# Single-step genomic selection: accommodating several key issues

Aidan McGarty<sup>1</sup> Brian Cullis<sup>1</sup>, Ahsan Asif<sup>2</sup> and Kristy Hobson<sup>2</sup> September 2, 2024

Mixed Models and Experiment Design Lab  $(MMaED)^1$ National Institute for Applied Statistics Research Australia University of Wollongong amcgarty@uow.edu.au

Chickpea Breeding Australia<sup>2</sup> NSW Department of Primary Industries | Agriculture



# **Overview**



• Genomic selection



- Genomic selection
- Motivating Example



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- Motivating Example
- Complexities





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	- These estimated relationships are then used to predict the performance of a set of MMaED | 20W test individuals























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- For the additive VE effects  $\mathbf{u}_a$ , covariance between varieties is also modelled via a known relationship matrix denoted G





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- The matrix formed via ancestral records is denoted  $\bf{A}$  and known as the numerator relationship matrix (NRM) [\(Oakey et al. \[2007\]](#page-79-1))





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- $\bullet$   $\psi$ <sub>a</sub> is a diagonal matrix with the site specific variances  $\psi$ <sub>a;</sub> on the diagonals



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- The final model fit included  $k_a = 2$  additive and  $k_b = 1$  non-additive factors



# Motivating example





# **Complexities**



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- Several strategies exist to overcome this complexity however the method we have used is a blending of the  $K$  and  $A$  relationship matrices





• Combining genomic information with pedigree information has been suggested by several authors [\(Legarra et al. \[2009\]](#page-79-0)[,Vitezica et al. \[2011\]](#page-80-0), [Meyer et al. \[2018\]](#page-79-1))



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- This will always result in an invertible genetic variance matrix even if the GRM **K** is non-invertible as a positive definite matrix (as is guaranteed when forming  $\mathbf{A}$ ) plus a positive semi definite matrix will be positive definite [\(Harville \[1997\]](#page-79-2))



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• The added parameter  $\lambda$  controls the weighting from the two sources of relatedness, with genomic information dominating when  $\lambda \approx 1$  and pedigree information dominating when  $\lambda \approx 0$ 



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- However this can be shown graphically through profiling on the parameter of interest  $(\lambda)$  and assessing the residual log-likelihood under a range of values for this parameter



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\tilde{\boldsymbol{\mathsf f}}_{a_2} = (\boldsymbol{\mathsf G}_{21}\boldsymbol{\mathsf G}_{11}^{-1}\otimes \boldsymbol{\mathsf I}_{k_a})\tilde{\boldsymbol{\mathsf f}}_{a_1}
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## **Conclusion**



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- However many complexities arise in implementation, a few of which have been explored here, specifically singularities in the GRM  $(K)$
- Combining both ancestral and genomic relationship matrices both overcomes this issue
- Finally DWReml (Butler, pers comm) can handle the high computational demand when fitting complex single-step factor models which include genetic relatedness such as the motivating example



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