

# Package ‘methInheritSim’

June 26, 2025

**Type** Package

**Title** Simulating Whole-Genome Inherited Bisulphite Sequencing Data

**Description** Simulate a multigeneration methylation case versus control experiment with inheritance relation using a real control dataset.

**Version** 1.30.0

**Date** 2021-11-21

**Author** Pascal Belleau, Astrid Deschênes and Arnaud Droit

**Author@R** `c(person("`Pascal", "`Belleau",  
email="`pascal_belleau@hotmail.com", role=c("`cre", "`aut")),  
person("`Astrid", "`Deschênes", email="`adeschen@hotmail.com",  
role=c("`aut")), person("`Arnaud", "`Droit",  
email="`arnaud.droit@crchuq.ulaval.ca", role=c("`aut")))`

**Depends** R (>= 3.4)

**Imports** methylKit, GenomicRanges, GenomeInfoDb, parallel,  
BiocGenerics, S4Vectors, methods, stats, IRanges, msm

**Suggests** BiocStyle, knitr, rmarkdown, RUnit, methylInheritance

**Encoding** UTF-8

**License** Artistic-2.0

**URL** <https://github.com/belleau/methInheritSim>

**BugReports** <https://github.com/belleau/methInheritSim/issues>

**VignetteBuilder** knitr

**biocViews** BiologicalQuestion, Epigenetics, DNAMethylation,  
DifferentialMethylation, MethylSeq, Software, ImmunoOncology,  
StatisticalMethod, WholeGenome, Sequencing

**Maintainer** Pascal Belleau <pascal\_belleau@hotmail.com>

**RoxygenNote** 6.0.1

**git\_url** <https://git.bioconductor.org/packages/methInheritSim>

**git\_branch** RELEASE\_3\_21

**git\_last\_commit** 8055b10  
**git\_last\_commit\_date** 2025-04-15  
**Repository** Bioconductor 3.21  
**Date/Publication** 2025-06-25

**Contents**

methInheritSim-package . . . . .	2
calculateNbDiffCase . . . . .	3
createSampleID . . . . .	4
dataSimExample . . . . .	4
estBetaAlpha . . . . .	6
estBetaBeta . . . . .	7
fixSeed . . . . .	7
getDiffCase . . . . .	8
getDiffMeth . . . . .	9
getSim . . . . .	11
getSyntheticChr . . . . .	12
runOnEachSynCHR . . . . .	13
runSim . . . . .	17
samplesForChrSynthetic . . . . .	20
saveData . . . . .	21
simEachGeneration . . . . .	22
simInheritance . . . . .	24
testIfAlreadyDone . . . . .	28
validateRunSimDoubleParameters . . . . .	29
validateRunSimIntegerParameters . . . . .	30
validateRunSimLogicalParameters . . . . .	31
validateRunSimOtherParameters . . . . .	32
validateRunSimParameters . . . . .	34
<b>Index</b>	<b>37</b>

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methInheritSim-package	<i>methInheritSim: Simulating Whole-Genome Inherited Bisulphite Sequencing Data</i>
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**Description**

This package generates simulations of multigeneration of bisulfite data using a real control dataset.

**Author(s)**

Pascal Belleau, Astrid Deschênes and Arnaud Droit  
Maintainer: Pascal Belleau <pascal\_belleau@hotmail.com>

**See Also**

- [runSim](#) for simulating a multigeneration methylation experiment with inheritance

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calculateNbDiffCase	<i>Calculate the number of differentially methylated cases.</i>
---------------------	-----------------------------------------------------------------

---

**Description**

Identify the number of differentially methylated cases.

**Usage**

```
calculateNbDiffCase(nbCase, propDiff, propDiffSd)
```

**Arguments**

nbCase	a positive integer, the number of cases.
propDiff	a double superior to 0 and inferior or equal to 1, the mean value for the proportion of samples that will have, for a specific position, differentially methylated values. It can be interpreted as the penetrance.
propDiffSd	a non-negative double, the standard deviation associated to the propDiff

**Value**

a integer, the number of differentially methylated cases.

**Author(s)**

Pascal Belleau, Astrid Deschenes

**Examples**

```
## Fix seed to have reproducible results
set.seed(3122)

## Obtained the number of differential cases
methInheritSim::calculateNbDiffCase(nbCase = 8,
  propDiff = 0.8, propDiffSd = 0.2)
```

---

createSampleID	<i>Generate the samples ID for the simulated dataset.</i>
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---

### Description

Generate the samples ID for the simulated dataset. The standard format of the samples ID is :  
 "F[Number for the generation]\_[Number for the sample] \_[OC for case or C for control]"

### Usage

```
createSampleID(nbGeneration, nbSample)
```

### Arguments

nbGeneration	a positive integer, the number of generations simulated.
nbSample	a positive integer, the number of controls (CTRL) and cases in the simulated dataset.

### Value

a list containing a list of sample ID for each generation.

### Author(s)

Pascal Belleau, Astrid Deschenes

### Examples

```
## Create sample ID
methInheritSim::createSampleID(nbGeneration = 3, nbSample = 6)
```

---

dataSimExample	<i>A list containing methylation information used by some internal functions (for demo purpose).</i>
----------------	------------------------------------------------------------------------------------------------------

---

### Description

A list containing methylation information used by some internal functions (for demo purpose).

### Usage

```
data(dataSimExample)
```

**Format**

a list containing:

- stateInfo a GRanges, a synthetic chromosome as generated by getSyntheticChr function.
- stateDiff a list containing:
  - stateDiff a vector of integer (0 and 1) with length corresponding the length of stateInfo. The vector indicates, using 1, the positions where the CpG sites are differentially methylated.
  - stateInherit a vector of integer (0 and 1) with length corresponding the length of stateInfo. The vector indicates, using 1, the positions where the CpG values are inherited.
- treatment a vector of integer (0 and 1) with length corresponding the number of samples. The vector indicates which samples are control (0) which samples are case (1).
- sample.id a list of 3 list. Each entry of the list correspond to one generation (first entry = first generation, etc..). Each list contains a list of 12 entries each containing a string of character, the name of the sample.

**Value**

a list containing:

- stateInfo a GRanges, a synthetic chromosome as generated by getSyntheticChr function.
- stateDiff a list containing:
  - stateDiff a vector of integer (0 and 1) with length corresponding the length of stateInfo. The vector indicates, using 1, the positions where the CpG sites are differentially methylated.
  - stateInherit a vector of integer (0 and 1) with length corresponding the length of stateInfo. The vector indicates, using 1, the positions where the CpG values are inherited.
- treatment a vector of integer (0 and 1) with length corresponding the number of samples. The vector indicates which samples are control (0) which samples are case (1).
- sample.id a list of 3 list. Each entry of the list correspond to one generation (first entry = first generation, etc..). Each list contains a list of 12 entries each containing a string of character, the name of the sample.

**See Also**

- [runSim](#) for running a simulation analysis using methylKit info as input

**Examples**

```
## Loading dataset
data(dataSimExample)

## Identify differentially methylated sites and among those, the ones
## that are inherited
methInheritSim::getDiffMeth(stateInfo = dataSimExample$stateInfo,
```

```
rateDiff = 0.2, minRate = 0.3,propInherite = 0.3)
```

---

**estBetaAlpha***Estimate the alpha parameter of a Beta distribution*

---

### Description

Estimate the alpha parameter from the mean and the variance of a Beta distribution.

### Usage

```
estBetaAlpha(meanCtrl, varCtrl, minVal = 1e-06)
```

### Arguments

meanCtrl	a double, the mean of the controls (CTRL) at a specific CpG site.
varCtrl	a double, the variance of the controls (CTRL) at a specific CpG site.
minVal	a double, the minimum value accepted for the mean value. If meanCtrl is smaller than minVal, then minVal is used in the calculation of the alpha parameter. Default: 1e-06.

### Value

a double, the alpha parameter of a Beta distribution.

### Author(s)

Pascal Belleau, Astrid Deschenes

### Examples

```
## Estimate alpha parameters with mean = 0.5 and variance = 0.1  
methInheritSim::estBetaAlpha(meanCtrl = 0.5, varCtrl = 0.1)
```

---

estBetaBeta	<i>Estimate the beta parameter of a beta distribution</i>
-------------	-----------------------------------------------------------

---

**Description**

Estimate the beta parameter from the mean and the variance of a beta distribution.

**Usage**

```
estBetaBeta(meanCtrl, varCtrl, minVal = 1e-06)
```

**Arguments**

meanCtrl	a double, the mean of the controls (CTRL) at a specific CpG site.
varCtrl	a double, the variance of the controls (CTRL) at a specific CpG site.
minVal	a double, the minimum value accepted for the mean value. If meanCtrl is smaller than minVal, then minVal is used in the calculation of the beta paramter. Default: 1e-06.

**Value**

a double, the beta parameter of a Beta distribution.

**Author(s)**

Pascal Belleau, Astrid Deschenes

**Examples**

```
## Estimate beta parameters with mean = 0.5, variance = 0.1  
methInheritSim::estBetaBeta(meanCtrl=0.5, varCtrl=0.1)
```

---

fixSeed	<i>Fix seed value.</i>
---------	------------------------

---

**Description**

Fix seed value when specified value is -1.

**Usage**

```
fixSeed(vSeed)
```

**Arguments**

vSeed                    a integer, a seed used when reproducible results are needed. When a value inferior or equal to zero is given, a random integer is used.

**Value**

a double, the seed value.

**Author(s)**

Pascal Belleau, Astrid Deschenes

**Examples**

```
## Return vSeed value when value is not -1
methInheritSim::fixSeed(vSeed = 10)

## Return new value when value is -1
methInheritSim::fixSeed(vSeed = -1)
```

---

getDiffCase	<i>Get the C/T proportion at a selected site, differentially methylated or not, for all cases</i>
-------------	---------------------------------------------------------------------------------------------------

---

**Description**

Simulate the proportion of C/T for each case at a selected site, differentially methylated or not.

**Usage**

```
getDiffCase(ctrlMean, ctrlVar, selectedAsDM, nbCase, sDiff, nbDiffCase)
```

**Arguments**

ctrlMean                a double, the mean of the CTRL at the site.

ctrlVar                 a double, the variance of the CTRL at the site.

selectedAsDM           a integer, 1 if the site is selected as differentially methylated, otherwise 0.

nbCase                  a integer, the number of cases.

sDiff                   a non-negative double included in [0,1], the proportion of C/T for a case differentially methylated that follows a beta distribution where the mean is shifted of vDiff from the CTRL distribution.

nbDiffCase             an integer, the number of cases differentially methylated.



**Value**

a vector containing  $3 + \text{nbCase}$  entries of type double:

- The mean proportion of C/T of the differentially methylated cases
- The number of cases simulated using shifted distribution
- The number of cases simulated using the control distribution
- The proportion of C/T for each case

**Author(s)**

Pascal Belleau, Astrid Deschenes

**Examples**

```
## Fix seed to obtain replicable results
set.seed(2010)

## Get the proportion of C/T for each case at a specific site.
methInheritSim::getDiffCase(ctrlMean = 0.9814562, ctrlVar =
0.0003607153, selectedAsDM = 0, nbCase=6, sDiff = 0.8,
nbDiffCase = round(6 * 0.9))
```

---

getDiffMeth	<i>Identify differentially methylated sites and among those, the ones that are inherited.</i>
-------------	-----------------------------------------------------------------------------------------------

---

**Description**

Identify the site positions where the cases are differentially methylated and, among those, the one that are inherited.

**Usage**

```
getDiffMeth(stateInfo, rateDiff, minRate, propInherite, c = 1, b = -0.1,
  endLength = 1000)
```

**Arguments**

stateInfo	a GRanges that contains the CpG (or methylated sites). The GRanges have four metadata from the real dataset: <ul style="list-style-type: none"> <li>• chrOri, the chromosome from the real dataset</li> <li>• startOri, the position of the site in the real dataset</li> <li>• meanCTRL, the mean of the control in the real dataset</li> <li>• varCTRL, the variance of the control in the real dataset</li> </ul>
-----------	----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------

rateDiff	a positive double inferior to 1, the mean of the chance that a site is differentially methylated.
minRate	a non-negative double inferior to 1, the minimum rate for differentially methylated sites.
propInherite	a non-negative double inferior or equal to 1, the proportion of differentially methylated regions that are inherited.
c	a positive double, a factor in the formula to compute the probability of site to be differentially methylated in a differentially methylated region. The probability formula of site in differentially methylated region is $c * \exp(b * \log(\text{distance with the preceding sites}))$ Default: 1.0.
b	a negative double, a factor in the formula to compute the probability of site to be differentially methylated in a differentially methylated region. The probability formula of site in differentially methylated region is $c * \exp(b * \log(\text{distance with the preceding sites}))$ . Default: $-1e-01$ .
endLength	a positive integer, when the distance with the preceding sites in a differentially methylated region is larger than endLength, the differentially methylated region is finished. Default: 1000.

### Value

a list containing the 2 following elements:

- stateDiff a vector of integer (0 and 1) with length corresponding the length of stateInfo. The vector indicates, using 1, the positions where the CpG sites are differentially methylated.
- stateInherite a vector of integer (0 and 1) with length corresponding the length of stateInfo. The vector indicates, using 1, the positions where the CpG values are inherited.

### Author(s)

Pascal Belleau, Astrid Deschenes

### Examples

```
## Load dataset containing a list of objects used by
## methInheritSim internal functions
data(dataSimExample)

## Identify differentially methylated sites and among those, the ones
## that are inherited
methInheritSim::getDiffMeth(stateInfo =
  dataSimExample$stateInfo, rateDiff = 0.3, minRate = 0.3,
  propInherite = 0.3)
```

---

getSim	<i>Simulate the proportion of C/T at each site of synthetic CHR for each control and case</i>
--------	-----------------------------------------------------------------------------------------------

---

### Description

For each control and case, generate the proportion of C/T at each of the synthetic CHR.

### Usage

```
getSim(nbCtrl, nbCase, generation, stateInfo, stateDiff, stateInherite,
       diffValue, propDiff, propDiffsd, propInheritance, propHetero)
```

### Arguments

nbCtrl	a positive integer, the number of controls.
nbCase	a positive integer, the number of cases.
generation	a positive integer, the number of generations.
stateInfo	a GRanges object, the synthetic chromosome generated by getSyntheticChr function.
stateDiff	a vector of integer (0 and 1) with length corresponding the length of stateInfo. The vector indicates, using a 1, the positions where the CpG sites are differentially methylated.
stateInherite	a vector of integer (0 and 1) with length corresponding the length of stateInfo. The vector indicates, using a 1, the positions where the CpG values are inherited.
diffValue	a non-negative double between between [0,1], the proportion of C/T for a case differentially methylated following a beta distribution where the mean is shifted of diffValue from the CTRL distribution.
propDiff	a double superior to 0 and inferior or equal to 1, the mean value for the proportion of samples that will have, for a specific position, differentially methylated values. It can be interpreted as the penetrance.
propDiffsd	a non-negative double, the standard deviation associated to the propDiff.
propInheritance	a non-negative double between [0,1], the proportion of case that inherit differentially methylated sites.
propHetero	a non-negative double between [0,1], the reduction of vDiff for the second and following generations.

### Value

a GRangesList, the object contains information about the simulation. The file have four metadata related to the real dataset:

- meanDiff, the means of the shifted distribution.

- meanCTRL, the means of the control distribution.
- partitionCase, the number of cases simulated using the shifted distribution.
- partitionCtrl, the number of cases simulated using the control distribution and a metadata for each cases and controls the proportion of C/T.

### Author(s)

Pascal Belleau, Astrid Deschenes

### Examples

```
## Fix seed to have reproducible results
set.seed(312)

## Load dataset
data("samplesForChrSynthetic")

## Generate a stateInfo object using samples
stateInformation <- methInheritSim::getSyntheticChr(methInfo =
  samplesForChrSynthetic, nbBlock = 1, nbCpG = 3)

## Generate a stateDiff and stateInherit objects with length corresponding
## to nbBlock * nbCpG from stateInformation
stateDiff <- c(1, 0, 1)
stateInherit <- c(1, 0, 0)

## Create a simulation using stateInformation, stateDiff and stateInherit
methInheritSim::getSim(nbCtrl = 3, nbCase = 2, generation = 3,
  stateInfo = stateInformation, stateDiff = stateDiff,
  stateInherit = stateInherit, diffValue = 10,
  propDiff = 0.8, propDiffsd = 0.2, propInheritance = 0.8,
  propHetero = 0.1)
```

---

getSyntheticChr

---

*Create a synthetic chromosome with the CTRL genome*


---

### Description

Create a synthetic chromosome with the sampling of a specified number of blocks and a specified number of consecutive CpG.

### Usage

```
getSyntheticChr(methInfo, nbBlock, nbCpG)
```

**Arguments**

methInfo	is object of class methylBase, the CpG information from controls (CTRL) that will be used to create the sythetic chromosome. The object can also contain information from cases but only the controls will be used.
nbBlock	a positive integer, the number of blocks used for sampling.
nbCpG	a integer, the number of consecutive CpG positions used for sampling from methInfo.

**Value**

a GRanges object, the synthetic chromosome.

**Author(s)**

Pascal Belleau

**Examples**

```
## Load methyl information
data(samplesForChrSynthetic)

## Ensure results are reproducible
set.seed(32)

## Create synthetic chromosome
methInheritSim::getSyntheticChr(methInfo = samplesForChrSynthetic,
                                nbBlock = 10, nbCpG = 20)
```

---

runOnEachSynCHR	<i>Simulate a multigeneration methylation experiment with inheritance on each synthetic chromosome.</i>
-----------------	---------------------------------------------------------------------------------------------------------

---

**Description**

Simulate a multigeneration methylation case versus control experiment with inheritance relation using a real control dataset.

**Usage**

```
runOnEachSynCHR(methData, nbSynCHR, nbSimulation, nbBlock, nbCpG, nbGeneration,
                vNbSample, vpDiff, vpDiffsd, vDiff, vInheritance, rateDiff, minRate,
                propInherite, propHetero, keepDiff, outputDir, fileID, minReads, maxPercReads,
                meanCov, context, assembly, saveGRanges, saveMethylKit, runAnalysis, nbCores,
                vSeed)
```

**Arguments**

methData	an object of class methylBase, the CpG information from controls (CTRL) that will be used to create the synthetic chromosome. The methData object can also contain information from cases but only the controls are used.
nbSynCHR	a positive integer, the number of distinct synthetic chromosomes that will be generated.
nbSimulation	a positive integer, the number of simulations generated for each parameter (vNbSample, vpDiff, vDiff and vInheritance). The total number of simulation is $\text{nbSimulation} * \text{length}(\text{vNbSample}) * \text{length}(\text{vpDiff}) * \text{length}(\text{vInheritance})$
nbBlock	a positive integer, the number of blocks used for sampling.
nbCpG	a positive integer, the number of consecutive CpG positions used for sampling from methInfo.
nbGeneration	a positive integer, the number of generations simulated.
vNbSample	a vector of distinct positive integer, the number of controls (CTRL) and cases in the simulated dataset. In the simulated dataset, the number of CTRL equals the number of cases. The number of CTRL do not need to be equal to the number of Case in the real methData dataset.
vpDiff	a vector of distinct double superior to 0 and inferior or equal to 1, the mean value for the proportion of samples that will have, for a specific position, differentially methylated values. It can be interpreted as the penetrance. Note that vpDiff and vpDiffsd must be the same length.
vpDiffsd	a vector of a non-negative double, the standard deviation associated to the vpDiff. Note that vpDiff and vpDiffsd must be the same length.
vDiff	a vector of distinct non-negative double included in [0,1], the proportion of C/T for a case differentially methylated that follows a beta distribution where the mean is shifted by vDiff from the CTRL distribution.
vInheritance	a vector of distinct non-negative double included in [0,1], the proportion of cases that inherits differentially methylated sites.
rateDiff	a positive double inferior to 1, the mean of the chance that a site is differentially methylated.
minRate	a non-negative double inferior to 1, the minimum rate for differentially methylated sites.
propInherite	a non-negative double inferior or equal to 1, the proportion of differentially methylated regions that are inherited.
propHetero	a non-negative double between [0,1], the reduction of vDiff for the second and following generations.
keepDiff	a logical, when TRUE, the differentially methylated sites will be the same for all simulated datasets. Datasets generated using differents parameter values from vector parameters (vpDiff, vDiff and vInheritance) wil all have the same differentially methylated sites.
outputDir	a string of character or NULL, the path where the files created by the function will be saved. When NULL, the files are saved in a directory called "outputDir" that is located in the current directory.

fileID	<p>a string of character, a identifiant that will be included in each output file name. Each output file name is composed of those elements, separated by "_":</p> <ul style="list-style-type: none"> <li>• a type name, ex: methylGR, methylObj, etc..</li> <li>• a fileID</li> <li>• the chromosome number, a number between 1 and nbSynCHR</li> <li>• the number of samples, a number in the vNbSample vector</li> <li>• the mean proportion of samples that has, for a specific position, differentially methylated values, a number in the vpDiff vector</li> <li>• the proportion of C/T for a case differentially methylated that follows a shifted beta distribution, a number in the vDiff vector</li> <li>• the proportion of cases that inherits differentially sites, a number in the vInheritance vector</li> <li>• the identifiant for the simulation, a number between 1 and nbSimulation</li> <li>• the file extension ".rds"</li> </ul>
minReads	a positive integer, sites and regions having lower coverage than this count are discarded. The parameter corresponds to the lo.count parameter in the methylKit package.
maxPercReads	a double between [0,100], the percentile of read counts that is going to be used as upper cutoff. Sites and regions having higher coverage than maxPercReads are discarded. This parameter is used for both CpG sites and tiles analysis. The parameter correspond to the hi.perc parameter in the methylKit package.
meanCov	a positive integer, the mean of the coverage at the simulated CpG sites.
context	a string of character, the short description of the methylation context, such as "CpG", "CpH", "CHH", etc..
assembly	a string of character, the short description of the genome assembly, such as "mm9", "hg18", etc..
saveGRanges	a logical, when true, the package save two files type. The first generate for each simulation contains a list. The length of the list corresponds to the number of generation. The generation are stored in order (first entry = first generation, second entry = second generation, etc..). All samples related to one generations are contained in a GRangesList. The GRangeaList store a list of GRanges. Each GRanges stores the raw mehylation data of one sample. The second file a numeric vector denoting controls and cases (a file is generates by entry in the vector parameters vNbSample).
saveMethylKit	a logical, when TRUE, for each simulations save a file contains a list. The length of the list corresponds to the number of generation. The generation are stored in order (first entry = first generation, second entry = second generation, etc..). All samples related to one generations are contained in a S4 methylRawList object. The methylRawList object contains two Slots: 1. treatment: A numeric vector denoting controls and cases. 2. .Data: A list of methylRaw objects. Each object stores the raw methylation data of one sample.
runAnalysis	<p>a logical, if TRUE, two files are saved for each simulation:</p> <ul style="list-style-type: none"> <li>• 1. The first file is the methylObj... file formated with the methylkit package in a S4 methylBase object (using the methylKit functions: filterByCoverage, normalizeCoverage and unite).</li> </ul>

- 2. The second file contains a S4 `calculateDiffMeth` object generated using the `methyKit` functions `calculateDiffMeth` on the first file.
- `nbCores` a positive integer, the number of cores used when creating the simulated datasets. Default: 1 and always 1 for Windows.
- `vSeed` a integer, a seed used when reproducible results are needed. When a value inferior or equal to zero is given, a random integer is used. .

**Value**

0 indicating that the function have been successful.

**Author(s)**

Pascal Belleau, Astrid Deschenes

**Examples**

```
## Load dataset containing methyl information
data(samplesForChrSynthetic)

## Set the output directory where files will be created
temp_dir <- "test_runOnEachSynCHR"

## Create directory
if(!dir.exists(temp_dir)) {
  dir.create(temp_dir, showWarnings = TRUE)
}

## Create 2 simulated dataset (nbSimulation = 2)
## over 3 generations (nbGeneration = 3) with
## 6 cases and 6 controls (nNbSample = 6) using only one set
## of parameters (vpDiff = 0.9, vpDiffsd = 0.1, vDiff = 0.8)
methInheritSim::runOnEachSynCHR(methData = samplesForChrSynthetic,
  nbSynCHR = 1, nbSimulation = 2, nbBlock = 10, nbCpG = 20,
  nbGeneration = 3, vNbSample = c(6), vpDiff = c(0.9), vpDiffsd = c(0.1),
  vDiff = c(0.8), vInheritance = c(0.5), propInherite = 0.3,
  rateDiff = 0.3, minRate = 0.2, propHetero = 0.5, keepDiff = FALSE,
  outputDir = temp_dir, fileID = "F1", minReads = 10,
  maxPercReads = 99.9, meanCov = 80, context = "CpG", assembly="Rnor_5.0",
  saveGRanges = FALSE, saveMethyKit = FALSE,
  runAnalysis = FALSE, nbCores = 1, vSeed = 32)

## Delete the output directory and its content
if (dir.exists(temp_dir)) {
  unlink(temp_dir, recursive = TRUE, force = FALSE)
}
```



runSim

*Simulate a multigeneration methylation experiment with inheritance***Description**

Simulate a multigeneration methylation case versus control experiment with inheritance relation using a real control dataset.

The simulation can be parametrized to fit different models. The number of cases and controls, the proportion of the case affected by the treatment (penetrance), the effect of the treatment on the mean of the distribution, the proportion of sites inherited, the proportion of the differentially methylated sites from the precedent generation inherited, etc..

The function simulates a multigeneration dataset like a bisulfite sequencing experiment. The simulation includes the information about control and case for each generation. Simulation dataset are saved in multiple files created in the directory specified by the user.

**Usage**

```
runSim(methData, nbSynCHR = 1, nbSimulation = 10, nbBlock = 100,
      nbCpG = 50, nbGeneration = 3, vNbSample = c(3, 6), vpDiff = c(0.9),
      vpDiffsd = c(0.1), vDiff = c(0.8), vInheritance = c(0.5),
      rateDiff = 0.01, minRate = 0.01, propInherite = 0.3, propHetero = 0.5,
      keepDiff = FALSE, outputDir = NULL, fileID = "s", minReads = 10,
      maxPercReads = 99.9, meanCov = 80, context = "CpG",
      assembly = "Rnor_5.0", saveGRanges = TRUE, saveMethylKit = TRUE,
      runAnalysis = FALSE, nbCores = 1, vSeed = -1)
```

**Arguments**

methData	an object of class methylBase, the CpG information from controls (CTRL) that will be used to create the synthetic chromosome. The methData object can also contain information from cases but only the controls are used.
nbSynCHR	a positive integer, the number of distinct synthetic chromosomes that will be generated. Default: 1.
nbSimulation	a positive integer, the number of simulations generated for each parameter (vNbSample, vpDiff, vDiff and vInheritance). The total number of simulation is nbSimulation * length(vNbSample) * length(vpDiff) * length(vInheritance). Default: 10.
nbBlock	a positive integer, the number of blocks used for sampling. Default: 100.
nbCpG	a positive integer, the number of consecutive CpG positions used for sampling from methInfo. Default: 50.
nbGeneration	a positive integer, the number of generations simulated. Default: 3.
vNbSample	a vector of distinct positive integer, the number of controls (CTRL) and cases in the simulated dataset. In the simulated dataset, the number of CTRL equals the number of cases. The number of CTRL do not need to be equal to the number of Case in the real methData dataset. Default: c(3, 6).

vpDiff	a vector of distinct double superior to 0 and inferior or equal to 1, the mean value for the proportion of samples that will have, for a specific position, differentially methylated values. It can be interpreted as the penetrance. Note that vpDiff and vpDiffsd must be the same length. Default: <code>c(0.9)</code> .
vpDiffsd	a vector of a non-negative double, the standard deviation associated to the vpDiff. Note that vpDiff and vpDiffsd must be the same length. Default: <code>c(0.1)</code> .
vDiff	a vector of distinct non-negative double included in [0,1], the proportion of C/T for a case differentially methylated that follows a beta distribution where the mean is shifted by vDiff from the CTRL distribution. Default: <code>c(0.8)</code> .
vInheritance	a vector of distinct non-negative double included in [0,1], the proportion of cases that inherits differentially methylated sites. Default: <code>c(0.5)</code> .
rateDiff	a positive double inferior to 1, the mean of the chance that a site is differentially methylated. Default: <code>0.01</code> .
minRate	a non-negative double inferior to 1, the minimum rate for differentially methylated sites. Default: <code>0.01</code> .
propInherite	a non-negative double inferior or equal to 1, the proportion of differentially methylated regions that is inherited. Default: <code>0.3</code> .
propHetero	a non-negative double between [0,1], the reduction of vDiff for the second and following generations. Default: <code>0.5</code> .
keepDiff	a logical, when TRUE, the differentially methylated sites will be the same for all simulated datasets. Datasets generated using different parameter values from vector parameters (vpDiff, vDiff and vInheritance) will all have the same differentially methylated sites. Default: FALSE.
outputDir	a string of character or NULL, the path where the files created by the function will be saved. When NULL, the files are saved in a directory called "outputDir" that is located in the current directory. Default: NULL.
fileID	a string of character, a identifier that will be included in each output file name. Each output file name is composed of those elements, separated by "_": <ul style="list-style-type: none"> <li>• a type name, ex: methylGR, methylObj, etc..</li> <li>• a fileID</li> <li>• the chromosome number, a number between 1 and nbSynCHR</li> <li>• the number of samples, a number in the vNbSample vector</li> <li>• the mean proportion of samples that has, for a specific position, differentially methylated values, a number in the vpDiff vector</li> <li>• the proportion of C/T for a case differentially methylated that follows a shifted beta distribution, a number in the vDiff vector</li> <li>• the proportion of cases that inherits differentially sites, a number in the vInheritance vector</li> <li>• the identifier for the simulation, a number between 1 and nbSimulation</li> <li>• the file extension ".rds"</li> </ul>

Default: "s".

minReads	a positive integer, sites and regions having lower coverage than this count are discarded. The parameter corresponds to the <code>lo.count</code> parameter in the <code>methyKit</code> package. Default: 10.
maxPercReads	a double between [0,100], the percentile of read counts that is going to be used as upper cutoff. Sites and regions having higher coverage than <code>maxPercReads</code> are discarded. This parameter is used for both CpG sites and tiles analysis. The parameter correspond to the <code>hi.perc</code> parameter in the <code>methyKit</code> package. Default: 99.9.
meanCov	a positive integer, the mean of the coverage at the simulated CpG sites. Default: 80.
context	a string of character, the short description of the methylation context, such as "CpG", "CpH", "CHH", etc.. Default: "CpG".
assembly	a string of character, the short description of the genome assembly, such as "mm9", "hg18", etc.. Default: "Rnor_5.0".
saveGRanges	a logical, when true, the package save two files type. The first generate for each simulation contains a list. The length of the list corresponds to the number of generation. The generation are stored in order (first entry = first generation, second entry = second generation, etc..). All samples related to one generations are contained in a <code>GRangesList</code> . The <code>GRangesList</code> store a list of <code>GRanges</code> . Each <code>GRanges</code> stores the raw methylation data of one sample. The second file a numeric vector denoting controls and cases (a file is generated by entry in the vector parameters <code>vNbSample</code> ). Default: TRUE.
saveMethyKit	a logical, when TRUE, for each simulations save a file contains a list. The length of the list corresponds to the number of generation. The generation are stored in order (first entry = first generation, second entry = second generation, etc..). All samples related to one generations are contained in a <code>S4 methylRawList</code> object. The <code>methylRawList</code> object contains two Slots: 1. <code>treatment</code> : A numeric vector denoting controls and cases. 2. <code>Data</code> : A list of <code>methylRaw</code> objects. Each object stores the raw methylation data of one sample. Default: TRUE.
runAnalysis	a logical, if TRUE, two files are saved for each simulation: <ul style="list-style-type: none"> <li>1. The first file is the <code>methylObj...</code> file formatted with the <code>methyKit</code> package in a <code>S4 methylBase</code> object (using the <code>methyKit</code> functions: <code>filterByCoverage</code>, <code>normalizeCoverage</code> and <code>unite</code>).</li> <li>2. The second file contains a <code>S4 calculateDiffMeth</code> object generated using the <code>methyKit</code> functions <code>calculateDiffMeth</code> on the first file.</li> </ul> Default: FALSE.
nbCores	a positive integer, the number of cores used when creating the simulated datasets. Default: 1 and always 1 for Windows.
vSeed	a integer, a seed used when reproducible results are needed. When a value inferior or equal to zero is given, a random integer is used. Default: -1.

## Value

0 indicating that the function have been successful.

**Author(s)**

Pascal Belleau, Astrid Deschenes

**See Also**

the vignette for detail description of the files created by the simulation.

**Examples**

```
## Load dataset containing methyl information
data(samplesForChrSynthetic)

## Set the output directory where files will be created
temp_dir <- "test_runSim"

## Create 2 simulated dataset (nbSimulation = 2)
## over 3 generations (nbGeneration = 3) with
## 6 cases and 6 controls (nNbSample = 6) using only one set
## of parameters (vpDiff = 0.9, vpDiffsd = 0.1, vDiff = 0.8)
runSim(methData = samplesForChrSynthetic, nbSynCHR = 1, nbSimulation = 2,
      nbGeneration = 3, nbBlock = 10, nbCpG = 20, vNbSample = c(6),
      vpDiff = c(0.9), vpDiffsd = c(0.1), vDiff = c(0.8),
      vInheritance = c(0.5), rateDiff = 0.3, minRate = 0.2,
      propInherite = 0.3, prophetero = 0.5, outputDir = temp_dir,
      fileID = "F", nbCores = 1, vSeed = 32)

## Delete the output directory and its content
if (dir.exists(temp_dir)) {
  unlink(temp_dir, recursive = TRUE, force = FALSE)
}
```

---

`samplesForChrSynthetic`

*All samples information, formatted by methylKit, in a methylBase format (for demo purpose).*

---

**Description**

The object is a `methylBase`. There is 12 samples (6 controls and 6 cases). Each sample information is stored in a `methylRaw` object.

**Usage**

```
data(samplesForChrSynthetic)
```

**Format**

A `methylBase` object contains the information for one generation. Each sample information is stored in a `methylRaw` object. There is `methylRaw` objects (6 controls and 6 cases).

**Details**

This dataset can be used to test the runSim function.

**Value**

A methylBase contains the information for one generation. Each sample information is stored in a methylRaw object. There is methylRaw objects (6 controls and 6 cases).

**See Also**

- [runSim](#) for running a simulation analysis using methylKit info as input

**Examples**

```
## Loading dataset
data(samplesForChrSynthetic)

## Set the output directory where files will be created
temp_dir <- "test_samplesForChrSynthetic"

## Create 4 simulated dataset (nbSimulation)
## over 3 generations (nbGeneration = 3) with
## 6 cases and 6 controls (nNbSample = 6) using only one set
## of parameters (vpDiff = 0.85, vpDiffsd = 0.1, vDiff = 0.8)
runSim(outputDir = temp_dir, fileID = "F1", nbSynCHR = 1,
      methData = samplesForChrSynthetic, nbSimulation = 4,
      nbBlock = 10, nbCpG = 20,
      nbGeneration = 3, vNbSample = c(6), vpDiff = c(0.85),
      vpDiffsd = c(0.1), vDiff = c(0.8),
      vInheritance = c(0.5), propInherite = 0.3,
      rateDiff = 0.3, minRate = 0.2, propHetero = 0.5,
      nbCores = 1, vSeed = 32)

## Delete the output directory and its content
if (dir.exists(temp_dir)) {
  unlink(temp_dir, recursive = TRUE, force = FALSE)
}
```

---

saveData

---

*Save data created during the simulation*


---

**Description**

Save data created during the simulation.

**Usage**

```
saveData(pathOut, extension, gRanges, methylData, methUnit, diffData,
  saveGRanges, saveMethylKit, runAnalysis)
```

**Arguments**

pathOut	a string of character, the path where the files are saved.
extension	a string of character representing the extension that will be given to the saved files.
gRanges	a list of methylRawList TODO
methylData	a list of methylRawList, the results of the normalization of the coverage.
methUnit	a list of methylBase, the results of the base filtering for all samples.
diffData	a list of methylDiff, the results of the calculation of differential methylation statistics.
saveGRanges	a logical, when true, files containing GRangeaList are saved.
saveMethylKit	a logical, when TRUE, files methylRawList object are saved.
runAnalysis	a logical, when TRUE, two files related to the analysis are saved.

**Value**

0 indicating that the function has been successful.

**Author(s)**

Pascal Belleau, Astrid Deschenes

**Examples**

```
## TODO
```

---

simEachGeneration	<i>Simulate a multigeneration methylation experiment with inheritance</i>
-------------------	---------------------------------------------------------------------------

---

**Description**

Simulate a multigeneration methylation case versus control experiment with inheritance relation using a real control dataset.

The simulation can be parametrized to fit different models. The number of cases and controls, the proportion of the case affected by the treatment (penetrance), the effect of the treatment on the mean of the distribution, the proportion of sites inherited, the proportion of the differentially methylated sites from the precedent generation inherited, etc..

The function simulates a multigeneration dataset like a bisulfite sequencing experiment. The simulation includes the information about control and case for each generation.

**Usage**

```
simEachGeneration(simulation, nbCtrl, nbCase, treatment, sample.id, generation,
  stateInfo, minReads, maxPercReads, context, assembly, meanCov, saveGRanges,
  saveMethylKit, runAnalysis)
```

**Arguments**

nbCtrl	a positive integer, the number of controls.
nbCase	a positive integer, the number of cases.
treatment	a numeric vector denoting controls and cases
sample.id	a matrix the name of each samples for each generation (row) and each case and control (column).
generation	a positive integer, the number of generations simulated.
stateInfo	a GRanges that contains the CpG (or methylated sites). The GRanges have four metadata from the real dataset: <ul style="list-style-type: none"> <li>• chrOri a numeric, the chromosome from the real dataset</li> <li>• startOri a numeric, the position of the site in the real dataset</li> <li>• meanCTRL a numeric, the mean of the control in the real dataset</li> <li>• varCTRL a numeric, the variance of the control in the real dataset.</li> </ul>
minReads	a positive integer, sites and regions having lower coverage than this count are discarded. The parameter corresponds to the lo.count parameter in the methylKit package.
maxPercReads	a double between [0,100], the percentile of read counts that is going to be used as upper cutoff. Sites and regions having higher coverage than maxPercReads are discarded. This parameter is used for both CpG sites and tiles analysis. The parameter correspond to the hi.perc parameter in the methylKit package.
context	a string of character, the short description of the methylation context, such as "CpG", "CpH", "CHH", etc..
assembly	a string of character, the short description of the genome assembly, such as "mm9", "hg18", etc..
meanCov	a positive integer, the mean of the coverage at the simulated CpG sites.
saveGRanges	a logical, when true, the package save two files type. The first generate for each simulation contains a list. The length of the list corresponds to the number of generation. The generation are stored in order (first entry = first generation, second entry = second generation, etc..). All samples related to one generations are contained in a GRangesList. The GRangesList store a list of GRanges. Each GRanges stores the raw methylation data of one sample. The second file a numeric vector denoting controls and cases (a file is generated by entry in the vector parameters vNbSample).
saveMethylKit	a logical, when TRUE, the package save a file contains a list. The length of the list corresponds to the number of generation. The generation are stored in order (first entry = first generation, second entry = second generation, etc..). All samples related to one generations are contained in a S4 methylRawList object. The methylRawList object contains two Slots: 1. treatment: A numeric vector denoting controls and cases. 2. .Data: A list of methylRaw objects. Each object stores the raw methylation data of one sample.
runAnalysis	a logical, if TRUE, two files are saved : <ul style="list-style-type: none"> <li>• 1. The first file is the methylObj... file formatted with the methylkit package in a S4 methylBase object (with the methylKit functions: filterByCoverage, normalizeCoverage and unite).</li> </ul>

- 2. The second file contains a S4 calculateDiffMeth object generated with the methylKit functions calculateDiffMeth using the first file.

### Value

0 indicating that the function has been successful.

### Author(s)

Pascal Belleau, Astrid Deschenes

### Examples

```
## Load dataset
data("samplesForChrSynthetic")
data("dataSimExample")

## Generate a stateInfo object using samples
stateInformation <- methInheritSim::getSyntheticChr(methInfo =
  samplesForChrSynthetic, nbBlock = 1, nbCpG = 3)

## Generate a stateDiff and stateInherit objects with length corresponding
## to nbBlock * nbCpG from stateInformation
stateDiff <- c(1, 0, 1)
stateInherit <- c(1, 0, 0)

## Create simulation
sim <- methInheritSim::getSim(nbCtrl = 3, nbCase = 2,
  generation = 3, stateInfo = stateInformation, stateDiff = stateDiff,
  stateInherit = stateInherit, diffValue = 10,
  propDiff = 0.8, propDiffsd = 0.2, propInheritance = 0.8,
  propHetero = 0.1)

## TODO
methInheritSim::simEachGeneration(simulation = sim,
  nbCtrl = 3, nbCase = 2, treatment = c(0,0,0,1,1),
  sample.id = dataSimExample$sample.id,
  generation = 3, stateInfo = stateInformation, minReads = 10,
  maxPercReads = 99, context = "Cpg", assembly = "RNOR_5.0", meanCov = 80,
  saveGRanges = FALSE, saveMethylKit = FALSE, runAnalysis = FALSE)
```

---

simInheritance

*Simulate a multigenerational methylation experiment with inheritance*

---

### Description

Simulate a multigenerational methylation case versus control experiment with inheritance relation using a real control dataset.



The simulation can be parametrized to fit different models. The number of cases and controls, the proportion of the case affected by the treatment (penetrance), the effect of the treatment on the mean of the distribution, the proportion of sites inherited, the proportion of the differentially methylated sites from the precedent generation inherited, etc..

The function simulates a multigeneration dataset like a bisulfite sequencing experiment. The simulation includes the information about control and case for each generation.

## Usage

```
simInheritance(pathOut, pref, k, nbCtrl, nbCase, treatment, sample.id,
  generation, stateInfo, propDiff, propDiffsd, diffValue, propInheritance,
  rateDiff, minRate, propInherite, propHetero, minReads, maxPercReads, context,
  assembly, meanCov, diffRes, saveGRanges, saveMethylKit, runAnalysis)
```

## Arguments

pathOut	a string of character or NULL, the path where the files created by the function will be saved. When NULL, the files are saved in the current directory.
pref	a string of character representing the parameters of specific simulation the string is composed of those elements, separated by "_": <ul style="list-style-type: none"> <li>• a fileID</li> <li>• the chromosome number, a number between 1 and nbSynCHR</li> <li>• the number of samples, a number in the vNbSample vector</li> <li>• the mean proportion of samples that has, for a specific position, differentially methylated values, a number in the vpDiff vector</li> <li>• the proportion of C/T for a case differentially methylated that follows a shifted beta distribution, a number in the vDiff vector</li> <li>• the proportion of cases that inherits differentially sites, a number in the vInheritance vector</li> </ul>
k	a positive integer, an ID for the current simulation.
nbCtrl	a positive integer, the number of controls.
nbCase	a positive integer, the number of cases.
treatment	a vector of integer denoting controls and cases. The vector length must correspond to the sum of cases and controls.
sample.id	a matrix the name of each samples for each generation (row) and each case and control (column).
generation	a positive integer, the number of generations simulated.
stateInfo	a GRanges that contains the CpG (or methylated sites). The GRanges have four metadata from the real dataset: <ul style="list-style-type: none"> <li>• chrOri a numeric, the chromosome from the real dataset</li> <li>• startOri a numeric, the position of the site in the real dataset</li> <li>• meanCTRL a numeric, the mean of the control in the real dataset</li> <li>• varCTRL a numeric, the variance of the control in the real dataset.</li> </ul>

propDiff	a double superior to 0 and inferior or equal to 1, the mean value for the proportion of samples that will have, for a specific position, differentially methylated values. It can be interpreted as the penetrance.
propDiffSd	a non-negative double, the standard deviation associated to the vpDiff. Note that vpDiff and vpDiffSd must be the same length.
diffValue	a non-negative double included in [0,1], the proportion of C/T for a case differentially methylated that follows a beta distribution where the mean is shifted by vDiff from the CTRL distribution.
propInheritance	a non-negative double included in [0,1], the proportion of cases that inherits differentially methylated sites.
rateDiff	a positive double inferior to 1, the mean of the chance that a site is differentially methylated.
minRate	a non-negative double inferior to 1, the minimum rate for differentially methylated sites. Default: 0.01.
propInherite	a non-negative double inferior or equal to 1, the proportion of differentially methylated regions that are inherited.
propHetero	a non-negative double between [0,1], the reduction of vDiff for the second and following generations.
minReads	a positive integer, sites and regions having lower coverage than this count are discarded. The parameter corresponds to the lo.count parameter in the methylKit package.
maxPercReads	a double between [0,100], the percentile of read counts that is going to be used as upper cutoff. Sites and regions having higher coverage than maxPercReads are discarded. This parameter is used for both CpG sites and tiles analysis. The parameter correspond to the hi.perc parameter in the methylKit package.
context	a string of character, the short description of the methylation context, such as "CpG", "CpH", "CHH", etc..
assembly	a string of character, the short description of the genome assembly, such as "mm9", "hg18", etc..
meanCov	a positive integer, the mean of the coverage at the simulated CpG sites.
diffRes	a list with 2 entries: <ul style="list-style-type: none"> <li>• stateDiff a vector of integer (0 and 1) with length corresponding the length of stateInfo. The vector indicates, using a 1, the positions where the CpG sites are differentially methylated.</li> <li>• stateInherite a vector of integer (0 and 1) with length corresponding the length of stateInfo. The vector indicates, using a 1, the positions where the CpG values are inherited.</li> </ul> when is NULL generate a new ones with getDiffMeth.
saveGRanges	a logical, when true, the package save two files type. The first generate for each simulation contains a list. The length of the list corresponds to the number of generation. The generation are stored in order (first entry = first generation, second entry = second generation, etc..). All samples related to one generations are contained in a GRangesList. The GRangesList store a list

of GRanges. Each GRanges stores the raw methylation data of one sample. The second file a numeric vector denoting controls and cases (a file is generated by entry in the vector parameters vNbSample).

- saveMethylKit** a logical, when TRUE, the package save a file contains a list. The length of the list corresponds to the number of generation. The generation are stored in order (first entry = first generation, second entry = second generation, etc..). All samples related to one generations are contained in a S4 methylRawList object. The methylRawList object contains two Slots: 1. treatment: A numeric vector denoting controls and cases. 2. .Data: A list of methylRaw objects. Each object stores the raw methylation data of one sample.
- runAnalysis** a logical, if TRUE, two files are saved :
- 1. The first file is the methylObj... file formatted with the methylkit package in a S4 methylBase object (with the methylKit functions: filterByCoverage, normalizeCoverage and unite).
  - 2. The second file contains a S4 calculateDiffMeth object generated with the methylKit functions calculateDiffMeth using the first file.

## Value

0 indicating that the function has been successful.

## Author(s)

Pascal Belleau, Astrid Deschenes

## Examples

```
## Name of the directory that will contained the generated files
temp_dir <- "test_simInheritance"

## Load dataset
data(dataSimExample)

## Generate a stateDiff object with length corresponding to
## nbBlock * nbCpG from stateInformation
stateDiff <- list()
stateDiff[["stateDiff"]] <- c(1, 0, 1)
stateDiff[["stateInherite"]] <- c(1, 0, 0)

## Simulate multigenerational methylation experiment with inheritance
methInheritSim::simInheritance(pathOut = temp_dir,
  pref = "S1_6_0.9_0.8_0.5", k = 1, nbCtrl = 6, nbCase = 6,
  treatment = dataSimExample$treatment,
  sample.id = dataSimExample$sample.id,
  generation = 3, stateInfo = dataSimExample$stateInfo[1:3],
  propDiff = 0.9, propDiffsd = 0.1, diffValue = 0.8,
  propInheritance = 0.5, rateDiff = 0.3, minRate = 0.3,
  propInherite = 0.3, propHetero = 0.5, minReads = 10, maxPercReads = 99,
  assembly="RNOR_5.0", context="Cpg", meanCov = 40, diffRes = stateDiff,
  saveGRanges = FALSE, saveMethylKit = FALSE, runAnalysis = FALSE)
```

```
## Delete directory
if (dir.exists(temp_dir)) {
  unlink(temp_dir, recursive = TRUE, force = FALSE)
}
```

---

testIfAlreadyDone	<i>Test if a specific simulation has already be done.</i>
-------------------	-----------------------------------------------------------

---

## Description

Test if a specific simulation has already be done.

## Usage

```
testIfAlreadyDone(pathOut, preference, id, saveGRanges, saveMethylKit,
  runAnalysis)
```

## Arguments

pathOut	a string of character, the path where the files are saved.
preference	a string of character representing the parameters of specific simulation.
id	a positive integer, a ID for the current simulation.
saveGRanges	a logical, when true, files containing GRangeaList are saved.
saveMethylKit	a logical, when TRUE, files methylRawList object are saved.
runAnalysis	a logical, when TRUE, two files related to the analysis are saved.

## Value

logical indicating if the simulation has already done.

## Author(s)

Pascal Belleau, Astrid Deschenes

## Examples

```
## Return TRUE when the specified simulation has already be done;
## otherwise, return FALSE.
methInheritSim::testIfAlreadyDone(pathOut = ".",
  preference = "S1_6_0.9_0.8_0.5", id = 33,
  saveGRanges = FALSE, saveMethylKit = FALSE, runAnalysis = FALSE)
```

---

 validateRunSimDoubleParameters

*Parameters validation for the `runSim` function. Only double parameters are validated.*

---

## Description

Validation of all parameters needed by the public `runSim` function. Only double parameters are validated.

## Usage

```
validateRunSimDoubleParameters(vpDiff, vpDiffstd, vDiff, vInheritance,
  propInherite, rateDiff, minRate, propHetero, maxPercReads)
```

## Arguments

<code>vpDiff</code>	a double superior to 0 and inferior or equal to 1, the mean value for the proportion of samples that will have, for a specific position, differentially methylated values. It can be interpreted as the penetrance.
<code>vpDiffstd</code>	a non-negative double, the standard deviation associated to the <code>vpDiff</code> .
<code>vDiff</code>	a positive double between [0,1], the proportion of C/T for a case differentially methylated follow a beta distribution where the mean is shifted of <code>vDiff</code> from the CTRL distribution
<code>vInheritance</code>	a positive double between [0,1], the proportion of cases that inherited differentially sites.
<code>propInherite</code>	a non-negative double inferior or equal to 1, the proportion of differentially methylated site are inherited
<code>rateDiff</code>	a positive double inferior to 1, the mean of the chance that a site is differentially methylated.
<code>minRate</code>	a non-negative double inferior to 1, the minimum rate of differentially methylated sites.
<code>propHetero</code>	a positive double between [0,1], the reduction of <code>vDiff</code> for the second and following generations.
<code>maxPercReads</code>	a double between [0,100], the percentile of read counts that is going to be used as upper cutoff. Bases ore regions having higher coverage than this percentile are discarded. Parameter used for both CpG sites and tiles analysis. The parameter correspond to the <code>hi.perc</code> parameter in the <code>methylKit</code> package.

## Value

0 indicating that the function has been successful.

## Author(s)

Pascal Belleau, Astrid Deschenes

## Examples

```
## The function returns 0 when all paramaters are valid
methInheritSim::validateRunSimDoubleParameters(vpDiff = 0.2,
vpDiffsd = 0.3, vDiff = 0.4, vInheritance = 0.2, propInherite = 0.5,
rateDiff = 0.2, minRate = 0.1, propHetero = 0.2, maxPercReads = 99.1)
```

---

```
validateRunSimIntegerParameters
```

*Parameters validation for the `runSim` function. Only integer parameters are validated.*

---

## Description

Validation of all parameters needed by the public `runSim` function. Only integer parameters are validated.

## Usage

```
validateRunSimIntegerParameters(nbSynCHR, nbSimulation, nbBlock, nbCpG,
vNbSample, nbGeneration, minReads, meanCov, nbCores, vSeed)
```

## Arguments

<code>nbSynCHR</code>	a positive integer, the number of distinct synthetic chromosomes that will be generated.
<code>nbSimulation</code>	a positive integer, the number of simulations for each parameter ( <code>vNbSample</code> , <code>vpDiff</code> , <code>vDiff</code> and <code>vInheritance</code> ).
<code>nbBlock</code>	a positive integer, the number of blocks used for sampling.
<code>nbCpG</code>	a positive integer, the number of consecutive CpG positions used for sampling from <code>methInfo</code> .
<code>vNbSample</code>	a vector of positive integer, the number of <code>methData</code> (CTRL) and cases in the the simulation dataset. In the simulated dataset, the number of CTRL equals the number of Case. The number of CTRL do not need to be equal to the number of Case in the real dataset.
<code>nbGeneration</code>	a positive integer, the number of generations.
<code>minReads</code>	a positive integer Bases and regions having lower coverage than this count are discarded. The parameter correspond to the <code>lo.count</code> parameter in the <code>methyKit</code> package.
<code>meanCov</code>	a positive integer represent the mean of the coverage at the CpG site Default: 80.
<code>nbCores</code>	a positive integer, the number of cores to use when creating the simulated datasets. Default: 1 and always 1 for Windows.
<code>vSeed</code>	a integer, a seed used when reproducible results are needed. When a value inferior or equal to zero is given, a random integer is used. Default: -1.

**Value**

0 indicating that the function has been successful.

**Author(s)**

Pascal Belleau, Astrid Deschenes

**Examples**

```
## The function returns 0 when all paramaters are valid
methInheritSim::validateRunSimIntegerParameters(nbSynCHR = 1,
nbSimulation = 2, nbBlock = 10, nbCpG = 4, vNbSample = 10,
nbGeneration = 3, minReads = 10, meanCov = 80,
nbCores = 1, vSeed = -1)
```

---

```
validateRunSimLogicalParameters
```

*Parameters validation for the [runSim](#) function. Only logical parameters are validated.*

---

**Description**

Validation of all parameters needed by the public [runSim](#) function. Only logical parameters are validated.

**Usage**

```
validateRunSimLogicalParameters(keepDiff, saveGRanges, saveMethylKit,
runAnalysis)
```

**Arguments**

keepDiff	logical if true, the differentially methylated sites will be the same for each parameter (vpDiff, vDiff and vInheritance). Default: FALSE.
saveGRanges	a logical, when true, the package save two files type. The first generate for each simulation contains a list. The length of the list corresponds to the number of generation. The generation are stored in order (first entry = first generation, second entry = second generation, etc..). All samples related to one generations are contained in a GRangesList. The GRangesList store a list of GRanges. Each GRanges stores the raw methylation data of one sample. The second file a numeric vector denoting controls and cases (a file is generated by entry in the vector parameters vNbSample).
saveMethylKit	a logical, when TRUE, for each simulations save a file contains a list. The length of the list corresponds to the number of generation. The generation are stored in order (first entry = first generation, second entry = second generation, etc..). All samples related to one generations are contained in a S4

methylRawList object. The methylRawList object contains two Slots: 1. treatment: A numeric vector denoting controls and cases. 2. .Data: A list of methylRaw objects. Each object stores the raw methylation data of one sample.

runAnalysis      a logical, if TRUE, two files are saved for each simulation:

- 1. The first file is the methylObj... file formatted with the methylkit package in a S4 methylBase object (with the methylKit functions: filterByCoverage, normalizeCoverage and unite).
- 2. The second file contains a S4 calculateDiffMeth object generated with the methylKit functions calculateDiffMeth on the first file.

### Value

0 indicating that the function has been successful.

### Author(s)

Pascal Belleau, Astrid Deschenes

### Examples

```
## Load dataset
data("samplesForChrSynthetic")

## The function returns 0 when all paramaters are valid
methInheritSim::validateRunSimLogicalParameters(keepDiff = FALSE,
saveGRanges = TRUE, saveMethylKit = FALSE, runAnalysis = FALSE)
```

---

validateRunSimOtherParameters

*Parameters validation for the [runSim](#) function. Only parameters other than double, integer and logical are validated.*

---

### Description

Validation of all parameters needed by the public [runSim](#) function. Only parameters other than double, integer and logical are validated.

### Usage

```
validateRunSimOtherParameters(outputDir, fileID, methData, context, assembly)
```



**Arguments**

outputDir	a string of character or NULL, the path where the files created by the function will be saved. When NULL, the files are saved in the current directory.
fileID	a string of character, a identifiant that will be included in each output file name. Each output file name is composed of those elements, separated by "_": <ul style="list-style-type: none"> <li>• a type name, ex: methylGR, methylObj, etc..</li> <li>• a fileID</li> <li>• the chromosome number, a number between 1 and nbSynCHR</li> <li>• the number of samples, a number in the vNbSample vector</li> <li>• the mean proportion of samples that has, for a specific position, differentially methylated values, a number in the vpDiff vector</li> <li>• the proportion of C/T for a case differentially methylated that follows a shifted beta distribution, a number in the vDiff vector</li> <li>• the proportion of cases that inherits differentially sites, a number in the vInheritance vector</li> <li>• the identifiant for the simulation, a number between 1 and nbSimulation</li> <li>• the file extension ".rds"</li> </ul>
methData	an object of class methylBase, the CpG information from controls (CTRL) that will be used to create the sythetic chromosome. The methData object can also contain information from cases but only the controls will be used.
context	a string of character, the methylation context string, ex: CpG,CpH,CHH, etc.
assembly	a string of character, the short description of the genome assembly. Ex: mm9,hg18 etc.

**Value**

0 indicating that the function has been successful.

**Author(s)**

Pascal Belleau, Astrid Deschenes

**Examples**

```
## Load dataset
data("samplesForChrSynthetic")

## The function returns 0 when all paramaters are valid
methInheritSim::validateRunSimOtherParameters(
  outputDir = "test", fileID = "test", methData = samplesForChrSynthetic,
  context = "CpG", assembly = "Rnor_5.0")
```

---

```
validateRunSimParameters
```

*Parameters validation for the [runSim](#) function.*

---

## Description

Validation of all parameters needed by the public [runSim](#) function.

## Usage

```
validateRunSimParameters(vpDiff, vpDiffSd, vDiff, vInheritance, propInherite,
  rateDiff, minRate, propHetero, maxPercReads, nbSynCHR, nbSimulation, nbBlock,
  nbCpG, vNbSample, nbGeneration, minReads, meanCov, nbCores, vSeed, keepDiff,
  saveGRanges, saveMethylKit, runAnalysis, outputDir, fileID, methData, context,
  assembly)
```

## Arguments

vpDiff	a double superior to 0 and inferior or equal to 1, the mean value for the proportion of samples that will have, for a specific position, differentially methylated values. It can be interpreted as the penetrance.
vpDiffSd	a non-negative double, the standard deviation associated to the propDiff.
vDiff	a positive double between [0,1], the proportion of C/T for a case differentially methylated follow a beta distribution where the mean is shifted of vDiff from the CTRL distribution
vInheritance	a positive double between [0,1], the proportion of cases that inherited differentially sites.
propInherite	a non-negative double inferior or equal to 1, the proportion of differentially methylated site are inherited
rateDiff	a positive double inferior to 1, the mean of the chance that a site is differentially methylated.
minRate	a non-negative double inferior to 1, the minimum rate of differentially methylated sites.
propHetero	a positive double between [0,1], the reduction of vDiff for the second and following generations.
maxPercReads	a double between [0,100], the percentile of read counts that is going to be used as upper cutoff. Bases ore regions having higher coverage than this percentile are discarded. Parameter used for both CpG sites and tiles analysis. The parameter correspond to the <code>hi.perc</code> parameter in the <code>methylKit</code> package.
nbSynCHR	a positive integer, the number of distinct synthetic chromosomes that will be generated.
nbSimulation	a positive integer, the number of simulations for each parameter (vNbSample, vpDiff, vDiff and vInheritance).
nbBlock	a positive integer, the number of blocks used for sampling.

nbCpG	a positive integer, the number of consecutive CpG positions used for sampling from methInfo.
vNbSample	a vector of positive integer, the number of methData (CTRL) and cases in the the simulation dataset. In the simulated dataset, the number of CTRL equals the number of Case. The number of CTRL do not need to be equal to the number of Case in the real dataset.
nbGeneration	a positive integer, the number of generations.
minReads	a positive integer Bases and regions having lower coverage than this count are discarded. The parameter correspond to the lo.count parameter in the methylKit package.
meanCov	a positive integer represent the mean of the coverage at the CpG site Default: 80.
nbCores	a positive integer, the number of cores to use when creating the simulated datasets. Default: 1 and always 1 for Windows.
vSeed	a integer, a seed used when reproducible results are needed. When a value inferior or equal to zero is given, a random integer is used. Default: -1.
keepDiff	logical if true, the differentially methylated sites will be the same for each parameter (vpDiff, vDiff and vInheritance). Default: FALSE.
saveGRanges	a logical, when true, the package save two files type. The first generate for each simulation contains a list. The length of the list corresponds to the number of generation. The generation are stored in order (first entry = first generation, second entry = second generation, etc..). All samples related to one generations are contained in a GRangesList. The GRangesList store a list of GRanges. Each GRanges stores the raw methylation data of one sample. The second file a numeric vector denoting controls and cases (a file is generates by entry in the vector parameters vNbSample).
saveMethylKit	a logical, when TRUE, for each simulations save a file contains a list. The length of the list corresponds to the number of generation. The generation are stored in order (first entry = first generation, second entry = second generation, etc..). All samples related to one generations are contained in a S4 methylRawList object. The methylRawList object contains two Slots: 1. treatment: A numeric vector denoting controls and cases. 2. .Data: A list of methylRaw objects. Each object stores the raw methylation data of one sample.
runAnalysis	a logical, if TRUE, two files are saved for each simulation: <ul style="list-style-type: none"> <li>• 1. The first file is the methylObj... file formatted with the methylkit package in a S4 methylBase object (with the methylKit functions: filterByCoverage, normalizeCoverage and unite).</li> <li>• 2. The second file contains a S4 calculateDiffMeth object generated with the methylKit functions calculateDiffMeth on the first file.</li> </ul>
outputDir	a string of character or NULL, the path where the files created by the function will be saved. When NULL, the files are saved in the current directory.
fileID	a string of character, a identifiant that will be included in each output file name. Each output file name is composed of those elements, separated by "_": <ul style="list-style-type: none"> <li>• a type name, ex: methylGR, methylObj, etc..</li> </ul>

	<ul style="list-style-type: none"> <li>• a fileID</li> <li>• the chromosome number, a number between 1 and nbSynCHR</li> <li>• the number of samples, a number in the vNbSample vector</li> <li>• the mean proportion of samples that has, for a specific position, differentially methylated values, a number in the vpDiff vector</li> <li>• the proportion of C/T for a case differentially methylated that follows a shifted beta distribution, a number in the vDiff vector</li> <li>• the proportion of cases that inherits differentially sites, a number in the vInheritance vector</li> <li>• the identifiant for the simulation, a number between 1 and nbSimulation</li> <li>• the file extension ".rds"</li> </ul>
methData	an object of class methylBase, the CpG information from controls (CTRL) that will be used to create the sythetic chromosome. The methData object can also contain information from cases but only the controls will be used.
context	a string of character, the methylation context string, ex: CpG,CpH,CHH, etc.
assembly	a string of character, the short description of the genome assembly. Ex: mm9,hg18 etc.

### Value

0 indicating that the function has been successful.

### Author(s)

Pascal Belleau, Astrid Deschenes

### Examples

```
## Load dataset
data("samplesForChrSynthetic")

## The function returns 0 when all paramaters are valid
methInheritSim::validateRunSimParameters(vpDiff = 0.2,
vpDiffsd = 0.3, vDiff = 0.4, vInheritance = 0.2, propInherite = 0.5,
rateDiff = 0.2, minRate = 0.1, propHetero = 0.2, maxPercReads = 99.1,
nbSynCHR = 1, nbSimulation = 2, nbBlock = 10, nbCpG = 4, vNbSample = 10,
nbGeneration = 3, minReads = 10, meanCov = 80,
nbCores = 1, vSeed = -1, keepDiff = FALSE, saveGRanges = TRUE,
saveMethylKit = FALSE, runAnalysis = FALSE, outputDir = "test",
fileID = "test", methData = samplesForChrSynthetic,
context = "CpG", assembly = "Rnor_5.0")
```

# Index

- \* **datasets**
  - dataSimExample, [4](#)
  - samplesForChrSynthetic, [20](#)
- \* **internal**
  - calculateNbDiffCase, [3](#)
  - createSampleID, [4](#)
  - estBetaAlpha, [6](#)
  - estBetaBeta, [7](#)
  - fixSeed, [7](#)
  - getDiffCase, [8](#)
  - getDiffMeth, [9](#)
  - getSim, [11](#)
  - getSyntheticChr, [12](#)
  - runOnEachSynCHR, [13](#)
  - saveData, [21](#)
  - simEachGeneration, [22](#)
  - simInheritance, [24](#)
  - testIfAlreadyDone, [28](#)
  - validateRunSimDoubleParameters, [29](#)
  - validateRunSimIntegerParameters, [30](#)
  - validateRunSimLogicalParameters, [31](#)
  - validateRunSimOtherParameters, [32](#)
  - validateRunSimParameters, [34](#)
- \* **package**
  - methInheritSim-package, [2](#)

calculateNbDiffCase, [3](#)

createSampleID, [4](#)

dataSimExample, [4](#)

estBetaAlpha, [6](#)

estBetaBeta, [7](#)

fixSeed, [7](#)

getDiffCase, [8](#)

getDiffMeth, [9](#)

getSim, [11](#)

getSyntheticChr, [12](#)

methInheritSim  
(methInheritSim-package), [2](#)

methInheritSim-package, [2](#)

runOnEachSynCHR, [13](#)

runSim, [3](#), [5](#), [17](#), [21](#), [29–32](#), [34](#)

samplesForChrSynthetic, [20](#)

saveData, [21](#)

simEachGeneration, [22](#)

simInheritance, [24](#)

testIfAlreadyDone, [28](#)

validateRunSimDoubleParameters, [29](#)

validateRunSimIntegerParameters, [30](#)

validateRunSimLogicalParameters, [31](#)

validateRunSimOtherParameters, [32](#)

validateRunSimParameters, [34](#)