### Genomic Selection

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#### Introduction

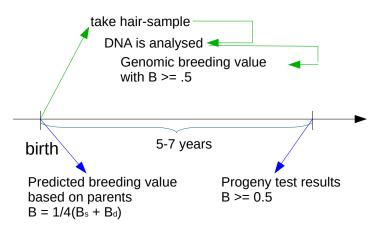
- Proposed in 2001
- ▶ Widely adopted in 2007/2008
- Costs of breeding program reduced due to shorter generation intervals
- In cattle: young sire selection versus selection based on sire proofs
- ▶ In pigs: early selection among full sibbs
- Inbreeding must be considered

## Terminology

- ► **Genomic Selection**: use of genomic Information for selection decisions
- Genomic Information is used to predict genomic breeding values

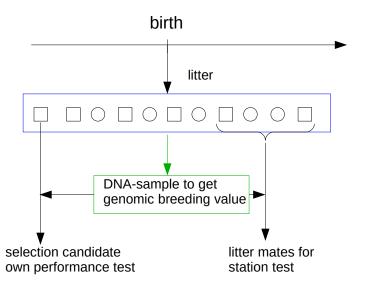
#### Benefits in Cattle

### With genomic information



Without genomic information

## Benefits in Pigs

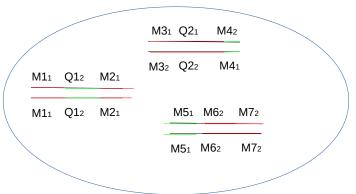


#### Genetic Model

- Recall: BLUP animal model is based on infinitesimal model
- Prediction of genomic breeding values is based on polygenic model
- In polygenic model: Single Nucleotide Polymorphisms (SNP) are used as markers
- Marker genotypes are expected to be associated with genotypes of Quantitative Trait Loci (QTL)

## Polygenic Model

### Distribution of SNP (M) and QTL (Q)



Non-Coding region

Coding region

#### Statistical Models

Two types of models are used

- 1. marker-effect models (MEM)
- 2. genomic-breeding-value based models (BVM)

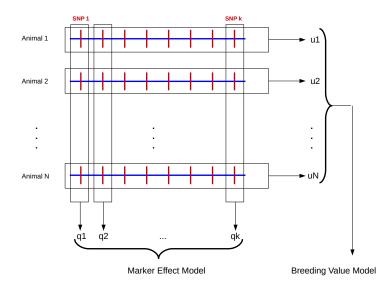
#### **MEM**

- marker effects (a-values) are fitted using
  - ightharpoonup a simple linear model ightharpoonup marker effects are fixed
  - ightharpoonup a linear mixed effects model ightharpoonup marker effects are random
- Problem of finding which markers are associated to QTL
- ► With high number of SNP compared to number of genotyped animals: very large systems of equations to solve

### **BVM**

- genomic breeding values as random effects
- similar to animal model
- genomic relationship matrix (G) instead of numerator relationship matrix (A)

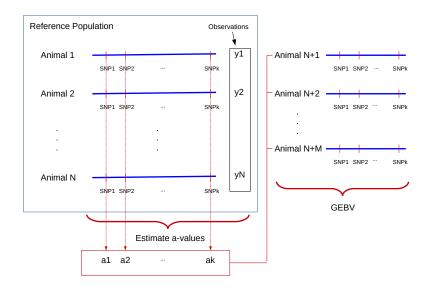
#### MEM versus BVM



### Logistic Procedures

- ► Two Step:
  - use reference population to get marker effects using MEM
  - use marker effects to get to genomic breeding values
- Single Step
  - MEM or BVM in a single evaluation
  - difficulty how to combine animals with and without genotypes

### Two Step Procedure



## Single Step GBLUP

- Use a mixed linear effect model
- ► Genomic breeding values *g* are random effects

$$y = Xb + Zg + e$$

with

- ► E(e) = 0,  $var(e) = I * \sigma_e^2$
- E(g) = 0,  $var(g) = G * \sigma_g^2$
- Genomic relationship matrix G

## Solution Via Mixed Model Equations

► All animals have genotypes and observations

$$\left[\begin{array}{cc} X^TX & X^TZ \\ Z^TX & Z^TZ + \lambda * G^{-1} \end{array}\right] \left[\begin{array}{c} \hat{b} \\ \hat{g} \end{array}\right] = \left[\begin{array}{c} X^Ty \\ Z^Ty \end{array}\right]$$

with  $\lambda = \sigma_e^2/\sigma_g^2$ .

#### Animals Without Observations

- Young animals do not have observations
- Partition  $\hat{g}$  into
  - $ightharpoonup \hat{g}_1$  animals with observations and
  - $ightharpoonup \hat{g}_2$  animals without observations
- lacktriangle Resulting Mixed Model Equations are (assume  $\lambda=1$ )

$$\begin{bmatrix} X^T X & X^T Z & 0 \\ Z^T X & Z^T Z + G^{(11)} & G^{(12)} \\ 0 & G^{(21)} & G^{(22)} \end{bmatrix} \begin{bmatrix} \hat{b} \\ \hat{g}_1 \\ \hat{g}_2 \end{bmatrix} = \begin{bmatrix} X^T y \\ Z^T y \\ 0 \end{bmatrix}$$

Predicted Genomic Breeding Values

Last line of Mixed model equations

$$G^{(21)} \cdot \hat{g}_1 + G^{(22)} \cdot \hat{g}_2 = 0$$

### Solutions

► Solving for  $\hat{g}_2$ 

$$\hat{g}_2 = -(G^{(22)})^{-1} \cdot G^{(21)} \cdot \hat{g}_1$$

### Genomic Relationship Matrix

- ▶ Breeding value model uses genomic breeding values *g* as random effects
- ▶ Variance-covariance matrix of g are proposed to be proportional to matrix G

$$var(g) = G * \sigma_g^2$$

where G is called **genomic relationship matrix** (GRM)

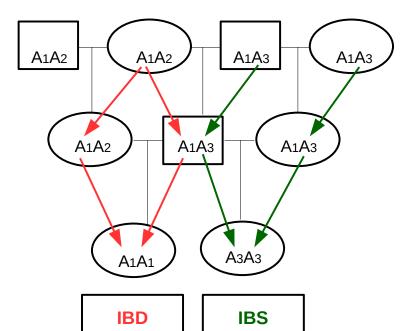
### Properties of *G*

- $\triangleright$  genomic breeding values g are linear combinations of q
- ightharpoonup g as deviations, that means E(g) = 0
- ightharpoonup var(g) as product between G and  $\sigma_g^2$  where G is the genomic relationship matrix
- G should be similar to A

## Change of Identity Concept

- ► A is based on identity by descent
- ► *G* is based on identity by state (including ibd), assuming that the same allele has the same effect
- ► IBS can only be observed with SNP-genotype data

## Identity



#### Linear Combination

- ► SNP marker effects (a values) from marker effect model are in vector q
- Genomic breeding values from breeding value model are determined by

$$g = U \cdot q$$

Matrix U is determined by desired properties of g

#### Deviation

 Genomic breeding values are defined as deviation from a certain basis

$$\rightarrow E(g) = 0$$

▶ How to determine matrix U such that E(g) = 0

### Equivalence Between Models

Decomposition of phenotypic observation  $y_i$  with

► Marker effect model

$$y_i = w_i^T \cdot q + e_i$$

Breeding value model

$$y_i = g_i + e_i$$

 $\triangleright$   $g_i$  and  $w_i^T \cdot q$  represent the same genetic effects and should be equivalent in terms of variability

## **Expected Values**

- ightharpoonup Required:  $E(g_i) = 0$
- ► Take *q* as constant SNP effects
- Assume w<sub>i</sub> to be the random variable with:

$$w_i = \left\{ egin{array}{ll} 1 & ext{with probability} & p^2 \ 0 & ext{with probability} & 2p(1-p) \ -1 & ext{with probability} & (1-p)^2 \end{array} 
ight.$$

 $\rightarrow E(w_i)$ : For a single locus

$$E(w_i) = 1*p^2 + 0*2p(1-p) + (-1)(1-p)^2 = p^2 - 1 + 2p - p^2 = 2p - 1 \neq 0$$

# Specification of g

Set

$$g_i = (w_i^T - s_i^T) \cdot q$$

with  $s_i = E(w_i) = 2p - 1$ 

Resulting in

$$g = U \cdot q = (W - S) \cdot q$$

with matrix S having columns j with all elements equal to  $2p_j-1$  where  $p_j$  is the allele frequency of the SNP allele associated with the positive effect.

#### Genetic Variance

- ► Requirement:  $var(g) = G * \sigma_g^2$
- ▶ Result from Gianola et al. (2009):

$$\sigma_g^2 = \sigma_q^2 * \sum_{i=1}^k (1 - 2p_j(1 - p_j))$$

From earlier:  $g = U \cdot q$ 

$$var(g) = var(U \cdot q) = U \cdot var(q) \cdot U^{\mathsf{T}} = UU^{\mathsf{T}} \sigma_q^2$$

Combining

$$var(g) = UU^{T}\sigma_{q}^{2} = G * \sigma_{q}^{2} * \sum_{i=1}^{k} (1 - 2p_{i}(1 - p_{i}))$$

## Genomic Relationship Matrix

$$G = \frac{UU^{T}}{\sum_{j=1}^{k} (1 - 2p_{j}(1 - p_{j}))}$$

### How To Compute *G*

- Read matrix W
- For each column j of W compute frequency  $p_j$
- ▶ Compute matrix S and  $\sum_{j=1}^{k} (1 2p_j(1 p_j))$  from  $p_j$
- Compute U from W and S
- Compute G