

Accuracy of genomic selection predictions for hip height in Brahman cattle using different relationship matrices

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Abstract – The objective of this work was to evaluate the effects of genomic information on the genetic evaluation of hip height in Brahman cattle using different matrices built from genomic and pedigree data. Hip height measurements from 1,695 animals, genotyped with high-density SNP chip or imputed from 50 K high-density SNP chip, were used. The numerator relationship matrix (NRM) was compared with the H matrix, which incorporated the NRM and genomic relationship (G) matrix simultaneously. The genotypes were used to estimate three versions of G: observed allele frequency (H_{GOF}), average minor allele frequency (H_{GMF}), and frequency of 0.5 for all markers (H_{G50}). For matrix comparisons, animal data were either used in full or divided into calibration (80% older animals) and validation (20% younger animals) datasets. The accuracy values for the NRM, H_{GOF} , and H_{G50} were 0.776, 0.813, and 0.594, respectively. The NRM and H_{GOF} showed similar minor variances for diagonal and off-diagonal elements, as well as for estimated breeding values. The use of genomic information resulted in relationship estimates similar to those obtained based on pedigree; however, H_{GOF} is the best option for estimating the genomic relationship matrix and results in a higher prediction accuracy. The ranking of the top 20% animals was very similar for all matrices, but the ranking within them varies depending on the method used.

Index terms: *Bos indicus*, beef cattle, genomics, rare alleles.

Acurácia da predição genômica para altura do quadril em bovinos Brahman com uso de diferentes matrizes de parentesco

Resumo – O objetivo deste trabalho foi avaliar os efeitos da informação genômica na avaliação genética para altura do quadril em bovinos da raça Brahman, por meio de diferentes matrizes construídas com dados genômicos e de pedigree. Utilizaram-se medidas de altura do quadril de 1.695 animais, genotipados com SNP chip de alta densidade ou imutados do 50 K SNP chip de alta densidade. A matriz de pedigree “numerator relationship matrix” (NRM) foi comparada à matriz H, a qual incorporou as matrizes NRM e de parentesco genômico (G) simultaneamente. Os genótipos foram utilizados para estimar três versões de G: frequência observada dos alelos (H_{GOF}), média da menor frequência alélica (H_{GMF}) e frequência de 0,5 para todos os marcadores (H_{G50}). Para a comparação das matrizes, foram utilizadas informações completas ou divididas em conjuntos de dados de calibração (80% dos animais mais velhos) e de validação (20% dos mais jovens). Os valores de acurácia para NRM, H_{GOF} e H_{G50} foram 0,776, 0,813 e 0,594, respectivamente. NRM e H_{GOF} foram semelhantes, com menores variâncias para os elementos da diagonal e fora da diagonal, bem como para os valores genéticos estimados. O uso de informações genômicas resultou em estimativas de parentesco semelhantes às obtidas com base em pedigree; entretanto, H_{GOF} é a melhor opção para estimar a matriz de parentesco genômico e resulta em maiores acurácias de predição. O ranking dos animais top 20% foi muito semelhante para as matrizes, mas a classificação dentro destas varia dependendo do método.

Termos para indexação: *Bos indicus*, gado de corte, genômica, alelos raros.



Introduction

Traditionally, animal selection studies target traits of interest and use the phenotype of individuals and information of kinship derived from pedigree records, which is the basis for building the numerator relationship matrix (NRM) (Henderson, 1975). This animal breeding and selection method is efficient, but the process can be slow, especially for traits measured only in one sex, such as milk production; measured after the slaughter of animals, including meat quality; or measured late in life, as, for example, longevity (Goddard & Hayes, 2009) and hip height. In breeding programs focused on economically important traits, researchers seek to identify genes, genetic markers associated to these traits, or methodologies to enhance or accelerate genomic selection. The application of a single-step genome-wide association approach in Brahman cattle has allowed identifying multiple regions scattered across the genome that influence weights at different ages. The most interesting regions found were connected to previously identified quantitative trait loci (QTLs) and to genes influencing growth and weight traits (Martínez et al., 2017).

The advancement of technology and the opportunity of genotyping a high number of individuals for numerous alleles make it possible to use information on alleles that can be shared through common ancestors in the pedigree, including ancestors that may be missing from the pedigree or not genotyped. This makes the use of a genomic relationship matrix (G) feasible (VanRaden, 2008), allowing to increase the accuracy of predicted breeding values in genetic evaluations. Genomic selection using the G matrix can increase the rate of genetic improvement and reduce the cost of progeny testing. This has been observed in the dairy system, although some practical barriers are still faced in its implementation in the beef sector, primarily due to the lack of short-term return on investment for most producers and to incremental and relatively small genetic gains (Rolf et al., 2014).

Breeding values are obtained, traditionally, using mixed model equations that adopt the NRM based on pedigree information. In one form of genomic selection, the NRM or the G matrix represent the additive genetic matrix. However, in most cases, the G matrix includes genomic information of fewer animals. Therefore, Legarra et al. (2009) and Misztal et al. (2009) proposed a method that integrates the

NRM and G matrix in a single H matrix, enabling genetic evaluation based on the best linear unbiased prediction (Blup), which was successfully applied to dairy cattle (Aguilar et al., 2010). Forni et al. (2011) used different strategies to create the G matrix and subsequently integrate it with the NRM by varying the allele frequencies of a population of pigs; the authors concluded that building the G matrix this way did not affect the estimated breeding values and variance components of the studied population. Brito et al. (2017) also evaluated alternative genomic relationship matrices, validation designs, and genomic prediction scenarios in a sheep program, and found that the G matrix, using the observed allele frequency, presented the highest average accuracies and is recommended for genomic predictions. These results show the importance of assessing the contribution of genomic information in genetic evaluation processes in different species and different population structures.

The objective of this work was to evaluate the effects of genomic information on the genetic evaluation of hip height in Brahman cattle using different matrices built from genomic and pedigree data.

Materials and Methods

The investigated population of Brahman cattle has 90% *Bos indicus* genetics (Bolormaa et al., 2013b). Hip height (HH, cm) measurements taken from 1,695 Brahman animals, between 15 and 18 months of age, were used in the present study. These cattle represent a subset of the extensively phenotyped population bred by the Cooperative Research Centre for Beef Genetic Technologies (Beef CRC, Armidale, Australia) and described previously in detail (Barwick et al., 2009; Fortes et al., 2011; Hawken et al., 2012). All individuals in this population have genotype information for 777,000 single nucleotide polymorphisms (SNPs), and these high-density SNP data were genotyped or imputed.

All these animals were genotyped using three different SNP chips (Illumina, Inc., San Diego, CA, USA): the BovineSNP50 beadchip, version 1, to genotype females; version 2, to genotype males, which, combined, are the 1,695 phenotyped animals; and the high-density SNP chip to genotype 917 samples from sires and selected representative animals of the Beef CRC populations, which were genotyped with the high-density SNP chip to allow for genotype

imputation using the Beagle program (Browning & Browning, 2011) with average of imputation accuracy of 0.90. For further detail on genotyping, imputation, and quality control see Bolormaa et al. (2013a).

In the quality control analysis, the SNP was excluded if the minor allele frequency was lower than 0.05 or the correlation between the SNPs was higher than 0.95. After quality control procedures, 569,620 SNPs remained to estimate the genomic relationship coefficients in the G matrices.

The pedigree information used to build the NRM was composed of 3,030 animals, including the genotyped ones, corresponding to 55.94% of the total population. Estimated breeding values for HH were calculated using the animal model with the fixed effects of sex, cohort (interaction between year of birth and farm), and age at HH measurement fitted as a covariate. To obtain the estimated breeding values, the NRM used a traditional method based on pedigree information.

The combined pedigree-genomic relationship matrix, denominated H, was calculated using both pedigree and genomic information according to Aguilar et al. (2010). The G matrix was obtained using the method of VanRaden (2008):

$$G = (M - P)(M - P)' / 2 \sum_{j=1}^m p_j (1 - p_j),$$

where M is the matrix that specifies which marker alleles each individual inherited with m columns (m is the total number of markers) and n rows (n is the total number of genotyped individuals); and P is the matrix with the frequency of the second allele (p_j), expressed as $2p_j$. The M matrix was filled by 0 if first homozygous, 1 if heterozygous, or 2 if second homozygous. The frequencies used to obtain P were those described by Forni et al. (2011): observed allele frequency of each SNP (GOF), average minor allele frequency (GMF), and frequency of 0.5 for all markers (G50). To avoid problems with inversion in the mixed model equations, the method proposed by VanRaden (2008), with weighting factor of 0.95 between the G matrix and the NRM, was used (Aguilar et al., 2010).

After obtaining the weighted G matrix (Gw), the inverse of H was calculated using the following equation (Aguilar et al., 2010; Christensen & Lund, 2010):

$$H^{-1} = NRM^{-1} + \begin{bmatrix} 0 & 0 \\ 0 & Gw^{-1} - NRM_{22}^{-1} \end{bmatrix},$$

where H^{-1} is the inverse of the pedigree-genomic relationship matrix; NRM^{-1} is the inverse of the pedigree-based numerator relationship matrix; G_w^{-1} is the inverse of the genomic matrix; and NRM_{22}^{-1} is the inverse of the pedigree-based numerator relationship matrix of the genotyped individuals. Considering the variations in the allele frequencies used to build the G matrices, three versions of the H matrix were also built: H_{GOF} , H_{GMF} , and H_{G50} .

To compare the accuracies of the genomic estimated breeding values (GEBVs) obtained with each H matrix, the mean accuracy was estimated using the prediction error variance (PEV), calculated by:

$$r_i = \sqrt{1 - \frac{PEV_j}{\sigma_{a_i}^2}}$$

where r_i is the accuracy of the mean additive value for each matrix i; $\sigma_{a_i}^2$ is the additive variance estimated for each matrix i; and PEV_j is the prediction error variance for each animal j estimated by matrix i.

To obtain the inversions of these matrices, as well as the estimates of the variance components, genetic parameters, and the PEV, the restricted maximum likelihood (REML) methods in the Wombat software were used (Meyer, 2007). The mean accuracies of the GEBVs based on 1,695 GEBVs were calculated using phenotypes of all the animals genotyped for the prediction (GEN) and 80% of the phenotypic information, i.e., the subset of data corresponding to the oldest animals (old) in the dataset.

To compare the accuracy of prediction, the old subset ($n = 1,356$ animals) was used to predict the GEBVs of 20% of the youngest animals (young subset, $n = 339$ animals); this accuracy was estimated by omitting the phenotypes of the younger animals from the prediction. This way, an alternative accuracy metric, denominated prediction accuracy (r), was calculated, based on the correlations between the adjusted phenotype ($Phen_{adj}$) and the GEBVs, as follows:

$$r = \text{cov}(Phen_{adj}, GEBV_j) / \sqrt{h_i^2},$$

where h_i^2 is the heritability estimated for HH for each matrix i (H_{GOF} , H_{GMF} , and H_{G50}). The correlation between the GEBVs estimated with and without including the phenotypes of young animals in the prediction was also determined.

The three versions of the H matrix were compared considering the ranking of the animals based on the estimated GEBVs. To compare the rankings, animals that had higher GEBVs for HH, i.e., top 20% of the population (top 20%, $n = 339$), were investigated. The Spearman rank coefficient (ρ) was used to compare the top 20%, defined as the Pearson correlation coefficient between ranked variables (Yitzhaki & Schechtman, 2013), using the alternative formula proposed by Conover (1999):

$$\rho = 1 - \frac{6 \sum d_i^2}{n(n^2 - 1)},$$

where d_i^2 is the difference between the ranks of each observation for two variables and n is the number of observations. The standard Pearson correlation between rankings of animals in different matrices was also estimated.

Results and Discussion

Minor variances and both diagonal and off-diagonal elements were obtained for H_{GOF} , H_{GMF} , H_{G50} , and the NRM (Table 1). For the diagonal elements, the NRM showed lower variance, probably because it is incomplete and the inbreeding value of the studied population is very low, indicating that there is a weak relationship between the evaluated families. Furthermore, the NRM calculates the probability of kinship, decreasing the variances of elements.

Table 1. Statistics of the relationship coefficients estimated using the full pedigree and genomic dataset of the genotyped Brahman cattle⁽¹⁾.

Matrix	Mean	Minimum	Maximum	Variance
Diagonal elements				
NRM	1.0003	1.0000	1.1250	3.7×10^{-5}
H_{GOF}	1.0281	0.8971	1.2588	3.4×10^{-3}
H_{GMF}	2.8420	2.5718	3.0816	3.6×10^{-3}
H_{G50}	1.3576	1.1979	1.5244	1.54×10^{-3}
Off-diagonal elements				
NRM	0.0086	0.0000	0.6250	1.4×10^{-3}
H_{GOF}	-0.0006	-0.1062	0.6614	1.9×10^{-3}
H_{GMF}	1.9121	1.5498	2.5818	5.7×10^{-3}
H_{G50}	0.6776	0.4453	1.1599	2.6×10^{-3}

⁽¹⁾NRM, pedigree-based numerator relationship matrix; H_{GOF} , genomic relationship matrix with observed allele frequency; H_{GMF} , genomic relationship matrix with average minor allele frequency; and H_{G50} , genomic relationship matrix with frequency of 0.5 for all markers.

However, when genomic information was used, these families did share common alleles and the estimated relationship coefficients were different. These results were expected because genomic relationships reflect a relationship that represents the actual gene fraction shared between individuals (Choi et al., 2017). The H_{GMF} and H_{G50} matrices used the same allele frequency for all markers: 0.27 average minor allele frequency or 0.50, respectively. Observed allele frequencies were distant from 0.5 for many markers (Figure 1), which may be an effect of the SNP chip development, based mostly on *Bos taurus* and not on *B. indicus* data (Gibbs et al., 2009). These frequency distributions, however, were not reported by Choi et al. (2017), who also studied *B. taurus* cattle.

The data used to compare variance components were either the full phenotype dataset of genotyped animals (GEN, $n=1,695$) or a subset that included 80% of the oldest animals (old, $n=1,356$). In both GEN and old datasets, the variance components were similar when the matrices estimated with the same methodology were compared (i.e., the NRM of GEN was similar to that of the old dataset) (Table 2). However, when matrices estimated with different methodologies were compared, the variance components were different.

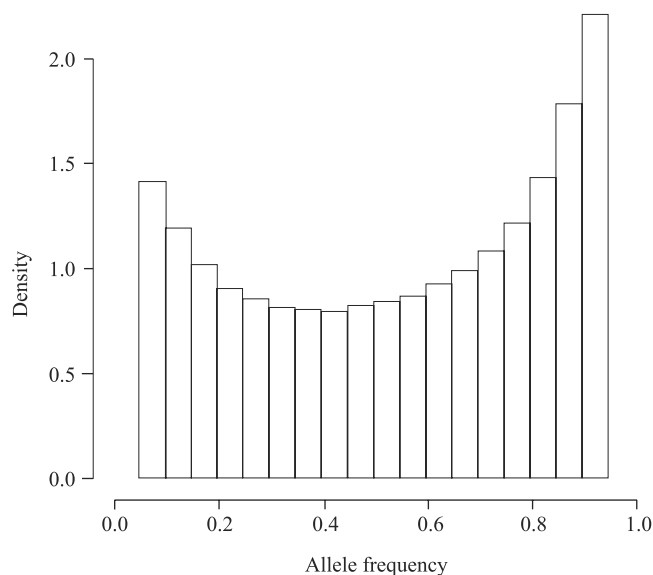


Figure 1. Distribution of observed frequencies for the second allele of the genotyped Brahman cattle.

These differences between matrices are in contrast with the data presented by Forni et al. (2011), who detected that the additive variance was higher when the difference between the average diagonal and off-diagonal elements of the matrix was lower. In the present study, the differences in the diagonal and off-diagonal elements estimated with the NRM, H_{GOF} , and H_{GMF} were not significant (0.99, 1.03, and 0.93 respectively), but the additive variances differed. Forni et al. (2011) found this relationship was true only for H_{G50} , which, in the present work, showed a difference of 0.68 between coefficients.

The variance components obtained using H_{GOF} and the NRM were quite similar in the present study, which is consistent with the findings of Riley et al. (2007). Variance components in H_{GMF} and H_{G50} were less similar to those in the NRM, compared with H_{GOF} , and may have been inflated with the use of fixed allele frequencies. Several researchers related problems

with inflated estimates of variance components due to false kinship coefficients (Aguilar et al., 2010; Forni et al., 2011); in this case, in H_{GMF} and H_{G50} matrices that showed higher values than the NRM or H_{GOF} .

When observed allele frequencies are distant from 0.5, “rare” alleles have greater influence on the relationship estimated, and this may be the underlying reason why H_{G50} and H_{GMF} were approximated to each other and distanced from the NRM and H_{GOF} (Figure 1). This difference between the NRM and H_{G50} or H_{GMF} was not verified by Forni et al. (2011), who tested the same variations of H in a population of pigs. The average minor allele frequency obtained in the studied population was similar to that reported by Forni et al. (2011): 0.24 and 0.27, respectively. However, the distribution of allele frequencies was different: while in the pig population allele frequencies were all close to 0.5, in the Brahman cattle population many markers had allele frequencies distant from 0.5. The presence of these “rare” markers may reflect the fact that the families in this population can be distinct. Estimates of genetic covariance in G matrices are influenced by allele frequencies in the population. Ideally, G matrices should be estimated using the allele frequencies from the unselected base population (Forni et al., 2011), which is not available. In a real situation, it is practically impossible to obtain this information, and the three methods tested alternative solutions. Relationships using the observed allele frequencies can provide more accurate predictions of genetic merit than those that use only pedigree-derived relationships in the NRM methodology. It is possible that this increased accuracy was a result of more precise estimates of genetic covariance between relatives, as suggested by Clark et al. (2012).

On average, the choice of a relationship matrix did not influence the GEBVs, as correlations were high (Figure 2). However, when validation phenotypes were omitted (20% young omitted), the GEBVs estimated for the youngest animals in the population varied and correlations between the GEBVs from the H matrices and the NRM were lower. Choi et al. (2017) also found lower correlations between pedigree-based estimated breeding values and the GEBVs when omitting phenotype values for intramuscular fat and marbling scores in Hanwoo cattle.

The GEBVs predicted for the GEN and old datasets in all matrices did not differ significantly (Table 3).

Table 2. Additive and residual variances and heritability estimates using pedigree and genomic matrices built with the full dataset of the genotyped Brahman animals (GEN) or with 80% (old animals) of the phenotypic information⁽¹⁾.

Matrix	GEN (n=1,695)	Old (n=1,356)
Additive variance		
NRM	7.96(±1.10)	7.91(±1.32)
H_{GOF}	8.52(±0.94)	8.57(±1.12)
H_{GMF}	9.40(±1.04)	9.45(±1.24)
H_{G50}	12.71(±1.40)	12.80(±1.67)
Residual variance		
NRM	6.47(±0.76)	6.96(±0.95)
H_{GOF}	5.84(±0.58)	6.26(±0.72)
H_{GMF}	5.82(±0.58)	6.24(±0.72)
H_{G50}	5.76(±0.58)	6.17(±0.73)
Heritability		
NRM	0.55(±0.06)	0.53(±0.07)
H_{GOF}	0.59(±0.05)	0.58(±0.06)
H_{GMF}	0.62(±0.05)	0.60(±0.06)
H_{G50}	0.69(±0.04)	0.67(±0.04)
Average prediction error variance for each animal		
NRM	3.168	3.338
H_{GOF}	2.890	3.100
H_{GMF}	16.200	16.657
H_{G50}	8.221	8.207

⁽¹⁾NRM, pedigree-based numerator relationship matrix; H_{GOF} , genomic relationship matrix with observed allele frequency; H_{GMF} , genomic relationship matrix with average minor allele frequency; and H_{G50} , genomic relationship matrix with frequency of 0.5 for all markers.

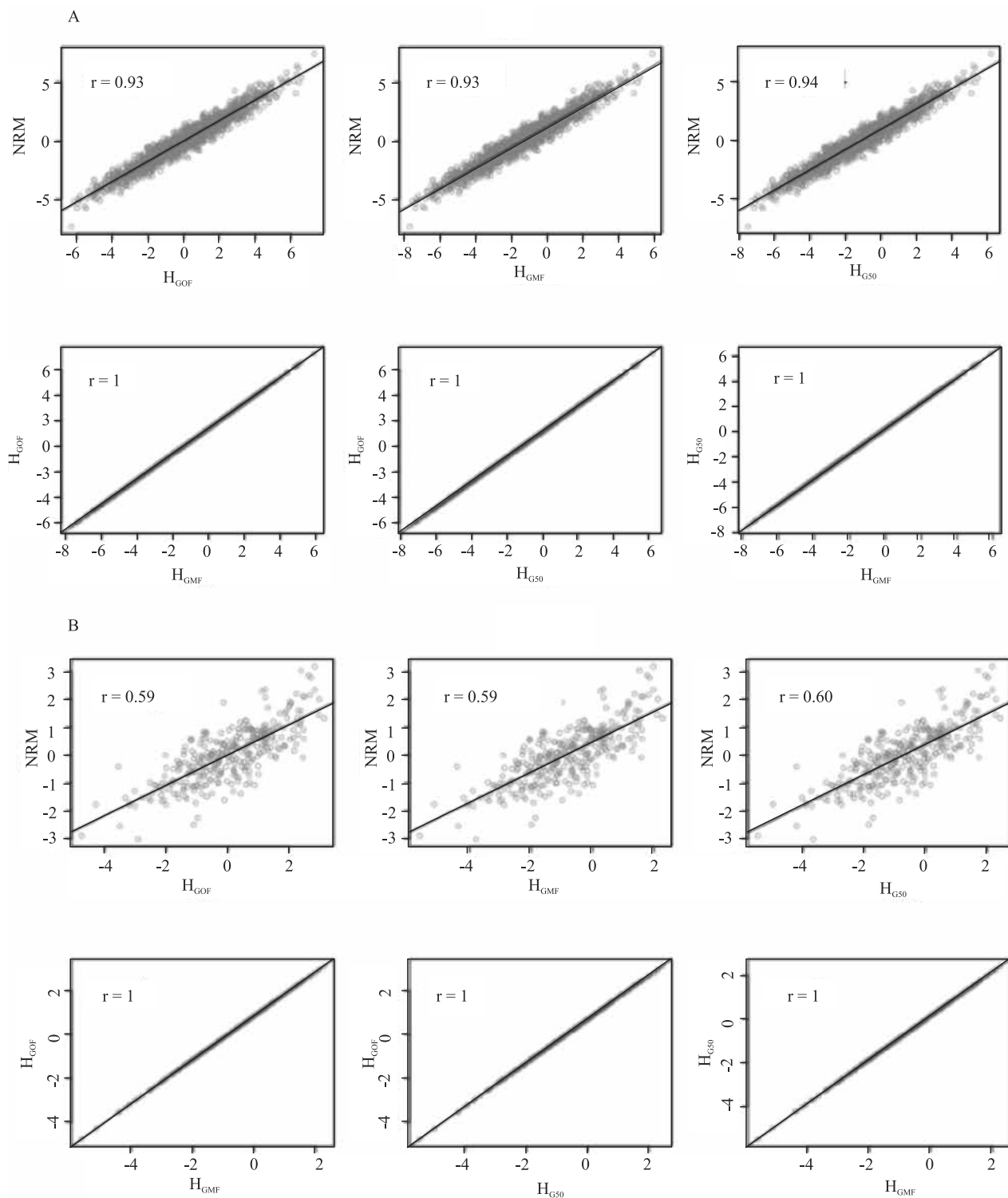


Figure 2. Correlations between estimated breeding values obtained by the pedigree-based numerator relationship matrix (NRM) and the genomic relationship coefficients with observed allele frequency (H_{GOF}), average minor allele frequency (H_{GMF}), and frequency of 0.5 for all markers (H_{G50}), using phenotypes from all 1,695 genotyped Brahman animals (A) and omitting 20% of the phenotypic information of 339 of the top 20% animals for validation purposes (B).

The accuracy of the GEBVs when young phenotypes were omitted decreased, which is in alignment with Forni et al. (2011) and Choi et al. (2017). This was expected because the G matrices include more related animals. For the young dataset, the accuracy was lower, approximately 8%, when using the NRM rather than the inclusion of genomic information. A closer relationship between the reference and validation set showed a higher accuracy of the GEBV (Choi et al., 2017). In the present study, the average accuracy reflects more variance component estimates than predictive ability; therefore, H_{GOF} provided a better rate PEV_i/σ_{ai}^2 than the other matrices. The average accuracy for H_{GOF} the highest in all population scenarios. The correlations shown in the young scenario for genomic matrices were higher than those in the NRM.

Besides the advantage of H_{GOF} presenting variance components similar to those of the NRM, it also resulted in higher accuracies for the predicted GEBVs, which may be an artefact of inflated additive variance as observed previously. It is possible that H_{GOF} was the best option in the present study for two reasons: the presence of extreme allele frequencies observed for many markers and the fact that the validation population was not independent from the calibration dataset. As the young animals used for validation are related to the old animals (calibration), it is expected that the observed allele frequencies be similar in both subgroups of this Brahman population. In practice, animals that are related to the reference population will benefit the most from genomic selection, with higher

chances of receiving accurate GEBVs. Mehrban et al. (2017) evaluated methods for genomic prediction in Hanwoo cattle and found low accuracies between the training and validation groups. These lower accuracies are mainly attributed to the small training population size and to the large effective population size.

Another difference between matrices is related to the ranking of individual animals. With the selection of 20% of the genotyped animals with higher GEBVs (top 20%, $n = 339$), 87% of them were the same (common animals) when the NRM was compared with any of the H matrices (Table 4). Between different H matrices, 99% of the top 20% animals were the same (Figure 3). However, the ranking of these top 20% animals was different between matrices, impacting the correlations between matrices (Figure 2 B). In the comparisons between the H matrices, almost all top 20% animals were the same and the Spearman coefficient between ranking positions was higher. In the comparisons between the NRM and H matrix, the correlations between animal rankings were also similar, around 0.83.

The top 20% animals were a similar group irrespective of which H matrix or NRM formulation was used. However, within this top 20%, the individual rankings of animals varied. Variations in the ranking of animals may be a problematic issue for the practical application of genomic selection because of commercial implications. In some countries, bull ranking is used as a marketing tool, and the bull ranked as number 1 could sell more doses of semen or achieve a higher price on an auction and sire a higher number

Table 3. Average accuracies of estimated breeding values (EBVs), as well as correlations between EBVs and adjusted phenotypes of the genotyped Brahman cattle⁽¹⁾.

Matrix	Accuracy			Correlation		
	GEN ⁽²⁾	Old ⁽³⁾	Young ⁽⁴⁾	GEN	Old	Young
NRM	0.776	0.699	0.457	0.969	0.900	0.479
H_{GOF}	0.813	0.746	0.536	0.938	0.868	0.613
H_{GMF}	-	-	-	0.916	0.853	0.612
H_{G50}	0.594	0.598	0.594	0.870	0.882	1.000

⁽¹⁾NRM, pedigree-based numerator relationship matrix; H_{GOF} , genomic relationship matrix with observed allele frequency; H_{GMF} , genomic relationship matrix with average minor allele frequency; and H_{G50} , genomic relationship matrix with frequency of 0.5 for all markers. ⁽²⁾All genotyped animals ($n=1,695$). ⁽³⁾Eighty percent of the population represented by the oldest animals ($n=1,356$). ⁽⁴⁾Twenty percent of the population represented by the youngest animals that had their phenotypes omitted for validation ($n=339$).

Table 4. Pearson correlations between estimated breeding values above the diagonal and Spearman coefficients calculated between the rank position of each animal below the diagonal for Brahman animals (top 20%, $n=339$) in common between the different matrices and that show the highest genomic estimated breeding values⁽¹⁾.

	NRM	H_{GOF}	H_{GMF}	H_{G50}
NRM	-	296 (0.996)	296 (0.996)	296 (0.996)
H_{GOF}	0.834	-	339 (0.999)	337 (0.999)
H_{GMF}	0.836	0.999	-	337 (0.999)
H_{G50}	0.837	0.999	0.999	-

⁽¹⁾NRM, pedigree-based numerator relationship matrix; H_{GOF} , genomic relationship matrix with observed allele frequency; H_{GMF} , genomic relationship matrix with average minor allele frequency; and H_{G50} , genomic relationship matrix with frequency of 0.5 for all markers.

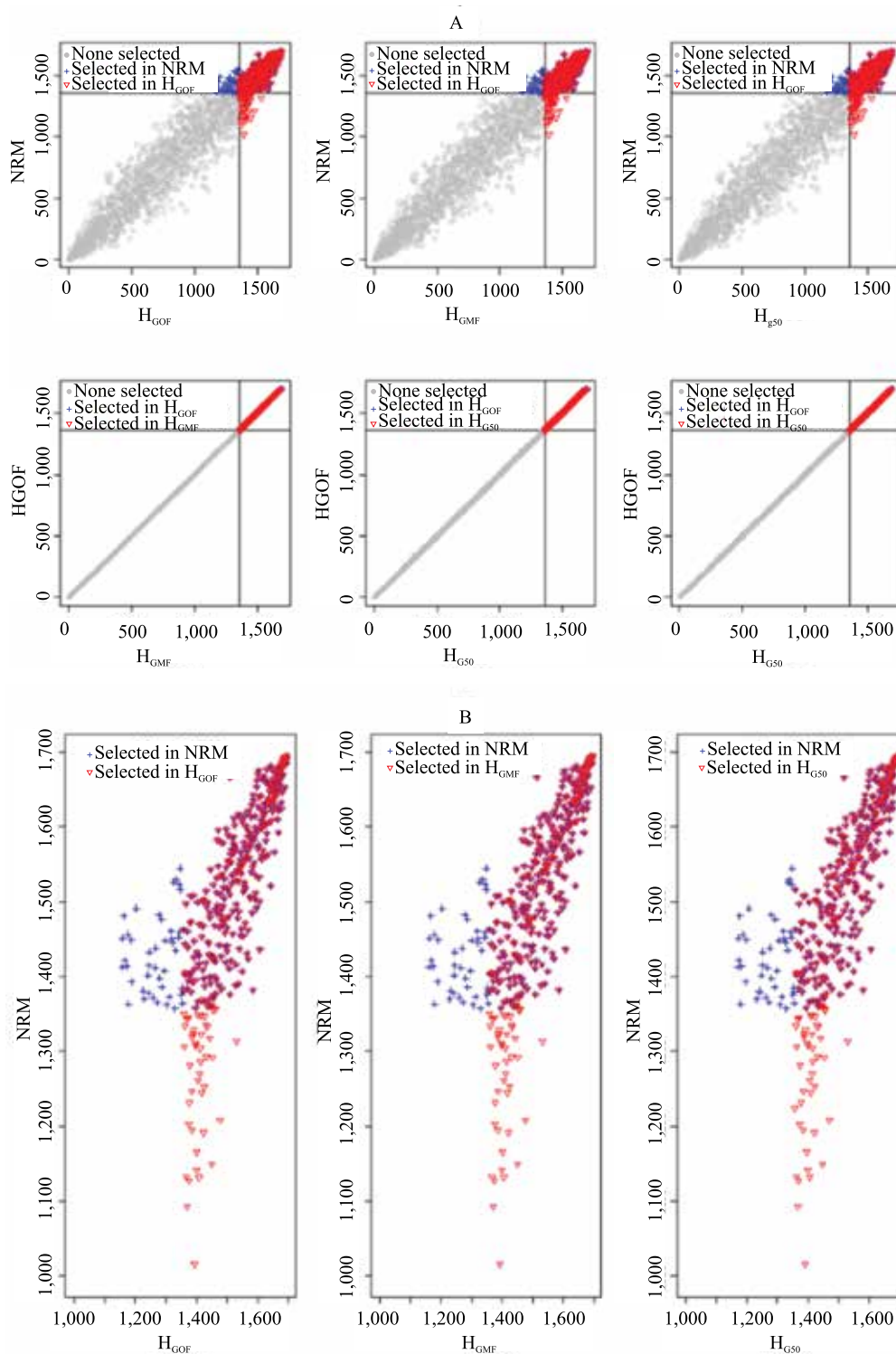


Figure 3. Correlations estimated with different relationship matrices for ranking between: A, all 1,695 genotyped Brahman animals; and B, 339 top 20% animals. Rankings were based on the estimated breeding values of all the 1,695 animals genotyped. NRM, pedigree-based numerator relationship matrix; H_{GOF} , genomic relationship matrix with observed allele frequency; H_{GMF} , genomic relationship matrix with average minor allele frequency; and H_{G50} , genomic relationship matrix with frequency of 0.5 for all markers.

of offspring in the following generation, for example. Evidently, if a different bull is ranked using different methods (NRM, H_{GOF} , H_{GMF} , and H_{G50}), there is room for discussion and conflict of interest.

Conclusions

1. The use of the observed allele frequency (H_{GOF}) is the best option for estimating the genomic relationship matrix and results in a higher accuracy of predictions.

2. The ranking of the top 20% animals is very similar for all matrices, but ranking within these varies depending on the method used.

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