Manual for

BLUPF90 family of programs

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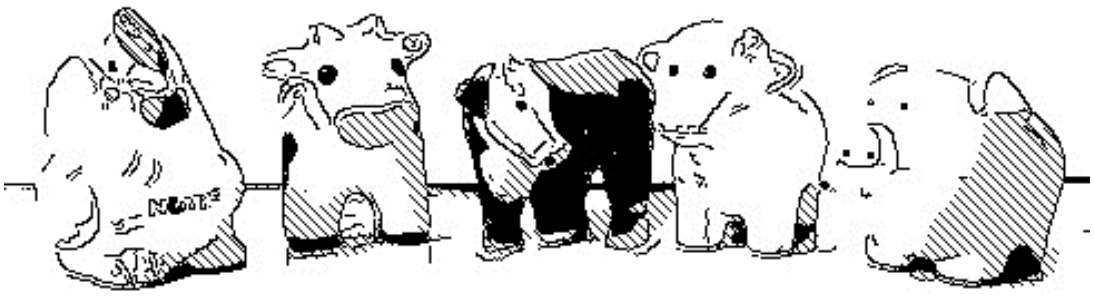

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Introduction

BLUPF90 is a family of programs for mixed-model computations with focus on animal breeding applications. The programs can do data conditioning, estimate variances using several methods, calculate BLUP for very large data sets, calculate approximate accuracy, and use SNP information for improved accuracy of breeding values + for genome-wide association studies (GWAS).
The programs have been designed with 3 goals in mind:

1. Flexibility to support a large set of models found in animal breeding applications.
2. Simplicity of software to minimize errors and facilitate modifications.
3. Efficiency at the algorithmic level.

Aside from being used in hundreds of studies, the programs are utilized for commercial genetic evaluation in dairy, beef, pigs and broiler chicken by major companies/institutions/associations in the US and beyond.

The programs are written in Fortran 90/95 and originated as exercises for a class taught by Ignacy Misztal at the University of Georgia. Over time, they have been upgraded and enhanced by many contributors. Details on programming and computing algorithms are available in an Interbull 1999 **paper** and as course notes. Nearly all programs are available in source code.

More information about the programs is available at **http://nce.ads.uga.edu/wiki/doku.php** as wiki pages.

Issues with the programs are discussed in group blupf90 at groups.yahoo.com.

List of programs from Wiki page

Latest versions available from website at
http://nce.ads.uga.edu/wiki/doku.php?id=application_programs
(Use latest versions. All applications for Linux, Mac OSX, and Windows have been updated frequently)

The programs support mixed models with multiple-correlated effects, multiple animal models and dominance.

- BLUPF90 - BLUP in memory blupf90.pdf
- REMLF90 - accelerated EM REML remlf90.pdf
- QXPAK - joint analysis of QTL and polygenic effects (M. Perez-Enciso) QxPak web page
- AIREMLF90 - Average Information REML with several options including EM-REML and heterogeneous residual variances (S. Tsuruta)
- CBLUP90 - solutions for bivariate linear-threshold models
- CBLUP90THR - as above but with thresholds computed and many linear traits (B. Auvray)
- CBLUP90REML - as above but with quasi REML (B. Auvray)
- GIBBSF90 - simple block implementation of Gibbs sampling
- GIBBS1F90 - as above but faster for creating mixed model equations only once
- GIBBS2F90 - as above but with joint sampling of correlated effects
- GIBBS3F90 - as above with support for heterogeneous residual variances
- POSTGIBBSF90 - statistics and graphics for post-Gibbs analysis (S. Tsuruta)
- THRGIBBSF90 - Gibbs sampling for any combination of categorical and linear traits (D. Lee)
- THRGIBBS1F90 – as above but simplified with several options (S. Tsuruta)
- RENUMF90 - a renumbering program that also can check pedigrees and assign unknown parent groups; supports large data sets
- INBUPGF90 - a program to calculate inbreeding coefficients with incomplete pedigree (I. Aguilar)

Available by request

- MRF90 - Method R program suitable for very large data sets; contact T. Druet.
- COXF90 – Bayesian Cox model - contact J. P. Sanchez (JuanPablo.Sanchez@irta.cat)
- BLUPF90HYP – BLUPF90 with hypothesis testing (F and Chi2 tests) - contact J. P. Sanchez as above

Available only under research agreement

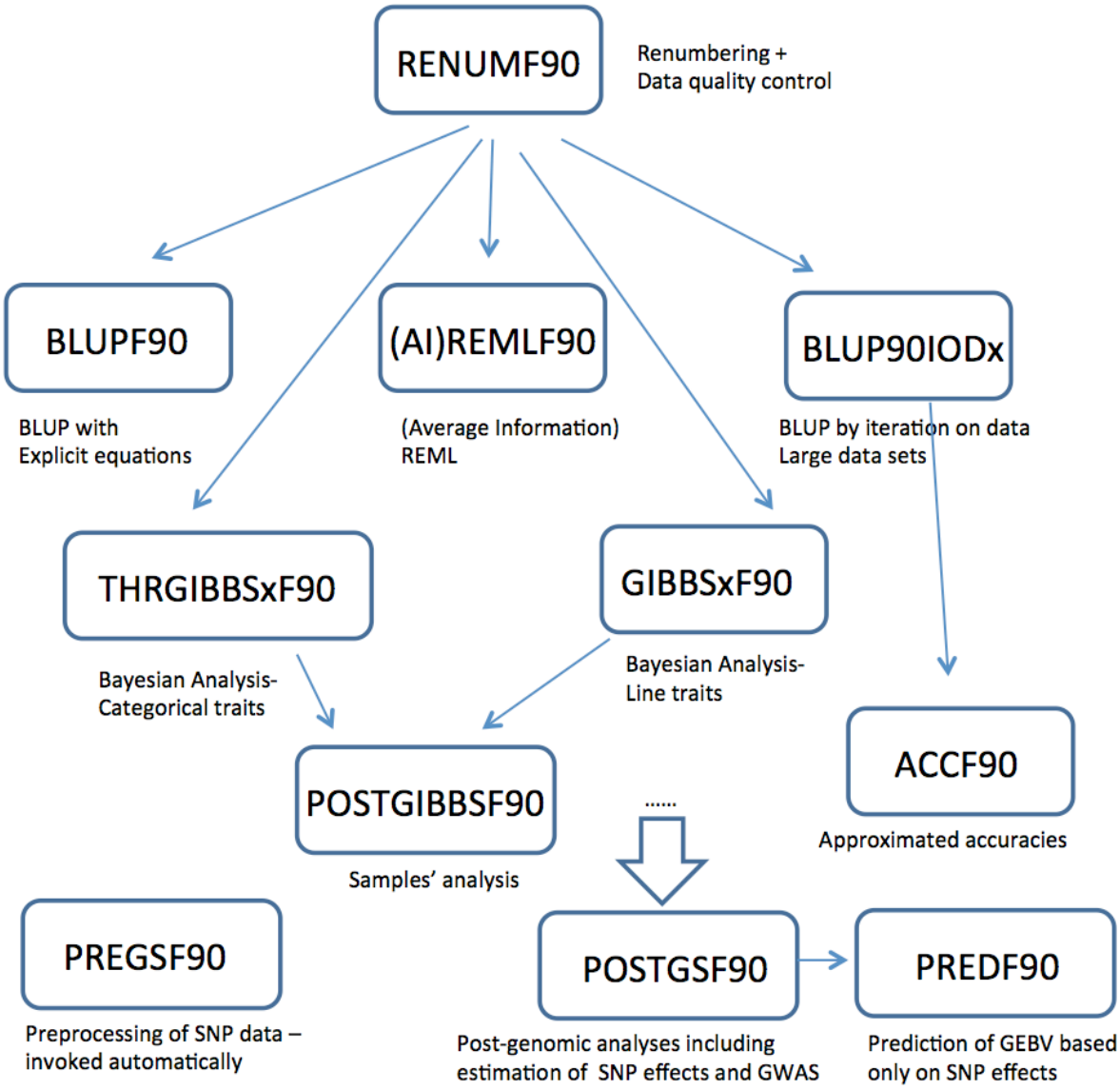
- BLUP90IOD2 - BLUP by iteration on data with support for very large models (S. Tsuruta)
- CBLUP90IOD - BLUP by iteration on data for threshold-linear models
- ACCF90 - approximation of accuracies for breeding values
- BLUP90MBE - BLUP by iteration on data with support for very large models for multi-breed evaluations
- BLUP90ADJ - BLUP data preadjustment tool

Included in application programs

- PREGSF90 – genomic preprocessor that combines genomic and pedigree relationships (I. Aguilar)
- POSTGSF90 – genomic postprocessor that extracts SNP solutions after genomic evaluations (single step, GBLUP) (I. Aguilar)

Other programming contributions were made by Miguel Perez-Enciso `(user_file)` and François Guillaume (Jenkins hashing functions).

Programs in a chart



Application programs (BLUP*, *REMLF90, THRGIBBS* and GIBBS*) are driven by parameter files and require data files with effects renumbered from 1 consecutively.

Renumbering and quality control can be done by RENUMF90, which is also driven by a parameter file. Separation of renumbering and application programs allows supporting complicated models.

Some models are not directly supported by RENUMF90 and require tweaking the parameter file in the application programs.

Parameter file for application programs

The parameter file has keywords that are fixed and cannot be changed followed by values, with the following structure:

Keywords*	Description
DATAFILE	Name of file with phenotypes; free format
file.dat	
NUMBER OF TRAITS	Number of traits
2	
NUMBER OF EFFECTS	Number of effects in a model except for residual
6	
OBSERVATIONS(S)	Position(s) of observations in data file
1 2	
WEIGHTS	Position of weight on observations if used; otherwise blank
2	"**2**" means that residual variance (R) is set to R/**2**.
EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]	
4 4 10 cross	**4 4** = crossclassified effect positions in data file for 2 traits; **10** = levels
5 0 100 cross	**5 0** = crossclassified effect, positions for 2 traits; **100** = levels
6 6 1 cov	**6 6** = covariable positions in data file
7 7 10 cov 4 4	**7 7** = covariable nested in effect position **4**; **10** = levels
8 8 1000 cross	**8 8** = crossclassified effect positions for 2 traits; **1000** = levels
0 9 1000 cross	**0 9** = crossclassified effect positions for 2 traits; **1000** = levels
RANDOM_RESIDUAL_VALUES	Residual variance or residual covariance matrix
10 1	For 2 trait model
1 10	
RANDOM_GROUP	List of effect numbers that form a group
5 6	For correlated random effects **5 6**
RANDOM_TYPE	Type of random effect (distribution)
add_animal	**diagonal**, **add_sire**, **add_an_upg**, **add_an_upginb**, **par_domin**, or **user_file**
FILE	Pedigree file or other file associated with random effect; blank if none
file.ped	
(CO)VARIANCES	(Co)variance matrix for each random effect
10 1	For 2 trait model
1 10	

***Misspelling causes the programs to crash.**

Description of effects

The effects are specified after the keyword:

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]

Each line contains the following:

- Position(s) of each effect in the data file; t positions for t traits
- Number of levels (assumed consecutive from 1)
- Type of effect: “cross” for crossclassified, and “cov” for covariable
    - crossclassified uses integer number from 1
    - covariable uses integer or real numbers
- For nested covariables, the following number (or t numbers for t traits) indicates the position of nesting in the data file
- Text after # can be used as a comment

Consider a data file (file.dat) with the following columns

i	j	k	y1	y2	x1
2	2	3	4.30	5.67	22.40
1	2	2	2.76	3.20	18.00
	.....				
3	1	1	2.20	5.30	7.25

Let i go from 1 to 50, j from 1 to 80, and k from 1 to 200. The model:

$$y1_{ij}=a_j+b_i+cX+e_{ij}$$

will be specified in the parameter file as:

```
DATAFILE
file.dat
NUMBER_OF_TRAITS
1
NUMBER_OF_EFFECTS
3
OBSERVATIONS(S)
4
WEIGHTS

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
2 80 cross   # position 2, 80 levels
1 50 cross   # position 1, 50 levels
3 1 cov      # covariable on position 3, one level
……
```

By definition, a regular covariable has one level (i.e., a slope as regression).

For a similar model but with a nested covariable:

$$y1_{ij}=a_j+b_i+c_iX+e_{ij}$$

The description will change to:

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
2 80 cross # position 2, 80 levels
1 50 cross # position 1, 50 levels
3 50 cov 1 # covariable on position 3 nested in position 1; 50 levels

Assume a two trait model:

$$y1_{ij}=a1_j+\quad c1_iX+e1_{ij}$$
$$y2_{ij}=\quad b2_i+c2_iX+e2_{ij}$$

This corresponds to:

……
NUMBER_OF_TRAITS
2
NUMBER_OF_EFFECTS
3
……
EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
2 0 80 cross # position 2 for trait 1 only, 80 levels
0 1 50 cross # position 1 for trait 2 only, 50 levels
3 3 50 cov 1 1 # covariable on position 3 for two traits nested in position 1

"0" in effect definitions means missing effect per trait.

Two effects above can be merged:

NUMBER_OF_EFFECTS
2
……
EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
2 1 80 cross # positions 2 and 1 for traits 1 and 2, 80 is max(50,80)levels
3 3 50 cov 1 1 # covariable on position 3 for two traits nested in position 1

Definition of random effects

RANDOM_GROUP defines one group of random effects. A group is one effect or multiple (correlated) effects that share the same covariance structure, e.g., direct-maternal effect or random regressions.

The structure of **RANDOM GROUP** is:

RANDOM_GROUP
5
or
5 6

Corresponding to the effect number specified above; "**5**" means that the 5th effect is random. Or "**5 6**" means that 5th and 6th are correlated random effects.

RANDOM_TYPE defines a covariance structure: diagonal var() = *s* ⊗ **I** or **G** where *s* is a variance and **G** is a covariance matrix. For other types, see "Random effects and Pedigree files"

Assume a model:

y = farm + animal_additive + animal_environment + error

with var(animal_additive) = 2.5⊗**A**, var(animal_environment) = 5.1⊗**I**; var(error) = 13.7⊗**I**

With these effects:

```
EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
3   100 cross    # effect 1: farm
2  1000 cross   # effect 2: additive genetic
2  1000 cross   # effect 3: permanent environment
RANDOM_RESIDUAL_VALUES
13.7
RANDOM_GROUP
2                    # this is for effect 2 on the effect list
RANDOM_TYPE
add_animal           # additive genetic
FILE
file.ped    # name of pedigree file
(CO)VARIANCES
2.5
RANDOM_GROUP
3                    # effect 3 on the effect list above
RANDOM_TYPE
diagonal  # permanent environment
FILE
                     # no file associated with diagonal structures
(CO)VARIANCES
5.1
```

Correlated effects

Assume a model:

y = farm + season + direct + maternal + error

var(direct,maternal)= $\begin{bmatrix} 5 & 1 \\ 1 & 6 \end{bmatrix} \otimes$ A

with the effects as specified:

```
EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
3    100 cross        # effect 1: farm
4      4 cross        # effect 2: season
2  1000 cross        # effect 3: direct
2  1000 cross        # effect 3: maternal
```

The distribution of the random effects are specified below:

```
…
RANDOM_GROUP
3  4                 # direct and maternal effects
RANDOM_TYPE
add_animal           # additive genetic
FILE
file.ped             # name of pedigree file
(CO)VARIANCES
5  1
1  6
…
```

Random regression models may have many correlated random effects. Assume a data file with the following positions:

1 to 4: polynomials

5: animal number (1000 levels)

6: herd year season (50 levels)

```
…
EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
6 50 cross           # herd year season
1 1000 cov 5         # first polynomial nested within the animal effect position 5
2 1000 cov 5         # second polynomial nested within the animal effect position 5
3 1000 cov 5         # third polynomial nested within the animal effect position 5
4 1000 cov 5         # fourth polynomial nested within the animal effect position 5
….
RANDOM_GROUP
2 3 4 5              # all covariables are correlated
RANDOM_TYPE
add_animal           # additive genetic
FILE
file.ped             # name of pedigree file
(CO)VARIANCES
  (4 x 4 matrix)
```

There are a few types of additive genetic effects, each with a different pedigree format.

a) additive sire (add_sire)

The pedigree file has the following format:
sire number, sire's sire number, sire's maternal grandsire (MGS) number
where unknown sire's sire and/or sire's MGS numbers are replaced by 0.

b) additive animal (add_animal)

The pedigree file has the following format:
animal number, sire number, dam number
where unknown sire and/or dam numbers are replaced by 0.

c) additive animal with unknown parent groups (add_an_upg)

The pedigree file has the following format:
animal number, sire number, dam number, parent code
where sire and/or dam numbers can be replaced by unknown parent group numbers
parent code = 3 - number of known parents:

1 (both parents known)
2 (one parent known)
3 (both parents unknown)

d) additive animal with unknown parent groups and inbreeding (add_an_upginb)

The pedigree file has the following format:
animal number, sire number, dam number, inb/upg code
where sire and/or dam numbers can be replaced by unknown parent group numbers
inb/upg code = 4000 / [(1+ms)(1-Fs) + (1+md)(1-Fd)]

where ms (md) is 0 whenever sire (dam) is known, and 1 otherwise, and Fs(Fd) is the coefficient of inbreeding of the sire (dam). For example, the inb/upg code for the animal with both parents known is 2000.

e) parental dominance (par_domin)

The pedigree class file has the following format:
s-d s-sd s-dd ss-d ds-d ss-sd ss-dd ds-sd ds-dd code
where x-y is a combination number of animals x and y, s is sire, d is dam, sd is sire of dam, etc.
Code is a number of 0 to 255 and refers to the combination of missing subclasses. If one line is:

p s8 s7 s6 s5 s4 s3 s2 s1 code

then code = sum(ai 2**i), where ai=0 if si=1 and 1 otherwise.
For example, the code for a line with all nonzero parental subclasses is 255. For a line with only zero parental subclasses, If classes are ordered so that lines with zero parental subclasses, code=0. If lines are ordered so that p for parental classes with code=0 are ordered last, they may be omitted and will added automatically. The parental dominance file can be created by program **RENDOMN**.

Data and Pedigree files

Transferring a file from Windows (DOS) to Linux environment

Use "dos2unix" to convert the DOS (Windows) format to the UNIX (Linux) format if the programs show an error message while reading a file.

Data file

a. Space(s) is a delimiter. At least one character space between columns is required.
b. Dot (.) is just one character but not a missing value (default missing value = 0).
c. Check the data again especially when converting from another format or software such as EXCEL, SAS, ...
d. For Gibbs sampling programs with "OPTION cont", copy the previous output files somewhere else just in case making mistakes and replacing those files.

Pedigree file

a. An original pedigree file for RENUMF90 can include alpha-numeric characters with free format.
b. Remove duplicates.
c. Use 0 for unknown parent(s).

Parameter file

a. Wrong data file name
   Check outputs for the data file name and the number of records on the screen. The program will not stop if the wrong file name already exists.
b. Wrong pedigree file name
   Check output for the pedigree file name and the number of animals on the screen. The program will not stop if the wrong file name exists.
c. Wrong positions or formats for observations and effects
   Program may not stop and may get wrong results. Check outputs for the number of levels for each effect on the screen.
d. Missing or skipping one or more fixed lines in the parameter file
   Program may stop. Check the missing line.
e. Misspelling
   Program may stop. Correct the wrong spelling.
f. Missing an empty last line
   Program may not stop. Parameter, data, and pedigree files may need one more extra line at the end of the file.
g. (Co)variance matrix is not symmetric, not positive definite, not right sized, ...
   Program may not stop.
h. A good result does not mean that your parameter file is correct. Always double-check!

RENUMF90 parameter file

RENUMF90 is a renumbering program to create input (data and pedigree) files for BLUPF90 programs and provide basic statistics.

Parameter file

DATAFILE

f_1 # data file name – input files cannot contain character # because it is used as a comment.

TRAITS

t_1 t_2 t_3 ... # positions of traits in data file

FIELDS_PASSED TO OUTPUT

p_1 p_2 .. p_m # positions that are not renumbered

WEIGHT(S)

w # position of weight - fraction to the residual variance

RESIDUAL_VARIANCE

R # matrix of residual (co)variances

EFFECT

e_1 e_2 e_3 ... type form # e_1 e_2 e_3 ... = position of this effect for each trait
type = 'cross' for crossclassified or 'cov' for covariables
form = 'alpha' for alphanumeric or 'numer' for numeric

EFFECT

d_1 d_2 d_3 ... cov # d_1 d_2 d_3 ... = positions of covariables nested in the following crossclassified effects

NESTED

e_1 e_2 e_3 ... form # e_1 e_2 e_3 ... = positions of crossclassified effects nested
form = 'alpha' for alphanumeric or 'numer' for numeric

RANDOM

random_type # 'diagonal', 'sire' or 'animal' for random effect

OPTIONAL

o_1 o_2 o_3 ... # 'pe' for permanent environment, 'mat' for maternal, and 'mped' for maternal permanent environment

FILE

fped # pedigree file name

FILE_POS

animal sire dam alt_dam yob # positions of animal, sire, dam, alternate dam (recipient dam), and year of birth in pedigree file (default 1 2 3 0 0).

SNP_FILE

fsnp # specify a SNP file with ID and SNP information; the relationship matrix will include the genomic information; a fsnp file should start with ID with the same format as fped, and SNP info needs to start from a fixed column and include digits 0, 1, 2 and 5;
ID and SNP info need to be separated by at least one space; see more information in **PREGSf90**.

PED_DEPTH
p # depth of pedigree search (default 3); all pedigrees are loaded if p = 0.
GEN_INT
min avg max # minimum, average and maximum generation interval; applicable only if year of birth present in pedigree file; minimum and maximum used for pedigree checks; average used to predict year of birth of parent with missing pedigree.
REC_SEX
sex # if only one sex has records, specifies which parent it is; used for pedigree checks.
UPG_TYPE
t # 'yob' = based on year of birth; if 'in_pedigrees', the value of a missing parent should be -x, where x is UPG number that this missing parent should be allocated to; in this option, all known parents should have pedigree lines, i.e., each parent field should contain either the ID of a real parent, or a negative UPG number. if it is 'internal', allocation is by a user-written function custom_upg (year_of_birth,sex,ID, parent_code).
(CO)VARIANCES
G # (co)variances for animal effects or animal + maternal effects
(CO)VARIANCES_PE
GPE # (co)variances for the PE effect
(CO)VARIANCES_MPE
GMPE # (co)variances for the MPE effect

The additive pedigree file renaddxx.ped has the following structure:
1) animal number (from 1)
2) parent 1 number or unknown parent group number for parent 1
3) parent 2 number or unknown parent group number for parent 2
4) 3 minus number of known parents
5) known or estimated year of birth (0 if not provided)
6) number of known parents
7) number of records
8) number of progenies as parent 1
9) number of progenies as parent 2
10) original animal id

Can we change the maximum size of character fields?
OPTION alpha_size nn # new size (default 20 characters)

How can we specify interactions?
Combining fields or interactions
Several fields in the data file can be combined into one using a COMBINE keyword.
COMBINE a b c .... # keywords COMBINE need to be on top of the parameter file, but possibly after comments.

For example:

COMBINE 7 2 3 4

combines content of fields 2 3 4 into field 7; the data file is not changed, only the program treats field 7 as fields 2 3 4 put together (without spaces). The combined fields can be treated as "numeric" with the total length is < 9 or "alpha".

Example

Input file - data

```
aa 1 10
aa 2 12
bb 1 11
cc 1 12
cc 2 14
dd 2 13
ee 2 14
```

Pedigree file - ped

```
aa  ff  ee 2004
bb  hh  gg 2004
cc  hh  ii 2004
dd  ff  0  2004
ee  ff  0  2002
ff  0   0  2002
gg  ff  0  2002
hh  0   0  2002
ii  0   0  2002
kk  0   0  2000
```

Parameter file - testpar1

```
# Parameter file for program renf90; it is translated to parameter
# file for BLUPF90 family f programs.
DATAFILE
data
TRAITS
3
FIELDS_PASSED TO OUTPUT
1
WEIGHT(S)

RESIDUAL_VARIANCE
1
EFFECT
2 cross num
EFFECT
1 cross alpha
RANDOM
animal
#OPTIONAL
#mat
FILE
```

ped
FILE_POS
1 2 3 0 4
PED_DEPTH
3
GEN_INT
1 2 10
UPG_TYPE
yob
2002 2003

Output log

```
RENUMF90 version 1.73
 name of parameter file?testpar1
 datafile:data
 traits:  3
 fields passed:  1
 R
   1.000
 Processing effect  1 of type cross
 item_kind=num

 Processing effect  2 of type cross
 item_kind=alpha
 pedigree file name  "ped"
 positions of animal, sire, dam, alternate dam and yob  1  2  3  0  4
 pedigree traced to generation   3
 Minimum, average and maximum generation intervals:   1  2  10
 Unknown parent groups separated by years:
   2002  2003

 Maximum size of character fields: 20

 hash tables for effects set up
 read   7  records
 table with   2  elements sorted
 added count
 Effect group   1  of column   1  with   2  levels
 table expanded from   10000  to   10000  records
 added count
 Effect group   2  of column   1  with   5  levels
 wrote statistics in file "renf90.tables"

 Basic statistics for input data  (missing value code is 0)
 Pos  Min         Max         Mean        SD                N
   2    1.0000      2.0000      1.5714     0.53452           7
   3    10.000      14.000      12.286      1.4960           7

 Correlation matrix
        2     3
  2   1.00  0.80
  3   0.80  1.00

 Counts of nonzero values (order as above)
       7      7
       7      7
```

```
 random effect   2
 type:animal
 opened output pedigree file "renadd02.ped"
 read   10  pedigree records
 loaded   4  parent(s) in round   1

 Pedigree checks
 ee: younger than parent   1  by   0  years
 gg: younger than parent   1  by   0  years

 Unknown parent group allocation
 Equation   Group       #Animals   Years
       10       1       0             0-   2001
       11       2       8          2002-   2002
       12       3       1          2003-

 Number of animals with records:  5
 Number of parents without records:  4
 Total number of animals:  9

 Wrote parameter file "renf90.par"
 Wrote renumbered data "renf90.dat"

Output data file - renf90.dat
observation, effect 1, effect 2, animal ID
 10 1 4 aa
 12 2 4 aa
 11 1 2 bb
 12 1 5 cc
 14 2 5 cc
 13 2 3 dd
 14 2 1 ee

Output pedigree file - renadd03.ped
Animal, sire, dam, 3-#unknown parents, birth year, #known parents, #records, #progeny of sire, #
progeny of dam, original animal ID
 1 6 11 2 2002 1 1 0 1 ee
 2 8 7 1 2004 2 1 0 0 bb
 7 6 11 2 2002 1 0 0 1 gg
 3 6 12 2 2004 1 1 0 0 dd
 9 11 11 3 2002 0 0 0 1 ii
 4 6 1 1 2004 2 2 0 0 aa
 6 11 11 3 2002 0 0 4 0 ff
 5 8 9 1 2004 2 2 0 0 cc
 8 11 11 3 2002 0 0 2 0 hh
```

Output parameter file - renf90.par

```
DATAFILE
 renf90.dat
NUMBER_OF_TRAITS
  1
NUMBER_OF_EFFECTS
  2
OBSERVATION(S)
  1
WEIGHT(S)
```

```
EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT[EFFECT NESTED]
 2        2 cross
 3       12 cross
RANDOM_RESIDUAL VALUES
  1.000
RANDOM_GROUP
  2
RANDOM_TYPE
add_an_upg
FILE
renadd02.ped
(CO)VARIANCES
  1.000
```

Output tables after renumbering - renf90.tables

```
Effect group 1 of column 1 with 2 levels Value  # consecutive number
      1     3           1
      2     4           2
```

When to use what program and computing limits

BLUP

BLUPF90 sets up equations in memory. It can support a few million equations with a simple model to much smaller with complicated models. BLUPF90 uses three solvers, chosen with options. PCG is the default solver and is usually the fastest one. SOR require less memory but usually converges slower. Sparse Cholesky (FSPAK) is usually the most accurate method but uses the most memory. The following options are available:

OPTION conv_crit 1e-12
Set convergence criteria (default 1e-10).
OPTION maxrounds 10000
Set maximum number of rounds (default 1000).
OPTION solv_method FSPAK
Selection of solving method: FSPAK, SOR or PCG (default PCG).
OPTION r_factor 1.6
Set relaxation factor for SOR (default 1.4).
OPTION sol se
Store solutions and s.e. If this option is used, the solving method will turn to FSPAK.
OPTION blksize 3
Set block size for preconditioner (default 1).

BLUP90IOD uses an iteration on data algorithm. It can handle hundreds of millions of equations with complicated models in a reasonable time. However, it is only available with a research contract or for research at UGA. The following options are available:

OPTION conv_crit 1e-12
Set convergence criteria (deault 1e-12).
OPTION maxrounds 10000
Set maximum number of rounds (default 5000).
OPTION blksize 3
Set block size for preconditioner (default 1). Usually **blksize** will be the same number of traits.
OPTION init_eq 10
Set the number of effects to be solved directly (default 0).
OPTION solv_method FSPAK
Solving method for initial equations (default DIRECT).
OPTION tol 1d-12
Tolerance to get a positive definite matrix (default 1d-12).
OPTION residual
y-hat and residuals will be included in “yhat_residual”.

OPTION avgeps 50

Using the last 50 average eps for convergence.

OPTION cont 1

To restart the program from the previous solutions.

OPTION missing -1

Set the missing value (default 0).

OPTION restart 100

Set the number of iteration to recompute residuals (default 100).

OPTION prior_solutions

Using the previous solution file to start the iteration. Additional software is required to use this option.

OPTION random_upg 1 2

Set the UPG random. "1" the weight for random UPG = 1. If the second number exists, the weight will be inverted (e.g., 1/2=0.5).

OPTION SNP_file snp

Specify the SNP file name snp to use genotype data.

Variance component estimation

There is not a single-best choice for variance component estimation. Programs below offer choices for simple and complicated models. For advice on what works best under your circumstances, google a paper "**Reliable computing in estimation of variance components**".

REMLF90 uses EM REML. For most problems it is the most reliable algorithm but can take hundreds of rounds of iteration. REMLF90 was found to have problems converging with random regression models. In this case, using starting variances that are too large than too small usually helps. Also, EM does not calculate standard errors. The following options are available:

OPTION conv_crit 1d-12
Convergence criterion (default 1d-10).
OPTION maxrounds 10000
Maximum rounds (default 5000).
OPTION sol se
Store solutions and se.
OPTION residual
y-hat and residuals will be included in "yhat_residual".
OPTION missing -999
Specify missing observations (default 0).
OPTION use_yams
Run the program with YAMS (modified FSPAK). The computing time can be dramatically improved.
OPTION constant_var 5 1 2
5: effect number, 1: first trait number, 2: second trait number implying the covariance between traits 1 and 2 for effect 5 is fixed.
OPTION SNP_file snp
Specify the SNP file name **snp** to use genotype data.

AIREMLF90 uses AI REML. It usually converges much faster but sometimes does not converge. Very slow convergence usually indicates that the model is over parameterized and there is insufficient information to estimate some variances. AI REML also calculates standard errors. The following options are available:

OPTION conv_crit 1d-12
Convergence criterion (default 1d-10).
OPTION maxrounds 500
Maximum rounds (default 5000). When it is negative, the program calculates BLUP without running REML.
OPTION EM-REML 10
Run EM-REML for the first 10 rounds to get initial variances within the parameter space (default 0).

OPTION tol 1d-18
Tolerance (or precision) for positive definite matrix and G-inverse subroutines (default 1d-14).
OPTION sol se
Store solutions and s.e.
OPTION missing -1
Set the missing observation (default 0).
OPTION constant_var 5 1 2
5: effect number, 1: first trait number, 2: second trait number implying the covariance between traits 1 and 2 for effect 5 is fixed.

Heterogeneous residual variances for a single trait
OPTION hetres_pos 10 11
Specify positions of covariables.
OPTION hetres_pol 4.0 0.1 0.1
Initial values of coefficients for heterogeneous residual variances. Use *ln*(a0, a1, a2, ...) to make these values. When the number of positions = the number of polynomials, the regressions do not include the intercept (e.g., linear spline).

Heterogeneous residual variances for multiple traits (the convergence will be very slow)
OPTION hetres_pos 10 10 11 11
Specify positions of covariables (trait first).
OPTION hetres_pol 4.0 4.0 0.1 0.1 0.01 0.01
Initial values of coefficients for heterogeneous residual variances using ln(a0, a1, a2, …) to make these values (trait first). "4.0 4.0" are intercept for first and second traits. "0.1 0.1" could be linear and "0.01 0.01" could be quadratic. To transform back to the original scale, use exp(a0+a1*X1+a2*X2).
OPTION SNP_file snp
Specify the SNP file name **snp** to use genotype data.

GIBBSxF90 programs implement Bayesian methods. These methods potentially have better statistical properties. Also they are more stable and use less memory for complicated models. After running any of the Gibbs sampling program, analyze samples with the POSTGIBBSF90 programs.
In practical cases, results from Gibbs samplers and REML are similar. Choose one or the other based on computing feasibility. If there are large differences beyond sampling errors, this indicates problems usually with the Gibbs sampler. Try longer chains or different priors.

Gibbs samplers may be slow to achieve convergence if initial values are far away from those at convergence, e.g., 100 times too low or too high. Before using more complicated models, Karin Meyer advocates using a series of simpler models.

GIBBS1F90 can run models with over 20 traits. However, if models are different per trait, the lines due to effects need to be modified. Also, with too many differences in models among traits, the program becomes increasingly slower.
GIBBS2F90 adds joint sampling of correlated effects. This results in faster mixing with random regression and maternal models.

Interactive inputs:
number of samples and length of burn-in?
In the first run, if you have no idea about the number of samples and burn-in, just type your guess (10000 or whatever) for samples and (0) for burn-in. You may need 2 or 3 runs to figure out the convergence.
Give n to store every n-th sample?
Gibbs samples are highly correlated, so you do not have to keep all samples (every 10th, 20th, 50th, ...).
The following options are available for **GIBBSxF90**:

OPTION fixed_var all 1 2 3
Store all solutions and posterior means and SD for effects for effects1, 2, and 3 are stored in "all_solutions" and in "final_solutions" every round using fixed variances. Without numbers, all solutions for all effects are stored.
OPTION fixed_var mean 1 2 3
Posterior means and SD for effects1, 2, and 3 in "final_solutions".
OPTION solution all 1 2 3
Store all solutions and posterior means and SD for effects1, 2, and 3 are stored in "all_solutions" and in "final_solutions" every round. Without numbers, all solutions for all effects are stored.
OPTION solution mean 1 2 3
Posterior means and SD for effects1, 2, and 3 in "final_solutions".
OPTION cont 10000
10000 is the number of samples run previously when restarting the program from the last run.
OPTION prior 5 2 -1 5
The (co)variance priors are specified in the parameter file. Degree of belief for all random effects should be specified using the following structure: OPTION prior eff1 db1 eff2 db2 … effn dbn -1 dbres effx correspond to the effect number and dbx to the degree of belief for this random effect, -1 corresponds to the degree of belief of the residual variance. In this example 2 is the degree of belief for the 5th effect, and 5 is the degree of belief for the residual.
OPTION seed 123 321
Two seeds for a random number generator can be specified.
OPTION SNP_file snp
Specify the SNP file name **snp** to use genotype data.

GIBBS3F90 adds estimation of heterogeneous residual covariances in classes. The computing costs usually increase with the number of classes.

OPTION hetres_int 5 10
The position (5) to identify the interval in the data file and the number of intervals (10) for heterogeneous residual variances.

Other options are the same as for **GIBBS1F90** and **GIBBS2F90**.

THRGIBBS1F90 is a Gibbs sampling program to analyze categorical and continuous traits simultaneously. The following options are available:

OPTION cat 0 0 2 5
"0" indicate that the first and second traits are linear. "2" and "5" indicate that the third and fourth traits are categorical with 2 (binary) and 5 categories.
OPTION thresholds 0.0 1.0 2.0
Set the fixed thresholds. No need to set 0 for binary traits.
OPTION residual 1
Set the residual variance = 1.
OPTION censored xx
Negative values of the last category in the data set indicate censored records. "xx" determines the lower and upper limit of the category + xx when sampling from the distribution.

Other options are the same as for **GIBBS1F90** and **GIBBS2F90**.

POSTGIBBSF90 is a program to calculate posterior means and SD and diagnose the convergence. The program reads "**gibbs_samples**" and "**fort.99**" files from Gibbs sampling programs.

Read 1000 samples from round 10 to 10000

Burn-in?
1000 # in the first run, type 0 for burn-in to include all samples

Give n to read every n-th sample? (1 means read all samples)
10 # Type the same number used with a Gibbs sampling program.

samples after burn-in = 9000

Input files:
gibbs_samples, fort.99, and other files used in a parameter file from **(THR)GIBBSxF90**
Output files:
postgibbs_samples, postout, postmean, postsd

postgibbs_samples

A text file containing all Gibbs samples from **gibbs_samples** for other software (EXCEL, SAS, …) to calculate posterior means and SD, and to create graphs.

postmean

Posterior means

postsd

Posterior standard deviations

postout

					******** Monte Carlo Error by Time Series ********						
Pos.	eff1	eff2	trt1	trt2	MCE	Mean	HPD Interval (95%)		Effective sample size	Median	Mode	Independent chain size
1	4	4	1	1	1.362E-02	0.9889	0.7788	1.215	70.4	0.9844	0.9861	18
2	4	4	1	2	1.288E-02	1.006	0.777	1.219	84.1	1.006	0.952	18
3	4	4	2	2	1.847E-02	1.66	1.347	1.987	80.3	1.652	1.579	25
4	0	0	1	1	9.530E-03	24.47	24.07	24.84	425.6	24.47	24.53	2
5	0	0	1	2	8.253E-03	11.84	11.54	12.18	395.8	11.83	11.82	2
6	0	0	2	2	1.233E-02	30.1	29.65	30.58	387.8	30.09	29.97	5

******** Posterior Standard Deviation ********

Pos.	eff1	eff2	trt1	trt2	PSD	Mean	PSD Interval (95%)		Geweke diagnostic	Autocorrelations lag: 1	10	50	Independent # batches
1	4	4	1	1	0.1144	0.9889	0.7648	1.213	-0.02	0.853	0.188	0.049	50
2	4	4	1	2	0.1182	1.006	0.7742	1.237	-0.11	0.828	0.111	-0.066	50
3	4	4	2	2	0.1656	1.66	1.335	1.984	0.06	0.828	0.108	-0.021	36
4	0	0	1	1	0.1967	24.47	24.09	24.86	-0.01	0.034	0.029	-0.062	450
5	0	0	1	2	0.1643	11.84	11.51	12.16	0.03	0.032	-0.006	-0.016	450
6	0	0	2	2	0.2429	30.1	29.62	30.57	-0.02	0.07	-0.014	0.037	180

where

"Pos."

position of each parameter in the parameter file

"eff1" and "eff2"

effect number in the parameter file

"trt1" and "trt2"

trait number in the parameter file

"MCE"

Monte Carlo Error

"Mean"

posterior means

"HPD interval (95%)"

95% Highest Probability Density

"Effective sample size"

at least > 10 is recommended. > 30 may be better.

"Median"

median of Gibbs samples

"Mode"

when the distribution of the samples is not normal, "Mean" and “Mode” could be different.

"Independent chain size"

number of dependent cycles of Gibbs samples

"PSD"

Posterior Standard Deviation

"PSD interval (95%)"

95% Posterior Standard Deviation interval

"Geweke diagnostic"

ratio between first half and second half of the samples should be < 1.0, but it is not useful because it is < 1.0 most of the time.

"Autocorrelations"

autocorrelations between two lags. High correlation implies samples are not independent.

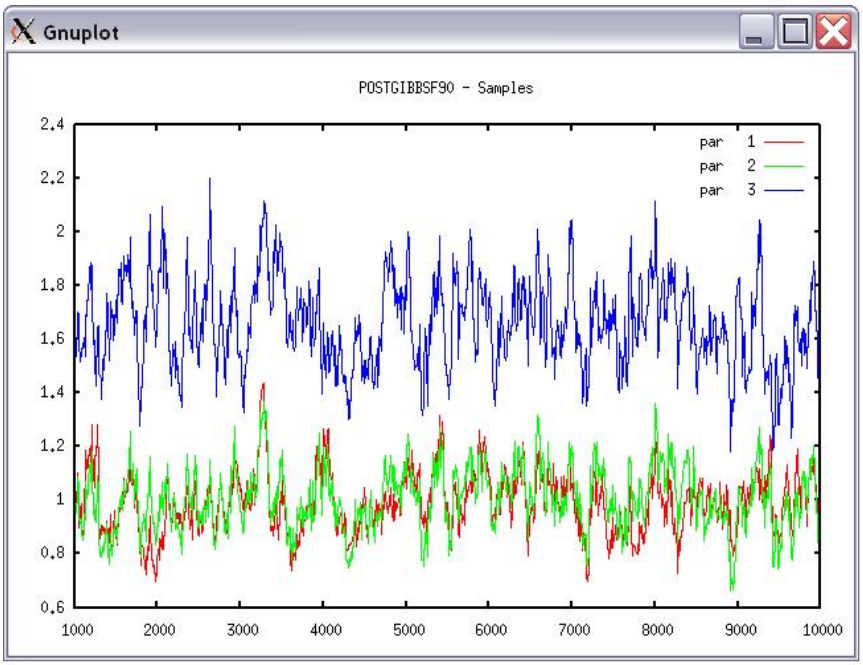
"Independent # batches"

Choose a graph for samples (= 1) or histogram (= 2); or exit (= 0)

1

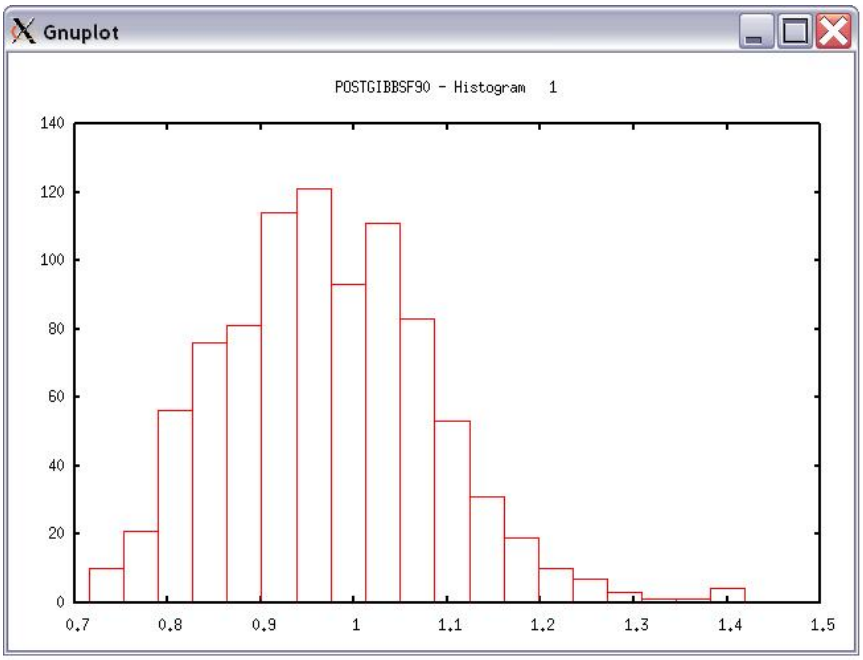
positions

1 2 3 # choose from the position numbers 1 through 6



If the graph is stable (not increasing or decreasing), the convergence is met. All samples before that point should be discarded as burn-in.

print = 1; other graphs = 2; or stop = 0
2
Choose a graph for samples (= 1) or histogram (= 2); or exit (= 0)
2
Type position and # bins
1 20



The distribution should be usually normal (Mean = Mode = Median).
print = 1; other graphs = 2; or stop = 0
0

*** Log Marginal Density for Bayes Factor ***
after 900 burn-in
log(p) = -179448.742766031

This value could be used when calculating Bayes Factor and/or DIC.

Genomic programs

The **PREGSF90** program constructs a genomic relationship matrix **G** and a relationship matrix $\mathbf{A}_{22}$ for genotyped animals. The relationship matrix **A** based on the pedigree information in mixed model equations is replaced by matrix **H**, which combines the pedigree and genomic information. The main difference between $\mathbf{A}^{-1}$ and $\mathbf{H}^{-1}$ is structure of $\mathbf{G}^{-1} - \mathbf{A}_{22}^{-1}$. Some of the options for **PREGSF90** can be also used with **BLUPF90**, (**AI)REMLF90**, **GIBBS1F90**, **GIBBS2F90**, **GIBBS3F90**, **THRGIBBS1F90**, and **BLUP90IOD2**.

OPTION SNP_file <file>
The SNP file should contain
Field 1 - animal ID with the same format as in pedigree file
Field 2 - genotypes with 0, 1, 2, and 5 (missing) or real values for gene content 0.12, …

Two Fields need to be separated by at least one space, and Field 2 should have fixed format (i.e., all rows of genotypes should start at the same column number or position).

```
 80   21101011002012011011010110111111211111210100
 8014 21110101511101120221110111511112101112210100
 516  21100101202252021120210121102111202212111101
 181  21110111112201120550200020101022212211111100
```

The renumbered ID file for genotypes named as the genotype file **name.XrefID** is created by RENUMF90, containing sequential ID renumbers and the original ID, which must be in the same order as in the SNP file as follows:

```
1732 80
8474 8014
406 516
9441 181
```

The pedigree file from RENUMF90 looks like

```
1732 11010 10584 1 3 12  1 0 0 80
8474  8691  9908 1 3 12  1 0 0 8014
 406  8691  9825 1 3 12  1 0 2 516
9441  8691  8829 1 3 12  1 0 0 181
```

Several optional files are available:

Allele frequencies (**OPTION FreqFilev <file>**)
Map file (**OPTION chrinfo <file>**)
Weight file (**OPTION weightedG <file>**)
G or its inverse, $\mathbf{A}_{22}$ or its inverse, etc, as specified by respective OPTIONs.
OPTION chrinfo <file>: read SNP map information from the file.

These files are useful to check for Mendelian conflicts and HWE (with also **OPTION sex_chr**) and for **POSTGSF90** (**ssGWAS**).

Format = all numeric variables: SNP order , chromosome, position (bp): the SNP order corresponds to the index number of the SNP, in the sorted map by chromosome and the position.
The first line in the file corresponds to the first SNP in the genotype file, and so on. Other alphanumeric fields are optional.

By default, **PREGSf90** always create GimA22i in binary format for use by later programs specifying **OPTION readGimA22i**. With **OPTION saveAscii**, this file can be stored as ASCII format: i, j, $\mathbf{G}^{-1} - \mathbf{A}_{22}^{-1}$.
"freqdata.count" contains allele frequencies in the original genotype file with the format: SNP number (related to the genotype file) and allele frequency.
"freqdata.count.after.clean" contains allele frequencies as used in calculations with the format: SNP number (related to the genotype file), allele frequency, and code of exclusion.

Exclusion codes:
1: Call Rate
2: MAF
3: Monomorphic
4: Excluded by request
5: Mendelian error
6: HWE
7: High Correlation with other(s) SNP

"Gen_call_rate" contains a list of animals excluded with call rate below the threshold.
"Gen_conflicts" contains a report of animals with Mendelian conflicts with their parents.
The program can store files such as **G** or its inverse, $\mathbf{A}_{22}$ or its inverse, or other reports from QC as specified by their respective OPTIONs.

Options for creation of genomic relationship Matrix (**G**)
The genomic relationship matrix G can be created in different ways.
OPTION whichG x
Specify how **G** is created.
The variable x can be
1: $\mathbf{G} = \frac{\mathbf{ZZ}'}{k}$; VanRaden, 2008 (default)
2: $\mathbf{G} = \frac{\mathbf{ZDZ}'}{n}$; Amin et al., 2007; Leuttenger et al., 2003; where $\mathbf{D} = \frac{1}{2p(1-p)}$
3: As 2 with modification UAR from Yang et al 2010

OPTION whichfreq x
Specify what frequency is used to create **G**.
The variable x can be
0: read from file "freqdata" or from the other file using **OPTION FreqFile**

1: 0.5
2: current calculated from genotypes (default)

OPTION FreqFile <file>
Read allele frequencies from a file. For example, based on allele frequencies calculated by estfreq.f90 (VanRaden, 2009) with format: SNP, frequency where SNP corresponds to the index of SNP based on the same order that are in the genotype file.
If **whichfreq** is set to 0, the default file name is "freqdata".

OPTION whichScale x
Specify how **G** is scaled.
The variable **x** can be
1: $2\sum\{p(1-p)\}$; VanRaden 2008 (default)
2: $\frac{tr(\mathbf{ZZ}')}{n}$; Legarra 2009, Hayes 2009
3: correction ; Gianola et al 2009

OPTION weightedG <file>
Read weights from a file to create weighted genomic relationship. Weighting Z* = Z sqrt(D) ⇒ **G** = Z*Z*' = ZDZ' (format: one column of weights in the same order as in the genotyped file). Weights can be extracted from output of the **POSTGSF90** program.

OPTION maxsnp x
Set the maximum length of string to read marker data from a file. It is only necessary if greater than default (400,000).

Quality Control (QC) for **G**
By default the following QC can be run:

- MAF
- Call rate (SNPs and animals)
- Monomorphic
- Parent-progeny conflicts (SNPs and animals)

Parameters can be modified with the following options:
OPTION minfreq x
Ignore all SNP with MAF < x (default value = 0.05).
OPTION callrate x
Ignore SNP with call rates < x (number of calls / number of individuals with genotypes). The default value is 0.90.
OPTION callrateAnim x
Ignore genotypes with call rates < x (number of calls / number of SNPs). Default value is 0.90.

OPTION monomorphic x

Ignore monomorphic SNPs. Optional parameter **x** can be used to enable (1) or disable (0) the check, default value 1.

OPTION hwe x

Check departure of heterozygous from Hardy-Weinberg equilibrium. By default this QC is not run. The optional parameter **x** can be the maximum difference between observed and expected frequency (default value = 0.15) as used in Wiggans et al. (2009) in JDS.

OPTION high_correlation x y

Check for high correlated SNP. By default this QC is not run. The optional parameter **x** can be the maximum difference in allele frequency to check a pair of locus. If no value is set, 0.025 is used. Decrease this value to speed up the calculation. A pair of loci is considered highly correlated if all genotypes are the same (0-0, 1-1, 2-2) or the opposite (0-2, 1-1, 2-0) (Wiggans et al., 2009. JDS). The optional parameter **y** can be used to set a threshold to check the number of identical samples out of the number of genotypes (default values: x=0.025, y=0.995).

OPTION verify_parentage x

Verify parent-progeny Mendelian conflicts and write report to a file "Gen_conflicts". The optional parameter **x** can be

0: no action

1: only detect

2: detect and search for an alternate parent; no change to any file. Not yet implemented

3: detect and eliminate progenies with conflicts (default)

OPTION exclusion_threshold x

Set the number of parent-progeny exclusions as percentage. All SNP are used to determine wrong relationships (default value = 2).

OPTION exclusion_threshold_snp x

Set the number of parent-progeny exclusions for each locus as percentage. A pair of genotyped animals is evaluated to exclude SNP from the analysis (default value = 10).

OPTION number_parent_progeny_evaluations x

Set the number of minimum pair of parent-progeny evaluations to exclude SNP due to parent-progeny exclusion (default value = 100).

OPTION outparent_progeny x

Create a full log file "Gen_conflicts_all" with all pairs of parent-progeny tested for Mendelian conflicts.

OPTION excludeCHR n1 n2 n3 …

Exclude all SNP from chromosomes n1, n2, n3, … A map file must be provided (see **OPTION chrinfo**).

OPTION sex_chr n

Set the chromosome number equal to or greater than **n** are not considered autosome. If this option is used, sex chromosomes will not be used for checking parent-progeny, Mendelian conflicts, and HWE. A map file must be provided (see **OPTION chrinfo**).

OPTION threshold_duplicate_samples x

Set the threshold to issue warning for possible duplicate samples if G(i,j) / sqrt(G(i,i) * G(j,j)) > **x** (default value = 0.9).

OPTION threshold_diagonal_g x

Check for extremely large diagonals in the genomic relationship matrix. If optional **x** is present, the threshold will be set (default value = 1.6).

OPTION plotca

Plot first two principal components to look for stratification in the population.

OPTION extra_info_pca <file> col

Read the column col to plot with different colors for different classes from the file. The file should contain at least one variable with different classes for each genotyped individual, and the order should match the order of the genotype file. Variables could be alphanumeric and separated by one or more spaces.

OPTION saveCleanSNPs *

Save clean genotype data with excluded SNP and animals based on the OPTIONS specified.

*_clean files are created:

- gt_clean
- gt_clean_XrefID

*_removed files are created.

- gt_SNPs_removed
- gt_Animals_removed

where "gt" is the genotype file.

OPTION no_quality_control

Turns off all quality control. It is useful to speed up computation when the QC was performed previously.

OPTION outcallrate

Print all call rate information for SNP and individuals. The files "callrate" for SNP and "callrate_a" for individuals are created.

Quality Control for Off-diagonal of $\mathbf{A}_{22}$ and **G**

OPTION thrWarnCorAG x

Set the threshold to issue warning if correlation between $\mathbf{A}_{22}$ and $\mathbf{G} <$ **x** (default value = 0.5).

OPTION thrStopCorAG x

Set the threshold to stop the analysis if correlation between $\mathbf{A}_{22}$ and $\mathbf{G} <$ **x** (default values = 0.3).

OPTION thrCorAG x

Set the threshold to calculate correlation between $\mathbf{A}_{22}$ and $\mathbf{G}$ for only $\mathbf{A}_{22}$, ≥ **x** (default values = 0.02).

Options for **H** including different weights to create $\mathbf{G}^{-1} - \mathbf{A}_{22}^{-1}$ as

(alpha **G** + beta $\mathbf{A}_{22}$ + gamma **I** + delta)$^{-1}$ - omega $\mathbf{A}_{22}^{-1}$

where the parameters are to scale the genomic info to be compatible with the pedigree information, to make matrices invertible in the presence of clones, and to control bias.

The defaults values are:

tau=1 alpha =0.95 beta = 0.05 gamma=0 delta=0 omega=1
Options to change these defaults are specified with:
OPTION TauOmega tau omega
OPTION AlphaBeta alpha beta
OPTION GammaDelta gamma delta
OPTION tunedG x
Scale **G** based on $\mathbf{A}_{22}$. The variable x can be:
0: no scaling
1: mean(diag(**G**))=1, mean(offdiag(**G**))=0
2: mean(diag(**G**))=mean(diag($\mathbf{A}_{22}$)), mean(offdiag(**G**))=mean(offdiag($\mathbf{A}_{22}$)) (default)
3: mean(**G**)=mean($\mathbf{A}_{22}$)
4: rescale **G** using the first adjustment as in Powell et al. (2010) or Vitezica et al. (2011).

OPTION nthreads n
Specify number of threads to be used with MKL-OpenMP for creation and inversion of matrices.
OPTION ntheadsiod n
Specify number of threads to be used with MKL-OpenMP in BLUP90IOD for matrix-vector multiplications in the PCG algorithm.
OPTION graphics s
Allows to generate plots with GNUPLOT. If optional parameter s is present, set the time in seconds to show the plot. Avoid using in batch programs!!!
OPTION msg x
Set the level of verbose; 0 minimal; 1 gives lots of diagnostics.

Save and Read options:
OPTION saveAscii
Save intermediate matrices (GimA22i, G, Gi, etc) files as ASCII (default = binary).
OPTION saveHinv
Save $\mathbf{H}^{-1}$ in "Hinv.txt" (format: i, j, val with i, j, the index level for the additive genetic effect).
OPTION saveAinv
Save $\mathbf{A}^{-}$ in "Ainv.txt" (format: i, j, val with i, j, the index level for the additive genetic effect).

The following options use the information of the original ID (alphanumeric) stored in the 10th column of the "renaddxx.ped" file created by **RENUMF90**.
OPTION saveHinvOrig
Save $\mathbf{H}^{-1}$ with original IDs
OPTION saveAinvOrig
Save $\mathbf{A}^{-1}$ with original IDs
OPTION saveDiagGOrig
Save diagonal of **G** in "DiagGOrig.txt" (format: id, val with id, original IDs).
OPTION saveGOrig
Save **G** in "G_Orig.txt" (format: id_i, id_j, val with id_i and id_j, the original IDs).

OPTION saveA22Orig

Save $\mathbf{A}_{22}$ in "A22_Orig.txt" (format: id_i, id_j, val with id_i and id_j, the original IDs).

OPTION readOrigId

Read information from "renaddxx.ped" file, original ID and possibly year of birth for its use in parent-progeny conflict. Only need unless the previous "save*Orig" is present.

OPTION savePLINK

Save genotypes in PLINK format files: toPLINK.ped and toPLINK.map.

Save and Read intermediate files:

OPTION readGimA22i <file>

This option can be used in analysis programs (BLUPF90, REMLF90, etc.) in order to use matrices stored in GimA22i file (default filename). In general, methods used to create and invert matrices in such programs don not use optimized version. For large number of genotyped animals, run first PREGSf90 and read stored matrices in analysis programs.

The optional file can be used to specify the other file name or path.

For example,

OPTION readGimA22i ../../pregsrun/GimA22i

Other intermediate matrices files can be stored for inspection or for use in BLUPF90 programs as **user_file** type of random effect. See **tricks** and **REMLF90** for details.

OPTION saveA22

OPTION saveA22Inverse

OPTION saveG all

If optional all is present, all intermediate matrices for **G** will be saved.

OPTION saveGInverse

OPTION saveGmA22

OPTION readG <file>

OPTION readGInverse <file>

OPTION readA22 <file>

OPTION readA22Inverse <file>

OPTION readGmA22 <file>

POSTGSF90

The following options for **POSTGSF90** (ssGWAS) are available:

OPTION Manhattan_plot

Plot using **GNUPLOT** the Manhattan plot (SNP effects) for each trait and correlated effect.

OPTION Manhattan_plot_R

Plot the Manhattan plot (SNP effects) for each trait and correlated effects using R. TIF images are created: manplot_Sft1e2.tif (note: t1e2 corresponds to trait 1, effect 2). **CAIRO** packaged is required.

OPTION plotsnp n

Control the values of SNP effects to use in Manhattan plots

1: plot regular SNP effects: abs(val)

2: plot standardized SNP effects: abs(val/sd) (default)

OPTION SNP_moving_average n
Solutions for SNP effects will be by moving average of n adjacent SNPs.
OPTION windows_variance n
Calculate the variance explained by n adjacent SNPs.
OPTION windows_variance_mbp n
Calculate the variance explained by n Mb window of adjacent SNPs.
OPTION windows_variance_type n
Set windows type for variances calculations
1: moving windows
2: exclusive windows
OPTION which_weight x
Generate a weight variable to be used in the creation of a weighted genomic relationship matrix G=ZDZ'
1: w = y^2 * (2(p(1-p)))
2: w = y^2
with scaled weight = w * nSnp/sum(w)

Output files for **POSTGSF90**:
"snp_sol" contains solutions of SNP and weights
1: trait
2: effect
3: SNP
4: Chromosome
5: Position
6: SNP solution
7: weight if **OPTION windows_variance** is used
8: variance explained by n adjacent SNP.

"chrsnp" contains data to create plot by GNUPLOT
1: trait
2: effect
3: values of SNP effects to use in Manhattan plots
4: SNP
5: Chromosome
6: Position

"chrsnpvar" contains data to create plot by GNUPLOT
1: trait
2: effect
3: variance explained by n adjacent SNP
4: SNP
5: Chromosome
6: Position

"snp_pred" contains gene frequencies + SNP effects

Graphic control files:
Several files are created to generate graphics using either GNUPLOT or R

File names rules
"Sft1e2.R". The first letter indicates "S" for solutions of SNP and "V" for variance explained.
"t1e2" indicates that the file is for the trait 1 and the effect 2.

Filename extension

xxx.gnuplot => GNUPLOT
xxx.R => R programs
xxx.tif => image

PREDF90 predicts GEBV for young animals based on only genotypes. The prediction is based on SNP effects obtained from **POSTGSF90**. For young animals that were not included in the previous analysis, GEBV can be calculated using the "snp_pred" file from **POSTGSF90**.
Input files:
"snp_pred"
- information about the random effect (number of traits + correlated effects)
- gene frequencies
- solutions of SNP effects

Prepare an updated genotype file in the same format as used in POSTGSf90.
Output file:
"SNP_predictions"
- ID, calling rate, and GEBV

Parameters**:**

1. alpha - fraction of G used (default=0.95); affects scale of prediction
2. callrate - to be used later for discarding genotypes with poor quality (default=0.7)

Sample run using example from our website

"renum.par" for **RENUMF90**

DATAFILE
phenotypes.txt
TRAITS
3
FIELDS_PASSED TO OUTPUT

```
WEIGHT(S)

RESIDUAL_VARIANCE  # variances are from airemlf90 results
 0.9038
EFFECT
1 cross alpha
EFFECT
2 cross alpha  #animal
RANDOM
animal
FILE
pedigree
SNP_FILE
marker.geno.clean
(CO)VARIANCES
   0.9951E-01
```

Run RENUMF90

```
RENUMF90 version 1.94
 name of parameter file?renum.par
 .....
 Number of animals with records:       15800
 Number of animals with genotypes:        1500
 .....
 Wrote renumbered data "renf90.dat"
```

"renf90.par" from RENUMF90

```
# BLUPF90 parameter file created by RENF90
DATAFILE
 renf90.dat
NUMBER_OF_TRAITS
       1
NUMBER_OF_EFFECTS
       2
OBSERVATION(S)
   1
WEIGHT(S)

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT[EFFECT NESTED]
 2       1 cross
 3    15800 cross
RANDOM_RESIDUAL VALUES
 0.9038
 RANDOM_GROUP
   2
 RANDOM_TYPE
 add_animal
 FILE
```

renadd02.ped
(CO)VARIANCES
0.9951E-01
OPTION SNP_file marker.geno.clean

Run BLUPF90

```
 name of parameter file?renf90.par
  .....
 round           67    convergence=  1.259204136398044E-012
 round           68    convergence=  9.025592858512443E-013
          68  iterations,   convergence criterion=  9.025592858512443E-013
 solutions stored in file: "solutions"


 $a/postGSf90
 name of parameter file?renf90.par

     postGS 1.11
.....
 Solutions read from file: "solutions"
 .....
 Files for pedictions by SNP effects in file: "snp_pred"


$head -5 snp_pred
        3000           1           0       15800
 0.751 0.382 0.569 0.680 0.184 0.298 0.392 0.380 0.597 0.352
 0.514 0.717 0.464 0.502 0.639 0.773 0.364 0.645 0.566 0.514
 0.622 0.673 0.238 0.556 0.606 0.590 0.477 0.341 0.523 0.525
 0.660 0.439 0.609 0.418 0.572 0.401 0.490 0.608 0.454 0.589
```

Run PREDF90

```
Predf90 1.00
 Predicts EBVs from genotypes based on results from single-step evaluation
 name of genotype file?
marker.geno.clean
Number of SNP:     3000
Number of traits:   1
number of correlated traits:   1
        3000  SNP
The genotype file contains    3000 SNP starting from position   7
  8002   0.1186204
  8014  -0.1033363
  8016   0.1308713
  8018  -0.1905423
  8024  -0.3675095
  8038   0.1939673
  8041  -0.1284970
  8063  -0.1314869
  8065  -2.8898019E-02
Processed  1500 genotypes
Average calling rate: 1.00

head -5 SNP_predictions
```

```
8002    1.00      0.1156
8014    1.00     -0.1007
8016    1.00      0.1276
8018    1.00     -0.1857
8024    1.00     -0.3582
```

PREDICTF90 is to calculate $\hat{\boldsymbol{y}}$ and residuals using the same parameter file and "solutions" and can be used to calculate predictive ability $r(y, \hat{\boldsymbol{y}})$.

Output files:

"yhat_residual"

Format: record #, original y, $\hat{\boldsymbol{y}}$, residual

"bvs.dat"

The same format as "solutions" including (G)EBV.

Examples for parameter files

Sire model without A

```
DATAFILE
test.dat
NUMBER_OF_TRAITS
1
NUMBER_OF_EFFECTS
2
OBSERVATION(S)
3
WEIGHT(S)

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
1 2 cross
2 3 cross
RANDOM_RESIDUAL VALUES
10
RANDOM_GROUP
2
RANDOM_TYPE
diagonal
FILE

(CO)VARIANCES
1
```

Sire model with A

```
DATAFILE
test.dat
NUMBER_OF_TRAITS
1
NUMBER_OF_EFFECTS
2
OBSERVATION(S)
3
WEIGHT(S)
EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
1 2 cross
2 3 cross
RANDOM_RESIDUAL VALUES
10
RANDOM_GROUP
2
RANDOM_TYPE
add_sire
FILE
sire.ped
(CO)VARIANCES
1
```

Multiple (2) trait sire model

```
DATAFILE
test.dat
NUMBER_OF_TRAITS
2
NUMBER_OF_EFFECTS
2
OBSERVATION(S)
3 4
WEIGHT(S)
EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
1 1 2 cross
2 2 3 cross
RANDOM_RESIDUAL VALUES
10 1
1 5
RANDOM_GROUP
2
RANDOM_TYPE
add_sire
FILE
sire.ped
(CO)VARIANCES
1 0.1
0.1 1
```

Animal model

```
DATAFILE
test.dat
NUMBER_OF_TRAITS
1
NUMBER_OF_EFFECTS
2
OBSERVATION(S)
3
WEIGHT(S)
EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
1 2 cross
5 10 cross
RANDOM_RESIDUAL VALUES
10
RANDOM_GROUP
2
RANDOM_TYPE
add_animal
FILE
animal.ped
(CO)VARIANCES
1
```

Multiple trait animal model

Example 1: 2 trait animal model

```
DATAFILE
test.dat
NUMBER_OF_TRAITS
2
NUMBER_OF_EFFECTS
2
OBSERVATION(S)
3 4
WEIGHT(S)
EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
1 1 2 cross
5 5 10 cross
RANDOM_RESIDUAL VALUES
10 1
1 5
RANDOM_GROUP
2
RANDOM_TYPE
add_animal
FILE
animal.ped
(CO)VARIANCES
1 0.1
0.1 1
```

Example 2: different model for each trait

```
DATAFILE
test.dat
NUMBER_OF_TRAITS
2
NUMBER_OF_EFFECTS
3
OBSERVATION(S)
3 4
WEIGHT(S)
EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
1 2 2 cross
5 5 10 cross
6 7 30 cross
RANDOM_RESIDUAL VALUES
10 1
1 5
RANDOM_GROUP
2
RANDOM_TYPE
add_animal
FILE
```

```
animal.ped
(CO)VARIANCES
1 0.1
0.1 1
RANDOM_GROUP
3
RANDOM_TYPE
diagonal
FILE

(CO)VARIANCES
1 0
0 1
```

Animal model with UPG

```
DATAFILE
test.dat
NUMBER_OF_TRAITS
2
NUMBER_OF_EFFECTS
2
OBSERVATION(S)
3 4
WEIGHT(S)
EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
1 1 2 cross
5 5 13 cross
RANDOM_RESIDUAL VALUES
10 1
1 5
RANDOM_GROUP
2
RANDOM_TYPE
add_an_upg
FILE
animal.ped
(CO)VARIANCES
1 0.1
0.1 1
```

Animal model with inbreeding

```
DATAFILE
test.dat
NUMBER_OF_TRAITS
2
NUMBER_OF_EFFECTS
2
OBSERVATION(S)
3 4
WEIGHT(S)
```

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
1 1 2 cross
5 5 13 cross
RANDOM_RESIDUAL VALUES
10 1
1 5
RANDOM_GROUP
2
RANDOM_TYPE
add_an_upginb
FILE
animal.ped
(CO)VARIANCES
1 0.1
0.1 1

Repeatability model 1

DATAFILE
test.dat
NUMBER_OF_TRAITS
1
NUMBER_OF_EFFECTS
3
OBSERVATION(S)
3
WEIGHT(S)
EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
1 2 cross
5 5 cross
5 10 cross
RANDOM_RESIDUAL VALUES
10
RANDOM_GROUP
2
RANDOM_TYPE
add_animal
FILE
animal.ped
(CO)VARIANCES
1
RANDOM_GROUP
3
RANDOM_TYPE
diagonal
FILE

(CO)VARIANCES
1

Repeatability model 2

```
DATAFILE
test.dat
NUMBER_OF_TRAITS
2
NUMBER_OF_EFFECTS
3
OBSERVATION(S)
3 4
WEIGHT(S)
EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
1 1 2 cross
5 5 5 cross
5 5 10 cross
RANDOM_RESIDUAL VALUES
10 1
1 5
RANDOM_GROUP
2
RANDOM_TYPE
add_animal
FILE
animal.ped
(CO)VARIANCES
1 0.1
0.1 1
RANDOM_GROUP
3
RANDOM_TYPE
diagonal
FILE

(CO)VARIANCES
1 0.1
0.1 1
```

Maternal effect model

```
DATAFILE
maternal.dat
NUMBER_OF_TRAITS
1
NUMBER_OF_EFFECTS
4
OBSERVATION(S)
4
WEIGHT(S)

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
3 946 cross
1 22473 cross
2 22473 cross
```

```
2 22473 cross
RANDOM_RESIDUAL VALUES
1050
RANDOM_GROUP
2 3
RANDOM_TYPE
add_animal
FILE
maternal.ped
(CO)VARIANCES
450 -100
-100 340
RANDOM_GROUP
4
RANDOM_TYPE
diagonal
FILE

(CO)VARIANCES
370
```

```
# For (THR)GIBBSxF90
# Example 1
DATAFILE
test.dat
NUMBER_OF_TRAITS
2
NUMBER_OF_EFFECTS
5
OBSERVATION(S)
3 4
WEIGHT(S)
EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
1 0 2 cross
0 2 2 cross
5 5 10 cross
6 0 30 cross
0 7 20 cross
RANDOM_RESIDUAL VALUES
10 1
1 5
RANDOM_GROUP
3
RANDOM_TYPE
add_animal
FILE
animal.ped
(CO)VARIANCES
1 0.1
0.1 1
```

```
RANDOM_GROUP
4
RANDOM_TYPE
diagonal
FILE

(CO)VARIANCES
1 0
0 0
RANDOM_GROUP
5
RANDOM_TYPE
diagonal
FILE

(CO)VARIANCES
0 0
0 1
```

Example 2

```
DATAFILE
test.dat
NUMBER_OF_TRAITS
2
NUMBER_OF_EFFECTS
5
OBSERVATION(S)
3 4
WEIGHT(S)
EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
1 0 2 cross
0 2 2 cross
5 5 10 cross
6 0 30 cross
0 7 30 cross
RANDOM_RESIDUAL VALUES
10 1
1 5
RANDOM_GROUP
3
RANDOM_TYPE
add_animal
FILE
animal.ped
(CO)VARIANCES
1 0.1
0.1 1
RANDOM_GROUP
4 5
RANDOM_TYPE
```

diagonal
FILE

(CO)VARIANCES
1 0 0 0
0 0 0 0
0 0 0 0
0 0 0 1

Dominance model

DATAFILE
dom.dat
NUMBER_OF_TRAITS
1
NUMBER_OF_EFFECTS
4
OBSERVATION(S)
3
WEIGHT(S)

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
1 1 cross
4 1 cov
2 30001 cross
5 10412 cross
RANDOM_RESIDUAL VALUES
100
RANDOM_GROUP
3
RANDOM_TYPE
add_an_upginb
FILE
add.ped
(CO)VARIANCES
10
RANDOM_GROUP
4
RANDOM_TYPE
par_dom
FILE
dom.ped
(CO)VARIANCES
2

Random regression model

Example 1

DATAFILE
data_score
NUMBER_OF_TRAITS
1

```
NUMBER_OF_EFFECTS
10
OBSERVATION(S)
9
WEIGHT(S)

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
 1    788 cross
 2     32 cross
 5      1 cov
 6      1 cov
 3  15097 cross
 5  15097 cov  3
 6  15097 cov  3
 3  81883 cross
 5  81883 cov  3
 6  81883 cov  3
RANDOM_RESIDUAL VALUES
 100
RANDOM_GROUP
5 6 7
RANDOM_TYPE
diagonal
FILE

(CO)VARIANCES
 100  1    1
 1    10   1
 1    1    10
RANDOM_GROUP
8 9 10
RANDOM_TYPE
add_an_upg
FILE
ped_score
(CO)VARIANCES
 100  1    1
 1    10   1
 1    1    10
```

Example 2

```
DATAFILE
test.dat1
NUMBER_OF_TRAITS
2
NUMBER_OF_EFFECTS
9
OBSERVATION(S)
3 4
WEIGHT(S)
```

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
1 1 2 cross
6 6 1 cov
7 7 1 cov
2 2 5 cross
6 6 5 cov 2 2
7 7 5 cov 2 2
2 2 10 cross
6 6 10 cov 2 2
7 7 10 cov 2 2
RANDOM_RESIDUAL VALUES
10 1
1 5
RANDOM_GROUP
4 5 6
RANDOM_TYPE
diagonal
FILE

(CO)VARIANCES
1 0.1 0.1 0.1 0.1 0.1
0.1 1 0.1 0.1 0.1 0.1
0.1 0.1 1 0.1 0.1 0.1
0.1 0.1 0.1 1 0.1 0.1
0.1 0.1 0.1 0.1 1 0.1
0.1 0.1 0.1 0.1 0.1 1
RANDOM_GROUP
7 8 9
RANDOM_TYPE
add_animal
FILE
animal.ped
(CO)VARIANCES
1 0.1 0.1 0.1 0.1 0.1
0.1 1 0.1 0.1 0.1 0.1
0.1 0.1 1 0.1 0.1 0.1
0.1 0.1 0.1 1 0.1 0.1
0.1 0.1 0.1 0.1 1 0.1
0.1 0.1 0.1 0.1 0.1 1

Example 3

DATAFILE
test.dat2
NUMBER_OF_TRAITS
2
NUMBER_OF_EFFECTS
10
OBSERVATION(S)
3 4
WEIGHT(S)

```
EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
1 1 2 cross
6 6 1 cov
7 7 1 cov
8 8 1 cov
6 6 5 cov 2 2
7 7 5 cov 2 2
8 8 5 cov 2 2
6 6 10 cov 2 2
7 7 10 cov 2 2
8 8 10 cov 2 2
RANDOM_RESIDUAL VALUES
10 1
1 5
RANDOM_GROUP
5 6 7
RANDOM_TYPE
diagonal
FILE

(CO)VARIANCES
1 0.1 0.1 0.1 0.1 0.1
0.1 1 0.1 0.1 0.1 0.1
0.1 0.1 1 0.1 0.1 0.1
0.1 0.1 0.1 1 0.1 0.1
0.1 0.1 0.1 0.1 1 0.1
0.1 0.1 0.1 0.1 0.1 1
RANDOM_GROUP
8 9 10
RANDOM_TYPE
add_animal
FILE
animal.ped
(CO)VARIANCES
1 0.1 0.1 0.1 0.1 0.1
0.1 1 0.1 0.1 0.1 0.1
0.1 0.1 1 0.1 0.1 0.1
0.1 0.1 0.1 1 0.1 0.1
0.1 0.1 0.1 0.1 1 0.1
0.1 0.1 0.1 0.1 0.1 1
```

Random regression model with heterogeneous residual variances

using airemlf90

Example 1: with intercept

```
DATAFILE
test.dat
NUMBER_OF_TRAITS
1
NUMBER_OF_EFFECTS
```

```
9
OBSERVATION(S)
3
WEIGHT(S)
EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
1 2 cross
6 1 cov
7 1 cov
5 5 cross
6 5 cov 5
7 5 cov 5
5 10 cross
6 10 cov 5
7 10 cov 5
RANDOM_RESIDUAL VALUES
10
RANDOM_GROUP
4 5 6
RANDOM_TYPE
diagonal
FILE

(CO)VARIANCES
1 0.1 0.1
0.1 1 0.1
0.1 0.1 1
RANDOM_GROUP
7 8 9
RANDOM_TYPE
add_animal
FILE
animal.ped
(CO)VARIANCES
1 0.1 0.1
0.1 1 0.1
0.1 0.1 1
OPTION hetres_pos 6 7
OPTION hetres_pol 4.0 1.0 0.1
```

 # Example 2: with no intercept

```
DATAFILE
test.dat
NUMBER_OF_TRAITS
1
NUMBER_OF_EFFECTS
7
OBSERVATION(S)
3
WEIGHT(S)
EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
```

1 2 cross
6 1 cov
7 1 cov
6 5 cov 5
7 5 cov 5
6 10 cov 5
7 10 cov 5
RANDOM_RESIDUAL VALUES
10
RANDOM_GROUP
4 5
RANDOM_TYPE
diagonal
FILE

(CO)VARIANCES
1 0.1
0.1 1
RANDOM_GROUP
6 7
RANDOM_TYPE
add_animal
FILE
animal.ped
(CO)VARIANCES
1 0.1
0.1 1
OPTION hetres_pos 6 7
OPTION hetres_pol 1.0 0.1

using GIBBS3F90
DATAFILE
test.dat
NUMBER_OF_TRAITS
1
NUMBER_OF_EFFECTS
9
OBSERVATION(S)
3
WEIGHT(S)
EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
1 2 cross
6 1 cov
7 1 cov
5 5 cross
6 5 cov 5
7 5 cov 5
5 10 cross
6 10 cov 5
7 10 cov 5

```
RANDOM_RESIDUAL VALUES
10
RANDOM_GROUP
4 5 6
RANDOM_TYPE
diagonal
FILE

(CO)VARIANCES
1 0.1 0.1
0.1 1 0.1
0.1 0.1 1
RANDOM_GROUP
7 8 9
RANDOM_TYPE
add_animal
FILE
animal.ped
(CO)VARIANCES
1 0.1 0.1
0.1 1 0.1
0.1 0.1 1
OPTION hetres_int 8 5
```

Competitive model

```
DATAFILE
competition.dat
NUMBER_OF_TRAITS
1
NUMBER_OF_EFFECTS
19
OBSERVATION(S)
24
WEIGHT(S)

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT[EFFECT NESTED]
2 88 cross
3 362 cross
21 2409 cross
4 8004 cross
22 0 cov 5
22 0 cov 6
22 0 cov 7
22 0 cov 8
22 0 cov 9
22 0 cov 10
22 0 cov 11
22 0 cov 12
22 0 cov 13
22 0 cov 14
```

22 0 cov 15
22 0 cov 16
22 0 cov 17
22 0 cov 18
22 8004 cov 19
RANDOM_RESIDUAL VALUES
1225.8
RANDOM_GROUP
4 5
RANDOM_TYPE
add_animal
FILE
renadd04.ped
(CO)VARIANCES
267.03 25.313
25.313 104.44
RANDOM_GROUP
2
RANDOM_TYPE
diagonal
FILE

(CO)VARIANCES
89.187
RANDOM_GROUP
3
RANDOM_TYPE
diagonal
FILE

(CO)VARIANCES
167.34

Appendix A (single trait animal model)

Single trait "USDA-type" animal model. This example is from the documentation of program JAA20.

$$y_{ijkl} = hys_i + hs_{ij} + p_k + a_k + e_{ijkl}$$

where

y_{ijkl} - production yield
hys_i - fixed herd year season
hs_{ij} - random herd x sire interaction
p_k- random permanent environment
a_k - random animal

and

var(hs_{ij}) = .05, var(p_k)=.1, var(a_k)=.5, var(e_{ijkl})=1

Data file (ic)

Format: animal/hys/p/hs/y

```
 1 1 1 1 10
 2 1 2 1 11
 3 2 3 2 15
 4 2 4 3 13
 5 3 5 4 14
 6 3 6 3 12
```

Relationship file (is)

Format: animal/dam/sire/code

```
 1  12  8 2
 2   1  8 1
 3   2  9 1
 4   7 10 1
 5  12 11 2
 6   1 10 1
 7  13 14 3
 8   5 11 1
 9  13  8 2
10   7 14 2
11 13 14 3
```

Parameter file

Example of single-trait animal model with one fixed effect

```
DATAFILE
ic
NUMBER_OF_TRAITS
1
NUMBER_OF_EFFECTS
4
OBSERVATION(S)
5
WEIGHT(S)
```

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT NESTED]
2 3 cross
3 6 cross
4 4 cross
1 14 cross
RANDOM_RESIDUAL VALUES
1
RANDOM_GROUP
2
RANDOM_TYPE
diagonal
FILE

(CO)VARIANCES
.1
RANDOM_GROUP
3
RANDOM_TYPE
diagonal
FILE

(CO)VARIANCES
.05
RANDOM_GROUP
4
RANDOM_TYPE
add_an_upg
FILE
is
(CO)VARIANCES
.5

Execution

```
/home/ignacy/f90/examples blupf90
 name of parameter file?exiap

     BLUPF90 1.00

 Parameter file:             exiap
 Data file:                  ic
 Number of Traits             1
 Number of Effects            4
 Position of Observations      5
 Position of Weight (1)        0
 Value of Missing Trait/Observation          0

EFFECTS
 #  type                position (2)        levels   [positions for nested]
 1  cross-classified      2                    3
 2  cross-classified      3                    6
 3  cross-classified      4                    4
 4  cross-classified      1                   14
```

```
 Residual (co)variance Matrix
     1.000
 Random Effect      2
 Type of Random Effect:      diagonal
 trait   effect    (CO)VARIANCES
  1       2       0.100

 Random Effect      3
 Type of Random Effect:      diagonal
 trait   effect    (CO)VARIANCES
  1       3       0.050

 Random Effect      4
 Type of Random Effect:      additive animal
 Pedigree File:              is
 trait   effect    (CO)VARIANCES
  1       4       0.500

 REMARKS
  (1) Weight position 0 means no weights utilized
  (2) Effect positions of 0 for some effects and traits means that such
      effects are missing for specified traits

 Data record length = 5
 original G
   0.10
 inverted G
  10.00
 original G
   0.05
 inverted G
  20.00
 original G
   0.50
 inverted G
   2.00
 solutions stored in file: "solutions"

/home/ignacy/f90/examples cat solutions
trait/effect level  solution
  1  1       1     11.8589
  1  1       2     13.7539
  1  1       3     14.7086
  1  2       1     -0.0088
  1  2       2      0.0088
  1  2       3     -0.0159
  1  2       4      0.0159
  1  2       5      0.0321
  1  2       6     -0.0321
  1  3       1      0.0000
  1  3       2     -0.0079
  1  3       3     -0.0081
  1  3       4      0.0161
  1  4       1     -1.7627
  1  4       2     -0.9553
  1  4       3      1.4288
  1  4       4     -0.9206
  1  4       5     -1.0781
  1  4       6     -2.3474
  1  4       7      0.8511
```

```
1  4       8     -0.1521
1  4       9      3.8926
1  4      10     -2.7717
1  4      11      0.8528
1  4      12     -3.1911
1  4      13      7.9976
1  4      14     -6.3340
```

Appendix B (multiple trait sire model)

Example of multiple trait sire model (from L.R. Schaeffer notes of 1985).

Models

Trait 1: $y_{1i}=h_i+s_{1j}+e_{1ijk}$

Trait 2: $y_{2i}=\mu+s_{2j}+e_{2jk}$

where

h - fixed herd

s - random sire

and

var(s)=A[8 6; 6 17], var(e)=I[10 10; 10 20]

Data file (lrsdat)

Format: h/μ/s/y_1/y_2

```
1 0 1 3.4 0
2 0 2 1.3 0
1 1 3 .8 50.3
2 1 4 4.5 52.6
0 1 5  0 55.0
```

Pedigree file (lrsrel)

Format: bull/sire/MGS

```
1 3 0
2 0 5
3 0 0
4 0 0
5 0 0
```

Parameter file (lrsex)

```
# Example of two trait sire model with unequal models
DATAFILE
lrsdat
NUMBER_OF_TRAITS
2
NUMBER_OF_EFFECTS
2
OBSERVATION(S)
4 5
WEIGHT(S)

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT  [EFFECT NESTED]
```

```
1 2 2 cross
3 3 5 cross
RANDOM_RESIDUAL VALUES
10 10
10 20
RANDOM_GROUP
2
RANDOM_TYPE
add_sire
FILE
lrsrel
(CO)VARIANCES
8 6
6 17
```

Execution

```
/home/ignacy/f90/examples blupf90
 name of parameter file?lrsex

     BLUPF90 1.00

 Parameter file:             lrsex
 Data file:                  lrsdat
 Number of Traits             2
 Number of Effects            2
 Position of Observations      4  5
 Position of Weight (1)        0
 Value of Missing Trait/Observation           0

EFFECTS
 #  type                position (2)        levels   [positions for nested]
 1  cross-classified      1  2                  2
 2  cross-classified      3  3                  5


 Residual (co)variance Matrix
    10.000    10.000
    10.000    20.000

 Random Effect      1
 Type of Random Effect:      additive sire
 Pedigree File:              lrsrel
 trait   effect    (CO)VARIANCES
  1       2       8.000     6.000
  2       2       6.000    17.000

 REMARKS
  (1) Weight position 0 means no weights utilized
  (2) Effect positions of 0 for some effects and traits means that such
      effects are missing for specified traits

 Data record length = 5
 original G
   8.00   6.00
   6.00  17.00
 inverted G
   0.17  -0.06
  -0.06   0.08
```

```
solutions stored in file: "solutions"

/home/ignacy/f90/examples cat solutions
trait/effect level  solution
  1  1       1      2.3877
  2  1       1     52.4449
  1  1       2      3.2180
  2  1       2      0.0000
  1  2       1      0.2243
  2  2       1     -0.0210
  1  2       2     -0.8217
  2  2       2     -0.3866
  1  2       3     -0.4969
  2  2       3     -0.7512
  1  2       4      0.6178
  2  2       4     -0.0769
  1  2       5      0.2217
  2  2       5      1.0851
```

Appendix C (test-day model)

This test-day model example comes from the paper of Schaeffer and Dekkers (WCGALP94 18:443)

Model

$$y_{ijkl} = h_i + \beta_1 X_{1j} + \beta_2 X_{2j} + a_k + \gamma_{1k} X_{1j} + \gamma_{2k} X_{2j} + e_{ijkl}$$

where

y_{ijkl} - yield of test day

h_i - test day effect

X_{1j} - days in milk

X_{2j} - log(days in milk)

β_1, β_2 - fixed regressions

a_k - random animal

γ_{1k}, γ_{2k} - random regressions for each animal

and

$$var(e_{ijkl}) = 1; \; var(a_k, \gamma_{1k}, \gamma_{2k}) = [\, 2.25 \; 4 \; -.7; \; 4 \; 1375 \; 12; \; -.7 \; 12 \; 94]^{-1}$$

Data file (lrsrrdat)

Format: h/a/X_1/X_2/y

```
1 1 73 1.42985 26
1 2 34 2.19395 29
1 3 8 3.64087 37
2 1 123 0.908127 23
2 2 84 1.28949 18
2 3 58 1.65987 25
2 4 5 4.11087 44
3 1 178 0.538528 21
3 2 139 0.785838 8
3 3 113 0.992924 19
3 4 60 1.62597 29
4 2 184 0.505376 1
4 3 158 0.657717 15
4 4 105 1.06635 22
4 5 14 3.08125 35
5 3 218 0.335817 11
5 4 165 0.614366 14
5 5 74 1.41625 23
5 6 31 2.28632 28
6 3 268 0.129325 7
6 4 215 0.349674 8
6 5 124 0.90003 17
6 6 81 1.32586 22
```

Relationship file (lrsrrrel)

Format: animal/sire/dam

```
1 9 7
2 10 8
3 9 2
4 10 8
```

```
5 11 7
6 11 1
7 0 0
8 0 0
9 0 0
10 0 0
11 0 0
```

Parameter file (exlrsrr)

```
# Example of single-trait random-regression model
DATAFILE
lrsrrdat
NUMBER_OF_TRAITS
1
NUMBER_OF_EFFECTS
6
OBSERVATION(S)
5
WEIGHT(S)

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT  [EFFECT NESTED]
1 6 cross
3 1 cov
4 1 cov
2 11 cross
3 11 cov 2
4 11 cov 2
RANDOM_RESIDUAL VALUES
1
RANDOM_GROUP
4 5 6
RANDOM_TYPE
add_animal
FILE
lrsrrrel
(CO)VARIANCES
 .447906  -0.001334   0.003506
-0.001334   0.000732  -0.000103
 0.003506  -0.000103   .010678
```

Execution

```
/home/ignacy/f90/examples blupf90
 name of parameter file?exlrsrr

     BLUPF90 1.00

 Parameter file:             exlrsrr
 Data file:                  lrsrrdat
 Number of Traits             1
 Number of Effects            6
 Position of Observations      5
 Position of Weight (1)        0
 Value of Missing Trait/Observation           0
```

```
EFFECTS
 #  type                position (2)        levels   [positions for nested]
 1  cross-classified      1                    6
 2  covariable            3                    1
 3  covariable            4                    1
 4  cross-classified      2                   11
 5  covariable            3                   11     2
 6  covariable            4                   11     2

 Residual (co)variance Matrix
     1.000

 correlated random effects     4  5  6
 Type of Random Effect:      additive animal
 Pedigree File:              lrsrrrel
 trait   effect    (CO)VARIANCES
  1       4       0.448    -0.001     0.004
  1       5      -0.001     0.001     0.000
  1       6       0.004     0.000     0.011

 REMARKS
  (1) Weight position 0 means no weights utilized
  (2) Effect positions of 0 for some effects and traits means that such
      effects are missing for specified traits

 Data record length = 5
 original G
   0.45   0.00   0.00
   0.00   0.00   0.00
   0.00   0.00   0.01
 inverted G
   2.25   4.00  -0.70
   4.001375.09  11.95
  -0.70  11.95  94.00
 solutions stored in file: "solutions"

/home/ignacy/f90/examples cat solutions
trait/effect level  solution
  1  1       1     19.9496
  1  1       2     20.3729
  1  1       3     20.6095
  1  1       4     19.7278
  1  1       5     18.6035
  1  1       6     17.8500
  1  2       1     -0.0498
  1  3       1      5.2912
  1  4       1     -0.4430
  1  4       2      0.2704
  1  4       3     -0.7288
  1  4       4      1.1019
  1  4       5     -0.1626
  1  4       6     -0.4828
  1  4       7     -0.0988
  1  4       8      0.4574
  1  4       9     -0.6288
  1  4      10      0.4574
  1  4      11     -0.1872
  1  5       1      0.0369
  1  5       2     -0.0661
  1  5       3      0.0068
  1  5       4     -0.0054
  1  5       5      0.0069
```

```
1  5       6      0.0167
1  5       7      0.0133
1  5       8     -0.0238
1  5       9      0.0350
1  5      10     -0.0238
1  5      11     -0.0008
1  6       1     -0.0370
1  6       2      0.0325
1  6       3     -0.0479
1  6       4      0.0767
1  6       5     -0.0149
1  6       6     -0.0377
1  6       7     -0.0103
1  6       8      0.0364
1  6       9     -0.0480
1  6      10      0.0364
1  6      11     -0.0145
```

Appendix D (multibreed maternal effect model)

This model was used for studies on multibreed evaluation in beef cattle. It is provided as an example of a model with maternal effect and different models per trait.

Model (in concise form, with most indices omitted)

$y_1 = cg_1 + bt + mbt + a + M + e$

$y_2 = cg_2 + bt + mbt + a + M + pe + e$

$y_3 = cg_3 + bt + mbt + a + e$

where

y_{1-3} - birth weight, weaning weight, and gain

cg_{1-3} - contemporary groups separate for each trait

br - breed type

mbt - maternal breed type

a - additive effect

m - maternal effect

pe - permanent environmental effect of the dam

Data file (data.out)

Format:

1. contemporary group for trait 1
2. contemporary group for trait 2
3. contemporary group for trait 3
4. animal breed type
5. maternal breed type
6. animal id
7. dam id
8. birth weight
9. weaning weight
10. gain

Relationship file (pedi.outok)

Format:

animal

sire or unknown parent group

dam or unknown parent group

"1 + number of missing parents"

Parameter file (exlrsrr)

```
DATAFILE
data.out
NUMBER_OF_TRAITS
3
NUMBER_OF_EFFECTS
6
OBSERVATION(S)
8 9 10
WEIGHT(S)


EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT [EFFECT
NESTED]
 1 2 3   133085 cross
 4 4 4     181 cross
 5 5 0     165 cross
 6 6 6  1724112 cross
 7 7 0  1724112 cross
 0 7 0  1724112 cross
RANDOM_RESIDUAL VALUES
26.3        0.0         0.0
0.0         1312.9      0.0
0.0         0.0         1246.3
RANDOM_GROUP
4 5
RANDOM_TYPE
add_an_upg
FILE
pedi.outok
(CO)VARIANCES
22.9        36.3        18.6        -4.6        0.0         0.0
36.6        500.2       110.8       0.0         -91.6       0.0
18.6        110.8       313.0       0.0         0.0         0.0
-4.6        0.0         0.0         10.1        0.0         0.0
0.0         -91.6       0.0         0.0         419.1       0.0
0.0         0.0         0.0         0.0         0.0         0.0
RANDOM_GROUP
2
RANDOM_TYPE
diagonal
FILE

(CO)VARIANCES
0.263       0.0         0.0
0.0         13.129      0.0
0.0         0.0         12.463
RANDOM_GROUP
```

3
RANDOM_TYPE
diagonal
FILE

(CO)VARIANCES
0.263 0.0 0.0
0.0 13.129 0.0
0.0 0.0 0.0
RANDOM_GROUP
6
RANDOM_TYPE
diagonal
FILE

(CO)VARIANCES
0.0 0.0 0.0
0.0 45.5 0.0
0.0 0.0 0.0

Appendix E (random regression model)

A single-trait random regression model for test-day milk is using cubic Legendre polynomials.

Model

func{

y_ijkl~=~hym_ij~+~sum from {m=1} to 4 alpha_m(l) h_im~+~

~sum from {m=1} to 4 alpha_m(l) u_km~+~

~sum from {m=1} to 4 alpha_m(l) p_im~+~e_ijkl

}

where

y_{ijkl} - test day milk

hym_{ij} - hear-year-test for herd i and year-test j

h_i - effects of herd i

$\alpha_m(l)$ - value of m-th Legendre polynomial at point corresponding to DIM=l

u - additive effects

pe - permanent environmental effects

Data file (datarr)

Format:

1.herd

2. hear-year-test

3-6. values of Legendre polynomials

7. weight for residuals: $100/var(e_{ijkl})$

8. test day

9. animal

Relationship file (pedirr)

Format:

animal

sire

dam

Parameter file (exrr3)

DATAFILE

datarr

NUMBER_OF_TRAITS

1

NUMBER_OF_EFFECTS

```
13
OBSERVATION(S)
8
WEIGHT(S)
7
EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT
2   3726 cross        #herd-year-test
3 84 cov 1                    #herd
4 84 cov 1
5 84 cov 1
6 84 cov 1
3 21874 cov 9                 #additive
4 21874 cov 9
5 21874 cov 9
6 21874 cov 9
3 21874 cov 9                 #pe
4 21874 cov 9
5 21874 cov 9
6 21874 cov 9
RANDOM_RESIDUAL VALUES
100
RANDOM_GROUP
6 7 8 9
RANDOM_TYPE
add_animal
FILE
pedirr
(CO)VARIANCES
(4 x 4 matrix)
RANDOM_GROUP
10 11 12 13
RANDOM_TYPE
diagonal
FILE

(CO)VARIANCES
(4 x 4 matrix)
```

Appendix F (terminal cross model)

A terminal cross model by Fernando et al. and Lo et al.

breed A: ya=cga + ua + ea
breed B: yb=cgb+ ub +eb
cross: yab=cgab+ uaab + ubab +eab

Data file (data_cross)
1. cg A (85 levels)
2. cg B (110 levels)
3. cg crossbred (87 levels)
4. animal - breed A (2400 animals) or parent from breed A
5. animal - breed B (3000 animals) or parent from breed B
7. ya
8. yb
9. yc

Pedigree files: pedig_A for breed A and pedig_B for breed B

Parameter file

```
# Example of a terminal-cross model
DATAFILE
data-cross
NUMBER_OF_TRAITS
3
NUMBER_OF_EFFECTS
3
OBSERVATION(S)
6 7 8
WEIGHT(S)

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT  [EFFECT NESTED]
1 2 3 110 cross
4 0 4 2400 cross
0 5 5 3000 cross
RANDOM_RESIDUAL VALUES
100 0 0
0 100 0
0 0 100
RANDOM_GROUP
2
RANDOM_TYPE
add_animal
FILE
pedig_A
```

(CO)VARIANCES
(9 x 9 matrix)
RANDOM_GROUP
3
RANDOM_TYPE
add_animal
FILE
pedig_B
(CO)VARIANCES
(9 x 9 matrix)

Appendix G (competitive model)

Example of a competitive model (a la Muir and Schinkel)

y=cg + a +c1 + c2 +..+c5 + e

ci is the effect of the i-th competitor; assumed pen size of up to 6.

Datafile (data_comp)

1. y
2. cg (max 120)
3. animal (max 3000)
4. competitor 1
5. c 2

...

8. c 5

If pen size is less than 10, unused fields set to 0.

Parameter file

```
# Example of a competitive model
DATAFILE
data_comp
NUMBER_OF_TRAITS
1
NUMBER_OF_EFFECTS
7
OBSERVATION(S)
1
WEIGHT(S)

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT  [EFFECT NESTED]
2 120 cross
3 3000 cross
4 0 cross
5 0 cross
6 0 cross
7 0 cross
8 3000 cross
RANDOM_RESIDUAL VALUES
50
RANDOM_GROUP
2 3
RANDOM_TYPE
```

The 2nd effect (position 3 in the data) is additive direct effect and 3rd to 7th effects (positions 4 to 8 in the data) are competitive effects (animal ID for competitors).

```
add_animal
FILE
pedig
(CO)VARIANCES
40 -10
-10 10
```

The covariance matrix contains variance for the second effect, variance for effects 3 to 7 (accumulated to 7), and covariance between direct and competitive effects.

Appendix H (genomic model)

Example of evaluation /variance component estimation using phenotypic, pedigree and genomic information in single-step evaluation

Files simulated by Huiyu Wang using program QMSim by Mehdi Sargolzaei.

Parameter file for renumbering program RENUMF90

```
DATAFILE
phenotypes.txt
TRAITS
3
FIELDS_PASSED TO OUTPUT

WEIGHT(S)

RESIDUAL_VARIANCE
 0.9038
EFFECT
1 cross alpha  #fixed effect
EFFECT
2 cross alpha  #animal
RANDOM
animal
FILE
pedigree
SNP_FILE
marker.geno.clean
 (CO)VARIANCES
   0.9951E-01
```

Phenotypes.txt – phenotype file
Single trait in position 3
Fixed effect in position 1 read as alphanumeric
Random animal effect in position 3
Pedigree file pedigrees
SNP file marker.geno.clean

Phenotype file

phenotypes.txt

```
1 1 4.16 0
1 2 3.47 0
1 3 4.5 0
1 4 4.97 0
1 5 5.98 0
1 6 6.63 0
1 7 3.32 0
1 8 5.85 0
1 9 4.77 0
1 10 4.22 0
```

Pedigree file

pedigree

```
1 0 0 0
2 0 0 0
3 0 0 0
4 0 0 0
5 0 0 0
```

```
6 0 0 0
7 0 0 0
8 0 0 0
9 0 0 0
10 0 0 0
```

SNP file for the first 50 SNP

```
cut -c1-50 marker.geno.clean|head -10
```

```
8002 21101011002012011011010110111111211111210100
8014 21110101111101120221110111111112101112210100
8016 21100101202202021120210121102111202212111101
8018 21110111112201120210200020101022212211111100
8024 21110102201201111202101111021222012211111111
8038 11110000102100120202121112120102211211121111
8041 22210001201201121110210121202111102102121001
8063 20110101202202020212211101101120222012120021
8065 21110101111112111221110101010220212001110012
8083 10111011110010111111111011210011112101101012
```

Run RENUMF90

```
RENUMF90 version 1.86
name of parameter file?renum.par
renum.par
datafile:phenotypes.txt
traits:            3
fields passed:            4
R
 0.9038

Processing effect  1 of type cross
item_kind=alpha

Processing effect  2 of type cross
item_kind=alpha
pedigree file name  "pedigree"
positions of animal, sire, dam, alternate dam and yob           1           2
          3           0           0
SNP file name  "marker.geno.clean"
all pedigrees to be included
Reading (CO)VARIANCES:           1 x           1

Maximum size of character fields: 20

Maximum size of record (max_string_readline): 800

Maximum number of fields innput file (max_field_readline): 100

hash tables for effects set up
table expanded from        10000  to        20000  records
table expanded from        20000  to        40000  records
read        15800  records
table with            1  elements sorted
added count
Effect group            1  of column            1  with            1  levels
table expanded from        10000  to        10000  records
added count
Effect group            2  of column            1  with        15800  levels
wrote statistics in file "renf90.tables"

Basic statistics for input data  (missing value code is 0)
Pos  Min         Max         Mean        SD                N
  3   0.73000      8.8300      4.9793      1.0069       15800
```

```
random effect with SNPs  2
type: animal
file: marker.geno.clean
read SNPs        1500  records
Effect group            2  of column            1  with        15800  levels

random effect   2
type:animal
opened output pedigree file "renadd02.ped"
read        15800  pedigree records

Pedigree checks

Number of animals with records:       15800
Number of animals with genotypes:        1500
Number of animals with records or genotypes:       15800
Number of animals with genotypes and no records            0
Number of parents without records or genotypes:            0
Total number of animals:       15800

Wrote cross reference IDs for SNP file "marker.geno.clean_XrefID"

Wrote parameter file "renf90.par"
Wrote renumbered data "renf90.dat"
```

Parameter file for application programs with renumbered fields

renf90.par

```
# BLUPF90 parameter file created by RENF90
DATAFILE
 renf90.dat
NUMBER_OF_TRAITS
       1
NUMBER_OF_EFFECTS
       2
OBSERVATION(S)
  1
WEIGHT(S)

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT[EFFECT NESTED]
 2       1 cross
 3    15800 cross
RANDOM_RESIDUAL VALUES
 0.9038
 RANDOM_GROUP
       2
 RANDOM_TYPE
 add_animal
 FILE
renadd02.ped
(CO)VARIANCES
 0.9951E-01
OPTION SNP_file marker.geno.clean
```

renf90.dat – phenotype file
Single trait in position 1
Two effects in model
Fixed effect in position 1 cross-classified with 1 level (μ)
Animal effect in position 3
Second effect (Random Group 2) is additive-animal with
renadd02.ped – pedigree file
SNP file marker.geno.clean

Renumbered pedigree file

```
renadd02.ped
```

```
1 5742 14705 1 0 2 1 0 0 14670
2 2302 1384 1 0 2 1 0 0 12367
3 4248 15309 1 0 12 1 0 2 9123
4 4241 3492 1 0 2 1 0 0 7455
5 14459 14202 1 0 2 1 0 0 5736
6 1029 1292 1 0 2 1 0 3 5877
7 10876 7596 1 0 2 1 0 0 9638
8 13589 12642 1 0 2 1 0 0 14136
9 7070 11562 1 0 2 1 0 0 6010
10 6449 2448 1 0 2 1 0 0 15498
```

Renumbered phenotype file

renf90.dat

```
4.16 1 5903 0
3.47 1 3628 0
4.5 1 1329 0
4.97 1 14808 0
5.98 1 12481 0
6.63 1 10205 0
3.32 1 7935 0
5.85 1 5639 0
4.77 1 3348 0
4.22 1 1951 0
```

Run BLUPF90

```
name of parameter file?renf90.par

* SNP file: marker.geno.clean
* SNP Xref file: marker.geno.clean_XrefID
* Frequency to Center Z=M-p to create G=ZZ'/k (default whichfreq = 2):
          2
    BLUPF90 1.42

Parameter file:             renf90.par
Data file:                  renf90.dat
Number of Traits             1
Number of Effects            2
Position of Observations      1
Position of Weight (1)        0
Value of Missing Trait/Observation           0

EFFECTS
 #  type                position (2)        levels   [positions for nested]
 1  cross-classified     2                                                1
 2  cross-classified     3                                            15800

 Residual (co)variance Matrix
 0.90380

 Random Effect(s)    2
 Type of Random Effect:      additive animal
 Pedigree File:              renadd02.ped
 trait   effect    (CO)VARIANCES
  1       2    0.9951E-01

 REMARKS
  (1) Weight position 0 means no weights utilized
  (2) Effect positions of 0 for some effects and traits means that such
      effects are missing for specified traits

 Data record length =            3
 # equations =        15801
 G
 0.99510E-01
```

```
read        15800  records in   3.5994001E-02  s,         31601  nonzeroes
 read        15800  additive pedigrees

*-------------------------------------------------------------*
*                 Setup Genomic: Version 1.76                 *
*                                                             *
*  Modified relationship matrix (H) created for effect:   2   *
*-------------------------------------------------------------*

Read 15800 animals from pedigree file
Pedigree was in not chronological order (parent first format), reodering will be performed!!!

Current OPTIONS

Genomic Matrix
  Make/Read   Which  Save Test File       StorageType
     Make        1     F     F   G  densem

Rel. Matrix A22
  Make/Read   Which  Save Test File       StorageType
     Make        4     F     F   A22  densem

Inv. Genomic Matrix
  Make/Read   Which  Save Test File       StorageType
     Make        9     F     F   Gi  densem

Inv. Rel. Matrix A22
  Make/Read   Which  Save Test File       StorageType
     Make        9     F     F   A22i  densem

Genomic - A22 Matrix
  Make/Read   Which  Save Test File       StorageType
     None        9     F     F   GmA22  densem

Inv. Genomic- A22 Matrix
  Make/Read   Which  Save Test File       StorageType
     Make        0     F     F   GimA22i  densem

Other options
   Allele Frequency file:    freqdata
   Center Allele Frequency:  2
   Scale  Allele Frequency:  2
   Scale  Method:            1
   Regression G on A:              F
   Tuned G Method:          2

   Creation of GimA22i
      tau inv(alpha G + beta A22 + gamma I + delta) - omega inv(A22)
            alpha,beta        0.950   0.050
            gamma,delta       0.000   0.000
            tau,omega         1.000   1.000

Number of Genotyped Animals        1500

Creating A22
   Extracting subset of: 3432 pedigrees from: 15800 elapsed time:     0.0000
   Calculating Inbreeding by M&L function.. elapsed time  1.0000020E-03
   Calculating A22 Matrix by Colleau ...elapsed time  0.3299500
```

Statistics for A22

```
Statistic of Rel. Matrix A22
                         N       Mean        Min        Max        Var
    Diagonal          1500      1.001      1.000      1.250      0.000
    Off-diagonal   2248500      0.003      0.000      0.750      0.001
```

Statistics for SNP file

```
Reading SNP file
```

```
   Column position in file for the first marker:            7
   Format to read SNP file: (6x,400000i1)
   Number of SNPs :        3000
   Number of Genotyped animals:         1500
   Reading SNP file elapsed time  0.4639290

Statistics of alleles frequencies in the current population
   N:           3000
   Mean:       0.501
   Min:        0.132
   Max:        0.890
   Var:        0.014
```

Several quality checks performed; no error messages as all files for this example have been simulated

```
Quality Control - Check call rate for animals


Quality Control - Check Parent-Progeny Mendelian conflicts
   Total animals: 15800 - Genotyped animals: 1500
   Number of Individual - Sire pairs: 470
   Number of Individual - Dams pairs: 256
   Number of Individual - Sire - Dam trios: 152

Checking SNPs for Mendelian conflicts
Total number of parent-progeny evaluations:          726
Number of SNPs with Mendelian conflicts: 0

Checking Animals for Mendelian conflicts

Statistics of alleles frequencies in the current population after
    Quality Control (MAF, monomorphic, call rate)
   N:           3000
   Mean:       0.501
   Min:        0.132
   Max:        0.890
   Var:        0.014

    Locus      Freq      0-2p      1-2p      2-2p
        1  0.751333 -1.502667 -0.502667  0.497333
        2  0.382333 -0.764667  0.235333  1.235333
        3  0.568667 -1.137333 -0.137333  0.862667
        4  0.680000 -1.360000 -0.360000  0.640000
        5  0.184333 -0.368667  0.631333  1.631333
        6  0.298333 -0.596667  0.403333  1.403333
        7  0.392000 -0.784000  0.216000  1.216000
        8  0.379667 -0.759333  0.240667  1.240667
        9  0.596667 -1.193333 -0.193333  0.806667
       10  0.352333 -0.704667  0.295333  1.295333
Genotypes missings (%):  0.0000000E+00

Average denom. (scale):   1415.90178466665
Center Matrix elapsed:   8.3986998E-02

Creating G Matrix

Calculating G Matrix
   Wall time: 08-05-2011  16h 57m 34s 213
MMP - OPTML
Elapsed time   18.47419
   Wall time: 08-05-2011  16h 58m 09s 371
```

Statistics of G calculated assuming current allele frequencies

```
Statistic of Genomic Matrix
                          N      Mean       Min       Max       Var
    Diagonal           1500     0.999     0.889     1.463     0.002
    Off-diagonal    2248500    -0.001    -0.147     0.830     0.002


Correlation of Genomic Inbreeding and Pedigree Inbreeding
```

```
    Correlation:     0.3220

All elements - Diagonal / Off-Diagonal
   Estimating Regression Coefficients G = b0 11' + b1 A + e
   Regression coefficients b0 b1 =     -0.004     0.997

   Correlation all elements G & A     0.644



Off-Diagonal
   Using 70386 elements from A22 >= 0.02000

   Estimating Regression Coefficients G = b0 11' + b1 A + e
   Regression coefficients b0 b1 =     -0.006     1.000

   Correlation Off-Diagonal elements G & A     0.660


Blend G as alpha*G + beta*A22: (alpha,beta)     0.950     0.050

Statistic of Genomic Matrix
                           N      Mean       Min       Max       Var
    Diagonal            1500     0.999     0.894     1.446     0.002
    Off-diagonal     2248500     0.000    -0.139     0.820     0.002

Frequency - Diagonal of G
   N:        1500
   Mean:        0.999
   Min:         0.894
   Max:         1.446
   Range:       0.028
   Class:     20

 #Class       Class   Count
      1   0.8942           9
      2   0.9218          86
      3   0.9494         343
      4   0.9770         480
      5    1.005         361
      6    1.032         139
      7    1.060          51
      8    1.087          16
      9    1.115           6
     10    1.142           2
     11    1.170           1
     12    1.198           1
     13    1.225           1
     14    1.253           1
     15    1.280           0
     16    1.308           0
     17    1.336           0
     18    1.363           2
     19    1.391           0
     20    1.418           1
     21    1.446           0


Scale G matrix according to A22 -  Method: 2
   Diagonal A:     1.001    Offdiagonal A:     0.003    All A:     0.004    Difference:    0.998
   Diagonal G:     0.999    Offdiagonal G:     0.000    All G:     0.000    Difference:    0.999
   Diff G Diag - G OffDiag:     0.999    (da-oa)/(dg-og):     0.998
   Diff A OffDiag - G OffDiag:     0.004
   Diff A all - G all:     0.004
   New Alpha:    0.948   New Beta:    0.050   :New Delta    0.004

------------------------------
 Final Pedrigree-Based Matrix
------------------------------

Statistic of Rel. Matrix A22
```

Correlations of off-diagonal elements of G and A22 is 0.660;
low numbers indicated genotyped mistakes or poor pedigrees

Diagonal elements of G should be 1± 0.2. Too large or too small elements indicate:
Genotyping mistakes
Mixed lines
See Simeone et al. (2011)

```
                          N      Mean       Min       Max       Var
    Diagonal           1500     1.001     1.000     1.250     0.000
    Off-diagonal    2248500     0.003     0.000     0.750     0.001
```

Statistics of G after scaling as in Chen et al (2011) or Vitezica et al. (2011)
Statistics should be same as for A22.

```
----------------------
 Final Genomic Matrix
----------------------

Statistic of Genomic Matrix
                          N      Mean       Min       Max       Var
    Diagonal           1500     1.001     0.896     1.447     0.002
    Off-diagonal    2248500     0.003    -0.134     0.822     0.002


Correlation of Genomic Inbreeding and Pedigree Inbreeding
    Correlation:     0.3363

All elements - Diagonal / Off-Diagonal
   Estimating Regression Coefficients G = b0 11' + b1 A + e
   Regression coefficients b0 b1 =      0.000     0.995

   Correlation all elements G & A     0.663

Off-Diagonal
   Using 70386 elements from A22 >= 0.02000

   Estimating Regression Coefficients G = b0 11' + b1 A + e
   Regression coefficients b0 b1 =     -0.001     0.998

   Correlation Off-Diagonal elements G & A     0.679

Creating A22-inverse
   Wall time: 08-05-2011  16h 58m 10s 866
Inverse using ginv2
elapsed time   3.54446100000000
   Wall time: 08-05-2011  16h 58m 17s 691
```

Statistics of A_{22}^{-1}

```
Statistic of Inv. Rel. Matrix A22
                          N      Mean       Min       Max       Var
    Diagonal           1500     1.607     1.056     9.221     0.575
    Off-diagonal    2248500    -0.001    -1.067     0.533     0.001


Creating G-inverse
   Wall time: 08-05-2011  16h 58m 17s 987
Inverse using ginv2
elapsed time   4.24635400000000
   Wall time: 08-05-2011  16h 58m 26s 044
```

Statistics of G^{-1}
2 x diag(G^{-1} - A_{22}^{-1}) is approx. measure of extra genomic info in terms of effective daughters

```
Statistic of Inv. Genomic Matrix
                          N      Mean       Min       Max       Var
    Diagonal           1500     8.007     3.597    64.893    21.055
    Off-diagonal    2248500    -0.005   -12.697     6.632     0.056


Creating GimA22i in file: "GimA22i"
Calculating GmA22/GimA22i Matrix Densem storage
Calculating GmA22/GimA22i Matrix...elapsed time  0.1269817
```

```
 Setup Genomic Done.
 wGimA22i   1.00000000000000
hash matrix increased from 100000 to 150000 % filled:     0.9000
hash matrix increased from 150000 to 225000 % filled:     0.9000
hash matrix increased from 225000 to 337500 % filled:     0.9000
hash matrix increased from 337500 to 506250 % filled:     0.9000
hash matrix increased from 506250 to 759375 % filled:     0.9000
hash matrix increased from 759375 to 1139062 % filled:     0.9000
hash matrix increased from 1139062 to 1708593 % filled:     0.9000
 finished peds in    30.68333      s,       1193064  nonzeroes
 round             1    convergence=  3.234776127905992E-004
 round             2    convergence=  1.615955145159698E-005
 round             3    convergence=  9.675137058360991E-006
 round             4    convergence=  6.533482675941447E-006
 round             5    convergence=  2.711751165983321E-006
.........
.........
 round            64    convergence=  2.721030958617683E-012
 round            65    convergence=  1.931029578758311E-012
 round            66    convergence=  1.610472992188148E-012
 round            67    convergence=  1.259204136643006E-012
 round            68    convergence=  9.025592862452768E-013
          68  iterations,   convergence criterion=  9.025592862452768E-013
 solutions stored in file: "solutions"
```

Solution file

```
solutions

trait/effect level  solution
   1   1         1      4.97591211
   1   2         1      0.10194865
   1   2         2      0.33749439
   1   2         3      0.04475742
   1   2         4     -0.31055520
   1   2         5      0.22368631
   1   2         6     -0.09454804
   1   2         7     -0.03186435
   1   2         8      0.18033163
```

Variance component estimation by AIREMLF90

```
 name of parameter file?renf90.par

 * SNP file: marker.geno.clean
 * SNP Xref file: marker.geno.clean_XrefID
 * Frequency to Center Z=M-p to create G=ZZ'/k (default whichfreq = 2):
           2
     AI-REMLF90 ver. 1.96

 Parameter file:             renf90.par
 Data file:                  renf90.dat
 Number of Traits             1
 Number of Effects            2
 Position of Observations      1
 Position of Weight (1)        0
 Value of Missing Trait/Observation           0

.........
.........

Statistic of Inv. Genomic Matrix
                            N      Mean       Min       Max       Var
     Diagonal            1500     8.007     3.597    64.893    21.055
     Off-diagonal     2248500    -0.005   -12.697     6.632     0.056


 Creating GimA22i in file: "GimA22i"
```

```
 Calculating GmA22/GimA22i Matrix Densem storage
 Calculating GmA22/GimA22i Matrix...elapsed time  0.1089821
 Setup Genomic Done.
 wGimA22i   1.00000000000000
hash matrix increased from 85428 to 128142 % filled:     0.9000
hash matrix increased from 128142 to 192213 % filled:     0.9000
hash matrix increased from 192213 to 288319 % filled:     0.9000
hash matrix increased from 288319 to 432478 % filled:     0.9000
hash matrix increased from 432478 to 648717 % filled:     0.9000
hash matrix increased from 648717 to 973075 % filled:     0.9000
hash matrix increased from 973075 to 1459612 % filled:     0.9000
hash matrix increased from 85428 to 128142 % filled:     0.9000
hash matrix increased from 128142 to 192213 % filled:     0.9000
hash matrix increased from 192213 to 288319 % filled:     0.9000
hash matrix increased from 288319 to 432478 % filled:     0.9000
hash matrix increased from 432478 to 648717 % filled:     0.9000
hash matrix increased from 648717 to 973075 % filled:     0.9000
hash matrix increased from 973075 to 1459612 % filled:     0.9000
 finished peds in    32.01313      s,       1193064  nonzeroes
 rank=       15801
              **************
              **** FSPAK ***
              **************
              MPE / IM / MAE
                  Jun 1994

             SPARSE STATISTICS
     DIMENSION OF MATRIX     =                   15801
     RANK                    =                   15801
     STORAGE AVAILABLE       =                 7061497
     MAXIMUM NEEDED          =                 7061497
     NZE IN UPPER TRIANGULAR =                 1208865
     NZE IN FACTOR           =                 1521840
     NO. OF CALLS NUM FACT   =                       1
     NO. OF CALLS SOLVE      =                       1
     NO. OF CALLS SPARS SOLV =                       0
     NO. OF CALLS DET / LDET =                       1
     NO. OF CALLS SPARS INV  =                       1
     TOTAL CPU TIME IN FSPAK =                9.465561
     TIME FOR FINDING ORDER  =                2.568611
     TIME FOR SYMBOLIC FAC   =                0.676899
     TIME FOR NUMERICAL FAC  =                2.017693
     TIME FOR SOLVE          =                0.008995
     TIME FOR SPARSE SOLVE   =                0.000000
     TIME FOR SPARSE INVERSE =                4.147369
-2logL =    43515.7413644011       : AIC =    43519.7413644011
 In round            1  convergence=  0.423851780381002
 delta convergence=  0.252173522062583
new R
 0.58510
new G
 0.28516
-2logL =    53013.2734486053       : AIC =    53017.2734486053
 In round            2  convergence=  0.141351613622645
 delta convergence=  0.117430758820623
new R
 0.52205
new G
 0.45696
-2logL =    52800.6601605267       : AIC =    52804.6601605267
 In round            3  convergence=  1.725330565925358E-002
 delta convergence=  4.769938966058494E-002
new R
 0.49575
new G
 0.52606
-2logL =    52785.2479463395       : AIC =    52789.2479463395
 In round            4  convergence=  1.101891763451498E-004
 delta convergence=  3.662497104484009E-003
new R
 0.49400
```

```
 new G
  0.53164
 -2logL =    52785.1635385807       : AIC =    52789.1635385807
  In round            5  convergence=  2.804695847240073E-009
  delta convergence=  1.777604045032979E-005
 new R
  0.49400
 new G
  0.53167
```

Estimates of variance components

```
Final Estimates
 Genetic variance(s) for effect  2
  0.53167
 Residual variance(s)
  0.49400
 inverse of AI matrix (Sampling Variance)
  0.40448E-03 -0.17367E-03
 -0.17367E-03  0.14702E-03
 Correlations from inverse of AI matrix
   1.0000     -0.71219
 -0.71219       1.0000
 SE for R
  0.12125E-01
 SE for G
  0.20112E-01
 solutions stored in file: "solutions"
```

Appendix I (complete genomic analysis)

Data files are available at http://nce.ads.uga.edu/wiki/doku.php?id=course_materials_-_from_uga_2014

Using **RENUMF90, PREGSF90, BLUPF90** (**BLUP**), **BLUPF90** (**ssGBLUP**), **PREDICTF90, POSTGSF90** (**ssGWAS**)

Simulated data

Single trait with heritability of 0.30 and phenotypic variance = 1.0
Five generations
Total of 994 parents from generations 1 to 4 were genotyped
Three hundred progeny from 5th generation had genotypes and pedigree, but phenotypes were removed for traditional and genomic evaluations

Data Structure:

#Animal Generation Sex Mu QTL Residual Phenotype (Phenotype = Mu + QTL + Residual)

```
1 0 1 1 -0.826104  1.586661 1.76056
2 0 1 1 -1.093034 -0.451821 -0.544855
3 0 1 1 -0.135824  0.984936 1.84911
4 0 1 1  0.044242 -0.802145 0.242097
5 0 1 1  0.342068  0.028434 1.3705
               .
               .
6095 5 1 1  1.801324 -0.494822 2.3065
6096 5 2 1  0.772964  0.791936 2.5649
6097 5 2 1  0.748241  0.285815 2.03406
6098 5 1 1  1.042522 -1.606656 0.435866
6099 5 1 1  0.891319  0.179843 2.07116
6100 5 1 1  0.745873  0.034715 1.78059
```

Pedigree: 6100 animals

#Animal Sire Dam

```
1 0 0
2 0 0
3 0 0
4 0 0
5 0 0
   .
   .
6095 4576 4403
6096 4576 4065
6097 4576 2263
6098 4576 4150
6099 4576 3690
6100 4576 4311
```

Genotypes: 1294 animals genotyped for 1000 SNP across 5 chromosomes

Animal $SNP_1SNP_2SNP_3SNP_4SNP_5…SNP_{1000}$

```
6100    22212…1
```

Map:

#SNP order chromosome position

```
1 1 0.00000
2 1 0.16722
3 1 0.33444
4 1 0.50166
5 1 0.66888
          .
          .
1000 5 49.99878
```

Parameter file for RENUMF90

DATAFILE
newdata.txt
TRAITS
7
FIELDS_PASSED TO OUTPUT
2
WEIGHT(S)

RESIDUAL_VARIANCE
0.70
EFFECT
4 cross alpha #mu
EFFECT
1 cross alpha #animal
RANDOM
animal
FILE
ped.txt
FILE_POS
1 2 3 0 0
SNP_FILE
snp.txt
PED_DEPTH
0
(CO)VARIANCES
0.30
OPTION chrinfo map.txt

Log file for RENUMF90

```
RENUMF90 version 1.104
 name of parameter file? renum.par
 datafile:newdata.txt
 traits:           7
 fields passed:           2
 R
  0.7000

 Processing effect  1 of type cross
 item_kind=alpha

 Processing effect  2 of type cross
```

```
item_kind=alpha
pedigree file name  "ped.txt"
positions of animal, sire, dam, alternate dam and yob    1    2    3    0    0
SNP file name  "snp.txt"
all pedigrees to be included
Reading (CO)VARIANCES:           1 x           1

Maximum size of character fields: 20

Maximum size of record (max_string_readline): 800

Maximum number of fields for input file (max_field_readline): 100

hash tables for effects set up
read         6100  records
table with            1  elements sorted
added count
Effect group            1  of column            1  with            1  levels
table expanded from        10000  to        10000  records
added count
Effect group            2  of column            1  with         6100  levels
wrote statistics in file "renf90.tables"

Basic statistics for input data  (missing value code is 0)
Pos  Min         Max         Mean        SD                N
  7   -2.8883      5.0863      1.0042     0.99034        6100

random effect with SNPs  2
type: animal
file: snp.txt
read SNPs        1294  records
Effect group            2  of column            1  with         6100  levels

random effect   2
type:animal
opened output pedigree file "renadd02.ped"
read         6100  pedigree records

Pedigree checks

Number of animals with records:        6100
Number of animals with genotypes:        1294
Number of animals with records or genotypes:        6100
Number of animals with genotypes and no records           0
Number of parents without records or genotypes:           0
Total number of animals:        6100

Wrote cross reference IDs for SNP file "snp.txt_XrefID"

Wrote parameter file "renf90.par"
Wrote renumbered data "renf90.dat"
```

Parameter file for PREGSF90 without quality control

```
DATAFILE
 renf90.dat
NUMBER_OF_TRAITS
      1
NUMBER_OF_EFFECTS
      2
```

```
OBSERVATION(S)
  1
WEIGHT(S)

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT[EFFECT NESTED]
 2      1 cross
 3    6100 cross
RANDOM_RESIDUAL VALUES
 0.70000
 RANDOM_GROUP
   2
 RANDOM_TYPE
 add_animal
 FILE
renadd02.ped
(CO)VARIANCES
 0.30000
OPTION SNP_file snp.txt
OPTION chrinfo map.txt
OPTION no_quality_control
```

Log file for PREGSF90 without quality control

```
name of parameter file?
renf90.par

     preGS 1.10

 Parameter file:             renf90.par
 Data file:                  renf90.dat
 Number of Traits             1
 Number of Effects            2
 Position of Observations      1
 Position of Weight (1)        0
 Value of Missing Trait/Observation           0

EFFECTS
 #  type                position (2)        levels   [positions for nested]
 1  cross-classified     2                                                  1
 2  cross-classified     3                                               6100

 Residual (co)variance Matrix
 0.70000

 Random Effect(s)    2
 Type of Random Effect:      additive animal
 Pedigree File:              renadd02.ped
 trait   effect    (CO)VARIANCES
  1       2    0.3000

 REMARKS
  (1) Weight position 0 means no weights utilized
  (2) Effect positions of 0 for some effects and traits means that such
      effects are missing for specified traits

Options read from parameter file:

 * SNP file: snp.txt
```

```
 * SNP Xref file: snp.txt_XrefID
 * Map file: map.txt
 * No Quality Control Checks !!!!! (default .false.):  T

 *-----------------------------------------------------------------*
 *                 Genomic Library: Version 1.164                  *
 *                                                                 *
 *                   Optimized OpenMP Version                      *
 *                                                                 *
 *  Modified relationship matrix (H) created for effect:   2     *
 *-----------------------------------------------------------------*

 Read 6100 animals from pedigree file: "renadd02.ped"
 Number of Genotyped Animals: 1294

Creating A22
    Extracting subset of: 2312 pedigrees from: 6100 elapsed time:     0.0150
    Calculating A22 Matrix by Colleau OpenMP...elapsed time: .0190
    Numbers of threads=8 16

 Reading SNP file
    Column position in file for the first marker: 8
    Format to read SNP file: (7x,400000i1)
    Number of SNPs: 1000
    Number of Genotyped animals: 1294
    Reading SNP file elapsed time: .06

 Statistics of alleles frequencies in the current population
    N:           1000
    Mean:       0.504
    Min:        0.043
    Max:        0.929
    Var:        0.032

 Reading MAP file: "map.txt" - 1000 SNPs out of 1000

    Min and max # of chromosome: 1 5

    Min and max # of SNP: 1 1000

 Genotypes missings (%):  0.000

 Calculating G Matrix
    Dgemm MKL #threads=     8   16 Elapsed omp_get_time:     0.7359

 Scale by Sum(2pq). Average:   435.221580281360

 Blend G as alpha*G + beta*A22: (alpha,beta)     0.950     0.050

 Frequency - Diagonal of G
    N:        1294
    Mean:        0.999
    Min:         0.895
    Max:         1.468
    Range:       0.029
    Class:     20

#Class       Class   Count
       1  0.8949          27
       2  0.9236         109
       3  0.9523         300
       4  0.9810         380
```

```
      5   1.010         287
      6   1.038         137
      7   1.067          33
      8   1.096          14
      9   1.124           3
     10   1.153           1
     11   1.182           0
     12   1.210           2
     13   1.239           0
     14   1.268           0
     15   1.296           0
     16   1.325           0
     17   1.354           0
     18   1.382           0
     19   1.411           0
     20   1.440           1
     21   1.468           0


 Check for diagonal of genomic relationship matrix

 Check for diagonal of genomic relationship matrix, genotypes not removed: 0

 ------------------------------
  Final Pedrigree-Based Matrix
 ------------------------------

 Statistic of Rel. Matrix A22
                          N      Mean       Min       Max       Var
    Diagonal           1294     1.001     1.000     1.250     0.000
    Off-diagonal    1673142     0.005     0.000     0.750     0.001

 ----------------------
  Final Genomic Matrix
 ----------------------

 Statistic of Genomic Matrix
                          N      Mean       Min       Max       Var
    Diagonal           1294     1.001     0.898     1.469     0.002
    Off-diagonal    1673142     0.005    -0.158     0.791     0.002

 Correlation of Genomic Inbreeding and Pedigree Inbreeding
    Correlation:     0.2177

All elements - Diagonal / Off-Diagonal
    Estimating Regression Coefficients G = b0 11' + b1 A + e
    Regression coefficients b0 b1 =      0.000     0.991

    Correlation all elements G & A     0.717

 Off-Diagonal
    Using 83426 elements from A22 >= .02000

    Estimating Regression Coefficients G = b0 11' + b1 A + e
    Regression coefficients b0 b1 =     -0.003     0.999

    Correlation Off-Diagonal elements G & A     0.777

 Creating A22-inverse
    Inverse LAPACK MKL dpotrf/i  #threads=    8   16 Elapsed omp_get_time:     0.1071

 ----------------------
```

```
 Final A22 Inv Matrix
 ----------------------

 Statistic of Inv. Rel. Matrix A22
                           N       Mean        Min        Max        Var
     Diagonal           1294      1.851      1.067      5.812      0.431
     Off-diagonal    1673142     -0.001     -1.200      0.600      0.001


 Creating G-inverse
    Inverse LAPACK MKL dpotrf/i  #threads=    8   16 Elapsed omp_get_time:     0.1050

 --------------------------
  Final Genomic Inv Matrix
 --------------------------

 Statistic of Inv. Genomic Matrix
                           N       Mean        Min        Max        Var
     Diagonal           1294     13.457      5.827     45.588     27.985
     Off-diagonal    1673142     -0.010    -13.500      6.896      0.226

 Check for diagonal of Inverse Genomic - Inverse of pedigree relationship matrix

 Saving GimA22i in file: "GimA22i"

------------------------------
  Final G Inv - A22 Inv Matrix
 ------------------------------

 Statistic of Inv. Genomic- A22 Matrix
                           N       Mean        Min        Max        Var
     Diagonal           1294     11.606      4.746     40.310     21.707
     Off-diagonal    1673142     -0.009    -12.500      6.396      0.211

*-------------------------*
* Setup Genomic Done !!! *
*-------------------------*
```

Parameter file for PREGSF90 with quality control

```
DATAFILE
 renf90.dat
NUMBER_OF_TRAITS
      1
NUMBER_OF_EFFECTS
      2
OBSERVATION(S)
  1
WEIGHT(S)

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT[EFFECT NESTED]
 2       1 cross
 3     6100 cross
RANDOM_RESIDUAL VALUES
 0.70000
 RANDOM_GROUP
   2
 RANDOM_TYPE
```

```
 add_animal
 FILE
renadd02.ped
(CO)VARIANCES
 0.30000
OPTION SNP_file snp.txt
OPTION chrinfo map.txt
```

Log file for PREGSF90 with quality control

```
name of parameter file?
renf90.par

     preGS 1.10

 Parameter file:             renf90.par
 Data file:                  renf90.dat
 Number of Traits             1
 Number of Effects            2
 Position of Observations      1
 Position of Weight (1)        0
 Value of Missing Trait/Observation           0

EFFECTS
 #  type                position (2)        levels    [positions for nested]
 1  cross-classified     2                                                   1
 2  cross-classified     3                                                6100

 Residual (co)variance Matrix
 0.70000

 Random Effect(s)    2
 Type of Random Effect:      additive animal
 Pedigree File:              renadd02.ped
 trait   effect    (CO)VARIANCES
  1       2    0.3000

 REMARKS
  (1) Weight position 0 means no weights utilized
  (2) Effect positions of 0 for some effects and traits means that such
      effects are missing for specified traits

Options read from parameter file:

 * SNP file: snp.txt
 * SNP Xref file: snp.txt_XrefID
 * Map file: map.txt

 *-----------------------------------------------------------------*
 *                 Genomic Library: Version 1.164                  *
 *                                                                 *
 *                   Optimized OpenMP Version                      *
 *                                                                 *
 *  Modified relationship matrix (H) created for effect:   2       *
 *-----------------------------------------------------------------*

 Read 6100 animals from pedigree file: "renadd02.ped"
 Number of Genotyped Animals: 1294

 Creating A22
```

```
   Extracting subset of: 2312 pedigrees from: 6100 elapsed time:     0.0160
   Calculating A22 Matrix by Colleau OpenMP...elapsed time: .0189
   Numbers of threads=8 16

Reading SNP file
   Column position in file for the first marker: 8
   Format to read SNP file: (7x,400000i1)
   Number of SNPs: 1000
   Number of Genotyped animals: 1294
   Reading SNP file elapsed time: .06

Statistics of alleles frequencies in the current population
   N:           1000
   Mean:       0.504
   Min:        0.043
   Max:        0.929
   Var:        0.032

Reading MAP file: "map.txt" - 1000 SNPs out of 1000

   Min and max # of chromosome: 1 5

   Min and max # of SNP: 1 1000

Quality Control - SNPs with Call Rate < callrate ( 0.90) will removed: 0

Quality Control - SNPs with MAF < minfreq ( 0.05) will removed: 1

Quality Control - Monomorphic SNPs will be removed: 0

Quality Control - Removed Animals with Call rate < callrate ( 0.90): 0

Quality Control - Check Parent-Progeny Mendelian conflicts

   Total animals: 6100 - Genotyped animals: 1294 - Effective: 1294

   Number of pairs Individual - Sire: 450
   Number of pairs Individual - Dam: 440
   Number of trios Individual - Sire - Dam: 206

   No sex Chromosome information is available
   Parent-progeny conflicts or HWE could eliminate SNPs in sex Chr
   Provide map information and sex Chr to checks using autosomes

Checking SNPs for Mendelian conflicts

   Total number of effective SNP: 999
   Total number of parent-progeny evaluations: 890
   Number of SNPs with Mendelian conflicts: 0

Checking Animals for Mendelian conflicts

   Total number of effective SNP for checks on Animals: 999

   Number of Parent-Progeny Mendelian Conflicts: 0

Number of effective SNPs (after QC): 999

Number of effective Indiviuals (after QC): 1294

Statistics of alleles frequencies in the current population after
Quality Control (MAF, monomorphic, call rate, HWE, Mendelian conflicts)
```

```
   N:            999
   Mean:       0.504
   Min:        0.051
   Max:        0.929
   Var:        0.032

Genotypes missings (%):  0.100

Genotypes missings after cleannig (%):  0.000

Calculating G Matrix
   Dgemm MKL #threads=     8   16 Elapsed omp_get_time:     0.9840

Scale by Sum(2pq). Average:   435.140185710293

Blend G as alpha*G + beta*A22: (alpha,beta)     0.950     0.050

Frequency - Diagonal of G
   N:        1294
   Mean:        0.999
   Min:         0.895
   Max:         1.469
   Range:       0.029
   Class:     20

 #Class       Class   Count
      1  0.8951          27
      2  0.9238         109
      3  0.9524         304
      4  0.9811         379
      5   1.010         285
      6   1.038         137
      7   1.067          32
      8   1.096          14
      9   1.125           3
     10   1.153           1
     11   1.182           0
     12   1.211           2
     13   1.239           0
     14   1.268           0
     15   1.297           0
     16   1.325           0
     17   1.354           0
     18   1.383           0
     19   1.411           0
     20   1.440           1
     21   1.469           0

Check for diagonal of genomic relationship matrix

Check for diagonal of genomic relationship matrix, genotypes not removed: 0

------------------------------
 Final Pedrigree-Based Matrix
------------------------------

Statistic of Rel. Matrix A22
                          N      Mean       Min       Max       Var
    Diagonal           1294     1.001     1.000     1.250     0.000
    Off-diagonal    1673142     0.005     0.000     0.750     0.001

----------------------
```

```
 Final Genomic Matrix
 ----------------------

 Statistic of Genomic Matrix
                           N      Mean       Min       Max       Var
     Diagonal           1294     1.001     0.898     1.470     0.002
     Off-diagonal    1673142     0.005    -0.158     0.791     0.002

 Correlation of Genomic Inbreeding and Pedigree Inbreeding
     Correlation:     0.2180

 All elements - Diagonal / Off-Diagonal
    Estimating Regression Coefficients G = b0 11' + b1 A + e
    Regression coefficients b0 b1 =      0.000     0.991

    Correlation all elements G & A     0.717

 Off-Diagonal
    Using 83426 elements from A22 >= .02000

    Estimating Regression Coefficients G = b0 11' + b1 A + e
    Regression coefficients b0 b1 =     -0.003     0.999

    Correlation Off-Diagonal elements G & A     0.777

 Creating A22-inverse
    Inverse LAPACK MKL dpotrf/i  #threads=    8   16 Elapsed omp_get_time:     0.1068
----------------------
 Final A22 Inv Matrix
 ----------------------

 Statistic of Inv. Rel. Matrix A22
                           N      Mean       Min       Max       Var
     Diagonal           1294     1.851     1.067     5.812     0.431
     Off-diagonal    1673142    -0.001    -1.200     0.600     0.001

 Creating G-inverse
    Inverse LAPACK MKL dpotrf/i  #threads=    8   16 Elapsed omp_get_time:     0.1047

 --------------------------
 Final Genomic Inv Matrix
 --------------------------

 Statistic of Inv. Genomic Matrix
                           N      Mean       Min       Max       Var
     Diagonal           1294    13.466     5.863    45.587    28.023
     Off-diagonal    1673142    -0.010   -13.521     6.897     0.227

 Check for diagonal of Inverse Genomic - Inverse of pedigree relationship matrix

 Saving GimA22i in file: "GimA22i"

 ------------------------------
 Final G Inv - A22 Inv Matrix
 ------------------------------

 Statistic of Inv. Genomic- A22 Matrix
                           N      Mean       Min       Max       Var
     Diagonal           1294    11.615     4.782    40.309    21.740
     Off-diagonal    1673142    -0.009   -12.521     6.397     0.211

*------------------------*
```

```
* Setup Genomic Done !!! *
*------------------------*
```

Parameter file for PREGSF90 with quality control, removing SNP from chromosome 5 and saving the clean SNP file

```
 DATAFILE
 renf90.dat
NUMBER_OF_TRAITS
      1
NUMBER_OF_EFFECTS
      2
OBSERVATION(S)
  1
WEIGHT(S)

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT[EFFECT NESTED]
 2      1 cross
 3    6100 cross
RANDOM_RESIDUAL VALUES
 0.70000
 RANDOM_GROUP
   2
 RANDOM_TYPE
 add_animal
 FILE
renadd02.ped
(CO)VARIANCES
 0.30000
OPTION SNP_file snp.txt
OPTION chrinfo map.txt
OPTION excludeCHR 5
OPTION saveCleanSNPs
```

Log file for PREGSF90 with quality control, removing SNP from chromosome 5 and saving the clean SNP file

```
name of parameter file?
renf90.par

     preGS 1.10

 Parameter file:             renf90.par
 Data file:                  renf90.dat
 Number of Traits             1
 Number of Effects            2
 Position of Observations      1
 Position of Weight (1)        0
 Value of Missing Trait/Observation           0

EFFECTS
 #  type                position (2)        levels    [positions for nested]
 1  cross-classified     2                                                  1
 2  cross-classified     3                                               6100
```

```
 Residual (co)variance Matrix
 0.70000

 Random Effect(s)    2
 Type of Random Effect:      additive animal
 Pedigree File:              renadd02.ped
 trait   effect    (CO)VARIANCES
  1       2    0.3000

 REMARKS
  (1) Weight position 0 means no weights utilized
  (2) Effect positions of 0 for some effects and traits means that such
      effects are missing for specified traits

Options read from parameter file:

 * SNP file: snp.txt
 * SNP Xref file: snp.txt_XrefID
 * Map file: map.txt
 * Save Clean SNP data to (SNP_file)_clean file (default .false.)
 * Exclude Chromosomes (default .false.): 5

 *-----------------------------------------------------------------*
 *                 Genomic Library: Version 1.164                  *
 *                                                                 *
 *                   Optimized OpenMP Version                      *
 *                                                                 *
 *  Modified relationship matrix (H) created for effect:   2     *
 *-----------------------------------------------------------------*

 Read 6100 animals from pedigree file: "renadd02.ped"
 Number of Genotyped Animals: 1294

 Creating A22
    Extracting subset of: 2312 pedigrees from: 6100 elapsed time:     0.0150
    Calculating A22 Matrix by Colleau OpenMP...elapsed time: .0190
    Numbers of threads=8 16

 Reading SNP file
    Column position in file for the first marker: 8
    Format to read SNP file: (7x,400000i1)
    Number of SNPs: 1000
    Number of Genotyped animals: 1294
    Reading SNP file elapsed time: .06

 Statistics of alleles frequencies in the current population
    N:           1000
    Mean:       0.504
    Min:        0.043
    Max:        0.929
    Var:        0.032

 Reading MAP file: "map.txt" - 1000 SNPs out of 1000

    Min and max # of chromosome: 1 5

    Min and max # of SNP: 1 1000

 Excluded 199 SNPs from 1 chromosomes: 5

 Quality Control - SNPs with Call Rate < callrate ( 0.90) will removed: 199
```

```
Quality Control - SNPs with MAF < minfreq ( 0.05) will removed: 1

Quality Control - Monomorphic SNPs will be removed: 0

Quality Control - Removed Animals with Call rate < callrate ( 0.90): 0

Quality Control - Check Parent-Progeny Mendelian conflicts

   Total animals: 6100 - Genotyped animals: 1294 - Effective: 1294

   Number of pairs Individual - Sire: 450
   Number of pairs Individual - Dam: 440
   Number of trios Individual - Sire - Dam: 206

   No sex Chromosome information is available
   Parent-progeny conflicts or HWE could eliminate SNPs in sex Chr
   Provide map information and sex Chr to checks using autosomes

Checking SNPs for Mendelian conflicts

   Total number of effective SNP: 801
   Total number of parent-progeny evaluations: 890
   Number of SNPs with Mendelian conflicts: 0

Checking Animals for Mendelian conflicts

   Total number of effective SNP for checks on Animals: 801

   Number of Parent-Progeny Mendelian Conflicts: 0

Number of effective SNPs (after QC): 801

Number of effective Indiviuals (after QC): 1294

Statistics of alleles frequencies in the current population after
Quality Control (MAF, monomorphic, call rate, HWE, Mendelian conflicts)
   N:            801
   Mean:       0.503
   Min:        0.051
   Max:        0.928
   Var:        0.032

List of SNPs removed in: "snp.txt_SNPs_removed"

Clean genotype file was created: "snp.txt_clean"

Cross reference ID file was created: "snp.txt_clean_XrefID"

Genotypes missings (%): 19.900

Genotypes missings after cleannig (%):  0.000

Calculating G Matrix
   Dgemm MKL #threads=     8   16 Elapsed omp_get_time:     0.8764

Scale by Sum(2pq). Average:   349.571560214902

Blend G as alpha*G + beta*A22: (alpha,beta)     0.950     0.050

Frequency - Diagonal of G
   N:        1294
   Mean:        1.000
```

Number of effective SNP was reduced to 801 after removing chromosome 5

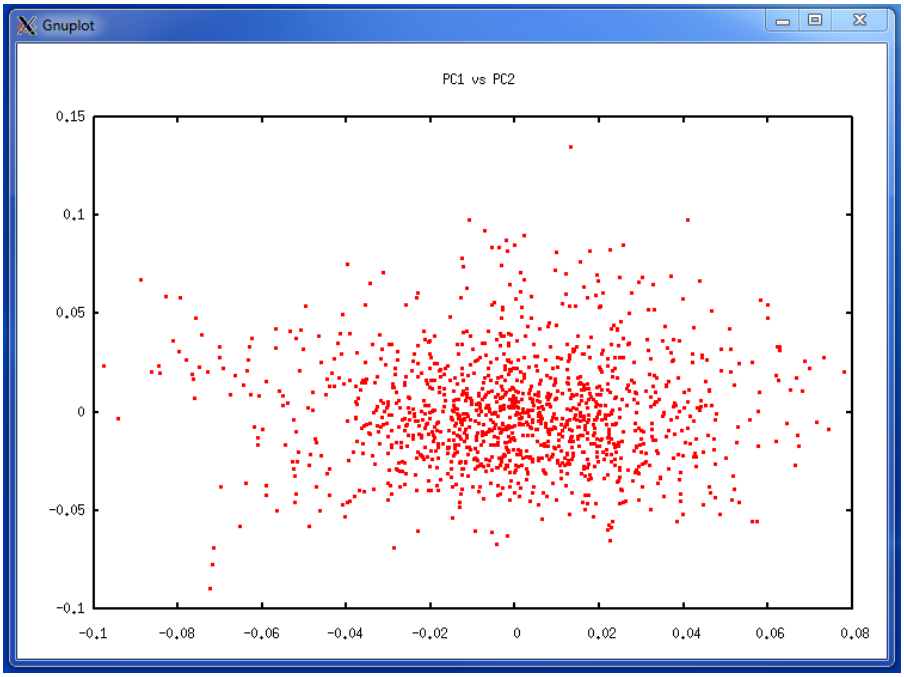
New files with clean genotypes

```
   Min:          0.874
   Max:          1.593
   Range:        0.036
   Class:     20

 #Class       Class   Count
      1  0.8741          17
      2  0.9100         107
      3  0.9460         341
      4  0.9819         419
      5   1.018         281
      6   1.054          98
      7   1.090          20
      8   1.126           4
      9   1.162           4
     10   1.198           1
     11   1.234           0
     12   1.270           1
     13   1.306           0
     14   1.342           0
     15   1.377           0
     16   1.413           0
     17   1.449           0
     18   1.485           0
     19   1.521           0
     20   1.557           1
     21   1.593           0

Check for diagonal of genomic relationship matrix

Check for diagonal of genomic relationship matrix, genotypes not removed: 0

------------------------------
 Final Pedrigree-Based Matrix
------------------------------

Statistic of Rel. Matrix A22
                          N      Mean       Min       Max       Var
    Diagonal           1294     1.001     1.000     1.250     0.000
    Off-diagonal    1673142     0.005     0.000     0.750     0.001

----------------------
 Final Genomic Matrix
----------------------

Statistic of Genomic Matrix
                          N      Mean       Min       Max       Var
    Diagonal           1294     1.001     0.876     1.593     0.002
    Off-diagonal    1673142     0.005    -0.169     0.861     0.003

Correlation of Genomic Inbreeding and Pedigree Inbreeding
    Correlation:     0.2092

All elements - Diagonal / Off-Diagonal
   Estimating Regression Coefficients G = b0 11' + b1 A + e
   Regression coefficients b0 b1 =      0.000     0.991

   Correlation all elements G & A     0.677

Off-Diagonal
   Using 83426 elements from A22 >= .02000
```

```
   Estimating Regression Coefficients G = b0 11' + b1 A + e
   Regression coefficients b0 b1 =     -0.002     0.996

   Correlation Off-Diagonal elements G & A     0.742

Creating A22-inverse
   Inverse LAPACK MKL dpotrf/i  #threads=    8   16 Elapsed omp_get_time:     0.1409

----------------------
 Final A22 Inv Matrix
----------------------

Statistic of Inv. Rel. Matrix A22
                          N      Mean       Min       Max       Var
    Diagonal           1294     1.851     1.067     5.812     0.431
    Off-diagonal    1673142    -0.001    -1.200     0.600     0.001

Creating G-inverse
   Inverse LAPACK MKL dpotrf/i  #threads=    8   16 Elapsed omp_get_time:     0.1370

--------------------------
 Final Genomic Inv Matrix
--------------------------

Statistic of Inv. Genomic Matrix
                          N      Mean       Min       Max       Var
    Diagonal           1294    17.075     7.840    56.092    43.645
    Off-diagonal    1673142    -0.013   -16.499     8.893     0.309

Check for diagonal of Inverse Genomic - Inverse of pedigree relationship matrix

Saving GimA22i in file: "GimA22i"

------------------------------
 Final G Inv - A22 Inv Matrix
------------------------------

Statistic of Inv. Genomic- A22 Matrix
                          N      Mean       Min       Max       Var
    Diagonal           1294    15.223     6.759    51.043    35.648
    Off-diagonal    1673142    -0.012   -15.499     8.393     0.289

*-------------------------*
* Setup Genomic Done !!! *
*-------------------------*
```

Parameter file for PREGSF90 with quality control and PCA analysis

Include extra option: **OPTION plotpca**



Parameter file for BLUPF90 without genomic information

```
DATAFILE
 renf90_5.dat
NUMBER_OF_TRAITS
        1
NUMBER_OF_EFFECTS
        2
OBSERVATION(S)
   1
WEIGHT(S)

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT[EFFECT NESTED]
  2       1 cross
  3    6100 cross
RANDOM_RESIDUAL VALUES
 0.70000
 RANDOM_GROUP
    2
 RANDOM_TYPE
 add_animal
 FILE
renadd02.ped
(CO)VARIANCES
 0.30000
OPTION conv_crit 1e-15
```

renf90_5.dat has phenotypes for all animals, but generation 5

Linux code to remove phenotypes for those animals:
awk '{ if ($4==5) print 0,$2,$3,$4; else print $1,$2,$3,$4}' renf90.dat > renf90_5.dat

Default convergence criteria = 1e-12

Log file for BLUPF90 without genomic information

```
name of parameter file?
```

```
renf90.par
 * convergence criterion (default=1e-12):  1.0000000E-15

     BLUPF90 1.48

 Parameter file:              renf90.par
 Data file:                   renf90_5.dat
 Number of Traits              1
 Number of Effects             2
 Position of Observations      1
 Position of Weight (1)        0
 Value of Missing Trait/Observation           0

EFFECTS
 #  type                 position (2)        levels    [positions for nested]
 1  cross-classified     2                                                   1
 2  cross-classified     3                                                6100

 Residual (co)variance Matrix
 0.70000

 Random Effect(s)    2
 Type of Random Effect:       additive animal
 Pedigree File:               renadd02.ped
 trait   effect    (CO)VARIANCES
  1       2    0.3000

 REMARKS
  (1) Weight position 0 means no weights utilized
  (2) Effect positions of 0 for some effects and traits means that such
      effects are missing for specified traits

 Data record length =            3
 # equations =         6101
 G
 0.30000
 read         6100  records in   1.4997000E-02  s,                   12201
  nonzeroes
  read         6100  additive pedigrees
 finished peds in   1.9996000E-02  s,                  27178  nonzeroes
round =     1  convergence =  0.1730E-03
round =     2  convergence =  0.7971E-03
round =     3  convergence =  0.5923E-04
round =     4  convergence =  0.6219E-04
round =     5  convergence =  0.2122E-04
        .
        .
        .
round =    40  convergence =  0.1230E-13
round =    41  convergence =  0.3164E-14
round =    42  convergence =  0.2804E-14
round =    43  convergence =  0.1081E-14
round =    44  convergence =  0.5761E-15
   44 iterations,   convergence criterion= 0.5761E-15
 solutions stored in file: "solutions"
```

Solutions for BLUPF90 without genomic information

```
trait/effect level  solution
   1   1         1      1.02176505
   1   2         1     -0.24665178
```

```
   1    2          2       0.16420973
   1    2          3       0.32371581
   1    2          4       0.00318130
   1    2          5      -0.13277100
```

The solution file (solutions) has 4 columns:

1) Trait [only 1 trait in this example]
2) Effect [we have 2 effects: overall mean (effect 1) and additive genetic direct (effect 2)]
3) Level [number of the level for each effect in the model]
4) Solution

EBV accuracy

If accuracy of EBV is desired, it can be calculated based on standard errors (se) for EBV.

BLUPF90 has an option for calculating se:

OPTION sol se

Solutions for BLUPF90 with option to calculate se

The solution file now includes a 5th column with EBV standard errors

```
trait/effect level  solution
s.e.
   1    1          1
1.02176504       0.02496866
   1    2          1      -0.24665117       0.39158195
   1    2          2       0.16421026       0.40488662
   1    2          3       0.32371755       0.29405286
   1    2          4       0.00318218       0.38229658
   1    2          5      -0.13277154       0.46566701
```

Parameter file for BLUPF90 with genomic information (ssGBLUP)

```
DATAFILE
 renf90_5.dat
NUMBER_OF_TRAITS
       1
NUMBER_OF_EFFECTS
       2
OBSERVATION(S)
   1
WEIGHT(S)

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT[EFFECT NESTED]
 2       1 cross
 3    6100 cross
RANDOM_RESIDUAL VALUES
 0.70000
 RANDOM_GROUP
    2
 RANDOM_TYPE
 add_animal
 FILE
renadd02.ped
(CO)VARIANCES
 0.30000
OPTION SNP_file snp.txt
```

OPTION chrinfo map.txt
OPTION conv_crit 1e-15

Log file for BLUPF90 with genomic information (ssGBLUP)

```
name of parameter file?
renf90.par
 * convergence criterion (default=1e-12):  1.0000000E-15


Options read from parameter file:

 * SNP file: snp.txt
 * SNP Xref file:snp.txt_XrefID
 * Map file: map.txt
     BLUPF90 1.48

 Parameter file:             renf90.par
 Data file:                  renf90_5.dat
 Number of Traits             1
 Number of Effects            2
 Position of Observations      1
 Position of Weight (1)        0
 Value of Missing Trait/Observation           0

EFFECTS
 #  type                position (2)        levels   [positions for nested]
 1  cross-classified     2                                                  1
 2  cross-classified     3                                               6100

 Residual (co)variance Matrix
 0.70000

 Random Effect(s)    2
 Type of Random Effect:      additive animal
 Pedigree File:              renadd02.ped
 trait   effect    (CO)VARIANCES
  1       2    0.3000

 REMARKS
  (1) Weight position 0 means no weights utilized
  (2) Effect positions of 0 for some effects and traits means that such
      effects are missing for specified traits

 Data record length =            3
 # equations =         6101
 G
 0.30000
 read         6100  records in   0.1499770      s,                   12201
  nonzeroes
  read         6100  additive pedigrees

 *-----------------------------------------------------------*
 *                 Genomic Library: Version 1.164            *
 *                                                           *
 *                   Optimized OpenMP Version                *
 *                                                           *
 *  Modified relationship matrix (H) created for effect:   2    *
 *-----------------------------------------------------------*

 Read 6100 animals from pedigree file: "renadd02.ped"
```

```
Number of Genotyped Animals: 1294

Creating A22
   Extracting subset of: 2312 pedigrees from: 6100 elapsed time:     0.0150
   Calculating A22 Matrix by Colleau OpenMP...elapsed time: .0346
   Numbers of threads=8 16

Reading SNP file
   Column position in file for the first marker: 8
   Format to read SNP file: (7x,400000i1)
   Number of SNPs: 1000
   Number of Genotyped animals: 1294
   Reading SNP file elapsed time: .06

Statistics of alleles frequencies in the current population
   N:           1000
   Mean:       0.504
   Min:        0.043
   Max:        0.929
   Var:        0.032

Reading MAP file: "map.txt" - 1000 SNPs out of 1000

   Min and max # of chromosome: 1 5

   Min and max # of SNP: 1 1000

Quality Control - SNPs with Call Rate < callrate ( 0.90) will removed: 0

Quality Control - SNPs with MAF < minfreq ( 0.05) will removed: 1

Quality Control - Monomorphic SNPs will be removed: 0

Quality Control - Removed Animals with Call rate < callrate ( 0.90): 0

Quality Control - Check Parent-Progeny Mendelian conflicts

   Total animals: 6100 - Genotyped animals: 1294 - Effective: 1294

   Number of pairs Individual - Sire: 450
   Number of pairs Individual - Dam: 440
   Number of trios Individual - Sire - Dam: 206

   No sex Chromosome information is available
   Parent-progeny conflicts or HWE could eliminate SNPs in sex Chr
   Provide map information and sex Chr to checks using autosomes

Checking SNPs for Mendelian conflicts

   Total number of effective SNP: 999
   Total number of parent-progeny evaluations: 890
   Number of SNPs with Mendelian conflicts: 0

Checking Animals for Mendelian conflicts

   Total number of effective SNP for checks on Animals: 999

   Number of Parent-Progeny Mendelian Conflicts: 0

Number of effective SNPs (after QC): 999

Number of effective Indiviuals (after QC): 1294
```

```
Statistics of alleles frequencies in the current population after
Quality Control (MAF, monomorphic, call rate, HWE, Mendelian conflicts)
   N:            999
   Mean:       0.504
   Min:        0.051
   Max:        0.929
   Var:        0.032

Genotypes missings (%):  0.100

Genotypes missings after cleannig (%):  0.000

Calculating G Matrix
   Dgemm MKL #threads=     8   16 Elapsed omp_get_time:     1.0240

Scale by Sum(2pq). Average:   435.140185710293

Blend G as alpha*G + beta*A22: (alpha,beta)     0.950     0.050

Frequency - Diagonal of G
   N:        1294
   Mean:        0.999
   Min:         0.895
   Max:         1.469
   Range:       0.029
   Class:     20

 #Class       Class   Count
      1  0.8951          27
      2  0.9238         109
      3  0.9524         304
      4  0.9811         379
      5   1.010         285
      6   1.038         137
      7   1.067          32
      8   1.096          14
      9   1.125           3
     10   1.153           1
     11   1.182           0
     12   1.211           2
     13   1.239           0
     14   1.268           0
     15   1.297           0
     16   1.325           0
     17   1.354           0
     18   1.383           0
     19   1.411           0
     20   1.440           1
     21   1.469           0


Check for diagonal of genomic relationship matrix


Check for diagonal of genomic relationship matrix, genotypes not removed: 0

------------------------------
 Final Pedrigree-Based Matrix
------------------------------

Statistic of Rel. Matrix A22
```

```
                          N      Mean       Min       Max       Var
    Diagonal           1294     1.001     1.000     1.250     0.000
    Off-diagonal    1673142     0.005     0.000     0.750     0.001


----------------------
 Final Genomic Matrix
----------------------

 Statistic of Genomic Matrix
                          N      Mean       Min       Max       Var
    Diagonal           1294     1.001     0.898     1.470     0.002
    Off-diagonal    1673142     0.005    -0.158     0.791     0.002


 Correlation of Genomic Inbreeding and Pedigree Inbreeding
    Correlation:     0.2180

All elements - Diagonal / Off-Diagonal
   Estimating Regression Coefficients G = b0 11' + b1 A + e
   Regression coefficients b0 b1 =      0.000     0.991

   Correlation all elements G & A     0.717

 Off-Diagonal
   Using 83426 elements from A22 >= .02000

   Estimating Regression Coefficients G = b0 11' + b1 A + e
   Regression coefficients b0 b1 =     -0.003     0.999

   Correlation Off-Diagonal elements G & A     0.777


 Creating A22-inverse
   Inverse LAPACK MKL dpotrf/i  #threads=    8   16 Elapsed omp_get_time:     0.1059

----------------------
 Final A22 Inv Matrix
----------------------

 Statistic of Inv. Rel. Matrix A22
                          N      Mean       Min       Max       Var
    Diagonal           1294     1.851     1.067     5.812     0.431
    Off-diagonal    1673142    -0.001    -1.200     0.600     0.001


 Creating G-inverse
   Inverse LAPACK MKL dpotrf/i  #threads=    8   16 Elapsed omp_get_time:     0.1093

--------------------------
 Final Genomic Inv Matrix
--------------------------

 Statistic of Inv. Genomic Matrix
                          N      Mean       Min       Max       Var
    Diagonal           1294    13.466     5.863    45.587    28.023
    Off-diagonal    1673142    -0.010   -13.521     6.897     0.227


 Check for diagonal of Inverse Genomic - Inverse of pedigree relationship matrix
------------------------------
  Final G Inv - A22 Inv Matrix
```

```
 ------------------------------

 Statistic of Inv. Genomic- A22 Matrix
                            N      Mean       Min       Max       Var
     Diagonal            1294    11.615     4.782    40.309    21.740
     Off-diagonal     1673142    -0.009   -12.521     6.397     0.211


*------------------------*
* Setup Genomic Done !!! *
*------------------------*

hash matrix increased from          131072 to          262144 % filled:     0.8000
hash matrix increased from          262144 to          524288 % filled:     0.8000
hash matrix increased from          524288 to         1048576 % filled:     0.8000
hash matrix increased from         1048576 to         2097152 % filled:     0.8000
 finished peds in    25.61810      s,                  861721  nonzeroes
round =     1  convergence =  0.6397E-03
round =     2  convergence =  0.4280E-03
round =     3  convergence =  0.3112E-03
round =     4  convergence =  0.9994E-04
round =     5  convergence =  0.8129E-04
        .
        .
        .
round =    90  convergence =  0.3590E-14
round =    91  convergence =  0.2549E-14
round =    92  convergence =  0.2022E-14
round =    93  convergence =  0.1453E-14
round =    94  convergence =  0.9599E-15
   94 iterations,   convergence criterion= 0.9599E-15
 solutions stored in file: "solutions"
```

Solutions for BLUPF90 with genomic information (ssGBLUP)

The solution file has the same format as in blupf90 without genomic information. The option for calculating se for EBV can also be used here.

Parameter file for PREDICTF90

Predictivity can be measured as correlation between corrected phenotypes and (G)EBV. In this example we show predictivity for all young animals and for all young genotyped animals.

1) Prectivity for traditional BLUP

As this program needs solution file, it can be run in the same folder as BLUP

Parameter file:

```
DATAFILE
 pred.dat
NUMBER_OF_TRAITS
      1
NUMBER_OF_EFFECTS
      2
OBSERVATION(S)
  1
WEIGHT(S)
```

pred.dat is the data file only for animals in 5th generation

Linux code to create pred.dat:

```
awk '$4==5' renf90.dat > pred.dat
```

For creating pred.dat only for genotyped animals in 5th generation:

```
awk 'BEGIN { for (i==0;i<1000;i++) print i+1}' > ind
paste -d " " pred.dat ind | sort +2 -3 > dat1.temp
awk '{print $1}' snp.txt_XrefID | sort +0 -1 > dat2.temp
join -1 +1 -2 +3 dat2.temp dat1.temp | sort -n +4 -5 | awk '{print $2,$3,$1,$4}' >
gen.dat
```

```
EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT[EFFECT NESTED]
 2      1 cross
 3    6100 cross
RANDOM_RESIDUAL VALUES
 0.70000
 RANDOM_GROUP
   2
 RANDOM_TYPE
 add_animal
 FILE
renadd02.ped
(CO)VARIANCES
 0.30000
OPTION include_effects 2
```

Log file for predicting all animals in 5th generation:

```
name of parameter file?
pred.par

 *** include effets to predict Yhat n, effects            1           2
     PREDICTF90 1.3

 Parameter file:             pred.par
 Data file:                  pred.dat
 Number of Traits             1
 Number of Effects            2
 Position of Observations      1
 Position of Weight (1)        0
 Value of Missing Trait/Observation           0

EFFECTS
 #  type                position (2)        levels   [positions for nested]
 1  cross-classified     2                                                 1
 2  cross-classified     3                                              6100

 Residual (co)variance Matrix
 0.70000

 Random Effect(s)    2
 Type of Random Effect:      additive animal
 Pedigree File:              renadd02.ped
 trait   effect    (CO)VARIANCES
  1       2    0.3000

 REMARKS
  (1) Weight position 0 means no weights utilized
  (2) Effect positions of 0 for some effects and traits means that such
      effects are missing for specified traits

 Data record length =            3
 # equations =         6101
 *** effets to include in Yhat (T/F):  F T
 solutions read from file: soltutions
 Animal Effect:            2
 y(s), yhat(s), residual(s) in written in "yhat_residual" file
        1000 records read
 Trait:            1        1000
```

```
   mean Y          2.803014134522528E-003  var Y          0.982498631522066
   mean Yhat      -1.204401846975088E-002  var Yhat       7.851567205278141E-002
   cov (Y,Yhat)    8.256236151221331E-002  corr (Y,Yhat)                  0.297261012857303
 wrote bvs for animals in data in file "bvs.dat"
```

Predictivity

Output files from PREDICTF90

yhat_residual

yhat_residual has 4 columns: animal | y | yhat | residual

```
4644     -0.266520      0.415535      0.339710
2176     -0.418925      0.094263      0.508577
2934      2.154195      0.157782      3.018178
 797      1.226565      0.328604      1.919726
6021      1.143225      0.430780      1.734210
```

Because OPTION include_effects 2 was used:
y is phenotype minus all effects other than animal
yhat receives the second effect, which is the animal effect
residual is phenotype minus animal effect

bvs.dat

bvs.dat has 4 columns: trait | effect | Animal | solution (EBV)

```
1  2    4644      0.415535
1  2    2176      0.094263
1  2    2934      0.157782
1  2     797      0.328604
1  2    6021      0.430780
```

Log file for predicting genotyped animals in 5th generation

```
name of parameter file?
pred.par

 *** include effets to predict Yhat n, effects            1           2
     PREDICTF90 1.3

 Parameter file:              gen.par
 Data file:                   gen.dat
 Number of Traits              1
 Number of Effects             2
 Position of Observations      1
 Position of Weight (1)        0
 Value of Missing Trait/Observation           0

EFFECTS
 #  type                position (2)        levels    [positions for nested]
 1  cross-classified     2                                                  1
 2  cross-classified     3                                               6100

 Residual (co)variance Matrix
 0.70000

 Random Effect(s)     2
 Type of Random Effect:       additive animal
 Pedigree File:               renadd02.ped
 trait    effect     (CO)VARIANCES
  1        2     0.3000
```

```
REMARKS
 (1) Weight position 0 means no weights utilized
 (2) Effect positions of 0 for some effects and traits means that such
     effects are missing for specified traits

Data record length =            3
# equations =         6101
*** effets to include in Yhat (T/F):  F T
solutions read from file: soltutions
Animal Effect:            2
y(s), yhat(s), residual(s) in written in "yhat_residual" file
        300 records read
Trait:            1         300
   mean Y        -5.204056186291079E-002  var Y          0.979795877964320
   mean Yhat     -1.187536126623551E-002  var Yhat       7.349890384221654E-002
   cov (Y,Yhat)   8.232182257800019E-002  corr (Y,Yhat)  0.306765659847626
wrote bvs for animals in data in file "bvs.dat"
```

Predictivity

2) Predictivity for ssGBLUP

Parameter file:

```
DATAFILE
 pred.dat
NUMBER_OF_TRAITS
      1
NUMBER_OF_EFFECTS
      2
OBSERVATION(S)
  1
WEIGHT(S)

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT[EFFECT NESTED]
 2       1 cross
 3     6100 cross
RANDOM_RESIDUAL VALUES
 0.70000
 RANDOM_GROUP
   2
 RANDOM_TYPE
 add_animal
 FILE
renadd02.ped
(CO)VARIANCES
 0.30000
OPTION SNP_file snp.txt
OPTION chrinfo map.txt
OPTION include_effects 2
```

Log file for predicting all animals in 5th generation

```
name of parameter file?
pred.par

 *** include effets to predict Yhat n, effects            1           2
```

```
    PREDICTF90 1.3

 Parameter file:             pred.par
 Data file:                  pred.dat
 Number of Traits             1
 Number of Effects            2
 Position of Observations     1
 Position of Weight (1)       0
 Value of Missing Trait/Observation          0

EFFECTS
 #  type                position (2)        levels    [positions for nested]
 1  cross-classified     2                                                  1
 2  cross-classified     3                                               6100

 Residual (co)variance Matrix
 0.70000

 Random Effect(s)    2
 Type of Random Effect:      additive animal
 Pedigree File:              renadd02.ped
 trait   effect    (CO)VARIANCES
  1       2    0.3000

 REMARKS
  (1) Weight position 0 means no weights utilized
  (2) Effect positions of 0 for some effects and traits means that such
      effects are missing for specified traits

 Data record length =            3
 # equations =         6101
 *** effets to include in Yhat (T/F):  F T
 solutions read from file: soltutions
 Animal Effect:            2
 y(s), yhat(s), residual(s) in written in "yhat_residual" file
       1000 records read
 Trait:            1        1000
    mean Y         1.729620725009590E-002  var Y          0.982498616557168
    mean Yhat     -1.458232188504189E-002  var Yhat       8.642763611542703E-002
    cov (Y,Yhat)   9.189071888926587E-002  corr (Y,Yhat)  0.315340214001692
```

Predictivity

Output files from PREDICTF90

Output files for predictf90 fallow the same pattern whether using genomic information or not.

Log file for predicting genotyped animals in 5th generation

```
name of parameter file?
pred.par

 *** include effets to predict Yhat n, effects           1           2
     PREDICTF90 1.3

 Parameter file:             gen.par
 Data file:                  gen.dat
 Number of Traits             1
 Number of Effects            2
 Position of Observations     1
 Position of Weight (1)       0
 Value of Missing Trait/Observation          0
```

```
EFFECTS
 #  type                position (2)        levels    [positions for nested]
 1  cross-classified     2                                                  1
 2  cross-classified     3                                               6100

 Residual (co)variance Matrix
 0.70000

 Random Effect(s)    2
 Type of Random Effect:      additive animal
 Pedigree File:              renadd02.ped
 trait   effect    (CO)VARIANCES
  1       2    0.3000

 REMARKS
  (1) Weight position 0 means no weights utilized
  (2) Effect positions of 0 for some effects and traits means that such
      effects are missing for specified traits

 Data record length =            3
 # equations =         6101
 *** effets to include in Yhat (T/F):  F T
 solutions read from file: soltutions
 Animal Effect:            2
 y(s), yhat(s), residual(s) in written in "yhat_residual" file
        300 records read
 Trait:            1         300
    mean Y        -3.754737233898292E-002  var Y          0.979795861954017
    mean Yhat     -1.863066248595715E-002  var Yhat       0.119326686734040
    cov (Y,Yhat)   0.117365728215231       corr (Y,Yhat)  0.343245384612940
 wrote bvs for animals in data in file "bvs.dat"
```

Predictivity

Parameter files for GWAS using ssGBLUP (ssGWAS)

Run BLUPF90 with genomic information and salve G^{-1} and A_{22}^{-1}

```
DATAFILE
 renf90.dat
NUMBER_OF_TRAITS
      1
NUMBER_OF_EFFECTS
      2
OBSERVATION(S)
   1
WEIGHT(S)

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT[EFFECT NESTED]
 2      1 cross
 3     6100 cross
RANDOM_RESIDUAL VALUES
 0.70000
 RANDOM_GROUP
   2
 RANDOM_TYPE
```

```
 add_animal
 FILE
renadd02.ped
(CO)VARIANCES
 0.30000
OPTION SNP_file snp.txt
OPTION chrinfo map.txt
OPTION no_quality_control
OPTION saveGInverse
OPTION saveA22Inverse
OPTION weightedG wei
```

Weights for SNP can be updated by an iterative process, where the initial weights are all equal to 1.

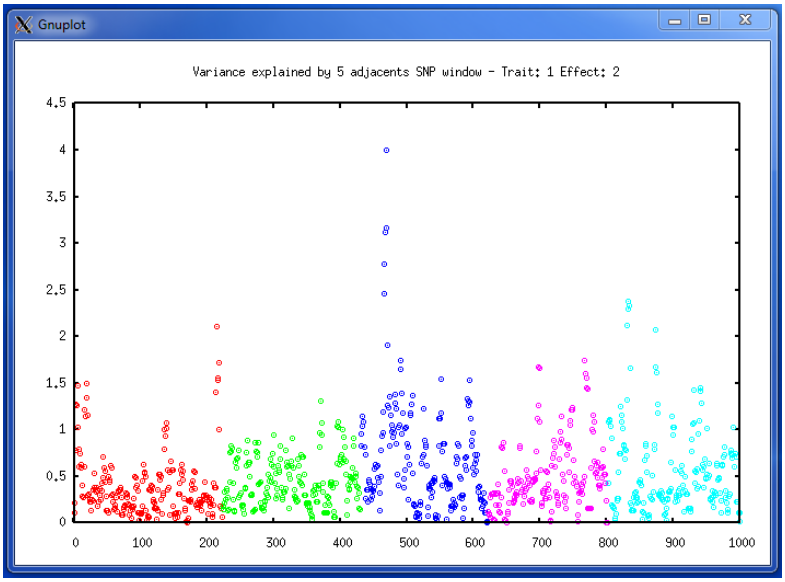
Linux code to get initial weights for 1000 SNP:
awk 'BEGIN { for (i==1;i<1000;i++) print 1}' > wei

Run POSTGSF90 and salve G^{-1} and A_{22}^{-1}

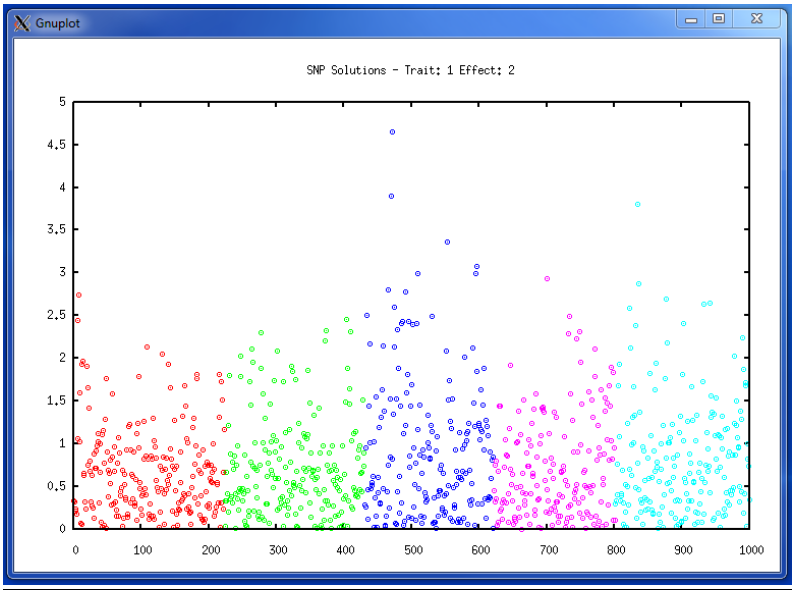
```
DATAFILE
 renf90.dat
NUMBER_OF_TRAITS
        1
NUMBER_OF_EFFECTS
        2
OBSERVATION(S)
   1
WEIGHT(S)

EFFECTS: POSITIONS_IN_DATAFILE NUMBER_OF_LEVELS TYPE_OF_EFFECT[EFFECT NESTED]
 2        1 cross
 3     6100 cross
RANDOM_RESIDUAL VALUES
 0.70000
 RANDOM_GROUP
    2
 RANDOM_TYPE
 add_animal
 FILE
renadd02.ped
(CO)VARIANCES
 0.30000
OPTION SNP_file snp.txt
OPTION chrinfo map.txt
OPTION no_quality_control
OPTION Manhattan_plot
OPTION readGInverse
OPTION readA22Inverse
OPTION weightedG wei
OPTION windows_variance 5
```

Moving average of SNP effects can be obtained by using the following option:
OPTION SNP_moving_average n
where n is the number of SNP

Manhattan plots for SNP windows variance



Manhattan plots for SNP effect using moving average of 2 SNP



Output files for ssGWAS

snp_sol

1	2	1	1	0	0.7001368E-02	0.2209213	0.1119293	0.1126648E-03
1	2	2	1	0	-0.1359349E-01	0.5065436	0.2104747	0.2118577E-03
1	2	3	1	0	0.8714214E-02	0.3917027	0.7757968	0.7808942E-03
1	2	4	1	0	-0.4223401E-02	0.6873333E-01	1.271113	0.1279465E-02
1	2	5	1	0	0.5471629E-03	0.1539137E-02	1.261010	0.1269296E-02

snp_sol has 9 columns:
trait | effect | SNP | chromosome | position | SNP_solution | weight | % of variance explained by n adjacent SNP | variance explained by n adjacent SNP

chrsnpvar

1	2	0.1119293459	1	1	0
1	2	0.2104747339	2	1	0
1	2	0.7757968029	3	1	0
1	2	1.2711127978	4	1	0
1	2	1.2610103595	5	1	0

chrsnpvar has 6 columns:
trait | effect | % of variance explained by n adjacent SNP | SNP | chromosome | position

This file is used by POSTGSF90 for Manhattan plots

Appendix J (selected programming details)

This section provides some programming insight into an early version of the blupf90 program. The model is completely described in the module MODEL.

```
module model
implicit none

!        Types of effects
integer,parameter::effcross=0,& !effects can be cross-classified
                   effcov=1     !or covariables

!        Types of random effects
integer,  parameter ::  g_fixed=1,&       ! fixed effect
                        g_diag=2, &       ! diagonal
                        g_A=3, &          ! additive animal
                        g_A_UPG=4, &      ! additive animal with unknown
                                          !       parent groups
                      & g_A_UPG_INB=5, &  ! additive animal with unknown
                                          !       parent groups and inbreeding
                      & g_As=6,&          ! additive sire
                        g_PD =7, &         ! parental dominance
                        g_last=8          ! last type

character (40)     ::   parfile, &     !name of parameter file
                        datafile       !name of data set

integer :: ntrait,&                  !number of traits
           neff,&                    !number of effects
           miss=0                      !value of missing trait/effect


integer,allocatable :: pos_y(:)        !positions of observations
integer ::             pos_weight      ! position of weight of records; zero if none


integer,allocatable :: pos_eff(:,:),&  !positions of effects for each trait
                       nlev(:),&       !number of levels
                       effecttype(:),& !type of effects
                       nestedcov(:,:),&!position of nesting effect for each trait
                                       ! if the effect is nested covariable
                     & randomtype(:),& ! status of each effect, as above
                       randomnumb(:)   ! number of consecutive correlated effects

character (40),allocatable::  randomfile(:)   ! name of file associated with given
                                                 ! effect

real, allocatable  ::  r(:,:),&        !residual (co)variance matrix
                       rinv(:,:),&     ! and its inverse
                       g(:,:,:)      ! The random (co)variance matrix for each trait
end module model
```

The core of the program is presented below.

```
program BLUPF90
use model;use sparsem; use sparseop
implicit none
real,allocatable :: y(:),&                     ! observation value
                    indata(:)                  ! one line of input data

real ::             weight_y                   ! weight for records

type (sparse_hashm)::xx              ! X'X in sparse hash form
type (sparse_ija):: xx_ija           ! X'X in IJA form, for use with FSPAK only
real, allocatable:: xy(:),sol(:)       !X'Y and solutions
```

```
real,allocatable :: weight_cov(:,:)
integer,allocatable:: address(:,:)      ! start and address of each effect
integer :: neq,io,&                     ! number of equations and io-status
           data_len,&                   ! length of data record to read
           i,j,k,l                      ! extra variables
real::  val, dat_eff

!
call read_parameters
call print_parameters
neq=ntrait*sum(nlev)
data_len=max(pos_weight,maxval(pos_y),maxval(pos_eff))
print*,'Data record length = ',data_len
allocate (xy(neq), sol(neq),address(neff,ntrait),&
          weight_cov(neff,ntrait),y(ntrait),indata(data_len))
call zerom(xx,neq); xy=0
!
call setup_g                    ! invert R matrices

open(50,file=datafile)         !data file

! Contributions from records
do
   read(50,*,iostat=io)indata
   if (io.ne.0) exit
   call decode_record
   call find_addresses
   call find_rinv
   do i=1,neff
       do j=1,neff
          do k=1,ntrait
             do l=1,ntrait
                 val=weight_cov(i,k)*weight_cov(j,l)*weight_y*rinv(k,l)
                 call addm(val,address(i,k),address(j,l),xx)
             enddo
          enddo
       enddo
       do k=1,ntrait
          do l=1,ntrait
              xy(address(i,k))=xy(address(i,k))+rinv(k,l)*y(l)*weight_cov(i,k) &
                                                              *weight_y
          enddo
       enddo
   enddo
enddo
!
!     Random effects' contributions
do i=1,neff
   select case (randomtype(i))
     case (g_fixed)
        continue                ! fixed effect, do nothing
     case (g_diag)
        call add_g_diag(i)
     case (g_A, g_As, g_A_UPG,g_A_UPG_INB)
        call add_g_add(randomtype(i),i)
     case (g_PD)
        call add_g_domin(i)
     case default
       print*,'unimplemented random type',randomtype(i)
   endselect
enddo

if (neq < 15) then
   print*,'left hand side'
   call printm(xx)
   print  '( '' right hand side:'' ,100f8.1)',xy
endif

 call solve_iterm(xx,xy,sol)

! Comment the line above and uncomments the lines below only if
```

```
! solutions by FSPAK are desired
!xx_ija=xx;
!call fspak90(‘solve’,xx_ija,xy,sol)


if (neq <15) print  '( '' solution:'' ,100f7.3)',sol

call store_solutions
```