





















Behavior of method LR to benchmark genetic evaluations

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# Summary

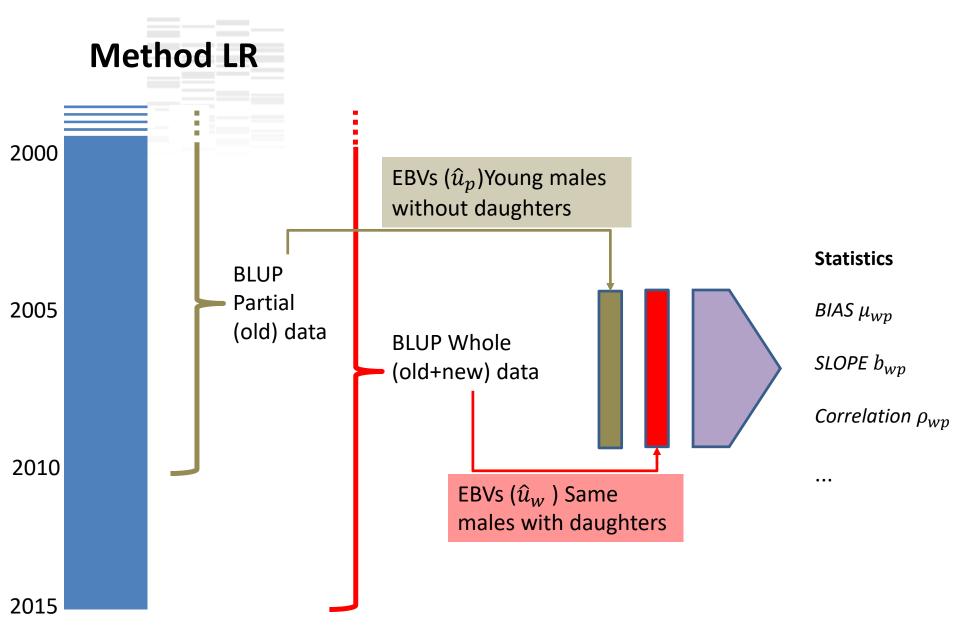
- Why another method to benchmark genetic evaluations?
- Method LR
- Simulations
- First results
- Conclusions



#### Why another method to benchmark genetic evaluations?

- In genomic evaluations cross validation is the most used tool for benchmarking
- All golden standard have problems:
  - Pre-corrected phenotypes may be not well corrected
  - Daughter Yield Deviations are not always available or might be inaccurate
  - Some traits (like maternal effects) don't have direct observation related to animals
- Need simple general tools for varied situations in animal breeding systems
- Legarra & Reverter (2017) proposed a new method based on comparisons of EBV from partial (old) data vs whole (old+new) data.
  - Does not require "true" breeding values
  - Does not require pre-corrected phenotypes







#### Method LR: Estimators.

#### **Bias**

 $\mu_{wp}=\overline{\widehat{u_p}}-\overline{\widehat{u_w}}$  . Expected value of 0 in absence of bias.

#### Slope of the regression EBVw on EBVp

$$b_{w,p}=rac{cov(\widehat{u_p},\widehat{u_w})}{var(\widehat{u_p})}$$
 . With a value of 1 in unbiased procedure.

#### Correlation between EBVp and EBVw.

Direct estimator of <u>relative increase of accuracy</u> from partial to whole.

$$\rho_{p,w} = \frac{cov(\widehat{u_w},\widehat{u_p})}{\sqrt{var(\widehat{u_w})var(\widehat{u_p})}}$$
. The expected value is  $E(\rho_{p,w}) \approx \frac{acc_p}{acc_w}$ .



#### **Objective**

Testing the estimators of bias, slope and accuracy using simulated selection schemes in several scenarios:

- 1. The genetic evaluation model is the correct one.
- 2. The genetic evaluation model is wrong.



#### Simulation details

Dairy sheep like scheme. Simulation was performed with QMSim software (version 1.10) (Sargolzaei & Schenkel, 2009).

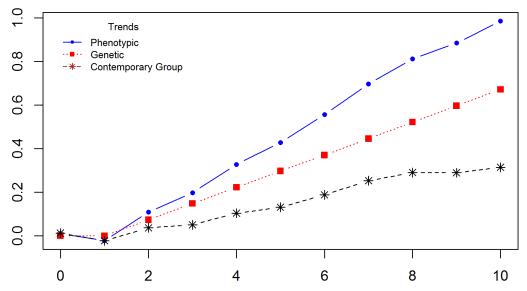
#### Parameters used:

- h<sup>2</sup> simulated: 0.05, 0.10, 0.25, 0.50
- 20 replicates for each h<sup>2</sup>
- Records only in females
- 10 generations
- Total animals in each replicate around 500,000
- Selection by higher EBV's



#### The genetic evaluation model is wrong

- 2 Strategies:
- 1. Contemporary groups with phenotypic trend
  - Around 90 CG/Generation with about 500 animals each.
  - Simulated: True effect of CG as random with time trend
  - Estimated: in BLUP as fixed effect



- 2. Using different h<sup>2</sup> in blup evaluations to those used for simulation (results not shown)
  - e.g. simulated h<sup>2</sup>=0.10 and evaluation h<sup>2</sup>=0.05



#### **Analyzed Datasets**

Within each replicate:

e.g.

At the end of generation 5: estimate EBV of young males (without progeny)  $\hat{u}_p$ At the end of generation 6: estimate EBV of the same males (with progeny)  $\widehat{u}_w$ Compute statistics:

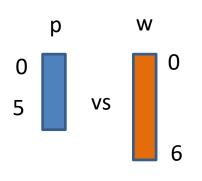
Bias

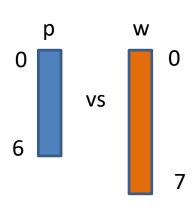
$$\mu_{wp} = \overline{\widehat{u_p}} - \overline{\widehat{u_w}}$$

Bias Slope 
$$\mu_{wp}=\overline{\widehat{u_p}}-\overline{\widehat{u_w}}$$
  $b_{w,p}=\frac{cov(\widehat{u_p},\widehat{u_w})}{var(\widehat{u_p})}$ 

$$\rho_{p,w} = \frac{ cov(\widehat{u_w}, \widehat{u_p}) }{ \sqrt{ var(\widehat{u_w}) var(\widehat{u_p}) } }$$

In this work we estimate the statistics for generations 5 to 9:







#### **First Results**

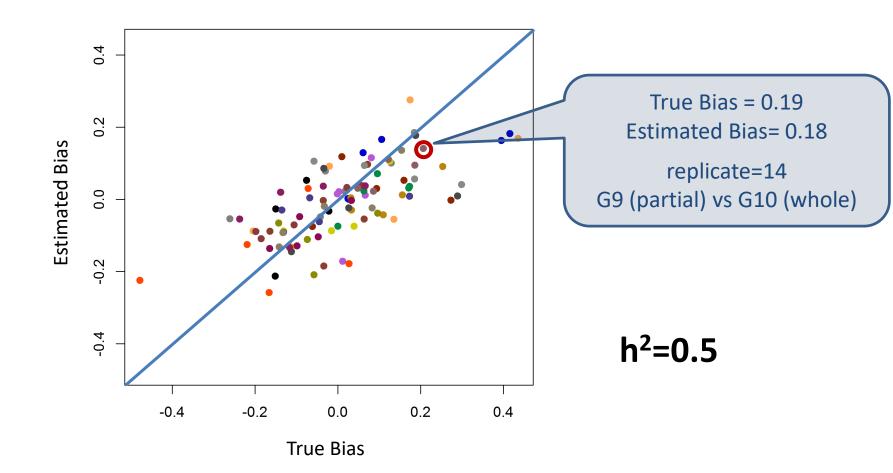
Using the correct evaluation model



#### **BIAS**

Estimated 
$$\Rightarrow \mu_{w,p} = \overline{\widehat{u_p}} - \overline{\widehat{u_w}}$$

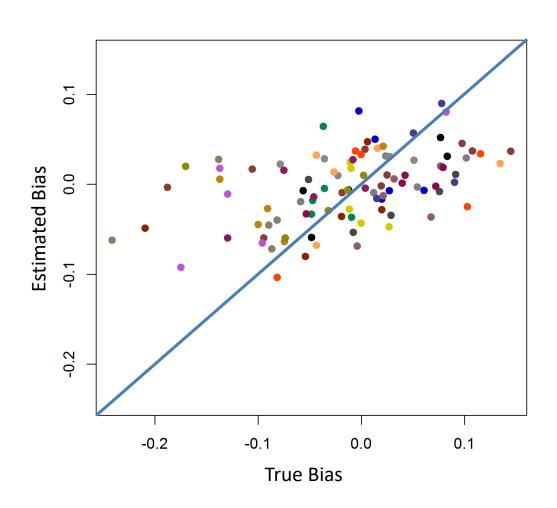
$$True \Rightarrow \mu_{u,p} = \overline{\widehat{u_p}} - \overline{u}$$



#### **BIAS**

Estimated Bias 
$$\Rightarrow \mu_{w,p} = \overline{\widehat{u_p}} - \overline{\widehat{u_w}}$$

True Bias 
$$\Rightarrow \mu_{u,p} = \overline{\widehat{u_p}} - \overline{u}$$

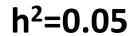


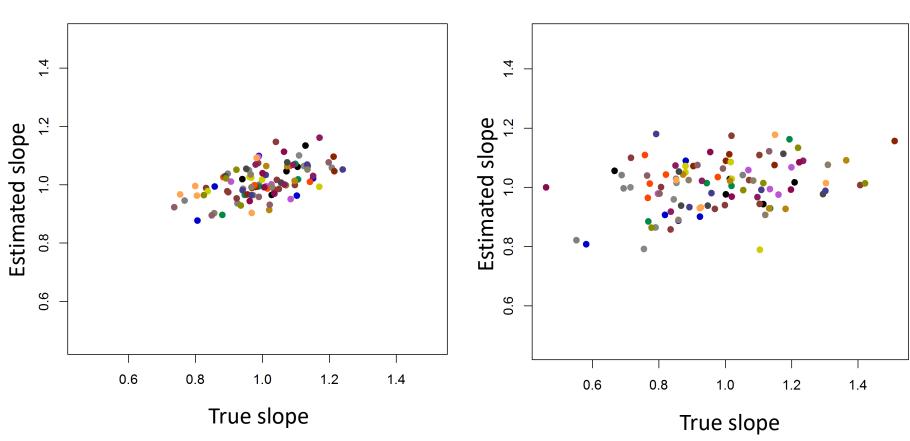
$$h^2=0.05$$

Estimated 
$$\Rightarrow b_{w,p} = \frac{cov(\widehat{u_p}, \widehat{u_w})}{var(\widehat{u_p})}$$

True 
$$\Rightarrow b_{u,p} = \frac{cov(\widehat{u_p}, u)}{var(\widehat{u_p})}$$

 $h^2=0.50$ 

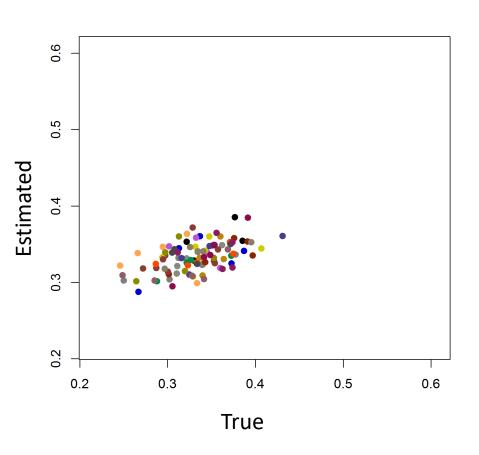


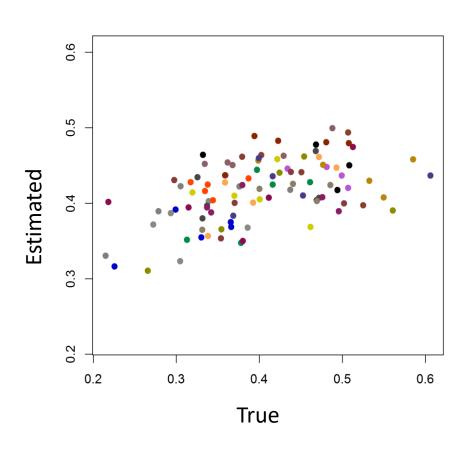


## Relative accuracy gain

$$h^2=0.50$$

$$h^2=0.05$$

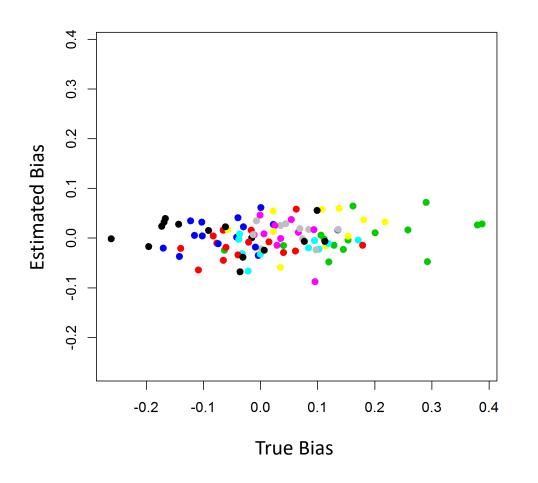




### Using the wrong evaluation model



#### Wrong evaluation fitting CG as fixed when they have a time trend



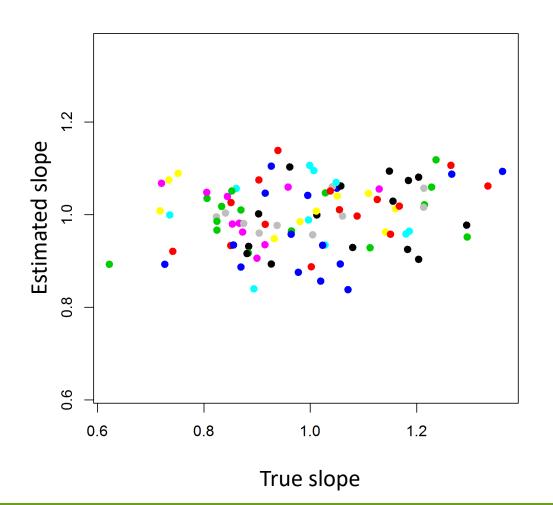
#### **BIAS**

$$Estimated \Rightarrow \mu_{w,p} = \overline{\widehat{u_p}} - \overline{\widehat{u_w}}$$

True 
$$\Rightarrow \mu_{u,p} = \overline{\widehat{u_p}} - \overline{u}$$

$$h^2=0.10$$

#### Wrong evaluation fitting CG as fixed when they have a time trend



## SLOPE b<sub>w,p</sub>

Estimated 
$$\Rightarrow b_{w,p} = \frac{cov(\widehat{u_p}, \widehat{u_w})}{var(\widehat{u_p})}$$

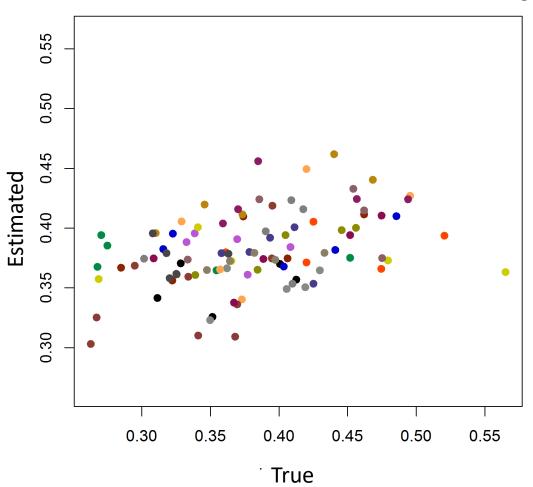
True 
$$\Rightarrow b_{u,p} = \frac{cov(\widehat{u_p}, u)}{var(\widehat{u_p})}$$

$$h^2=0.10$$



#### Wrong evaluation fitting CG as fixed when they have a time trend

#### Relative accuracy gain



Estimated 
$$\Rightarrow \rho_{p,w} = \frac{cov(\widehat{u_w}, \widehat{u_p})}{\sqrt{var(\widehat{u_w})var(\widehat{u_p})}}$$

$$\text{True} \Rightarrow \frac{acc_p}{acc_w}$$

$$h^2=0.10$$

#### **Conclusions**

 The proposed method LR estimates well bias, slope and accuracy when the model is in concordance with the reality.



But when the model has differences with the reality:

- With wrong model for contemporary groups
  - It is not possible to estimate bias or slope.
  - Accuracies can be estimated but not well
- With wrong heritabilities:
  - The bias could be under or over estimated
  - The slope is uninformative about the reality.
  - Accuracies can be estimated





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As well as to **Computing platform Bioinfo-Genotoul** for providing bioinformatics support.





#### Thank you for your attention!

Questions or comments?



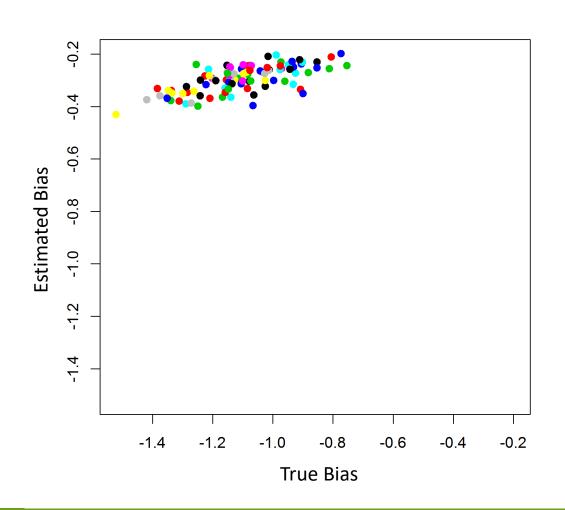








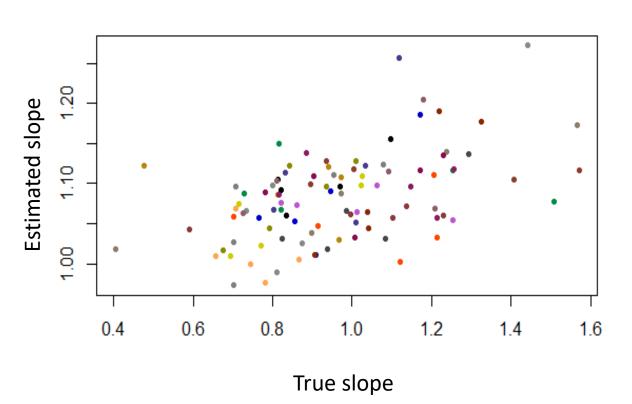
## Wrong h<sup>2</sup> in genetic evaluations simulated with h<sup>2</sup> 0.10 and evaluated with h<sup>2</sup> 0.05



#### **BIAS**

Estimated 
$$\Rightarrow \mu_{w,p} = \overline{\widehat{u_p}} - \overline{\widehat{u_w}}$$
  
 $True \Rightarrow \mu_{u,p} = \overline{\widehat{u_p}} - \overline{u}$ 

## Wrong h<sup>2</sup> in genetic evaluations simulated with h<sup>2</sup> 0.10 and evaluated with h<sup>2</sup> 0.05



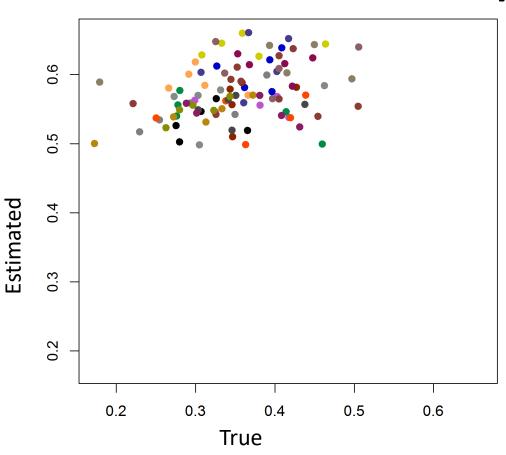
#### SLOPE b<sub>w,p</sub>

Estimated 
$$\Rightarrow b_{w,p} = \frac{cov(\widehat{u_p}, \widehat{u_w})}{var(\widehat{u_p})}$$

True 
$$\Rightarrow b_{u,p} = \frac{cov(\widehat{u_p}, u)}{var(\widehat{u_p})}$$

## Wrong h<sup>2</sup> in genetic evaluations simulated with h<sup>2</sup> 0.10 and evaluated with h<sup>2</sup> 0.05

#### Relative accuracy gain



Estimated 
$$\Rightarrow \rho_{p,w} = \frac{cov(\widehat{u_w}, \widehat{u_p})}{\sqrt{var(\widehat{u_w})var(\widehat{u_p})}}$$

True 
$$\Rightarrow \frac{acc_p}{acc_w}$$