

# Whole-population relationship matrix including pedigree and markers for genomic selection

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**Rational: use of DYD's or deregressed proofs for genomic selection is problematic**

Loss of information

- Bias (selection)
- Tricky covariances and weights, in particular for females (YD's)
- Difficult for many species
- Family information ignored

**OBJECTIVE: build up complete relationship matrix**

Genomic relationship matrix

- As good as BayesB from experience

If the relationship matrix includes all animals

- Raw records can be used for evaluation
- All animals benefit from increased accuracies from genomic selection
- Avoid selection bias

## Conclusion

- Joint relationship matrix **H** gives a unified framework for including all data, pedigree and genotypes
- More elegant, simple and possibly accurate
- Useful for populations where DYDs are inaccurate (sheep, swine...)

## Joint relationships

Let  $\mathbf{u}_1$  be ungenotyped animals and  $\mathbf{u}_2$  be genotyped animals. Let **A** be the numerator relationship matrix.

Let **G** be the *genomic relationship matrix* based on SNP's (VanRaden 2008, Habier et al. 2007)

- **G** contains information on relationships among ancestors and descendants of genotyped animals
- Need to account for **G** in Var ( $\mathbf{u}_1$ ) and in Cov ( $\mathbf{u}_1, \mathbf{u}_2$ )

Genomic information  $\mathbf{u}_2 \sim N(0, \mathbf{G})$

Now, condition ungenotyped animals in genotyped animals by the selection index:

Selection index

$$p(\mathbf{u}_1 | \mathbf{u}_2) = N(\mathbf{A}_{12}\mathbf{A}_{22}^{-1}\mathbf{u}_2, \mathbf{A}_{11} - \mathbf{A}_{12}\mathbf{A}_{22}^{-1}\mathbf{A}_{21})$$

$$\text{Var}(\mathbf{u}_1 | \mathbf{G}) = \mathbf{A}_{12}\mathbf{A}_{22}^{-1}\mathbf{G}\mathbf{A}_{22}^{-1}\mathbf{A}_{21} + \mathbf{A}_{11} - \mathbf{A}_{12}\mathbf{A}_{22}^{-1}\mathbf{A}_{21} = \mathbf{A}_{11} + \mathbf{A}_{12}\mathbf{A}_{22}^{-1}(\mathbf{G} - \mathbf{A}_{22})\mathbf{A}_{22}^{-1}\mathbf{A}_{21}$$

With this, we can create a joint distribution

Joint distribution **H**  $\begin{pmatrix} \mathbf{u}_1 \\ \mathbf{u}_2 \end{pmatrix} \sim N(0, \mathbf{H}), \mathbf{H} = \begin{bmatrix} \mathbf{H}_{11} & \mathbf{H}_{12} \\ \mathbf{H}_{21} & \mathbf{H}_{22} \end{bmatrix}$

$$\mathbf{H} = \mathbf{A} + \begin{bmatrix} \mathbf{A}_{12}\mathbf{A}_{22}^{-1}(\mathbf{G} - \mathbf{A}_{22})\mathbf{A}_{22}^{-1}\mathbf{A}_{21} & \mathbf{A}_{12}\mathbf{A}_{22}^{-1}(\mathbf{G} - \mathbf{A}_{22}) \\ (\mathbf{G} - \mathbf{A}_{22})\mathbf{A}_{22}^{-1}\mathbf{A}_{21} & \mathbf{G} - \mathbf{A}_{22} \end{bmatrix}$$

Inverse of **H**  $\mathbf{H}^{-1} = \mathbf{A}^{-1} + \begin{bmatrix} 0 & 0 \\ 0 & \mathbf{G}^{-1} - \mathbf{A}_{22}^{-1} \end{bmatrix}$

## Notes:

- **H** = classical relationships + a difference depending on the difference between « numerator » ( $\mathbf{A}_{22}$ ) and « genomic » (**G**) relationship matrices.

- The selection index transmits genomic information in **G** backwards and forwards to all individuals in the pedigree to form **H**

- **H** can be used in genetic evaluations, even of very large size (millions). See Ignacy Misztal (this meeting).

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