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

NeedsCompilation yes

R topics documented:

LEA-package	*LEA: an R package for Landscape and Ecological Associations studies.*

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 num-
ber 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

Package:	LEA
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Author(s)

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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 fatent 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 inflatino 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 th 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 Frichot

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

`input.file`	A character string containing a path to the input file, a genotypic matrix in the ancestrymap 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 as 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 Frichot

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

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

output.file   A character string containing a path to the output file, a genotypic matric 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 matric 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
                _I.geno extension.

seed   A seed to initialize the random number generator. By default, the seed is ran-
                domly chosen.

percentage   A numeric value between 0 and 1 containing the percentage of masked geno-
                types.

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

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

Value

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

masked.ce	The value of the cross-entropy criterion of the masked genotypes.
all.ce	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 polymor-
phisms 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

input.file	A character string containing a path to the input file, a genotypic matrix in the lfmm{lfmm_fomat} format.
environment.file	
	A character string containing a path to the environmental file, an environmental data matrix in the [env](env) format.
K	An integer corresponding to the number of latent factors.
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.
d	An integer corresponding to the fit of lfmm model with the d-th variable only from environment.file. By default (if NULL and all is FALSE), fit lfmm with each variable from environment.file sequentially and independently.
all	A boolean option. If true, fit lfmm with all variables from environment.file at the same time. This option is not compatible with the d option.
missing.data	A boolean option. If true, the input.file contains missing genotypes.
CPU	A number of CPUs to run the parallel version of the algorithm. By default, the number of CPUs is 1.
iterations	The total number of iterations in the Gibbs Sampling algorithm.
burnin	The burnin number of iterations in the Gibbs Sampling algorithm.

seed	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.
repetitions	The number of repetitions of each run.
epsilon.noise	Prior on the different variances.
epsilon.b	Prior on the variance of the correlation coefficients.
random.init	A boolean option. If true, the Gibbs Sampler is initiliazed 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:

show	Display information about the analyses.
summary	Summarize the analyses.
z.scores	Return the lfmm output vector of zscores for the chosen runs with K latent factors, the d-th variable and the all option.
p.values	Return the lfmm output vector of p-values for the chosen runs with K latent factors, the d-th variable and the all option.
adjusted.pvalues	
	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.
mlog10p.values	Return the lfmm output vector of -log10(p-values) for the chosen runs with K latent factors, the d-th variable and the all option.
load.lfmmProject (file = "character")	
	Load the file containing an lfmmProject objet and return the lfmmProject object.
remove.lfmmProject (file = "character")	
	Erase a lfmmProject object. Caution: All the files associated with the object will be removed.
export.lfmmProject(file.lfmmProject)	
	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 force == TRUE.
import.lfmmProject(file.lfmmProject)	
	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 force == TRUE.
combine.lfmmProject(file.lfmmProject, toCombine.lfmmProject)	
	Combine to.Combine.lfmmProject into file.lfmmProject. 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 force == TRUE.

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.lfmmProject"
```

```
# 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   *Input file for* lfmm

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   *Convert from* lfmm *to* geno *format*

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 Frichot

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 -log10(p-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 -log10(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 -log10(p-values) for all runs for K = 3
mp = mlog10p.values(project, K = 3)

# get the -log10(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)
```

---

pca   *Principal Component Analysis*

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

input.file	A character string containg the path to the genotype input file, a genotypic matrix in the [lfmm]() format.
K	An integer corresponding to the number of principal components calculated. By default, all principal components are calculated.
center	A boolean option. If true, the data matrix is centered (default: TRUE).
scale	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:

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

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

plot	Plot the eigenvalues.
show	Display information about the analysis.
summary	Summarize the analysis.
tracy.widom	Perform Tracy-Widom tests on the eigenvalues.
load.pcaProject(file.pcaProject)	
	Load the file containing a pcaProject object and return the pcaProject object.
remove.pcaProject(file.pcaProject)	
	Erase a pcaProject object. Caution: All the files associated with the object will be removed.
export.pcaProject(file.pcaProject)	
	Create a zip file containing the full pcaProject 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 force == TRUE.

```
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 Frichot

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 #
##############################


# Perfom 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")
```

---

ped   ped *format description*

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 Frichot

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

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 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 .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 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 matricx 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 matricx 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)
```

Q   *Admixture coefficients from a snmf run*

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 Frichot

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* env*format*

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 individ-
                ual 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 Frichot

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 Frichot

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   *Read the output files of* lfmm

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 -log10(p-value). The third column is the
               p-value.

Author(s)

Eric Frichot

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	*Estimates individual ancestry coefficients and ancestral allele frequencies.*

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.estimation](#)).
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 entropy == TRUE (see [cross.entropy](#)).
I	The number of SNPs to initialize the algorithm. It starts the algorithm with a run of snmf using a subset of nb.SNPs random SNPs. If this option is set with nb.SNPs, the number of randomly chosen SNPs is the minimum between 10000 and 10 % of all SNPs. This option can considerably speeds up snmf 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:

plot	Plot the minimal cross-entropy in function of K.
show	Display information about the analyses.
summary	Summarize the analyses.
[Q](#)	Return the admixture coefficient matrix for the chosen run with K ancestral populations.
[G](#)	Return the ancestral allele frequency matrix for the chosen run with K ancestral populations.
[cross.entropy](#)	Return the cross-entropy criterion for the chosen runs with K ancestral populations.

load.snmfProject(file.snmfProject)

        Load the file containing an snmfProject objet and return the snmfProject object.

remove.snmfProject(file.snmfProject)

        Erase a snmfProject object. Caution: All the files associated with the object will be removed.

export.snmfProject(file.snmfProject)

        Create a zip file containing the full snmfProject 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 force == TRUE.

import.snmfProject(file.snmfProject)

        Import and load an snmfProject 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.

```
combine.snmfProject(file.snmfProject, toCombine.snmfProject)
```
> Combine to.Combine.snmfProject into file.snmfProject. 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 force == TRUE.

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 significative 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 descreasing 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 #
#############################
```

```
# Perfom 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 invdividuals.

---

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 Frichot

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

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

See Also

[vcf](#) [geno](#) [ancestrymap2lfmm](#) [ancestrymap2geno](#) [ped2lfmm](#) [ped2geno](#) [lfmm2geno](#) [geno2lfmm](#)

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

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

[vcf](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 formt.

Value

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

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*

Description

Write a file in the geno format.

Usage

```
write.geno(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
               geno 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.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 geno2lfmm 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   *z-scores from a lfmm run*

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 -log10(pvalue), the third value is the p-value (separated by spaces or
tabulations).

Author(s)

Eric Frichot

See Also

lfmm lfmm.data env

Index