

Package ‘TSDFGS’

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Type Package

Title Training Set Determination for Genomic Selection

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Description We propose an optimality criterion to determine the required training set, r-score, which is derived directly from Pearson's correlation between the genomic estimated breeding values and phenotypic values of the test set <[doi:10.1007/s00122-019-03387-0](https://doi.org/10.1007/s00122-019-03387-0)>. This package provides two main functions to determine a good training set and its size.

License GPL (>= 3)

Encoding UTF-8

Imports dplyr, ggplot2, latex2exp, lifecycle, parallel, Rcpp (>= 1.0.8.3)

LinkingTo Rcpp, RcppEigen

RoxygenNote 7.2.0

URL <https://github.com/oumarkme/TSDFGS>

BugReports <https://github.com/oumarkme/TSDFGS/issues>

Depends R (>= 2.10)

LazyData true

NeedsCompilation yes

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cd_score	<i>CD-score</i>
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Description

This function calculate CD-score [doi:10.1186/1297-9686-28-4-359](https://doi.org/10.1186/1297-9686-28-4-359) by given training set and test set.

Usage

```
cd_score(X, X0)
```

Arguments

- X A numeric matrix. The training set genotypic information matrix can be given as genotype matrix (coded as -1, 0, 1) or principle component matrix (row: sample; column: marker).
- X0 A numeric mareix. The test set genotypic information matrix can be given as genotype matrix (coded as -1, 0, 1) or principle component matrix (row: sample; column: marker).

Value

A floating-point number, CD score.

Author(s)

Jen-Hsiang Ou

Examples

```
data(geno)
## Not run: cd_score(geno[1:50, ], geno[51:100])
```

FGCM

Fit logistic growth curve model

Description

A function for fitting logisti growth model

Usage

```
FGCM(geno, nt = NULL, n_iter = NULL, multi.threads = TRUE)
```

Arguments

geno	Genotype information saved as a dataframe. Columns represent variants (SNPs or PCs).
nt	A numerical vector of training set sample size for estimating logistic growth curve parameters
n_iter	Number of simulation of each training set size. Automatically gave a suitable number by default.
multi.threads	Default: TRUE. Set as FALSE if you just want to run it by single thread.

Value

Estimation of parameters.

Examples

```
data(geno)
## Not run: FGCM(geno)
```

geno

Genotype information

Description

A PCA matrix of rice genotype information. This data was published by Zhao et al. (2011) [doi: 10.1038/ncomms1467](https://doi.org/10.1038/ncomms1467)

Usage

```
geno
```

Format

A numeric matrix (PCA) with 404 rows (sample) and 404 columns (PCs).

Source

<http://www.ricediversity.org/data/>

Examples

```
data(geno)
```

nt2r

Simulate r-scores of each training set size

Description

Calculate r-scores (un-target) by in parallel.

Usage

```
nt2r(geno, nt, n_iter = 30, multi.threads = TRUE)
```

Arguments

geno	A numeric dataframe of genotype, column represent sites (genotype coding as 1, 0, -1)
nt	Numeric. Number of training set size
n_iter	Times of iteration. (default = 30)
multi.threads	Default: TRUE

Value

A vector of r-scores of each iteration

Examples

```
data(geno)
## Not run: nt2r(geno, 50)
```

`optTrain`*Optimal training set determination*

Description

This function is designed for determining optimal training set.

Usage

```
optTrain(  
  geno,  
  cand,  
  n.train,  
  subpop = NULL,  
  test = NULL,  
  method = "rScore",  
  min.iter = NULL  
)
```

Arguments

<code>geno</code>	A numeric matrix of principal components (rows: individuals; columns: PCs).
<code>cand</code>	An integer vector of which rows of individuals are candidates of the training set in the <code>geno</code> matrix.
<code>n.train</code>	The size of the target training set. This could be determined with the help of the <code>ssdfgp</code> function provided in this package.
<code>subpop</code>	A character vector of sub-population's group name. The algorithm will ignore the population structure if it remains <code>NULL</code> .
<code>test</code>	An integer vector of which rows of individuals are in the test set in the <code>geno</code> matrix. The algorithm will use an un-target method if it remains <code>NULL</code> .
<code>method</code>	Choices are <code>rScore</code> , <code>PEV</code> and <code>CD</code> . <code>rScore</code> will be used by default.
<code>min.iter</code>	Minimum iteration of all methods can be appointed. One should always check if the algorithm is converged or not. A minimum iteration will set by considering the candidate and test set size if it remains <code>NULL</code> .

Value

This function will return 3 information including `OPTtrain` (a vector of chosen optimal training set), `TOPscore` (highest scores of before iteration), and `ITERscore` (criteria scores of each iteration).

Author(s)

Jen-Hsiang Ou

Examples

```
data(geno)
## Not run: optTrain(geno, cand = 1:404, n.train = 100)
```

pev_score	<i>PEV score</i>
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Description

This function calculate prediction error variance (PEV) score [doi:10.1186/s12711-015-0116-6](https://doi.org/10.1186/s12711-015-0116-6) by given training set and test set.

Usage

```
pev_score(X, X0)
```

Arguments

X	A numeric matrix. The training set genotypic information matrix can be given as genotype matrix (coded as -1, 0, 1) or principle component matrix (row: sample; column: marker).
X0	A numeric matrix. The test set genotypic information matrix can be given as genotype matrix (coded as -1, 0, 1) or principle component matrix (row: sample; column: marker).

Value

A floating-point number, PEV score.

Author(s)

Jen-Hsiang Ou

Examples

```
data(geno)
## Not run: pev_score(geno[1:50, ], geno[51:100])
```

r_score

r-score

Description

This function calculate r-score [doi:10.1007/s00122-019-03387-0](https://doi.org/10.1007/s00122-019-03387-0) by given training set and test set.

Usage

```
r_score(X, X0)
```

Arguments

X	A numeric matrix. The training set genotypic information matrix can be given as genotype matrix (coded as -1, 0, 1) or principle component matrix (row: sample; column: marker).
X0	A numeric mareix. The test set genotypic information matrix can be given as genotype matrix (coded as -1, 0, 1) or principle component matrix (row: sample; column: marker).

Value

A floating-point number, r-score.

Author(s)

Jen-Hsiang Ou

Examples

```
data(geno)
## Not run: r_score(geno[1:50, ], geno[51:100])
```

SSDFGS

Sample size determination for genomic selection

Description

This function is designed to generate an operating curve for sample size determination

Usage

```
SSDFGS(geno, nt = NULL, n_iter = NULL, multi.threads = TRUE)
```

Arguments

<code>geno</code>	A numeric data frame carried genotype information (column: PCs, row: sample)
<code>nt</code>	A numeric vector carried training set sizes for r-score simulation.
<code>n_iter</code>	Number of iterations for estimating parameters.
<code>multi.threads</code>	Default (<code>multi.threads = TRUE</code>) use 75% of threads if the computer has more than 4 threads.

Value

An operating curve and its information.

Author(s)

Jen-Hsiang Ou & Po-Ya Wu

Examples

```
data(geno)
## Not run: SSDFGS(geno)
```

subpop	<i>Sub-population information</i>
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Description

Sub-population information of samples. This data was published by Zhao et al. (2011) [doi:10.1038/ncomms1467](https://doi.org/10.1038/ncomms1467)

Usage

```
subpop
```

Format

A character vector.

Source

<http://www.ricediversity.org/data/>

Examples

```
data(subpop)
```


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