The Indian Journal of Animal Reproduction; 23(2): 135 - 137; December 2002

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Effect of prostaglandin F₂α, oxytocin and butylated hydroxytoluene on fertility of crossbred (HF x Hariana) bull spermatozoa*

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> Receiped : August 4, 2001 Accepted : June 14, 2002

ABSTRACT

Semen samples collected from two crossbred bulls (HF x Hariana) were diluted in tris with three combinations of additives i.e., prostaglandin $F_2\alpha$ (PGF₂ α) (1.5 ng/ml), oxytocin (0.1 unit/ml) and butylated hydroxytoluene (BHT) (2 mM/ml) and frozen in liquid fitrogen. A total number of 176 oestrus cows were inseminated with all the additives treated and control semen samples, the pregnancy diagnosis was carried out 50-60 days post insemination. The results, revealed overall conception rate 40.48% 48.28% 42.31% and 42.91% in control, PGF₂ α , oxytocin and .BHT treated groups. No significant difference was observed among the treated groups separately. However, the conception rate was higher with PGF₂ α treated semen when compared to control.

Key words : Bull, spermatozoa, conception rate. PGF2a, oxytocin. BHT

Improvement in the keeping quality, freezability and fertility of bovine semen by incorporating various additives is the subject of study for the last few years. Different additives have been tried to enhance the motility of spermatozoa with the view to improve the conception rate (Balasubramanian, 1979). The addition of prostaglandins to semen prior to freezing was observed to improve the fertility rate in sheep (Gustafsson et al., 1975) and buffaloes (Muralinath, 1988). Oxytocin is known to increase uterine and oviductal contractions (Reeves, 1987) and intrauterine infusion of oxytocin increases sperm transport (Baker et al., 1968). BHT added to semen extenders prior to freezing may improve the fertility of bull semen and reduce the risk of transmitting viral diseases to cows during A.I. (Anderson et al., 1994). Present investigation is an attempt to enhance the conception rate (CR) with the frozen semen containing added PGF2 a, oxytocin and BHT additives.

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MATERIALS AND METHODS

Twenty four ejaculates (12 from each bull) from two crossbred bulls (HF x Hariana) maintained under identical feeding and managemental regimen were used for the study. Semen was collected in A.V. twice a week. The ejaculates were split into 4 parts. Each part was diluted separately with Tris+additive combinations @(1.5 ng/ml PGF₂a, 0.1 unit/ml oxytocin) as per the method described by Lokanathan, (1993) and BHT 2 mM /ml as per Killian et al. (1989), respectively and the 4th aliquot was kept as control. The four split samples were Frozen in medium french straws after providing combined cooling cum equilibration period of 4 hrs. A total number of 176 oestrous cows were inseminated with the frozen semen of treated and control groups (after thawing at 37°C for 30 seconds). Pregnancy verification was carried out 50-60 days post insemination. Statistical analysis of the data was done according to the method described by Snedecor and Cochran, (1989).

RESULTS AND DISCUSSION

The results obtained in this study are presented in Table 1. The mean conception rate was recorded higher

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Treatment Total no. of animals of inseminated f		Total no. of animals followed up	Total no. of animals pregnant	Conception rate (%) at 1st insemination	Overall mean mean CR (%)	
		I	Buli A			
Control	25	23	10	43.47	40.48	
Prostaglandin F ₂ α 25 Dxytocin 25		23	12	52.17	48.28 42.31	
		24	11	45.83		
BHT	25	24	11	45.83	42.91	
		I	Bull B			
Control	19	16	6	37.50		
Prostaglandin F _o a	19	18	8	44.40		
Oxytocin 2	19	18	7	38.80		
BHT	19 15		6	40.00		

Table	1.	Conception	rate (post	AI)	in	different	additive	combinations
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in PGF₂ α treated group (48.28%) followed by BHT (42.91%). oxytocin (42.31%) and control (40.48%). However, the difference in mean values among the traeted groups did not reach any statistical significance.

The CR in PGF₂ a treated semen was higher (48.28%) than control (40.48%) in this study, which is well comparable to (Gokeen et al., 1985) ram and (Radosalvov et al., 1990) bull semen. However, in this study CR was higher than that reported in buffaloes (Reddy et al., 1992). The trend in improved CR of our study was similar to that reported in ewes (Gustafsson et al., 1977) and rabbit (Edquist et al., 1975), using PGF₂a treated semen. This improvement possibly because of an increased sperm number in the oviduct following A.I. in rabbit (Edquist et al., 1975) and ewe (Gustafsson et al., 1977). The CR in oxytocin treated semen was higher (42.31%) as compared to control (40.48%) in this study. But, this is lower than cows (Drume, 1975 and Shubin et al., 1982) and sows (Schlegel and Loebel, 1972). The difference in the CR among studies might be due to the variation in the concentration of oxytocin used in such studies. Addition of oxytocin increases uterine motility and sperm transport to oviduct (Reeves, 1987). Probably favoured the CR in the present study as it was in sows (Martinek et al., 1978). The CR obtained in the BHT treated semen was relatively higher (42.91%) than control (40.48%) but failed to reach statistical significance. Similar response was reported elsewhere (Anderson et al., 1994). It can be concluded that the addition of PGF₂ α , oxytocin and BHT prior to semen freezing, PGF2a performed much better than other additives and control, but statistically it is not significant with control. Gu

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ACKNOWLEDGEMENT

The authors thank the Director, Joint Director (Academic), Head. Animal Reproduction Division, IVRI for providing facilities, and ICAR, New Delhi for the providing fellowship during the study.

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