

A Comparative Study on Semen Quality of Stud Bulls

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Abstract—A study was conducted to evaluate the effect of genetic group, season, temperature, humidity and temperature - humidity index (THI) on the semen quality {semen volume (ml), sperm concentration (million/ml), initial motility (%), post-thaw motility (%) and crenellation pattern (noted in code)} of 25 stud bulls (five bulls from each genetic groups of Sahiwal, Gir, Jersey cross, H.F. cross and Murrah buffalo bulls) maintained at the Frozen Semen Bull Station, Nadia, Haringhata, West Bengal, India in two seasons (i.e. winter and summer). Meteorological data was obtained from Bidhan Chandra Krishi Vishwavidyalaya, Mohanpur, Nadia, West Bengal, India. Significant variation among genetic group and genetic group x season interaction was found on semen volume, sperm concentration, initial motility, post thaw motility and crenellation pattern of semen of stud bulls ($P < 0.01$). The effect of season was found on semen volume and crenellation pattern ($P < 0.05$), initial motility and post thaw motility ($P < 0.01$). Only the, effect of air temperature was found significant ($P < 0.01$) on semen volume whereas, the effect of THI was significant statistically ($P < 0.01$) on semen volume and post- thaw motility of semen of stud bulls.

Keywords: semen quality, stud bulls, genetic group, season, temperature, humidity.

1. INTRODUCTION

Stud bull plays a unique role in cattle breeding. It has large impact on both production and profitability of commercial dairy operation. The fertility and genetic potential of bulls significantly affect the overall productivity and production efficiency of dairy herds. The two main purposes of breeding bulls are to contribute high reproduction and genetic performance to the herd [1]. In addition to being such a resource, bulls have been treated as companions and viewed with affection by those whose job it is to care for them [2]. It is the animal which provides gainful employment as well as round the year income to millions of people in India because today India has largest breeding infrastructure in the world. Artificial insemination using frozen semen is now the most widespread tool employed nationwide for improving the genetic potential of livestock [3]. Therefore, semen quality is

an important aspect of stud bull characteristics because breeding behaviour of male is closely related with its conventional semen profile (i.e., semen volume, sperm concentration, initial motility, post-thaw motility and crenellation pattern). Evaluation of semen quality and freezability of semen is the important aspect of successful breeding performance result [4]. Variation in quality of semen according to different genetic group, species and season has been found. Good management practices have positive impact on semen quality and quantity. Thus, present study has been carried out to evaluate the effect of genetic group, season, temperature, humidity and temperature - humidity index (THI) on the semen quality of bulls, so that better management practices could be adopted in the farm for exhibiting better performances by the bulls that have positive impact on its seminal quality.

2. MATERIALS AND METHOD

The study was conducted on twenty five stud bulls (five bulls of each five genetic groups viz., Sahiwal, Gir, Holstein cross, Jersey cross and Murrah Buffalo) at Frozen Semen Bull Station, Haringhata Farm under Paschim Banga Go Sampad Bikash Sangsthan, ARD Department, Government of West Bengal in two seasons, viz., winter and summer. All the animals were maintained as per routine management practices of farm under uniform environment and identical conditions. Meteorological data pertaining to both experimental seasons were collected from the Observatory managed by the Bidhan Chandra Krishi Viswavidyalaya, Mohanpur, Nadia. Temperature humidity Index (THI) (morning and afternoon) was calculated as, $THI = 0.72 (Cdb + Cwb) + 40.6$, where, Cdb and Cwb = dry and wet bulb temperature in centigrade respectively [5].

Procedure of semen collection

Semen were collected twice weekly from the same bull in artificial vagina. The site and the personnel taking collection were kept the same for all the bulls during the period of study. Collections were taken in the morning between 7.30 to 9.00 A.M under all possible sterile conditions.. Generally two ejaculates were collected from most of the bulls in a day [6, 7] with a gap of 20-30 minutes. Semen samples soon after collection were diluted in Egg Yolk Citrate diluter [8] in the ratio of 1:1. One ml. of diluted semen from each bull in separate sterilized test tube was brought to the laboratory in a thermo flask filled up with ice up to ¾ capacity. The samples brought to the laboratory were kept in a beaker containing water at approximately 32 °c for 20 to 30 minutes and were subjected to different tests as:

Semen volume was noted directly from the graduated collecting conical test tubes just after collection [9]. Sperm concentration was determined by direct cell count method [10, 11]. Neubaur Haemocytometer is used to count the sperm concentration [8] . Initial motility of sperm was determined after an hour of collection. The initial motility of fresh semen and was expressed in terms of percentage as described by [12] . Freezability of semen includes thawed motility of semen [13] which gives relationship between the number of viable sperm and fertility after artificial insemination. It was used to calculate post thaw motility of semen and was expressed in percentage. Crenellation Pattern is a method used in assessment of semen quality. It is significantly correlated with other seminal characteristics [14]. One drop of semen was taken on dry glass slide just after collection. Then it showed a characteristic pattern, called as crenellation pattern. By visualizing the pattern, the quality of semen was noted as good, moderate and bad and scored as 1, 2 and 3 respectively.

The data were analyzed using mixed model least squares analysis for fitting constants [15]. The mathematical model used to study the effect of different factors was as follows -

$Y_{ijk} = \mu + B_i + S_j + (B \times S)_{ij} + b_1(X_{ijk} - \bar{X}) + b_2(X_{ijk} - \bar{X}) + b_3(X_{ijk} - \bar{X}) + e_{ijk}$ Where, Y_{ijk} is the record for the k^{th} animal. μ is the overall mean. B_i is the random effect of the i^{th} genetic group. S_j is the effect of the j^{th} season of sampling. $(B \times S)_{ij}$ is the interaction effect of the genetic group of animal and season of sampling. b_1 is the linear regression coefficient for temperature during sampling. b_2 is the linear regression coefficient for humidity during sampling. b_3 is the linear regression coefficient for temperature humidity index (THI) during sampling. X_{ijk} is the temperature, humidity and THI, resp. corresponding to Y_{ijk} . \bar{X} is the arithmetic mean of temperature, humidity THI, resp. during sampling. e_{ijk} is the residual error element with standard assumptions. Other statistical analysis like correlation etc. was calculated as per standard statistical techniques [16].

3. RESULTS AND DISCUSSION

Mean \pm S.E. of various seminal parameters are presented in Table 1.

Table 1: Mean \pm S.E. of semen quality parameters in different groups of genetic groups

Genetic groups	Semen Volume (ml)	Sperm Concentration (million/ml)	Initial Motility (%)	Post -thaw Motility (%)	Crenellati on Pattern (score)
Sahiwal	4.75 \pm 0.14 ^{ACDE}	1630.4 \pm 0.02 ^{ACEG}	66.0 \pm 0.9 ^{ABCE}	68.92 \pm 1.5 ^{ACEG}	1.89 \pm 0.07 ^{ABCD}
Gir	5.04 \pm 0.14 ^{DILN}	1460.2 \pm 0.02 ^{FJNQ}	61.6 \pm 0.9 ^{DIMP}	30.4 \pm 1.4 ^{FLPS}	2.03 \pm 0.07 ^{CHEM}
Jersey cross	4.89 \pm 0.14 ^{CGLM}	1086.4 \pm 0.02 ^{DIMO}	65.8 \pm 0.9 ^{BGLN}	49.8 \pm 0.7 ^{DJOQ}	2.11 \pm 0.07 ^{CFJK}
H.F. cross	3.96 \pm 0.14 ^{BFHJ}	1263.9 \pm 0.02 ^{BHJK}	66.7 \pm 0.9 ^{AGHJ}	61.6 \pm 1.1 ^{BIKM}	1.77 \pm 0.07 ^{AEIG}
Murrah	4.15 \pm 0.14 ^{EKMO}	1584.9 \pm 0.02 ^{GLPR}	70.5 \pm 0.9 ^{FKOQ}	26.8 \pm 1.5 ^{HNRS}	1.65 \pm 0.07 ^{DILN}

Semen Volume

It could be seen from the result that the overall semen volume irrespective of season were highest in Gir (5.04 \pm 0.14 ml.) followed by those in Jersey cross (4.89 \pm 0.14 ml.), Sahiwal (4.75 \pm 0.14 ml.), Murrah (4.35 \pm 0.14 ml.) and H.F cross (3.96 \pm 0.14 ml.) in that order (Table 1). The effect of genetic groups, season, and genetic group x season interaction, air temperature and THI were found to be statistically significant at 1, 5, 1, 1 and 1% level respectively. Seasonal variation were observed in Sahiwal, H.F cross, Jersey cross and Murrah buffalo bulls between winter and summer. Higher volume was recorded during winter than summer season (Fig. 1). These findings coincided with some researchers in Sahiwal bulls [17, 18] and in H.F. cross bulls [19, 20] but some were in contrast [21-23] with the present findings. Comparatively lower ejaculate volume were reported in crossbred bulls [24], Sahiwal bulls and Murrah bulls respectively [19, 20]. Comparatively higher value than the present result was found in H.F cross bulls [25]. Higher values in winter season were also reported in Murrah buffalo bulls [26, 3].

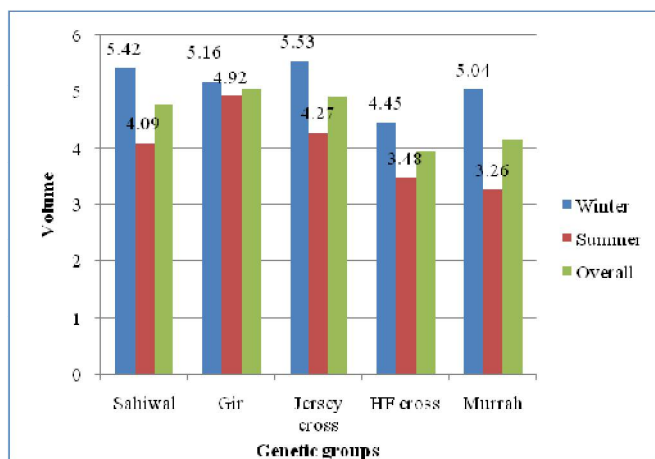


Figure 1: Semen volume of different genetic groups in different seasons

Sperm Concentration

Overall sperm concentration irrespective of season were found to be highest in Sahiwal (1630.4 ± 0.02 million/ml.) followed by those in Murrah (1584.9 ± 0.02 million/ml.), Gir (1460.2 ± 0.02 million/ml.), H.F.cross (1263.9 ± 0.02 million/ml.) and Jersey cross bulls (1086.4 ± 0.02 million/ml.) in that order (Table 1). The effect of genetic group and genetic group x season were found significant statistically ($P < 0.01$). Only the differences in sperm concentration between H.F. cross and Jersey cross in winter and summer season were significant statistically ($P < 0.05$). The present findings were higher than the present values in Sahiwal bulls [17, 22, 24, 27-29]. Similar value was reported in Jersey cross bull [25] and in H.F. cross bull [29], whereas lower value was found in H.F cross bull [25].

Initial motility

Overall initial motility of sperm irrespective of season were highest in Murrah ($70.5 \pm 0.9\%$) followed by those in H.F. cross ($66.7 \pm 0.9\%$), Sahiwal ($66.0 \pm 0.9\%$), Jersey cross ($65.8 \pm 0.9\%$) and Gir ($61.6 \pm 0.9\%$) in that order (Table 1). The effect of genetic group, season and genetic group x season were found significant statistically ($P < 0.01$). Initial motility was found to be lower in summer season in the present finding (Fig. 2). The differences in initial sperm motility between Sahiwal, Gir, Jersey cross, HF cross and Murrah buffalo bulls in winter and summer season were significant ($P < 0.05$). The differences in initial sperm motility between Gir and Sahiwal, Murrah and Sahiwal, H.F. cross and Murrah, Jersey cross and Murrah and Gir and Murrah bulls were significant statistically ($P < 0.05$). The present findings were in accordance with the observations of various researchers [22, 30, 27], but comparatively higher sperm motility in crossbred cattle and Murrah buffalo bulls than the present values were reported [31, 32, 24, 19, 20]. Significant seasonal variation in initial motility found in this study was in agreement with the reports

of some research workers [33, 34]. This variation might be due to genetic differences, management and type of animal and adaptability of animal towards environment.

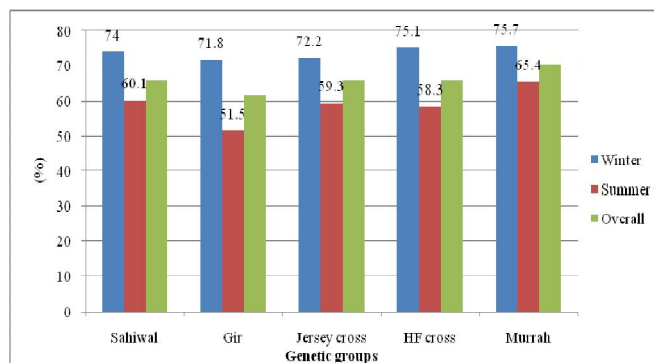


Figure 2: Initial motility of different genetic groups in different seasons

Post - thaw motility

Overall post-thaw motility of sperm irrespective of season were highest in Sahiwal ($68.92 \pm 1.5\%$) followed by those in H.F. cross ($61.6 \pm 1.1\%$), Jersey cross ($49.8 \pm 0.7\%$), Gir ($30.4 \pm 1.4\%$) and Murrah ($26.8 \pm 1.5\%$) in that order (Table 1). Only the effects of genetic group, season, genetic group x season and THI were found significant statistically ($P < 0.01$). Post-thaw motility was less in summer season (Fig.3). The differences between post-thaw motility of sperm between H.F.cross and Jersey cross. in winter and summer season was found significant statistically ($P < 0.05$). Similar values were found in Friesian cross bulls [28]. Lower values in comparison to present findings were reported in Kundhi buffalo and crossbred bulls respectively [35, 20], whereas higher value was reported in Murrah buffalo bulls [20]. This variation might be due to the variation in climatic conditions of these two seasons and the uniformity in managerial practices under which these bulls had been kept.

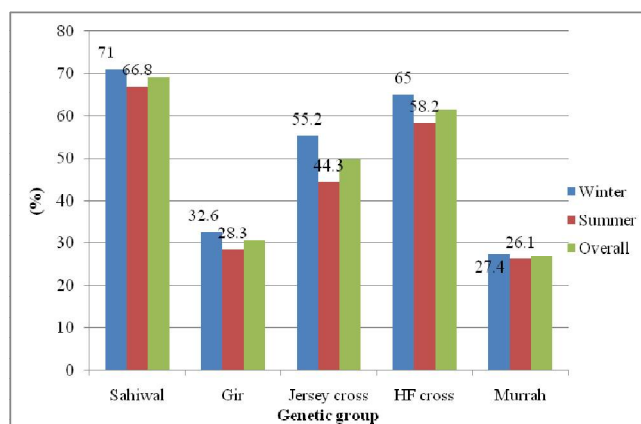


Figure 3: Post - thaw motility of different genetic groups in different seasons

Crenellation Pattern

Overall crenellation pattern of semen irrespective of season were lowest in Murrah (1.65 ± 0.07) followed by those in H.F.cross (1.77 ± 0.07), Sahiwal (1.89 ± 0.07), Gir (2.03 ± 0.07) and Jersey cross (2.11 ± 0.07) in that order (Table 1). The effect of genetic groups, season and genetic group x season interaction were found statistically significant at 1, 5 and 1% level respectively. Good crenellation pattern of semen was found during winter in comparison to that in summer season (Fig.4). The differences in crenellation pattern of semen between Sahiwal, Gir, H.F. cross and Jersey cross in winter and summer season were found to be significant ($P < 0.01$). Similar variation was observed in Karan Fries bulls [36] and in Murrah and crossbred bulls [37]. All these variations found might be due to differences in genetic constitution of bulls, variation in management and adaptability of the animal to particular situation.

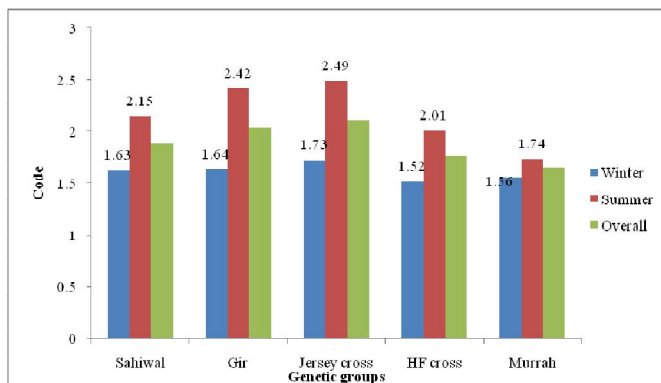


Figure 4: Crenellation pattern of different genetic groups in different seasons

CONCLUSION

Variation in semen quality was shown by different genetic groups of stud bulls. Better semen quality (semen volume, initial motility, post-thaw motility and crenellation pattern) was observed during winter than summer season. The differences found might be due to the species specificity of the bulls or due to the different expression of the trait as a result of genotype-environmental interactions, adaptability towards environment and management condition.

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