



Synchronization of oestrus by 'buck effect' and PGF₂α treatment in local indigenous goats

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Received: 14 October 2010; Accepted: 23 February 2012

ABSTRACT

The study was carried out to investigate the influence of buck effect and PGF₂α treatment on oestrus synchronization in local indigenous goats. Apparently healthy normally cycling nulliparous goats (16) were divided into 3 groups considering their age, body weight and serum progesterone concentration indicating their similar stage of oestrous cycle prior to treatment. The group 1 (control, n=4) was kept separate without treatment, the group 2 (T₁, n=6) was treated with 2 injections of PGF₂α at 11 days apart @ 7.5mg/animal/injection while the group 3 (T₂, n=6) was teased with a sexually-active-aproned buck. The proportion of females showing behavioural oestrus was successfully synchronized by double injection of PGF₂α at 11 days apart, as well as by male introduction in T₁ and T₂ groups, respectively. The time interval between start of treatment and onset of oestrus were significantly shorter in T₁ (64.00±3.35 h) and T₂ (65.00±18.50 h) than in control (264±30.24 h) group but was similar in both T₁ and T₂. However, the duration of oestrus was 30.5±1.5 h, 34.00±3.69 h and 30.00±4.38 h for control, T₁ and T₂ groups, respectively. There was no significant difference in conception rates amongst the groups. All the goats conceived after second service irrespective of treatment groups. It was found that the buck effect appeared to be as effective as conventional PGF₂α treatment in indigenous goats for synchronization of oestrus.

Key words: Buck effect, Goat, PGF₂α, Synchronization

Oestrus synchronization has been successfully used in enhancing reproductive efficiency, particularly in small ruminants (Kusina *et al.* 2000). Oestrus synchronization in does can be achieved either by reducing the length of the luteal phase of oestrous cycle with prostaglandin F₂α (PGF₂α) and its analogues (Jainudeen *et al.* 2000) or by extending it artificially with exogenous progesterone (Karatzas *et al.* 1997). These hormonal treatments are not easily available or are too much expensive for the farmers in developing countries like India. Therefore, it requires development of managemental strategies such as pheromones and biostimulations. Induction of synchronous oestrus behaviour and ovulation in anovulatory females after introduction of buck is called the buck effect (Ungerfeld *et al.* 2004). This response is widely used to advance and synchronize breeding in goats (Veliz *et al.* 2002). The 'buck effect' is now one of the well established means of

manipulating goat reproduction without using much criticized hormonal therapy. The objective of the present study was to determine the influence of 'buck effect' and PGF₂α administration on oestrus synchronization in local indigenous goats.

MATERIALS AND METHODS

Apparently healthy, normally cycling nulliparous goats (16) were selected on the basis of age (15.38±0.96 months) and body weight (13.63±0.58 kg). There was no significant difference of age and body weight among the groups. Pre-treatment serum progesterone concentration was estimated to ensure similar stage of oestrous cycle in all experimental groups. The experimental animals were maintained under stall-fed condition and housed in separate sheds, each attached with open paddock. Experimental animals were fed with cultivated green fodders and concentrate feed. Does and buck were daily fed with 250 g and 300 g of concentrate, respectively as per routine feeding system of the farm. The breeding of animals is usually done during Nov-Jan at this farm. Experimental animals were divided into 3 groups of 4 in control and 6 each in first treatment (T₁) and second treatment (T₂). The control group was kept separate without any treatment; T₁ group was treated with 2 injections of

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PGF₂α (dinoprost tromethamine) at 11 days apart @ 7.5mg/animal/injection, while in T₂ group, a sexually active buck was introduced and allowed to stay with does continuously day and night till all the does shown behavioural oestrus. The buck was aproned to prevent undesired mating. Heat detection was done by teaser buck in control and T₁ group and by the same buck kept with does in T₂ group daily at 6 h interval, i.e., at 04:30 h, 10:30 h, 16:30 h and 22:30 h. Onset of oestrus and duration of oestrus were recorded for each doe in all groups and 3 bucks were allowed to mate with the does. The time interval (h) between onset of oestrus and the treatment was calculated by taking difference between the recorded date and time of starting the experiment and first incidence of one or more behavioural activity characteristic to females in oestrus. While the duration of oestrus (h) was calculated by taking difference between the recorded time of onset and end of oestrus. The does were considered to be at oestrus by observing two main signs i.e. does allowing buck to mount and vigorous wagging of tail. Similarly, the end of oestrus was considered when mounting by male was not allowed and there was no wagging of tail. Conception rates were calculated on the basis of non-return rate and serum progesterone level.

Blood samples were collected daily starting from a day prior to treatment till exhibition of oestrus and at an interval of 4 days for 10 consecutive days thereafter i.e. till 40 days post-mating. Sampling was done between 8 AM to 11 AM daily in all the experimental females. Serum samples were separated and stored at -20°C till hormonal assay. Serum progesterone was estimated by RIA using standard diagnostic kit.

The general linear model of one way ANOVA was used for analyzing continuous data as per the procedure outlined in SPSS® 11.00 statistical packages. The difference among the percentages and frequencies were tested using nonparametric test (Chi-square test).

RESULTS AND DISCUSSION

The effect of treatments on synchronization of oestrus, oestrus onset, oestrus duration and conception rates were shown in Table.1. The results revealed that the proportion of females shown behavioural oestrus was successfully synchronized by double injection of PGF₂α at 11 days apart as well as by buck effect in T₁ and T₂ groups respectively. The PGF₂α treatment and buck effect induced behavioural oestrus, respectively in 83.3% (5 out of 6) and 50.0% (3 out of 6) cyclic does within 72 h of treatments. Remaining 1 doe in T₁ group and 3 does in T₂ group shown behavioural oestrus between 72–120 h of treatment while all the does (4 out of 4) in control group shown behavioural oestrus after 120 h. The oestrus synchronization response in T₁ group was consistent with observations reported by others (Alvarez *et al.* 2010, Vazquez *et al.* 2010 and Ahmed *et al.* 1998). However, Perera *et al.* (1978) and Taylor (1978) reported 83.33% and 100% synchronization of oestrus by PGF₂α

Table 1. Effect of PGF₂α treatment and buck introduction on oestrus response and synchronization in cyclic does

Attributes	Control	T ₁	T ₂
No of does treated	4	6	6
Does showing behavioural oestrus (χ ² value 0.02)	4 ^a (100)	6 ^b (100)	6 ^b (100)
<i>Degree of oestrus synchronization</i>			
Up to 72 h (χ ² value 0.25)	-	5 ^a (83.33)	3 ^b (50.00)
72 to 120 h (χ ² value 0.25)	#	1 ^a (16.66)	3 ^b (50.00)
<i>Oestrus response</i>			
Onset of oestrus(h)	264 ^b ±30.24	64.00 ^a ±3.35	65.00 ^a ±18.50
Duration of oestrus(h)	30.5±1.5	34.00±3.69	30.00±4.38
<i>Conception rate</i>			
At first service (χ ² value 0.33)	3 ^a (75)	4 ^b (66.66)	5 ^c (83.33)
At second service (χ ² value 0.33)	1 ^a (25)	2 ^b (33.33)	1 ^a (16.66)

Figures with different superscripts in a row differ significantly (P<0.05) for oestrus response and against respective χ² value (in parentheses) for frequency data. Figures in parenthesis indicate percentages. # All the control females exhibited oestrus after 120 h of start of treatments in other groups.

treatment in very short time with a range of 18–23 and 24–48 h, respectively. In T₂ group, 100% synchronization of oestrus (between 14–116 h) was in agreement with Keskin (2003). However, the maximum time taken to synchronize oestrus in all the does was lower in the present study. The buck effect had been shown as a multi-sensory phenomenon involving olfactory, visual, tactile and auditory cues. High% of females responds to male stimulation when all cues are active (Chemineau 1987). The gonadotropin-releasing hormone (GnRH) pulse generator generated intermittent GnRH discharges into the portal vessels and thereby regulated the pulsatile LH secretion into the peripheral circulation (Karsch 1984). Therefore, the central target of the buck effect pheromone was thought to be the hypothalamic GnRH pulse generator.

The average time interval between start of treatment and onset of oestrus was significantly shorter in T₁ (64.00±3.35 h) and T₂ (65.00±18.50 h) than in control (264.5±30.24 h) group. The mean duration for onset of oestrus in T₁ group was similar to reports of others (Vazquez *et al.* 2010, Alvarez *et al.* 2010 and Amaranditis *et al.* 2004). However, the interval reported by Pandey *et al.* (1985) and Greyling and Van Niekerk (1991) was longer than the present findings. The onset of oestrus in T₂ group was in agreement with Keskin (2003).

Onset and end of oestrus was considered to be 3 h less than the first time of two consecutive detections and non-detections, respectively (Dhali *et al.* 2005). The duration of oestrus was 30.5±1.5 h, 34.00±3.69 h and 30.00±4.38 h for

control, T₁ and T₂ groups, respectively (Table 1). Although there was no significant difference between the groups, the oestrus duration was more in T₁ group which might be due to exogenous hormonal treatment. The results obtained in the present study corroborated with the earlier findings (Sahni and Roy 1976, Greyling and Van Niekerk 1991). However, the duration of oestrus was observed shorter by Pandey *et al.* (1985), Ishwar and Pandey (1992) and Akusu and Egbunike (1984) while longer duration was observed by Amaranditis *et al.* (2004). The nonsignificant difference in the oestrus duration among all the groups in present study suggested that PGF₂ α or buck effect did not interfere with the duration of oestrus. The progesterone concentration remained at basal levels (<1ng/ml) during the oestrus, as observed by others (Pathak *et al.* 1992, Nandy *et al.* 2001 and Khadiga *et al.* 2005).

There was no significant difference in conception rates between the groups. The conception rate was calculated on the basis of two services. After first service, the maximum conception rate was observed in T₂ group (83.3%) followed by control group (75%) and T₁ group (66.7%) while the remaining 25%, 33.3% and 16.7% in control, T₁ and T₂ groups, respectively conceived after second service. Overall 100% does in all the groups conceived after second service. The conception rate of T₁ group (66.7%) was in agreement with Hearnshaw *et al.* (1974) and Serna *et al.* (1978) who got 56 and 71.4% conception rate, respectively, following synchronization of oestrus by PGF₂ α analogue. However, the higher conception rates were observed by Pandey *et al.* (1985) and Akusu and Egbunike (1984). The results of present investigation showed 83.33% conception rate, which was in agreement with Chemineau (1983) who reported 81% conception rate in goats cycling before introduction of male. Higher conception rate after first service was observed in T₂ group which might be due to continuous presence of male, which prepared the female to get ready to mate and conceive under the most natural social conditions. Overall serum progesterone concentration for control, T₁ and T₂ group over 40-days post-mating was 6.88 $\pm$ 0.61, 7.34 $\pm$ 0.52 and 6.05 $\pm$ 0.57 ng/ml, respectively, which confirmed the conception rates.

Thus, it can be concluded from the present study that the buck effect appears to be as effective as conventional PGF₂ α treatment in indigenous goats for synchronization of oestrus without any extra input.

ACKNOWLEDGEMENT

Authors are highly thankful to the Director, Indian Veterinary Research Institute for extending help in taking up such research. The financial help from Indian Council of Agricultural Research is duly acknowledged.

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