

Package ‘milorGWAS’

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Type Package

Title Mixed Logistic Regression for Genome-Wide Analysis Studies
(GWAS)

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Author Hervé Perdry and Jacqueline Milet

Maintainer Hervé Perdry <herve.perdry@u-psud.fr>

Description Fast approximate methods for mixed logistic regression in genome-wide analysis studies (GWAS). Milet et al 2020 <[doi:10.1101/2020.01.17.910109](https://doi.org/10.1101/2020.01.17.910109)>.

License GPL-3

Imports Rcpp (>= 1.0.2)

Depends gaston (>= 1.5.6)

LinkingTo Rcpp, RcppEigen, gaston

Suggests knitr, rmarkdown, png

VignetteBuilder knitr

NeedsCompilation yes

RoxygenNote 7.0.2

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```
association.test.logistic
```

Mixed logistic regression for GWAS

Description

Mixed logistic regression for GWAS

Usage

```
association.test.logistic(
  x,
  Y = x@ped$pheno,
  X = matrix(1, nrow(x)),
  K,
  beg = 1,
  end = ncol(x),
  algorithm = c("offset", "amle"),
  eigenK,
  p = 0,
  ...
)
```

Arguments

x	a bedmatrix
Y	phenotype vector. Default is column pheno of x@ped
X	A matrix of covariates (defaults to a column of ones for the intercept)
K	A genetic relationship matrix (or a list of such matrices)
beg	Index of the first SNP tested for association
end	Index of the last SNP tested for association
algorithm	Algorithm to use
eigenK	eigen decomposition of K (only if p > 0)
p	Number of principal components to include in the model
...	Additional parameter for <code>gaston::logistic.mm.aireml</code>

Details

Tests the association between the phenotype and requested SNPs in x. The phenotype Y is a binary trait. A Wald test is performed using an approximate method defined by the parameter `algorithm`. All other arguments are as in `gaston::association.test`.

Value

A data frame giving for each SNP the association statistics.

See Also[association.test](#)**Examples**

```
data(TTN)
x <- as.bed.matrix(TTN.gen, TTN.fam, TTN.bin)
## Simulation data ##
set.seed(1)
# some covariables
X <- cbind(1, runif(nrow(x)))
# A random GRM
ran <- random.pm( nrow(x))
# random effects (tau = 1)
omega <- lmm.simu(1, 0, eigenK=ran$eigen)$omega
# linear term of the model
lin <- X %*% c(0.1,-0.2) + omega
# vector of probabilities
pi <- 1/(1+exp( -lin ))
# vector of binary phenotypes
y <- rbinom(nrow(x), 1, pi)
# testing association with 1) the score test, 2) the offset algorithm, 3) the 'amle' algorithm
a1 <- association.test(x, y, X, K = ran$K, method = "lmm", response = "bin")
a2 <- association.test.logistic(x, y, X, K = ran$K, algorithm = "offset")
a3 <- association.test.logistic(x, y, X, K = ran$K, algorithm = "amle")
```

`qqplot.pvalues`*Stratified QQ-plot of p-values*

Description

Draws a QQ plot of p-values

Usage

```
qqplot.pvalues(
  p,
  snp.cat,
  col.cat,
  col.abline = "red",
  CB = TRUE,
  col.CB = "gray80",
  CB.level = 0.95,
  thinning = TRUE,
  ...
)
```

Arguments

p	vector of p-values, or a data.frame with a column named p
snp.cat	(optional) A factor giving the SNP categories.
col.cat	(optional) A vector of colors used to plot the SNP categories.
col.abline	Color of the line of slope 1. Set to NA to suppress.
CB	Logical. If TRUE, a confidence band is included in the plot.
col.CB	The color of the confidence band.
CB.level	The level of the confidence band.
thinning	Logical. If TRUE, not all points are displayed.
...	Graphical parameters to be passed to plot and points

Details

This function draws a QQ plot of p -values, stratified by categories. If the parameter `snp.cat` is missing, the function falls back on `gaston::qqplot.pvalues`.

See Also

[SNP.category](#), [qqplot.pvalues](#) (in `gaston`)

Examples

```
# a random vector of categories
ca <- sample(c("A","B","C"), 1e6, TRUE, c(0.05, 0.9, 0.05))
# a vector of p-values, with different distribution depending on the strata
p <- runif(1e6)**ifelse(ca == "A", .8, ifelse(ca == "B", 1, 1.2))
qqplot.pvalues(p, ca)
```

SNP.category

SNP.category

Description

SNP.category

Usage

```
SNP.category(bed, Z, threshold = 0.8)
```

Arguments

bed	A bed matrix
Z	A vector of length <code>nrow(bed)</code>
threshold	Variance thresholds

Details

This function determines a SNP Category from a covariable Z, which can be for example an indicator variable for a population strata, or the first genomic principal component.

See Also

[qqplot.pvalues](#)

Examples

```
# a random vector of categories
ca <- sample(c("A","B","C"), 1e6, TRUE, c(0.05, 0.9, 0.05))
# a vector of p-values, with different distribution depending on the strata
p <- runif(1e6)**ifelse(ca == "A", .8, ifelse(ca == "B", 1, 1.2))
qqplot.pvalues(p, ca)
```

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