

Package: CodataGS (via r-universe)

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Type Package

Title Genomic Prediction Using SNP Codata

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Description Computes genomic breeding values using external information on the markers. The package fits a linear mixed model with heteroscedastic random effects, where the random effect variance is fitted using a linear predictor and a log link. The method is described in Mouresan, Selle and Ronnegard (2019) <doi:10.1101/636746>.

License GPL

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NeedsCompilation no

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 CodataGS-package *Genomic Prediction Using SNP Codata*

Description

Computes genomic breeding values using external information on the markers. The package fits a linear mixed model with heteroscedastic random effects, where the random effect variance is fitted using a linear predictor and a log link. The method is described in Mouresan, Selle and Ronnegard (2019) <doi:10.1101/636746>.

Details

The DESCRIPTION file:

```
Package:           CodataGS
Type:              Package
Title:             Genomic Prediction Using SNP Codata
Version:           1.43
Date:              2019-05-17
Author:            Lars Ronnegard
Maintainer:        Lars Ronnegard <lrn@du.se>
Description:        Computes genomic breeding values using external information on the markers. The package fits a linear
License:           GPL
Depends:           Matrix
NeedsCompilation: no
```

Index of help topics:

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CodataGS-package   Genomic Prediction Using SNP Codata
MME                 Mixed model equations
Transform           Transforms hat values
compute_GL          Computes genomic relationship matrix
compute_phitau      Computes models for the variance components
genomicEBV.w.codata Performs genomic prediction based on SNP
                    codata.
hat.transf          Transforms hat values
scaleZ              Scales the genotype matrix.
summary.CodataGS    Summary method for CodataGS objects
```

This package performs genomic prediction based on SNP codata. The main function is genomicEBV.w.codata.

Author(s)

Lars Ronnegard
 Maintainer: Lars Ronnegard <lrn@du.se>

compute_GL

Computes genomic relationship matrix

Description

This function computes the genomic relationship matrix, G, together with its matrix square root, L.

Usage

```
compute_GL(Z, w)
```

Arguments

Z	Scaled matrix with genotype information
w	weights

Value

L	Square root matrix of G
svdVec	Vectors in the Single Value Decomposition of G
svdD	Diagonal elements in the Single Value Decomposition of G
wZt	weights times the transpose of Z

Author(s)

Lars Ronnegard

Examples

```
set.seed(1234)
N <- 20 #Number of individuals
k <- 30 #Number of SNPs with all marker positions including a QTL
Z1 <- matrix(0, N, k )
Z2 <- matrix(0, N, k )
Z1[1:N, 1] <- rbinom(N, 1, 0.5) #Simulated phased SNP matrices
Z2[1:N, 1] <- rbinom(N, 1, 0.5)
LD.par <- 0.2 #A parameter to simulate LD. 0 gives full LD, and 0.5 no LD
for (j in 2:k) {
  Z1[1:N, j] <- abs( Z1[1:N, j-1] - rbinom(N, 1, LD.par) )
  Z2[1:N, j] <- abs( Z2[1:N, j-1] - rbinom(N, 1, LD.par) )
}
Z <- Z1 + Z2 #Genotypic SNP matrix
sim.res <- compute_GL(Z, w = rep(1,k))
```

compute_phitau *Computes models for the variance components*

Description

This function computes the residual variance, the SNP variances and the linear predictor for the SNP variance model.

Usage

```
compute_phitau(dev, hv, devu, hvu, X.rand.disp)
```

Arguments

dev	Deviance values
hv	Hat values for the observed response values
devu	Deviance values computed for the random effects
hvu	Hat values for the random effects
X.rand.disp	Design matrix used in the linear predictor for the SNP variance model.

Value

var.e	Residual variance
phi	Vector of SNP variances
coef	Fitted coefficients for the linear predictor in the SNP variance model

Author(s)

Lars Ronnegard

Examples

```
set.seed(1234)
N <- 20 #Number of individuals
k <- 30 #Number of SNPs with all marker positions including a QTL
#Simulated deviances and hat values
dev <- rnorm(N)^2
hv <- runif(N, 0.1, 0.5)
devu <- rnorm(k)^2
hvu <- runif(k, 0.1, 0.85)
X.rand.disp <- matrix(1, k, 1)
sim.res <- compute_phitau(dev, hv, devu, hvu, X.rand.disp)
```

genomicEBV.w.codata *Performs genomic prediction based on SNP codata.*

Description

The main function of the package. The input includes response values, a design matrix for the fixed effects, a matrix with SNP genotype data and a design matrix for the SNP codata.

Usage

```
genomicEBV.w.codata(y, X, Z, X.SNPcodata, Z.test = NULL, max.iter = 100, conv.crit = 1e-5)
```

Arguments

y	Response values
X	Design matrix for the fixed effects
Z	Genotype matrix with element values of 0, 1 or 2
X.SNPcodata	Design matrix for the linear predictor of the SNP variances.
Z.test	An optional genotype matrix for a test data set.
max.iter	The maximum number of iterations
conv.crit	The value of the convergence criterion.

Details

By specifying the matrix `Z.test` in the input, the function computes predicted genomic breeding values for an out-of-sample data set.

Value

gEBV	Genomic breeding values
predicted.gEBV	Genomic breeding values based on the genotypes in <code>Z.test</code>
w	Computed SNP weights
u	Fitted SNP effects
beta	Fitted fixed effects
disp.beta	Fitted coefficients in the linear predictor for the SNP variance model
Converge	Shows whether the algorithm has converged or not
iter	The number of iterations used

Author(s)

Lars Ronnegard

Examples

```
#####
#Simulation part
set.seed(1234)
N <- 200 #Number of individuals
k <- 300 #Number of SNPs with all marker positions including a QTL
Z1 <- matrix(0, N, k )
Z2 <- matrix(0, N, k )
Z1[1:N, 1] <- rbinom(N, 1, 0.5) #Simulated phased SNP matrices
Z2[1:N, 1] <- rbinom(N, 1, 0.5)
LD.par <- 0.2 #A parameter to simulate LD. 0 gives full LD, and 0.5 no LD
for (j in 2:k) {
  Z1[1:N, j] <- abs( Z1[1:N, j-1] - rbinom(N, 1, LD.par) )
  Z2[1:N, j] <- abs( Z2[1:N, j-1] - rbinom(N, 1, LD.par) )
}
Z <- Z1 + Z2 #Genotypic SNP matrix
x1 <- c(rep(1,k/2), rep(0,k/2)) #An indicator for the SNPs.
#The first k/2 SNPs and the last k/2 have different variances
#Simulate linear predictor for the random effect variance
lin.pred <- 0 + 2*x1
X.snp <- model.matrix( ~ x1 ) #Corresponding design matrix
u <- rnorm(k, 0 , sqrt( exp(lin.pred) ))
#Took the square root here because it is the SD that is specified.
#and exp() because we are modelling a log link.
u.scaled <- u/as.numeric( sqrt( var( crossprod(t(Z), u) ) ) )
#Scaled by the variance of the breeding values
e <- rnorm(N) #A residual variance
mu <- 0
y <- mu + crossprod(t(Z),u.scaled) + e
#####
#Estimation part
mod1 <- genomicEBV.w.codata(y = as.numeric(y),
  X = matrix(1, N, 1), Z = Z, X.SNPcodata = X.snp)
#To fit gBLUP just specify X.SNPcodata = matrix(1, k, 1)
cat("Correlation between true and estimated BV for the codata model:")
cat(cor(crossprod(t(Z),u.scaled), mod1$gEBV), "\n")
```

hat.transf

Transforms hat values

Description

Transforms hat values between the SNP-BLUP model and the gBLUP model.

Usage

```
hat.transf(C22, transf, vc, k, N, w)
```

Arguments

C22	Submatrix of the inverse of the LHS in the MME
transf	A transformation matrix.
vc	Genetic variance
k	Number of SNPs
N	Number of individuals
w	SNP weights

Value

Transformed hat values

Author(s)

Lars Ronnegard

MME

Mixed model equations

Description

A fast version of the Henderson's mixed model equations (MME)

Usage

MME(y, X, Z, var.e, var.u)

Arguments

y	Response
X	Design matrix for fixed effects
Z	Design matrix for the random effects
var.e	Residual variance
var.u	Genetic variance

Value

beta	Estimates of fixed effects
v	Fitted random effects
hv	Hat values
dev	Deviances

Author(s)

Lars Ronnegard

scaleZ *Scales the genotype matrix.*

Description

Scales the genotype matrix so that ZZ' gives the genomic relationship matrix.

Usage

```
scaleZ(Z, freq1)
```

Arguments

Z	Genotype matrix with element values 0, 1 and 2
freq1	Optional input parameter with allele frequencies. A vector of length equal to the number of columns in Z.

Value

Z	Scaled genotype matrix
---	------------------------

Author(s)

Lars Ronnegard

Examples

```
#####
#Simulation part
set.seed(1234)
N <- 200 #Number of individuals
k <- 300 #Number of SNPs with all marker positions including a QTL
Z1 <- matrix(0, N, k )
Z2 <- matrix(0, N, k )
Z1[1:N, 1] <- rbinom(N, 1, 0.5) #Simulated phased SNP matrices
Z2[1:N, 1] <- rbinom(N, 1, 0.5)
LD.par <- 0.2 #A parameter to simulate LD. 0 gives full LD, and 0.5 no LD
for (j in 2:k) {
  Z1[1:N, j] <- abs( Z1[1:N, j-1] - rbinom(N, 1, LD.par) )
  Z2[1:N, j] <- abs( Z2[1:N, j-1] - rbinom(N, 1, LD.par) )
}
Z <- Z1 + Z2 #Genotypic SNP matrix
sim.res <- scaleZ(Z)
```

summary.CodataGS *Summary method for CodataGS objects*

Description

A summary method for the object class CodataGS

Usage

```
## S3 method for class 'CodataGS'
summary(object, ...)
```

Arguments

object	A CodataGS object
...	arguments not used

Details

Provides a concise summary of CodataGS objects.

Examples

```
#####
#Simulation part
set.seed(1234)
N <- 200 #Number of individuals
k <- 300 #Number of SNPs with all marker positions including a QTL
Z1 <- matrix(0, N, k )
Z2 <- matrix(0, N, k )
Z1[1:N, 1] <- rbinom(N, 1, 0.5) #Simulated phased SNP matrices
Z2[1:N, 1] <- rbinom(N, 1, 0.5)
LD.par <- 0.2 #A parameter to simulate LD. 0 gives full LD, and 0.5 no LD
for (j in 2:k) {
  Z1[1:N, j] <- abs( Z1[1:N, j-1] - rbinom(N, 1, LD.par) )
  Z2[1:N, j] <- abs( Z2[1:N, j-1] - rbinom(N, 1, LD.par) )
}
Z <- Z1 + Z2 #Genotypic SNP matrix
x1 <- c(rep(1,k/2), rep(0,k/2)) #An indicator for the SNPs.
#The first k/2 SNPs and the last k/2 have different variances
#Simulate linear predictor for the random effect variance
lin.pred <- 0 + 2*x1
X.snp <- model.matrix( ~ x1 ) #Corresponding design matrix
u <- rnorm(k, 0 , sqrt( exp(lin.pred) ))
#Took the square root here because it is the SD that is specified.
#and exp() because we are modelling a log link.
u.scaled <- u/as.numeric( sqrt( var( crossprod(t(Z), u) ) ) )
#Scaled by the variance of the breeding values
e <- rnorm(N) #A residual variance
```

```

mu <- 0
y <- mu + crossprod(t(Z),u.scaled) + e
#####
#Estimation part
mod1 <- genomicEBV.w.codata(y = as.numeric(y),
                           X = matrix(1, N, 1), Z = Z, X.SNPcodata = X.snp)
summary(mod1)

```

Transform

Transforms hat values

Description

The function calls the `hat.transf` function.

Usage

```
Transform(X, L, var.e, var.u, v, svdVec, svdD, wZt, w)
```

Arguments

<code>X</code>	Design matrix for the fixed effects
<code>L</code>	Square root matrix of the genomic relationship matrix, G
<code>var.e</code>	Residual variance
<code>var.u</code>	Genetic variance
<code>v</code>	Random effects
<code>svdVec</code>	Vector from the Single Value Decomposition of G
<code>svdD</code>	Diagonal elements of the Single Value Decomposition of G
<code>wZt</code>	Weights times the transpose of the scaled genotype matrix
<code>w</code>	Fitted SNP weights

Value

<code>u</code>	SNP effects
<code>qu</code>	Hat values for the SNP effects

Author(s)

Lars Ronnegard

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