

# Package ‘predhy’

August 16, 2021

**Type** Package

**Title** Genomic Prediction of Hybrid Performance

**Version** 1.2.0

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**Description** Performs genomic prediction of hybrid performance using eight GS methods including GBLUP, BayesB, RKHS, PLS, LASSO, Elastic net, Random forest and XG-Boost. It also provides fast cross-validation and mating design scheme for training population (Xu S et al (2016) <[doi:10.1111/tpj.13242](https://doi.org/10.1111/tpj.13242)>; Xu S (2017) <[doi:10.1534/g3.116.038059](https://doi.org/10.1534/g3.116.038059)>).

**License** GPL-3

**Encoding** UTF-8

**LazyData** true

**RoxygenNote** 7.1.1

**Depends** R (>= 3.6.0)

**Imports** BGLR, pls, glmnet, randomForest, xgboost, foreach, doParallel, parallel

**NeedsCompilation** no

**Repository** CRAN

**Date/Publication** 2021-08-16 16:40:02 UTC

## R topics documented:

convertgen . . . . .	2
crodesign . . . . .	3
cv . . . . .	4
cv_fast . . . . .	5
hybrid_phe . . . . .	6
infergen . . . . .	7
input_geno . . . . .	8
input_geno1 . . . . .	8
kin . . . . .	8
mixed . . . . .	9
predhy.predict . . . . .	10

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convertgen	<i>Convert Genotype</i>
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### Description

Convert genotypes in HapMap format or in numeric format for hypred package.

### Usage

```
convertgen(
  input_geno,
  type = c("hmp1", "hmp2", "num"),
  missingrate = 0.05,
  impute = TRUE
)
```

### Arguments

input_geno	genotype in HapMap format or in numeric format. The names of individuals should be provided. Missing (NA) values are allowed.
type	the type of genotype. There are three options: "hmp1" for genotypes in HapMap format with single bit, "hmp2" for genotypes in HapMap format with double bit, and "num" for genotypes in numeric format.
missingrate	max missing percentage for each SNP, default is 0.05.
impute	logical. If TRUE, imputation. Default is TRUE.

### Value

A matrix of genotypes in numeric format, coded as 1, 0, -1 for AA, Aa, aa. Each row represents an individual and each column represents a marker. The rownames of the matrix are the names of individuals.

### Examples

```
## load genotype in HapMap format with double bit
data(input_geno)

## convert genotype for hypred package
inbred_gen <- convertgen(input_geno, type = "hmp2")

## load genotype in numeric format
data(input_geno1)
head(input_geno1)

## convert genotype for hypred package
```

```
inbred_gen1 <- convertgen(input_gen01, type = "num")
```

---

crodesign

*Generate Mating Design*

---

## Description

Generate a mating design for a subset of crosses based on a balanced random partial rectangle cross-design (BRPRCD) (Xu et al. 2016).

## Usage

```
crodesign(d, male_name, female_name, seed = 123)
```

## Arguments

**d** an integer denoting 1/d percentage of crosses to be evaluated in the field.  
**male\_name** a character string for the names of male parents.  
**female\_name** a character string for the names of female parents.  
**seed** the random number, default is 123.

## Value

A data frame of mating design result with three columns. The first column is "crossID", the second column is the "male\_Name" and the third column is the "female\_Name".

## References

Xu S, Xu Y, Gong L and Zhang Q. (2016) Metabolomic prediction of yield in hybrid rice. *Plant J.* 88, 219-227.

## Examples

```
## generate a mating design with 100 male parents and 150 female parents
## for 1/d = 1/50 percentage of crosses to be evaluated in the field.
## the total number of potential crosses is 100*150 = 15000.
## The number of crosses to be field evaluated is 15000*(1/50) = 300.

male_name <- paste("m", 1:100, sep = "")
female_name <- paste("f", 1:150, sep = "")
design <- crodesign(d = 50, male_name, female_name)
```

**Description**

The `cv` function evaluates trait predictability based on eight GS methods via k-fold cross validation. The trait predictability is defined as the squared Pearson correlation coefficient between the observed and the predicted trait values.

**Usage**

```
cv(  
  fix = NULL,  
  gena,  
  gend = NULL,  
  y,  
  method = "GBLUP",  
  nfold = 5,  
  nTimes = 1,  
  seed = 123,  
  CPU = 1  
)
```

**Arguments**

<code>fix</code>	a design matrix of the fixed effects.
<code>gena</code>	a matrix (n x m) of additive genotypes for the training population.
<code>gend</code>	a matrix (n x m) of dominance genotypes for the training population. Default is NULL.
<code>y</code>	a vector(n x 1) of the phenotypic values.
<code>method</code>	eight GS methods including "GBLUP", "BayesB", "RKHS", "PLS", "LASSO", "EN", "XGBOOST", "RF". Users may select one of these methods or all of them simultaneously with "ALL". Default is "GBLUP".
<code>nfold</code>	the number of folds. Default is 5.
<code>nTimes</code>	the number of independent replicates for the cross-validation. Default is 1.
<code>seed</code>	the random number. Default is 123.
<code>CPU</code>	the number of CPU.

**Value**

Trait predictability

## Examples

```
## load example data from hybred package
data(hybrid_phe)
data(input_genos)

## convert original genotype
inbred_gen <- convertgen(input_genos, type = "hmp2")

##additive model infer the additive and dominance genotypes of hybrids
gena <- infergen(inbred_gen, hybrid_phe)$add
gend <- infergen(inbred_gen, hybrid_phe)$dom

##additive model
R2<-cv(fix=NULL,gena,gend=NULL,y=hybrid_phe[,3],method="GBLUP",nfold=5,nTimes=1,seed=123,CPU=1)

##additive-dominance model
R2<-cv(fix=NULL,gena,gend,y=hybrid_phe[,3],method="GBLUP",nfold=5,nTimes=1,seed=123,CPU=1)
```

---

 cv\_fast

*Evaluate Trait Predictability via the HAT Method*


---

## Description

The HAT method is a fast algorithm for the ordinary cross validation. It is highly recommended for large dataset (Xu et al. 2017).

## Usage

```
cv_fast(fix = NULL, y, kk, nfold = 5, seed = 123)
```

## Arguments

fix	a design matrix of the fixed effects. If not passed, a vector of ones is added for the intercept.
y	a vector of the phenotypic values.
kk	a list of one or multiple kinship matrices.
nfold	the number of folds, default is 5. For the HAT Method, nfold can be set as the sample size (leave-one-out CV) to avoid variation caused by random partitioning of the samples, but it is not recommended for <a href="#">cv</a> .
seed	the random number, default is 123.

## Value

Trait predictability

## References

Xu S. (2017) Predicted residual error sum of squares of mixed models: an application for genomic prediction. *G3 (Bethesda)* 7, 895-909.

## Examples

```
## load example data from hypred package
data(hybrid_phe)
data(input_geno)

## convert original genotype
inbred_gen <- convertgen(input_geno, type = "hmp2")

## infer the additive and dominance genotypes of hybrids
gena <- infergen(inbred_gen, hybrid_phe)$add
gend <- infergen(inbred_gen, hybrid_phe)$dom

## calculate the additive and dominance kinship matrix
ka <- kin(gena)
kd <- kin(gend)

##for the additive model
predictability <- cv_fast(y = hybrid_phe[,3], kk = list(ka))

##for the additive-dominance model
predictability <- cv_fast(y = hybrid_phe[,3], kk = list(ka,kd))
```

---

hybrid\_phe

*Phenotypic data of hybrids*

---

## Description

This dataset contains phenotypic data of 410 hybrids for grain yield in maize.

## Usage

```
hybrid_phe
```

## Format

A data frame with 410 rows and 3 variables:

M The names of male parents.

F The names of female parents.

GY The grain yield of hybrids.

---

infergen

*Infer Genotype of Hybrids*

---

## Description

Infer additive and dominance genotypes of hybrids based on their parental genotypes.

## Usage

```
infergen(inbred_gen, hybrid_phe)
```

## Arguments

inbred_gen	a matrix for genotypes of parental lines in numeric format, coded as 1, 0 and -1. The row.names of inbred_gen must be provided. It can be obtained from the original genotype using <a href="#">convertgen</a> function.
hybrid_phe	a data frame with three columns. The first column and the second column are the names of male and female parents of the corresponding hybrids, respectively; the third column is the phenotypic values of hybrids. The names of male and female parents must match the rownames of inbred_gen. Missing (NA) values are not allowed.

## Value

A list with following information is returned:

\$add additive genotypes of hybrids

\$dom dominance genotypes of hybrids

## Examples

```
## load example data from hypred package
data(hybrid_phe)
head(hybrid_phe)
data(input_gen)

## convert original genotype
inbred_gen <- convertgen(input_gen, type = "hmp2")

gena <- infergen(inbred_gen, hybrid_phe)$add
gend <- infergen(inbred_gen, hybrid_phe)$dom
```

---

input_geno	<i>Genotype in Hapmap Format</i>
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---

**Description**

Genotypic data of 348 maize inbred lines in Hapmap format with double bit.

**Usage**

input\_geno

**Format**

A data frame with 4979 rows and 359 columns.

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input_geno1	<i>Genotype in Numeric Format</i>
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**Description**

Genotypic data of 50 rice inbred lines with 1000 SNPs.

**Usage**

input\_geno1

**Format**

A data frame with 1000 rows and 50 variables.

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kin	<i>Calculate Kinship Matrix</i>
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**Description**

Calculate the additive and dominance kinship matrix.

**Usage**

kin(gen)

**Arguments**

gen	a matrix for genotypes, coded as 1, 0, -1 for AA, Aa, aa. Each row represents an individual and each column represents a marker.
-----	--



**Value**

a kinship matrix

**Examples**

```
## random population with 100 lines and 1000 markers
gen <- matrix(rep(0,100*1000),100,1000)
gen <- apply(gen,2,function(x){x <- sample(c(-1,0,1), 100, replace = TRUE)})

## generate 100*100 kinship matrix
k <- kin(gen)
```

---

mixed

*Solve Mixed Model*

---

**Description**

Solve linear mixed model using restricted maximum likelihood (REML). Multiple variance components can be estimated.

**Usage**

```
mixed(fix = NULL, y, kk)
```

**Arguments**

**fix** a design matrix of the fixed effects. If not passed, a vector of ones is added for the intercept.

**y** a vector of the phenotypic values.

**kk** a list of one or multiple kinship matrices.

**Value**

A list with following information is returned:

**\$v\_i** the inverse of the phenotypic variance-covariance matrix

**\$var** estimated variance components of genetic effects

**\$ve** estimated residual variance

**\$beta** estimated fixed effects

**References**

Xu S, Zhu D and Zhang Q. (2014) Predicting hybrid performance in rice using genomic best linear unbiased prediction. Proc. Natl. Acad. Sci. USA 111, 12456-12461.

## Examples

```
## load example data from hybred package
data(hybrid_phe)
data(input_genos)

## convert original genotype
inbred_gen <- convertgen(input_genos, type = "hmp2")

## infer the additive and dominance genotypes of hybrids
gena <- infergen(inbred_gen, hybrid_phe)$add
gend <- infergen(inbred_gen, hybrid_phe)$dom

## calculate the additive and dominance kinship matrix
ka <- kin(gena)
kd <- kin(gend)

## for the additive model
parm <- mixed(y = hybrid_phe[,3], kk = list(ka))

## for the additive-dominance model
parm <- mixed(y = hybrid_phe[,3], kk = list(ka, kd))
```

---

predhy.predict

*Predict the Performance of Hybrids*

---

## Description

Predict all potential crosses of a given set of parents using a subset of crosses as the training sample.

## Usage

```
predhy.predict(
  inbred_gen,
  hybrid_phe,
  method = "GBLUP",
  model = "A",
  select = "top",
  number = "100"
)
```

## Arguments

**inbred\_gen** a matrix for genotypes of parental lines in numeric format, coded as 1, 0 and -1. The row.names of inbred\_gen must be provided. It can be obtained from the original genotype using [convertgen](#) function.

hybrid_phe	a data frame with three columns. The first column and the second column are the names of male and female parents of the corresponding hybrids, respectively; the third column is the phenotypic values of hybrids. The names of male and female parents must match the rownames of inbred_gen. Missing (NA) values are not allowed.
method	eight GS methods including "GBLUP", "BayesB", "RKHS", "PLS", "LASSO", "EN", "XGBOOST", "RF". Users may select one of these methods. Default is "GBLUP".
model	the prediction model. There are two options: model = "A" for the additive model, model = "AD" for the additive-dominance model. Default is model = "A".
select	the selection of hybrids based on the prediction results. There are three options: select = "all", which selects all potential crosses. select = "top", which selects the top n crosses. select = "bottom", which selects the bottom n crosses. The n is determined by the param number.
number	the number of selected top or bottom hybrids, only when select = "top" or select = "bottom".

**Value**

a data frame of prediction results with two columns. The first column denotes the names of male and female parents of the predicted hybrids, and the second column denotes the phenotypic values of the predicted hybrids.

**Examples**

```
## load example data from hypred package
data(hybrid_phe)
data(input_geno)
inbred_gen <- convertgen(input_geno, type = "hmp2")

## infer the additive and dominance genotypes of hybrids
gena <- infergen(inbred_gen, hybrid_phe)$add
gend <- infergen(inbred_gen, hybrid_phe)$dom

pred<-predhy.predict(inbred_gen,hybrid_phe,method="LASSO",model="A",select="top",number="100")
pred<-predhy.predict(inbred_gen,hybrid_phe,method="LASSO",model="AD",select="all")
```

# Index

## \* datasets

- hybrid\_phe, 6
- input\_gen0, 8
- input\_gen01, 8

convertgen, 2, 7, 10

crodesign, 3

cv, 4, 5

cv\_fast, 5

hybrid\_phe, 6

infergen, 7

input\_gen0, 8

input\_gen01, 8

kin, 8

mixed, 9

predhy.predict, 10