

Package ‘msgbsR’

October 18, 2022

Type Package

Title msgbsR: methylation sensitive genotyping by sequencing (MS-GBS)
R functions

Version 1.20.0

Date 2021-11-21

Author Benjamin Mayne

Maintainer Benjamin Mayne <benjamin.mayne@adelaide.edu.au>

Depends R (>= 3.4), GenomicRanges, methods

Imports BSgenome, easyRNASeq, edgeR, GenomicAlignments,
GenomicFeatures, GenomeInfoDb, ggbio, ggplot2, IRanges,
parallel, plyr, Rsamtools, R.utils, stats,
SummarizedExperiment, S4Vectors, utils

Suggests roxygen2, BSgenome.Rnorvegicus.UCSC.mn6

biocViews ImmunoOncology, DifferentialMethylation, DataImport,
Epigenetics, MethylSeq

Description Pipeline for the analysis of a MS-GBS experiment.

License GPL-2

LazyLoad yes

Collate 'msgbsR.R' 'rawCounts.R' 'checkCuts.R' 'plotCounts.R'
'diffMeth.R' 'plotCircos.R'

RoxygenNote 5.0.1

git_url <https://git.bioconductor.org/packages/msgbsR>

git_branch RELEASE_3_15

git_last_commit acb23f0

git_last_commit_date 2022-04-26

Date/Publication 2022-10-18

R topics documented:

checkCuts	2
cuts	3
diffMeth	4
msgbsR	4
plotCircos	5
plotCounts	6
ratdata	6
ratdata2	7
rawCounts	8
Index	9

checkCuts	<i>checkCuts</i>
-----------	------------------

Description

Determines the sequence around a cut site using a fasta file or BSgenome

Usage

```
checkCuts(cutSites, genome, fasta = FALSE, seq)
```

Arguments

cutSites	A GRanges object containing the locations of the cut sites to be checked for sequence match. The names of the correct cut sites will be returned as a GRanges object.
genome	The path to a fasta file or a BSgenome object to check for genomic sequences.
fasta	TRUE if a fasta file has been supplied. Default = FALSE
seq	The desired recognition sequence that the enzyme should have cut.

Value

A GRanges object containing the names of the sites that had the correct sequence.

Author(s)

Benjamin Mayne

Examples

```
library(GenomicRanges)
library(SummarizedExperiment)
library(BSgenome.Rnorvegicus.UCSC.rn6)
# Load the positions of possible MspI cut sites
data(ratdata)
# Extract the cut sites
cutSites <- rowRanges(ratdata)
# Adjust the cut sites to overlap recognition site on each strand
start(cutSites) <- ifelse(test = strand(cutSites) == '+',
                          yes = start(cutSites) - 1, no = start(cutSites) - 2)
end(cutSites) <- ifelse(test = strand(cutSites) == '+',
                        yes = end(cutSites) + 2, no = end(cutSites) + 1)
correctCuts <- checkCuts(cutSites = cutSites, genome = "rn6", seq = "CCGG")
```

cuts	<i>A GRanges object of differentially methylated MspI cut sites on chromosome 20 in Rat from a MS-GBS experiment.</i>
------	---

Description

The GRanges object was created from a list of differentially methylated cut sites from a MS-GBS experiment between two groups of rats that were fed either a control diet or a high fat diet.

Usage

```
data(cuts)
```

Format

A GRanges object of length 10.

Details

- Positions of MspI cut sites differentially methylated in the prostate on chromosome 20 in Rats.

The data set contains 10 differentially methylated sites in the prostate between rats fed a control or high fat diet.

Value

A GRanges object of length 10.

diffMeth	<i>diffMeth</i>
----------	-----------------

Description

Determines differential methylated sites from a RangedSummarizedExperiment

Usage

```
diffMeth(se, category, condition1, condition2,
         block = NULL, cpmThreshold, thresholdSamples)
```

Arguments

se	A RangedSummarizedExperiment containing meta data of the samples.
category	The heading name in the sample data to be tested for differential methylation.
condition1	The reference group within the category.
condition2	The experimental group within the category.
block	The heading name in the sample data if differential methylation is to be tested with a blocking factor. Default is NULL.
cpmThreshold	Counts per million threshold of read counts to be filtered out of the analysis.
thresholdSamples	Minimum number of samples to contain the counts per million threshold.

Value

A data frame containing which cut sites that are differentially methylated.

Author(s)

Benjamin Mayne

Examples

```
# Load data
data(ratdata2)
top <- diffMeth(se = ratdata2, category = "Group",
               condition1 = "Control", condition2 = "Experimental",
               cpmThreshold = 1, thresholdSamples = 1)
```

msgbsR	<i>msgbsR</i>
--------	---------------

Description

msgbsR

plotCircos	<i>plotCircos</i>
------------	-------------------

Description

Plot a circos representing the cut site locations

Usage

```
plotCircos(cutSites, seqlengths, cutSite.colour, seqlengths.colour)
```

Arguments

cutSites A GRanges object containing the locations of the cut sites to be plotted.
seqlengths An integer with the lengths of the chromosomes.
cutSite.colour The colour of the cut sites.
seqlengths.colour The colour of the chromosomes

Value

A circos plot showing the locations of the cut sites.

Author(s)

Benjamin Mayne

Examples

```
# load example cut site positions
data(cuts)
# Obtain the length of chromosome 20 in rn6
library(BSgenome.Rnorvegicus.UCSC.rn6)
chr20 <- seqlengths(BSgenome.Rnorvegicus.UCSC.rn6)["chr20"]
plotCircos(cutSites = cuts, seqlengths = chr20,
           cutSite.colour = "red", seqlengths.colour = "blue")
```

plotCounts	<i>plotCounts</i>
------------	-------------------

Description

Plots the total number of reads vs total number of cut sites per sample

Usage

```
plotCounts(se, category)
```

Arguments

se	A RangedSummarizedExperiment containing meta data of the samples.
category	The heading name in the sample data to distinguish groups.

Value

Produces a plot showing the total number reads vs total number of cut sites per sample.

Author(s)

Benjamin Mayne

Examples

```
data(ratdata2)
plotCounts(se = ratdata2, category = "Group")
```

ratdata	<i>Read counts of potential MspI cut sites from a MS-GBS experiment of prostates from rats</i>
---------	--

Description

A RangedSummarizedExperiment containing read counts generated from a MS-GBS experiment using the restriction enzyme MspI, focusing on chromosome 20 of Rat.

Usage

```
data(ratdata)
```

Format

RangedSummarizedExperiment

Details

- ratdata A RangedSummarizedExperiment with 16047 potential MspI cut sites on chromosome 20 in Rat and six samples (3 Control and 3 Experimental).

This dataset contains six prostate samples from rats: 3 control and 3 experimental high fat diet.

Value

RangedSummarizedExperiment

ratdata2	<i>Read counts of correct MspI cut sites from a MS-GBS experiment of prostates from rats</i>
----------	--

Description

A RangedSummarizedExperiment containing read counts generated from a MS-GBS experiment using the restriction enzyme MspI, focusing on chromosome 20 of Rat. The sites have been checked for the correct recognition site.

Usage

```
data(ratdata2)
```

Format

RangedSummarizedExperiment

Details

- ratdata2 A RangedSummarizedExperiment containing data for 13983 MspI cut sites on chromosome 20 in Rat and six samples (3 Control and 3 Experimental).

This dataset contains six prostate samples from rats: 3 control and 3 experimental high fat diet. The data can be used for differential methylation analyses.

Value

RangedSummarizedExperiment

`rawCounts`*rawCounts*

Description

Imports the raw read counts from sorted and indexed bam file(s)

Usage

```
rawCounts(bamFilepath, threads = 1)
```

Arguments

`bamFilepath` The path to the location of the bam file(s).
`threads` The total number of usable threads to be used. Default is 1.

Value

Produces a `RangedSummarizedExperiment`. Columns are samples and the rows are cut sites. The cut site IDs are in the format `chr:position-position:strand`.

Author(s)

Benjamin Mayne, Sam Buckberry

Examples

```
my_path <- system.file("extdata", package = "msgbsR")  
my_data <- rawCounts(bamFilepath = my_path)
```


Index

* datasets

cuts, [3](#)

ratdata, [6](#)

ratdata2, [7](#)

checkCuts, [2](#)

cuts, [3](#)

diffMeth, [4](#)

msgbsR, [4](#)

msgbsR-package (msgbsR), [4](#)

plotCircos, [5](#)

plotCounts, [6](#)

ratdata, [6](#)

ratdata2, [7](#)

rawCounts, [8](#)