

# Package: mbmixture (via r-universe)

November 28, 2024

**Version** 0.6

**Date** 2024-11-27

**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. (2021) [doi:10.1093/g3journal/jkab308](https://doi.org/10.1093/g3journal/jkab308).

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**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** 7.2.3

**NeedsCompilation** no

**Repository** CRAN

**Date/Publication** 2024-11-27 23:20:02 UTC

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

*Bootstrap to get standard errors*

---

**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 <a href="#">parallel::detectCores()</a> .) Alternatively, this can be links to a set of cluster sockets, as produced by <a href="#">parallel::makeCluster()</a> .
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>
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**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)
```

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mle_e	<i>MLE of e for fixed p</i>
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**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)
```

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