

Package ‘mbmixture’

October 22, 2020

Version 0.2-5

Date 2020-10-13

Title Microbiome Mixture Analysis

Description Evaluate whether a microbiome sample is a mixture of two samples, by fitting a model for the number of read counts as a function of single nucleotide polymorphism (SNP) allele and the genotypes of two potential source samples.
Lobo et al. (2019) <doi:10.1101/529040>.

Author Karl W Broman [aut, cre] (<<https://orcid.org/0000-0002-4914-6671>>)

Maintainer Karl W Broman <broman@wisc.edu>

Depends R (>= 3.1.0)

Imports stats, parallel, numDeriv

Suggests knitr, rmarkdown, testthat, devtools, roxygen2

License MIT + file LICENSE

URL <https://github.com/kbroman/mbmixture>

BugReports <https://github.com/kbroman/mbmixture/issues>

VignetteBuilder knitr

LazyData true

Encoding UTF-8

ByteCompile true

RoxygenNote 7.1.1

NeedsCompilation no

Repository CRAN

Date/Publication 2020-10-22 09:10:05 UTC

R topics documented:

bootstrapNull	2
bootstrapSE	3
mbmixdata	4
mbmix_loglik	5
mle_e	5
mle_p	6
mle_pe	7
Index	8

bootstrapNull	<i>Bootstrap to assess significance</i>
---------------	---

Description

Perform a parametric bootstrap to assess whether there is significant evidence that a sample is a mixture.

Usage

```
bootstrapNull(
  tab,
  n_rep = 1000,
  interval = c(0, 1),
  tol = 0.000001,
  check_boundary = TRUE,
  cores = 1,
  return_raw = TRUE
)
```

Arguments

tab	Dataset of read counts as 3d array of size 3x3x2, genotype in first sample x genotype in second sample x allele in read.
n_rep	Number of bootstrap replicates
interval	Interval to which each parameter should be constrained
tol	Tolerance for convergence
check_boundary	If TRUE, explicitly check the boundaries of interval.
cores	Number of CPU cores to use, for parallel calculations. (If 0, use <code>parallel::detectCores()</code> .) Alternatively, this can be links to a set of cluster sockets, as produced by <code>parallel::makeCluster()</code> .
return_raw	If TRUE, return the raw results. If FALSE, just return the p-value. Unlink <code>bootstrapSE()</code> , here the default is TRUE.

Value

If return_raw=FALSE, a single numeric value (the p-value). If return_raw=TRUE, a vector of length n_rep with the LRT statistics from each bootstrap replicate.

See Also

[bootstrapSE\(\)](#)

Examples

```
data(mbmixdata)
# just 100 bootstrap replicates, as an illustration
bootstrapNull(mbmixdata, n_rep=100)
```

bootstrapSE	<i>Bootstrap to get standard errors</i>
-------------	---

Description

Perform a parametric bootstrap to get estimated standard errors.

Usage

```
bootstrapSE(
  tab,
  n_rep = 1000,
  interval = c(0, 1),
  tol = 0.000001,
  check_boundary = FALSE,
  cores = 1,
  return_raw = FALSE
)
```

Arguments

tab	Dataset of read counts as 3d array of size 3x3x2, genotype in first sample x genotype in second sample x allele in read.
n_rep	Number of bootstrap replicates
interval	Interval to which each parameter should be constrained
tol	Tolerance for convergence
check_boundary	If TRUE, explicitly check the boundaries of interval.
cores	Number of CPU cores to use, for parallel calculations. (If 0, use parallel::detectCores() .) Alternatively, this can be links to a set of cluster sockets, as produced by parallel::makeCluster() .
return_raw	If TRUE, return the raw results. If FALSE, just return the estimated standard errors.

Value

If return_raw=FALSE, a vector of two standard errors. If return_raw=TRUE, an matrix of size n_rep x 2 with the detailed bootstrap results.

See Also

[bootstrapNull\(\)](#)

Examples

```
data(mbmixdata)
# just 100 bootstrap replicates, as an illustration
bootstrapSE(mbmixdata, n_rep=100)
```

mbmixdata

Example dataset for mbmixture package

Description

Example dataset for mbmixture package.

Usage

```
data(mbmixdata)
```

Format

Dataset of read counts as 3d array of size 3x3x2, genotype in first sample x genotype in second sample x allele in read.

Examples

```
data(mbmixdata)
mle_pe(mbmixdata)
```

mbmix_loglik	<i>log likelihood function for microbiome mixture</i>
--------------	---

Description

Calculate log likelihood function for microbiome sample mixture model at particular values of p and e .

Usage

```
mbmix_loglik(tab, p, e = 0)
```

Arguments

tab	Dataset of read counts as 3d array of size $3 \times 3 \times 2$, genotype in first sample x genotype in second sample x allele in read.
p	Contaminant probability (proportion of mixture coming from the second sample).
e	Sequencing error rate.

Value

The log likelihood evaluated at p and e .

Examples

```
data(mbmixdata)
mbmix_loglik(mbmixdata, p=0.74, e=0.002)
```

mle_e	<i>MLE of e for fixed p</i>
-------	-----------------------------

Description

Calculate the MLE of the sequencing error rate e for a fixed value of the contaminant probability p .

Usage

```
mle_e(
  tab,
  p = 0.05,
  interval = c(0, 1),
  tol = 0.000001,
  check_boundary = FALSE
)
```

Arguments

tab	Dataset of read counts as 3d array of size 3x3x2, genotype in first sample x genotype in second sample x allele in read.
p	Assumed value for the contaminant probability
interval	Interval to which each parameter should be constrained
tol	Tolerance for convergence
check_boundary	If TRUE, explicitly check the boundaries of interval.

Value

A single numeric value, the MLE of e , with the log likelihood as an attribute.

Examples

```
data(mbmixdata)
mle_e(mbmixdata, p=0.74)
```

mle_p

MLE of p for fixed e

Description

Calculate the MLE of the contaminant probability p for a fixed value of the sequencing error rate e .

Usage

```
mle_p(
  tab,
  e = 0.002,
  interval = c(0, 1),
  tol = 0.000001,
  check_boundary = FALSE
)
```

Arguments

tab	Dataset of read counts as 3d array of size 3x3x2, genotype in first sample x genotype in second sample x allele in read.
e	Assumed value for the sequencing error rate
interval	Interval to which each parameter should be constrained
tol	Tolerance for convergence
check_boundary	If TRUE, explicitly check the boundaries of interval.

Value

A single numeric value, the MLE of p , with the log likelihood as an attribute.

Examples

```
data(mbmixdata)
mle_p(mbmixdata, e=0.002)
```

mle_pe

Find MLEs for microbiome mixture

Description

Find joint MLEs of p and e for microbiome mixture model

Usage

```
mle_pe(
  tab,
  interval = c(0, 1),
  tol = 0.000001,
  check_boundary = FALSE,
  SE = FALSE
)
```

Arguments

tab	Dataset of read counts as 3d array of size $3 \times 3 \times 2$, genotype in first sample x genotype in second sample x allele in read.
interval	Interval to which each parameter should be constrained
tol	Tolerance for convergence
check_boundary	If TRUE, explicitly check the boundaries of interval.
SE	If TRUE, get estimated standard errors.

Value

A vector containing the estimates of p and e along with the evaluated log likelihood and likelihood ratio test statistics for the hypotheses $p=0$ and $p=1$.

Examples

```
data(mbmixdata)
mle_pe(mbmixdata)
```

Index

* datasets

- mbmixdata, 4

- bootstrapNull, 2
- bootstrapNull(), 4
- bootstrapSE, 3
- bootstrapSE(), 2, 3

- mbmix_loglik, 5
- mbmixdata, 4
- mle_e, 5
- mle_p, 6
- mle_pe, 7

- parallel::detectCores(), 2, 3
- parallel::makeCluster(), 2, 3