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Palm genomics and genetics Workshop

Factors controlling accuracy of genomic selection in oil palm (*Elaeis guineensis*)

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The oil palm

- world major oil crop
- one cultivated species, *Elaeis guineensis*
- **allogamous**
(monoecious with male and female cycles)
- vegetative multiplication difficult



Selection criteria:



Average bunch weight (ABW),
Bunch number (BN)

Fruits to bunch ratio (F/B),
Pulp to fruits ratio (P/F),
Oil to pulp ratio (O/P)

Height increment (INC)

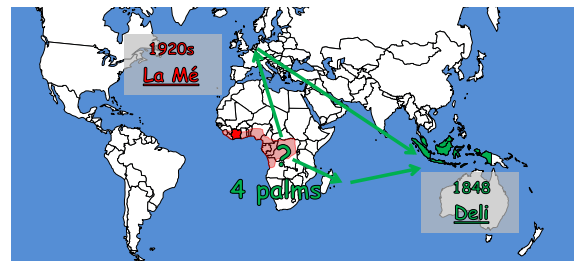
Bunch production (FFB)

Oil extraction rate (OER)

Breeding populations:

Distant populations with narrow genetic bases

- Deli
- La Mé



Genomic selection

Method of MAS (Meuwissen et al 2001):

- Training population phenotyped and genotyped
- Dense genotyping of the whole genome
- All markers effects estimated simultaneously
- No test of significance of marker effects
- Selection on markers alone (GEBV) in test population

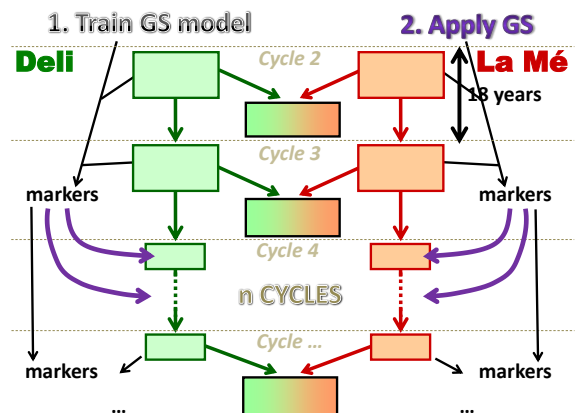
“Selection on genetic values predicted from markers could substantially increase the rate of genetic gain in animals and plants”

Genetics 137: 1820-1829 (April 2001)

Prediction of Total Genetic Value Using Genome-Wide Dense Marker Maps

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Could we increase the rate of genetic gain in oil palm breeding with genomic selection ?

Hypothesis: Progeny-tested individuals could be used to train a GS model that could be applied to predict breeding values of individuals of the same populations

→ Will be checked by measuring the **accuracy of GS in a cross-validation study with real data**

$$\text{Genetic gain per year} = \frac{\text{Intensity} * \text{Accuracy} * \sigma_a}{\text{Generation interval}}$$

With accuracy = $r(\text{TBV}, \text{EBV})$ and current accuracy ~ 0.8

Hypothesis: Progeny-tested individuals could be used to train a GS model that could be applied to predict breeding values of individuals of the same populations

With:

- Narrow genetic base / Low effective size
- Small training populations
- Small number of markers
- Multiallelic markers

Materials and methods

Plant material:

- **Deli:** 131 individuals
- **La Mé:** 93 individuals



Materials and methods

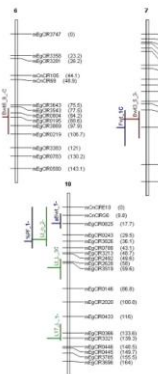
Molecular data:

235 SSR (Billotte et al 2005; Tranbarger et al 2011)

$\sim 1 \text{ SSR} / 7.4 \text{ cM}$

Phenotypic data:

1. Progeny tests, **10 quantitative traits**
2. **Estimated breeding values (BLUP)**
3. **Deregressed** and used in a **weighted analysis** to derive genomic estimated breeding values (Garrick et al 2009)



Materials and methods

5-fold cross-validation:

- 1/ Individuals (genotyped and phenotyped) divided into 5 groups to make training population (4 groups) and test population (5th group)
→ 5 replicates
- 2/ Estimation of allelic effects
- 3/ Calculation of GEBV for test individuals
- 4/ Calculation of accuracy of genomic selection in test population

Materials and methods

Definition of groups for training and test populations:

2 methods, in order to get a range of accuracy of GS:

1. Lower bound: **CLUSTERING** (Saatchi et al 2011)

- Calculate matrix of additive genetic relationships between individuals,
- Use K-means clustering to make 5 groups

→ **Increases within-group relationships / Decreases between-group relationships** (groups represent subpopulations)

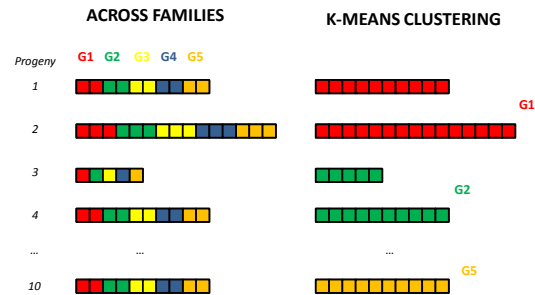
2. Upper bound: **ACROSS FAMILIES**

- Each family is randomly divided into 5 groups

→ **Maximizes relationships between training and test populations**

Materials and methods

Definition of groups for training and test populations:



Materials and methods

Statistical methods to estimate GEBV:

| Group of method | Method | Marker effects | Comments | Reference |
|-----------------|--------|----------------|-------------------------------------|--|
| Mixed Model | ABLUP | no | Control | Henderson 1975 |
| Mixed Model | BLUP | yes | $gi \sim N(0, Vm)$ | Meuwissen et al 2001 |
| Mixed Model | GBLUP | no | Estimate GEBV | Henderson 1975, Eding and Meuwissen 2001 |
| Bayes | BRR | yes | $gi \sim N(0, \sigma^2_{aa})$ | Perez et al 2010 |
| Bayes | BL | yes | $gi \sim N(0, \tau_i * \sigma^2_e)$ | Perez et al 2010 |
| Semi-parametric | RKHS | no | Estimate GEBV | Gianola et al 2006, Heslot et al 2012 |

→ Some methods better suited for traits with many small effect genes, others for traits with major genes + small effect genes

Accuracy:
$$r_{GEBV, TBV} = \frac{\hat{\sigma}_{DEBV, GEBV}}{\sqrt{\sigma_a^2 + \sigma_{GEBV}^2}} \quad (\text{Saatchi et al 2011})$$

Results

➤ Factors with the strongest effect on accuracy:

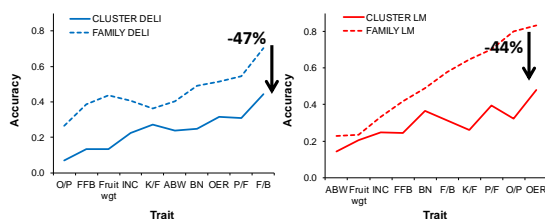
- (1) TRAINING MODE,
- (2) TRAIT, POPULATION
- (3) TRAIT * POPULATION INTERACTION

➤ No effects of statistical method, no statistical method * trait interaction

Results

Effect of training mode on accuracy:

→ Range of accuracy for GS



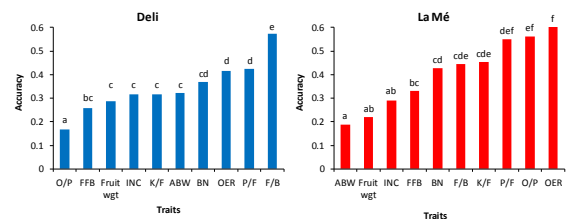
... Effect of training mode related to a_{max} :

family → cluster, -22% a_{max} family → cluster, -18%
1.1 0.86 0.93 0.77

Results

Effect of trait on accuracy:

Accuracy varies with a factor 3 according to trait (very low to very high)

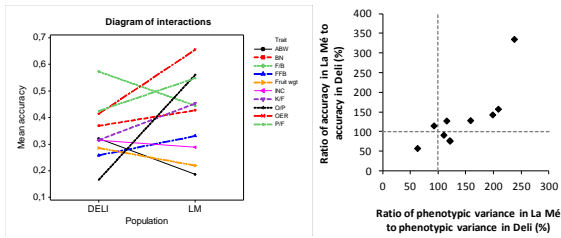


Due to genetic architecture of each trait ?

(number of QTLs, distribution of QTL effects, distribution of QTL along genome versus distribution of SSRs, LD between markers and QTLs)

Results

Effect of trait * population interaction on accuracy:



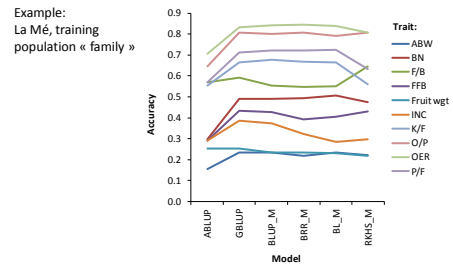
Due to differences in genetic architecture between populations and traits ?

Related to differences in phenotypic variance

Results

Effect of statistical method on accuracy:

No effect, no interaction with trait



... contradictory with trait effect and trait * population interaction
→ Too small number of phenotypic records ?

Conclusions

Some traits with very low accuracy

→ bigger training populations / more markers

More studies required before implementing GS in our oil palm breeding program...

- Effect of increasing training population size ?
- Rate of decrease of accuracy over generations ?
- Accuracy between experimental designs ?
- Genetic architecture of traits in each population ?

...

Some answers in 2013 (simulations)

and 2014 (more real data: 2 experimental designs, 2 generations + GBS genotyping)

