

Command Language Interface of MiX99 (CLIM)

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MiX99 course: test-day models and single step genomic prediction

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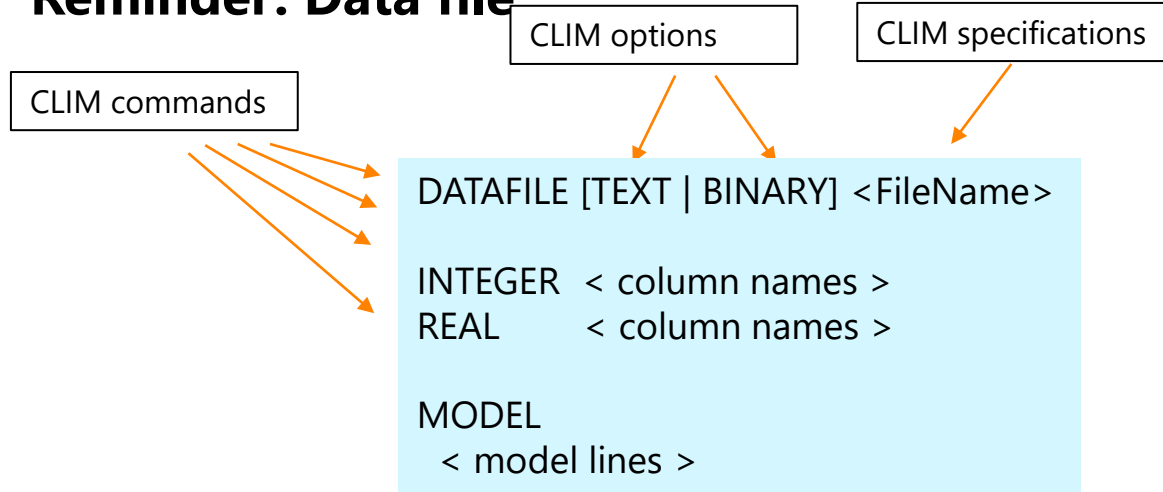
Introduction

Assumed that basic concepts of CLIM known

Contents

- Reminder: simplest CLIM
- GBLUP: many ways to achieve the same
- SNPBLUP and RegIndex to easy computations
- Giving variance components by CLIM
- Complex models and macros

Reminder: Data file



- Default format is “text format”, where columns are separated by space (alternative: binary format)
- There must not be a header in the file
- Sorting of records can reduce solving time

• Example:

```

DATAFILE  data.dat

INTEGER   id herd year
REAL     xlin x2 pheno1 pheno2

MODEL
  pheno1 = xlin x2 year herd
  pheno2 = xlin x2 year herd

```

	id	herd	year	xlin	x2	pheno1	pheno2
1	1	1	0.17001745	0.028905933	-0.7079236	-0.7079236	
2	1	2	0.81993105	0.672286934	25.1489620	25.1489620	
3	1	1	0.10122364	0.010246224	1.3475760	1.3475760	
4	1	2	0.89622887	0.803226186	27.0999594	27.0999594	
5	1	1	0.33690867	0.113507451	2.7174209	2.7174209	
...							

- The **data file** contains observations and model effect information (classification effects or regression coefficients)
- Each record ,i.e., line has two parts
 - 1st part: **integer numbers** (classification effects, sorting variables, index values)
 - 2nd part: **real numbers** (observations, covariables, weights)
- Integer numbers ≤ 0 are assumed as missing
- Missing real values are coded by a missing variable code (e.g. -9999.0)

Some ways to compute a simple two trait GBLUP model

```
DATAFILE data.dat
INTEGER id herd year
REAL xlin x2 pheno1 pheno2
```

```
PEDFILE LOWER iG.dat
PEDIGREE id FILE
RANDOM id
PARFILE vc_2tr.par
```

Equivalent in MiX99

```
DATAFILE data.dat
INTEGER id herd year
REAL xlin x2 pheno1 pheno2
```

```
GBLUP id LOWER iG.dat
RANDOM id
PARFILE vc_2tr.par
```

Different in MiX99

COVFILE approach

```
DATAFILE data.dat
INTEGER id herd year
REAL xlin x2 pheno1 pheno2
```

```
COVFILE id LOWER iG.dat
RANDOM id
PARFILE vc_2tr.par
```

But results are the same

```
MODEL
phenol = xlin x2 year herd id
phenol2 = xlin x2 year herd id
```

```
MODEL
phenol = xlin x2 year herd id
phenol2 = xlin x2 year herd id
```

```
MODEL
phenol = xlin x2 year herd id
phenol2 = xlin x2 year herd id
```

Solreg:

Trt	Reg-No	Solution	Trait	Covariable
1	1	9.6142	phenol	xlin
1	2	5.1683	phenol	x2
2	1	9.6142	phenol2	xlin
2	2	5.1683	phenol2	x2

Solfix:

Fact.	Trt	Level	N-Obs	Solution
1	1	1	10	-1.7419
1	2	1	10	-1.7419
1	1	2	10	13.427
1	2	2	10	13.427
2	1	1	10	0.49912
2	2	1	10	0.49912
2	1	2	10	11.186
2	2	2	10	11.186

```
Factor Trait
year phenol
year phenol2
year phenol
year phenol2
herd phenol
herd phenol2
herd phenol
herd phenol2
```

```
iG.dat
5 5
1 2 3 4 5
27.5
-60.0 137.5
-15.0 35.0 10.0
25.0 -57.5 -15.0 25.0
-47.5 107.5 27.5 -45.0 85.0
```

Solani:

1	1	4	-0.85062E-01	-0.85062E-01
2	1	4	-0.63589E-01	-0.63589E-01
3	1	4	0.37509E-01	0.37509E-01
4	1	4	-0.98786E-01	-0.98786E-01
5	1	4	-0.30458E-01	-0.30458E-01

Genomic breeding values

Solr01:

1	4	-0.85062E-01	-0.85062E-01
2	4	-0.63589E-01	-0.63589E-01
3	4	0.37509E-01	0.37509E-01
4	4	-0.98786E-01	-0.98786E-01
5	4	-0.30458E-01	-0.30458E-01



Note: different formats of inverse G in GBLUP can lead to same computations

DATAFILE data.dat

INTEGER id herd year
REAL xlin x2 pheno1 pheno2

GBLUP id LOWER iG.dat

RANDOM id

PARFILE vc_2tr.par

MODEL

pheno1 = xlin x2 year herd id
pheno2 = xlin x2 year herd id

Inverse of G matrix in lower triangle dense format:

iG.dat

```
5 5
1 2 3 4 5
27.5
-60.0 137.5
-15.0 35.0 10.0
25.0 -57.5 -15.0 25.0
-47.5 107.5 27.5 -45.0 85.0
```

By default, the solver reads matrices to RAM
→ These formats lead to same computations.

Lower dense is more efficient (storage).

DATAFILE data.dat

INTEGER id herd year
REAL xlin x2 pheno1 pheno2

GBLUP id iGijv.dat

RANDOM id

PARFILE vc_2tr.par

MODEL

pheno1 = xlin x2 year herd id
pheno2 = xlin x2 year herd id

Inverse of G matrix in lower ijvalue format:

iGijv.dat

```
1 1 27.5
2 1 -60.0
3 1 -15.0
4 1 25.0
5 1 -47.5
2 2 137.5
3 2 35.0
4 2 -57.5
5 2 107.5
3 3 10.0
4 3 -15.0
5 3 27.5
4 4 25.0
5 4 -45.0
5 5 85.0
```

→ Some approaches may look different but solver may treat them the same.

For large models, "bin" or "raw" formats are more efficient.

We'll cover that later.

SNPBLUP.clm: DATAFILE data.dat

Example: SNPBLUP model by REGMATRIX

INTEGER id herd year
 REAL xlin x2 pheno1 pheno2

DATASORT PEDIGREECODE=id

REGFILE RM_geno.dat
 REGMATRIX Random SNP id=1 first=2 last=6
 REGPARFILE snp_2tr.par # snp variance
 PARFILE res_2tr.par

This SNPBLUP model produces marker solutions, not directly breeding values.

MODEL
 pheno1 = xlin x2 year herd
 pheno2 = xlin x2 year herd

Solreg:

Trt	Reg-No	Solution	Trait	Covariable
1	1	9.6142	pheno1	xlin
1	2	5.1683	pheno1	x2
2	1	9.6142	pheno2	xlin
2	2	5.1683	pheno2	x2

Solfix:

Fact.	Trt	Level	N-Obs	Solution	Factor Trait
1	1	1	10	-1.7949	year pheno1
1	2	1	10	-1.7949	year pheno2
1	1	2	10	13.374	year pheno1
1	2	2	10	13.374	year pheno2

Solf01:

Trt	N-Obs	Solution
1	10	0.44614
2	10	11.133

Solreg_mat:

Trt	Matrix	Effect	Solution	Mat-Name
1	1	1	0.59477E-01	SNP
1	1	2	0.33264E-01	SNP
1	1	3	0.89845E-02	SNP
1	1	4	0.34570E-01	SNP
1	1	5	-0.30324E-01	SNP
2	1	1	0.59477E-01	SNP
2	1	2	0.33264E-01	SNP
2	1	3	0.89845E-02	SNP
2	1	4	0.34570E-01	SNP
2	1	5	-0.30324E-01	SNP

Solutions similar but No breeding values directly by this SNPBLUP model and preparing data is awkward.

GBLUP.clm

Trt	Reg-No	Solution	Trait	Covariable
1	1	9.6142	pheno1	xlin
1	2	5.1683	pheno1	x2
2	1	9.6142	pheno2	xlin
2	2	5.1683	pheno2	x2

Fact.	Trt	Level	N-Obs	Solution	Factor Trait
1	1	1	10	-1.7419	year pheno1
1	2	1	10	-1.7419	year pheno2
1	1	2	10	13.427	year pheno1
1	2	2	10	13.427	year pheno2
2	1	1	10	0.49912	herd pheno1
2	2	1	10	0.49912	herd pheno2
2	1	2	10	11.186	herd pheno1
2	2	2	10	11.186	herd pheno2

Solani:

Trt	Reg-No	Solution	Trait	Covariable
1	1	4 -0.85062E-01	-0.85062E-01	
2	1	4 -0.63589E-01	-0.63589E-01	
3	1	4 0.37509E-01	0.37509E-01	
4	1	4 -0.98786E-01	-0.98786E-01	
5	1	4 -0.30458E-01	-0.30458E-01	



Example: Using REGINDEX in REGMATRIX

SNPBLUP.clm: DATAFILE data.dat

INTEGER id herd year
REAL xlin x2 pheno1 pheno2

DATASORT PEDIGREECODE=id

```
REGFILE RM_geno.dat
REGMATRIX Random SNP id=1 first=2 last=6
REGPARFILE snp_2tr.par # snp variance
```

```
PARFILE res_2tr.par
```

MODEL

```
pheno1 = xlin x2 year herd
pheno2 = xlin x2 year herd
```

data_geno.dat

```
1 1 1 0.17001745 0.028905933 -0.7079236
2 1 2 0.81993105 0.672286934 25.1489620
3 1 1 0.10122364 0.010246224 1.3475760
4 1 2 0.89622887 0.803226186 27.0999594
5 1 1 0.33690867 0.113507451 2.7174209
1 1 2 0.07535836 0.005678883 14.0722251
2 1 1 0.24611983 0.060574969 0.7791845
3 1 2 0.08260870 0.006824198 15.0323770
4 1 1 0.80893255 0.654371868 9.1249718
5 1 2 0.61464561 0.377789222 22.3646796
1 2 1 0.99636615 0.992745498 25.7296024
2 2 2 0.68082035 0.463516349 35.1481721
3 2 1 0.55235644 0.305097638 16.8500291
4 2 2 0.78372758 0.614228920 35.3142666
5 2 1 0.92074811 0.847777080 22.2667699
1 2 2 0.34093483 0.116236556 28.2127363
2 2 1 0.83466677 0.696668623 20.1699458
3 2 2 0.90065598 0.811181186 35.3930001
4 2 1 0.25502135 0.065035889 11.9655357
5 2 2 0.58140852 0.338035871 31.7120296
```

RM_geno.dat

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

RegMatrix data and DataFile data must align exactly!

→ When an individual has several observations, genotypes in the RegMatrix will be repeated.

- 1) In this case, a lot of repetition of the same data
- 2) REGFILE needs to be sorted in the same order as DATAFILE

It is simple to solve these using a **REGINDEX** in REGMATRIX.

Example: Using REGINDEX in REGMATRIX

SNPBLUP.clm DATAFILE data.dat

INTEGER id herd year
REAL xlin x2 pheno1 pheno2

DATASORT PEDIGREECODE=id

```
REGFILE RM_geno.dat
REGMATRIX Random SNP id=1 first=2 last=6
REGPARFILE snp_2tr.par # snp variance
```

```
PARFILE res_2tr.par
```

MODEL

```
pheno1 = xlin x2 year herd
pheno2 = xlin x2 year herd
```

DATAFILE data.dat

INTEGER id herd year
REAL xlin x2 pheno1 pheno2

DATASORT PEDIGREECODE=id

```
REGFILE RMi_geno.dat
REGMATRIX Random SNP id=1 RegIndex=id first=2 last=6
REGPARFILE snp_2tr.par # snp variance
```

```
PARFILE res_2tr.par
```

MODEL

```
pheno1 = xlin x2 year herd
pheno2 = xlin x2 year herd
```

data_geno.dat

```
1 1 1 0.17001745 0.028905933 -0.7079236
2 1 2 0.81993105 0.672286934 25.1489620
3 1 1 0.10122364 0.010246224 1.3475760
4 1 2 0.89622887 0.803226186 27.0999594
5 1 1 0.33690867 0.113507451 2.7174209
1 1 2 0.07535836 0.005678883 14.0722251
2 1 1 0.24611983 0.060574969 0.7791845
3 1 2 0.08260870 0.006824198 15.0323770
4 1 1 0.80893255 0.654371868 9.1249718
5 1 2 0.61464561 0.377789222 22.3646796
1 2 1 0.99636615 0.992745498 25.7296024
2 2 2 0.68082035 0.463516349 35.1481721
3 2 1 0.55235644 0.305097638 16.8500291
4 2 2 0.78372758 0.614228920 35.3142666
5 2 1 0.92074811 0.847777080 22.2667699
1 2 2 0.34093483 0.116236556 28.2127363
2 2 1 0.83466677 0.696668623 20.1699458
3 2 2 0.90065598 0.811181186 35.3930001
4 2 1 0.25502135 0.065035889 11.9655357
5 2 2 0.58140852 0.338035871 31.7120296
```

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

RM_geno.dat

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

RMi_geno.dat

RegIndex gives the owner of the row in the data file.

- 1) Each owner can have only one line in REGFILE
- 2) There is no need to sort the REGFILE file.



Example: SNPBLUP model

```

DATAFILE data.dat
SNPBLUPi.clm

INTEGER id herd year
REAL xlin x2 pheno1 pheno2

DATASORT PEDIGREECODE=id

REGFILE RMi_geno.dat
REGMATRIX Random SNP id=1 RegIndex=id first=2 last=6
REGPARFILE snp_2tr.par # snp variance

PARFILE res_2tr.par

MODEL
  pheno1 = xlin x2 year herd
  pheno2 = xlin x2 year herd
  
```

Results remain the same!

Solreg:	Trt	Reg-No	Solution	Trait	Covariable
	1	1	9.6142	pheno1	xlin
	1	2	5.1683	pheno1	x2
	2	1	9.6142	pheno2	xlin
	2	2	5.1683	pheno2	x2

Solfix:	Fact.	Trt	Level	N-Obs	Solution	Factor	Trait
	1	1	1	10	-1.7949	year	pheno1
	1	2	1	10	-1.7949	year	pheno2
	1	1	2	10	13.374	year	pheno1
	1	2	2	10	13.374	year	pheno2

Solf01:					
	1	10	0.44614	0.44614	
	2	10	11.133	11.133	

Solreg_mat:	Trt	Matrix	Effect	Solution	Mat-Name
	1	1	1	0.59477E-01	SNP
	1	1	2	0.33264E-01	SNP
	1	1	3	0.89845E-02	SNP
	1	1	4	0.34570E-01	SNP
	1	1	5	-0.30324E-01	SNP
	2	1	1	0.59477E-01	SNP
	2	1	2	0.33264E-01	SNP
	2	1	3	0.89845E-02	SNP
	2	1	4	0.34570E-01	SNP
	2	1	5	-0.30324E-01	SNP

New:
SolIDGV01:

	1	8	0.20909E-01	0.20909E-01	
	2	8	0.42382E-01	0.42382E-01	
	3	8	0.14348	0.14348	
	4	8	0.71849E-02	0.71849E-02	
	5	8	0.75513E-01	0.75513E-01	

GBLUP.clm					Trt	Reg-No	Solution	Trait	Covariable
					1	1	9.6142	pheno1	xlin
					1	2	5.1683	pheno1	x2
					2	1	9.6142	pheno2	xlin
					2	2	5.1683	pheno2	x2

Fact.	Trt	Level	N-Obs	Solution	Factor	Trait
1	1	1	10	-1.7419	year	pheno1
1	2	1	10	-1.7419	year	pheno2
1	1	2	10	13.427	year	pheno1
1	2	2	10	13.427	year	pheno2
2	1	1	10	0.49912	herd	pheno1
2	2	1	10	0.49912	herd	pheno2
2	1	2	10	11.186	herd	pheno1
2	2	2	10	11.186	herd	pheno2

Solani:						
	1	1	4	-0.85062E-01	-0.85062E-01	
	2	1	4	-0.63589E-01	-0.63589E-01	
	3	1	4	0.37509E-01	0.37509E-01	
	4	1	4	-0.98786E-01	-0.98786E-01	
	5	1	4	-0.30458E-01	-0.30458E-01	



Example: SNPBLUP model

SNPBLUPi.clm breeding values

SoldGV01:

1	8	0.20909E-01	0.20909E-01
2	8	0.42382E-01	0.42382E-01
3	8	0.14348	0.14348
4	8	0.71849E-02	0.71849E-02
5	8	0.75513E-01	0.75513E-01

GBLUP.clm breeding values

Solani:

1	1	4	-0.85062E-01	-0.85062E-01
2	1	4	-0.63589E-01	-0.63589E-01
3	1	4	0.37509E-01	0.37509E-01
4	1	4	-0.98786E-01	-0.98786E-01
5	1	4	-0.30458E-01	-0.30458E-01

Correlation is one but there is a level difference of 0.10597: $SoldGV01 = Solani + 0.10597$
This is due to not centering REGMATRIX as is done in the **G** matrix.

DATAFILE data.dat

INTEGER id herd year
REAL xlin x2 pheno1 pheno2

DATASORT PEDIGREECODE=id

REGFILE Rmi_gen0.dat
REGMATRIX Random SNP id=1 RegIndex=id first=2 last=6 **CENTER=1**
REGPARFILE snp_2tr.par # snp variance

PARFILE res_2tr.par

MODEL
pheno1 = xlin x2 year herd
pheno2 = xlin x2 year herd

Centering as for **G** in \mathbf{G}^{-1} of iG.dat gives SoldGV01:

1	8	-0.85062E-01	-0.85062E-01
2	8	-0.63589E-01	-0.63589E-01
3	8	0.37509E-01	0.37509E-01
4	8	-0.98786E-01	-0.98786E-01
5	8	-0.30458E-01	-0.30458E-01

Other useful options in REGMATRIX: SCALE & FORMAT

SCALE: allows using the same variances as for GBLUP model as scaling done like for the G matrix
FORMAT: allows SNP marker packing (FORMAT=m) or not space separated SNPs (FORMAT=s)

Similar options for single-step SNPBLUP considered later.

SNPBLUPi.clm

DATAFILE data.dat

INTEGER id herd year
REAL xlin x2 pheno1 pheno2

DATASORT PEDIGREECODE=id

REGFILE Rmi_gen0.dat
REGMATRIX Random SNP id=1 **RegIndex=id** first=2 last=6
REGPARFILE snp_2tr.par # snp variance

PARFILE res_2tr.par

MODEL
pheno1 = xlin x2 year herd
pheno2 = xlin x2 year herd



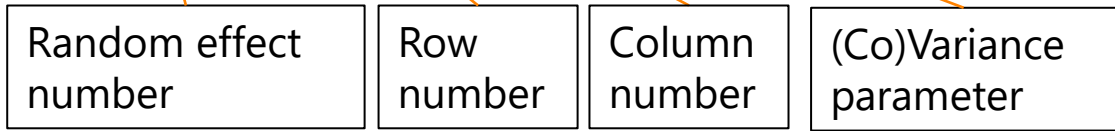
Variance component parameter file

GBLUP model

PARFILE Vc_2tr.par

```
1 1 1 1.0
1 2 2 1.0
2 1 1 2.0
2 2 2 2.0
```

Random effect number



- Genetic variance is 1.0
- Residual variance is 2.0

Alternatively (in new MiX99)

PARFILE CLIM

```
1 LOWER
  1.0
  0.0 1.0
2 DIAGONAL
  2.0
  2.0
```

It is possible to mix formats after "PARFILE CLIM":

LOWER
DIAGONAL
IDENTITY
SPARSE

Lower dense matrix
A diagonal matrix
Identity matrix
The typical parfile format

Special option 1: PARFILE IDENTITY → all variance matrices set to identity matrices

Special option 2: PARFILE MIXED <parfile> → parfile has the LOWER/DIAGONAL ... commands



Variance component parameter file

GBLUP model

PARFILE vc_2tr.par

vc_2tr.par:

```
1 1 1 1.0
1 2 2 1.0
2 1 1 2.0
2 2 2 2.0
```

Random effect
number

PARFILE MIXED vc_2tr.mix

vc_2tr.mix:

```
1 IDENTITY
2 DIAGONAL
  2.0
  2.0
```

Or in CLIM equivalently:

PARFILE CLIM

```
1 IDENTITY
2 DIAGONAL
  2.0
  2.0
```

Random effect
number

Row
number

Column
number

(Co)Variance
parameter

→ Genetic variance is 1.0

→ Residual variance is 2.0

Reminder: MODEL instructions with component names

Single trait repeatability model with a fixed regression function

$$\text{milk} = c_1 * \text{cov1} + c_2 * \text{cov2} + c_3 * \text{cov3} + c_4 * \text{cov4} + \text{HYS} + p + a + e$$

- p = nonhereditary animal effect
- a = random additive genetic effect
- where the regression function could be a 3rd order Legendre on days in milk

```

DATAFILE  RM.dat  # Data file
PEDFILE   RM.ped  # Pedigree file
PARFILE   RM.var  # Variance component file

INTEGER   animal HYS DIM          # Integer column names
REAL      milk  cov1 cov2 cov3 cov4 # Real column names

MODEL
milk = cov1 cov2 cov3 cov4 HYS P(animal) G(animal) # The model

PEDIGREE  G am    # Genetics associated with animal code: am=animal model
RANDOM     P      # Random nonhereditary animal effect

WITHINBLOCKORDER G P HYS # Random effects are within block effects

```

- General regression effects are always given at the beginning of the model line
- Only the covariables are specified

User-defined component names: P and G

Example: two trait GBLUP model (PARFILE CLIM available in new MiX99)

```
DATAFILE small_GBLUP_with_misses.dat
```

```
INTEGER ID ones HerdYear
REAL Cov1 Y1 Y2
```

```
MISSING -99999.
```

```
DATASORT PEDIGREECODE=ID
```

```
GBLUP DGV LOWER iGL.dat
```

```
PARFILE CLIM
```

```
1 DIAGONAL
```

```
2.0
```

```
2.1
```

```
2 LOWER
```

```
1.0
```

```
0.5 0.8
```

```
0.1 0.01 0.2
```

```
3 LOWER
```

```
3.0
```

```
0.8 3.1
```

```
Tmpfile.par
```

```
1 1 1 2.0
```

```
1 2 2 2.1
```

```
2 1 1 1.0
```

```
2 2 1 0.5
```

```
2 2 2 0.8
```

```
2 3 1 0.1
```

```
2 3 2 0.01
```

```
2 3 3 0.2
```

```
3 1 1 3.0
```

```
3 2 1 0.8
```

```
3 2 2 3.1
```

DGV variance of trait 1, component 1

DGV covariance between 1st components in traits 1 & 2

DGV variance of trait 2

DGV covariance of trait 1 components 1 and Cov1

DGV covariance of Cov1 of trait 1 and trait 2

DGV variance of Cov1 of trait 1

PE
DGV
Residual

```
RANDOM PE DGV
```

```
MODEL
```

```
Y1 = HerdYear - PE(ID) DGV(1 Cov1 | ID)
```

```
Y2 = - ones PE(ID) DGV(1 - | ID)
```

Variance counting is column down from left to right



Macros can simplify models

- Some model parts are replaced by a short name
- Less errors because same macro used in different traits
- Large models look simpler

Example model:

- 6 trait model with maternal and direct effects
- Showing the model is not possible
- Following are given steps to simpler macro based model

Macro: DEFINE <name> <what this name means>

NOTE: macro name should not be the same as an already used CLIM command or column name or component name.
Depending on where macro has been defined, this rule may be broken (not recommended!)

Using CLIM having macros:
mix99i --usemacros AMBLUP_macros.clm
Where AMBLUP_macros.clm has the CLIM file.

```

DEFINE HOFR      HO FR
DEFINE OTHER     JE MO NO RR ID
DEFINE MIS5      - - - - -
DEFINE BREEDS    AA AU BA BB CH HE LM PI PT SA SH SI IB
DEFINE MAT7      damHO damFR damJE damMO damNO damRR damID
DEFINE MAT2      damHO damFR - - - - -
DEFINE DAMH      damDY1 damDY2 damAA damAU damBA damBB damCH damHE damLM damPI damPT damSA damSH damSI damIB
DEFINE DAMM      - - - - -
DEFINE HFxA      FRxJEh FRxH0h H0xJEh DYxDYh
DEFINE HFxM      - - - - -
DEFINE OTxB      TRxTRh C0xC0h OBx0Bh C0xDYh C0xTRh C0x0Bh DYxTRh DYx0Bh TRx0Bh
DEFINE OTxM      - - - - -
DEFINE HYDH      hyDH - - - -
DEFINE HYDC      - hyDC - - - -
DEFINE HYBH      - - hyBH - - -
DEFINE HYBC      - - - hyBC - -
DEFINE HYcalv    - - - - hycalv

MODEL
DH   = HOFR OTHER - - BREEDS MAT7 DAMM HFxA OTxM agedam - - byr sex - HYDH G(anim dam)
DC   = HOFR OTHER - - BREEDS MAT7 DAMM HFxA OTxM - qq(agedam|dampar) byr sex damp HYDC G(anim dam)
BH   = HOFR MIS5 DY1 DY2 BREEDS MAT2 DAMH HFxM OTxB agedam - - byr sex - HYBH G(anim dam)
BC   = HOFR MIS5 DY1 DY2 BREEDS MAT2 DAMH HFxM OTxB - qq(agedam|dampar) byr sex damp HYBC G(anim dam)
bsize = HOFR MIS5 DY1 DY2 BREEDS MAT2 DAMH HFxM OTxB - qq(agedam|dampar) byr sex damp HYcalv G(anim dam)
bwt  = HOFR MIS5 DY1 DY2 BREEDS MAT2 DAMH HFxM OTxB - qq(agedam|dampar) byr sex damp HYcalv G(anim dam)

```

These effects are all the same for all traits

These effects can differ by trait: macro makes a substitution



Remember: Order of effects have to follow the required order: regression, fixed class effects, random effects, additive genetics

Macros using macros possible

```

DEFINE HOFR      HO FR
DEFINE OTHER    JE MO NO RR ID
DEFINE MIS5     - - - - -
DEFINE BREEDS   AA AU BA BB CH HE LM PI PT SA SH SI IB
DEFINE MAT7     damHO damFR damJE damMO damNO damRR damID
DEFINE MAT2     damHO damFR - - - - -
DEFINE DAMH     damDY1 damDY2 damAA damAU damBA damBB damCH damHE damLM damPI damPT damSA damSH damSI damIB
DEFINE DAMM     - - - - -
DEFINE HFxA     FRxJEh FRxHOh HOxJEh DYxDYh
DEFINE HFxM     - - - - -
DEFINE OTxB     TRxTRh COxC0h OBxOBh COxDYh COxTRh COxOBh DYxTRh DYxOBh TRxOBh
DEFINE OTxM     - - - - -
DEFINE HYDH     hyDH - - -
DEFINE HYDC     - hyDC - -
DEFINE HYBH     - - hyBH -
DEFINE HYBC     - - - hyBC -
DEFINE HYcalv   - - - - hycalv
DEFINE COVHC    HOFR OTHER - - BREEDS MAT7 DAMM HFxA OTxM
DEFINE COVOTH   HOFR MIS5 DY1 DY2 BREEDS MAT2 DAMH HFxM OTxB

```

Using CLIM having macros:

```
mix99i --usemacros AMBLUP_macros2.clm
```

Where AMBLUP_macros2.clm has the CLIM file.

Macros having macros are defined after the macros they include

```

MODEL
DH   = COVHC   agedam - - byr sex - HYDH G(anim dam)
DC   = COVHC   - qq(agedam|dampar) byr sex damp HYDC G(anim dam)
BH   = COVOTH  agedam - - byr sex - HYBH G(anim dam)
BC   = COVOTH  - qq(agedam|dampar) byr sex damp HYBC G(anim dam)
bsize = COVOTH - qq(agedam|dampar) byr sex damp HYcalv G(anim dam)
bwt  = COVOTH  - qq(agedam|dampar) byr sex damp HYcalv G(anim dam)

```

A simplified model

```

DEFINE H0FR      H0 FR
DEFINE OTHER    JE M0 N0 RR ID
DEFINE MIS5     - - - - -
DEFINE BREEDS   AA AU BA BB CH HE LM PI PT SA SH SI IB
DEFINE MAT7     damH0 damFR damJE damM0 damN0 damRR damID
DEFINE MAT2     damH0 damFR - - - - -
DEFINE DAMH     damDY1 damDY2 damAA damAU damBA damBB damCH damHE damLM damPI damPT damSA damSH damSI damIE
DEFINE DAMM     - - - - -
DEFINE HFxA     FRxJEh FRxH0h H0xJEh DYxDYh
DEFINE HFxM     - - - - -
DEFINE OTxB     TRxTRh C0xC0h 0Bx0Bh C0xDYh C0xTRh C0x0Bh DYxTRh DYx0Bh TRx0Bh
DEFINE OTxM     - - - - -
DEFINE HYDH     hyDH - - - -
DEFINE HYDC     - hyDC - - -
DEFINE HYBH     - - hyBH - -
DEFINE HYBC     - - - hyBC -
DEFINE HYcalv   - - - - hycalv
DEFINE COVHC    H0FR OTHER - - BREEDS MAT7 DAMM HFxA OTxM
DEFINE COV0TH  H0FR MIS5 DY1 DY2 BREEDS MAT2 DAMH HFxM OTxB
DEFINE FIXC1    agedam - - byr sex -
DEFINE FIXC2    - qq(agedam|dampar) byr sex damp
DEFINE GENETIC  G(anim dam)

```

```

MODEL
DH   = COVHC  FIXC1 HYDH  GENETIC
DC   = COVHC  FIXC2 HYDC  GENETIC
BH   = COV0TH FIXC1 HYBH  GENETIC
BC   = COV0TH FIXC2 HYBC  GENETIC
bsize = COV0TH FIXC2 HYcalv GENETIC
bwt   = COV0TH FIXC2 HYcalv GENETIC

```

Using CLIM having macros:

```
mix99i --usemacros AMBLUP_macros2.clm
```

Where AMBLUP_macros2.clm has the CLIM file.



Using macros may give uninformative error messages

```
***** CLIM Syntax Error *****
Error:                               Time: 18:01:20.5 08.04.2025
  Syntax error: Expecting left parenthesis. Possibly incorrectly written effect name.

On line: 17
Near token: HOFE -
Complete line information:
  BH   = HOFE - - - - - DY1 DY2 AA AU BA BB CH HE LM PI PT SA SH SI IB damHO damFR
damSA damSH damSI damIB - - - - - TRxTRh C0xC0h 0Bx0Bh C0xDYh C0xTRh C0x0Bh DYx
)
```

**Should have used
HOFR**

```
***** CLIM Syntax Error *****
Error:                               Time: 18:17:20.7 08.04.2025
  Syntax error: Expecting left parenthesis. Possibly incorrectly written effect name.

On line: 15
Near token: HOFR OTHER
Complete line information:
  DH   = HOFR OTHER - - BREEDS MAT7 DAMM HFxA OTxM agedam -
```

Should define macros using macros later

```
DEFINE COVHC   HOFR OTHER - - BREEDS MAT7 DAMM HFxA OTxM
DEFINE COV0TH  HOFR MIS5 DY1 DY2 BREEDS MAT2 DAMH HFxM OTxB
DEFINE HOFR    HO FR
DEFINE OTHER   JE MO NO RR ID
```

Error message usually looks like this and seem not very informative!

Here OTHER is not present in the original DATAFILE columns suggesting macro was not replaced by the string intended.

DEFINE MODEL HO FR → MODEL command is changed to be 'HO FR' → ERROR: Syntax error or unknown command.



Questions?

