

# Package ‘epigraHMM’

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**Description** epigraHMM provides a set of tools for the analysis of epigenomic data based on hidden Markov Models. It contains two separate peak callers, one for consensus peaks from biological or technical replicates, and one for differential peaks from multi-replicate multi-condition experiments. In differential peak calling, epigraHMM provides window-specific posterior probabilities associated with every possible combinatorial pattern of read enrichment across conditions.

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addOffsets	<i>Add offsets to epigraHMMDataset</i>
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**Description**

This function adds model offsets to epigraHMMDataset

**Usage**

```
addOffsets(object, offsets)
```

**Arguments**

object	an epigraHMMDataset
offsets	a matrix with model offsets

**Details**

To be added

**Value**

An epigraHMMDataset with an 'offsets' assay filled in.

## References

<https://github.com/plbaldoni/epigraHMM>

## Examples

```
# Creating dummy object
countData <- list('counts' = matrix(rpois(4e5,10),ncol = 4),
'controls' = matrix(rpois(4e5,5),ncol = 4))
colData <- data.frame(condition = c('A','A','B','B'), replicate = c(1,2,1,2))
object <- epigraHMMDatasetFromMatrix(countData,colData)

# Adding pre-computed offsets
object <- addOffsets(object = object,
                    offsets = matrix(rnorm(4e5),ncol = 4))
```

---

callPatterns	<i>Extract posterior probabilities (or combinatorial patterns) associated with differential regions</i>
--------------	---

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## Description

Given results from epigraHMM's differential peak caller, this function will output either posterior probabilities or combinatorial patterns associated with the mixture components of the embedded mixture model.

## Usage

```
callPatterns(
  object,
  peaks,
  hdf5 = metadata(object)$output,
  type = "all",
  fdr = NULL,
  pattern = NULL,
  ranges = NULL
)
```

## Arguments

object	an epigraHMMDataset
peaks	a GRanges object with differential peaks from 'callPeaks'
hdf5	a character with the location of the epigraHMM HDF5 output file
type	a character string that defines which output will be given (see details; default is 'all')
fdr	the desired fdr thresholding level to define combinatorial patterns
pattern	a string that explicitly specifies the combinatorial pattern to be output
ranges	a GRanges object with the genomic ranges to subset the output

## Details

The output of ‘callPatterns’ is always restricted to genomic windows intersecting peaks.

If ‘type = ‘all’’, all windows’ posterior probabilities associated with all differential combinatorial patterns are returned. If ‘type = ‘fdr’’, users must also specify the input argument ‘pattern’ and this function will output windows which are associated with the given ‘pattern’ that pass a particular fdr threshold level. If ‘type = ‘max’’, this function will output the combinatorial pattern which has the maximal posterior probability for each window. If ‘type = ‘ranges’’, the windows that are output are restricted to those that intersect the ‘ranges’ input argument.

## Value

A GRanges object with metadata

## Author(s)

Pedro L. Baldoni, <pedrobaldoni@gmail.com>

## References

<https://github.com/plbaldoni/epigraHMM>

## Examples

```
# Creating dummy object
countData <- cbind(rbind(matrix(rnbinom(1e2, mu = 1, size = 10), ncol = 1),
  matrix(rnbinom(1e2, mu = 10, size = 5), ncol = 1),
  matrix(rnbinom(1e2, mu = 1, size = 10), ncol = 1),
  matrix(rnbinom(1e2, mu = 10, size = 5), ncol = 1),
  matrix(rnbinom(1e2, mu = 1, size = 10), ncol = 1),
  matrix(rnbinom(1e2, mu = 1, size = 10), ncol = 1),
  matrix(rnbinom(1e2, mu = 1, size = 10), ncol = 1)),
  rbind(matrix(rnbinom(1e2, mu = 1, size = 10), ncol = 1),
  matrix(rnbinom(1e2, mu = 1, size = 10), ncol = 1),
  matrix(rnbinom(1e2, mu = 1, size = 10), ncol = 1),
  matrix(rnbinom(1e2, mu = 10, size = 5), ncol = 1),
  matrix(rnbinom(1e2, mu = 1, size = 10), ncol = 1),
  matrix(rnbinom(1e2, mu = 10, size = 5), ncol = 1),
  matrix(rnbinom(1e2, mu = 1, size = 10), ncol = 1)))

colData <- data.frame(condition = c('A','B'), replicate = c(1,1))
rowRanges <- GenomicRanges::GRanges('chrA',
  IRanges::IRanges(start = seq(1,by = 500,
  length.out = nrow(countData)),width = 500))

object <- epigraHMMDataSetFromMatrix(countData,colData,rowRanges = rowRanges)

# Initializing
object <- initializer(object,controlEM())

# Running epigraHMM
object <- epigraHMM(object,controlEM(),type = 'differential',dist = 'nb')
```

```
# Calling peaks
peaks <- callPeaks(object = object,
                  hdf5 = S4Vectors::metadata(object)$output,
                  method = 'viterbi')

# Extracting posterior probabilities
patterns <- callPatterns(object = object, peaks = peaks, type = 'max')
```

---

callPeaks	<i>Summarize peak calls and optionally create a BED 6+3 file in broad-Peak format for visualization</i>
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---

### Description

This function imports the output from ‘epigraHMM’ and outputs a set of peaks (consensus or differential) for a given FDR control threshold or Viterbi sequence.

### Usage

```
callPeaks(
  object,
  hdf5 = metadata(object)$output,
  method = "viterbi",
  saveToFile = FALSE,
  control = NULL
)
```

### Arguments

object	an epigraHMMDataset
hdf5	a character with the location of the epigraHMM HDF5 output file
method	either ‘viterbi’ or a numeric FDR control threshold (e.g. 0.05). Default is ‘viterbi’.
saveToFile	a logical indicating whether or not to save the results to file. Output files are always saved with peaks of interest defined on the region level. Default is FALSE.
control	list of control arguments from controlEM(). This is an optional parameter and it is only required when ‘saveToFile = TRUE’ so that the output directory can be obtained. Default is NULL.

### Value

A GRanges object with differential peak calls in BED 6+3 format

### Author(s)

Pedro L. Baldoni, <pedrobaldoni@gmail.com>

## References

<https://github.com/plbaldoni/epigraHMM>

## Examples

```
# Creating dummy object
countData <- rbind(matrix(rnbinom(1e3,mu = 2,size = 10),ncol = 1),
                   matrix(rnbinom(2e3,mu = 7.5,size = 5),ncol = 1),
                   matrix(rnbinom(1e3,mu = 2,size = 10),ncol = 1))

colData <- data.frame(condition = 'A', replicate = 1)

rowRanges <- GenomicRanges::GRanges('chrA',
IRanges::IRanges(start = seq(from = 1, length.out = 4e3,by = 250),width = 250))

object <- epigraHMMDatasetFromMatrix(countData,colData,rowRanges)

# Initializing
object <- initializer(object,controlEM())

# Running epigraHMM
object <- epigraHMM(object,controlEM(),type = 'consensus',dist = 'nb')

# Calling peaks
peaks <- callPeaks(object = object,
                   hdf5 = S4Vectors::metadata(object)$output,
                   method = 'viterbi')
```

---

cleanCounts

*Remove effects from covariates of interest*

---

## Description

This function removes the effect from covariates of interest (such as GC content) from experimental counts

## Usage

```
cleanCounts(object, effectNames, byNames = NULL, log = TRUE)
```

## Arguments

object	an epigraHMMDataset
effectNames	a character vector with the names of assays for which the effect will be removed from the experimental counts. Names in ‘effectNames’ must be assays stored in the epigraHMMDataset ‘object’.

byNames	a character vector with the name of an assay containing stratification variables which will be used to define stratum-specific effects. Examples of byNames assays include the 'peaks' assay from 'initializer()'. In this case, models will be fit separately for peaks and non-peaks regions. This can be useful for effects such as GC content, which are known to have a differential effect between peaks and non-peak regions. Default is NULL, i.e., effects will be removed without stratification.
log	a logical indicating if the effect from 'effectNames' should be log-transformed in the regression model (default is TRUE)

**Value**

An epigraHMMDataset with an 'offset' assay filled in.

**Author(s)**

Pedro L. Baldoni, <pedrobaldoni@gmail.com>

**References**

<https://github.com/plbaldoni/epigraHMM>

**Examples**

```
# Creating dummy object
gc <- rbeta(3e3,50,50)

countData <- list('counts' = rbind(matrix(rnbinom(2e3,mu = 7.5,size = 10),ncol = 1),
                                   matrix(rnbinom(3e3,mu = exp(0.5 + 8*gc),size = 5),ncol = 1),
                                   matrix(rnbinom(2e3,mu = 7.5,size = 10),ncol = 1)),
                 'gc' = matrix(c(rbeta(2e3,50,50),gc,rbeta(2e3,50,50)),ncol = 1))

colData <- data.frame(condition = 'A', replicate = 1)
object <- epigraHMMDatasetFromMatrix(countData,colData)

# Initializing
object <- initializer(object = object,controlEM())

# Cleaning counts
object <- cleanCounts(object = object,effectNames = 'gc',byNames = 'peaks')

# Plotting the cleaned data
#par(mfrow = c(2,1))
#smoothScatter(log1p(assay(object))~assay(object,'gc'),xlab = 'gc',ylab = 'log counts')
#smoothScatter(as.numeric(log(assay(object)+1) - assay(object,'offsets'))~assay(object,'gc'),
#              xlab = 'gc',ylab = 'log cleaned counts')
```

controlEM

*Control parameters for the EM algorithm from epigraHMM***Description**

This function passes controlling parameters for the EM algorithm implemented in the epigraHMM package.

**Usage**

```
controlEM(
  epsilonEM = c(MRCPE = 0.001, MACPE = 0.001, ARCEL = 0.001),
  maxIterEM = 500,
  minIterEM = 3,
  gapIterEM = 3,
  maxCountEM = 3,
  maxDisp = 1000,
  criterion = "all",
  minZero = .Machine$double.xmin,
  probCut = 0.05,
  quiet = TRUE,
  maxIterInnerEM = 5,
  epsilonInnerEM = 0.001,
  trimOffset = 3,
  pattern = NULL,
  tempDir = tempdir(),
  fileName = "epigraHMM",
  pruningThreshold = NULL,
  quietPruning = TRUE
)
```

**Arguments**

epsilonEM	a named vector of positive values specifying up to four possible convergence criterion tolerances for the EM algorithm (see 'criterion' below). Default is c('MRCPE' = 1e-3, 'MACPE' = 1e-3, 'ARCEL' = 1e-3).
maxIterEM	a positive integer giving the maximum number of EM iterations. Default is 500.
minIterEM	a positive integer giving the minimum number of EM iterations to start evaluating the convergence. Default is 3.
gapIterEM	a positive integer giving the number of EM iterations apart to compute the convergence criterion. Default is 3.
maxCountEM	a positive integer giving the number of consecutive EM iterations satisfying the convergence criterion in order to stop the algorithm. Default is 3.
maxDisp	a positive value for the upper limit constraint of the dispersion parameters. Default is 1000.



critereon	a character specifying the convergence criterion. Either "MRCPE" (maximum absolute relative change in parameter estimates), "MACPE" (maximum absolute change of parameter estimates), "ARCEL" (absolute relative change of the Q-function), or "all" (simultaneously check for MRCPE, MACPE, and ARCEL). Default is "all".
minZero	a positive value for the minimum positive value allowed in computations to avoid having zeros. Default is <code>.Machine\$double.xmin</code> .
probCut	a number between 0 and 1 for the cutoff of the rejection controlled EM algorithm. Default 0.05.
quiet	a logical indicating whether to print messages. Default is TRUE.
maxIterInnerEM	a positive integer giving the maximum number of inner EM iterations. Default is 5.
epsilonInnerEM	a positive value with the convergence tolerance value for the inner EM algorithm. The criterion for the inner EM is "MRCPE". Default is 1e-3.
trimOffset	either NULL or a positive integer indicating the number of decimal places to be used in the offset. Default is 3.
pattern	either NULL (the default) or a list with length equal to the number of differential patterns to be modeled by the differential HMM state. See Details section below.
tempDir	a string where results will be saved. Default is <code>'tempdir()'</code> .
fileName	a string with the name of the result files. Default is <code>'epigraHMM'</code> .
pruningThreshold	a numeric value between 0 and 1 to consider when pruning rare combinatorial patterns. Default is NULL (see Details).
quietPruning	a logical indicating whether to print messages during the pruning step. Default is TRUE.

### Details

If `pattern` is NULL, every possible combinatorial pattern will be considered. If `pattern` is a list, elements of it should specify the differential patterns to be modeled by each mixture component. For instance, if `pattern = list(2,c(1,3))` the mixture model will have two components that will represent the enrichment of condition 2 alone and the enrichment of conditions 1 and 3 together.

If `pruningThreshold` is a value between 0 and 1, say 0.05, `epigraHMM` will sequentially remove differential combinatorial patterns of enrichment from any mixture model component with associated posterior mixture proportion less than 0.05.

### Value

A list with components equal to the arguments

### Author(s)

Pedro L. Baldoni, <[pedrobaldoni@gmail.com](mailto:pedrobaldoni@gmail.com)>

### References

<https://github.com/plbaldoni/epigraHMM>

## Examples

```
# No more than 100 EM iterations
control <- controlEM(maxIterEM = 100)
```

---

epigraHMM

*Perform peak calling of epigenomic data sets*

---

## Description

This function runs either consensus (one condition, multiple samples) or differential (multiple conditions and samples) peak callers for epigenomic data.

## Usage

```
epigraHMM(object, control, type, dist = "nb")
```

## Arguments

object	an epigraHMMDataset
control	list of control arguments from <code>controlEM</code>
type	character, either "consensus" or "differential"
dist	character, either "zinb" or "nb" (default)

## Value

An epigraHMMDataset object with the results from epigraHMM

## Author(s)

Pedro L. Baldoni, <pedrobaldoni@gmail.com>

## References

<https://github.com/plbaldoni/epigraHMM>

## Examples

```
# Creating dummy object
countData <- rbind(matrix(rnbinom(1e3,mu = 2,size = 10),ncol = 1),
                  matrix(rnbinom(2e3,mu = 7.5,size = 5),ncol = 1),
                  matrix(rnbinom(1e3,mu = 2,size = 10),ncol = 1))

colData <- data.frame(condition = 'A', replicate = 1)
object <- epigraHMMDatasetFromMatrix(countData,colData)

# Initializing
object <- initializer(object,controlEM())
```

```
# Running epigraHMM
object <- epigraHMM(object,controlEM(),type = 'consensus',dist = 'nb')
```

---

```
epigraHMMDatasetFromBam
```

*Create a epigraHMMDataset from a set of BAM files*

---

## Description

This function creates a [RangedSummarizedExperiment](#) object from of a set of BAM files. It is used to store the input data, the model offsets, and the results from the peak calling algorithms.

## Usage

```
epigraHMMDatasetFromBam(
  bamFiles,
  colData,
  genome,
  windowSize,
  gapTrack = TRUE,
  blacklist = TRUE
)
```

## Arguments

bamFiles	a string vector (or a list of string vectors) with the path for BAM files. If bamFiles is a list of string vectors, vectors must be named, have the same dimension, and, at least, a vector with name 'counts' must exist (see details).
colData	a data.frame with the experimental data. It must contain the columns condition and replicate. condition refers to the experimental condition identifier (e.g. cell line name). replicate refers to the replicate identification number (unique for each condition).
genome	either a single string with the name of the reference genome (e.g. 'hg19') or a GRanges object with ranges to be tiled into a set of non-overlapping windows.
windowSize	an integer specifying the size of genomic windows where read counts will be computed.
gapTrack	either a logical (TRUE, the default, or FALSE) or a GRanges object with gap regions of the genome to be excluded. If TRUE, the function will discard genomic coordinates overlapping regions present in the UCSC gap table of the respective reference genome (if available). See Details section below.
blacklist	either a logical (TRUE, the default, or FALSE) or a GRanges object with blacklisted regions of the genome to be excluded. If TRUE, the function will discard ENCODE blacklisted regions from selected reference genomes (if available). See Details section below.

## Details

The index ".bai" files must be stored in the same directory of their respective BAM files. The index files must be named after their respective BAM files with the additional ".bai" suffix.

'epigraHMMDatasetFromBam' will store experimental data (e.g. ChIP-seq counts) from bamFiles (or bamFiles[['counts']], if a list is provided). Additional data (e.g. input control counts) will be stored similarly with their respective list names.

By default, the function computes read counts using csaw's estimated fragment length via cross correlation analysis. For experimental counts (e.g. ChIP-seq), sequencing reads are shifted downstream half of the estimated fragment length. For additional counts (e.g. input control), sequencing reads are not shifted prior to counting.

Additional columns included in the colData input will be passed to the resulting epigraHMM-Dataset assay and can be accessed via colData() function.

The genome argument will call GenomeInfoDb::Seqinfo() to fetch the chromosome lengths of the specified genome. See ?GenomeInfoDb::Seqinfo for the list of UCSC genomes that are currently supported.

If gapTrack = TRUE and the name of a reference genome is passed as input through genome (e.g. 'hg19'), the function will discard any genomic coordinate overlapping regions specified by the respective UCSC gap table. If gapTrack is a GRanges object, the function will discard any genomic coordinate overlapping regions from gapTrack.

If blacklist = TRUE and the name of a reference genome is passed as input through genome (e.g. 'hg19'), The function will fetch the manually curated blacklist tracks (Version 2) from <https://github.com/Boyle-Lab/Blacklist/tree/master/lists>. Current available genomes are ce10, dm3, hg19, hg38, and mm10. If blacklist is a GRanges object, the function will discard any genomic coordinate overlapping regions from blacklist.

## Value

An epigraHMMDataset object with sorted colData regarding conditions and replicates. Experimental counts will be stored in the 'counts' assay in the resulting epigraHMMDataset object. Additional experimental data will be stored with their respective names from the list bamFiles.

## Author(s)

Pedro L. Baldoni, <pedrobaldoni@gmail.com>

## References

<https://github.com/plbaldoni/epigraHMM> DOI: 10.1093/nar/gkv1191 DOI: 10.1038/s41598-019-45839-z DOI: 10.1038/nature11247

## Examples

```
bamFiles <- system.file("extdata", "euratrans",
                       "lv-H3K27me3-SHR-male-bio2-tech1.bam",
                       package="chromstaRData")
```

```
colData <- data.frame(condition = 'SHR', replicate = 1)
```

```
object <- epigraHMMDatasetFromBam(bamFiles = bamFiles,
                                   colData = colData,
                                   genome = 'rn4',
                                   windowSize = 25000,
                                   gapTrack = TRUE,
                                   blacklist = TRUE)
```

---

epigraHMMDatasetFromMatrix

*Create a epigraHMMDataset from matrices of counts*

---

## Description

This function creates a [RangedSummarizedExperiment](#) object from matrices of counts. It is used to store the input data, the model offsets, and the results from the peak calling algorithms.

## Usage

```
epigraHMMDatasetFromMatrix(countData, colData, rowRanges = NULL)
```

## Arguments

countData	a matrix (or a list of matrices). If countData is a list of matrices, matrices must be named, have the same dimensions, and, at least, a matrix with name 'counts' must exist (see details).
colData	a data.frame with columns condition and replicate. condition refers to the experimental condition identifier (e.g. cell line name). replicate refers to the replicate identification number (unique for each condition).
rowRanges	an optional GRanges object with the genomic coordinates of the countData

## Details

Additional columns included in the colData input will be passed to the resulting epigraHMMDataset assay and can be accessed via colData() function.

## Value

An epigraHMMDataset object with sorted colData regarding conditions and replicates. Experimental counts will be stored in the 'counts' assay in the resulting epigraHMMDataset object. If 'countData' is a list of matrices, the resulting 'counts' assay will be equal to 'countData[['counts']]'.

Additional matrices can be included in the epigraHMMDataset. For example, if one wants to include counts from an input control experiment from 'countData[['controls']]', an assay 'control' will be added to the resulting epigraHMMDataset..

**Author(s)**

Pedro L. Baldoni, <pedrobaldoni@gmail.com>

**References**

<https://github.com/plbaldoni/epigraHMM>

**Examples**

```
countData <- list('counts' = matrix(rpois(4e5,10),ncol = 4),
  'controls' = matrix(rpois(4e5,5),ncol = 4))
colData <- data.frame(condition = c('A','A','B','B'), replicate = c(1,2,1,2))
object <- epigraHMMDataSetFromMatrix(countData,colData)
```

---

estimateTransitionProb

*Estimate transition probability from a sequence of integers*

---

**Description**

This function estimates the transition probabilities for a k-state Markov chain based on a sequence of integers that represent states of the chain

**Usage**

```
estimateTransitionProb(chain, numStates)
```

**Arguments**

chain	a vector of integers
numStates	an integer, the number of states in the Markov chain

**Value**

A k-by-k matrix of transition probabilities, such that k is the number of states of the chain

**References**

<https://github.com/plbaldoni/epigraHMM>

**Examples**

```
trueMat <- matrix(c(0.9,0.1,0.1,0.9),2,2)
simChain <- simulateMarkovChain(trueMat,1e3)
estMat <- estimateTransitionProb(simChain,2)

# estMat should be close to trueMat
estMat
```

---

expStep	<i>E-step of HMM (forward-backward probability + posterior probability calculation)</i>
---------	---

---

## Description

E-step of HMM (forward-backward probability + posterior probability calculation)

## Usage

```
expStep(pi, gamma, logf, hdf5)
```

## Arguments

pi	a vector of probabilities (sum of probabilities should sum to one)
gamma	a matrix of transition probabilities (row sums should be one)
logf	a matrix of observed log-likelihood values. Columns represent hidden states, rows represent genomic regions
hdf5	path to where the hdf5 is saved

## Examples

```
#Creating dummy object
countData <- rbind(matrix(rnbinom(1e3,mu = 2,size = 10),ncol = 1),
                  matrix(rnbinom(2e3,mu = 7.5,size = 5),ncol = 1),
                  matrix(rnbinom(1e3,mu = 2,size = 10),ncol = 1))

colData <- data.frame(condition = 'A', replicate = 1)
object <- epigraHMMDataSetFromMatrix(countData,colData)

#Initializing
object <- initializer(object,controlEM())

#Running epigraHMM
object <- epigraHMM(object,controlEM(),type = 'consensus',dist = 'nb')

#Example
expStep(pi = c(0.99,0.02),
        gamma = matrix(c(0.99,0.01,0.01,0.99),nrow = 2),
        logf = cbind(dnbinom(rnbinom(100,mu = 2,size = 10),mu = 2,size = 10,log = TRUE),
                    dnbinom(rnbinom(100,mu = 7.5,size = 5),mu = 7.5,size = 5,log = TRUE)),
        hdf5 = file.path(tempdir(),'tmp.h5'))
```

---

helas3

*ENCODE ChIP-seq broad data from Helas3 cell line*


---

### Description

Data from EZH2, H3K27me3, and H3K36me3 ChIP-seq data from Helas3 cell line. For illustrative purposes, the data has been subset to chromosome 19. The dataset contains two replicates from each mark.

### Usage

```
data(helas3)
```

### Format

An object of class "epigraphMMDataSet".

### Source

[ENCODE Broad Histone](#)

### References

Davis et al. (2018) NAR 46(D1):D794-D801. ([PubMed](#))

### Examples

```
## # The data 'helas3' was created as follows.
# options(timeout=9999999)
#
# url <- 'http://hgdownload.soe.ucsc.edu/goldenPath/hg19/encodeDCC/wgEncodeBroadHistone/'
# samples <- c('wgEncodeBroadHistoneHelas3H3k36me3StdAlnRep1.bam',
#             'wgEncodeBroadHistoneHelas3H3k36me3StdAlnRep2.bam',
#             'wgEncodeBroadHistoneHelas3H3k27me3StdAlnRep1.bam',
#             'wgEncodeBroadHistoneHelas3H3k27me3StdAlnRep2.bam',
#             'wgEncodeBroadHistoneHelas3Ezh239875A1nRep1.bam',
#             'wgEncodeBroadHistoneHelas3Ezh239875A1nRep2.bam')
#
# input <- paste0(url,samples)
# output <- paste0(tempdir(),samples)
#
# for(idx in seq_len(length(input))){
#   download.file(url = input[idx],destfile = output[idx])
#   download.file(url = paste0(input[idx],'.bai'),
#                 destfile = paste0(output[idx],'.bai'))
# }
#
# gr <- segmentGenome(genome = 'hg19',
#                     window = 1000,rm.gap = TRUE,rm.blacklist = TRUE)
```



```
#
# cData <- data.frame(condition = rep(c('H3K36me3','H3K27me3','EZH2'),each = 2),
#                       replicate = rep(c(1,2),times = 3))
#
# subGr <- gr[seqnames(gr) == 'chr19' & start(gr) >= 40e6 & end(gr) <= 50e6]
#
# helas3 <-
#   epigraHMMDatasetFromBam(bamFiles = output,colData = cData,
#                           genome = subGr>windowSize = 1000)

data(helas3)
helas3
```

---

info

*Get information about peak calling results*

---

## Description

This function returns the BIC and expected log-likelihood function of the model, with respect to the last conditional distribution of unknown enrichment peaks given the data. The latter is also known as 'Q-function' in the EM context.

## Usage

```
info(object)
```

## Arguments

object            an epigraHMMDataset

## Value

A list with BIC, and expected log-likelihood function of the model. If the input object contains results from a differential analysis, 'info' will also output the enrichment patterns associated with each mixture component used in the mixture model.

## Author(s)

Pedro L. Baldoni, <pedrobaldoni@gmail.com>

## References

<https://github.com/plbaldoni/epigraHMM>

**Examples**

```

# Creating dummy object
countData <- rbind(matrix(rnbinom(1e3,mu = 2,size = 10),ncol = 1),
                   matrix(rnbinom(2e3,mu = 7.5,size = 5),ncol = 1),
                   matrix(rnbinom(1e3,mu = 2,size = 10),ncol = 1))

colData <- data.frame(condition = 'A', replicate = 1)
object <- epigraHMMDataSetFromMatrix(countData,colData)

# Initializing
object <- initializer(object,controlEM())

# Running epigraHMM
object <- epigraHMM(object,controlEM(),type = 'consensus',dist = 'nb')

# Get info
info(object)

```

---

 initializer

*Initializer of epigraHMM*


---

**Description**

This function call enriched windows individually for each sample in an epigraHMMDataSet. These are then used for initializing purposes in epigraHMM. By default, the Viterbi algorithm is used to determine enriched windows. Input controls and normalizing offsets are not utilized in this initialization step.

**Usage**

```
initializer(object, control)
```

**Arguments**

object	an epigraHMMDataSet
control	list of control arguments from controlEM()

**Details**

To be added

**Value**

An epigraHMMDataSet with a 'peaks' assay filled in.

**Author(s)**

Pedro L. Baldoni, <pedrobaldoni@gmail.com>

**References**

<https://github.com/plbaldoni/epigraHMM>

**Examples**

```
# Creating dummy object
countData <- rbind(matrix(rnbinom(1e3,mu = 2,size = 10),ncol = 1),
                  matrix(rnbinom(2e3,mu = 7.5,size = 5),ncol = 1),
                  matrix(rnbinom(1e3,mu = 2,size = 10),ncol = 1))

colData <- data.frame(condition = 'A', replicate = 1)
object <- epigraHMMDataSetFromMatrix(countData,colData)

# Initializing
object <- initializer(object,controlEM())

# Visualizing initialization peaks
#plot(assay(object),type = 'l')
#lines(7.5*assay(object,'peaks'),col = 'red')
```

---

maxStepProb

*M-step (maximization w.r.t. initial and transition probabilities)*


---

**Description**

M-step (maximization w.r.t. initial and transition probabilities)

**Usage**

```
maxStepProb(hdf5)
```

**Arguments**

hdf5                    path to where the hdf5 is saved

**Examples**

```
#Creating dummy object
countData <- rbind(matrix(rnbinom(1e3,mu = 2,size = 10),ncol = 1),
                  matrix(rnbinom(2e3,mu = 7.5,size = 5),ncol = 1),
                  matrix(rnbinom(1e3,mu = 2,size = 10),ncol = 1))

colData <- data.frame(condition = 'A', replicate = 1)
object <- epigraHMMDataSetFromMatrix(countData,colData)

#Initializing
```

```
object <- initializer(object,controlEM())

#Running epigraHMM
object <- epigraHMM(object,controlEM(),type = 'consensus',dist = 'nb')

#Example
maxStepProb(hdf5 = S4Vectors::metadata(object)$output)
```

---

normalizeCounts	<i>Normalize counts</i>
-----------------	-------------------------

---

## Description

This function performs a non-linear normalization of counts with respect to a reference sample (geometric mean)

## Usage

```
normalizeCounts(object, control, span = 1, ...)
```

## Arguments

object	an epigraHMMDataset
control	list of control arguments from controlEM()
span	the span parameter of <code>loessFit</code> (default is 1)
...	arguments to be passed to <code>loessFit</code> for loess calculation

## Details

This function 'limma::loessFit', which simply a wrapper for the 'stats::lowess' smoother.

## Value

An epigraHMMDataset with an 'offsets' assay filled in.

## Author(s)

Pedro L. Baldoni, <pedrobaldoni@gmail.com>

## References

<https://github.com/plbaldoni/epigraHMM>

**Examples**

```
# Creating dummy object
countData <- list('counts' = matrix(rpois(1e5,10),ncol = 2),
  'controls' = matrix(rpois(1e5,5),ncol = 2))
colData <- data.frame(condition = c('A','A'), replicate = c(1,2))
object <- epigraHMMDataSetFromMatrix(countData,colData)

# Normalizing counts
object <- normalizeCounts(object = object,control = controlEM(), span = 1)
```

---

plotCounts	<i>Create a plot with the results from epigraHMM</i>
------------	--

---

**Description**

'plotCounts()' plots read counts and peak regions from 'epigraHMM()'

**Usage**

```
plotCounts(
  object,
  ranges,
  hdf5 = metadata(object)$output,
  peaks = NULL,
  annotation = NULL
)
```

**Arguments**

object	an epigraHMMDataSet
ranges	a GRanges object or a pair of integers with the genomic coordinates/windows to be plotted
hdf5	an optional character string with the hdf5 file path from 'epigraHMM'
peaks	an optional parameter with a GRanges object or a vector of logicals (with length equal to the number of rows in 'object') specifying the genomic coordinates/windows with peaks
annotation	an optional parameter with a GRanges object or a vector of logicals (with length equal to the number of rows in 'object') specifying the genomic coordinates/windows of an annotation track

**Details**

If the input object contains the assay 'offset', reads will be normalized prior to plotting (e.g. counts/exp(offset)). Reads from replicates pertaining to the same condition are aggregated prior to plotting.

**Value**

A ggplot

**Author(s)**

Pedro L. Baldoni, <pedrobaldoni@gmail.com>

**References**

<https://github.com/plbaldoni/epigraHMM>

**Examples**

```
countData <- rbind(matrix(rnbinom(1e3,mu = 2,size = 10),ncol = 1),
                   matrix(rnbinom(1e3,mu = 7.5,size = 5),ncol = 1),
                   matrix(rnbinom(1e3,mu = 7.5,size = 5),ncol = 1),
                   matrix(rnbinom(1e3,mu = 2,size = 10),ncol = 1))

colData <- data.frame(condition = 'A', replicate = 1)

object <- epigraHMMDataSetFromMatrix(countData,colData)

plotCounts(object,ranges = c(500,3500))
```

---

plotPatterns

*Create a plot of differential patterns posterior probabilities from epigraHMM*

---

**Description**

'plotPatterns()' plots the posterior probabilities associated with differential patterns from a differential analysis of 'epigraHMM()'

**Usage**

```
plotPatterns(
  object,
  ranges,
  peaks,
  hdf5 = metadata(object)$output,
  colors = NULL
)
```

**Arguments**

object	an epigraHMMDataSet
ranges	a GRanges object or a pair of integers with the genomic coordinates/windows to be plotted
peaks	either a GRanges object or a vector of logicals (with length equal to the number of rows in 'object') specifying the genomic coordinates/windows with peaks
hdf5	a character string with the hdf5 file path from 'epigraHMM'
colors	an optional argument that specifies the colors for each differential combinatorial pattern

**Value**

A pheatmap

**Author(s)**

Pedro L. Baldoni, <pedrobaldoni@gmail.com>

**References**

<https://github.com/plbaldoni/epigraHMM>

**Examples**

```
# Creating dummy object
countData <- cbind(rbind(matrix(rnbinom(1e2, mu = 1, size = 10), ncol = 1),
  matrix(rnbinom(1e2, mu = 10, size = 5), ncol = 1),
  matrix(rnbinom(1e2, mu = 1, size = 10), ncol = 1),
  matrix(rnbinom(1e2, mu = 10, size = 5), ncol = 1),
  matrix(rnbinom(1e2, mu = 1, size = 10), ncol = 1),
  matrix(rnbinom(1e2, mu = 1, size = 10), ncol = 1),
  matrix(rnbinom(1e2, mu = 1, size = 10), ncol = 1)),
  rbind(matrix(rnbinom(1e2, mu = 1, size = 10), ncol = 1),
  matrix(rnbinom(1e2, mu = 1, size = 10), ncol = 1),
  matrix(rnbinom(1e2, mu = 1, size = 10), ncol = 1),
  matrix(rnbinom(1e2, mu = 10, size = 5), ncol = 1),
  matrix(rnbinom(1e2, mu = 1, size = 10), ncol = 1),
  matrix(rnbinom(1e2, mu = 10, size = 5), ncol = 1),
  matrix(rnbinom(1e2, mu = 1, size = 10), ncol = 1)))

colData <- data.frame(condition = c('A','B'), replicate = c(1,1))
rowRanges <- GenomicRanges::GRanges('chrA',
  IRanges::IRanges(start = seq(1,by = 500,
  length.out = nrow(countData)),width = 500))

object <- epigraHMMDataSetFromMatrix(countData,colData,rowRanges = rowRanges)

# Initializing
object <- initializer(object,controlEM())
```

```
# Running epigraHMM
object <- epigraHMM(object,controlEM(),type = 'differential',dist = 'nb')

# Calling peaks
peaks <- callPeaks(object = object,
                  hdf5 = S4Vectors::metadata(object)$output,
                  method = 'viterbi')

# Plotting patterns
plotPatterns(object,
             ranges = peaks[1],
             peaks = peaks)
```

---

segmentGenome

*Segmentation of a genome in non-overlapping windows*

---

## Description

This function segments a genome into non-overlapping windows.

## Usage

```
segmentGenome(genome, window, rm.gap = TRUE, rm.blacklist = TRUE)
```

## Arguments

genome	a string with the name of the genome (e.g. 'hg19')
window	an integer with the window size
rm.gap	a logical indicating gap regions should be removed
rm.blacklist	a logical indicating blacklisted regions should be removed

## Value

a GRanges object with the binned genome

## Author(s)

Pedro L. Baldoni, <pedrobaldoni@gmail.com>

## References

<https://github.com/plbaldoni/epigraHMM>

## Examples

```
gr <- segmentGenome(genome = 'mm10', window = 500)
```



---

```
simulateMarkovChain  Simulates a Markov Chain of length 'n' given a matrix of transition
                      probabilities P
```

---

**Description**

Simulates a Markov Chain of length 'n' given a matrix of transition probabilities P

**Usage**

```
simulateMarkovChain(P, n)
```

**Arguments**

P                    a matrix of transition probabilities (row sums should be 1)  
n                    an integer specifying the length of the simulated sequence

**Examples**

```
#Example
simulateMarkovChain(matrix(c(0.99,0.01,0.01,0.99),2,2),100)
```

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