}$ signature(object = &quot;MEDMEset&quot;)</td>
<td>extracts the probe chromosomal assignment</td>
</tr>
<tr>
<td>$\text{org}$ signature(object = &quot;MEDMEset&quot;)</td>
<td>extracts the organism</td>
</tr>
<tr>
<td>$\text{initialize}$ signature(.Object = &quot;MEDMEset&quot;)</td>
<td>automatically generates smoothed, AMS and RMS matrix when only the logR slot is filled</td>
</tr>
<tr>
<td>$\text{logR}$ signature(object = &quot;MEDMEset&quot;)</td>
<td>extracts the matrix of MeDIP un-transformed logRatios</td>
</tr>
<tr>
<td>$\text{pos}$ signature(object = &quot;MEDMEset&quot;)</td>
<td>extracts the probe genomic position</td>
</tr>
<tr>
<td>$\text{RMS}$ signature(object = &quot;MEDMEset&quot;)</td>
<td>extracts the Relative Methylation Score from the RMS slot</td>
</tr>
<tr>
<td>$\text{show}$ signature(object = &quot;MEDMEset&quot;)</td>
<td>prints a summary of the object content</td>
</tr>
<tr>
<td>$\text{smoothed}$ signature(object = &quot;MEDMEset&quot;)</td>
<td>extracts the Absolute Methylation Score from the AMS slot</td>
</tr>
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Author(s)

Mattia Pelizzola

References

http://genome.cshlp.org/cgi/content/abstract/gr.080721.108v1

See Also

MEDME.readFiles, MEDME.writeFiles

Examples

showClass("MEDMEset")

smooth Determining weighted MeDIP data

Description

MeDIP data from tiling arrays are smoothed by determining for each probe $i$ the weighted average of the probes within a window of size $w_{size}$ centered at $i$

Usage

smooth(data, wsize=1000, wFunction='linear')
Arguments

data An object of class MEDMEset
wsize number; the size of the smoothing window, in bp
wFunction string; the type of weighting function, to choose among linear, exp, log or none

Details

The un-smoothed data are read from the slot logR of the data MEDMEset and the resulting smoothed data are saved on the smoothed slot.

Value

An object of class MEDMEset. In particular, the smoothed data are saved on the smoothed slot.

Examples

data(testMEDMEset)
# just an example with the first 1000 probes
testMEDMEset = smooth(data = testMEDMEset[1:1000,])

Description

This dataset contains a subset of the data reported in references. It contains normalized un-smoothed probe-level MeDIP enrichment for almost 50000 probes. This is a random subset of a custom Nimblegen chromosome X tiling array. It is a two channels array with an resolution of 100bp and oligos of 60nt. The M value is reported only. The fullyMet column of the logR slot contains data from a calibration experiments where MeDIP has been applied to a fully methylated sample. The last two columns NBMEL and YUSAC2 contain DNA-methylation experimental data for two cell strains: NBMEL are newborn normal melanocytes cells and YUSAC2 a melanoma strain. Data was processed with within and between array normalization. The full dataset contains almost 380K probes. See references for details. Chromosome, genomic position and logR of probes can be accessed with the methods chr, pos and logR respectively.

Please note that the original genomic coordinates were mapped to the hg17 human genome. These have been converted to hg18 using the LiftOver USCS tool available online for batch conversion.

Usage

data(testMEDMEset)

Format

MEDMEset

References

http://genome.cshlp.org/cgi/content/abstract/gr.080721.108v1
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