

# Package ‘radmixture’

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**Title** Calculate Population Stratification

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**Maintainer** Beilei Bian <bianbeilei@wegene.com>

**Description** Implementation of ADMIXTURE for individual ancestry inference in R. Specifically, ADMIXTURE is a software tool for maximum likelihood estimation of individual ancestries from multilocus SNP genotype datasets, see <<https://www.genetics.ucla.edu/software/admixture/>>. Users can use 'radmixture' to calculate ancestry components with different public datasets. It is very convenient and fast for personal genotype data. For more details, see <<https://github.com/wegene-llc/radmixture/blob/master/README.md>>.

**Depends** R (>= 3.1.0)

**Imports** quadprog, plyr, magrittr, MCMCpack

**Suggests** rmarkdown, knitr, testthat

**License** MIT + file LICENSE

**URL** <https://github.com/wegene-llc/radmixture>

**BugReports** <https://github.com/wegene-llc/radmixture/issues>

**Encoding** UTF-8

**LazyData** true

**RoxygenNote** 6.0.1

**NeedsCompilation** yes

**VignetteBuilder** knitr

**ByteCompile** true

**Author** Beilei Bian [aut, cre],  
Dajun Luo [ctb] (R's C API),  
Gang Chen [ctb],  
Senwei Tang [ctb],  
WeGene [cph]

**Repository** CRAN

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br	<i>Block Relaxation for parameters estimation</i>
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### Description

This function is also used for estimating Q and F but faster than EM.

### Usage

```
br(g, q, f, acc, max.iter, tol, model)
```

### Arguments

g	Genotype matrix with dimensions $np$ , where n is sample size and p is the number of SNPs.
q	Ancestry coefficient matrix with dimensions $nK$ , where n is sample size and K is the number of populations.
f	Minor allele frequency matrix with dimensions $Kp$ , where K is the number of populations and p is the number of SNPs.
acc	a logical value indicating whether use quasi-Newton accelerated BR or not.
max.iter	If acc = T, max.iter must be set, the default is 3. max.iter should greater than 1.
tol	Tolerance, if acc = F, tolerance must be set, the default is 1e-4.
model	Choose which model you want to use. Supervised learning or unsupervised learning.

### Value

Estimation results of q, f and the loglikelihood value of each iteration.

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em *Do ancestry analysis with EM algorithm*

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### Description

The EM algorithm could be used for estimating the Q and F matrix.

### Usage

```
em(g, q, f, acc, max.iter, tol, model)
```

### Arguments

g	Genotype matrix with dimensions $np$ , where n is sample size and p is the number of SNPs.
q	Ancestry coefficient matrix with dimensions $nK$ , where n is sample size and K is the number of populations.
f	Minor allele frequency matrix with dimensions $Kp$ , where K is the number of populations and p is the number of SNPs.
acc	a logical value indicating whether use accelerated EM or not.
max.iter	an integer. If acc is TRUE, the number of iterations must be set. max.iter should greater than 1.
tol	Tolerance. If acc is FALSE, tol must be set. The default is 1e-4.
model	Choose which model you want to use. Supervised learning or unsupervised learning.

### Value

Estimation results of q, f and the loglikelihood value of each iteration.

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fFixBr *Block relaxation when f is fixed*

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### Description

This function can be used for ancestry analysis when frequency matrix is fixed.

### Usage

```
fFixBr(gnew, qnew, f, acc, max.iter, tol, pubdata)
```

**Arguments**

gnew	Genotype matrix. The number of row present in gnew is 1 and the number of column is the number of SNPs.
qnew	Initial q used in calculation. A vector. Sum(q) must be 1.
f	Allele frequencies matrix learned from the reference panels.
acc	a logical value indicating whether use quasi-Newton accelerated BR or not.
max.iter	If acc = T, max.iter must be set, the default is 3. max.iter should greater than 1.
tol	If acc = F, tolerance must be set, the default is 1e-4.
pubdata	You can choose a public dataset here, E11, K13, K4, K12b, K7b, World9. You also can use other public dataset which is not in this package.

**Value**

Estimation results of q and the loglikelihood value of each iteration.

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fFixEm	<i>EM when f is fixed</i>
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**Description**

This function can be used for ancestry analysis when frequency matrix is fixed.

**Usage**

```
fFixEm(gnew, qnew, f, acc, max.iter, tol = 1e-4, pubdata)
```

**Arguments**

gnew	Genotype matrix. The number of row present in gnew is 1 and the number of column is the number of SNPs.
qnew	Initial q used in calculation. A vector. sum(q) must be 1.
f	Allele frequencies learned from the reference panels.
acc	a logical value indicating whether use quasi-Newton accelerated EM or not.
max.iter	an integer. If acc is TRUE, the number of iterations must be set. max.iter should greater than 1.
tol	Tolerance. If acc is FALSE, tol must be set. The default is 1e-4.
pubdata	You can choose a public dataset here, E11, K13, K4, K12b, K7b, World9. You also can use other public dataset which is not in this package.

**Value**

Estimation results of q and the loglikelihood value of each iteration.

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fFixQN	<i>quasi-Newton when f is fixed</i>
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**Description**

quasi-Newton for ancestry analysis when F is fixed

**Usage**

```
fFixQN(gnew, qnew, f, tol, method, pubdata)
```

**Arguments**

gnew	Integer which length is the number of SNPs used in calculation.
qnew	Initial q used in calculation. A vector. sum(q) must be 1.
f	Allele frequencies learned from the reference panels.
tol	Tolerance, the default value is 1e-4.
method	Choose which algorithm you want to use. EM or BR.
pubdata	You can choose a public dataset here, E11, K13, K4, K12b, K7b, World9. You also can use other public dataset which is not in this package.

**Value**

Estimation results of q and the loglikelihood value of each iteration.

**Examples**

```
## res <- tfrdpub(genotype, 4, globe4.alleles, globe4.4.F)
## ances <- fFixQN(res$g, res$q, res$f, tol = 1e-4, method = 'BR', pubdata = 'K4')
```

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generateG	<i>Transfer ped file to genotype matrix</i>
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**Description**

This function can be used to transfer a ped file to g matrix

**Usage**

```
generateG(rawped)
```

**Arguments**

rawped	A data.frame. Standard ped format. Genotype should be transferred to 1,2,3,4 from A,C,G,T. 0 represents missing. '-' , '_' , 'I' , 'D' should be replaced by 0 by yourself.
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**Value**

genotype matrix

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initQF	<i>Initialize Q and F</i>
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**Description**

This function could help you initialize Q and F matrix conveniently especially when you intend to use supervised learning.

**Usage**

```
initQF(g, pop = NULL, alpha = NULL, K = NULL, model)
```

**Arguments**

g	genotype matrix
pop	A data.frame. If you intend to do supervised learning, you must specify the ancestries of the reference individuals.
alpha	Parameter for dirichlet distribution. Vector of shape parameters, or matrix of shape parameters corresponding to the number of draw.
K	If you intend to do unsupervised learning, set the number of populations you will use.
model	Choose supervised or unsupervised learning.

**Value**

A list contains q and f matrix.

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qn	<i>quasi-Newton algorithm for ancestry analysis</i>
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**Description**

Use quasi-Newton algorithm to accelerate EM or block relaxation.

**Usage**

```
qn(g, q, f, tol = 1e-4, method, model)
```

**Arguments**

g	Genotype matrix with dimensions $np$ , where n is sample size and p is the number of SNPs.
q	Ancestry coefficient matrix with dimensions $nK$ , where n is sample size and K is the number of populations.
f	Minor allele frequency matrix with dimensions $Kp$ , where K is the number of populations and p is the number of SNPs.
tol	Tolerance, the default value is 1e-4.
method	Choose which algorithm you want to use. EM or BR.
model	Choose which model you want to use. Supervised learning or unsupervised learning.

**Value**

Estimation results of q, f and the loglikelihood value of each iteration.

**Examples**

```
## qn(g, q, f, tol = 1e-4, method = 'BR', model = 'supervised')
```

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radmixture	<i>radmixture</i>
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**Description**

radmixture is an R package for ancestry calculation. It provides both supervised and unsupervised learning with several algorithms for researchers and DNA customers. see README on [GitHub](#)

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tfrdpub	<i>Transfer personal genotype raw data according public dataset</i>
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**Description**

Transfer personal genotype raw data to g matrix which the number of row is 1 and the number of column is the number of SNPs used here.

**Usage**

```
tfrdpub(genotype, K, map, f)
```

**Arguments**

genotype	A data.frame contains your genotype information.
K	The number of populations
map	A data.frame, it should contain rsid, major allele and minor allele information for both plus and minus strands. You should download datasets from GitHub.
f	Frequency matrix learned from reference panel. You should download datasets from GitHub.

**Details**

Please download datasets from [GitHub](#) See README.

**Value**

A list contains g, q, f which can be used for calculation.

**Examples**

```
## download.file(url = 'https://github.com/wegene-llc/radmixture/  
## raw/master/data/globe4.alleles.RData', destfile = 'K4.RData')  
## download.file(url = 'https://github.com/wegene-llc/radmixture/  
## raw/master/data/globe4.4.F.RData', destfile = 'K4f.RData')  
## load('K4.RData')  
## load('K4f.RData')  
## res <- tfrdpub(genotype, 4, globe4.alleles, globe4.4.F)
```

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