# Package 'LACE'

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<b>Date</b> 2020-02-29
Title Longitudinal Analysis of Cancer Evolution (LACE)
Maintainer Daniele Ramazzotti <daniele.ramazzotti@yahoo.com></daniele.ramazzotti@yahoo.com>
<b>Depends</b> R (>= $3.6.0$ )
Imports parallel, Rfast, stats
Suggests BiocGenerics, BiocStyle, testthat, knitr
Name LACE: an R package for the inference of longitudinal cancer evolution models
<b>Description</b> LACE is an algorithmic framework that processes single-cell somatic mutation profiles from cancer samples collected at different time points and in distinct experimental settings, to produce longitudinal models of cancer evolution. The approach solves a Boolean Matrix Factorization problem with phylogenetic constraints, by maximizing a weighed likelihood function computed on multiple time points.
Encoding UTF-8
LazyData TRUE
License file LICENSE
<pre>URL https://github.com/BIMIB-DISCo/LACE</pre>
BugReports https://github.com/BIMIB-DISCo/LACE
biocViews BiomedicalInformatics
RoxygenNote 7.0.2
VignetteBuilder knitr
NeedsCompilation no
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2 inference

data	mutation data from Rambow, Florian, et al. "Toward minimal residual
	disease-directed therapy in melanoma." Cell 174.4 (2018): 843-855.

# Description

mutation data from Rambow, Florian, et al. "Toward minimal residual disease-directed therapy in melanoma." Cell 174.4 (2018): 843-855.

## Usage

data(data)

### **Format**

list of mutation data for four time points

#### Value

list of mutational data for a total of 475 single cells

#### **Source**

Rambow, Florian, et al. "Toward minimal residual disease-directed therapy in melanoma." Cell 174.4 (2018): 843-855.

inference	results obtained with the function LACE on the provided input data
	from Rambow, Florian, et al. "Toward minimal residual disease-
	directed therapy in melanoma." Cell 174.4 (2018): 843-855.

## Description

results obtained with the function LACE on the provided input data from Rambow, Florian, et al. "Toward minimal residual disease-directed therapy in melanoma." Cell 174.4 (2018): 843-855.

# Usage

data(inference)

## Format

results obtained with the function LACE on the provided input data

#### Value

results obtained with the function LACE on the provided input data

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## Description

Perform inference of the maximum likelihood clonal tree from longitudinal data.

# Usage

```
LACE(
  D,
  lik_w = NULL,
  alpha = NULL,
  beta = NULL,
  initialization = NULL,
  keep_equivalent = TRUE,
  num_rs = 50,
  num_iter = 10000,
  n_{try_bs} = 500,
  learning_rate = 1,
  marginalize = FALSE,
  num_processes = Inf,
  seed = NULL,
  verbose = TRUE,
  log_file = ""
)
```

## Arguments

D	Mutation data from multiple experiments for a list of driver genes.	
	Weight for each data point. If not provided, weights to correct for sample sizes are used.	
·	False positive error rate provided as list of elements; if a vector of alpha (and beta) is provided, the inference is performed for multiple values and the solution at maximum-likelihood is returned.	
	False negative error rate provided as list of elements; if a vector of beta (and alpha) is provided, the inference is performed for multiple values and the solution at maximum-likelihood is returned.	
initialization	Starting point of the mcmc; if not provided, a random starting point is used.	
keep_equivalent		
	Boolean. Shall I return results (B and C) at equivalent likelihood with the best returned solution?	
num_rs	Number of restarts during mcmc inference.	
num_iter	Maximum number of mcmc steps to be performed during the inference.	
n_try_bs	Number of steps without change in likelihood of best solution after which to	

stop the mcmc.

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learning\_rate Parameter to tune the probability of accepting solutions at lower values during

mcmc. Value of learning rate = 1 (default), set a probability proportional to the difference in likelihood; values of learning\_rate greater than 1 inclease the chance of accepting solutions at lower likelihood during mcmc while values

lower than 1 decrease such probability.

Boolean. Shall I marginalize C when computing likelihood? marginalize

Number of processes to be used during parallel execution. To execute in single num\_processes

process mode, this parameter needs to be set to either NA or NULL.

seed Seed for reproducibility.

Boolean. Shall I print to screen information messages during the execution? verbose log\_file

log file where to print outputs when using parallel. If parallel execution is dis-

abled, this parameter is ignored.

#### Value

A list of 8 elements: B, C, clones prevalence, relative likelihoods, joint likelihood, clones summary and error rates. Here, B returns the maximum likelihood longitudinal clonal tree, C the attachment of cells to clones and clones\_prevalence clones' prevalence; relative\_likelihoods and joint\_likelihood are respectively the likelihood of the solutions at each individual time points and the joint likelihood; clones summary provide a summary of association of mutations to clones. In equivalent solutions, solutions (B and C) with likelihood equivalent to the best solution are returned. Finally error\_rates provides the best values of alpha and beta among the considered ones.

### **Examples**

```
data(data)
inference = LACE(D = data,
                  lik_w = c(0.2308772, 0.2554386, 0.2701754, 0.2435088),
                  alpha = list(c(0.10, 0.05, 0.05, 0.05)),
                  beta = list(c(0.10, 0.05, 0.05, 0.05)),
                  keep_equivalent = FALSE,
                  num_rs = 5,
                  num_iter = 10,
                  n_{try_bs} = 5,
                  num_processes = NA,
                  seed = 12345,
                  verbose = FALSE)
```

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