Package ‘BGLR’

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The BGLR ('Bayesian Generalized Linear Regression') function fits various types of parametric and semi-parametric Bayesian regressions to continuous (censored or not), binary and ordinal outcomes.

**Usage**

```r
BGLR(y, response_type = "gaussian", a==NULL, b==NULL, ETA = NULL, nIter = 1500, burnIn = 500, thin = 5, saveAt = "", S0 = NULL, df0 = 5, R2 = 0.5, weights = NULL, verbose = TRUE, rmExistingFiles = TRUE, groups=NULL)
```

**Arguments**

- **y** (numeric, n) the data-vector (NAs allowed).
- **response_type** (string) admits values "gaussian" or "ordinal". The Gaussian outcome may be censored or not (see below). If response_type="gaussian", y should be coercible to numeric. If response_type="ordinal", y should be coercible to character, and the order of the outcomes is determined based on the alphanumeric order A<1<RNN<a<bNNI. For ordinal traits the probit link is used.
- **a, b** (numeric, n) only required for censored outcomes, a and b are vectors specifying lower and upper bounds for censored observations, respectively. The default value, for non-censored and ordinal outcomes, is NULL (see details).
- **ETA** (list) This is a two-level list used to specify the regression function (or linear predictor). By default the linear predictor (the conditional expectation function in case of Gaussian outcomes) includes only an intercept. Regression on covariates and other types of random effects are specified in this two-level list. For instance:

```r
ETA=list(list(X=W, model="FIXED"), list(X=Z,model="BL"), list(K=G,model="RKHS")),
```

specifies that the linear predictor should include: an intercept (included by default) plus a linear regression on W with regression coefficients treated as fixed effects (i.e., flat prior), plus regression on Z, with regression coefficients modeled as in the Bayesian Lasso of Park and Casella (2008) plus a random effect with covariance structure G.

For linear regressions the following options are implemented: FIXED (Flat prior), BRR (Gaussian prior), BayesA (scaled-t prior), BL (Double-Exponential prior).
BGLR implements a Gibbs sampler for a Bayesian regression model. The linear predictor (or regression function) includes an intercept (introduced by default) plus a number of user-specified regression components (X) and random effects (u), that is:

$$\eta = 1 + X_1\beta_1 + ... + X_p\beta_p + u_1 + ... + u_q$$

The components of the linear predictor are specified in the argument ETA (see above). The user can specify as many linear terms as desired, and for each component the user can choose the prior density to be assigned. The distribution of the response is modeled as a function of the linear predictor.
For Gaussian outcomes, the linear predictor is the conditional expectation, and censoring is allowed. For censored data points the actual response value \( y_i \) is missing, and the entries of the vectors a and b (see above) give the lower an upper vound for \( y_i \). The following table shows the configuration of the triplet \((y, a, b)\) for un-censored, right-censored, left-censored and interval censored.

<table>
<thead>
<tr>
<th>Type</th>
<th>a</th>
<th>y</th>
<th>b</th>
</tr>
</thead>
<tbody>
<tr>
<td>Un-censored</td>
<td>NULL</td>
<td>( y_i )</td>
<td>NULL</td>
</tr>
<tr>
<td>Right censored</td>
<td>( a_i )</td>
<td>NA</td>
<td>( \infty )</td>
</tr>
<tr>
<td>Left censored</td>
<td>( -\infty )</td>
<td>NA</td>
<td>( b_i )</td>
</tr>
<tr>
<td>Interval censored</td>
<td>( a_i )</td>
<td>NA</td>
<td>( b_i )</td>
</tr>
</tbody>
</table>

Internally, censoring is dealt with as a missing data problem.

*Ordinal outcomes* are modelled using the probit link, implemented via data augmentation. In this case the linear predictor becomes the mean of the underlying liability variable which is normal with mean equal to the linear predictor and variance equal to one. In case of only two classes (binary outcome) the threshold is set equal to zero, for more than two classess thresholds are estimated from the data. Further details about this approach can be found in Albert and Chib (1993).

**Value**

A list with estimated posterior means, estimated posterior standard deviations, and the parameters used to fit the model. See the vignettes in the package (or [http://genomics.cimmyt.org/BGLR-extdoc.pdf](http://genomics.cimmyt.org/BGLR-extdoc.pdf)) for further details.

**Author(s)**

Gustavo de los Campos, Paulino Perez Rodriguez,

**References**


Examples

```r
## Not run:
#Demos
library(BGLR)

#BayesA
demo(BA)

#BayesB
demo(BB)

#Bayesian LASSO
demo(BL)

#Bayesian Ridge Regression
demo(BRR)

#BayesCpi
demo(BayesCpi)

#RKHS
demo(RKHS)

#Binary traits
demo(Bernoulli)

#Ordinal traits
demo(ordinal)

#Censored traits
demo(censored)

## End(Not run)
```

---

Bayesian Linear Regression

Description

The BLR (‘Bayesian Linear Regression’) function was designed to fit parametric regression models using different types of shrinkage methods. An earlier version of this program was presented in de los Campos et al. (2009).

Usage

```r
BLR(y, XF, XR, XL, GF, prior, nIter, burnIn, thin, thin2, saveAt, minAbsBeta, weights)
```
Arguments

\( y \) (numeric, \( n \)) the data-vector (NAs allowed).

\( X_F \) (numeric, \( n \times p_F \)) incidence matrix for \( \beta_F \), may be NULL.

\( X_R \) (numeric, \( n \times p_R \)) incidence matrix for \( \beta_R \), may be NULL.

\( X_L \) (numeric, \( n \times p_L \)) incidence matrix for \( \beta_L \), may be NULL.

\( GF \) (list) providing an \$ID (integer, \( n \)) linking observations to groups (e.g., lines or sires) and a (co)variance structure \( \$A \) (numeric, \( p_U \times p_U \)) between effects of the grouping factor (e.g., line or sire effects). Note: ID must be an integer taking values from 1 to \( p_U \); \$ID[i]=q indicates that the ith observation in \( y \) belongs to cluster \( q \) whose (co)variance function is in the qth row (column) of \( A \). GF may be NULL.

weights (numeric, \( n \)) a vector of weights, may be NULL.

\( niter, burnin, thin \) (integer) the number of iterations, burn-in and thinning.

\( saveAt \) (string) this may include a path and a pre-fix that will be added to the name of the files that are saved as the program runs.

\( prior \) (list) containing the following elements,

- \( prior$varE, prior$varBR, prior$varU \) (list) each providing degree of freedom \( (df) \) and scale \( (S) \). These are the parameters of the scaled inverse-\( \chi^2 \) distributions assigned to variance components, see Eq. (2) below. In the parameterization used by BLR() the prior expectation of variance parameters is \( S/(df - 2) \).

- \( prior$lambda \) (list) providing \$value (initial value for \( \lambda \)); \$type (‘random’ or ‘fixed’) this argument specifies whether \( \lambda \) should be kept fixed at the value provided by \$value or updated with samples from the posterior distribution; and, either \$shape and \$rate (this when a Gamma prior is desired on \( \lambda^2 \)) or \$shape1, \$shape2 and \$max, in this case \( p(\lambda|\max, \alpha_1, \alpha_2) \propto Beta(\frac{\max}{\lambda}, \alpha_1, \alpha_2) \). For detailed description of these priors see de los Campos et al. (2009).

\( thin2 \) This value controls whether the running means are saved to disk or not. If \( thin2 \) is greater than \( niter \) the running means are not saved (default, \( thin2=1 \times 10^{10} \)).

\( minAbsBeta \) The minimum absolute value of the components of \( \beta_L \) to avoid numeric problems when sampling from \( \tau^2 \), default \( 1 \times 10^{-9} \).

Details

The program runs a Gibbs sampler for the Bayesian regression model described below.

Likelihood. The equation for the data is:

\[ y = 1\mu + X_F\beta_F + X_R\beta_R + X_L\beta_L + Zu + \varepsilon \quad (1) \]

where \( y \), the response is a \( n \times 1 \) vector (NAs allowed); \( \mu \) is an intercept; \( X_F, X_R, X_L \) and \( Z \) are incidence matrices used to accommodate different types of effects (see below), and; \( \varepsilon \) is a vector of
model residuals assumed to be distributed as \( \varepsilon \sim N(0, Diag(\sigma^2_\varepsilon/w_i^2)) \), here \( \sigma^2_\varepsilon \) is an (unknown) variance parameter and \( w_i \) are (known) weights that allow for heterogeneous-residual variances. Any of the elements in the right-hand side of the linear predictor, except \( \mu \) and \( \varepsilon \), can be omitted; by default the program runs an intercept model.

**Prior.** The residual variance is assigned a scaled inverse-\( \chi^2 \) prior with degree of freedom and scale parameter provided by the user, that is, \( \sigma^2_\varepsilon \sim \chi^{-2}(\sigma^2_\varepsilon|df_\varepsilon, S_\varepsilon) \). The regression coefficients \( \{\mu, \beta_F, \beta_R, \beta_L, u\} \) are assigned priors that yield different type of shrinkage. The intercept and the vector of regression coefficients \( \beta_F \) are assigned flat priors (i.e., estimates are not shrunk). The vector of regression coefficients \( \beta_R \) is assigned a Gaussian prior with variance common to all effects, that is, \( \beta_{R,j} \overset{iid}{\sim} N(0, \sigma^2_{\beta_R}) \). This prior is the Bayesian counterpart of Ridge Regression. The variance parameter \( \sigma^2_{\beta_R} \) is treated as unknown and it is assigned a scaled inverse-\( \chi^2 \) prior, that is, \( \sigma^2_{\beta_R} \sim \chi^{-2}(\sigma^2_{\beta_R}|df_{\beta_R}, S_{\beta_R}) \) with degrees of freedom \( df_{\beta_R} \) and scale \( S_{\beta_R} \) provided by the user. The vector of regression coefficients \( \beta_L \) is treated as in the Bayesian LASSO of Park and Casella (2008). Specifically,

\[
p(\beta_L, \tau^2, \lambda|\sigma^2_\varepsilon) = \prod_k N(\beta_{L,k}|0, \sigma^2_{\beta_L}^2) Exp\left(\frac{\tau^2}{\lambda}\right) \quad p(\lambda),
\]

where, \( Exp(\cdot) \) is an exponential prior and \( p(\lambda) \) can either be: (a) a mass-point at some value (i.e., fixed \( \lambda \)); (b) \( p(\lambda^2) \sim Gamma(r, \delta) \) this is the prior suggested by Park and Casella (2008); or, (c) \( p(\lambda) \propto \frac{1}{\lambda} \) (\( r > 1 \), \( \delta > 0 \)).

It can be shown that the marginal prior of regression coefficients \( \beta_{L,k} \), \( \int N(\beta_{L,k}|0, \sigma^2_{\beta_L}^2) Exp\left(\frac{\tau^2}{\lambda}\right) \partial \tau^2 \), is Double-Exponential. This prior has thicker tails and higher peak of mass at zero than the Gaussian prior used for \( \beta_R \), inducing a different type of shrinkage.

The vector \( u \) is used to model the so called ‘infinitesimal effects’, and is assigned a prior \( u \sim N(0, A\sigma^2_u) \), where, \( A \) is a positive-definite matrix (usually a relationship matrix computed from a pedigree) and \( \sigma^2_u \) is an unknown variance, whose prior is \( \sigma^2_u \sim \chi^{-2}(\sigma^2_u|df_u, S_u) \).

Collecting the above mentioned assumptions, the posterior distribution of model unknowns, \( \theta = \{\mu, \beta_F, \beta_R, \sigma^2_{\beta_R}, \beta_L, \tau^2, \lambda, u, \sigma^2_u, \sigma^2_\varepsilon\} \), is,

\[
p(\theta|y) \propto N\left(y|1\mu + X_F\beta_F + XR\beta_R + XL\beta_L + Zu; Diag\left(\frac{\sigma^2_\varepsilon}{w_i^2}\right)\right) \\
\times \prod_j N\left(\beta_{R,j}|0, \sigma^2_{\beta_R}\right) \chi^{-2}\left(\sigma^2_{\beta_R}|df_{\beta_R}, S_{\beta_R}\right) \\
\times \prod_k N\left(\beta_{L,k}|0, \sigma^2_{\beta_L}^2\right) Exp\left(\frac{\tau^2}{\lambda}\right) p(\lambda) \\
\times N(\mu|0, A\sigma^2_u)\chi^{-2}(\sigma^2_u|df_u, S_u) \chi^{-2}(\sigma^2_\varepsilon|df_\varepsilon, S_\varepsilon)
\]

**Value**

A list with posterior means, posterior standard deviations, and the parameters used to fit the model:

\$yHat \quad \text{the posterior mean of } 1\mu + X_F\beta_F + XR\beta_R + XL\beta_L + Zu + \varepsilon.

\$SD.yHat \quad \text{the corresponding posterior standard deviation.}

\$mu \quad \text{the posterior mean of the intercept.
the posterior mean of $\sigma^2_E$.

the posterior mean of $\beta_R$.

the corresponding posterior standard deviation.

the posterior mean of $\sigma^2_{\beta_R}$.

the posterior mean of $\beta_L$.

the corresponding posterior standard deviation.

the posterior mean of $\tau^2$.

the posterior mean of $\lambda$.

the posterior mean of $u$.

the corresponding posterior standard deviation.

the posterior mean of $\sigma^2_u$.

a list with evaluations of effective number of parameters and DIC (Spiegelhalter et al., 2002).

a vector indicating which entries in $y$ were missing.

a list containing the priors used during the analysis.

vector of weights.

list containing the following elements,

- $\logLikAtPostMean$: log-likelihood evaluated at posterior mean.
- $\postMeanLogLik$: the posterior mean of the Log-Likelihood.
- $pD$: estimated effective number of parameters, Spiegelhalter et al. (2002).
- $DIC$: the deviance information criterion, Spiegelhalter et al. (2002).

the number of iterations made in the Gibbs sampler.

the number of iterations used as burn-in.

the thin used.

original data-vector.

The posterior means returned by BLR are calculated after burnIn is passed and at a thin as specified by the user.

Save. The routine will save samples of $\mu$, variance components and $\lambda$ and running means (rm*.dat). Running means are computed using the thinning specified by the user (see argument thin above); however these running means are saved at a thinning specified by argument thin2 (by default, thin2=1 $\times$ $10^{10}$ so that running means are computed as the sampler runs but not saved to the disc).

Author(s)

Gustavo de los Campos, Paulino Perez Rodriguez,
References


Examples

```r
### Not run:
#.startTime

#Example 1:

rm(list=ls())
setwd(tempdir())
library(BGLR)
data(wheat)  #Loads the wheat dataset

y=wheat[,1]

## Creates a testing set with 100 observations
whichNa<sample(1:length(y),size=100,replace=FALSE)
yNa=y
yNa[whichNa]<NA

## Runs the Gibbs sampler
fm<-BLR(y=yNa, Xl=wheat.X, GF=list(ID=1:nrow(wheat.A), A=wheat.A),
prior=list(varE=list(df=3,S=0.25),
varU=list(df=3,S=0.63),
lambda=list(shape=0.52,rate=1e-4,
type="random",value=30)),
nIter=5500,burnIn=500,thin=1)

MSE.tst<mean((fm$yHat[whichNa]-y[whichNa])^2)
MSE.tst
MSE.trn<mean((fm$yHat[-whichNa]-y[-whichNa])^2)
MSE.trn
COR.tst<cor(fm$yHat[whichNa],y[whichNa])
COR.tst
COR.trn<cor(fm$yHat[-whichNa],y[-whichNa])
COR.trn

plot(fm$yHat,y,xlab="Phenotype",
     ylab="Pred. Gen. Value",cex=.8)
points(x=y[whichNa],y=fm$yHat[whichNa],col=2,cex=.8,pch=19)
x11()
plot(scan('var.E.dat'),type="o",
```
### Example 2: Ten fold, Cross validation, environment 1,

```r
rm(list=ls())
setwd(tempdir())
library(BGLR)
data(wheat)  #Loads the wheat dataset
nIter<1500   #For real data sets more samples are needed
burnIn<500
thin<10
folds<10
y<-wheat.Y[,1]
A<-wheat.A

priorBL<-list(
  varE=list(df=3,S=2.5),
  varU=list(df=3,S=0.63),
  lambda = list(shape=0.52,rate=1e-5,value=20,type='random')
)

set.seed(123)  #Set seed for the random number generator
sets<-rep(1:10,60)[-1]
sets<-sets[order(runif(nrow(A)))]
COR.CV<-rep(NA,n=nrow(COR.CV))
names(COR.CV)<c(paste('fold=',1:folds,sep='')),'Pooled')
w<-rep(1/nrow(A),folds)  ## weights for pooled correlations and MSE
yHatCV<-numeric()

for(fold in 1:folds)
{
  yNA<-y
  whichNa<-which(sets==fold)
yNA[whichNa]<-NA
  prefix<-paste('pm_bl','_fold_','fold','_sep=',sep='')
  fm<BLR(y=yNA,XL=wheat.X,GF=list(ID=(1:nrow(wheat.A)),A=wheat.A),prior=priorBL,
    nIter=nIter,burnIn=burnIn,thin=thin)
  yHatCV[whichNa]<-fm$yHat[fm$whichNa]
w[fold]<w[fold]*length(fm$whichNa)
  COR.CV[fold]<cor(fm$yHat[fm$whichNa],y[whichNa])
}

COR.CV
```

# End(Not run)
getVariances

Description
Computes the sample-variance (var()) for sets of markers as well as the total variance.

Usage

getVariances(X, B, sets, verbose=TRUE)

Arguments

- **X** (numeric, \(n \times p\)) incidence matrix for \(\beta\).
- **B** (numeric), object returned by the function readBinMat().
- **sets** (numeric).
- **verbose** (logical), if TRUE it shows progress information in the console.

Value
A matrix with variances for markers as well as the total.

Author(s)
Gustavo de los Campos.

Examples

```r
## Not run:
#Demos

library(BGLR)  
data(wheat)  
y=wheat.Y[,1]; X=scale(wheat.X)  
dir.create('test_saveEffects')  
setwd('test_saveEffects')  
fm=BGLR(y,y.ETA=list(list(X=X,model='BayesB',saveEffects=TRUE)),nIter=12000,thin=2,burnIn=2000)  
B=readBinMat('ETA_1_b.bin')  
plot(B[,1],type='o',col=4)  
VAR=getVariances(B=B,X=X,sets=sample(1:20,size=1279,replace=T))  
head(VAR)  
plot(VAR[,"total"])

## End(Not run)
```
Description

The mice data comes from an experiment carried out to detect and locate QTLs for complex traits in a mice population (Valdar et al. 2006a; 2006b). This data has already been analyzed for comparing genome-assisted genetic evaluation methods (Legarra et al. 2008). The data file consists of 1814 individuals, each genotyped for 10,346 polymorphic markers. The trait here here is body mass index (BMI), and additional information about body weight, season, month and day.

Usage

data(mice)

Format

Matrix mice.A contains the pedigree. The matrix mice.X contains the markes information and mice.pheno contains phenotypical information.

References


Description

Is a numerator relationship matrix (1814 x 1814) computed from a pedigree that traced back many generations.

References

mice.pheno

**Phenotypical data for the mice dataset**

**Description**

A data frame with phenotypical information related to diabetes. The data frame has several columns: SUBJECT.NAME, PROJECT.NAME, PHENOTYPE.NAME, Obesity.BMI, Obesity.BodyLength, Date.Month, Date.Year, Date.Season, cDate.StudyStartSeconds, Date.Hour, Date.StudyDay, GENDER, EndNormalBW, CoatColour, CageDensity, Litter, cage.

The phenotypes are described in http://gscan.well.ox.ac.uk.

mice.X

**Molecular markers**

**Description**

Is a matrix (1814 x 10346) with SNP markers.

plot.BGLR

**Plots for BGLR Analysis**

**Description**

Plots observed vs predicted values for objects of class BGLR.

**Usage**

```r
## S3 method for class 'BGLR'
plot(x, ...)```

**Arguments**

- `x` An object of class BGLR.
- `...` Further arguments passed to or from other methods.

**Author(s)**

Gustavo de los Campos, Paulino Perez Rodriguez,

**See Also**

BGLR.
Examples

## Not run:

```R
setwd(tempdir())
library(BGLR)
data(wheat)
out=BLR(y=wheat.Y[,1],X1=wheat.X)
plot(out)
```

## End(Not run)

### predict.BGLR

#### Model Predictions

**Description**

extracts predictions from the results of BGLR function.

**Usage**

```R
## S3 method for class 'BGLR'
predict(object, newdata, ...)
```

**Arguments**

- `object`: An object of class BGLR.
- `newdata`: Currently not supported, for new data you should assing missing value indicator (NAs) to the corresponding entries in the response vector (y).
- `...`: Further arguments passed to or from other methods.

**Author(s)**

Gustavo de los Campos, Paulino Perez Rodriguez,

**See Also**

BGLR.
Examples

## Not run:

```r
setwd(tempdir())
library(BGLR)
data(wheat)
out=BLR(y=wheat.Y[,1], XL=wheat.X)
predict(out)

## End(Not run)
```

### Description

Function to read effects saved by BGLR when ETA[[j]]$saveEffects=TRUE.

### Usage

```r
readBinMat(filename, byrow=TRUE)
```

### Arguments

- `filename` (string), the name of the file to be read.
- `byrow` (logical), if TRUE the matrix is created by filling its corresponding elements by rows.

### Value

A matrix with samples of regression coefficients.

### Author(s)

Gustavo de los Campos.

### Examples

## Not run:

```r
#Demos

library(BGLR)
data(wheat)
```

readBinMat
y=wheat.V[,1]; X=scale(wheat.X)
dir.create('test_saveEffects')
setwd('test_saveEffects')
fm=BGLR(y=y,ETA=list(X=X,model='BayesB'),saveEffects=TRUE),nIter=12000,thin=2,burnIn=2000)
B=readBinMat('ETA_1_b.bin')

## End(Not run)

### read_bed

**Description**

This function reads genotype information stored in binary PED (BED) files used in plink. These files save space and time. The pedigree/phenotype information is stored in a separate file (*.fam) and the map information is stored in an extended MAP file (*.bim) that contains information about the allele names, which would otherwise be lost in the BED file. More details [http://zzz.bwh.harvard.edu/plink/binary.shtml](http://zzz.bwh.harvard.edu/plink/binary.shtml).

### Usage

```r
read_bed(bed_file, bim_file, fam_file, na.strings, verbose)
```

### Arguments

- **bed_file**: binary file with genotype information.
- **bim_file**: text file with pedigree/phenotype information.
- **fam_file**: text file with extended map information.
- **na.strings**: missing value indicators, default=c("0","-9").
- **verbose**: logical, if true print hex dump of bed file.

### Value

The routine will return a vector of dimension n*p (n=number of individuals, p=number of snps), with the snps(individuals) stacked, depending whether the BED file is in SNP-major or individual-major mode.

The vector contains integer codes:

<table>
<thead>
<tr>
<th>Integer code</th>
<th>Genotype</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>00 Homozygote &quot;1&quot;/&quot;1&quot;</td>
</tr>
<tr>
<td>1</td>
<td>01 Heterozygote</td>
</tr>
<tr>
<td>2</td>
<td>10 Missing genotype</td>
</tr>
<tr>
<td>3</td>
<td>11 Homozygote &quot;2&quot;/&quot;2&quot;</td>
</tr>
</tbody>
</table>
Description

This function reads genotype information stored in PED format used in plink.

Usage

read_ped(ped_file)

Arguments

ped_file     ASCII file with genotype information.

Details

The PED file is a white-space (space or tab) delimited file: the first six columns are mandatory:
Family ID Individual ID Paternal ID Maternal ID Sex (1=male; 2=female; other=unknown) Pheno-
type
The IDs are alphanumeric: the combination of family and individual ID should uniquely identify a
person. A PED file must have 1 and only 1 phenotype in the sixth column. The phenotype can be
either a quantitative trait or an affection status column.

Value

The routine will return a vector of dimension n*p (n=number of individuals, p=number of snps),
with the snps stacked.

The vector contains integer codes:

<table>
<thead>
<tr>
<th>Integer code</th>
<th>Genotype</th>
</tr>
</thead>
<tbody>
<tr>
<td>0</td>
<td>00 Homozygote &quot;1/1&quot;</td>
</tr>
</tbody>
</table>
residuals.BGLR

1 01 Heterozygote
2 10 Missing genotype
3 11 Homozygote “2”/”2”

Author(s)

Gustavo de los Campos, Paulino Perez Rodriguez,

Examples

```r
## Not run:
library(BGLR)
demo(read_ped)

## End(Not run)
```

residuals.BGLR Extracts models residuals

Description

extracts model residuals from objects returned by BGLR function.

Usage

```r
## S3 method for class 'BGLR'
residuals(object, ...)
```

Arguments

- `object` An object of class BGLR.
- `...` Further arguments passed to or from other methods.

Author(s)

Gustavo de los Campos, Paulino Perez Rodriguez,

See Also

BGLR.
Examples

```r
## Not run:
setwd(tempdir())
library(BGLR)
data(wheat)
out=BLR(y=wheat.Y[,1],XL=wheat.X)
residuals(out)

## End(Not run)
```

summary.BGLR summary for BGLR fitted models

Description
Gives a summary for a fitted model using BGLR function.

Usage
```r
## S3 method for class 'BGLR'
summary(object, ...)
```

Arguments
- `object`: An object of class BGLR.
- `...`: Further arguments passed to or from other methods.

Author(s)
Gustavo de los Campos, Paulino Perez Rodriguez,

See Also
- BGLR.

Examples

```r
## Not run:
setwd(tempdir())
library(BGLR)
data(wheat)
out=BLR(y=wheat.Y[,1],XL=wheat.X)
summary(out)
```
Description

Information from a collection of 599 historical CIMMYT wheat lines. The wheat dataset is from CIMMYT’s Global Wheat Program. Historically, this program has conducted numerous international trials across a wide variety of wheat-producing environments. The environments represented in these trials were grouped into four basic target sets of environments comprising four main agro-climatic regions previously defined and widely used by CIMMYT’s Global Wheat Breeding Program. The phenotypic trait considered here was the average grain yield (GY) of the 599 wheat lines evaluated in each of these four mega-environments.

A pedigree tracing back many generations was available, and the Browse application of the International Crop Information System (ICIS), as described in http://repository.cimmyt.org/xmlui/bitstream/handle/10883/3488/72673.pdf (McLaren et al. 2000, 2005) was used for deriving the relationship matrix A among the 599 lines; it accounts for selection and inbreeding.

Wheat lines were recently genotyped using 1447 Diversity Array Technology (DArT) generated by Triticarte Pty. Ltd. (Canberra, Australia; https://www.diversityarrays.com). The DArT markers may take on two values, denoted by their presence or absence. Markers with a minor allele frequency lower than 0.05 were removed, and missing genotypes were imputed with samples from the marginal distribution of marker genotypes, that is, $x_{ij} = \text{Bernoulli}(\hat{p}_j)$, where $\hat{p}_j$ is the estimated allele frequency computed from the non-missing genotypes. The number of DArT MMs after edition was 1279.

Usage

data(wheat)

Format

Matrix Y contains the average grain yield, column 1: Grain yield for environment 1 and so on. The matrix A contains additive relationship computed from the pedigree and matrix X contains the markers information.

Source

International Maize and Wheat Improvement Center (CIMMYT), Mexico.
References


---

**wheat.A**

Pedigree info for the wheat dataset

**Description**

Is a numerator relationship matrix (599 x 599) computed from a pedigree that traced back many generations. This relationship matrix was derived using the Browse application of the International Crop Information System (ICIS), as described in [http://repository.cimmyt.org/xmlui/bitstream/handle/10883/3488/72673.pdf](http://repository.cimmyt.org/xmlui/bitstream/handle/10883/3488/72673.pdf) (McLaren et al. 2000, 2005).

**Source**

International Maize and Wheat Improvement Center (CIMMYT), Mexico.

**References**


---

**wheat.sets**

Sets for cross validation (CV)

**Description**

Is a vector (599 x 1) that assigns observations to 10 disjoint sets; the assignment was generated at random. This is used later to conduct a 10-fold CV.

**Source**

International Maize and Wheat Improvement Center (CIMMYT), Mexico.
wheat.X

Molecular markers

Description

Is a matrix (599 x 1279) with DArT genotypes; data are from pure lines and genotypes were coded as 0/1 denoting the absence/presence of the DArT. Markers with a minor allele frequency lower than 0.05 were removed, and missing genotypes were imputed with samples from the marginal distribution of marker genotypes, that is, \( x_{ij} = \text{Bernoulli}(\hat{p}_j) \), where \( \hat{p}_j \) is the estimated allele frequency computed from the non-missing genotypes. The number of DArT MMs after edition was 1279.

Source

International Maize and Wheat Improvement Center (CIMMYT), Mexico.

wheat.Y

Grain yield

Description

A matrix (599 x 4) containing the 2-yr average grain yield of each of these lines in each of the four environments (phenotypes were standardized to a unit variance within each environment).

Source

International Maize and Wheat Improvement Center (CIMMYT), Mexico.

write_bed

write_bed

Description

This function writes genotype information into a binary PED (BED) file used in plink. For more details about this format see [http://zzz.bwh.harvard.edu/plink/binary.shtml](http://zzz.bwh.harvard.edu/plink/binary.shtml).

Usage

```
write_bed(x, n, p, bed_file)
```
write_bed

Arguments

n integer, number of individuals.
p integer, number of SNPs.
x integer vector that contains the genotypic information coded as 0,1,2 and 3 (see details below). The information must be in snp major order. The vector should be of dimension n*p with the snps stacked.

bed_file output binary file with genotype information.

Details

The vector contains integer codes:

<table>
<thead>
<tr>
<th>Integer code</th>
<th>Genotype</th>
</tr>
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<tbody>
<tr>
<td>0</td>
<td>00 Homozygote &quot;1&quot;/&quot;1&quot;</td>
</tr>
<tr>
<td>1</td>
<td>01 Heterozygote</td>
</tr>
<tr>
<td>2</td>
<td>10 Missing genotype</td>
</tr>
<tr>
<td>3</td>
<td>11 Homozygote &quot;2&quot;/&quot;2&quot;</td>
</tr>
</tbody>
</table>

Author(s)

Gustavo de los Campos, Paulino Perez Rodriguez,

Examples

```r
## Not run:

library(BGLR)
demo(write_bed)

## End(Not run)```
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