ELPGV Manual

Ensemble learning for integrative prediction of genetic values with genomic variants.

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1 Brief introduction

Whole genome variants offer sufficient information for genetic prediction of human disease, and animal/plant breeding values that has been widely accepted for animal breeding. Many sophisticate statistical algorithms have been developed for enhancing the prediction accuracy; however, each method has its own advantages and disadvantages, so far no method can beat anther. We herein propose a new ensemble learning strategy, called ELPGV (*Ensemble Learning of Prediction for Genetic Value*) for genetic prediction, which assembles predictions from several basic methods, to more accurate predictions.

2 ELPGV function

ELPGV

Ensemble learning for integrative prediction of genetic values

2.1 Description

The ELPGV (Ensemble Learning of Prediction for Genetic Value) function is ensemble learning for integrative prediction of genetic values with genomic variants.

2.2 Usage

ELPGV(rep_times = 100, interation_times=20, weight_min=0, weight_max=1, weight_dimension=ncol(test_PredMat), rate_min=-0.01, rate_max=0.01, paticle_number=20, CR = 1.0, train_PredMat=as.matrix(train_PredMat), test_PredMat=as.matrix(test_PredMat), R = 0.5, IW = 1, AF1 = 2, AF2 = 2, type = "pcc", select="auto", index=NULL)

2.3 Arguments

rep_times	Number of repetitions of ELPGV algorithm, default 100.				
interation_times	The number of ELPGV iterations within each repetition, default 20.				
weight_min	Minimum value of training weight range. The range is from 0 to 1, the default value is 0.				
weight_max	Maximum value of training weight range. The range is from 0 to 1, the default value is 1.				
weight_dimension	The number of basic learners for ensemble learning.				
rate_min	The minimum value of the rate range of particle swarm optimization, the default value is -0.01.				

rate_max The maximum value of the rate range of particle swarm

optimization, the default value is 0.01.

paticle_number Initial population number of particle swarm optimization

and differential evolution algorithm. The range is 5N to 10N, where N is the number of basic learners for ensemble

learning.

CR Crossover probability of differential evolution algorithm,

the default value is 1.

train_PredMat Prediction matrix for training group, with Phenotype.

test_PredMat Prediction matrix for testing group, without Phenotype.

R Mutation probability of differential evolution algorithm.

Inertia weight of particle swarm optimization, the default

value is 0.5.

AF1 The first acceleration factor of particle swarm

optimization, the default value is 2.

AF2 The second acceleration factor of particle swarm

optimization, the default value is 2.

type The evaluating indicator, "pcc" is Pearson correlation

coefficient.

select User chooses the way to generate reference phenotype

values, setting as either 'auto' or 'manu', if 'auto' is chosen, parameter index (see below) should be chosen

accordingly.

index Indicating which prediction is taken as reference

phenotype, f.i., if 3th method (3th columns) is chosen, parameter select='manu' and index=3; otherwise, setting

"auto" select: the index is NULL.

2.4 Value

IW

The pred Prediction value of all models.

3 Build in data

An example datasets 'exdata' is a list that including the milk yield datasets. The mkg list including the prediction matrix of Training set(train) and the prediction matrix of Testing set(test). The train list including the real phenotype value for each individual(obs), the predicted phenotype value of the BayesA model for each individual(BayesA), the predicted phenotype value of the BayesCpai model for each individual(BayesCpai) and the predicted phenotype value of the GBLUP model for each individual(GBLUP). The test list structure is the same as the train list, 'exdata'

can be loaded with data(exdata).

```
List of 2
train: 'data.frame': 452 obs. of 5 variables:
..$ obs : num [1:452] 0.755 1.606 -0.805 1.431 0.401 ...
..$ BayesA : num [1:452] -0.182 0.752 -0.554 1.379 0.192 ...
..$ BayesB : num [1:452] -0.265 0.742 -0.478 1.425 0.187 ...
..$ BayesCpai: num [1:452] -0.176 0.813 -0.467 1.459 0.184 ...
..$ GBLUP : num [1:452] -0.169 0.77 -0.652 1.409 0.171 ...
test :'data.frame': 50 obs. of 5 variables:
..$ obs : num [1:50] -1.0159 0.3105 0.0128 0.4025 0.9542 ...
..$ BayesA : num [1:50] -0.943 0.8 -0.449 -0.259 0.202 ...
..$ BayesB : num [1:50] -0.971 0.675 -0.393 -0.142 0.167 ...
..$ BayesCpai: num [1:50] -0.985 0.729 -0.41 -0.137 0.222 ...
..$ GBLUP : num [1:50] -1.177 0.793 -0.41 -0.363 0.178 ...
```

3.1 Running build-in data

library("ELPGV") data(exdata) train_PredMat = mkg\$train View(train_PredMat)

obs [‡]	BayesA [‡]	BayesB [‡]	BayesCpai [‡]	GBLUP [‡]
0.754697	-0.181631580	-0.264827571	-0.176131795	-0.168798907
1.605698	0.752134605	0.742498663	0.813063593	0.769769482
-0.805472	-0.554030372	-0.478379983	-0.466988176	-0.651766990
1.431134	1.379331103	1.425064252	1.458779803	1.409240915
0.400892	0.191647279	0.186696803	0.184477310	0.170519354
-0.627791	-0.914932829	-1.023727364	-1.000427061	-0.899369627

test PredMat = mkg test[,-1]

BayesA

TBV = mkg stest[,1]View(test_PredMat)

True breeding value of testing group

GBLUP

BayesCpai -0.94305106 -0.97133040 -0.98532235 -1.17745944 0.79996260 0.67490873 0.72949490 0.79319091 -0.44917172 -0.39293983 -0.40961475 -0.41031868 -0.25904775 | -0.14195774 | -0.13735587 -0.36286804

BayesB

0.20197808 0.16738521 0.22187015 0.17752197 0.30662630 0.25471571 0.25984270 0.14490990

ELPGV_pred = ELPGV(rep_times = 100, interation_times=20, weight_min=0, weight_max=1, weight_dimension=ncol(test_PredMat), rate_min=-0.01, rate_max=0.01, paticle_number=20,

CR = 1.0, train_PredMat=as.matrix(train_PredMat), test_PredMat=as.matrix(test_PredMat), R = 0.5, IW = 1, AF1 = 2, AF2 = 2, type ="pcc", select="auto", index=NULL)

View(ELPGV_pred)

BayesA [‡]	BayesB [‡]	BayesCpai [‡]	GBLUP [‡]	ELPGV [‡]
-0.94305106	-0.97133040	-0.98532235	-1.17745944	-0.98287785
0.79996260	0.67490873	0.72949490	0.79319091	0.70533835
-0.44917172	-0.39293983	-0.40961475	-0.41031868	-0.40295220
-0.25904775	-0.14195774	-0.13735587	-0.36286804	-0.16423402
0.20197808	0.16738521	0.22187015	0.17752197	0.18320268
0.30662630	0.25471571	0.25984270	0.14490990	0.25486829

head(PredMat)

```
> PredMat<-cbind(TBV,ELPGV_pred)
> colnames(PredMat)<-c("obs","BayesA", "BayesB", "BayesCpai", "GBLUP", "ELPGV")</p>
> head(PredMat)
         obs
                BayesA
                         BayesB BayesCpai
                                             GBLUP
[1,] -1.015885 -0.9430511 -0.9713304 -0.9853224 -1.1774594 -0.9828779
[2,] 0.310493 0.7999626 0.6749087 0.7294949 0.7931909 0.7053383
    [4,]
   0.954199 0.2019781 0.1673852 0.2218702 0.1775220 0.1832027
[5,]
[6,] -0.004347 0.3066263 0.2547157
                                0.2598427
                                          0.1449099 0.2548683
```

If the observable value of testing population is given, f.i. in the situation of cross validation, one can evaluates the prediction accuracy by calculating the correlation coefficient between prediction and observable value.

```
> PreMat <- cbind(TBV,ELPGV_pred)</p>
> colnames(PreMat) <- c("obs","BayesA","BayesB","BayesCpai","GBLUP","ELPGV")</p>
> cor(PreMat)
                obs
                                 BayesB BayesCpai
                       BayesA
          1.0000000 0.7811417 0.7813917 0.7777354 0.7680360 0.7815003
obs
          0.7811417 1.0000000 0.9913966 0.9936144 0.9883919 0.9946540
BayesA
          0.7813917 0.9913966 1.0000000 0.9966254 0.9776841 0.9993514
BayesB
BayesCpai 0.7777354 0.9936144 0.9966254 1.0000000 0.9848915 0.9985654
          0.7680360 0.9883919 0.9776841 0.9848915 1.0000000 0.9835819
          0.7815003 0.9946540 0.9993514 0.9985654 0.9835819 1.0000000
ELPGV
```

3.2 Quick running your data

We also provide external dataset for testing, which can be accessed from "exa mpleFile" of ELPGV package.

```
library("ELPGV")
```

#loading training predictions

train_PredMat = read.table(system.file("exampleFile/exampleTrainingFile.txt",

```
head(train_PredMat)
 library(ELPGV)
 library("ELPGV")
#loading training predictions
train_PredMat <- read.table(system.file("exampleFile/exampleTrainingFile.txt", package = "ELPGV"),header = T)
 head(train_PredMat)

        obs
        BayesA
        BayesB
        BayesCpai
        GBLUP

        -0.656353
        0.03053361
        -0.121156067
        -0.27331072
        -0.1710210

        -2.290264
        -1.20102098
        -1.129217026
        -1.01576577
        -1.1698250

        0.728920
        0.40946229
        0.426954496
        0.40397526
        0.4170243

 -1.153630 -0.97216542 -0.988925786 -1.21599601 -1.0371759
 0.444761 0.91462040 0.805147269 0.53904085
0.515801 0.16333984 -0.003664033 -0.04268064
                                                  0.6166968
                                                  0.3519748
#loading testing predictions
test_PredMat = read.table(system.file("exampleFile/exampleTestingFile.txt",
                                                           package = "ELPGV"), header = T)
head(test_PredMat)
  test_PredMat <- read.table(system.file("exampleFile/exampleTestingFile.txt", package = "ELPGV"),header = T)
 head(test_PredMat)
      BayesA
                 BayesB BayesCpai
                                             GBLUP
  -0.5767065 -0.3802978 -0.3672191 -0.87568361
  -0.3767065 -0.3802978 -0.3672191 -0.87368361
-1.0239310 -0.9226303 -0.7153346 -0.62825170
0.8821122 0.8496787 0.9319635 0.78240236
-0.2768977 -0.2753204 -0.2430209 0.01691417
-0.3853227 -0.5162605 -0.5731072 -0.14592593
   0.6471890 0.5502385 0.3541639 0.30823536
ELPGV_pred = ELPGV(rep_times = 100, interation_times=20, weight_min=0,
                                    weight max=1, weight dimension=ncol(test PredMat),
                                    rate min=-0.01, rate max=0.01, paticle number=20,
                                    CR = 1.0, train_PredMat=as.matrix(train_PredMat),
                                    test_PredMat=as.matrix(test_PredMat), R = 0.5,
                                    IW = 1, AF1 = 2, AF2 = 2, type ="pcc", select="auto",
                                    index=NULL)
colnames(ELPGV_pred) = c("BayesA", "BayesB", "BayesCpai",
                                         "GBLUP", "ELPGV")
head(ELPGV_pred)
 > colnames(ELPGV_pred)<-c("BayesA","BayesB","BayesCpai","GBLUP","ELPGV")</pre>
 > head(ELPGV_pred)
                                 BayesB BayesCpai
                                                                      GBLUP
               BayesA
                                                                                       ELPGV
  [1,] -0.5767065 -0.3802978 -0.3672191 -0.87568361 -0.4281360
  [2,] -1.0239310 -0.9226303 -0.7153346 -0.62825170 -0.8926898
  [3,] 0.8821122 0.8496787 0.9319635 0.78240236 0.8679455
  [4,] -0.2768977 -0.2753204 -0.2430209 0.01691417 -0.2600380
  [5,] -0.3853227 -0.5162605 -0.5731072 -0.14592593 -0.4917666
         0.6471890 0.5502385 0.3541639 0.30823536 0.5233226
```

4 Additional explanation of parameter 'select' and 'index'

ELPGV needs to compare the prediction accuracies of training set of different methods first and chooses the best one as reference phenotypic values, by which we are able to evaluate the prediction accuracies for testing set. We herein provide two ways to generate reference phenotype values, (I) auto (automatic) and (II) manu (manual). The "auto" function chooses the best prediction method as reference automatically, whereas the 'manu' function requires users to choose the best one manually, which needs user's experience. 'auto' function is recommended for general users, and "manu" is for advanced users. The example is below, if we need to ensemble four basic models, such as BayesA, BayesB, BayesC π and GBLUP, we will

have the predictions of four methods for training set. Among these predictions, users need to choose the best method with the most accurate predictions as input (reference phenotypic value) for prediction of test set. In this example, the prediction accuracy of BayesCpai is the best and chosen as reference phenotype values. If "auto" function is selected, ELPGV will choose the best method (BayesCpai in this case) automatically (select="auto" and index="NULL"); but if "manu" function is selected, users can choose any method freely (select="manu" and index=3, if the third method or the third column is the chosen). It is noted that the performance of ELPGV will rely on the "best method" for training, users should choose the function with caution, if not sure which method is the best, "auto" function is recommended.

I. auto

```
ELPGV(rep_times = 100, interation_times=20, weight_min=0, weight_max=1, weight_dimension=ncol(test_PredMat), rate_min=-0.01, rate_max=0.01, paticle_number=20, CR = 1.0, train_PredMat=as.matrix(train_PredMat), test_PredMat=as.matrix(test_PredMat), R = 0.5, IW = 1, AF1 = 2, AF2 = 2, type = "pcc", select="auto", index=NULL)
```

II. manu

```
ELPGV(rep_times = 100, interation_times=20, weight_min=0, weight_max=1, weight_dimension=ncol(test_PredMat), rate_min=-0.01, rate_max=0.01, paticle_number=20, CR = 1.0, train_PredMat=as.matrix(train_PredMat), test_PredMat=as.matrix(test_PredMat), R = 0.5, IW = 1, AF1 = 2, AF2 = 2, type = "pcc", select="manu", index=3)
```

5 Code availability

The source code of ELPGV is freely available on request.

6 How to access help

If users have any bugs or issues or any suggestions are available, feel free to contact: Linlin Gu: gulinlin 141006@163.com

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