

# Package ‘rhierbaps’

November 18, 2022

**Type** Package

**Title** Clustering Genetic Sequence Data Using the HierBAPS Algorithm

**Version** 1.1.4

**Description** Implements the hierarchical Bayesian analysis of populations structure (hierBAPS) algorithm of Cheng et al. (2013) <[doi:10.1093/molbev/mst028](https://doi.org/10.1093/molbev/mst028)> for clustering DNA sequences from multiple sequence alignments in FASTA format. The implementation includes improved defaults and plotting capabilities and unlike the original 'MATLAB' version removes singleton SNPs by default.

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**Encoding** UTF-8

**Imports** ape, purrr, utils, ggplot2, matrixStats, patchwork, methods

**RoxygenNote** 7.2.1

**Suggests** knitr, rmarkdown, ggtree, phytools, testthat, formatR

**VignetteBuilder** knitr

**URL** <https://github.com/gtonkinhill/rhierbaps>

**BugReports** <https://github.com/gtonkinhill/rhierbaps/issues>

**NeedsCompilation** no

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**Repository** CRAN

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`calc_change_in_ml`      *calc\_change\_in\_ml*

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**Description**

Calculate the change in the log marginal likelihood after moving index to each possible cluster

**Usage**

```
calc_change_in_ml(snp.object, partition, indexes)
```

**Arguments**

- |            |                                                                   |
|------------|-------------------------------------------------------------------|
| snp.object | A snp.object containing the processed SNP data.                   |
| partition  | An integer vector indicating a partition of the isolates.         |
| indexes    | Indexes of the isolates to be moved (must come from one cluster.) |

**Value**

the best cluster to move indexes to.

---

`calc_log_ml`      *calc\_log\_ml*

---

**Description**

Calculate the log marginal likelihood assuming a Multinomial-Dirichlet distribution

**Usage**

```
calc_log_ml(snp.object, partition)
```

**Arguments**

- |            |                                                           |
|------------|-----------------------------------------------------------|
| snp.object | A snp.object containing the processed SNP data.           |
| partition  | An integer vector indicating a partition of the isolates. |

**Value**

The log marginal likelihood of the given partition.

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**hierBAPS***hierBAPS*

---

**Description**

Runs the hierBAPS algorithm of Cheng et al. 2013

**Usage**

```
hierBAPS(  
  snp.matrix,  
  max.depth = 2,  
  n.pops = floor(nrow(snp.matrix)/5),  
  quiet = FALSE,  
  n.extra.rounds = 0,  
  assignment.probs = FALSE,  
  n.cores = 1  
)
```

**Arguments**

snp.matrix	Character matrix of aligned sequences produced by <a href="#">load_fasta</a> .
max.depth	Maximum depth of hierarchical search (default = 2).
n.pops	Maximum number of populations in the data (default = number of isolates/5)
quiet	Whether to suppress progress information (default=FALSE).
n.extra.rounds	The number of additional rounds to perform after the default hierBAPS settings (default=0). If set to Inf it will run until a local optimum is reached (this might take a long time).
assignment.probs	whether or not to calculate the assignment probabilities to each cluster (default=FALSE)
n.cores	The number of cores to use.

**Value**

A list containing a dataframe indicating an assignment of each sequence to hierarchical clusters as well as the log marginal likelihoods for each level.

**Author(s)**

Gerry Tonkin-Hill

## References

Cheng, Lu, Thomas R. Connor, Jukka Sirén, David M. Aanensen, and Jukka Corander. 2013. “Hierarchical and Spatially Explicit Clustering of DNA Sequences with BAPS Software.” Molecular Biology and Evolution 30 (5): 1224–28.

## Examples

```
snp.matrix <- load_fasta(system.file("extdata", "small_seqs.fa", package = "rhierbaps"))
hb <- hierBAPS(snp.matrix, max.depth=2, n.pops=20, quiet=FALSE)
```

```
snp.matrix <- load_fasta(system.file("extdata", "seqs.fa", package = "rhierbaps"))
system.time({hb <- hierBAPS(snp.matrix, max.depth=2, n.pops=20, quiet=FALSE)})
```

*join\_units\_2*

*join\_units\_2*

## Description

Perform an iteration of the second move in the algorithm. That is combine two clusters to improve the marginal likelihood.

## Usage

```
join_units_2(
  snp.object,
  partition,
  threshold = 1e-05,
  n.cores = 1,
  comb.chache = NULL
)
```

## Arguments

<code>snp.object</code>	A <code>snp.object</code> containing the processed SNP data.
<code>partition</code>	An integer vector indicating an initial partition of the isolates.
<code>threshold</code>	The increase in marginal log likelihood required to accept a move.
<code>n.cores</code>	The number of cores to use.
<code>comb.chache</code>	a matrix recording previous marginal llks of combining clusters

## Value

The best partition after combining two clusters as well as a boolean value indicating whether a move increased the marginal likelihood.

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<code>load_fasta</code>	<i>load_fasta</i>
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**Description**

Loads a fasta file into matrix format ready for running the hierBAPS algorithm.

**Usage**

```
load_fasta(msa, keep.singletons = FALSE)
```

**Arguments**

- |                              |                                                                                                                          |
|------------------------------|--------------------------------------------------------------------------------------------------------------------------|
| <code>msa</code>             | Either the location of a fasta file or ape DNAbin object containing the multiple sequence alignment data to be clustered |
| <code>keep.singletons</code> | A logical indicating whether to consider singleton mutations in calculating the clusters                                 |

**Value**

A character matrix with filtered SNP data

**Examples**

```
msa <- system.file("extdata", "seqs.fa", package = "rhierbaps")
snp.matrix <- load_fasta(msa)
```

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<code>log_stirling2</code>	<i>log_stirling2</i>
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**Description**

`log_stirling2`

**Usage**

```
log_stirling2(n, k)
```

**Arguments**

- |                |                      |
|----------------|----------------------|
| <code>n</code> | number of objects    |
| <code>k</code> | number of partitions |

**Value**

log of the Stirling number of the second kind

`model_search_parallel` *model\_search\_parallel*

### Description

Clusters DNA alignment using independent loci model

### Usage

```
model_search_parallel(
  snp.object,
  partition,
  round.types,
  quiet = FALSE,
  n.extra.rounds = 0,
  n.cores = 1
)
```

### Arguments

<code>snp.object</code>	A <code>snp.object</code> containing the processed SNP data.
<code>partition</code>	An integer vector indicating an initial starting partition.
<code>round.types</code>	A vector indicating which series of moves to make.
<code>quiet</code>	Whether to suppress progress information (default=FALSE).
<code>n.extra.rounds</code>	The number of additional rounds to perform after the default hierBAPS settings (default=0). If set to Inf it will run until a local optimum is reached (this might take a long time).
<code>n.cores</code>	The number of cores to use.

### Value

an optimised partition and marginal llk

`move_units_1`

*move\_units\_1*

### Description

Perform an iteration of the first move in the algorithm. That is move units from one cluster to another to improve the marginal likelihood

**Usage**

```
move_units_1(  
  snp.object,  
  partition,  
  threshold = 1e-05,  
  frac.clust.searched = 0.3,  
  min.clust.size = 20,  
  n.cores = 1  
)
```

**Arguments**

snp.object	A snp.object containing the processed SNP data.
partition	An integer vector indicating an initial partition of the isolates.
threshold	The increase in marginal log likelihood required to accept a move.
frac.clust.searched	The percentage of a large cluster that will be moved.
min.clust.size	All isolates in clusters less than or equal to min.clus.size will be searched.
n.cores	The number of cores to use.

**Value**

The best partition after moving units from one cluster to another as well as a boolean value indicating whether a move increased the marginal likelihood.

---

*plot\_sub\_cluster*      *plot\_sub\_cluster*

---

**Description**

Creates a zoom plot using ggtree focusing on a cluster.

**Usage**

```
plot_sub_cluster(hb.object, tree, level, sub.cluster)
```

**Arguments**

hb.object	The resulting object from running hierBAPS
tree	A phylo tree object to plot
level	The level of the subcluster to be considered.
sub.cluster	An integer representing the subcluster to be considered.

## Examples

```
snp.matrix <- load_fasta(system.file("extdata", "seqs.fa", package = "rhierbaps"))
newick.file.name <- system.file("extdata", "seqs.fa.treefile", package = "rhierbaps")
tree <- phytools::read.newick(newick.file.name)
hb.result <- hierBAPS(snp.matrix, max.depth=2, n.pop=20)
plot_sub_cluster(hb.result, tree, level = 1, sub.cluster = 9)
```

**preproc\_alignment**      *preproc\_alignment*

## Description

Preprocessed the snp matrix for hierBAPS.

## Usage

```
preproc_alignment(snp.matrix)
```

## Arguments

**snp.matrix**      A matrix containing SNP data. Rows indicate isolates and columns loci.

## Value

an snp.object

**reallocates\_units\_4**      *reallocates\_units\_4*

## Description

Perform an iteration of the fourth move in the algorithm. That is split cluster into n subclusters and re-allocate one sub-cluster.

## Usage

```
reallocates_units_4(
  snp.object,
  partition,
  threshold = 1e-05,
  min.clust.size = 20,
  split = FALSE,
  n.cores = 1
)
```

**Arguments**

snp.object	A.snp.object containing the processed SNP data.
partition	An integer vector indicating an initial partition of the isolates.
threshold	The increase in marginal log likelihood required to accept a move.
min.clust.size	Clusters smaller than min.clust.size will not be split.
split	Whether to split only into two clusters (for move type 3).
n.cores	The number of cores to use.

**Value**

The best partition after splitting a cluster and re-allocating as well as a boolean value indicating whether a move increased the marginal likelihood.

save_lml_logs	save_lml_logs
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**Description**

Saves the log marginal likelihoods to a text file.

**Usage**

```
save_lml_logs(hb.object, file)
```

**Arguments**

hb.object	The resulting object from running hierBAPS
file	The file you would like to save the log output to.

**Examples**

```
snp.matrix <- load_fasta(system.file("extdata", "small_seqs.fa", package = "rhierbaps"))
hb.result <- hierBAPS(snp.matrix, max.depth=2, n.pops=20)
save_lml_logs(hb.result, file.path(tempdir(), "output_file.txt"))
```

---

split\_clusters\_3      *split\_clusters\_3*

---

### Description

Peform an iteration of the third move in the algorithm. That is split cluster in two and re-allocate one sub-cluster.

### Usage

```
split_clusters_3(  
  snp.object,  
  partition,  
  threshold = 1e-05,  
  min.clust.size = 20,  
  n.cores = 1  
)
```

### Arguments

snp.object	A snp.object containing the processed SNP data.
partition	An integer vector indicating an initial partition of the isolates.
threshold	The increase in marginal log likelihood required to accept a move.
min.clust.size	Clusters smaller than min.clust.size will not be split.
n.cores	The number of cores to use.

### Value

The best partition after splitting a cluster and re-allocating as well as a boolean value indicating whether a move increased the marginal likelihood.

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