

Pedigree *p*-rep Designs:

A class of designs for early stage variety trials

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Breeders Equation

$$RGG = \frac{i\sigma_g r}{t}$$

- ► RGG: Rate of Genetic Gain
- ► *i* : selection intensity
- σ_g^2 : genetic variance of trait (yield)
- ▶ r: accuracy of selection the correlation between the true and predicted genetic effects. Note that r^2 is the reliability (Mrode 1995)
- ► t : generation interval



Breeders Equation

$$RGG = \frac{i\sigma_g r}{t}$$

- ► For fixed i, σ_g^2 and t, an increase in the accuracy of EBLUPs increases the RGG
- ► This requires implementing procedures that are (and remain) best practice
- ▶ We can contribute statistically by
 - Optimal experimental design
 - Appropriate construction of a MET dataset (contemporary groups, co-located trials, etc.)
 - ► Sophisticated analysis of the dataset (Smith et. al. 2014)
 - Summary and dissemination of results (selection tools (Smith & Cullis 2018), etc.)

p-rep Designs



- ► Since Cullis et. al. (2006), adoption of *p*-rep designs in plant breeding and crop research has replaced traditional grid-plot designs
- ► The 25% *p*-rep threshold advocated by Cullis et. al. (2006) was based on the ratio of test plots to grid plots in traditional grid-plot designs
- ► Cullis et. al. (2006) stated "At present pedigrees are not generally used in routine analyses of EGVTs"
- ► From 2017 we incorporated pedigree data in MET analysis of PBA breeding programs



Pedigree *p*-rep designs

- ► Historically, the design of PBA early generation variety trials (EGVT) were generated in-house using DiGGer (Coombes 2002)
- ► From 2017 we transitioned to od (Butler & Cullis 2018) and exploited known genetic relationships for the allocation of varieties to both plots and sites
- ► This gives rise to two areas of research
 - ▶ Is the 25% level of *p*-rep necessary to achieve a given level of accuracy (*r*)
 - Is there an advantage to using pedigree information in the design of EGVT



Outline of the simulation study

- ► A simulation study was designed to primarily investigate
 - ► Various levels of *p*-rep
 - ► The advantage (if any) of including pedigree information in the design of the PBA Southern Faba Bean S1 trial that consisted of 256 breeding lines and 4 check varieties
- ► The random component of the linear mixed model for the analysis of a single PBA trial is:

 random ~ vm(Line, A) + ide(Line) + Block + Column + Row residual ~ ar1(Column) : ar1(Row)
- ▶ Where **A** is the (pedigree derived) relationship matrix
- ▶ Variance parameters: $\sigma_g^2 = \bar{a}\sigma_a^2 + \sigma_i^2$, σ_b^2 , σ_c^2 , σ_r^2 , ρ_c and ρ_r



Outline of the simulation study

- Simulation treatment factors
 - ▶ 7 levels of *p*-rep: p = 0, 5, 10, 15, 25, 50, 100%
 - ▶ 3 levels of proportion of additive genetic variance to the total: k = 0.5, 0.7, 0.9
 - ▶ 3 levels of null (baseline) reliability: $r_0^2 = 0.33, 0.5, 0.66$
 - ▶ 3 design types $d = od_{od}, od_{gg}, od_{\alpha}$
 - od_{od} od design with pedigree information included Butler and Cullis (2018)
 - ► od_{gg} DiGGer style row-column design Coombes (2002)
 - od_{α} augmented α design for single site Williams et. al. (2011)
- ▶ $7 \times 3 \times 3 = 63$ treatments $\{T\} \times 3$ designs



Outline of the simulation study

- ▶ For each $\{T_i\}$ generate n=4000 datasets
- Values of non-genetic variance components chosen from previous analyses of data
- $ightharpoonup \sigma_a^2$ and σ_i^2 chosen to realise pre-specified values of r_0^2 for a design with no replication and sub-optimal allocation of varieties to plots
- ▶ $T_{(id)_j}$ → the j^{th} simulation allocated to plots by design strategy d



Outline of the experiment designs

► With reference to the LMM used for analyses, the following indicates a) if a term was fitted and b) whether it was fitted as fixed or random for each of the three design types d

Term	od _{od}	od_{gg}	od_{lpha}
Line		F	F
Additive	R		
Non-Additive	R		
Block	R	R	F
Column	R	R	
Row	R	R	
Column:Row (Plot)	R	R	

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Simulations

- ▶ The simulated data from each $\{T_{(id)_j}\}$ were analysed in ASReml-R, Butler et. al. (2018).
- \blacktriangleright For T_i we have

$$\mathbf{u}_{g}, \mathbf{u}_{a}, \mathbf{u}_{i}$$

and for $T_{(id)}$

$$\tilde{u}_{\textit{gd}}, \tilde{u}_{\textit{ad}}, \tilde{u}_{\textit{id}}$$

For d in $\{od_{od}, od_{gg}, od_{\alpha}\}$, where each \mathbf{u} is of length 260.

- ► Summarise
 - ▶ Bias associated with variance parameter estimates for each design method d for the 63 combinations of p, k and r_0^2
 - Correlation between the "true" (simulated) value and corresponding predicted value for for each design method d and 63 combinations of p, k and r₀²

Correlation of total genetic effects k=0.7 and $r^2=0.3$ k=0.7, r^2=0.3 0.55 -0.50 -Design Alpha Digger 0.40 -25 100 50 Partial Replication (% p-rep)

Correlation of total genetic effects for each combination of k = 0.5, 0.7, 0.9 and r = 0.3, 0.5, 0.6 k=0.5, r^2=0.3 k=0.5, r^2=0.5 k=0.5, r^2=0.6 0.60 -0.75 -0.85 -0.55 -0.70 -0.80 -0.50 -0.65 -0.75 -0.45 -0.60 k=0.7, r^2=0.6 k=0.7, r^2=0.3 k=0.7, r^2=0.5 Correlation between true and predicted 0.70 -0.80 -Design Alpha 0.65 -Digger od od 0.75 -0.60 k=0.9, r^2=0.3 k=0.9, r^2=0.5 k=0.9, r^2=0.6 0.50 -0.800 -0.45 -0.65 -0.775 -0.40 -0.750 -0.60 -0.35 - 🧶 0.725 -0.30 0 25 50 75 100 Partial Replication (% p-rep)



Correlation between true and predicted: Total

- ► There is little to no distinction between the performance of od_{gg} and od_{α} allocations for any level of p, k, and r_0^2 for total correlations
- ► This suggests that spatial models (i.e. fitting AR1×AR1 to the residual) are not as critical in the design of trials such as these
- As k increases for fixed p and r_0^2 the advantage of od_{od} to od_{gg} and od_{α} designs increases
- ► For fixed r_0^2 and p, as k increases, the total correlation decreases as the bias associated with the non-additive genetic variance increases



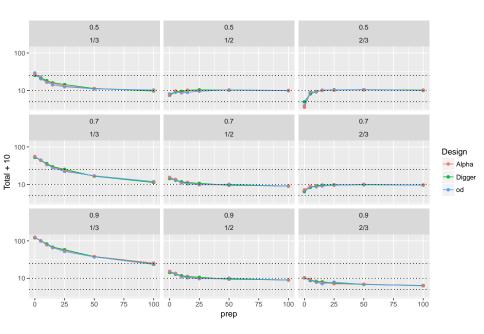
Correlation between true and predicted

- ► For k = 0.7 and $r_0^2 = 0.3$
- ▶ Using pedigree information, we can reduce the level of *p* and still maintain the same level of accuracy as other allocations in this situation.

Design	0	5	10	15	25	50	100
od_{lpha}	0.370	0.396	0.411	0.421	0.440	0.482	0.532
od_{gg}	0.369	0.391	0.408	0.418	0.440	0.481	0.534
$od_{lpha} \ od_{gg} \ od_{od}$	0.386	0.412	0.430	0.443	0.464	0.502	0.553



- ▶ There is still recovery of information for p=0?!
- \blacktriangleright BUT the bias of variance parameter estimation for low p is large
- ▶ Bias associated with σ_i^2 particularly large
- ▶ Bias associated with σ_g^2 significantly affected by the bias with σ_i^2 (if it is large) $\forall k$





- ► For each level of p and across all levels of k and r_0^2
 - od_{od} achieved the highest accuracy of prediction out of all three design strategies
 - od_{od} had the lowest design criterion (A-value) i.e. smallest average pairwise variance of variety contrasts
- ► For all combinations of p and r_0^2
 - od_{od} achieves a given level of accuracy with smaller p than od_{gg} and od_{α}
- ▶ od_{od} is the best design strategy for the true model
- ▶ It can be shown the **A** value is proportion to the expected level of accuracy of the predictions in the subsequent analysis



- ▶ Is the 25% level of *p*-rep the gold-standard for EGVTs?
- No... however, this is up to the discretion of the breeder taking into account
 - ► cost(\$)/plot i.e. how large can the trial be
 - desired level of accuracy
 - ▶ population diversity (σ_g^2) and proportion (k) of additive genetic variance
- Is there an advantage to using pedigree data in the design of EGVTs?
- Yes... results indicate for every level of p and all combinations of k and r_0^2
 - the accuracy of the predicted values and
 - relative response to selection (not presented today) are higher for allocations using od_{od} compared to od_{gg} and od_{α} designs (Cullis et. al. in prep.)





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