

Package ‘LEA’

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Title LEA: an R package for Landscape and Ecological Association Studies

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Depends R (>= 3.0.2), methods, stats, utils

Description LEA is an R package dedicated to landscape genomics and ecological association tests. LEA can run analyses of population structure and genome scans for local adaptation. It includes statistical methods for estimating ancestry coefficients from large genotypic matrices and evaluating the number of ancestral populations (snmf, pca); and identifying genetic polymorphisms that exhibit high correlation with some environmental gradient or with the variables used as proxies for ecological pressures (lfmm), and controlling the false discovery rate. LEA is mainly based on optimized C programs that can scale with the dimension of very large data sets.

License GPL-3

biocViews Software, StatisticalMethod, Clustering, Regression

URL <http://membres-timc.imag.fr/Olivier.Francois/lea.html>

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R topics documented:

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LEA-package	<i>LEA: an R package for Landscape and Ecological Associations studies.</i>
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Description

LEA is an R package dedicated to landscape genomics and ecological association tests. LEA can run analyses of population structure and genome scans for local adaptation. It includes statistical methods for estimating ancestry coefficients from large genotypic matrices and evaluating the number of ancestral populations (snmf, pca); and identifying genetic polymorphisms that exhibit high correlation with some environmental gradient or with the variables used as proxies for ecological pressures (lfmm). LEA is mainly based on optimized C programs that can scale with the dimension of very large data sets.

Details

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Author(s)

Eric Frichot Maintainer: Eric Frichot <eric.frichot@imag.fr>

adjusted.pvalues *adjusted p-values from a lfmm run*

Description

Return the lfmm output vector of adjusted p-values and the genomic inflation factor using the genomic control method or the lambda inflation factor parameter for the chosen runs with K latent factors, the d-th variable and the all option. For an example, see [lfmm](#).

Usage

```
adjusted.pvalues (object, genomic.control, lambda, K, d, all, run)
```

Arguments

object	A lfmmProject object.
genomic.control	A boolean option. If true, the p-values are automatically calibrated using the genomic control method. If false, the p-values are calculated using the lambda inflation factor parameter.
lambda	the lambda inflation factor used to calibrate the p-value if genomic.control = FALSE (default: 1.0).
K	The number of latent factors.
d	The d-th variable.
all	A Boolean option. If true, the run with all variables at the same time. If false, the runs with each variable separately.
run	A list of chosen runs.

Value

res A matrix containing a vector of p.values for the chosen runs per column.

Author(s)

Eric Frichot

See Also

[lfmm.data](#) [lfmm p.values](#) [mlog10p.values](#)

Examples

```
### Example of analyses using lfmm ###

data("tutorial")
# creation of the genotype file, genotypes.lfmm.
# It contains 400 SNPs for 50 individuals.
write.lfmm(tutorial.R, "genotypes.lfmm")
# creation of the environment file, gradient.env.
# It contains 1 environmental variable for 40 individuals.
write.env(tutorial.C, "gradients.env")

#####
# runs of lfmm #
#####

# main options, K: (the number of latent factors),
#           CPU: the number of CPUs.

# Toy runs with K = 3 and 2 repetitions.
# around 15 seconds per run.
project = NULL
project = lfmm("genotypes.lfmm", "gradients.env", K = 3, repetitions = 2,
              iterations = 6000, burnin = 3000, project = "new")

# get the adjusted p-values using the genomic control method
res = adjusted.pvalues(project, K = 3)

hist(res$p.values, col = "yellow3")

# get the adjusted p-values with the genomic inflation factor
res = adjusted.pvalues(project, genomic.control = FALSE,
                      lambda = res$genomic.inflation.factor, K = 3)

hist(res$p.values, col = "yellow3")
```

ancestrymap

ancestrymap *format description*

Description

Description of the ancestrymap format. The ancestrymap format can be used as an input format for genotypic matrices in the functions [pca](#), [lfmm](#) and [snmf](#).

Details

The ancestrymap format has one row for each genotype. Each row has 3 columns: the 1st column is the SNP name, the 2nd column is the sample ID, the 3rd column is the number of alleles. Genotypes for a given SNP name are written in consecutive lines. The number of alleles can be the number of reference alleles or the number of derived alleles. Missing genotypes are encoded by the value 9.

Here is an example of a genotypic matrix using the ancestrymap format with 3 individuals and 4 SNPs:

```

rs0000    SAMPLE0    1
rs0000    SAMPLE1    1
rs0000    SAMPLE2    2
rs1111    SAMPLE0    0
rs1111    SAMPLE1    1
rs1111    SAMPLE2    0
rs2222    SAMPLE0    0
rs2222    SAMPLE1    9
rs2222    SAMPLE2    1
rs3333    SAMPLE0    1
rs3333    SAMPLE1    2
rs3333    SAMPLE2    1

```

Author(s)

Eric Fritchot

See Also

[ancestrymap2lfmm](#) [ancestrymap2geno](#) [geno](#) [lfmm](#).data [ped](#) [vcf](#)

ancestrymap2geno

*Convert from [ancestrymap](#) to [geno](#) format***Description**

A function that converts from the [ancestrymap](#) format to the [geno](#) format.

Usage

```
ancestrymap2geno(input.file, output.file = NULL, force = TRUE)
```

Arguments

<code>input.file</code>	A character string containing a path to the input file, a genotypic matrix in the ancestrymap format.
<code>output.file</code>	A character string containing a path to the output file, a genotypic matrix in the geno format. By default, the name of the output file is the same name as the input file with a <code>.geno</code> extension.
<code>force</code>	A boolean option. If <code>FALSE</code> , the input file is converted only if the output file does not exist. If <code>TRUE</code> , convert the file anyway.

Value

<code>output.file</code>	A character string containing a path to the output file, a genotypic matrix in the geno format.
--------------------------	-----------------------------------------------------------------------------------------------------------------

Author(s)

Eric Fritchot

See Also

[ancestrymap](#) [geno](#) [read.geno](#) [ancestrymap2lfmm](#) [geno2lfmm](#) [ped2lfmm](#) [ped2geno](#) [vcf2geno](#)
[lfmm2geno](#)

Examples

```
# Creation of of file called "example.ancestrymap"
# a file containing 4 SNPs for 3 individuals.
data("example_ancestrymap")
write.table(example_ancestrymap,"example.ancestrymap",
col.names = FALSE, row.names = FALSE, quote = FALSE)

# Conversion from the ancestrymap format ("example.ancestrymap")
# to the geno format ("example.geno").
# By default, the name of the output file is the same name
# as the input file with a .geno extension.
# Create file: "example.geno".
output = ancestrymap2geno("example.ancestrymap")

# Conversion from the ancestrymap format (example.ancestrymap)
# to the geno format with the output file called plop.geno.
# Create file: "plop.geno".
output = ancestrymap2geno("example.ancestrymap", "plop.geno")

# As force = false and the file "example.geno" already exists,
# nothing happens.
output = ancestrymap2geno("example.ancestrymap", force = FALSE)
```

ancestrymap2lfmm *Convert from ancestrymap to lfmm format*

Description

A function that converts from the [ancestrymap](#) format to the [lfmm](#) format.

Usage

```
ancestrymap2lfmm(input.file, output.file = NULL, force = TRUE)
```

Arguments

<code>input.file</code>	A character string containing a path to the input file, a genotypic matrix in the ancestrymap format.
<code>output.file</code>	A character string containing a path to the output file, a genotypic matrix in the lfmm format. By default, the name of the output file is the same name as the input file with a .lfmm extension.
<code>force</code>	A boolean option. If FALSE, the input file is converted only if the output file does not exist. If TRUE, convert the file anyway.

Value

output.file A character string containing a path to the output file, a genotypic matrix in the [lfmm](#) format.

Author(s)

Eric Frichot

See Also

[ancestrymap lfmm.data](#) [ancestrymap2geno](#) [geno2lfmm](#) [ped2lfmm](#) [ped2geno](#) [vcf2geno](#) [lfmm2geno](#)

Examples

```
# Creation of a file called "example.ancestrymap"
# containing 4 SNPs for 3 individuals.
data("example_ancestrymap")
write.table(example_ancestrymap,"example.ancestrymap",
col.names = FALSE, row.names = FALSE, quote = FALSE)

# Conversion      from the ancestrymap format ("example.ancestrymap")
#                    to the lfmm format ("example.lfmm").
# By default,      the name of the output file is the same name
#                    as the input file with a .lfmm extension.
# Create file:      "example.lfmm".
output = ancestrymap2lfmm("example.ancestrymap")

# Conversion      from the ancestrymap format (example.ancestrymap)
#                    to the geno format with the output file called plop.lfmm.
# Create file:      "plop.lfmm".
output = ancestrymap2lfmm("example.ancestrymap", "plop.lfmm")

# As force = false and the file "example.lfmm" already exists,
# nothing happens.
output = ancestrymap2lfmm("example.ancestrymap", force = FALSE)
```

create.dataset

create a data set with masked data

Description

[create.dataset](#) creates a data set with a given percentage of masked data from the original data set. It is used to calculate the [cross.entropy](#) criterion.

Usage

```
create.dataset (input.file, output.file, seed = -1, percentage = 0.05)
```

Arguments

input.file	A character string containing a path to the input file, a genotypic matrix in the geno format.
output.file	A character string containing a path to the output file, a genotypic matrix in the geno format. The output file is the input file with masked genotypes. By default, the name of the output file is the same name as the input file with a <code>_I.geno</code> extension.
seed	A seed to initialize the random number generator. By default, the seed is randomly chosen.
percentage	A numeric value between 0 and 1 containing the percentage of masked genotypes.

Details

This is an internal function, automatically called by [snmf](#) with the entropy option.

Value

output.file	A character string containing a path to the output file, a genotypic matrix in the geno format.
-------------	-----------------------------------------------------------------------------------------------------------------

Author(s)

Eric Frichot

See Also

[geno snmf cross.entropy](#)

Examples

```
# Creation of tuto.geno
# A file containing 400 SNPs for 50 individuals.
data("tutorial")
write.geno(tutorial.R,"genotypes.geno")

# Creation of the masked data file
# Create file: "genotypes_I.geno"
output = create.dataset("genotypes.geno")
```

cross.entropy

Cross-entropy criterion from snmf runs

Description

Return the cross-entropy criterion for the chosen runs with K ancestral populations. For an example, see [snmf](#). The cross-entropy criterion is a value based on the prediction of masked genotypes to evaluate the error of ancestry estimation. The criterion will help to choose the best number of ancestral population (K) and the best run among a set of runs in [snmf](#). A smaller value of cross-entropy means a better run in terms of prediction capacity. The cross-entropy criterion can be automatically calculated by the [snmf](#) function with the entropy option.

Usage

```
cross.entropy(object, K, run)
```

Arguments

object	A snmfProject object.
K	The number of ancestral populations.
run	A list of chosen run number.

Value

res	A list containing the cross-entropy criterion for the chosen runs with K ancestral populations.
-----	-------------------------------------------------------------------------------------------------

Author(s)

Eric Frichot

See Also

[geno snmf G Q](#)

Examples

```
### Example of analyses using snmf ###

# creation of the genotype file, genotypes.geno.
# It contains 400 SNPs for 50 individuals.
data("tutorial")
write.geno(tutorial.R, "genotypes.geno")

#####
# runs of snmf #
#####

# main options, K: (the number of ancestral populations),
# entropy: calculate the cross-entropy criterion,
# CPU: the number of CPUs.

# Runs with K = 3 with cross-entropy and 2 repetitions.
project = NULL
project = snmf("genotypes.geno", K = 3, entropy = TRUE, repetitions = 2,
              project = "new")

# get the cross-entropy for all runs for K = 3
ce = cross.entropy(project, K = 3)

# get the cross-entropy for the 2nd run for K = 3
ce = cross.entropy(project, K = 3, run = 2)
```

cross.entropy. estimation

compute the cross-entropy criterion

Description

Calculate the cross-entropy criterion. This is an internal function, automatically called by `snmf`. The cross-entropy criterion is a value based on the prediction of masked genotypes to evaluate the error of ancestry estimation. The criterion will help to choose the best number of ancestral population (K) and the best run among a set of runs in `snmf`. A smaller value of cross-entropy means a better run in terms of prediction capacity. The `cross.entropy. estimation` function displays the cross-entropy criterion estimated on all data and on masked data based on the input file, the masked data file (created by `create. dataset`, the estimation of the ancestry coefficients Q and the estimation of ancestral genotypic frequencies, G (calculated by `snmf`). The cross-entropy estimation for all data is always lower than the cross-entropy estimation for masked data. The cross-entropy estimation useful to compare runs is the cross-entropy estimation for masked data. The cross-entropy criterion can also be automatically calculated by the `snmf` function with the entropy option.

Usage

```
cross.entropy. estimation (input. file, K, masked. file, Q. file, G. file,
                          ploidy = 2)
```

Arguments

<code>input. file</code>	A character string containing a path to the input file without masked genotypes, a genotypic matrix in the <code>geno</code> format.
<code>K</code>	An integer corresponding to the number of ancestral populations.
<code>masked. file</code>	A character string containing a path to the input file with masked genotypes, a genotypic matrix in the <code>geno</code> format. This file can be generated with the function, <code>create. dataset</code>). By default, the name of the masked data file is the same name as the input file with a <code>_I. geno</code> extension.
<code>Q. file</code>	A character string containing a path to the input ancestry coefficient matrix Q. By default, the name of this file is the same name as the input file with a <code>K. Q</code> extension.
<code>G. file</code>	A character string containing a path to the input ancestral genotype frequency matrix G. By default, the name of this file is the same name as the input file with a <code>K. G</code> extension (<code>input_ file. K. G</code>).
<code>ploidy</code>	1 if haploid, 2 if diploid, n if n-ploid.

Value

`cross.entropy. estimation` returns a list containing the following components:

<code>masked. ce</code>	The value of the cross-entropy criterion of the masked genotypes.
<code>all. ce</code>	The value of the cross-entropy criterion of all the genotypes.

Author(s)

Eric Frichot

References

Frichot E, Mathieu F, Trouillon T, Bouchard G, Francois O. (2014). *Fast and Efficient Estimation of Individual Ancestry Coefficients*. *Genetics*, 194(4) : 973–983.

See Also

[geno create.dataset snmf](#)

Examples

```
# Creation of tuto.geno
# A file containing 400 SNPs for 50 individuals.
data("tutorial")
write.geno(tutorial.R,"genotypes.geno")

# The following command are equivalent with
# project = snmf("genotypes.geno", entropy = TRUE, K = 3)
# cross.entropy(project)

# Creation of the masked data file
# Create file: "genotypes_I.geno"
output = create.dataset("genotypes.geno")

# run of snmf with genotypes_I.geno and K = 3
project = snmf("genotypes_I.geno", K = 3, project = "new")

# calculate the cross-entropy
res = cross.entropy. estimation("genotypes.geno", K = 3, "genotypes_I.geno",
    "./genotypes_I.snmf/K3/run1/genotypes_I_r1.3.Q",
    "./genotypes_I.snmf/K3/run1/genotypes_I_r1.3.G")

# get the result
res$masked.ce
res$all.ce

#remove project
remove.snmfProject("genotypes_I.snmfProject")
```

env

Environmental input file format for [lfmm](#)

Description

Description of the env format. The env format can be used as an input format for the environmental variables in the [lfmm](#) function.

Details

The env format has one row for each individual. Each row contains one value for each environmental variable (separated by spaces or tabulations).

Here is an example of an environmental file using the env format with 3 individuals and 2 variable:

```
0.252477 0.95250639
0.216618 0.10902647
-0.47509 0.07626694
```

Author(s)

Eric Frichot

See Also

[lfmm read.env write.env](#)

G

Ancestral allele frequencies from a snmf run

Description

Return the snmf output matrix of ancestral allele frequency matrix for the chosen run with K ancestral populations. For an example, see [snmf](#).

Usage

G(object, K, run)

Arguments

object	A snmfProject object.
K	The number of ancestral populations.
run	A chosen run.

Value

res	A matrix containing the ancestral allele frequencies for the chosen run with K ancestral populations.
-----	-------------------------------------------------------------------------------------------------------

Author(s)

Eric Frichot

See Also

[geno snmf Q cross.entropy](#)

Examples

```
### Example of analyses using snmf ###

# creation of the genotype file, genotypes.geno.
# It contains 400 SNPs for 50 individuals.
data("tutorial")
write.geno(tutorial.R, "genotypes.geno")

#####
# runs of snmf #
#####

# main options, K: (the number of ancestral populations),
#     entropy: calculate the cross-entropy criterion,
#     CPU: the number of CPUs.

# Runs with K between 1 and 5 with cross-entropy and 2 repetitions.
project = NULL
project = snmf("genotypes.geno", K = 3, repetitions = 2, project = "new")

# get the ancestral genotype frequency matrix, G, for the 2nd run for K = 3.
res = G(project, K = 3, run = 2)
```

geno

Input file for snmf

Description

Description of the geno format. The geno format can be used as an input format for genotypic matrices in the functions [snmf](#), [lfmm](#), and [pca](#).

Details

The geno format has one row for each SNP. Each row contains 1 character for each individual: 0 means zero copy of the reference allele. 1 means one copy of the reference allele. 2 means two copies of the reference allele. 9 means missing data.

Here is an example of a genotypic matrix using the geno format with 3 individuals and 4 loci:

```
112
010
091
121
```

Author(s)

Eric Frichot

See Also

[geno2lfmm](#) [lfmm2geno](#) [ancestrymap2geno](#) [ped2geno](#) [vcf2geno](#) [read.geno](#) [write.geno](#)

geno2lfmm *Convert from [geno](#) to [lfmm](#) format*

Description

A function that converts from the [geno](#) format to the [lfmm](#) format.

Usage

```
geno2lfmm(input.file, output.file = NULL, force = TRUE)
```

Arguments

input.file	A character string containing a path to the input file, a genotypic matrix in the geno format.
output.file	A character string containing a path to the output file, a genotypic matrix in the lfmm format. By default, the name of the output file is the same name as the input file with a .lfmm extension.
force	A boolean option. If FALSE, the input file is converted only if the output file does not exist. If TRUE, convert the file anyway.

Value

output.file	A character string containing a path to the output file, a genotypic matrix in the lfmm format.
-------------	-----------------------------------------------------------------------------------------------------------------

Author(s)

Eric Frichot

See Also

[lfmm.data](#) [geno](#) [ancestrymap2lfmm](#) [ancestrymap2geno](#) [ped2lfmm](#) [ped2geno](#) [vcf2geno](#) [lfmm2geno](#)
[read.geno](#) [write.geno](#)

Examples

```
# Creation of a file called "genotypes.geno" in the working directory
# with 400 SNPs for 50 individuals.
data("tutorial")
write.geno(tutorial.R, "genotypes.geno")

# Conversion     from the geno format ("genotypes.geno")
#               to the lfmm format ("genotypes.lfmm").
# By default,   the name of the output file is the same name
#               as the input file with a .lfmm extension.
# Create file:   "genotypes.lfmm".
output = geno2lfmm("genotypes.geno")

# Conversion     from the geno format ("genotypes.geno")
#               to the lfmm format with the output file called "plop.lfmm".
# Create file:   "plop.lfmm".
```

```

output = geno2lfmm("genotypes.geno", "plop.lfmm")

# As force = false and the file "genotypes.lfmm" already exists,
# nothing happens.
output = geno2lfmm("genotypes.geno", force = FALSE)

```

lfmm

*Fitting Latent Factor Mixed Models***Description**

`lfmm` is used to fit Latent Factor Mixed Models. The goal of `lfmm` is to identify genetic polymorphisms that exhibit high correlation with some environmental gradient or with the variables used as proxies for ecological pressures.

Usage

```

lfmm(input.file, environment.file, K,
      project = "continue",
      d = 0, all = FALSE,
      missing.data = FALSE, CPU = 1,
      iterations = 10000, burnin = 5000,
      seed = -1, repetitions = 1,
      epsilon.noise = 1e-3, epsilon.b = 1000,
      random.init = TRUE)

```

Arguments

<code>input.file</code>	A character string containing a path to the input file, a genotypic matrix in the <code>lfmm{lfmm_fomat}</code> format.
<code>environment.file</code>	A character string containing a path to the environmental file, an environmental data matrix in the <code>env</code> format.
<code>K</code>	An integer corresponding to the number of latent factors.
<code>project</code>	A character string among "continue", "new", and "force". If "continue", the results are stored in the current project. If "new", the current project is removed and a new one is created to store the result. If "force", the results are stored in the current project even if the input file has been modified since the creation of the project.
<code>d</code>	An integer corresponding to the fit of <code>lfmm</code> model with the <code>d</code> -th variable only from <code>environment.file</code> . By default (if <code>NULL</code> and <code>all</code> is <code>FALSE</code>), fit <code>lfmm</code> with each variable from <code>environment.file</code> sequentially and independently.
<code>all</code>	A boolean option. If true, fit <code>lfmm</code> with all variables from <code>environment.file</code> at the same time. This option is not compatible with the <code>d</code> option.
<code>missing.data</code>	A boolean option. If true, the <code>input.file</code> contains missing genotypes.
<code>CPU</code>	A number of CPUs to run the parallel version of the algorithm. By default, the number of CPUs is 1.
<code>iterations</code>	The total number of iterations in the Gibbs Sampling algorithm.

<code>burnin</code>	The burnin number of iterations in the Gibbs Sampling algorithm.
<code>seed</code>	A seed to initialize the random number generator. By default, the seed is randomly chosen. The seed is initialized at each repetition. If you want to set a seed, please provide a seed per repetition.
<code>repetitions</code>	The number of repetitions of each run.
<code>epsilon.noise</code>	Prior on the different variances.
<code>epsilon.b</code>	Prior on the variance of the correlation coefficients.
<code>random.init</code>	A boolean option. If true, the Gibbs Sampler is initialized randomly. Otherwise, it is initialized with zeros.

Value

lfmm returns an object of class lfmmProject.

The following methods can be applied to the object of class lfmmProject:

<code>show</code>	Display information about the analyses.
<code>summary</code>	Summarize the analyses.
<code>z.scores</code>	Return the lfmm output vector of zscores for the chosen runs with K latent factors, the d-th variable and the all option.
<code>p.values</code>	Return the lfmm output vector of p-values for the chosen runs with K latent factors, the d-th variable and the all option.
<code>adjusted.pvalues</code>	Return the output vector of adjusted p-values using the genomic control method or the provided lambda inflation factor for the chosen runs with K latent factors, the d-th variable and the all option.
<code>mlog10p.values</code>	Return the lfmm output vector of $-\log_{10}(\text{p-values})$ for the chosen runs with K latent factors, the d-th variable and the all option.
<code>load.lfmmProject (file = "character")</code>	Load the file containing an lfmmProject object and return the lfmmProject object.
<code>remove.lfmmProject (file = "character")</code>	Erase a lfmmProject object. Caution: All the files associated with the object will be removed.
<code>export.lfmmProject(file.lfmmProject)</code>	Create a zip file containing the full lfmmProject object. It allows to move the project to a new directory or a new computer (using import). If you want to overwrite an existing export, use the option <code>force == TRUE</code> .
<code>import.lfmmProject(file.lfmmProject)</code>	Import and load an lfmmProject object from a zip file (made with the export function) into the chosen directory. If you want to overwrite an existing project, use the option <code>force == TRUE</code> .
<code>combine.lfmmProject(file.lfmmProject, toCombine.lfmmProject)</code>	Combine <code>to.Combine.lfmmProject</code> into <code>file.lfmmProject</code> . Caution: Only projects with runs coming from the same input file can be combined. If the same input file has different names in the two projects, use the option <code>force == TRUE</code> .

Author(s)

Eric Frichot

References

Frichot E, Schoville SD, Bouchard G, Francois O. (2013). *Testing for associations between loci and environmental gradients using latent factor mixed models*. *Molecular biology and evolution*, 30(7), 1687-1699.

See Also

[lfmm.data z.scores p.values adjusted.pvalues mlog10p.values pca lfmm tutorial](#)

Examples

```
### Example of analyses using lfmm ###

data("tutorial")
# creation of the genotype file, genotypes.lfmm.
# It contains 400 SNPs for 50 individuals.
write.lfmm(tutorial.R, "genotypes.lfmm")
# creation of the environment file, gradient.env.
# It contains 1 environmental variable for 40 individuals.
write.env(tutorial.C, "gradients.env")

#####
# runs of lfmm #
#####

# main options, K: (the number of latent factors),
#           CPU: the number of CPUs.

# Runs with K = 9 and 5 repetitions.
# The runs are composed of 6000 iterations including 3000 iterations
# for burnin.
# around 30 seconds per run.
project = NULL
project = lfmm("genotypes.lfmm", "gradients.env", K = 6, repetitions = 5,
              project = "new")

# get the adjusted p-values using the genomic control method
res = adjusted.pvalues(project, K = 6)

for (alpha in c(.05,.1,.15,.2)) {
  # expected FDR
  print(paste("expected FDR:", alpha))
  L = length(res$p.values)
  # return a list of candidates with an expected FDR of alpha.
  w = which(sort(res$p.values) < alpha * (1:L) / L)
  candidates = order(res$p.values)[w]

  # estimated FDR and True Positif
  estimated.FDR = length(which(candidates <= 350))/length(candidates)
  estimated.TP = length(which(candidates > 350))/50
  print(paste("FDR:", estimated.FDR, "True Positive:", estimated.TP))
}

#####
# Post-treatments #
#####
```

```

# show the project
show(project)

# summary of the project
summary(project)

# get the z-scores for the 2nd run for K = 6
z = z.scores(project, K = 6, run = 2)

# get the p-values for the 2nd run for K = 6
p = p.values(project, K = 6, run = 2)

# get the adjusted p-values for for K = 6
res = adjusted.pvalues(project, K = 6)

# get the -log10(p-values) for the 2nd run for K = 6
mp = mlog10p.values(project, K = 6, run = 2)

#####
# Manage an lfmm project #
#####

# All the runs of lfmm for a given file are
# automatically saved into a lfmm project directory and a file.
# The name of the lfmmProject file is a combination of
# the name of the input file and the environment file
# with a .lfmmProject extension ("genotypes_gradient.lfmmProject").
# The name of the lfmmProject directory is the same name as
# the lfmmProject file with a .lfmm extension ("genotypes_gradient.lfmm/")
# There is only one lfmm Project for each input file including all the runs.

# An lfmmProject can be load in a different session.
project = load.lfmmProject("genotypes_gradients.lfmmProject")

# An lfmmProject can be exported to be imported in another directory
# or in another computer
export.lfmmProject("genotypes_gradients.lfmmProject")

dir.create("test", showWarnings = TRUE)
#import
newProject = import.lfmmProject("genotypes_gradients_lfmmProject.zip", "test")

# combine projects
combinedProject = combine.lfmmProject("genotypes_gradients.lfmmProject", "test/genotypes_gradients.lfmmProj

# remove
remove.lfmmProject("test/genotypes_gradients.lfmmProject")

# An lfmmProject can be erased.
# Caution: All the files associated with the project will be removed.
remove.lfmmProject("genotypes_gradients.lfmmProject")

```

lfmm.data	<i>Input file for lfmm</i>
-----------	----------------------------

Description

Description of the lfmm format. The lfmm format can be used as an input format for genotypic matrices in the functions [snmf](#), [lfmm](#), and [pca](#).

Details

The lfmm format has one row for each individual. Each row contains one value at each loci (separated by spaces or tabulations) corresponding to the number of alleles. The number of alleles corresponds to the number of reference alleles or the number of derived alleles. Missing genotypes are encoded by the value -9 or 9.

Here is an example of a genotypic matrix using the lfmm format with 3 individuals and 4 loci:

```
1 0 0 1
1 1 9 2
2 0 1 1
```

Author(s)

Eric Frichot

See Also

[lfmm](#) [geno2lfmm](#) [lfmm2geno](#) [ancestrymap2lfmm](#) [ped2lfmm](#) [read.lfmm](#) [write.lfmm](#)

lfmm2geno	<i>Convert from lfmm to geno format</i>
-----------	-----------------------------------------

Description

A function that converts from the [lfmm](#) format to the [geno](#) format.

Usage

```
lfmm2geno(input.file, output.file = NULL, force = TRUE)
```

Arguments

input.file	A character string containing a path to the input file, a genotypic matrix in the lfmm format.
output.file	A character string containing a path to the output file, a genotypic matrix in the geno format. By default, the name of the output file is the same name of the input file with a .geno extension.
force	A boolean option. If FALSE, the input file is converted only if the output file does not exist. If TRUE, convert the file anyway.

Value

output.file A character string containing a path to the output file, a genotypic matrix in the [geno](#) format.

Author(s)

Eric Fritchot

See Also

[lfmm.data](#) [geno](#) [ancestrymap2lfmm](#) [ancestrymap2geno](#) [geno2lfmm](#) [ped2lfmm](#) [ped2geno](#) [vcf2geno](#)

Examples

```
# Creation of a file called "genotypes.lfmm" in the working directory,
# with 400 SNPs for 50 individuals.
data("tutorial")
write.lfmm(tutorial.R, "genotypes.lfmm")

# Conversion      from the lfmm format ("genotypes.lfmm")
#                    to the geno format ("genotypes.geno").
# By default,      the name of the output file is the same name
#                    as the input file with a .geno extension.
# Create file:      "genotypes.geno".
output = lfmm2geno("genotypes.lfmm")

# Conversion      from the lfmm format ("genotypes.lfmm")
#                    to the geno format with the output file called "plop.geno".
# Create file:      "plop.geno".
output = lfmm2geno("genotypes.lfmm", "plop.geno")

# As force = false and the file "genotypes.geno" already exists,
# nothing happens.
output = lfmm2geno("genotypes.lfmm", force = FALSE)
```

mlog10p.values

-log10(p-values) from a lfmm run

Description

Return the lfmm output matrix of $-\log_{10}(p\text{-values})$ for the chosen runs with K latent factors, the d -th variable and the all option. For an example, see [lfmm](#).

Usage

```
mlog10p.values (object, K, d, all, run)
```

Arguments

object A lfmmProject object.
 K The number of latent factors.
 d The d -th variable.

all A Boolean option. If true, the run with all variables at the same time. If false, the runs with each variable separately.

run A list of chosen runs.

Value

res A matrix containing a vector of $-\log_{10}(\text{p-values})$ for the chosen runs per column.

Author(s)

Eric Frichot

See Also

[lfmm.data lfmm p.values adjusted.pvalues z.scores](#)

Examples

```
### Example of analyses using lfmm ###

data("tutorial")
# creation of the genotype file, genotypes.lfmm.
# It contains 400 SNPs for 50 individuals.
write.lfmm(tutorial.R, "genotypes.lfmm")
# creation of the environment file, gradient.env.
# It contains 1 environmental variable for 40 individuals.
write.env(tutorial.C, "gradients.env")

#####
# runs of lfmm #
#####

# main options, K: (the number of latent factors),
#            CPU: the number of CPUs.

# Toy runs with K = 3 and 2 repetitions.
# around 15 seconds per run.
project = NULL
project = lfmm("genotypes.lfmm", "gradients.env", K = 3, repetitions = 2,
              iterations = 6000, burnin = 3000, project = "new")

# get the  $-\log_{10}(\text{p-values})$  for all runs for K = 3
mp = mlog10p.values(project, K = 3)

# get the  $-\log_{10}(\text{p-values})$  for the 2nd run for K = 3
mp = mlog10p.values(project, K = 3, run = 2)
```

p.values

p-values from a lfmm run

Description

Return the lfmm output matrix of p-values for the chosen runs with K latent factors, the d-th variable and the all option. For an example, see [lfmm](#).

Usage

p.values (object, K, d, all, run)

Arguments

object	A lfmmProject object.
K	The number of latent factors.
d	The d-th variable.
all	A Boolean option. If true, the run with all variables at the same time. If false, the runs with each variable separately.
run	A list of chosen runs.

Value

res A matrix containing a vector of p.values for the chosen runs per column.

Author(s)

Eric Frichot

See Also

[lfmm.data](#) [lfmm](#) [mlog10p.values](#) [adjusted.pvalues](#) [z.scores](#)

Examples

```
### Example of analyses using lfmm ###

data("tutorial")
# creation of the genotype file, genotypes.lfmm.
# It contains 400 SNPs for 50 individuals.
write.lfmm(tutorial.R, "genotypes.lfmm")
# creation of the environment file, gradient.env.
# It contains 1 environmental variable for 40 individuals.
write.env(tutorial.C, "gradients.env")

#####
# runs of lfmm #
#####

# main options, K: (the number of latent factors),
#            CPU: the number of CPUs.

# Toy runs with K = 3 and 2 repetitions.
# around 15 seconds per run.
project = NULL
project = lfmm("genotypes.lfmm", "gradients.env", K = 3, repetitions = 2,
              iterations = 6000, burnin = 3000, project = "new")

# get the p-values for all runs for K = 3
p = p.values(project, K = 3)

# get the p-values for the 2nd run for K = 3
p = p.values(project, K = 3, run = 2)
```

Description

The function `pca` performs a Principal Component Analysis of a genotypic matrix using the `lfmm`, `geno`, `ancestrymap`, `ped` or `vcf` format. The function computes eigenvalue, eigenvector, and standard deviation for each principal component and the projection of each individual on each component. The function `pca` returns an object of class "pcaProject" containing the output data and the input parameters.

Usage

```
pca (input.file, K, center = TRUE, scale = FALSE)
```

Arguments

<code>input.file</code>	A character string containing the path to the genotype input file, a genotypic matrix in the <code>lfmm</code> format.
<code>K</code>	An integer corresponding to the number of principal components calculated. By default, all principal components are calculated.
<code>center</code>	A boolean option. If true, the data matrix is centered (default: TRUE).
<code>scale</code>	A boolean option. If true, the data matrix is centered and scaled (default: FALSE).

Value

`pca` returns an object of class `pcaProject` containing the following components:

<code>eigenvalues</code>	The vector of eigenvalues.
<code>eigenvectors</code>	The matrix of eigenvectors (one column for each eigenvector).
<code>sdev</code>	The vector of standard deviations.
<code>projections</code>	The matrix of projections (one column for each projection).

The following methods can be applied to the object of class `pcaProject` returned by `pca`:

<code>plot</code>	Plot the eigenvalues.
<code>show</code>	Display information about the analysis.
<code>summary</code>	Summarize the analysis.
<code>tracy.widom</code>	Perform Tracy-Widom tests on the eigenvalues.
<code>load.pcaProject(file.pcaProject)</code>	Load the file containing a <code>pcaProject</code> object and return the <code>pcaProject</code> object.
<code>remove.pcaProject(file.pcaProject)</code>	Erase a <code>pcaProject</code> object. Caution: All the files associated with the object will be removed.
<code>export.pcaProject(file.pcaProject)</code>	Create a zip file containing the full <code>pcaProject</code> object. It allows to move the project to a new directory or a new computer (using <code>import</code>). If you want to overwrite an existing export, use the option <code>force == TRUE</code> .

```
import.pcaProject(file.pcaProject)
    Import and load an pcaProject object from a zip file (made with the export
    function) into the chosen directory. If you want to overwrite an existing project,
    use the option force == TRUE.
```

Author(s)

Eric Fritchot

See Also

[lfmm.data snmf lfmm tutorial](#)

Examples

```
# Creation of the genotype file "genotypes.lfmm"
# with 1000 SNPs for 165 individuals.
data("tutorial")
write.lfmm(tutorial.R,"genotypes.lfmm")

#####
# Perform a PCA #
#####

# run of PCA
# Available options, K (the number of PCs calculated),
# center and scale.
# Creation of genotypes.pcaProject - the pcaProject object.
# a directory genotypes.pca containing:
# Create files: genotypes.eigenvalues - eigenvalues,
# genotypes.eigenvectors - eigenvectors,
# genotypes.sdev - standard deviations,
# genotypes.projections - projections,
# Create a pcaProject object: pc.
pc = pca("genotypes.lfmm", scale = TRUE)

#####
# Display Information #
#####

# Display information about the analysis.
show(pc)

# Summarize the analysis.
summary(pc)

#####
# Graphical outputs #
#####

par(mfrow=c(2,2))

# Plot eigenvalues.
plot(pc, lwd=5, col="red",xlab=("PCs"),ylab="eigen")

# PC1-PC2 plot.
plot(pc$projections)
```

```

# PC3-PC4 plot.
plot(pc$projections[,3:4])

# Plot standard deviations.
plot(pc$sdev)

#####
# Perform Tracy-Widom tests #
#####

# Perform Tracy-Widom tests on all eigenvalues.
# Create file: genotypes.tracyWidom - tracy-widom test information,
#           in the directory genotypes.pca/.
tw = tracy.widom(pc)

# Plot the percentage of variance explained by each component.
plot(tw$percentage)

# Display the p-values for the Tracy-Widom tests.
tw$pvalues

#####
# Manage an pca project #
#####

# All the file of pca for a given file are
# automatically saved into a pca project directory and a file.
# The name of the pcaProject file is the same name as
# the name of the input file with a .pcaProject extension
# ("genotypes.pcaProject").
# The name of the pcaProject directory is the same name as
# the name of the input file with a .pca extension ("genotypes.pca/")
# There is only one pca Project for each input file including all the runs.

# An pcaProject can be load in a different session.
project = load.pcaProject("genotypes.pcaProject")

# An pcaProject can be exported to be imported in another directory
# or in another computer
export.pcaProject("genotypes.pcaProject")

dir.create("test", showWarnings = TRUE)
#import
newProject = import.pcaProject("genotypes_pcaProject.zip", "test")
# remove
remove.pcaProject("test/genotypes.pcaProject")

# An pcaProject can be erased.
# Caution: All the files associated with the project will be removed.
remove.pcaProject("genotypes.pcaProject")

```

Description

Description of the ped format. The ped format can be used as an input format for genotypic matrices in the functions [snmf](#), [lfmm](#), and [pca](#).

Details

The ped format has one row for each individual. Each row contains 6 columns of information for each individual, plus two genotype columns for each SNP. Each column must be separated by spaces or tabulations. The genotype format must be either 0ACGT or 01234, where 0 means missing genotype. The first 6 columns of the genotype file are: the 1st column is the family ID, the 2nd column is the sample ID, the 3rd and 4th columns are the sample IDs of parents, the 5th column is the gender (male is 1, female is 2), the 6th column is the case/control status (1 is control, 2 is case), the quantitative trait value or the population group label.

The ped format is described [here](#).

Here is an example with 3 individuals and 4 SNPs:

```
1 SAMPLE0 0 0 2 2 1 2 3 3 1 1 2 1
2 SAMPLE1 0 0 1 2 2 1 1 3 0 4 1 1
3 SAMPLE2 0 0 2 1 2 2 3 3 1 4 1 2
```

Author(s)

Eric Fritchot

See Also

[ped2lfmm](#) [ped2geno](#) [geno](#) [lfmm.data](#) [ancestrymap](#) [vcf](#)

ped2geno

Convert from [ped](#) to [geno](#) format

Description

A function that converts from the [ped](#) format to the [geno](#) format.

Usage

```
ped2geno(input.file, output.file = NULL, force = TRUE)
```

Arguments

<code>input.file</code>	A character string containing a path to the input file, a genotypic matrix in the ped format.
<code>output.file</code>	A character string containing a path to the output file, a genotypic matrix in the geno format. By default, the name of the output file is the same name as the input file with a <code>.geno</code> extension.
<code>force</code>	A boolean option. If <code>FALSE</code> , the input file is converted only if the output file does not exist. If <code>TRUE</code> , convert the file anyway.

Value

output.file A character string containing a path to the output file, a genotypic matrix in the [geno](#) format.

Author(s)

Eric Frichot

See Also

[ped geno ancestrymap2lfmm](#) [ancestrymap2geno](#) [geno2lfmm](#) [ped2lfmm](#) [vcf2geno](#) [lfmm2geno](#)

Examples

```
# Creation of a file called "example.ped"
# with 4 SNPs for 3 individuals.
data("example_ped")
write.table(example_ped,"example.ped",
            col.names = FALSE, row.names = FALSE, quote = FALSE)

# Conversion      from the ped format ("example.ped")
#                    to the geno format ("example.geno").
# By default,      the name of the output file is the same name
#                    as the input file with a .geno extension.
# Create file:      "example.geno".
output = ped2geno("example.ped")

# Conversion      from the ped format ("example.ped")
#                    to the geno format with the output file called "plop.geno".
# Create file:      "plop.geno".
output = ped2geno("example.ped", "plop.geno")

# As force = false and the file "example.geno" already exists,
# nothing happens.
output = ped2geno("example.ped", force = FALSE)
```

ped2lfmm

Convert from [ped](#) to [lfmm](#) format

Description

A function that converts from the [ped](#) format to the [lfmm](#) format.

Usage

```
ped2lfmm(input.file, output.file = NULL, force = TRUE)
```

Arguments

input.file A character string containing a path to the input file, a genotypic matrix in the [ped](#) format.

`output.file` A character string containing a path for the output file, a genotypic matrix in the `lfmm` format. By default, the name of the output file is the same name as the input file with a `.lfmm` extension.

`force` A boolean option. If `FALSE`, the input file is converted only if the output file does not exist. If `TRUE`, convert the file anyway.

Value

`output.file` A character string containing a path for the output file, a genotypic matrix in the `lfmm` format.

Author(s)

Eric Frichot

See Also

[ped lfmm.data](#) [ancestrymap2lfmm](#) [ancestrymap2geno](#) [geno2lfmm](#) [ped2geno](#) [vcf2geno](#) [lfmm2geno](#)

Examples

```
# Creation of a file called "example.ped"
# with 4 SNPs for 3 individuals.
data("example_ped")
write.table(example_ped,"example.ped",
            col.names = FALSE, row.names = FALSE, quote = FALSE)

# Conversion    from the ped format ("example.ped")
#              to the lfmm format ("example.lfmm").
# By default,  the name of the output file is the same name
#              as the input file with a .lfmm extension.
# Create file: "example.lfmm".
output = ped2lfmm("example.ped")

# Conversion    from the ped format ("example.ped")
#              to the geno format with the output file called "plop.lfmm".
# Create file:  "plop.lfmm".
output = ped2lfmm("example.ped", "plop.lfmm")

# As force = false and the file "example.lfmm" already exists,
# nothing happens.
output = ped2lfmm("example.ped", force = FALSE)
```

Description

Return the snmf output matrix of admixture coefficients for the chosen run with K ancestral populations. For an example, see [snmf](#).

Usage

`Q(object, K, run)`

Arguments

object	A snmfProject object.
K	The number of ancestral populations.
run	A chosen run.

Value

res	A matrix containing the admixture coefficients for the chosen run with K ancestral populations.
-----	-------------------------------------------------------------------------------------------------

Author(s)

Eric Fritchot

See Also

[geno snmf G cross.entropy](#)

Examples

```
### Example of analyses using snmf ###

# creation of the genotype file, genotypes.geno.
# It contains 400 SNPs for 50 individuals.
data("tutorial")
write.geno(tutorial.R, "genotypes.geno")

#####
# runs of snmf #
#####

# main options, K: (the number of ancestral populations),
#   entropy: calculate the cross-entropy criterion,
#   CPU: the number of CPUs.

# Runs with K between 1 and 5 with cross-entropy and 2 repetitions.
project = NULL
project = snmf("genotypes.geno", K = 3, repetitions = 2, project = "new")

# get the ancestry coefficients for the 2nd run for K = 3.
res = Q(project, K = 3, run = 2)

# plot the 2nd run for K = 3 (ancestry coefficients).
barplot(t(Q(project, K = 3, run = 2)))
```

read.env

Read environmental file in the envformat

Description

Read a file in the [env](#) format.

Usage

```
read.env(input.file)
```

Arguments

`input.file` A character string containing a path to the input file, an environmental data matrix in the `env` format.

Value

R
A matrix containing the environmental variables with one line for each individual and one column for each environmental variable.

Author(s)

Eric Frichot

See Also

[env write.env lfmm](#)

Examples

```
# Creation of an environmental matrix, C
# containing 2 environmental variables for 3 individuals.
# C contains one line for each individual and one column for each variable.
C = matrix(runif(6), ncol=2, nrow=3)

# Write C in a file called "example.env".
# Create file: "example.env".
write.env(C,"example.env")

# Read the file "example.env".
C = read.env("example.env")
```

read.geno

read a file in the [geno](#) format

Description

Read a file in the `geno` format.

Usage

```
read.geno(input.file)
```

Arguments

`input.file` A character string containing a path to the input file, a genotypic matrix in the `geno` format.

Value

R A matrix containing the genotypes with one line for each individual and one column for each SNP.

Author(s)

Eric Fritchot

See Also

[write.geno](#) [geno.snmf](#) [geno2lfmm](#) [lfmm2geno](#) [ancestrymap2geno](#) [ped2geno](#) [vcf2geno](#)

Examples

```
# tutorial contains a matrix of genotypes R with 1000 SNPs for 165 individuals.
# and a matrix with an environmental variable C.
data("tutorial")

# Write R in a file called "genotypes.geno".
# Create file: "genotypes.geno".
write.geno(tutorial.R, "genotypes.geno")

# Read the file "genotypes.geno".
R = read.geno("genotypes.geno")
```

read.lfmm *Read files in the lfmm format*

Description

Read a file in the [lfmm](#) format.

Usage

```
read.lfmm(input.file)
```

Arguments

`input.file` A character string containing a path to the input file, a genotypic matrix in the [lfmm](#) format.

Value

R A matrix containing the genotypes with one line per individual and one column per SNP.

Author(s)

Eric Fritchot

See Also

[write.lfmm](#) [lfmm.data](#) [lfmm](#) [geno2lfmm](#) [lfmm2geno](#) [ancestrymap2lfmm](#) [ped2lfmm](#)

Examples

```
# tutorial contains a matrix of genotypes R with 1000 SNPs for 165 individuals.
# and a matrix with an environmental variable C.
data("tutorial")

# write R in a file called "genotypes.lfmm"
# Create file: "genotypes.lfmm".
write.lfmm(tutorial.R, "genotypes.lfmm")

# read the file "genotypes.lfmm".
R = read.lfmm("genotypes.lfmm")
```

read.zscore	<i>Read the output files of lfmm</i>
-------------	--------------------------------------

Description

Read the output file from [lfmm](#). This is an internal function. Zscores of a run can be accessed using the function [z.scores](#).

Usage

```
read.zscore(input.file)
```

Arguments

input.file a character string containing a path to the output of [lfmm](#).

Value

R A matrix containing the [lfmm](#) results with one line per SNP. The first column is the zscore. The second column is the $-\log_{10}(\text{p-value})$. The third column is the p-value.

Author(s)

Eric Fritchot

See Also

[zscore.format lfmm](#)

Examples

```
### Example of analyses using lfmm ###

data("tutorial")
# creation of the genotype file, genotypes.lfmm.
# It contains 400 SNPs for 50 individuals.
write.lfmm(tutorial.R, "genotypes.lfmm")
# creation of the environment file, gradient.env.
# It contains 1 environmental variable for 40 individuals.
write.env(tutorial.C, "gradients.env")
```

```
#####
# runs of lfmm #
#####

# main options, K: (the number of latent factors),
#           CPU: the number of CPUs.

# Toy runs with K = 3 and 2 repetitions.
# around 15 seconds per run.
project = NULL
project = lfmm("genotypes.lfmm", "gradients.env", K = 3,
              iterations = 6000, burnin = 3000, project = "new")

res = read.zscore("./genotypes_gradients.lfmm/K3/run1/genotypes_r1_s1.3.zscore")
```

snmf	<i>Estimates individual ancestry coefficients and ancestral allele frequencies.</i>
------	-------------------------------------------------------------------------------------

Description

[snmf](#) estimates admixture coefficients using sparse Non-Negative Matrix Factorization algorithms, and provide STRUCTURE-like outputs.

Usage

```
snmf (input.file, K,
      project = "continue",
      repetitions = 1, CPU = 1,
      alpha = 10, tolerance = 0.00001, entropy = FALSE, percentage = 0.05,
      I, iterations = 200, ploidy = 2, seed = -1, Q.input.file)
```

Arguments

input.file	A character string containing a the path to the input file, a genotypic matrix in the geno format.
K	An integer vector corresponding to the number of ancestral populations for which the snmf algorithm estimates have to be calculated.
project	A character string among "continue", "new", and "force". If "continue", the results are stored in the current project. If "new", the current project is removed and a new one is created to store the result. If "force", the results are stored in the current project even if the input file has been modified since the creation of the project.
repetitions	An integer corresponding with the number of repetitions for each value of K.
CPU	A number of CPUs to run the parallel version of the algorithm. By default, the number of CPUs is 1.
alpha	A numeric value corresponding to the snmf regularization parameter. The results can depend on the value of this parameter, especially for small data sets.
tolerance	A numeric value for the tolerance error.

entropy	A boolean value. If true, the cross-entropy criterion is calculated (see create.dataset and cross.entropy.estimate).
percentage	A numeric value between 0 and 1 containing the percentage of masked genotypes when computing the cross-entropy criterion. This option applies only if <code>entropy == TRUE</code> (see cross.entropy).
I	The number of SNPs to initialize the algorithm. It starts the algorithm with a run of <code>snmf</code> using a subset of <code>nb.SNPs</code> random SNPs. If this option is set with <code>nb.SNPs</code> , the number of randomly chosen SNPs is the minimum between 10000 and 10 % of all SNPs. This option can considerably speed up <code>snmf</code> estimation for very large data sets.
iterations	An integer for the maximum number of iterations in algorithm.
ploidy	1 if haploid, 2 if diploid, n if n-ploid.
seed	A seed to initialize the random number generator. By default, the seed is randomly chosen.
Q.input.file	A character string containing a path to an initialization file for Q, the individual admixture coefficient matrix.

Value

`snmf` returns an object of class `snmfProject`.

The following methods can be applied to the object of class `snmfProject`:

<code>plot</code>	Plot the minimal cross-entropy in function of K.
<code>show</code>	Display information about the analyses.
<code>summary</code>	Summarize the analyses.
<code>Q</code>	Return the admixture coefficient matrix for the chosen run with K ancestral populations.
<code>G</code>	Return the ancestral allele frequency matrix for the chosen run with K ancestral populations.
<code>cross.entropy</code>	Return the cross-entropy criterion for the chosen runs with K ancestral populations.
<code>load.snmfProject(file.snmfProject)</code>	Load the file containing an <code>snmfProject</code> object and return the <code>snmfProject</code> object.
<code>remove.snmfProject(file.snmfProject)</code>	Erase a <code>snmfProject</code> object. Caution: All the files associated with the object will be removed.
<code>export.snmfProject(file.snmfProject)</code>	Create a zip file containing the full <code>snmfProject</code> object. It allows to move the project to a new directory or a new computer (using <code>import</code>). If you want to overwrite an existing export, use the option <code>force == TRUE</code> .
<code>import.snmfProject(file.snmfProject)</code>	Import and load an <code>snmfProject</code> object from a zip file (made with the <code>export</code> function) into the chosen directory. If you want to overwrite an existing project, use the option <code>force == TRUE</code> .
<code>combine.snmfProject(file.snmfProject, toCombine.snmfProject)</code>	Combine <code>to.Combine.snmfProject</code> into <code>file.snmfProject</code> . Caution: Only projects with runs coming from the same input file can be combined. If the same input file has different names in the two projects, use the option <code>force == TRUE</code> .

Author(s)

Eric Frichot

References

Frichot E, Mathieu F, Trouillon T, Bouchard G, Francois O. (2014). *Fast and Efficient Estimation of Individual Ancestry Coefficients*. *Genetics*, 194(4): 973–983.

See Also

[geno pca lfmm tutorial](#)

Examples

```
### Example of analyses using snmf ###

# creation of the genotype file, genotypes.geno.
# It contains 400 SNPs for 50 individuals.
data("tutorial")
write.geno(tutorial.R, "genotypes.geno")

#####
# runs of snmf #
#####

# main options, K: (the number of ancestral populations),
#     entropy: calculate the cross-entropy criterion,
#     CPU: the number of CPUs.

# Runs with K between 1 and 5 with cross-entropy and 2 repetitions.
project = NULL
project = snmf("genotypes.geno", K=1:10, entropy = TRUE, repetitions = 10,
              project = "new")

# plot cross-entropy criterion of all runs of the project
plot(project, lwd = 5, col = "red", pch=1)

# get the cross-entropy of each run for K = 4
ce = cross.entropy(project, K = 4)

# select the run with the lowest cross-entropy
best = which.min(ce)

# plot the best run for K = 4 (ancestry coefficients).
barplot(t(Q(project, K = 4, run = best)))

#####
# Post-treatments #
#####

# show the project
show(project)

# summary of the project
summary(project)
```

```

# get the cross-entropy for all runs for K = 4
ce = cross.entropy(project, K = 4)

# get the cross-entropy for the 2nd run for K = 4
ce = cross.entropy(project, K = 4, run = 2)

# get the ancestral genotype frequency matrix, G, for the 2nd run for K = 4.
res = G(project, K = 4, run = 2)

#####
# Advanced snmf run options #
#####

# Q.input.file: init a run with a given ancestry coefficient matrix Q.
# Here, it is initialized with the Q matrix from the first run with K=4
project = snmf("genotypes.geno", K = 4,
  Q.input.file = "./genotypes.snmf/K4/run1/genotypes_r1.4.Q")

# I: init the Q matrix of a run from a smaller run with 100 randomly chosen
# SNPs.
project = snmf("genotypes.geno", K = 4, I = 100)

# CPU: run snmf with 2 CPUs.
project = snmf("genotypes.geno", K = 4, CPU=2)

# percentage: run snmf and calculate the cross-entropy criterion with 10% of
# masked genotypes, instead of 5% of masked genotypes.
project = snmf("genotypes.geno", K = 4, entropy= TRUE, percentage = 0.1)

# seed: choose the seed to init the randomization.
project = snmf("genotypes.geno", K = 4, seed=42)

# alpha: choose the regularization parameter.
project = snmf("genotypes.geno", K = 4, alpha = 100)

# tolerance: choose the tolerance parameter.
project = snmf("genotypes.geno", K = 4, tolerance = 0.0001)

#####
# Manage an snmf project #
#####

# All the runs of snmf for a given file are
# automatically saved into a snmf project directory and a file.
# The name of the snmfProject file is the same name as
# the name of the input file with a .snmfProject extension
# ("genotypes.snmfProject").
# The name of the snmfProject directory is the same name as
# the name of the input file with a .snmf extension ("genotypes.snmf/")
# There is only one snmf Project for each input file including all the runs.

# An snmfProject can be load in a different session.
project = load.snmfProject("genotypes.snmfProject")

# An snmfProject can be exported to be imported in another directory
# or in another computer
export.snmfProject("genotypes.snmfProject")

```

```

dir.create("test", showWarnings = TRUE)
#import
newProject = import.snmfProject("genotypes_snmfProject.zip", "test")
# combine projects
combinedProject = combine.snmfProject("genotypes.snmfProject", "test/genotypes.snmfProject")
# remove
remove.snmfProject("test/genotypes.snmfProject")

# An snmfProject can be erased.
# Caution: All the files associated with the project will be removed.
remove.snmfProject("genotypes.snmfProject")

```

tracy.widom

Tracy-Widom test for eigenvalues

Description

Perform tracy-widom tests on a set of eigenvalues to determine the number of significant eigenvalues and calculate the percentage of variance explained by each principal component. For an example, see [pca](#).

Usage

```
tracy.widom (object)
```

Arguments

object a pcaProject object.

Value

tracy.widom returns a list containing the following components:

eigenvalues	The sorted input vector of eigenvalues (by decreasing order).
twstats	The vector of tracy-widom statistics.
pvalues	The vector of p-values associated with each eigenvalue.
effecn	The vector of effective sizes.
percentage	The vector containing the percentage of variance explained by each principal component.

Author(s)

Eric Frichot

References

Tracy CA and Widom H. (1994). *Level spacing distributions and the bessel kernel*. Commun Math Phys. 161 :289–309. Patterson N, Price AL and Reich D. (2006). *Population structure and eigenanalysis*. PLoS Genet. 2 :20.

See Also

[pca lfmm.data lfmm](#)

Examples

```
# Creation of the genotype file "genotypes.lfmm"
# with 1000 SNPs for 165 individuals.
data("tutorial")
write.lfmm(tutorial.R,"genotypes.lfmm")

#####
# Perform a PCA #
#####

# run of PCA
# Available options, K (the number of PCs calculated),
# center and scale.
# Creation of genotypes.pcaProject - the pcaProject object.
# a directory genotypes.pca containing:
# Create files: genotypes.eigenvalues - eigenvalues,
# genotypes.eigenvectors - eigenvectors,
# genotypes.sdev - standard deviations,
# genotypes.projections - projections,
# Create a pcaProject object: pc.
pc = pca("genotypes.lfmm", scale = TRUE)

#####
# Perform Tracy-Widom tests #
#####

# Perform Tracy-Widom tests on all eigenvalues.
# Create file: genotypes.tracyWidom - tracy-widom test information,
# in the directory genotypes.pca/.
tw = tracy.widom(pc)

# Plot the percentage of variance explained by each component.
plot(tw$percentage)

# Display the p-values for the Tracy-Widom tests.
tw$pvalues

# remove pca Project
remove.pcaProject("genotypes.pcaProject")
```

tutorial

Example tutorial data sets

Description

This dataset is composed of a genotypic matrix called tutorial.R with 50 individuals for 400 SNPs. The last 50 SNPs are correlated with an environmental variable called tutorial.C. This dataset is a subset of the dataset displayed in the note associated with the package.

Usage

tutorial

Value

tutorial.R A genotypic matrix with 50 individuals for 400 SNPs. The last 50 SNPs are correlated with an environmental variable called tutorial.C.

tutorial.C An environmental variable for the 50 individuals.

vcf *vcf format description*

Description

Description of the vcf format. The vcf format can be used as an input format for genotypic matrices in the functions [snmf](#), [lfmm](#), and [pca](#).

Details

The vcf format is described [here](#).

Here is an example of a genotypic matrix using the vcf format with 3 individuals and 4 loci:

```
##fileformat=VCFv4.1
##FORMAT=<ID=GM,Number=1,Type=Integer,Description="Genotype meta">
##INFO=<ID=VM,Number=1,Type=Integer,Description="Variant meta">
##INFO=<ID=SM,Number=1,Type=Integer,Description="SampleVariant meta">
#CHROM POS ID REF ALT QUAL FILTER INFO FORMAT SAMPLE0 SAMPLE1 SAMPLE2
1 1001 rs0000 T C 999 . VM=1;SM=100 GT:GM 1/0:1 0/1:2 1/1:3
1 1002 rs1111 G A 999 . VM=2;SM=101 GT:GM 0/0:6 0/1:7 0/0:8
1 1003 notres G AA 999 . VM=3;SM=102 GT:GM 0/0:11 ./.:12 0/1:13
1 1004 rs2222 G A 999 . VM=3;SM=102 GT:GM 0/0:11 . 1/0:13
1 1003 notres GA A 999 . VM=3;SM=102 GT:GM 0/0:11 ./.:12 0/1:13
1 1005 rs3333 G A 999 . VM=3;SM=102 GT:GM 1/0:11 1/1:12 0/1:13
```

Author(s)

Eric Fritchot

See Also

[vcf2geno](#) [vcf2lfmm](#) [geno](#) [lfmm](#) [ped](#) [ancestrymap](#)

vcf2geno

*Convert from vcf to geno format***Description**

A function that converts from the [vcf](#) format to the [geno](#) format.

Usage

```
vcf2geno(input.file, output.file = NULL, force = TRUE)
```

Arguments

<code>input.file</code>	A character string containing a path to the input file, a genotypic matrix in the vcf format.
<code>output.file</code>	A character string containing a path to the output file, a genotypic matrix in the geno format. By default, the name of the output file is the same name as the input file with a <code>.geno</code> extension.
<code>force</code>	A boolean option. If <code>FALSE</code> , the input file is converted only if the output file does not exist. If <code>TRUE</code> , convert the file anyway.

Value

<code>output.file</code>	A character string containing a path to the output file, a genotypic matrix in the geno format.
--------------------------	-----------------------------------------------------------------------------------------------------------------

Author(s)

Eric Fritchot

See Also

[vcf](#) [geno](#) [ancestrymap2lfmt](#) [ancestrymap2geno](#) [ped2lfmt](#) [ped2geno](#) [lfmt2geno](#) [geno2lfmt](#)

Examples

```
# Creation of a file called "example.vcf"
# with 4 SNPs for 3 individuals.
data("example_vcf")
write.table(example_vcf,"example.vcf",col.names =
  c("#CHROM", "POS", "ID", "REF", "ALT", "QUAL", "FILTER", "INFO",
    "FORMAT", "SAMPLE0", "SAMPLE1", "SAMPLE2"),
  row.names = FALSE, quote = FALSE)

# Conversion from the vcf format ("example.vcf")
# to the geno format ("example.geno").
# By default, the name of the output file is the same name
# as the input file with a .geno extension.
# Create files: "example.geno",
# "example.vcfsnp" - SNP informations,
# "example.removed" - removed lines.
output = vcf2geno("example.vcf")
```

```

# Conversion    from the vcf format ("example.vcf")
#              to the geno format with the output file called "plop.geno".
# Create files: "plop.geno",
#              "plop.vcfsnp" - SNP informations,
#              "plop.removed" - removed lines.
output = vcf2geno("example.vcf", "plop.geno")

# As force = false and the file "example.geno" already exists,
# nothing happens.
output = vcf2geno("example.vcf", force = FALSE)

```

vcf2lfmm

*Convert from vcf to lfmm format***Description**

A function that converts from the [vcf](#) format to the [lfmm](#) format.

Usage

```
vcf2lfmm(input.file, output.file = NULL, force = TRUE)
```

Arguments

<code>input.file</code>	A character string containing a path to the input file, a genotypic matrix in the vcf format.
<code>output.file</code>	A character string containing a path to the output file, a genotypic matrix in the lfmm format. By default, the name of the output file is the same name as the input file with a <code>.lfmm</code> extension.
<code>force</code>	A boolean option. If <code>FALSE</code> , the input file is converted only if the output file does not exist. If <code>TRUE</code> , convert the file anyway.

Value

<code>output.file</code>	A character string containing a path to the output file, a genotypic matrix in the lfmm format.
--------------------------	-----------------------------------------------------------------------------------------------------------------

Author(s)

Eric Frichot

See Also

[vcf lfmm.data](#) [ancestrymap2lfmm](#) [ancestrymap2geno](#) [ped2lfmm](#) [ped2geno](#) [vcf2geno](#)

Examples

```
# Creation of a file called "example.vcf"
# with 4 SNPs for 3 individuals.
data("example_vcf")
write.table(example_vcf,"example.vcf",col.names =
  c("#CHROM", "POS", "ID", "REF", "ALT", "QUAL", "FILTER", "INFO",
    "FORMAT", "SAMPLE0", "SAMPLE1", "SAMPLE2"),
  row.names = FALSE, quote = FALSE)

# Conversion    from the vcf format ("example.vcf")
#              to the lfmm format ("example.lfmm").
# By default,  the name of the output file is the same name
#              as the input file with a .lfmm extension.
# Create files: "example.lfmm",
#              "example.vcfsnp" - SNP informations,
#              "example.removed" - removed lines.
output = vcf2lfmm("example.vcf")

# Conversion    from the vcf format ("example.vcf")
#              to the lfmm format with the output file called "plop.lfmm".
# Create files: "plop.lfmm",
#              "plop.vcfsnp" - SNP informations,
#              "plop.removed" - removed lines.
output = vcf2lfmm("example.vcf", "plop.lfmm")

# As force = false and the file "example.lfmm" already exists,
# nothing happens.
output = vcf2lfmm("example.vcf", force = FALSE)
```

write.env

Write files in the env format

Description

Write a file in the [env](#) format.

Usage

```
write.env(R, output.file)
```

Arguments

R	A matrix containing the environmental variables with one line for each individual and one column for each environmental variable. The missing genotypes have to be encoded with the value 9.
output.file	A character string containing a path to the output file, an environmental data matrix in the env format.

Value

output.file	A character string containing a path to the output file, an environmental data matrix in the env format.
-------------	----------------------------------------------------------------------------------------------------------

Author(s)

Eric Frichot

See Also[read.env](#) [env](#) [lfmm](#)**Examples**

```
# Creation of an environmental matrix C
# containing 2 environmental variables for 3 individuals.
# C contains one line for each individual and one column for each variable.
C = matrix(runif(6), ncol=2, nrow=3)

# Write C in a file called "tuto.env".
# Create file: "tuto.env".
write.env(C,"tuto.env")

# Read the file "tuto.env".
C = read.env("tuto.env")
```

`write.geno`*Write files in the [geno](#) format*

DescriptionWrite a file in the [geno](#) format.**Usage**`write.geno(R, output.file)`**Arguments**

<code>R</code>	A matrix containing the genotypes with one line for each individual and one column for each SNP. The missing genotypes have to be encoded with the value 9.
<code>output.file</code>	A character string containing a path to the output file, a genotypic matrix in the geno format.

Value

<code>output.file</code>	A character string containing a path to the output file, a genotypic matrix in the geno format.
--------------------------	-----------------------------------------------------------------------------------------------------------------

Author(s)

Eric Frichot

See Also[read.geno](#) [geno](#) [snmf](#) [geno2lfmm](#) [lfmm2geno](#) [ancestrymap2geno](#) [ped2geno](#) [vcf2geno](#)

Examples

```
# Creation of a file called "genotypes.geno" in the working directory,
# with 1000 SNPs for 165 individuals.
data("tutorial")

# Write R in a file called "genotypes.geno".
# Create file: "genotypes.geno".
write.geno(tutorial.R, "genotypes.geno")

# Read the file "genotypes.geno".
R = read.geno("genotypes.geno")
```

write.lfmm

Write files in the lfmm format

Description

Write a file in the [lfmm](#) format.

Usage

```
write.lfmm(R, output.file)
```

Arguments

R	A matrix containing the genotypes with one line for each individual and one column for each SNP. The missing genotypes have to be encoded with the value 9.
output.file	A character string containing a path to the output file, a genotypic matrix in the lfmm format.

Value

output.file	A character string containing a path to the output file, a genotypic matrix in the geno format.
-------------	-------------------------------------------------------------------------------------------------

Author(s)

Eric Frichot

See Also

[read.lfmm](#) [lfmm.data](#) [lfmm](#) [lfmm2lfmm](#) [lfmm2geno](#) [ancestrymap2lfmm](#) [ped2lfmm](#)

Examples

```
# Creation of a file called "genotypes.geno" in the working directory,
# with 1000 SNPs for 165 individuals.
data("tutorial")

# write R in a file called "genotypes.lfmm"
# Create file: "genotypes.lfmm".
write.lfmm(tutorial.R, "genotypes.lfmm")
```

```
# read the file "genotypes.lfmm".
R = read.lfmm("genotypes.lfmm")
```

z.scores	<i>z-scores from a lfmm run</i>
----------	---------------------------------

Description

Return the lfmm output matrix of zscores for the chosen runs with K latent factors, the d-th variable and the all option. For an example, see [lfmm](#).

Usage

```
z.scores (object, K, d, all, run)
```

Arguments

object	A lfmmProject object.
K	The number of latent factors.
d	The d-th variable.
all	A Boolean option. If true, the run with all variables at the same time. If false, the runs with each variable separately.
run	A list of chosen runs.

Value

res	A matrix containing a vector of z-scores for the chosen runs per column.
-----	--------------------------------------------------------------------------

Author(s)

Eric Frichot

See Also

[lfmm lfmm.data](#)

Examples

```
### Example of analyses using lfmm ###

data("tutorial")
# creation of the genotype file, genotypes.lfmm.
# It contains 400 SNPs for 50 individuals.
write.lfmm(tutorial.R, "genotypes.lfmm")
# creation of the environment file, gradient.env.
# It contains 1 environmental variable for 40 individuals.
write.env(tutorial.C, "gradients.env")

#####
# runs of lfmm #
#####
```

```
# main options, K: (the number of latent factors),
#           CPU: the number of CPUs.

# Toy runs with K = 3 and 2 repetitions.
# around 15 seconds per run.
project = NULL
project = lfmm("genotypes.lfmm", "gradients.env", K = 3, repetitions = 2,
              iterations = 6000, burnin = 3000, project = "new")

# get the z-scores for all runs for K = 3
z = z.scores(project, K = 3)

# get the z-scores for the 2nd run for K = 3
z = z.scores(project, K = 3, run = 2)

# remove
remove.lfmmProject("genotypes_gradients.lfmmProject")
```

zscore.format

Output file format for lfmm

Description

Description of the zscore output format of [lfmm](#).

Details

The zscore format has one row for each SNP. Each row contains three values: The first value is the zscore, the second value is the $-\log_{10}(\text{pvalue})$, the third value is the p-value (separated by spaces or tabulations).

Author(s)

Eric Fritchot

See Also

[lfmm lfmm.data env](#)

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