OL. 6 No. 1 : JUNE 1985



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The Indian Journal of Animal Reproduction

JOURNAL OF THE INDIAN SOCIETY FOR THE STUDY OF ANIMAL REPRODUCTION

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Editarial

Embryo Transfer in Bovines as a Potential Tool for Increased Milk Production by 2000 A.D.*

Traditionally, the bull is said to be half the herd. In last few decades, there has been widespread use of artificial insemination, thereby exploiting the potential of available proven or good quality sires to increase the milk production. The great potential left to exploit the remaining half of the herd i.e. the female's participation in herd improvement and genetic conservation initiated the embryo transfer experiment in the cattle. The coming years will witness the embryo transfer technique as a routine. It is based on the principle of inducing superovulation in superior donor cows and buffaloes and later inseminating with pedigreed bull's semen. A donor animal can be subjected for embryo transfer five to six times a year or till there are no abnormalities, lesions or adhesions in the uterus and fallopian tubes that may develop from constant or improper flushing. Once recovered, the embryos can be either transfered immediately into recipients or frozen or subjected to micro-manipulation before transferr. In case of immediate transfer the embryos are individually placed non-surgically into the uterus of suitably synchronised recipients to complete the term of pregnancy. The principal advantage of non-surgical method is that the elaborate facilities, costs, risks and postoperative adhesions associated with surgery are avoided. On the farm collections are possible, lactating cows that are bad surgical subjects can be used and most importantly collections can be repeated from the same donor.

Although, the offspring born would be fostered by the recipient, its genetic make-up would still be the combination of traits from its original dam (donor) and sire. Embryos can be subjected to micro-manipulation such as splitting by micro-surgery (cloning) and sexing by identification of sex chromatin or karyotyping. A technique recently developed for sex determination looks more promising, with the use of H-Y antigen which is male determinant. Techniques similar to the production of frozen semen are now available for freezing embryos in liquid nitrogen for long term storage and transport.

Another major possibility is twin pregnancy, either 2 embryos of same sex can be transferred or a single embryo split into two, which will have same sex can be transferred. The other side of embryo transfer potential for increasing milk production is the increase in reproductive rate, increase in accuracy of progeny testing, increase in accuracy of selection procedures, better genetic control, establishing gene banks and effective utilisation of low valued poor germ plasm as recipient mothers. Even calves from good pedigreed prepuberal heifers can be obtained and age at maturity drastically reduced.

Embryo transfer has made a great impact on international livestock trade and movement of genetically superior germ plasm. This reduces high transportation costs. Quarantine requirements are met more easily. Neither transit mortality nor fatigue occurs nor difficulty in acclamatisation is faced. Embryo transfer will open avenues for contract mating especially at the metro politian city farms where still large number of good quality animals are brought for only one lactation and culled. The trend of these animals which eventually end-up at slaughter houses instead of returning to their breeding tract can be brought to an end.

Success rate in non-surgical transfer of embryo varies with potency and response to superovulating agents, season, lactating stage of the donor, age of the donor, progesterone level during the stage of follicular stimulation, experience and knowledge of the performer, reproductive state of the recipient, individual variation, hygienic standards and various other factors. The uterine environmental changes at different stages of estrous cycle is essential for the development of the embryo and it is imperative to meet the environmental requirment of the embryo in the recipient. If the difference is large, embryo is often rejected by the recipient resulting in unsuccessful transfer. Hence controlling all the variable factors and synchronising the estrous of donor and recipient is a necessity, which is not difficult any more with the advent of prostaglandin F2a or its synthetic analogue cloprostenol.

ETT being an area of reproduction has excited the imagination of researcher and farmers alike by pointing towards near future in which cows of superior genetic merit would be effectively superovulated and embryo taken one by one and transferred for subsequent development in the uteri of foster mothers of promiscous genetic merit.

Should AI technology be described as the first wave of revolution in dairy husbandry industry the second wave is indeed imminent with the development of ETT. With deep freezing of the male gametes (sperms) it is possible to widely use the genetically superior sire component. Similar use of genetically superior female gametes however, has been limited to life time production of ova, its normal in-vitro fertilization, its successful implementation and embryonic development thereafter. With rapid development of embryo transfer technology applicable to domestic animals, the new vista for greater utilisation of genetically superior female component has been opened up to make rapid strides in milk production.

Dr. Kurien, Chairman N.D.D.B. & Vice Chancellor, G.A.U. observed in his article that keeping in view the application of advanced technology to the dairy insustry, tomorrow's professionals would have to possess knowledge or appreciation of computers and a system approach in addition to their skills in classical disciplines of dairy science; namely animal husbandry, dairy technology and engineering etc. Our teaching and research institutions shall have to be fine-tuned to these realities of tomorrow.

Editorial Board

* SOURCE: 1. Dairy India 1985, 2nd edition, published by P.R. Gupta, New Delhi. , 2. News item entitled "Milk Production to be doubled by 2000 AD" in Times of India, Ahmedabad edition dated 22nd April, 1985.

Histological Changes In Male Gonads From Birth To Sexual Maturity In Buffaloes

LALITA V. DESHPANDE AND K. JANAKIRAMAN

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Reproductive Biology Research Unit Gujarat Agricultural University Anand Campus, Anand

ABSTRACT

Histological changes in male gonads from birth to sexual maturity in buffaloes have been described.

* * *

The buffalo plays an important role in the Indian agricultural economy. With the realization of their importance more and more research on the species is taking place. However, the studies on male aspect are limited. With this study an attempt is made to understand the developmental changes in male gonad from birth to sexual maturity. Postnatal histological changes in testis are reported here; which indicate that the animal has a potential for development as early as 15-16 month of age.

Materials and Methods

The testicular tissue from 4 male buffalo calves was sampled by repeated incision biopsies (as per Hotehkiss, 1944) over 17 stages from birth to 600 days of age. The incision biopsies did not have any deteriorating effect on the testicular growth and development. The testicular tissue sampled at biopsy was fixed in Bouin's fixative, was processed as per Luna (1968) and sectioned at 5-7 microns thickness and stained with Haematoxylin and Eosin.

Seminiferous tubule diameter:-The seminiferous tubule diameter was measured at 2 different places (maximum and minimum) by ocular linear micrometer scale. Their average was worked out to give the mean diameter of that particular section of the tubule. The overall mean was an average of 150 such observations. The magnification used for this measurement was 10×45 . The size was expressed as units which when multiplied by the least count 3.30 gives the measurements in microns.

Tubular count:—The number of the cross sections of the seminiferous tubules present in a microscopic field were counted to give tubular count. Such countings were done at 10 different location of the same sections at a given stage in a given animal, and then the mean of 10 observations was taken as tubular count. The magnification used was 10×10 . Each mean represents the number of tubular cross sections occupying 400 square in the net micrometer. One square of the net micrometer comes to 16C0 square microns of the section (under 10×10 magnification).

Interstitial space:—Initially the interstitial space was measured microscopically by the number of squares occupied by interstitial space out of 100 squares of net micrometer $(10 \times 45 \text{ magnification})$. When multiplied by 20.25 least count, square micron area occupied by interstitial space can be known.

Histological changes in male gonads from birth to sexual maturity in buffaloes.

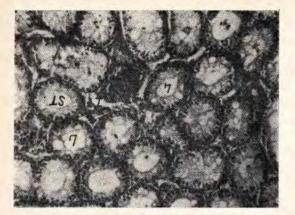


Fig. 1: Cross section (cs) of seminiferous tubulos (ST) in Day old animal. ST are solid and are separated by interstitial space (I) (Magnification 10×10).

Initiation of lumen formation:—This fact was observed qualitatively and the age at which the tubuler were first canalized was noted. The degree of lumenation was also recorded.

Results and Discussion

The growth of seminiferous tubule is the primary feature of testicular development. The size of seminiferous tubule increases from birth to sexual maturity. The mean diameter of the tubules was the criteria adopted to measure growth.

SEMINIFEROUS TUBULE DIAMETER:

The diameter of the tubules increased four fold from birth to 600 days of age. The diameter on Day one was as low as 11.70 ± 0.38 units (Fig. 1), and was maintained on Day 28. Thereafter it kept gradually increasing and till day 240 the rate of increase was very slow. On Day 360 the tubular diameter got doubled the value at birth. The increase in diameter was much spectacular from 390 to 450 days (Fig. 2) again from Day 570 to 600 great enlarge-

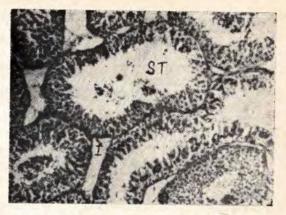


Fig. 2: Cross section of seminiferous tubule (ST) of 420 day old animal, showing lurmn (L) formation in all tubules (10×10 magnificat ion). Increased ST size limits the interstitialspace (I) into triangles.

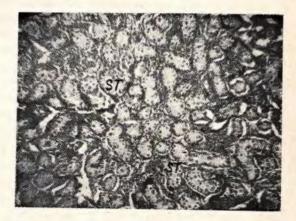


Fig. 3: Cross section of ST in low fener (from 690 day old animal) showing fully developed ST.

ment in tubular diameter was observed (Fig. 3). On Day 600 the tubules measured 47.50 units in diameter. Such sigmoid growth curve was also observed by Goyal and Dhingra (1973) in buffalo; by Abdel Raouf (1960) in Swedish Red and White breed of cattle and by MacMillan and Hafs (1969) in Holstein bulls calves. TUBULAR COUNT:

The tubular count was inversely related with the seminiferous tubule diameter,

when the later increases, the count decreases. The number of cross section of tubules in a given microscopic field halved at 240 days of age (mean+SE 202.5 +25.94) than the one present at birth i.e. 398.0+6.15. Reduction in the tubular count was slow till Day 125 and faster between 125 to 240 Days of age. On Day 300 the tubular count was a high as 233.2+18.66 which got reduced to 134.00+15.08 on Day 420. Finally, the mean value of tubular count was found to be only 49.40+23.22 on Day 600. The overall was a declining pattern, regarding tubular count in correlation with semini ferous tubule diameter and interstitiaspace.

INTERSTITIAL SPACE:

At birth the tubules were widely separated apart from each other by the interstitial space, which on crude estimates accounted for 29.62 per cent of the total space area observed, rest was occupied by the tubules. The interstitial space increased slightly till day 125 when it occupied 35.6 percent area against 64.4 per cent occupied by tubules. The area occupied by tubules increased in subsequent stages, leading to decreased intertubular space, thus a reverse relationship between two characteristics existed. In animals older than 300 days the interstitial space was reduced to "triangles" in between the seminiferous tubules, thus bringing the tubules adjacent to each other and creating a compactness. Such triangle formation was observed by Abdel Raouf (1960) in 4-5 month old bull ealves. The interstitial space kept reducing constantly and on Day 600 it was only 11.3 percent. Similar data is reported for adult Murrah buffalo bull by Sharma and Gupta, (1980).

Findings of the present study, till Day 125 support the report by Goyal and Dhingra (1973) that in Murrah calves interstitial space increases with age from 4 to 52 weeks. However, the interstitial space started decreasing after 200 days of age in the present study. The results clearly indicate a reduction in intertubular space with increasing age in buffalo testes, which suggests that testicular growth is primarily tubular in nature.

LUMEN FORMATION:

At birth the tubules were not patent (Fig. 1) the centre of tubules was filled with homogenous ground substance. Slight clearing of central matrix was observed in few peripheral tubules on Day 125, the lumen started appearing by 200 days in our experimental animals in many tubules. The initiation of lumen was marked by lysis of ground tissue in the centre. The central region stained very lightly. Sometimes degenerating clumps of cells staining darkly were also seen in the centre of tubules. In this study the lumen was seen earlier than the age reported by Goyal and Dhingra (1973) in Murrah buffalo calves, probably because their sampling interval was much larger than ours. In the present study clear vacuolization in biopsy samples occurred by 330 days and the lumenation continued till 480 days of age in different animals. All tubules did not undergo simultaneously lumenation, but were in different phases of canalization. Peripheral tubules under the capsule are the first to form lumen. The presence of fully formed lumen was noted by 360 days in one animal and by 450-480 days in all animals.

In puberal and sexually matured animals the lumen is lined by spermatids and spermatezea. Our results are supported by the findings of Igboeli and Rakha (1971), in beef bulls, and Aire and Akpokdje (1975) in Fulani bull calves. They reported lumen formation as late as 11 months of age. The study helped to trace the developmental stages of the testis from birth to maturity and reveal the potential for sperm formation and transport in buffalo calves much earlier than reported. There is a good scope to management and endocrine status is perhaps not adverse to bring about maturity in male calves towards fertile service by 16 months onwards.

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Cytological Changes In Male Gonads From Birth To Sexual Maturity In Buffaloes

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ABSTRACT

Cytological changes in male gonad from birth to sexual maturity in buffaloes have been described.

The male buffalo calves of Surati breed were studied for the cytological changes occurring in the developing gonad. As little literature is available on this aspect on the species. An attempt is made to know the various cell types in male calves from birth through puberty and sexual maturity. The data revealed that the species possesses the potential of early breeding.

Materials and Methods

The testicular tissue from 4 male buffalo calves of Surati breed was sampled by repeated incision biopsies (Hotchkiss, 1944), over 17 stages from birth to 600 days of age. The incision biopsies did not have any deteriorating effect on the testicular growth and develpment. The testicular tissue was fixed in Bouin's fluid, was processed as per Luna (1968) and was sectioned at 5-7 micron thickness. Staining was done with Haematoxylin and Eosin. Microscopical studies on intratubular cells and interstitial cells were made as given below:

The study of intratubular elements was done by identifying and counting their numbers in 15×45 magnification. The different types of cells were counted per cross section (c.s.) of the tubule. Overall mean of 25 seminiferous tubule c.s. per stage per animal was taken for different cell types.

The various cell types were classified as follows:

(i) Basal indifferent cells:—These are generally present close to the basement membrane and are small. The cells are of various shapes. They have few nucleoli in their nucleoplasm. These line the basement membrane singly.

- (ii) Gonocytes:—These cells are present between the small basal indifferent cell layer and also sometimes towards the centre of tubule. Their size is three to six times larger than the basal indifferent cells. These have a centrally placed nucleolus and few lightly stained, fine chromatin granules.
- (iii) Spermatogonia:—These are large cells. The nucleus is large in size and exhibits either fine chromatin (type A spermatogonia) or coarse chromatin (type B spermatogonia). Type A and type B spermatogonia were counted separately.

(iv) Spermatocytes:— Primary spermatocytes:— The spermatocytes in early leptotene

Fig. 1: Cytological changes in male gonads from birth to sexual maturity in buffaloes

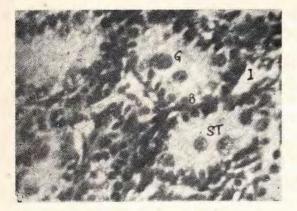


FIG. 1 High power 10×40 view of the cross section (C.S.) of seminiferous tubules (ST) of day old animal, which are separated by interstitial space (I) The solid ST shows gonocytes (G) and basal indifferent cells (B). Intestitial tissue contains mesenchynimal cells (M).

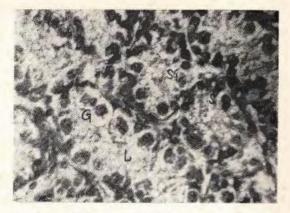


FIG. 3 Day 200 — gonocytes (G) are in division. Few spermatogonia (SG) and sertoli cells (S) are also seen. Lumen formation (L) has been initiated (10×40 magni.)

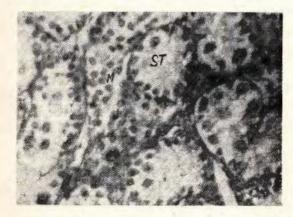


FIG. 2 Cross section of seminiferous Tubules at 125 days of age (Magnification 10×25). Interstitial space (I) occupied by mesenchymal cells (M) and fibroblasts (F).

phase resemble type B spermatogonia. The primary spermatocytes were observed in leptotene and zygotene phases of division. The cells in two phases were counted separately. Secondary spermatocytes:—

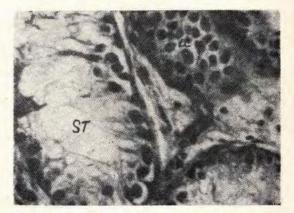


FIG. 4. CS of seminiferous tubules at 420 days of age (Magnification 10×40). Triangular interstitial space contains groups of Leydig cells (LG)

These cells have a spherical nucleus without nucleolus and have scattered chromatin mostly on nuclear envelope. These cells undergo second maturation division to produce spermatids. These were seen rarely and their number was not counted.

(v) Spermatids:—Spermatids assume various shapes round or elongated during their transformation to spermatozoa. The round spermatids are smaller than the secondary spermatocytes and appear as tiny round cells under light microscope. Round and elongated spermatids were counted separately,

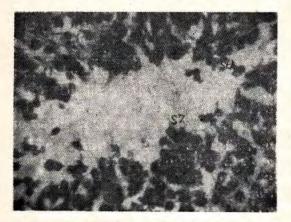


FIG. 5. High Power view of CS of seminiferous tubules of 600day old animal in active spermatogenesis. Spermatozoa (SZ) are seen attached to Sertoli cells (S) with their tails projecting into humen. Spermatids (SPD) are seen in groups.

- (vi) Spermatozoa:—Many a spermatozoa transforming from spermatids are seen in later stages of development. These are also attached to Sertoli cells in earlier stages and are later released free into the lumen of the tubule.
- (vii) Sertoli cells:—The cytoplasmic boundaries are not distinct, but the cell possesses a typical pyramidal or oval nucleus. The cell is attached vertically to the basement membrane.

Interstitial cells or cells of Leydig were studied for the morphological changes in their cytoplasm, nucleus and shape.

Results and Discussion

BASAL INDIFFERENT CELLS AND GONOCYTES:-In the present study in the newborn animals the non-patent seminiferous tubules were lined mainly by 2 types of cells-gonocytes and basal indifferent cells (Fig. 1). At birth the number of basal indifferent cells and gonocytes per tubnlar cross section was found to be 15.19 and 2.71 respectively. The number of these cells of embryonic origin kept fluctuating from day 1 to 240. Mitotic figures in both cell types were observed right from Day one. The gonocyte number was unchanged till Day 240, after that it decreased fast and on Day 390 these were only 0.2 cells per tubular c.s. From Day 420 onwards gonocytes could not be observed any more. Gonocytes were observed till 12 weeks in Swedish Red and White calves (Abdel Raouf, 1960). Goyal and Dhingra (1973) reported an increase in the number of gonocytes from 1 to 12 months of age which is just opposite to the data reported in the present study. The basal indifferent cells in the present study decreased very little till day 390, but their number got reduced drastically on Day 420 when these figured only 3.4 cells per tubular c.s. thereafter these were observed. Goyal and Dhingra (1973) also observed a trend in the number of declining basal indifferent cells, but they reported 67.3 percent cells of the type in one year old animals whereas in the present work, only 41.8 percent were bsal indifferent cells in animals of same age.

SPERMATOGONIA:—In the present study the primary spermatogonia was first observed at the age of 200 days (Fig. 3). At 300 days of age the number of spermatogonia increased to 4.3 per tubular cross section. The cell number kept increasing constantly, so as to reach a number of 26.7 cells per tubular c.s. at 17 months of age. In Swedish Red and White bull presence of spermatogonia was noted as early as 8 weeks' age (Abdel Roouf, 1960). Type B spermatogonia were noticed at the age of 330 days, with highest number (8.29 cells per tubular c.s.) on Day 420. Thereafter it fluctuated.

SPERMATOCYTES:-The primary spermatocytes appeared at the age of 330 days. Although the average number of cells increased consistently, prominent raise in cell number was noted Day 510 and Day 600. The on spermatocytes (including leptotene and zygotene both types) averaged about 100 cells per tubuler c.s. on Day 600. Contrary to the present study Goyal and Dhingra (1973) did not observe any spermatocytes in one year old Murrah calves. In cattle quite early appearance of spermatocytes has been reported by Santamarina and Reece, 1957 (21 months) and Abdel Raouf, 1960 (5 months) slightly late appearance is reported by Hooker, 1944 (6 months) and by Igboeli and Rakha (1971), 9 months.

SPERMATIDS AND SPERMATO-ZOA:—In the present study round spermatid was first noted at the age of 450 days. The elongated spermatid appear one month later. The spermatozoa were first observed at the age of 510 days. A gradual increase in the number of spermatids and spermatozoa was noted till Day 600 when these were in abundance in the seminiferous tubules (Fig. 5). spermatogenesis in cattle is reported at an earlier age by Knudsen, 1954 and Abdel Raouf, 1960 (7 months), Hooker 1944 (8[‡] to 9[‡] months) and Igboeli and Rakha, 1971 (11 months). Residual bodies during spermiogenesis were observed in large number in late developmental stages-around 540 days and onwards.

SERTOLI CELLS:—In the present study the Sertoli cells could be identified right from Day one, their number and size increased from birth to puberty. In day old animals the number of Sertoli cells was less than one which increased to reach 13.5 cells per tubular c.s. on Day 600.

mesenchymal Cells: - The Leydig cells started transforming from Day 125 onwards (Fig. 2). Metamorphosis involved increased cell and nuclear size, acquiring nucleoli, losing the processes appearance of granules in the cytoplasm and vacuolization of cytoplasm in late developmental stages (Albert, 1961 and Hooker, 1970). Till Day 360 more and more Leydig cells were being converted from mesenchymal cells. At this stage the interstitial space was reduced to "Triangles" between tubules, the Ledyig cells in these spaces were in the form of pockets, sheaths, rows or singly also. Our data verifies the results of Goyal observed and Dhingra (1973) who Leydig cells around 15-16 weeks in Murrah calves. Fully developed secretory cells in large number appeared by 390 to 420 days (Fig. 4). From the foregoing data it is evident that the establishment of spermatogenesis is a very long and progressive phenomena. Based on the above data, in buffalo the whole process can be categorised as below:

- 1. Impuberal phase—when only gonocytes and basal indifferent cells are present. The phase stretches from birth to 200 days postnatally.
- 2. Prepuberal phase-from Day 200 to 420 when spermatogonia appeared

and gave rise to spermatocytes mitotically. Lumenation was initiated and the Leydig cell number increased greatly during this period.

- 3. Puberal phase—from Day 450 to 510. The spermatids and spermatozoa were observed during this stage along with fully active Leydig cells.
- 4. Postpubertal phase—from 510 onwards. It evinced a great increase in

the number of various spermatogenic elements.

It can be safely concluded that contrary to popular misconcept, the buffalo male matures at a comparatively early age and can be used successfully for breeding to facilitate extensive use of proven sires in a shorter span and the life time use can also be more.

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Antigenic Analysis Of Spermatozoa And Seminal Plasma of Exotic and Indigenous Rams By Geldiffusion and Immunoelectrophoresis

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ABSTRACT

Analysis of Antigens of seminal plasma and spermatozoa of Chokla (10), Russian Merino and Rambouillet (6 each) rams by gel-diffusion and immunoelectrophoresis revealed the presence of three precipitin lines in chokla spermatozoa and 5 to 6 in seminal plasma antigens of all rams, against their hyperimmune serum raised in rabbits. Two common antigenic components between seminal plasma and spermatozoa were observed in chokla rams, whereas the seminal plasma of Russian Merino and Rambouillet showed three common antigens against the hyperimmune serum raised in rabbits. The immunoelectrophoretic studies further indicated that the seminal plasma of Russian Merino, Rambouillet and chokla rams with their hyperimmune serum gave 13,14 and 10 precipitin lines, whereas the antigens of chokla spermatozoa gave only 3 precipitin lines. The study suggests that some antigenic differences exist in the spermatozoa and seminal plasma of different breeds of rams.

* *

The present study was aimed to study the fractionization of the semiral plasma and spermatozoal antigens by double gel diffusion and immunoelectrophoresis in Russian Merino, Rambouillet and chokala rams.

Materials and Methods

male Russian Healthy Merino, Rambouillet (6 each) and chokla rams (10) were selected for the study. The semen was collected in the morning hours and the seminal plasma and the spermatozoa were separated by centrifuging and washing three times with PBS buffer (0.1M pH 7.00). The antigens were kept at -20°C pending antigenic analysis. The procedure of Cruickshan-K. (1965) with some modifications was followed for the preparation of agar gel (Noble agar Difco 1%). The motten agar 4 C.C. was poured over the clean microslides and allowed to solidify. The agar gel slides were kept in refrigerator immersed in veronal buffer (pH 8.6) for subsequent use. The wells were made with the help of a pipette attached to a vaccum pump. The bottom of the wells was sealed with a drop of motten agar to prevent the sepage. The wells were charged with the following material for gel diffusion.

Central wells: (i) Seminal plasma antigen of chokla, Rambouillet and Russian Merino rams (ii) Spermatozoal antigen of chokla rams.

Peripheral wells: (i) Rabbit hyperimmune serum prepared against the seminal plasma of chokla, Rambouillet and Russian Merino rams in three peripheral wells and one well was kept as control.

Group		Rabbit number	Total No. of Inj. given	Interval	Route	Nature and amount of antigen	
	r	1493	14	Alternate	Subcutaneo-	Rambouillet pooled Se	minal
Rambouillet I	1			days	usly	plasma —	0.2 ml
	l	1494	14	**	>>	Incomplete Fruend's	
						Adjuvant -	0.2 ml
	Y					Antibiotic mixture -	0.1 ml
	r	1497	73	35	>7	Russian Merino	
Russian Merino II	1					Pooled seminal	
Merino II	l	1490	22	37	33	plasma —	0.2 ml
						Incomplete Fruend's	
						Adjuvant-	0.2 ml
						Antibiotic mixture -	0.1 ml
	r	1495	>>	33		Chokla pooled	0.2 ml
Chokla III	1					seminal plasma	
	L	1496	>>		33	Incomplete Fruend's	
						Adjuvant -	0.2 ml
						Antibiotic mixture	0.1 ml
Chokla		1491	7	22	22	Chokla Washed	
						spermatozoa —	0.4 ml
						Incomplete Fruend's	
				-		Adjuvant	0.4 ml
						Antibiotic mixture	0.2 ml

TABLE: THE SCHEDULE OF INJECTIONS GIVEN FOR THE PREPARATION OF RABBIT HYPER IMMUNE SERUM AGAINST SPERMATOZOA AND SEMINAL PLASMA

N.B. A pause of 7 days was given after the last injection before collecting the blood from each group.

ii) Rabbit hyperimmune serum prepared against chokla ram spermatozoa in three peripheral wells and one well as control.

After charging with respective antigens and hyperimmune serum, the slides were transferred to a moist chamber for 7 days for the diffusion to occur. Every day the slides were examined for the appearance of any precipitin lines. In some cases the agar wells were again charged with the antiscra and antigen to hasten the precipitation.

Immunoelectrophoresis: The following combinations were used:

- i) Spermatozoa antigen of chokla ram with its hyperimmune serum.
- ii) Electrophorised seminal plasma antigens of chokla, Rambouillet and Russian Merino rams with

their corresponding hyperimmune serum.

Preparation of hyperimmune serum:

Albino rabbits were used to produce antibodics against the seminal plasma and spermatozoa antigens. The schedule of immunization for each group of rabbits is shown in Table 1. 5 ml of blood from each group of rabbits were collected from the ear vein. Serum was separated and kept in screw-capped vials at -20°C pending antigenic analysis.

Washing, staining and drying of slides:

Antigen solution was leached from the agar gel by soaking in a saline solution (2% pH 7.4) and kept undisturbed to avoid floating of agar for 2 days. The slides were stained with bromophenol blue solution (0.1%) and then destained with acetic acid solution (5%).

Results and Discussion

Analysis of antigens of seminal plasma and spermatozoa of chokla, Russian Merino and Rambouillet by gel diffusion and immunoelectrophoresis.

- 1. The chokla, spermatozoal antigen with its corresponding rabbit hyperimmune serum produced three precipitin lines whereas the seminalplasma antigen produced five to six precipitin lines.
- 2. The antigens of seminal plasma of Russian Merino and Rambouillet rams produced five precipitin lines with their hyperimmune serum produced in rabbits.
- 3. Two common antigenic components between seminal plasma and-spermatozoa were observed by gel diffusion in chokla rams, whereas three common antigenic precipitin lines among the seminal plasma of chokla, Rambouillet and Russian Merino were observed. The reaction indicated that the common components were more between Russian Merino and Rambouillet than chokala rams.
- 4. The Immuno-electrophoretic studies further indicated that the seminal plasma of Russian Merino, Rambouillet and chokla rams with their hyperimmune serum gave 13,14 and 10 precipitin lines, whereas the antigen of chokla spermatozoa gave only 3 precipitin lines.

Our results are similar to Dikov and Tornov (1970) who observed two antigens in the ram spermatozoa. However, Kulangara (1969) observed 6 to 11 seminal plasma antigens and 5 to 10 in spermatozoa antigens in different individuals of Black-face and Merino rams, and 12 precipitin lines in semen and 10 in seminal plasma in indigenous rams (Agar, 1965) whereas, Harthaway and Hartree (1963) reported 4 antigens in the extracts of spermatozoa in rams. The variations in the antigenic components may be due to the existance of difference in the antigenecity of spermatozoa in rams (Hunter, 1963).

The double gel diffusion and immunoelectrophoresis of the spermatozoal and seminal plasma antigens of the different breeds of rams (exotic and indigenous) revealed the different seminal antigen pattern in rams and reflects that genetic difference among the different breed of the same species. Immunologic finger prints can threfore be obtained of the ram semen for the first time, which could be of great importance in breeding practices.

Acknowledgement

The authors acknowledge the I.C.A.R., New Delhi for providing financial assistance and the Dean, College of Veterinary and Animal Sciences, Bikaner for providing necessary facilities. We are also very much thankful to Dr. P. R. Jhatkar for the advice and technical help in conducting this study.

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Plasma Testosterone During Oestrous Cycle And Postpartum Period Among Buffaloes

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ABSTRACT

Investigating buffalo endocrinology during oestrous cyclicity and postpartum period, plasma testosterone (T) was estimated in jugular plasma using RIA techniques. During oestrous cycle, blood samples were collected daily and at regular 4 hourly intervals, 3 days before and three days after behavioural oestrus. For postpartum animals, analysis was carried out from sample drawn every 7-8 days postpartum. The mean values of testosterone from different animals on the day of oestrus was 0.054+0.0206 ng/ml. The mean testosterone levels remained same up to day 6 of oestrous cycle and then rose gradually upto day 9 during diestrus, with values of 0.1035+ 0.0424 ng/ml. Testosterone levels were 0.110, 0.093, 0.080 ng/ml on day 3,2 and 1 before oestrus. Individual animals showed conspicuous pulses during oestrous cycle. In all the animals the levels of testosterone were high during luteal phase with a number of variable peaks. The mean concentration of testosterone during the post partum period was 0.038, 0.063, 0.038, 0.074 and 0.038 ng/ml on day 7, 15, 22, 30, 37 and 45, respectively. The mean postpartum testosterone level was not significantly different over this period.

and South East Asian countries is predominantly buffalo oriented. However, information on hormonal factors responsible for ocstrus and ovulation is scarce particularly in light of the fact that the problems related to buffalo reproduction, such as late maturity, repeat breeding, delay in post-partum estrus and anestrum, limit the productivity of the animal. Limited information regarding the peripheral plasma progestercne, estrogens and LH profile is available for buffaloes during cyclicity (Bachlaus et al., 1980; Arora and Pandey, 1982) and early post-partum (Perera et al., 1981; Naqvi, 1982; Prakash, 1983).

Evidence is now available that testosterone, besides being a precursor for oestradiol, is also involved in triggering the mechanism for the onset of luteolysis in the cow. Peaks of testosterone have been recorded around days 11-14 of the oestrus cycle in cows (Shemesh and Hansel, 1974; Herriman *et al.*, 1979; Kesler *et al.*, 1979) and also preceding oestrus behaviour (Shemesh and Hansel, 1975; Kanchev *et al.*, 1976; Kesler *et al.*, 1979). The objectives of the present study were to determine the plasma testosterone profile in buffaloes during cyclicity and early post-partum.

Materials and Methods

Five regular cycling Murrah buffalo heifers ranging from 42 to 54 months of

The dairy industry in many South

age, and five pregnant Murrah buffaloes in their second or third lactations were selected from the National Dairy Research Institute herd for experimentatiou. All animals were maintained under uniform feeding and managemental conditions existing in the herd. Estrus detection for cycling heifers was carried out by trained animal attendants with the help of a teaser bull paraded three times over 24 hours. Behavioral symptoms of heat, including the presence of vaginal mucus was confirmed by rectal palpation and examination.

TABLE 1: Testosterone entiserum characterisation

Steroid	% cross reactivity			
Testosterone		100		
$5 - \alpha - dihydroxytestosterone$		6.65		
Androstenedione	-	1.80		
11-B-hydroxyandrostenedione		0.00		

Blood samples were collected at noon daily by jugular vein puncture in heparinized tubes from each heifer for a minimum of two estrous cycles. The pregnant buffaloes were allowed to calve and jugular blood samples were collected from them subsequently on days 7, 15, 22, 30, 37 and 45 postpartum. All blood samples thus collected were centrifuged within 1/2 hr of collection, plasma separated and stored in deep freeze at -20° C till further analysis for testosterone by radioimmunoassay (RIA)

RIA of testosterone: Testosterone was estimated by RIA standardized in the laboratory.

Duplicate 0.2 ml of plasma aliquots were taken in 12×75 mm tubes and extracted with 2 ml of ether for 1 min twice. The lower aqueous layer was frozen in an ice-salt mixture and upper organic layer was decanted in 12×75 mm tubes (Corning). The ether was evaporated to dryness in a waterbath at 35-40°C. The residue was dissolved in 0.2 ml of PBS (pH 7.5, 0.1M), vortexed and this was followed by addition of 0.1 ml of testosterone antiserum used at a

TABLE 2:	Mean \pm SEM plasma testosterone
	ng/ml during estrous cycle of
	buffalo heifers.

Days from estrus	Testosterone (ng/ml)
-4	0.080±0.010
-3	0.110±0.042
-2	0.093 ± 0.030
-1	0.080 ± 0.010
0	0.54 ± 0.020
1	0.072 ± 0.023
2	0.066±0.022
3	0.053 ± 0.021
4	0.054±0.020
5	0.059 ± 0.010
6	0.071 ± 0.030
7	0.080 ± 0.032
8	0.100±0.040
9	0.103±0.042
10	0.052 ± 0.020
11	0.092 ± 0.031
12	0.053±0.021
13	0.072 ± 0.023
14	0.061 ± 0.050
15	0.071±0.030
16	0.074±0.010
17	0.073±0.011

dilution of 1:60,000 and then 0.1 ml of testosterone tracer having 10,000 CPM was added in each tube. The resulting mixture was vortexed and incubated at 4° C in refrigerator overnight. Following incubation, 0.5 ml of fresh prepared activated charcoal-dextran suspension (1% activated characoal, 0.5% dextran) under constant stirring condition with magnetic stirrer, was added to each tube, vortexed and centrifuged at 4°C just after charcoal addition at 4°C in ice water. The supernatant containing the bound testosterone was decanted into scintillation vials and counted in 5 ml of scintillation fluid (1000 ml toluene, PPO-4 g, POPOP-0.1 g). In addition to above unknown tubes, four other sets of tubes were also run with each assay, as follows:

i) Standard tubes containing 0.1 ml of series of concentration in duplicate.

ii) Blank tubes in duplicate containing 0.3 ml PBS and 0.1 ml tracer (10,000 cpm) to observe non-specific binding of charcoal separation.

iii) Four tubes containing 0.1 ml tracer and 0.1 ml antisera and 0.2 ml PBS to obtain maximum binding of tracer by the antibody.

iv) Duplicate tubes containing 0.1 ml of tracer, diluted with 0.3 ml PBS to obtain total counts of tracer added.

The sensitivity of the assay, defined as the lowest amount of hormone that can be significantly detected from zero concentration, was found to be 5 pg/tube. The Accuracy of the method estimated by determining the recovery of 50, 125, 250 and 500 pg of testosterone added to charcoal stripped plasma, was found to be 105, 92, 105 and 102% respectively. The cross reactivity of the antisera with different steroids indicated high specificity for testosterone (Table 1). The intra and interassay coefficient of variation for the method was 5.0 and 8.9% respectively.

Results

The mean testosterone levels in cycling heifers from day 4 pre-oestrus upto day 17 post-oestrus (day of oestrus being depicted as day 0) is presented in Table 2. The mean testosterone levels in buffaloes on days 7, 15, 22, 30, 37 and 45 postpartum is presented in table 3.

The mean testosterone value recorded at oestrus was 54 pg/ml and remained relatively constant upto day 5 of the oestrous cycle rising gradually to 103 pg/ml ($P \angle 0.05$) on day 9 of the cycle. Thereafter the mean hormone level again dropped ($P \angle 0.05$) and fluctuated between 52 and 92 pg/ml.

The mean postpartum testosterone levels among buffaloes remained low on days 7, 15 and 22 postpartum showing a small though non-significant ($P \angle 0.05$) increase on days 30 and 37.

Discussion

The results for testosterone pattern in cycling Murrah buffaloes indicate an increase in the jugular testosterone levels on day 9 of oestrus which roughly coincides with the period when the luteolytic activity has set in. Similar observations have also been recorded for in cattle though the timing of the peak was slightly different (Shemesh and Hansel, 1974; Herriman *et al.*, 1979; Kesler *et al.* 1979). However, no testosterone peak

TABLE 3: Postpartu	m Testosterone	(Pg/ml)	among	buffaloes.	
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Animal		Days Postpartum					
No.	7	15	22	30	37	45	
231	0.052	0.142	0.069	0.19	0.17	0.050	
551	0.034	0.039	0.035	0.037	0.062	0.039	
97	0.034	0.029	0.019	0.034	0.027	0.019	
1168	0.033	0.041	0.028	0.056	0.038	0.044	
Av.±SEM	0.038±0.005	0.063 ± 0.027	0.038±0.011	0.079±0.037	0.074 ± 0.033	0.038±0.007	

preceding oestrus was obtained in buffaloes, which was in contrast to the observations of Shemesh and Hansel (1975), Kanchev et. al. (1976) and Kesler et al. (1979). In the absence of any information in buffaloes, this variance could be attributed to the species difference. The present observations is suggestive of testosterone playing a possible role in luteolysis in buffaloes

The post-partum testosterone pattern in buffaloes did not confirm to any distinct trend. However, the small though nonsignificant increase in the hormone level on day 30 and 37 postpartum might probably have been due to the fact that 3 of the 5 animals had started showing luteal activity during that period. A more detailed close sampling than the one carried out in this study for postpartum buffaloes will therefore be more helpful in evaluating the testosterone profile during the postpartum period.

Acknowledgement

The authors are indebted to Dr. I.S. Verma, Director, N.D.R.I., Karnal for providing the necessary facilities to conduct this research. They are also grateful to Dr. P.N. Rao, Director, Department of Organic and Biological Chemistry, South-West Foundation for Research and Education, Texas, U.S.A. for his generous gift of testosterone antiserum.

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Serum FSH Levels During And After Synchronized Oestrus Of Murrah Buffaloes (Bubalus bubalis)

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ABSTRACT

Circulating FSH profile of 9 Murrah buffaloes during synchronized oestrus with MGA feeding and subsequent oestrus were investigated. Only 3 buffaloes exhibited peak FSH levels during synchronized oestrus. Mean basal FSH levels recorded on different days after estrus fluctuated between 7.79 ± 1.38 and 24.93 ± 3.29 ng/ml. All the 9 buffaloes returned to next estrus, but only 4 buffaloes were available for studying serum FSH profile.

Melengestrol acetate (MGA), a potent progestagen has been attempted for oestrus synchronization in buffaloes by a few workers (Shukla *et al.*, 1971; Mithuji *et al.*, 1972). A high percentage of treated animals show oestrous behaviour during the short period following withdrawal of MGA but fertility at the synchronized oestrns has been reported to be very low. The present investigation was undertaken to study the circulating Follicle Stimulating Hormone (FSH) profile of MGA fed buffaloes during synchronized estrus and subsequent oestrus.

Materials and Methods

Animals: Nine healthy Murrah buffaloes, 30 days after calving, were selected from the University Farm and were divided randomly into 2 groups. Groups 1 (5 animals) and 2 (4 animals) were fed with 1 and 2 mg/animal/day melengestrol acetate 60 premix (MGA), respectively, for 18 days continuously. All the animals were closely watched for oestrous behaviour before and during MGA feeding, and following MGA withdrawal. This was done, thrice daily using a trained teaser bull. The animals which were detected in oestrus, were put to natural service.

Blood sampling: Blood samples were collected 24 h before the start of MGA feeding, on days 7 and 14 of the MGA feeding and 24 h after the withdrawal of the treatment. Further blood samples were collected at the time of detection of oestrus (0 h) and thereafter every 4 h for 24 h, followed by sampling on every third day up to 23 days. Serum was separated by centrifugation at 3,000 rpm (1000 g) for 30 min at 5°C and stored frozen at -20° C till used for assay.

FSH assay: The double antibody radioimmunoassay, using a complete bovine system, was followed for measuring serum FSH levels. Details of the technique were reported earlier (Razdan *et al.*, 1982).

Results and Discussion

The mean plasma FSH levels of buffaloes in Groups 1 and 2 prior to start and during treatment are presented in Table 1. FSH levels during treatment ranged from 8.41 ± 2.33 to 15.73 ± 4.22 ng/ml serum among two groups, which are considered as fluctuating basal levels.

Time	Group 1 (1 mg)	Group 2 (2 mg)
24 h pre-treatment	14.65±2.60	12.65 ± 3.55
7 days after the start of treatment	8.41 ± 2.33	12.54 ± 7.70
14 days after the start of treatment	14.85 ± 3.39	15.73±4.22

Table 1. Mean (±S.E.) serum FSH levels before and during MGA feeding.

These values were not different from the pre-treatment values. Serum FSH values of individual buffalo for the first 24 h after the detection of synchronized estrus under both gronps are given in Table 2. The mean values observed on different days after estrus detection are presented in Table 3. In group 1, only 1 out of 5 animals showed peak FSH levels of 69.93 ng/ml at oestrus detection. In group 2, only 2 out of 4 animals exhibited high FSH values. Buffalo No 7. showed highest concentration of 73.26 ng/ml at the oestrus detection. Buffalo No. 6 had 48.28 ng/ml at 4 h of estrus detection (Table 2).

The mean FSH levels measured on different days after oestrus fluctuated within normal range of basal FSH levels when compared to normal cycling Murrah buffaloes reported earlier (Razdan *et al.*, 1982).

All the 9 animals returned to next oestrus but blood samples from only 4

TABLE 2. Serum FSH concentration (ng/ml) at different intervals after detection of synchronized oestrus-

Time after detection of cestrus		-		Buff	alo No.			-	
	Group 1 (1 mg)					Group 2 (2 mg)			
(h)	1	2	3	4	5	6	7	8	6
-48 to	4.99	15.56	21.64	11.32	22.31	15.32	13.32	27.64	11.65
72									-
0	69.73	27.97	-	20.97	19.31	17.65	73.26	12.32	11.09
44	39.96	16.65	28.64	15.32	18.65	48.28	6.60	8.899	9.66
8	18.98	19.58	18,32	13.32	18.65	24.38	_	20.65	16.65
12	8.66	11.32	16.65	15.32	15.98	15.32	-	17.98	18.85
16	10.65	6.99	3.99	10.65	26.64	18.98	-	13.32	10.65
20	10.65	14.65	13.99	14.32	16.65	16.98	36.63	14.98	11.09

 TABLE 3. Mean (±S.E.) serum FSH concentration at different intervals during the synchronized oestrus cycle.

Time after detection of estrus (days)	Group 1 (1 mg)	Group 2 (2 mg)
2	12.05+2.29	13.73±2.86
5	12.78 ± 2.73	24.93 ± 3.29
8	10.72 ± 1.70	17.73±1.77
11	7.79±1.38	16.56 ± 1.77
14	8.38 ± 1.33	14.48±2.18
17	0.49 ± 1.93	20.97±6.61
20	9.34±0.76	20.16 ± 6.46
23	8.10±0.18	22.47 ± 5.82

Time after detection		Buffalo No) .		
of oestrus	Group	1 (1 mg)	Group 2 (2 m		
(h) ·	1	2	7	8	
0	10.32	21.98	16.65	73,26	
4	7.49	23.98	14.65	63.27	
8	66-60	17.32	17.64	16.65	
12	34.96	16.65	29.30	18.32	
16	18.65	11.65	21.64	11.99	
20	14.65	13.32	16.98	14.65	
24	6.99	10.98	11.65	-	

TABLE 4. Serum FSH concentration (ng/ml) at four hourly intervals after detection of estrus during oestrus subsequent to synchronization.

TABLE 5. Mean $(\pm S.E.)$ serum FSH concentration at different intervals after detection of oestrus during oestrous cycle subsequent to synchronization.

Time after detection of estrus (days)	Group 1 (1 mg)	Group 2 (2 mg)
2	11.15±1.49	19.47±0.83
5	13.48 ± 3.16	14.65 ± 0.50
8	8.40±2,57	16.81 ± 3.16
11	4.66±0.50	10.98 ± 1.66
14	7.32 ± 0.06	15.31 ± 4.99
17	10.32 ± 0.66	13.98 ± 0.66
20	9.49±2.07	12.40 ± 5.24
23	6.63 ± 0.69	9.66 ± 0.00

animals could be collected for FSH assay. The FSH levels observed during first 24 h of oestrus are given in Table 4 and the mean values observed on different days are given in Table 5. Buffalo No. 1 showed peak FSH level of 66.60 ng/ml at 8 h of oestrus detection. Buffalo No 2 did not show marked elevated FSH values during first 24 h of estrus, although FSH levels during first 4 h were slightly higher than rest of the period. It is possible that FSH peak value might have been missed due to late detection of oestrus in this animal. Buffalo No. 8 showed FSH peak level of 73.26 ng/ml at oestrus detection. This animal had not shown any peak level during synchronized oestrus.

Buffalo No 7 had elevated FSH levels of (29.30 ng/ml) at 12 h of oestrus detection than at other hours of estrus (Table 4).

The mean FSH values observed during different days after estrus detection were obviously basal levels. All the 4 buffaloes conceived at this oestrus.

None of the buffaloes conceived at the synchronized oestrus. Out of the 9 buffaloes in two groups only 3 buffaloes showed peak levels of FSH during estrus. This suggested that the inhibitory effect of high level of progesterone continued for some time after withdrawal of MGA. This prolonged inhibitory effect may be responsible for the suppression of FSH peak in most of the buffaloes which in turn could be one of the contributing factors responsible for poor fertility at the first synchronized oestrus. Lamond et al. (1971) and Britt and Ulberg (1972) recorded higher concentration of progesterone during follicular phase in MGA treated cows than the levels observed in the controls. Poor fertility at the synchronised estrns in cows had also been recorded by Zimbelman (1966) and Randel et al. (1972).

Acknowledgement

This work was carried out under a

research project sponsored by the Indian Council of Agricultural Research, New Delhi. We thank Drs. L.E. Reichert and D.J. Bolt for the generous gift of standard FSH and RIA grade FSH, respectively. Our thanks are also due to the Department of Livestock Production and Management, Haryana Agricultural University, Hissar, India, for allowing to collect blood samples from the experimental animals.

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Studies On Some Factors Affecting Gestation Period In Cattle

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ABSTRACT

Effect of heritage, parity, sex and birth wieght on gestation period were studied in 1401 calves. For correlation and regression studies 814 pairs of gestation period and birth weight were analysed. The calves with 50% Brown Swiss heritage were significantly heavy $(\frac{1}{B} + H + G)$ $(P \angle 0.01)$ at birth $(30.71\pm1.17 \text{ kg})$ and carried for the maximum duration in the womb (284.27+1.41 days) as compared to all other groups. The male calves (23.94±0.23 kg) were heavy in weight and carried in the womb for longer duration than the female (278.17 +0.22 days). The sex ratio was 50.25%. There was no significant effect of parity of dam on birth weight and gestation period. Correlation between gestation period and birth weight of calves was 0.316, which was significantly positive (P ∠ 0.01).

The length of gestation in cattle is considered from fertile service to parturition. Number of factors are said to influence the gestation period. Systematic studies pertaining to the factors influencing gestation period in cattle appears to be lacking (Salisbury *et al.*, 1978). The present investigation was aimed to envisage the effect of breed, parity, sex and birth weight on the gestation period in cattle.

Materials and Methods

The study was conducted at the All India Co-ordinated Research Project on Cattle, Jabalpur, It included indigenous Gir (G) cows as foundation stock. Frozen semen of Holstein Friesian (H), Jersey (J) and Brown Swiss (B) exotic breeds procured from U.S.A. was utilised for cross-breeding. After the desired number of $\frac{1}{2}$ and $\frac{1}{4}$ exotic crossbred progenies were generated, *inter-se* mating was followed using liquid semen of $\frac{3}{4}$ th exotic cross-bred bulls on the same genetic group of females. Close mating was avoided.

Total 711 cows were employed to generate 1401 claves in various genetic groups. In all, 814 calves were weighed immediately after birth. The number of calves studied in each genetic group are presented in Table-1. For correlation and regression studies, 814 pairs of gestation period and birth weight were analysed statistically.

Results and Discussion

The mean gestation period in the herd was 277.69 ± 0.16 days with a range of 275.13 ± 0.38 to 284.27 ± 1.41 days in different genetic groups (Table 1) and parities (Table-2). There was a significant difference (P $\angle 0.01$) in the gestation period between various genetic groups (Table-3), the minimum in

Geneti	c group	Male ca	lves	Female	calves	Overall n	nean	Sex
Dam	Calves	Duration in womb(days)	Birth weight (kg)	Duration in womb(days)	Birth weight (kg)	Duration in womb(days)	Birth weight (kg) (N	ratio Iale%
G	f]fC	278.55±0.58 (101)	17.68±1.16 (17)	278.16±0.50 (106)	20.36±0.44 (68)	278.35±0.38 (207)	19.82 ± 0.43 (85)	48.79
G	<u></u> ¹ ₂ H ¹ ₂ G	277.70±0.37 (209)	22.61±0.96 (87)	276.90±0.39 (226)	23.96±0.45 (53)	277.29±0.27 (435)	23.13±0.29 (140)	48.04
₽HĨC	∦] ∦Н ‡ G	277.80±0.52 (126)	22.96±0.42 (95)	277.76±0.50 (108)	22.19±0.36 (103)	277.78±0.36 (234)	22.56±0.28 (198)	53.84
į́Нł́С	³ B ¹ H ¹ G	281.07±0.56 (108)	27.66±0.47 (89)	279.31±0.57 (110)	26.50±045. (75)	280.18±0.40 (218)	27.13±0.36 (164)	49.54
₫.J ł G	[‡] H [‡] J [‡] G	076.15±0.49 (129)	23.66±0.42 (92)	274.02±0.58 (113)	22.48±0.37 (89)	275.13±0.38 (236)	23.08±0.28 (181)	52.11
iliHiG	≱Jł́Нł́С	277.44±1.42 (16)	20.56±1.34 (8)	276.83±1.33 (12)	19.91±1.06 (11)	277.18±1.00 (28)	20.18±0.81 (19)	57.14
[‡] H [†]] [†] C	₽HŢĴŦĊ	278.64±1.09 (14)	22.45±1.29 (11)	276.00±1.64 (14)	23.78±1.70 (9)	277.32±1.00 (28)	24.15±1.02 (20)	50.00
₿₽₽₽₽₽	₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽₽	284.86±2.02 (7)	30.60±1.66 (5)	283.75±2.08 (8)	31.00 ± 1.00 (2)	284.27±1.41 (15)	30.71±1.17 (7)	46.66

TABLE: 1 Gestation period and birth weight of male and female calves with sex ratio in different genetic groups (Mean±SE).

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Note: Figures in parenthesis indicate number of calves studied in each group.

TABLE 2:	Gestation	period a	und birth	weight of	male and	female calves	with sex	ratio in different
	parities	(Mean±S	SE).					

	Ma	ale calves	Female	calves	Overall m	nean	Sex ratio
Parity	Duration in womb (days)	Birth weight (kg)	Duration in womb (days)	Birth weight (kg)	Duration in womb (days)	Birth weight (kg)	(Male %)
İst	277.95±0.98 (295)	22.95±0.33 (213)	277.14±0.38 (286)	23.36±0.31 (194)	277.55±0.27 (581)	23.14±0.23 (407)	50.94
2nd	278.18±0.33 (244)	25.15±0.42 (104)	277.92±0.36 (227)	22.37±0.99 (130)	278.05±0.24 (451)	23.61±0.30 (234)	54.10
3rd	278.90±0.51 (116)	24.84±0.58 (60)	276.11±0.57 (117)	23.25±0.59 (55)	277.50 ± 0.40 (233)	24.08±0.42 (115)	49.78
4th	278.29±0.63 (62)	25.44±0.82 (25)	277.02±0.75 (57)	22.87±0.68 (26)	277.68±0.49 (119)	24.13±0.56 (51)	52.10
5th	27 4,00 ±1.54 (7)	21.50 ± 1.50 (2)	276.20±2.40 (10)	22.20 ± 1.89 (5)	279.29±1.53 (17)	22.00±1.35 (7)	41.17
Overall mean	278.17±0.22 (704)	23.94±0.23 (404)	277.20±0.23 (697)	22.98±0.21 (410)	277.69±0.16 (1401)	23.46±0.16 (814)	50.25

Note: Figures in parenthesis indicate number of calves studied in each group.

 $\frac{1}{4}H_{4}J_{4}G$ and maximum in $\frac{1}{2}B_{4}H_{4}G$ calves. The gestation period for which the male calves were carried $(278.17\pm$ 0.22 days) was significantly more (P \angle 0.01) than those of female (277.20 ± 0.23) days). The mean sex ratio ranged between 41.17 to 57.14% amongst various genetic groups and parities with overall mean as 50.25%. The calves were carried longest in the second parity of dam (278.05+0.24 days) and for shortest duration in the fifth parity (275.29 \pm 1.53 days). However, the effect of parity on gestation period was nonsignificant.

Comparable results with respect to genetic groups studied in the present investigation in relation to gestation period were not available. However, Salisbury et al. (1978) described gestation length for various dairy breeds to vary from 278 to 284 days with maximum gestation length in the Brown Swiss breed as found in the present investigation also. Jondet and Arias (1973) believed that breed of sire had a significant effect on gestation length. As observed in the present findings, Manrique and Wilcox (1978) also reported that the gestation length of the cows carrying male foetuses was longer than those carrying female foetuses. Chapman et al. (1938) described

that the sex ratio in female calves is variable, but the trend is towards a higher proportion of males in the younger foctuses. This evidence suggests that more males than females are conceived, but a greater embryonic death rate among males tends to narrow the ratio by the time of birth. Fisher and Williams (1978) observed that duration of pregnancy increases with increasing parity. However, in the present study no such relationship could be established as the effect of parity on the gestation length was found to be non-significant.

The mean birth weight of the calves was 23.46+0.16 kg with a range of 19.82 +0.43 to 30.71+1.17 kg. The lightest calves were in \$ JIG group, while heaviest in 1B1H1G. A significant difference $(P \angle 0.01)$ was observed in the birth weight calves from various genetic groups. A significant difference (P ∠ 0.01) was also observed in the weight of male (23.94+0.23 kg) and female (22.98 ± 0.21) kg) calves. The weight of the calves born to primiparous cows (23.14+0.23 kg) increased upto 4th parity (24.13±0.56 kg), which declined in the 5th parity cows $(22.00\pm1.35 \text{ kg})$. However, the difference was non-significant.

The calves with 50% Brown Swiss heritage $(\frac{1}{4}B\frac{1}{4}H\frac{1}{4}G)$ were found heaviest

Trait	Source of Variation	df	MS
Gestation Period	Between genetic group	7	532.20**
	Between parity	4	43.94
	Between sex	1	333.00**
	Error	1389	32.21
Birth weight	Between genetic group	7	587.89**
	Between parity	4	31.96
	Between sex	1	187.96**
100	Error	802	15.30

TABLE 3: Analysis of variance for gestation period and birth weight of calves in relation to various genetic groups, sex and parity.

** P < 0.01

in the present study. Shrivastava et al., (1978) also reported mean birth weight of Brown Swiss half-breds to be more than Holstein and Jersey half-breds. It is obvious that the birth weight of calves vary from breed to breed and within breed from dam to dam. Rathore (1949) pointed out that the heterosis of the dam was accomplished by heavy weights of calves at birth but the heteroiss of the calf itself was associated with greater gain in the later life.

The correlation between gestation period and birth weight was 0.316, which was significantly positive (P \angle 0.01) with the regression value of birth weight on gestation period being 0.79. It was evident from the observation that the calves belonging to $\frac{1}{4}B\frac{1}{4}H\frac{1}{4}G$ genetic group were heaviest and carried in the womb for longest period as compared to all other groups. Similarly, the male calves were heavier than female and carried longer. The lightest calves were delivered during 5th parity, hence, minimum gestation period as compared to other parity groups. The findings are in close agreement to those of Zavertyaev (1979).

Acknowledgement

We thank Dr. H.K.B. Parekh, Senior Scientiest, All India Co-ordinated Research Project on Cattle, Jabalpur, for providing facilities.

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Induction of Parturition in Cattle

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ABSTRACT

Parturition was induced in 9 cows by administration of Dexamethazone alone 20 mgm. intramuscular and in 19 cows by giving a combination of Dexamethazone 20 mgms. and Stilboestrol at an average gestation period of 266 and 263 days respectively keeping 9 cows as control. All the induced cows calved at an average interval of 45.7 hours after the administration of the drug. Parturition occurred at 262.65 day of gestation in induced cows compared to 275.56 day of gestation in control. Those given a combination of Dexamethazone and stilboestrol calved 3.31 days earlier than those given Dexamethazone alone. Average birth weight of calves in the control and treated group was 27.65 kg. and 23.98 kg. respectively. Incidence of retained placenta was higher in the treated cows compared to control. Administration of Dexamethazone and oestrogen could reduce the incidence of retained placenta. Post Partum oestrus was delayed in the treated COWS compared to control animals.

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The ability of synthetic Cortico steroids to induce parturition in cattle has practical application in dairy industry especially because of the ability to synchronise the calving period with availability of labour, for therapeutic termination of pregnancy in prolonged gestation and

also for various clinical reasons. Since the initial reports of the use of Cortico steroids in the induction of parturition in cattle (Adams, 1969) several successful reports have appeared in the literature (Carrol, 1974; Thomas, 1975; O Farrel and Langley, 1975). However, Winter et al. (1974) Barth et al. (1978) and Davis et al. (1979) reported high incidence of retention of foetal membranes in induced calving. According to Garverick et al., (1974) administration of oestrogen in combination with dexamethaxone reduced incidence of retained placenta. Based on the above reports, a trial was carried out to study the effect of dexamethazone aloneanda combination of dexamethazone and oestrogen on the induction of parturition in cattle.

Materials and Methods

Twentyseven cross bred cows belonging to the Livestock Farm attached to the Kerala Agricultural University in advanced stage of gestation and not reported to have any difficulties in the previous calvings were selected for the study. These cows between 255 to 270 days of gestation were randomnly allotted in three groups of nine each. Cows in gorup I was not given any treatment and kept as control. Cows in group II were given Dexamethazone (9 alpha fluro 16 alpha methyl prednisolone) 20 mgms. intra muscularly. Cows in group III were given 25 mgms of Dexamethazone and Effect of treatment with Dexamethazone and a combination of Dexamethazone and Stilboestrol

		Control	Experimental			
	Item		Dexametha zone alone	a- Dexametha- zone and Mea stilboest-rol		
		Group I	Group II	Group II	I	
1.	No. of cows	9	9	9	-	
2.	No. responded to treatment		9	9		
3.	Average gestation period before treatment (days)		266	263	264.50	
	Average time response after drug administration to calving					
	an hour		48.6	42.8	45.7	
i.	Gestation length in days	275.56	268.06	264.75	266.40	
i	Average birth weight of calves in kg. /	27.65	24.52	23.62	23.98	
7.	Calving difficulty	nil	1	1	1	
3.	Retained foetal membrane	1 -	4	2	3	
9.	Onset of Post partum oestrous in days	58.2	78.2	66.51	72.38	

50 mgms of stilboestrol intramuscularly. Time from injection of the drug to calving was recorded. The calving difficulties if any were recorded and birth weight of calves immediately after birth was noted. Cows that had not expelled placenta by 24 hours were considered to have retained foetal membranes. Onset of post partum oestrum in all the cows in each group was recorded and avegrage post partum oestrous interval in each group was calculated.

Results and Discussion

Data presented in the table would reveal that the gestation length could be reduced to 266.40 days by induction of parturition compared to 275.56 days in the control. All the cows in which parturition was induced responded to treatment with all of them calving within 45.7 hours after injection of the drug. The interval from treatment of calving averaged 48.6 hours and 42.8 hours in group II and III respectively. Thus treatment with Dexamethazone and oestrogen resulted earlier calving than with Dexamethazone alone. Garverick et al., (1974) reported a shorter response time in beef cows given Dexamethazone plus oestradiol compared to dexamethazone alone. The mean birth weight of calves born to cows in group I, II and III was 27.65 kg., 24.52 kg. and 23.64 kg. respectively. Induction of calving resulted in average reduction of 3.67 kg. in their birth weight, the average birth weight of calves born with induced calving being 23.98 kg. Wagner et al., (1974) opined average increase of 0.5 kg. per day in the weight of foetus during the last few days of gestation. The study also revealed none of the cows in the control group had difficult calving while one each in the two treatment groups had difficult birth. Beardsley et al., (1974) and Carrol (1974) also reported that though, the calves in the induced birth was smaller in size, calving difficulty was greater in them compared to normal birth. Winter et al., (1974) and Bearsdley et al. (1974) also reported that induction of calving would result in greater incidence of difficult birth. However, Kelly et al. (1973) reported that by reducing gestation length by inducing calving the birth weight of calves would be reduced and the dystocia would be minimum.

According to Mc Diarmid (1980) incidence of Dystocia due to maternal foetal disproportion is not great in induced calving and the calving difficulty might be increased as a result of incomplete preparation of the birth canal.

Perusal of data in the table would reveal that incidence of retained placenta was higher in induced calving compared to control. It was also revealed that dexamethazone in combination with decreased oestrogen the incidence of retained placenta compared with those which were given dexamethazone alone. Garverick et al. (1974) reported that the failure to increase the serum oestrogen to quantities similar to those of natural calving prior to parturition might explain the high incidence of retained placenta in induced calving. He found that 6 mgm. of estradiol benzoate given with dexamethazone elevated blood oestrogen in beef cows and reduced the incidence of retention of foetal membrane. However, Davis et al. (1979) did not find decreased incidence of retained placenta in cows given 40 mgms of non conjugated oestrone or oestradiol 17.13. Schmitt et al. (1975) observed that a dose of 8 mgm of oestradiol benzoate to dairy cows failed to reduce the incidence of retained foetal membrane and failed to increase either circulatory plasma oestradiol or excretion to total oestrogen in urine to that of control.

The treatment also revealed that there was delay in the onset of post partum ocstrum in induced calving (72.35 days) compared to control (58.2 days). It could also be seen that post partum oestrum occurred earlier in group III than in group II. This might be attributed to the higher incidence of retained placenta. Arthur (1979) and Sandals et al. (1979) reported that the delay in the onset of post partum oestrum might be due to mild uterine infection which would follow retained foetal membrane. However, several workers claimed that induction of parturition with subsequent reproductive health management would not affect the future reproductive performance (Lauderdale, 1972; Beardsley et al. 1974; Carrol, 1974; Bolte et al. 1976).

Acknowledgement

The authors are thankful to the Dean, College of Veterinary and Animal Sciences, Mannuthy for permission to publish this paper.

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Biochemical Polymorphic Effect On Reproductive Status In Crossbred Cattle

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ABSTRACT

Sixty blood and serum samples from three equal groups viz., prepuberal, anestrus and cycling animals maintained under rural conditions were subjected to haemoglobin and serum albumin typing. Three haemoglobin phenotypes viz., AA, AB and BB were found in this present study. Similarly three albumin types AB, AA and BB were observed. Significant difference was not observed due to reproductive status both in Hb and Alb variants. However, Hb gene frequencies in prepuberal animals appeared differently from anestrus and cycling animals. Low incidence of Alb A and higher incidence of Alb B was observed in anestrus animals.

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Investigation of biochemical polymorphism in cattle especially genetic typing of haemoglobin and albumin is of recent development in animal production (Harpreet Singh and Khanna, 1971; Khanna and Singh, 1972, Krishna Singh and Nair, 1981 and Harpreet Singh, 1981). The application of biochemical polymorphism in generating exotic gene flow in our animal development was limited to few studies on production parameters (Krishna Singh and Nair, 1981; Harpreet Singh et al., 1983 and Jairam and Nair, 1983). No study was made so far applying the genetic typing of blood proteins especially haemoglobin

and albumin variants on reproductive status of crossbred animals. A study was therefore made for understanding the reproductive status of the crossbred animals as maintained by the farmers in relation to genetic variants of blood proteins.

Materials and Methods

Blood samples of 60 crossbred animals comprising of three equal groups of prepuberal, anestrus and cycling animals were collected. These animals having varying level of Jersey blood (25 to 87.5%) were maintained under rural conditions in Chittoor area of Andhra Pradesh. Haemoglobin types were studied on agar gel electrophoresis as described by Balakrishnan and Nair (1966). The separation of albumin was based on a discontinuous buffer system on starch gel electrophoresis as described by Poulik (1957).

Results and Discussion

Three Hb phenotypes viz. AA, AB and BB as reported by Harpreet Singh and Khanna (1971) and Nandakumaran et al. (1982) were observed in the present study. The gene frequencies of Hb A and Hb B were found to be respectively 0.30 and 0.70 in prepuberal, 0.53 and 0.47 in anestrus and 0.50 and 0.50 in cycling animals. There was no significant variation in gene frequencies between groups as shown in Table 1. However, low

Reproductive Status	No. of		Pheno	typic freque	encies	X2	Gene fre	quencies
	observations		AB	В	A		B	Α
Prepuberal heifers	20	0	12	8	_		0.70	0.30
		%	60	40	_			
nestrue animale		E	8.4	9.8	1.8	1.6		
Anestrus animals	20	0	13	3	4		0.47	0.53
		%	65	15	20			
		E	10.0	4.4	5.6	0.66		
Cycling animals	20	0	8	6	6		0.50	0.50
		%	40	30	30			
		E	10.0	5.0	5.0	0.54		

TABLE 1 Distribution of Haemoglobin Variants in Various Reproductive Conditions in Crossbred Cattle

TABLE 2 Distribution of Albumin Variants in Serum of Various Reproductive Conditions in Crossbred Cattle

Reproductive status	No. of		Phe	Phenotypic frequencies				Gene frequencies	
	observations		AB		A	B		A	B
Prepuberal heifers	20	0	12		6	2		0.60	0.40
		%	60		30	10			
		E	9.6		7.2	3.2	1.25		
Anestrus animals	20	0	10		1	9		0.30	0.70
		%	50		5	45			
		E	8.4		1.8	9.8	0.71		
Cycling animals	20	0	14		4	2		0.55	0.45
		%	70		20	10			
		E	9.9	Y	6.05	4.05	3.41		

O = Observed frequency E = Expected frequency A = Fast moving

B = Slow moving

incidence of allele B (0.47) in anestrus and allele A (0.30) in prepuberal heifers was observed. Absence of Hb A type in homozygus form was found in prepuberal heifers. Perhaps this represents nondiffusion of gene A in its crosses. In all the three groups higher frequencies of heterozygotes (AB) were observed. The gene frequencies in prepuberal animals appeared differently from anestrus and cycling animals.

Similarly three albumin type AB, AA

and BB were observed in this study as reported by Harpreet Singh (1981) and Nandakumaran *et al.* (1982). The frequencies of A1b A and A1b B genes were recorded as 0.60 and 0.40 in prepuberal, 0.30 and 0.70 in anestrus and 0.55 and and 0.45 in cycling animals as shown in Table 2. There was no significant difference in gene frequencies between groups. However, in anestrus animals low incidence of Alb A allele (0.30) and higher incidence of Alb B allele (0.70) was noticed. In all the three groups higher incidence of heterozygotes (AB) was observed. Since limited studies were made with few samples, further detailed studies are necessary to throw more light and confirm the present findings on haemoglobin and albumin polymorphism in crossbred cattle.

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Effect Of Forced Exercise On Seminal Characteristics And Sexual Behaviour Of Buffalo Bulls

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ABSTRACT

Investigations were conducted on eight healthy Murrah buffalo bulls (5-13 years old) to study the effect of exercise on seminal characteristics and sexual behaviour of buffalo bulls. On an average exercise and non exercise groups donated 3.29 and 2.25ml of semen and the differences were statistically significant (P Z 0.(5). But the effect of the exercise on semen consistency, mass activity, initial motility and sperm concentration was not significant. The study further revealed a significant $(P \angle 0.01)$ difference in reaction time of the two groups i.e. 247 and 94 seconds for exercise and non exercise groups respectively. On the whole, higher reaction time followed an increase in semen volume but similar consistency, sperm concentration/ml, mass activity and intitial motility were obtained in exercise group as against the non-exercise group.

Forced exercise has been considered to be an important pre-requisite for the management of breeding males Observations made by Bartlett and Perry (1939) on bulls used for artificial insemination suggested that forced exercise was of advantage in improving the quality of semen. But the evidences, thereafter, indicated that the forced exercise in case of bulls kept in loose pens, did not have significant effect on semen quality and quantity (Lepard, et al. 1941; Snyder and Ralston, 1955). No such studies seem to have been made in buffalo bulls. The present investigation was therefore, undertaken to study the effect of forced exercise on semen quantity, quality and sexual behaviour in buffalo bulls kept in loose pens.

Materials and Methods

Eight healthy Murrah buffalo bulls of 5-13 years of age maintained at the Punjab Agricultural University, Dairy Farm were used for the present investigation. Fulls were housed in half walled pucka bull pens. The bulls were divided into two groups of four each keeping in view their seminal characteristics and age. Group-I was given forced exercise for half an hour daily with the help of mannual bull exerciser whereas Group-II was kept as control. Semen was collected twice a week by artificial vagina technique. A total number of 48 collections were made. Sexual behaviour comprising of reaction time, libido index and service behavionr were noted at the time of semen collection. Semen was examined for physical characteristics such as volume, colour, consistency, mass activity, initial motility and sperm concentration. Total sperm production was calculated as volume x sperms/ml. The analysis of data was done according to Snedecor and Cochran (1967) after transfering

Bull No.	Volume	Consistancy (0-3)	Mass activity (0-4)	Sperm concen- tration millions/ml	Total sperm production ×10 ^s	Initial motility (Percent)
			Group-I (Exer	cise)		
952	3.21 ± 0.76	2.33 ± 0.16	3.16 ± 0.27	1365 ± 118.5	26461	64.16+3.52
586	1.8 ± 0.45	2.66 ± 0.16	3.5 ± 0.25	1275 ± 522	14558	67.5+282
558	5.0 ± 0.92	2.6 ± 0.16	3.25 ± 0.38	1480 ± 173	and the second	
156	3.14±0.48 2.5±0.15 3.7		3.7 ± 0.19	1375 ± 73	26172	69.1±2.39 70.5±2.14
Average	3.29±0.40	2.54±0.39	3.39±0.69	1360±290	111941*	67.59±7.0
		G	roup-II (N-Exerc	cise)		
962	2.1 ± 0.43	2.41 ± 0.23	2.91 ± 0.08	1308±200	18654	60.00+2.2
771	2.4 ± 0.49	2.1 ± 0.10	3.35 ± 0.16	1196 ± 67.6	16464	64.16+2.0
589	3.18 ± 0.27	2.8 ± 0.10	3.6 ± 0.24	1601 ± 64.3	30074	66.66 + 2.0
888	1.3 ± 0.53	2.6 ± 1.06	3.8 ± 0.16	1538 ± 109	12253	73.3±3.0
Average	2.25 ± 0.22	2.52 ± 0.08	3.37±0.12	1402±65.21	77545*	66.04+1.50

TABLE 1 Effect of forced exercise on semen characteristics of buffalo bull

* Total sperm production during the experimental period.

the percent scores of motility into arc since values.

Results and Discussion

Findings of the experiment on semen characteristics in relation to forced exercise are presented in table 1.

Semen characteristics

On an average exercise and nonexercise group of buffalo bulls donated 3.29 and 2.25 ml semen, respectively. The analysis of variance revealed a significant (P \angle 0.05) difference in semen volume of the two groups. Bartlett and Perry (1939) also observed an increase in semen volume as a result of exercise. The results obtained in the present investigation are also in agreement with Poroshin and Oboskalov (1976) who has reported 83, 34, 5, 20, 23, 12 and 17 per cent increase in ejaculate volume for bulls aged 3,4, 5, 6, 7, 9 and 10 years respectively due to exercise. The increase in semen volume in the exercise group may also be attributed due to the higher

reaction time encountered in this group which might have led to more excitement leading to increased semen volume. However, Lepard, Shuart and Foster (1941) and Snyder and Ralston (1955) found that exercise had no significant effect on semen quantity. No effect of forced exercise on the semen consistency, mass activity intitial motility and sperm concentration/ml was observed. This was in agreement with previous studies (Lepard et al. 1941 and Snyder and Halston, 1955) in cattle. Total sperm production was observed to be higher in case of exercise group as compared to non-exercise group, but the difference was found to be non-significant. An increase of 1.5 to 23.2 per cent sperm concentration due to exercise has also been reported by Poroshin and Oboskalov (1976).

Libido and Service behaviour

On an average reaction time of 247 ± 33.39 and 94 ± 26.41 seconds was observed in exercise and non exercise

Character	Observation	expressed in centage
	(Group-I) Exercise	(Group-II) Non exercise
Approach to the dummy.	-	
Very keen	9.0	66,6
keen	91.0	33.4
Chin rest	86.0	87.5
Election of penis with seath	99.0	91.7
Protruction of penis out side the seath	22.7	29.1
Curling of lips	45.5	25.0
Salivation at the time of ejaculation	22.7	12.5
Licking of dummy	72.7	58.3
Shniffing of urine	13.1	Nil
Reaction time (Seconds)	247 ± 33.39	94 ± 26.41

TABLE 2a Libido index and service behaviour of buffalo bulls in relation to forced exercise

TABLE 2b Service behaviour of buffalo bulls at the time of ejaculation in relation to forced exercise

Characteristics	and the second se	f a character in centage
	Group-I (Exercise)	Group-II (Non exercise)
Approach to the dummy		
Eager	63.7	12.5
Very eager	36.3	87.5
Location of forelegs		
Between pin and hook bone	86.9	91.7
in front of hook bone	13.1	8.3
Copulatory thrust		
Strong	42.0	46.0
Medium	59.0	54.0
Lifting of hind legs from the ground		
Complete	41.0	50.0
Partial	59.0	50.0

troups, respectively. These differences were statistically significant $(P \angle 0.01)$.

The results on libido index and service behaviour are presented in table 2a and 2b. Approach to the dummy was observed to be very keen and keen in 9.0 and 91.0 per cent of observations in exercise group, whereas it was 66.6 and 33.4 per cent of the observations in non-exercise groups. It is obvious from the observations that all the bulls were observed to be keen at the time of approaching dummy, which may be due to the reason that study was conducted on bulls trained for artificial collection of semen and more over during the favourable season of buffalo breeding. No difference was observed in respect of chin rest character in exercise and non exercise groups, table 2a. Similarly, there was not much differences in exercise and non-exercise bulls for erection of penis with sheath and protrusion of penis out side the sheath at the time of first mount on the dummy, table 2a.

Curling of lips and salivation were observed to be 45.5 and 22.7 per cent in exercise groups, somewhat higher as compared with non-exercise group in which the values were 25.0 and 12.5 percent Licking of dnmmy was expressed in 72.7% and 58.3% cases in exercise and non-exercise groups respectively. Sniffing of urine and vocalization were shown by 13.1 per cent cases only in exercise group.

Approach to the dummy by the bulls to serve the vagina was observed to be eager in 63.7 and 12.5% cases in exercise and non exercise groups, respectively but non exercise bulls were observed to be more eager to serve the vagina than the exercise bulls (Table 2b). Not much differences were noted as regard location of fore legs, between pin and hook bone in exercise and non exercise bulls, but the location of the fore legs in front of hook bone was obtained in 13.1% cases in exercise group where as it was only in 8.3% cases in non exercise group.

A strong copulatory thrust at the time of ejaculation was expressed in 42% in exercise and 46% cases in non exercise bulls. The copulatory thrust was observed to be medinm in 59.0 and 54.0% cases in exercise and non exercise bulls, respectively. A slight difference in regard of lifting hind legs from the ground at the time of ejaculation was noticed in exercise group only. (Table 2b).

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Non-Specific Genital Infections In Repeat Breeder Buffaloes

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ABSTRACT

A total of 69 repeat-breeder buffaloes were investigated for primary isolation of bacteria from cervical mucus samples. Out of them 57 (82.60%) buffaloes were found positive. The overall frequencies of different isolates in buffaloes were: gram negative bacilli 44.57 per cent, Corynebacterium 24.09 per cent, gram positive cocci 24.09 per cent and Anthracoids 7.22 per cent.

The antibiotic sensitivity test conducted with isolated organisms by paper disc sensitivity test, revealed a wide variation in the sensitivity pattern. The sensitivity test conducted in the composite cervical culture against the 8 antibacterial drugs viz., Penicillin, Streptomycin, Tetracycline, Furadentin, Neomycin, Ampicillin, Chloramphenicol and Gentamycin. The majority of mucus samples were found sensitive to Gentamycin (80.70%) and Chloramphenicol (63.16%). It could be seen that in repeat breeding animals non-visible genital infections (low-grade) played an important role for lowering the breeding efficiency.

The intra-uterine infusions of Dicrysticine, Mastalone-U, Phenivet and Gentavet were carried out after drug sensitivity test. The success of intrauterine infusions in repeat breeder buffaloes was 53.84 per cent with Dicrysticine, 64.28 per cent with Mastalone-U, 73.33 per cent with Phenivet and 73.33 percent with Gentavet. The overall conception rate in repeated and control group of buffaloes were 66.67 and 27.27 per cent respectively. Chi-square analysis to know the differences in the conception rate between treated and control group was found to be highly significant ($P \ge 0.01$).

Intensive dairy cattle and buffalo breeding under marginal feeding and maintainence has resulted into a great stress on these animals from the point reproduction and of production. Besides, the presence of specific venerial infections or low grade non-specific infections introduced during natural or artificial breeding or at the time of previous breedings and calvings may be responsible for unfavourable uterine environment. Various studies have indicated that non-specific bacterial infections could be clinically visible or non-visible, palpable or non-palpable changes may be responsible for the hostile environment in the tubular genitalia for fertilization and subsequent survival of zygote. This is more economically importtant type of problem affecting the dairy breeds of Surti buffaloes. On clinical examination there is no visible or palpable evidence or pathology.

Materials and Methods

A total of 69 repeat breeder buffaloes

and heifers were investigated for bacteriological examination. While 33 buffaloes were kept as control.

Mucus samples were collected asceptically by aspiration using sterilized glass pipette (10 ml Cap.), the pointed end of which was connected to a syringe with rubber junction, by recto-vaginal technique (Panagala *et al..*, 1978). After collection the samples were sent to laboratory and processed immediately.

Isolation work was done as per Cruickshank (1965). The isolates were subjected to in-vitro antihiotic sensitivity test as per the method recommended by Bauer *et al.*, (1966). Antihiotic discs supplied by Pasteur Biological Laboratories (India) were used. The isolates were tested for their sensitivity with 8 antihiotics viz., Penicillin, Streptomycin, Tetracycline, Furadentin, Neomycin, Ampicillin, Chloramphenicol and Gentamycin.

Out of 69 repeat breeder buffaloes, whose cervical mucus samples were investigated bacteriologically, 12 samples were found negative for presence of organisms. Remaining 57 animals were treated with antibiotic preparation as per the drug sensitivity tests. In this group 33 repeat breeder buffaloes which were not tested for cultural examination and drug sensitivity test were kept as control.

Results and Discussion

Under the present study genital microflora were isolated from 69 (15 heifers, 54 buffaloes) repeating buffaloes. Out of 15 samples in heifers, 4 samples (26.66%) did not reveal any bacteria and 8 (14.81%) samples from buffaloes were sterile. In all 11 (73.33%) heifers and 46 (85.18%) buffaloes had the presence of bacteria in the genital tract (Table 1.)

Repeat breeding animals showed the evidence of non-specific infections caused by varieties of micro-organisms. Perusal of literature indicated that non-specific bacterial infections either clinically diagnosible or not may be responsible for reduced fertility.

The results obtained presently, indicated that 73.33% heifers and 85.18% buffaloes harboured micro-organisms and these were in conformity with the observations made by Krishnamurthy *et al.*, (1974), and Verma and Tyagi (1974).

TABLE 1:	Genital	microflora	isolated	from	repeat	breeder	buffaloes
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	No. of animals	Microf	lora	Positive isolatio	bacterial on	-	Different	Different bacterial organisms				
		+ve samples	ve samples	Total No.	Single type	Mixed type	Gram negative	Coryne- bacterium bacilli	Anthra-	Gram positive cocci		
Heifers	15	73.33	26.66		54.54	45.45	43.75	25.00	12.50	18.75		
		(11)	(4)	16	(6)	(5)	(7)	(4)	(2)	(3)		
Buffaloes	54	85.18	14.81		65.21	34.78	44.77	23.88	5.97	25.37		
		(46)	(8)	67	(30)	(16)	(30)	(16)	(4)	(17)		
Total	69	82.60	17.39	1 2	43.37	25.30	44.57	24.09	7.22	24.09		
	224	(57)	(12)	83	(36)	(21)	(37)	(20)	(6)	(20)		

Figures in parenthesis indicate number of observation.

	No. of	Insemi	ination	number	Pregnan	cy per cen	t	Fertile	Overall
Drug used	animals	Cycle I	Cycle II	Cycle III	Cycle I	Cycle II	Cycle III	oestrus interval (days)	conception rate
Dicrysticin-S	13	13	10	8	23.07	15.38	15.38	32.57	53.84
					(3)	(2)	(2)	± 5.81	(7)
Mastalone-U	14	14	10	7	28.57	21.42	14.28	31.88	64.28
					(4)	(3)	(2)	±	(9)
	+							6.25	
Phenivet 15	15	15	9	5	37.50	26.66	7.14	27.54	73.33
					(6)	(4)	(1)	. ±	(11)
								4.12	
Gentavet	15	15	7	4	53.33	20.00		22.36	73.33
					(8)	(3)		+	(21)
30								2.61	
Total	57	57	36	24	36.84	21.05	8.77	28.00	66.67
					(21)	(12)	(5)	±	(38)
(Treated group)								2.27	
Control group	33	33	31	29	6.06	6.06	15.15	47.66	27.27
			10.0		(2)	(2)	(5)	+	(9)
-						4.4		5.98	

TABLE 2 Treatment of repeat breeder buffaloes with antibiotics and resulting pregnancies.

Figures in parenthesis indicate number of animals.

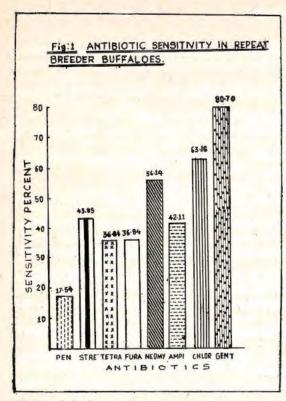
Besides, many other workers have also stressed that non-specific infection of uterus results in breeding difficulties in cattle (Jainudeen, 1965; Jerkovic *et al.*, 1971). The present findings do not support the findings of Hawak *et al.* (1958), Nunn (1970) and Hartigen *et al.* (1972) who found low occurrence of infection in the repeat breeders.

Under the present study, the genital microflora were invariably isolated from repeat breeder buffaloes With the non-visible type of infection, there have been single and mixed type of isolations to the tune of 43.37% (36) & 25.30%(21) respectively.

Antibiotic Sensitivity:

The sensitivity test of micro-organisms with antibacterial agents is felt to be of paramount importance, in effective check of various infections, because many organisms, in the process of their multiplication in vivo, develop resistance to the antibiotics due to their indiscriminate usage. The information about the sensitivity test and to choose the appropriate antibiotic for treatment is meagre in literature. However, in the present study, primary isolation of organism along with their sensitivity test was conducted to find out a suitable drug for successful treatment for the repeaters. When the composite samples were tested with different antibiotics viz., Penicillin, Streptomycin, Tetracycline Furadentin, Neomycin, Ampicillin, Chloramphenicol and Gentamycin isolates were found to be sensitive to the tune of 17.54, 43.86, 36.84, 36.84, 56.14, 42.11, 63.16 and 80.70 per cent in buffaloes respectively. (Fig.: 1).

It was observed that the sensitivity was limited to higher antibiotics range only.



This might be perhaps due to undue usage of lower range antibiotics for various ailments for a considerable period. The present findings are in agreement with those of Gragg (1964) and Nunn (1970). While they were not in conformity with the observations of Polakova and Labduska (1962). Thus, it may be concluded that the choice of drug for treatment of repeat breeders should be so made which may be effective even in lower concentration.

Antibiotic Therapy:

In the present study, microflora in the repeat breeder buffaloes was isolated from cervical mucus samples and the specific antibiotics were used for the treatment. Out of 57 buffaloes treated with Dicrysticine, Mastalone-U, Phenivet, and Gentavet for bacterial infection, 38 (67 67%) conceived. In 33 buffaloes kept as control 9 (27.27%) conceived. The conception rate differed highly significantly between treated and control group ($p \angle 0.01$) (Table - 2).

General clinical practitioners have employed intra-uterine infusion of antibiotics with variable results in increasing fertility in repeat breeders. However, significant differences could not be seen between treated and untreated control (Paufler, 1969; Buckstrom, 1970). On the contrary some reports of good success have been claimed with Penicillin Plus Streptomycin in a limited trial, having suitable control (Luktuke et al., 1958 and Khan and Luktuke, 1967). The present findings suggest that antibiotics should not be used arbitrarily for intrauterine therapy. However, usage of specific antibiotic resulted intoencouraging pregnancy rates in repeat breeder buffaloes.

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Mycoflora Of The Genital Tract Of Surti Buffaloes

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ABSTRACT

The present investigation reports on the isolation of fungal organisms from the cervicovaginal mucus samples of 100 Surti buffaloes under different reproductive status viz. normal animals, animals with endometrtis and anoestrus animals. Fungi were isolated from 47% of animals irrespective of their reproductive status. The fungi isolated include Mucor sp., Aspergillus sp., Fusarium'sp. Microsporon sp. and Geotricum sp. Aspergillus sp. (23.8%) predominated in endometritis cases.

*

Indiscriminate use of broad spectrum antibiotics and corticosteroids has probably led to an overall increase in the incidence of fungal infections. Fungal infections of the bovine genital tract resulting into infertility have been described by Jungherr (1935) and Saxena and Pathak (1972). Fungi have been isolated from cervical mucus of infertile as well as healthy cows by Rollinson and Haq (1948), Ainsworth and Austwick (1955) and Wawrzkiewicks and Galeza (1972). In the present study attempts were made to isolate and identify fungi from cervicovaginal mucus of Surti buffaloes under different reproductive status.

Materials and Methods

The cervico vaginal mucus samples from 100 Surti buffaloes, brought for gynaecological check up to the A.I. Centre of this College were examined. Animals chosen for study included those which were: (1) in oestrus and having normal discharge, (2) having mucopurulent discharge (cases of mild endometritis) (3) in anoestrum.

The samples were collected aseptically by aspiration using sterilized glass pipette (10 ml. cap.), the pointed end of which was connected to a syringe with rubber junction by rectovaginal technique (Pattabhiraman *et al*, 1967; Panangala *et al*, 1978). In anestrous animals, sterile saline was infused into the enital tract and aspirated back. After collection, the samples were brought to Bacteriology laboratory and processed immediately.

Isolation of fungi was carried out as per Cruickshank *et al.* (1975) using Saboraud's dextrose agar medium and cultures were characterized as per Sigurd Funder (1961).

Results and Discussion

Out of 100 samples of cervicovaginal mucus examined, 47 samples were found to be positive for fungal organisms. All groups of animals yielded almost similar percentage of isolates irrespective of their reproductive status. The fungi isolated were *Mucor* sp., *Aspergillus* sp., *Fusarium sp.*, *Microsporon* sp. and *Geotricum* sp. (Table -1).

Jerkovick et. al. (1971) found Aspergillus sp. to be most common in endometritis cases which confirms the findings of

Sr. No. Name of Fungi	Normal fertile animals in oe		Animals endometritis		Animals in anestrus (22)		
	No. of isolates	%	No. of isolates	%	No. of isolates	%	
1. Mucor sp.	11	30.55	6	14.26	8	36.36	
2. Aspergillus sp.	4	11.11	10	23.80	2	9.09	
3. Fusarium sp.	2	5.55	1	2.38	0	00.00	
4. Microsporon sp.	0	00.00	2	4.76	0	00.00	
5. Geotricum sp.	0	00.00	1	2.38	0	00.00	
Total	17	47.22	20	47.61	10	45.45	

TABLE 1. Fungal isolates obtained from cervicovaginal mucus of Surti buffaloes.

present study. Lecklark et al. (1972) found Penicillum sp. more frequently in infertile animals, however in the present study, Penicillum sp. was not obtained. Sinha et al. (1980) isolated fungi in 33% of 146 cervical mucus samples from infertile cows and buffaloes. They found A. fumigatus, A. niger, Absidia sp. and Rhizopus sp. as predominant species while the other isolates were Curvularia sp., Trichosporon sp. and Candida sp.

It is likely that subsequent to bacterial infection and resultant lowering of uterine resistance, fungi might be playing the roles as secondary invader or it may be predisposing cause for setting up of bacterial infection. Though as many as 47% of samples yielded fungi, more work is required to be done for assessing the pathogenic role of fungi in infertility conditions among animals particularly as fungi are ubiquitous in nature.

Acknowledgement

We are grateful to the Principal for the facilities provided and to the Department of Gynaecology for the help in collection of samples.

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Studies On Mycotic Abortion Caused By Aspergillus Fumigatus Fresenius

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ABSTRACT

Role of Aspergillus fumigatus in the etiology of abortions was studied in 31 buffaloes and 22 cows. Of the 53 animals investigated, mycotic abortion due to A. fumigatus could be recorded in 2 buffaloes and 1 cow giving a prevalence rate of 5.1 per cent. Abortions in all the animals occurred at 6-7 months of gestation period. Diagnosis was based on the isolation of A. fumigatus from the placental tissues, stomach contents, skin scrapings and lungs of the aborted foetuses, detection of 2-4 µ thick septate, dichotomously branched fungal hyphae compatible with Aspergillus under direct microscopy by potassium hydroxide technique and demonstration of A. fumigatus antibodies in the sera of one buffalo and one cow by immunodiffusion method. The isolates of A. fumigatus recovered from clinical and environmental samples were found pathogenic to male Swiss albino mice when inoculated intravenously into caudal vein. The epidemiological findings revealed the presence of the fungus in the saprobic environment of the milch animals.

Abortions in dairy animals are important both from economic as well as public health point of view. A number of infectious agents such as viruses, bacteria, fungi, protozoa, chlamydia, mycoplasma and rickettsia are incriminated in the etiology of abortions in animals (Blood et al., 1979). In recent years due to reduction in the incidence of bacterial abortions, mycotic abortions are receiving more interest and importance (Ainsworth and Austwick, 1955; Hillman, 1969; Ainsworth and Austwick, 1973).

Mycotic abortion is a sporadic infection of genital tract of animals particularly of cows (Ainsworth and Austwick, 1973). The incidence of abortion varies from locality to locality and is probably associated with the concentration of fungi in the saprobic materials and surroundings of the milch animals (Hillman, 1969; Kremlev, 1971; Carter et al., 1973; Venev, 1974; Siddique et al., 1976). Although a number of fungi and actinomycetes are associated with the etiology of abortions in animals, Aspergillus fumigatus is considered as the chief organism responsible for a substantial number of abortions (Hugh-Jones and Austwick, 1967; Carter et al., 1973; Konig and Nicolet, 1974; Jerrett, et al., (1984) Mycotic abortion due to A. fumigatus has been recorded in cows, sheep, goat and buffaloes (Adamesteanu and Baba, 1973; Ainsworth and Austwick, 1973; Osman and Gabel, 1978). There appears to be little information on the clinicoepidemiology and mycoserolegy of mycotic abortions in milch animals. The present paper describes the clinical, mycological, serological and epidemiological observations in the naturally

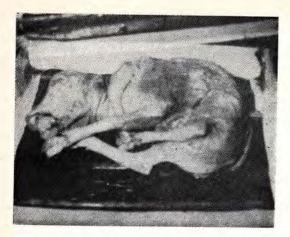


FIG. 1. Six month old foetus aborted from an 8 year-old Murrah she buffalo due to natural infection with Aspergillus fumigatus.

occurring abortions due to A. fumigatus in cows and buffaloes.

Materials and Methods

The study was conducted on 58 dairy animals which comprised of 31 buffaloes, 22 cows and 5 goats. A total of 91 clinical samples which included 27 foetal stomach contents (15 buffaloes, 9 cows, 3 goats), 24 foetal skin scrapings (12 buffaloes, 8 cows, 4 goats), 21 foetal lungs (11 buffaloes, 9 cows, 1 goat) and 19 placentas (10 buffaloes, 6 cows, 3 goats) were collected aseptically to process for mycological investigation. All these organs were examined for gross lesions, if any. A portion of the material was treated with 10 per cent potassium hydroxide for 10 minutes for direct microscopy. Each specimen was liberally inoculated on to slants of Sabouraud's dextrose agar (Emmons et al., 1977) with chloramphenicol (0.05 mg/ml) and incubated at 37°C for one week. The blood was collected by jugular vein from a positive cow and a buffalo for A. fumigatus precipitins by immunodiffusion

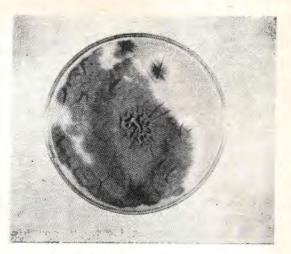


FIG. 2. Four day old primary culture of Aspergillus fumigatus from the stomach contents of a six-month old aborted foctus on Sabourauds dextrose agar with chloramphenicol after 3 days of incubation at 37°C.

technique. Environmental prevalence of A. fumigatus was studied in 12 samples of fodder, 12 animal excreta and 12 soil (Pal and Mehrotra, 1982). In order to detect the presence of A. fumigatus in the air, 12 petri plates of Rose Bengal agar were exposed for 3-4 minutes in the inside environment of animal sheds. The morphology of the fungus was studied in lactophenol cotton blue mounts as per the description given by Raper and Fennell (1965). The pathogenicity test was conducted with 8 isolates of A. fumigatus in male Swiss albino mice (Pal, 1983).

Results

Of the 58 animals investigated, mycotic abortion could be diagnosed in 3 animals giving a prevalence of 5.1 per cent. The positive animals were 7 and 8 years old she-buffaloes and 8 year old cow (Table 1). The isolation of Aspergillus fumigatus was successful from 3 stomach contents, 3 skin scrapings, 2 lungs of the

Species of animal examined	Number of cases screened	Number found positive A. fumigatus	for
Buffalo	31	2 (6.4)*	
Cow	22	1 (4.5)	
Goat	5	0 (0.0)	
Total	58	3 (5.1)	

TABLE 1: Association of Aspergillus fumigatus with abortion in dairy animals.

* Figures in parenthesis indicate percentage

Types of clinical specimen investigated	Number of samples examined	Animals yielding A. fumigatus				
		Buffalo	Cow	Goat		
Stomach contents	27	2	1	0		
Skin scrapings	24	2	1	0		
Lungs	21	1	1	0		
Placentas	19	2 -	1	0		
Total	91	7	4	0		

TABLE 2: Isolation of Aspergillus fumigatus from various clinical materials of aborted foetuses and placentas.

foetus besides 3 placentas (Table 2). No isolation could be achieved from any of the clinical specimens of goats. The details of the clinical, mycological and serological observations are summarised in Table 3. Abortions in all the 3 animals occurred during 6-7 months of pregnancy. The placentas in most of the cases were found thickened, necrotic, haemorrhagic and oedematous. The aborted foetus from a eight-year-old Murrah she-buffalo showed discrete, raised areas on the skin of head and back (Fig. 1). Aspergillus fumigatus was isolated from the stomach contents of a 6-month-old foetus aborted from a 8-year-old Murrah buffalo (Fig. 2). Branched, septate hyphae, 2-4 µ thick compatible with Aspergillus could be detected in the 2 skin scrapings and 3 placental tissues by potassium hydroxide techniques. Precipitins against A. fumigatus was demonstrated in two sera samples obtained from a buffalo and a cow. The fungus was highly prevalent in the environment of all the 3 dairy animal (Table 4). All the 8 isolates of A. fumigatus proved to be pathogenic to Swiss albino mice as evidenced by the death of the laboratory animals and also by reisolation of the pathogen from most of the visceral organs.

Discussion

The results of this study suggested that Aspergillus fumigatus may be considered as an important causative agent in the etiology of infectious abortion. This is evident that 3 animals out of 58 were positive for mycotic abortions giving a incidence rate of 5.1 per cent. All the

Animal	Age of	Stage of	Lesion	Gross	Clinical	Is	olation of A	. fumig	gatus	Prevalence
species	s	on skin placenta	signs	Lung	Stomach contents			of A. <i>fumigatus</i> antibodies in sera		
1. Buffalo	8yrs	6 month	Discreate, raised areas on skin of head and neck	necrotic and haem orrhagic	temp. - 40.4°C,	+	+	÷	+	+
2. Cow	8yrs	6 month	Circular, ringworm type lesions on the head region	hagic, oedema- tous suppura- tive, necrotic	temp. 40.6°C, vaginal discharge, anorexia, retention of	+	+	+	+	+
3. Buffalo	7yrs	7 month	No skin O lesions obsered	dematous, thickened necrotic	Rectal	-	+	-	+	sera was not tested

TABLE 3: Clinical, mycological and serological findings in mycotic abortion of animals due to Aspergillus fumigatus.

three abortions (2 in buffalo and 1 cow) occurred during six and seventh month of pregnancy. This observation is similar with Adamesteanu and Baba (1973) who reported mycotic abortion in cows due to A. fumigatus in late gestation.

Although macroscopic lesions on foetus and placenta are not very characteristic to warrent a diagnosis of mycotic abortion, a tentative diagnosis can be made by observing central necrosis, leather-like consistency, thickening and haemorrhages on the cotyledons and ringworm type circular lesions on the skin of head and neck of the foetus (Ainsworth and Austwick, 1973). We have observed similar gross lesions on the foetal skin as well as cotyledons of one buffalo and a cow. Further, 2-4 μ thick, dichomotously branched, septate hyphae could be demonstrated in the placental tissues and skin scrapings of one cow and a buffalo when examined by potassium hydroxide techniques.

The fungi have been frequently recovered from the placenta, the amniotic fluid, foetal stomach contents and foetal skin lesions (Ainsworth and Austwick, 1973). Very rarely isolations of the fungi have been made from the other organs

Sr. No.	Species of animal	Age of animal	Isolation of A. fumigatus from the saprobic materials					
			Fodder	Animal excreta:	Soil	Air		
1.	Buffalo	8 yrs	3/4	1/4	2/4	3/4		
2	Cow	8 yrs	2/4	2/4	3/4	3/4		
3.	Buffalo	7 yrs	1/4	1/4	2/4	2/4		
		Total	6/12	4/12	7/12	7/12		

TABLE 4: Prevalence of Aspergillus fumigatus in the environment of animal sheds of animal positive for mycotic abortion.

of the foetus such as the lungs and liver. However, Austwick and Venn (1961) have found that the best isolations can be achieved from the stomach contents of the foetus on mycological agar containing antibiotics. In this study A. fumigatus was isolated in pure and heavy growth on Sabouraud's dextrose agar with chloramphenicol at 37°C from the foetal stomach contents, skin lesions and placental tissues besides lungs of the foetus. It is pertinent to mention in this context that the isolation of the pathogen from within 24 hours from the placental tissues on mycological media may be significant considered provided the organism isolated showed the similar morphology in the clinical specimen under direct microscopy (Ainsworth and Austwick, 1973). Interestingly both the criteria were confirmed in the present investigation.

The sera from the two aborted dams (cow and buffalo) when tested by immunodiffusion technique on the method of Oucheerlony (1967) revealed the presence of precipitins against Aspergillus fumigatus. However, the significance of agar gel precipitation test in the sero diagnosis of mycotic abortions due to A fumigatus has been discussed by Corbel (1972) and Uppal et al. (1978).

The pathogenesis of mycotic abortion due to Aspergillus fumigatus is still controversial. Most of the investigators believe that the primary infection occurs in the lungs as the result of the inhalation of the fungus from the mouldy fodder (Cordes et al., 1964; Korotochencko et al., 1974; Veney, 1974) and the infection may spread to the genital organs by means of blood stream. However, a recent experimental study made by Kremlev (1977) in cows to elucidate the role of A. fumigatus and other fungi in the pathogenesis of mycotic abortion reported that possibly the fungi and their toxins penetrated the uterus and foetus by haematogenous route. Likewise Weney (1975) has stated that cows in heat if inseminated with mould contaminated semen may develop sexual cycle disorders. In order to substantiate any of the above views, a detailed systematic study tracing the path of the organism would be rewarding.

Acknowledgement

We are grateful to Dr. I. D. Sharma and Dr. H. S. Jain, Animal Husbandry Department, Delhi for their kind help and cooperation in the collection of clinical material and detailed history of the cases. Technical help of Mr. Ram Prakash is also thankfully acknowledged.

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Neoplasms In The Genital System of Cows

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ABSTRACT

As the genital neoplasms are one of the important causes of infertility and sterility in bovines, the present study is of special significance. Out of 10,000 genitalia of cows examined in the Calcutta Slaughter Houses, only 28 (0.28%) cases of neoplastic growths were observed. Of these, 2 (0.02%) fibropapillomas and 6 (0.06%) squamous cell carcinomas in the vulva, 3 (0.03%) leiomyomas in uterus and vagina,1 (0.01%) adenoma in uterus, 2 (0.02%) adenocarcinomas in uterus and cervix, 2 (0.02%) lymphosarcomas in uterus, 2 (0.02%) fibromas in cervix and vagina, 8 (0.08%) dermoid cysts and 2 (0.02%) granulosa cell tumour in the ovary were observed, during pathological investigations.

Neoplasms in the genital organs of cows are grave concern due to reproductive failure in the form of infertility or sterility and mortality. The reproductive failure is caused by destruction of normal architecture and narrowing or obstructing the lumina of tracts which are concerned for the secretion and transport of ovum.

The neoplastic growths in the genitalia are rather difficult to detect clinically unless they are well pronounced. Literature on the subject is scanty, based only on case records and thus it is proposed to describe below the prevalence and pathology of neoplasms of the genitalia of cows.

Materials and Methods

In the course of examination of 10,000 genital organs of cows slaughtered for beef at Calcutta slaughter houses, 28 genitalia showing abnormal growths were collected during January, 1982 to December, 1984. The respective organs showing growths were preserved in 10% neutral formol-saline, processed through conventional techniques, sectioned at 4-5 μ in thickness and stained with haematoxylin (Lillie Mayer's) and cosin. Some of these were also stained by Masson's trichrome and Van Gieson's picrofuchsin for connective tissue.

Results and Discussion

Out of a total of 10,000 genital organs examined, 28(0.28%) were found to have neoplastic conditions.

Fibropapilloma: It was observed in 2 (0.02%) cases involving the skin of the vulva. Jones and Hunt (1983) also opine that it is a rare tumour of cows.

Pathologically, it was cauliflower-like in appearance, grayish-white in colour, variable in size and projecting on the surface. Histopathological lesions consisted of matured connective tissue extremely covered with stratified squamous epithelium of limited thickness.



Fig. 1: Microphotograph showing oval masses of Keratinised epithelial cells arranged concentrically, known as cell nest. H & E X 100.

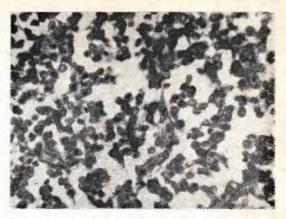


Fig. 3: Microphotograph showing massive proliferation of lymphocytes with hyperchromatism and distinct mitotic figures. H & E X 400.



Fig. 2: Microphotograph showing spindle-shaped muscle fibers, separated by connective tissue. H & E X 80.

A viral etiology is attributed for this condition (Runnells et al. 1976).

Squamous cell carcinoma: This neoplasm involved the skin of 6 (0.6%) vulva. Among the excellent reviews on the subject could be mentioned those of Nair and Sastri (1954), Murray (1968), and Damodaran *et al.* (1975).

This tumour was soft, grayish in colour

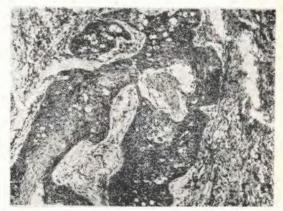


Fig. 4: Microphotograph showing multilayed neoplastic cells arranged both solid and papillomatous forms, which were supported by fibrous stroma and blood vessels. H & E X 80.

with haemorrhagic foci and ulceration. Histologic sections featured a squamous cell carcinoma which consisted of rapid proliferation of prickle cell layer running random in all directions. The cells were embryonic as evidenced by larger size, numerous mitotic figures, hyperchromasia and lack of differentiation. There were round or oval masses of keratinised epithelial cells arranged concentrically, known as cellnest (Fig. 1). The connective tissue was abundant with numerous thinwalled blood vessels. The pathological lesions observed had a close resemblance to those described earliar in bovine (Nair and Sastri, 1954; Murray, 1968 and Damodaran *et al.* 1975).

It is believed that the high intensity of biologically potent ultraviolet radiation that occurred in countries near the equator, such as India, Ceylon, Kenya etc. possibly induced cancerous changes in the unpigmented vulvular skin of cattle (Wettimuny et al. 1974). Discussing the possibility of a hereditory basis for the incidence of vulvular cancer, Vandegraaff (1976) considered it probable that a hereditory predisposition resulted from the amount of white coat inherited and hence the likelyhood of the offspring having partly or wholly unpigmented vulvas.

Fibroma: It was encountered in 1 case of each in the cervix (0.01%) and vagina (0.01%) which are in agreement of earlier reports in cow (Chenna Reddy, 1980) and she-buffaloes (Sane and Purohit, 1958; Rama Rao and Rajya, 1976; Sharma et al., 1977).

Grossly, it was hard, smooth and encapsulated. Histologically, it consisted of interlacing bundles of fibrous tissue running in all directions and the nuclei of the cells were spindle shaped. Fibromas situated in the female genitalia blocked the passage thereby inducing infertility.

Leiamyoma: This tumour was recorded in 3 (0.03%) cases — 1 (0.01%) in vagina and 2 (0.02%) in uterus. A case of uterine leiomyoma was also reported by Chakraborty and Kwatra (1984).

It was pink in colour, firm in consistency, spherical in shape, dry in appearance and protruding in the respective organs. Histopathologically, the muscle fibres were spindle shaped with elliptical nuclei having chromatin. The bundles of muscles were separated by connective tissue with abundant blood supply (Fig. 2). This tumour in the vagina and uterus caused obstruction to the passage of secretion and transport of ova resulting infertility or sterility.

Lymphosarcoma: It was noticed in 2 (0.02%) uterus. Olson (1974) and Bhowmik and Iyer (1978) also reported that this neoplasm is 2.0 to 2.5 per 10,000 buffaloes, which bears similarity to the present records.

The neoplasm was grayish-white in colour and appeared as solitary tumors. The histologic sections consisted of massive proliferation of lymphocytes and their precursors which completely replaced the architecture of the organ. Neoplastic cells were either diffusely, scattered or developed ill-developed follicles. The cells were either oval in shape with vesicular nuclei or the nuclei were hyper-chromatic with distinct mitotic figures (Fig. 3).

Lymphosarcoma is sporadic in adult cattle and undoubtedly fatal (Gresham and Jennings, 1964; Bhowmik and Iyer, 1978). The viral etiology of this neoplasm was attributed by Jarrett (1964).

Adenoma: This benign tumour was encountered in 1 (0.01%) uterus which is in agreement of Kumar and Singh (1984) in a she-buffalo.

Pathologically, it was soft, pink in colour and encapsulated. There was excessive proliferation of glandular epithelial cells with faint granular cytoplasm and oval nucleus. The cells were grouped into masses separated by stroma forming lobules.

Adenocarcinoma: It was recorded in 2 (0.02%) cases - 1 (0.01%) in cervix and 1 (0.01%) in uterus. The prevalence of this condition in the genitalia was also described by Runnells *et al.* (1976) and Jones and Hunt (1983).

Pathologically, it was firm, grey or vellow in colour, nodular in appearance and projecting on the surface. The proliferating epithelial cells were single or multilayed and replaced the parenchymal architectures. Sometimes they broke through the basement membrane and infiltrated in the underlying stroma. The cells were arranged both solid and papillomatous forms (Fig. 4) and had the usual characteristics of malignancy, i.e. vesicular nuclei with large nuclei, hyperchromatism and distinct mitotic figures. The growth was supported by fibrons stroma and thick-walled blood vessels (Fig. 4).

Adenocarcinoma is rare tumour in the uterus of the aged cattle and is always serious (Runnells et al., 1976; Jones and Hunt, 1983). Hormonal imbalance is believed for the development of this tumour in the genitalia (Meissner et al., 1957). It is not known how frequently sterility is associated with this neoplasm since the clinical data is lacking.

Dermoid cyst: It was observed in 8 (0.08%) ovaries. The tumour is very common in the ovaries of she-buffaloes (Rama Rao and Rajya, 1976; Kumar and Singh, 1984).

Pathologically, the ovarian architecture was replaced by dermoid cyst which was creamy-white in and lined by skin and its colour and appendages.

Granulosa cell tumour: Two (0.02%) ovaries showed granulosa cell tumour, which corroborated the findings of Mukherjee (1969). Its low incidence can be attributed to factors associated with the ageing.

The tumour was oval, yellow in colour, lobulated, encapsulated and projecting on the surface. Cut surface revealed cystic spaces filled with clear fluid. Histologic sections consisted of masses of large polyhydral cells with granular cytoplasm and centrally placed nuclei. The cells were either arranged in solid masses or follicles around central lumen containing acidophilic materials. Hyperchromatism and mitotic figures were plentiful. The pathological lesions are in agreement with Mukherjee (1969) and Jones and Hunt (1983).

This tumour is mostly observed in 2-5 years old cows and associated with hyperestrinism (Runnells *et al.*, 1976) which invariably causes nymphomania resulting to infertility or sterility.

The results showed the lowest incidence (0.28%) of genital neoplasms in cows, which are in agreement with the statement of Laing (1970). The low incidence can be attributed either to a species difference in susceptibility or to factors associated with endocrine functions.

As the present materials were collected from the slaughterhouse, history of cases, clinical signs associated with tumours, fertility examination of the cases etc. could not be ascertained.

Acknowledgement

Thanks are due to the Head, Deptt. of Veterinary Pathology and the Vice-Chancellor of this Viswavidyalaya for the funds and facilities provided.

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Uterine Torsion In Buffaloes I-Incidence

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ABSTRACT

A survey of obstetrical disorders treated over the last 12 years at the Veterinary Clinic, H.A.U. Hissar revealed that uterine torsion comprises the single largest cause of dystocia in buffaloes. Its incidence was high during the months of July to October and was more common in pleuriparous buffaloes. A consistant feature observed was the close association of uterine torsion with parturition. The most common form of rotation was 180 degrees and the direction was clock-wise. Post-cervical uterine torsion occurred in a much higher frequency than pre-cervical uterine torsion.

* *

It is estimated that out of the worlds approximately 130 million domestic buffaloes, about one half exists in India and a quarter in China (FAO, 1975). In India, buffaloes contribute about 55 percent of the milk and 60 percent of the meat produced in the country. In view of the significant contribution of buffalocs towards national economy, it has become necessary that these animals reproduce fairly regularly with minimum intercalving periods for maintaining optimum reproductive efficiency. This envisage a study into the incidence, causes and remedial measures of the various reproductive disorders in buffaloes. Some of these aspects have been recently studied and reviewed by Gupta et al (1981). It was briefly reported in that

study that abnormal parturition and uterine torsion in particular constitute a serious reproductive disorder affecting a fairly large number of buffaloes. In this paper we report the result of a detailed analysis of the incidence of uterine torsion in buffaloes.

Materials and Methods

The overall incidence of uterine torsion and its relation to season and parity of the animal was studied from clinical records of obstetrical disorders maintained at the Veterinary Clinic, H.A.U. Hissar over a period of 12 years ranging from January 1970—January 1983. In addition, the incidence of uterine torsion in relation to the stage of gestation, degree, direction and position was studied from 17 clinical cases presented for treatment at the same clinic during the last year.

Results and Discussion

The analysis of the data on the incidence of difficult calving in buffaloes and cows revealed that uterine torsion comprises the single largest cause of dystocia in buffaloes (Table 1). The incidence of 43.44% recorded in the present study is in close agreement with the reports of Mannari and Tadkod (1976) and Pattabiraman et al (1979). However, a relatively much higher incidence (Jit Singh et al 1978). and a marginally lower incidence of uterine torsion have also been reported. We subserve with the

TABLE:	Incidence of uterine torsion in buffaloes and cows at
	Veterinary Clinics, Haryana Agricultural University,
	Hissar during the period Jan. 1970 - Jan. 1983.

	Buffaloes	Cows	Total
Total recorded Dystocias	145	50	195
	(74.35)	(25.65)	
Uterine torsion	63	7	70
	(43,44)	(14)	(359)

Figures in parenthesis indicate percentage.

view of Pearson (1971) that the breed and geographical location has a significant influence on the incidence of uterine torsion. Nevertheless, the incidence in buffaloes seems to be relatively much higher than the reported incidence in Cows (Morten and Cox 1978, Pearson 1971). Besides species differences, limited studies made so far have attributed the voluminous abdomen and the wallowing habbit in buffaloes, to the high incidence of uterine torsion in this species (Jit Singh et al 1978).

The present study revealed a definite seasonal pattern in the incidence of uterine torsion. The incidence was highest during the months of July to October and lowest during the months of March to June. The incidence also followed a similar month-wise pattern with highest incidence in the months of August and September. Its high incidence between July and October corresponds closely with the calving season in the species.

The incidence of uterine torsion was

higher in pleuriparous (73.02) than in primiparous buffaloes (26.98%). This has been previously reported by Mannari and Tadkod (1976) and Jit Singh et al (1978). This may perhaps be due to the progressive stretching of the broad ligament associated with successive pregnancies.

The incidence of uterine torsion with the stage of gestation, degree, direction and position was related in fourteen clinical cases treated during the last year (Table II). A consistent feature observed was its close association with parturition as all the cases were presented near term or in the first stage of labour. The association of uterine torsion with parturition, particularly the first stage of labour has been earlier reported by Fouad and El-Sawaf (1964), Thangaraj et al (1972), Jit Singh et al (1979) and Pattabiraman et al (1979). Although the reason for its close association with labour has not been completely understood, Wright (1958) contended that it is due to exciting causes which operate maxi-

TABLE. II:	Incidence of different degrees,	direction and position of uterine torsion in buffaloes.	
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	Degress				Direction		Position	
	90°	180°	270°_	360°	Clockwise	Anticlock wise.	Precer- vical	Postcer- vical.
Numbers of animals	2	9	2	1	14	,	1	13
Percentage	14.28		14.28	7.4	100.0		7.14	92.86

mally during the first stage of labour. On the other hand, pearson (1971) speculated the excessive foetal movement that occur during the first stage of labour as a direct exciting cause for uterine torsion.

The present study also revealed that 180 degree rotation of the uterus was the most common degree of uterine torsion and 90 and 270 degrees torsion occurred with almost similar frequency. 360 degree rotation was recorded in a solitary case. Fouad and El-Sawaf (1964) and Jit Singh et al (1978) also reported 180 degrees torsion to occur most frequently in buffaloes.

The direction of torsion was clockwise in all the 14 clinical cases. Similar observations have been made by Jit Singh et al (1978) and Pattabiraman et al (1979). In cows on the other hand uterine torsion has been reported to occur predominantly on the left side (Pearson 1971). Although the reasons for the variation are not known, it is probable that in buffaloes, a more capacious abdomen and a distended rumen in the left side may prevent the more frequent rotation of the gravid horn in the anticlockwise direction.

The incidence of postcervical uterine torsion was much higher than that of precervical uterine torsion. According to Pearson (1971), the uterus in torsion rotates on its vaginal and mesometrial attachments and therefore, in most cases the actual twist affected the anterior vagina.

Acknowledgement

Author's thanks are due to Dr. R. C. Gupta, Professor and Head, Department of Veterinary Gynaecology & Obstetrics for his valuable suggestion and providing the facilities in the present study.

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Profiles Of Some Plasma Biochemical Constituents Associated with Uterine Torsion And Following Its Correction By Laparohysterotomy In Buffaloes

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ABSTRACT

The objective of this study was to monitor the profiles of certain plasma biochemical constituents associated with uterine torsion and following its surgical by laparohysterotomy correction in buffaloes. Blood samples were obtained from fourteen affected animals before surgical intervention for the delivery of foetus and correction of uterine torsion and from further eight animals on days 5 and 10 post-operation. Blood plasma samples were analysed for the concentrations of calcium, inorganic phosphorus, total proteins, sodium, potassium, copper, zinc and cortisol. Six of the 14 affected animals died within 24-48 after surgery and the pre-operation biochemical profiles in these animals were similar to those which surviced after surgery. In comparision to normal buffaloes at term, the affected animals had significantly lower plasma levels of inorganic phosphorus, total proteins, copper and zinc and significantly higher levels of plasma cortisol. Post-operatively, only the plasma levels of inorganic phosphorus and total proteins showed significant fluctuations, but all the constituents studied were within the range of normal values for non-pregnant non-lactating animals by day 10 postoperation. However, plasma cortisol had not returned to basal levels suggesting

that the stressful factors continued to operate by 10th day following surgical correction of uterine torsion in buffaloes.

Several earlier reports have emphasized uterine torsion as the single largest cause of maternal dystocia in buffaloes (Fouad and EL-Sawaf, 1964; Mannari and Tadkod, 1976, EL-Naggar, 1978). In a recent survey we found that uterine torsion constitutes nearly 44 percent of all maternal dystocias in buffaloes (Manju and Verma, in press). The economic impact of such a high prevalence among buffaloes could be tremendous as uterine torsion may affect subsequent lactation and reproductive efficiency and may be even fatal. Although several techniques of relieving uterine torsion are now available (Fouad and EL-Sawaf, 1964; EL-Naggar, 1978, Jit Singh et al, 1978) the success rate generaly depends upon the time-lapse between the occurence of uterine torsion and its presentation for treatment, the techniques of treatment and the post-treatment measures. An improved rate of success may perhapse be obtained if the animal is evaluated before and after correction of uterine torsion for predicting prognosis and to supplement the post-treatment regimens. These objectives have been achieved with success by the use of a number of metabolic profile tests in a variety of affections in man and domestic animals. However, their use in buffaloes affected with uterine torsion has received vertually no attention. We studied the profiles of some biochemical constituents associated with uterine torsion in buffaloes and the results presented in this paper is a part of a longterm study on the various aspects of uterine torsion in buffaloes.

Materials and Methods

This study utilized 14 murrah buffaloes presented for treatment of uterine torsion at our clinic. The details of the degree of torsion and the position and direction of torsion in these animals have been reported earlier (Manju and Verma, 1985). All the affected animals were presented at term and an attempt had been invariably made by the local veterinarian to relieve uterine torsion by rolling. We made no further attempt to relieve uterine torsion by rolling of the animal and uniformly adopted laparohysterotomy to deliver the foetus and correct the uterine torsion through an incision made lateral and parallel to the milk vein. During the course of surgery each animal received 4000 ml of 5% Dextrose saline i.v., 40 ml chromostat, i.v. (Life Plasma), Hostacortin, (Hoechst) 10 ml, i.m. and 540 ml Unimezol i.v. (Unichem). Post-operatively each animal received 60 i.u. oxytocin i.v., Dicrystacin 5 g i.m. (Sarabhai), Calborol, 540 mli.v. (May and Baker) and Dextrose saline 2 liters i.v. Calboral and Dextrose saline therapy was restricted for two days, while the antibiotic therapy was given daily for 5 days. Blood samples were drawn into heparinized test tubes from each animal immediately after the completion of clinical examination and before any treatment being instituted. Blood samples were also obtained on 5th and 10th day post-operation from 8 animals which surived following surgical intervention for correction of uterine torsion. Blood samples were also drawn, once, from a group of 6 normal buffaloes

TABLE 1: Mean plasma biochemical concentrations in normal buffaloes at term and in buffaloes before and after correction of uterine torsion by laparohysterotomy.

		PLAS	MA CONC	ENTRAT	IONS			
	Calcium (mg %)		Total pro- teins.(g%)		Potassium (Meq/L)	Copper (ug %)	Zinc (ug %)	Cortisol (ng/ml)
Normal-at term	10.72	5.95	9.38	117,66	2.60	89.58	243.0	1.11
(N=6)	±	±	土	+	土	±	±	±
	0.11	0.20	0.11	0.61	0.02	6.79	1.50	0.48
Pre-operation	9.51	3.87	7.40	108.00	2.91	57.14	148.0	63.30
(N = 14) .	±	±	±	±	±	±	± -	±
	0.75	0.48	0.36	5.27	0.47	3,62	15.0	13.70
Post-operation (N-8)								
DAY-5	11.11	6.87	8.70	116.00	2.48	71.87	93.0	61.74 "
	±	+	±	±	±	土	±	±
	0.27	0.39 -	0.22	3.02	0.14	8.09	21.0	30,54
DAY-10	10.83	5.66	7.83	107.00	2.21	81.25	148.0	11.72
1	±	±	土	±	±	+	+	±
	0.29	0.28	0.13	3.02	0.14	8,49	19.0	4.22
				12 march	1			

near term (-5 to -1 d) for comparision with the pretreatment biochemical profiles in the affected animals. We preferred to use normal buffaloes at term for this purpose because the affected animals presented to us were invariably at the completion of their gestation period.

All the plasma samples were analysed for the levels of calcium, inorganic phosphorus, sodium, potassium, total proteins, copper and zinc using standard analytical procedures, plasma cortisol levels were estimated by Radioimmunoassay using 1251 labelled kits procured from Biodata, Switzerland. The procedure followed was as advised by the manufacturers.

Results and Discussion

Of the 14 cases subjected to laparohysterotomy for the delivery of foetus and surgical correction of uterine torsion, 6 animals died within 24-48 after surgery. Four of the 6 animals showed extensive peritonitis on necropsy. These animals had earliar exhibited signs of severe dehydration and toxaemia when presented for treatment. The cause of death in the remaining two animals was probably shock associated with rolling and/or laparohysterotomy. However, the pre-operation plasma biochemical profiles of the animals which died following laperohysterotomy were similar to the pre-operation biochemical profiles of the animals which surived after surgical treatment. Therefore, we were unable to speculate the prognosis of the case based on plasma biochemical profiles. Perhaps, the interval between the occurance of uterine torsion and its presentation for treatment and the number of rollings given to the animal prior to its presentation to us might have had a bearing on the plasma biochemical profiles in both the groups of animals. Additionaly, some treatments given by the local veterinarian may also have had an influence on the plasma biochemical profiles. Unfortunately, we could not obtain the details of rolling and various treatment measures received by the animal before the cases were presented to us.

When the pre-operation levels of each biochemical constituents in the animals irrespective of whether the animal died or survived after laparohysterotomy were compared with normal buffaloes at term, the affected group of animals were found to have significantly lower values of plasma inorganic phosphorus, total proteins, copper and zinc and extremely high levels of cortisol. However, this study could not substantiate the observations of Pattabiramn and Pandit (1980) that buffaloes affected with uterine torsion had lower levels of plasma calcium. This was probably because they used animals in late gestation for comparision and plasma calcium levels are known to be higher at late gestation than at term (Muttalli, 1980). We are however, not aware of the stage of gestation at which buffaloes affected with uterine torsion were used in their study.

The low plasma inorganic phosphorus observed in the affected animals has also been recorded in cases of dystocia in cows (Bostedt 1974, Ramanarayana 1978) and tuffaloes (Muttalli, 1980). That buffaloes affected with uterine torsion had lower levels of plasma proteins has also been previously reported by Pattabiraman and Pandit (1980). Similarly Pryor (1976) speculated the possibility of an association between the zinc status of the animal and the events occuring at parturition and Dufty et al (1977) showed that cows with parturient abnormalities exhibited a dramatic fall

in plasma zinc levels. Although the present study recorded lower levels of plasma inorganic phosphorus, total proteins, copper and zinc in the affected group of animals, we are unble to state whether these differences were as a result of different managemental conditions under which each affected animal was maintained since the control group of animals used for the purpose of comparision were all from a single herd maintained under uniform conditions of feeding and management. Nevertheless, animals affected with uterine torsion had consistantly higher plasma levels of cortisol than in the control group of animals. Therefore, we suggest that uterine torsion induces considerable stress on the animal and the lower levels of plasma inorganic phosphorus and zinc in the affected animals are perhaps an indirect reflection of the stressful events occuring in uterine torsion as there is some evidence that their levels are influenced by cortisol (Wagner, 1973; Wilkinson, 1980).

The plasma levels of calcium, sodium, potassium, copper and zinc did not exhibit significant fluctuations during the post-operative period. Therefore this study could not confirm the observations of Bestedt (1974) that cows with parturient abnormalities had low levels of plasma calcium which sustained for a longer duration and that of Mousely and Oxford (1971) who demonstrated a 10 percent fall in plasma calcium levels within one hour after minor surgery in sheep. Perhaps, a more frequent sampling schedule than the one used in this study may have provided a more clear picture of the alterations in the plasma profiles of calcium following laparohysteromy in buffaloes affected with uterine torsion. The non-significant differences in plasma

calcium values before and after laparohysteromy may also be due to the fact that the affected animals received calcium borogluconote intravenously in 2-3 divided doses during the first 3 days post-operation. A almost steady levels of plasma calcium, sodium and potassium recorded during the post-operative period may suggest that the fluid and electrolyte therapy given to these animals was adequate.

Among other plasma constituents studied, inorganic phosphorus showed a dramatic increase by 5th day post-operation and although the levels were marginally lower by 10th day, its concentration was still higher in comparision to the pre-operation values. On the otherhand, the plasma total proteins had increased to significantly higher levels by day 5 post-operation, but fell significantly by day 10. We have no explanations for these fluctuations, but considering the fact that none of the affected animals were lactating Post-operatively the plasma levels of inorganic phosphorus and total proteins were within the range of values for non pregnant nonlactating animals (Kaneko, 1980).

By far, the most dramatic change observed was in the plasma levels of cortisol. Its level by day 5 post-operation, was similar to the pre-operation concentrations suggesting that the affected animals continued to be under a spell of stress. By day 10, however, the plasma cortisol levels had appreciably reduced suggesting that the animals were on the course of recovery. However, the plasma cortisol remained slightly elevated and did not approach the basal values for non-pregnant, non-lactating animals suggesting that stressful factors continued to operate by day 10 post-operation.

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Studies On The General Characteristics Of Caprine Foetal Fluids

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ABSTRACT

Foetal fluids from twelve pregnant goats during different stages of gestation were studied for their general characteristics like colour, consistency, pH. and specific gravity. The light yellow colour of the allantoic fluid changed to intense brown, whereas the amniotic fluid became colourless, with the advancement of gestation. The allantoic fluid remained watery throughout the gestation, however the amniotic fluid became turbid and mucoid in the second half of gestation. The pH of allantoic fluid decreased with the advancement of gestation and chauged finally to acidic from alkaline in the beginning. The pH of amniotic fluid also followed the same trend but it again became lightly alkaline in the last stage of gestation. There was no significant change in the specific gravity of both the fluids.

The caprine foetus is surrounded by two fluids: i.e. an inner amniotic fluid and an outer allantoic fluid throughout the gestation. These foetal fluids play an important role in the efficient handling of the foetal waste products and in preventing mechanimal shock to the developing foetus during the entire gestational length. Furthermore, the fluids are also helpful during the process of parturition. The characteristics of the allantoic and amniotic fluid have been studied in Cattle (Morgan and Whitehair, 1943; Arthur, 1965) and Sheep (Medougall, 1949; Mellor and Slater, 1971). However the goat has remained an exception in this respect for a long time.

Materials and Methods

The study was carried out on a total of 12 pregnant goats of different gestational lengths. Stage I had animals of 1-2 months of gestation, Stage II had animals of 2-3 months of gestation, Stage III had animals of 3-4 months of gestation and Stage IV had animals of 4-5 months of gestation. Each had three animals.

The pregnant uteri were exposed by mid ventral laprotomy after inducing local anesthesia (2% Gesicain) at mid ventral line just anterior to the udder. A small incision in the uterus was made and using a sterile 20 ml glass syrige, attached with a 18 gauze needle about 20 ml of fluid was collected separately from each placental sac.

The physical characteristics i.e. colonr, consistency, pH and specific gravity of both the fluids were studied immediately after their collection.

Results

The colour of the allantoic fluid changed considerably during different stages of pregnancy. During stage I of gestation the colour was light yellow and there after the colour changed to light brown during stage II and intense brown during the III and IV stages of gestation. The Amniotic fluid was also slightly yellowish during stage I of gestation, but changed to a colourless fluid during the II, III, and IV stages.

TABLE 1. Mean values for pH of allantoic and amniotic fluid

Stage of gestation	Allantoic fluid	Amniotic fluid
1	7.35 ± 0.06	7.31±0.07
II	6.84 ± 0.32	6.93 ± 0.10
IH	6.48 ± 0.18	6.82 ± 0.08
IV	6.84 ± 0.09	7.06 ± 0.17

Data are expressed as values \pm standard error.

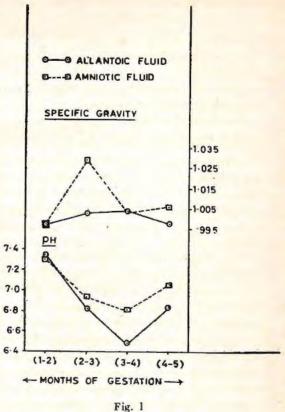
The consistency of allantoic fluid remained watery throughout gestation. However, the amniotic fluid became turbid and mucoid in the second half of gestation.

The pH of allantoic fluid was 7.35 during the I stage of gestation and thereafter there was a fall in the pH to 6.84 and 6.48 during II and III stages respectively. In the last stage of gestation, it however increased to 6.84 (Table-1). However, this fall in the pH of allantoic fluid during III and IV stage was significant (P \angle 0.05) when compared with the stage I of gestation. (Fig. 1)

 TABLE 2: Mean values for specific gravity of allantoic and amniotic fluid.

Stage of gestation	Allantoic fluid	Amniotic fluid
I	0.997±3.21	0.997 ± 4.09
II	1.003 ± 5.01	1.029 ± 0.02
III	1.004 ± 0.009	$1.003 \pm .008$
IV	0.998 ± 4.48	1.006 ± 0.02

Data are expressed as values ± standard error.



rig. I

The pH of amniotic fluid also decreased with advancement of gestation from 7.31 during I stage to 6.93 and 6.82 during II and III stages of gestation, respectively. The decrease in the pH in stage III was highly significant (P \angle 0.01) in comparison to stage I. The pH however rose to 7.06 during the IV stage of gestation. (Table 1, fig. 1).

There was no significant change in the specific gravity of allantoic as well as amniotic fluids throughout gestation (Table-III).

Discussion

The general characteristics of bovine foetal fluids are influenced by exchange of biochemical constituents through the placenta, foetal urine formation and its flow through the urachus or urethra and foetal secretions from lungs and salivary glands (Baetz et al, 1976).

Arthur (1965) oberved that the foctal fluids of bovine were like pale urine, and as the gestation advanced the allantoic fluid became more urine like whereas the amniotic fluid turned to a colourless and slightly viscous fluid.

In the present study the amber coloured allantoic fluid of early gestation changed to light brown and intense brown in late gestation, whereas the amniotic fluid changed to a colourless, mucoid and gelatinous fluid from yellowish in the early gestation. Similar changes in goats were also reported by Bongso et al. (1979).

Arthur (1965) observed that in cattle the changes in the colour of both the fluids and consistency of amniotic fluid with advancing gestation may be due to the onset of salivary secretion and increased mucous secretion from the alimentry tract and/or to decreased flow of urine in to the amniotic cavity as the sphincter of the urinary bladder began to function between $6\frac{1}{2}$ to $7\frac{1}{2}$ months of gestation. Similar onset of the function of the bladder sphincter probably occurs earlier in case of goat (Bongso et al, 1979). The mean pH of the amniotic fluid for all stages was 7.03. Initially during the I stage, the pH was alkaline (7.31). However, during the II and III stages the pH was acidic which again changed to alkaline during the IV stage of gestation (fig. 1). The pattern of change of pH was similar to the findings reported in Cows (Morgan and Whitehair, 1943).

The initial alkaline pH of the allantoic fluid (7.35) during I stage, changed to acidic during the subsequent stages of gestation. Meller & Slater (1971) also observed the amniotic fluid to be alkaline (pH 7-7.50) in comparison to allantoic fluid (pH 6.00 - 7.40).

McDougall (1949) reported that in sheep amniotic fluid had a lower specific gravity than the allantoic fluid. In the present study although there was no significant change in the specific gravity of both the fluids across gestation yet the mean specific gravity of allantoic fluid for all stages (1.001) was slightly lower than for the amniotic fluid (1.009). This may be due to the change in the consistency of the amniotic fluid and/or to the higher solnte concentration in the amniotic fluid than in the allantoic fluid.

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Studies On Heteroplasmic Preservation Of Buck Semen

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ABSTRACT

Addition of 50% bull seminal plasma to buck semen increased the percentage of progressively motile spermatozoa and live sperms. Abnormal sperm percentage was also decreased. It increased the "heteroplasmic vigour" of buck spermatozoa, the range was between 11.15 (reduction in percentage of abnormal spermatozoa) and 32.33 (percent increase in progressive motility) in Black Bengal bucks and in Barbari 10.85 (live sperm percentage) and 28.32 (progressive motility percentage). A rough estimate of overall superiority of treatment over control in Black Bengal and Barbari was 23.40 and 19.59 respectively.

* *

Successful preservation of semen is essential for the implementation of any breeding programme. Heterospermic preservation has been reported to increase the keeping quality of semen (Narayan and Singh, 1971 and Sinha, 1981). This study reports the results of an experiment conducted to estimate the effect of bull seminal plasma used as an additive in the buck semen.

Materials and Methods

Eight bucks (Four of Black Bengal and four of Barbari) and two Holstin Friesian bulls maintained at Ranchi Veterinary college, Ranchi were used in this study. Semen was collected from the bulls on alternate day at about 6.30 A.M. by the artificial Vagina method. Soon after collection the seminal plasma was separated by centrifuging it for 15 minutes at the rate of 3000 RPM.

The Egg-yolk-citrate (Salisbury et al, 1941) was used as semen dilutor.

Bull seminal plasma (BSP) and Eggyolk-citrate dilutor in the ratio of 1:1 (50% BSP) were mixed. This was prepared by mixing 4 ml each of bull seminal plasma and Egg-yolk-citrate.

Each prepared sample was divided into two equal halves in order to dilute the semen of two bucks (Black Bengal and Barbari). Egg-yolk-citrate constituted the control for both the bucks.

Bucks were maintained under the similar managemental conditions throughout the period of the experiment. Collections were obtained from one Black Bengal and one Barbari buck, on every alternate day at about 7.30 A.M. by means of artificial vagina method. After evaluating the quality of the ejaculates collected from bucks, the semen was diluted with the prepared dilutors. Thus, there were two samples from each collection i.e. control (Containing EYC as dilutor) and treatment (containing 50% BSP and 50% EYC). The samples were strored in 8 pint wide mouth thermoflask filled with ice. Stored semen samples were evaluted at 24,48,72 & 96 hours of preservation for the following parameters:

Breed and		Progress	sive motili	ty percent	age		live sp	erm perce	ntage		Percent	age of abn	ormal spe	rmatozoa	
Treatment	24 Hr.	48 hr.	72 hr.	96 hr.	overall	24 hr.	48 hr.	72 hr.	96 hr.	overall	24 hr.	48 hr.	72 hr.	96 hr.	overall
Black Benga	al	-													
Control	64.71a	53.48a	45.02a	34.73a	49.48	84.42ª	78.80a	74.18ª	66.624	76.00	12.67	14.39a	15.714	17.86ª	15.15
	±0.889	±1.175	±0.979	±1.253	± 1.411 (32.33)	±0.685	±0.985	±0.958	±1.369	±0.922 (14.46)	±0.677	±0.696	±0.734	±0.751	±0.412 (11.15)
Treatment	73.586	68.73b	64.776	60.77b	66.96	90.30%	88.16b	86.06b	83.44b	86,99	11.64	12.716	14.016	15.510	13.46
	±0.494	± 0.515	±0.505	± 0.396	±0.068	±0.604	±0.522	± 0.528	±0.446	±0.387	± 0.602	± 0.559	± 0.577	± 0.583	±0.332
Barbari Control	64.86¢	56.65¢	48.39¢	37.32¢	51.80	85.480	82.130	77.410	70.44¢	78.85	12.27	13.700	15.59¢	18.15¢	14.92
	±0.684	±0.688	±0.715	±1.150	± 1.273 (28.32)	±0.713	±0.858	±0.751	±2.207	0.916 (10.85)	±0.686	±0.730	±0.772	±0.871	±0.458 (11.05)
Treatment	t 73.31d	68.67d	63.84d	60.08d	66.47	90.71d	88.44d	86.28d	84.23d	87.41	11.42	12.68d	13.75d	15.25d	13.27
	±0.496	±0.528	±0.563	±0.526	±0.640	±0.512	±0.503	±0.470	±0.474	± 0.360	± 0.718	± 0.662	± 0.679	± 0.694	± 0.374

TABLE 1. Average values of seminal attributes in black bengal and barbari bucks at different hours of preservation.

Figures bearing different superscripts in a column within a bred differed significantly. Values in parentheses indicated percent superiority of treatment over control.

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x

- 1. Motility percentage of spermatozoa. "Un
- 2. Live percentage of spermatozoa and
- 3. Percentage of abnormal spermatozoa.

Motility percentage of the diluted and stored semen samples was studied according to slightly modified method described by Lasley (1951). The live percentage of spermatozoa in each of the samples was estimated by using Eosin-Nigrosin staining technique mentioned by Swanson and Bearden (1959). Percentage of abnormal spermatozoa was determined from the same slides used for determination of live spermatozoa percentage.

The data recorded on different parameters were subjected to statistical analysis according to Snedecor and Cocharan (1968).

Results and Discussion

Average values of seminal attributes in Black Bengal and Barbari bucks at different hours of preservation have been shown in Table I.

Addition of bull seminal plasma increased the percentage of progressively motile spermatozoa and live sperms in the semen during different hours of preservation. It also decreased the abnormal sperm percentage, thus increasing the quality of semen in both the breeds. Analysis of variance indicated significant effect of treatmen on all the attributes.

Addition of 50% BSP increased the "heteroplasmic vigonr" of buck spermatozoa, the range was between 11.15 (reduction in percentage of abnormal spermatozoa) and 32.33 (percent increase in progressive motility) in Black Bengal bucks. The corresponding values in Barbari were 10.85 (live sperm percentage) and 28.32 (progressive motility percentage).

A rough estimate of overall superiority (Taking average of the percent superiority of treatment group over control for progressive motility and live sperm content) of treatment over control in Black Bengal and Barbari was 23.40 and 19.59 respectively.

These results indicate that increased quantity of bull seminal plasma possibly supplemented both quantitatively and the biochemical ingredients required for enhancing and maintaining the activity and life span of buck spermatozoa for a longer period, as well it checked an increase in their morphological abnormality during preservation. Biochemical analysis of semien, seminal plasma and sperm cells of different species/breeds of livestock separately and in different combinations may be helpful in explaining the heteroplasmic superiority of "heteroplasmic vigour".

Acknowledgement

The anthors are very much thankful to the Dean, Ranchi Veterinary College and ICAR for utilising the data from AICRP on goats, chotanagpur unit, Ranchi, The first author is thankful to the government of Jammu and Kashmir for financial assistance.

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Studies On Inter-Lambing Period In Indian Breeds Of Sheep

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ABSTRACT

A study on inter-lambing period was carried out to determine the effects of seasons and breeds on this trait in four breeds of sheep-Bikaneri, Nali and Hissar Dale at Hissar farm and Muzaffarnagri at the I.V.R.I., Izatnagar farm. Interlambing period averaged 335.73+7.17 days in Bikaneri, 341.31+4.62 days in Nali, 434.70+13.92 days in Hissar Dale and 330.48+8.34 days in Muzaffarnagri. Hissar Dale had significantly longer interlambing period than that of the other 3 breeds which were not significantly different from each other. Analysis of variance further revealed that seasons had significant effect on interlambing period in Bikaneri and Nali breeds. Seasonality did not seem to have significant effect on inter-lambing period in Hissar Dale and Muzaffarnagri breeds.

Inter-lambing period is an important trait which has a bearing on the reproductive and productive life of sheep. Studies on factors affecting inter-lambing period in sheep are important but such studies are few. Sahni and Roy (1967) reported that tropical sheep, Bikaneri is nonseasonal in their sexual activity. In 1972 they further reported that the monthly distribution of lamb births in a flock and of Bikaneri ewes in which rams were run continuously for several years showed lack of seasonal variation in concentration of lamb births in tropics. Taparia

(1972) reported non-seasonality of oestrus activity in Sonadi sheep.

Further studies were required on other Indian breeds of sheep on inter-lambing period. In the present investigation, these have been carried out in four breeds of sheeps, namely Bikaneri, Nali, Hissar Dale and Muzaffarnagri.

Materials and Methods

Inter-lambing period was studied on 202 Bikaneri, 176 Hissar Dale and 581 Nali ewes at Hissar Farm. Two hundred fifteen ewes of Muzaffarnagri breed maintained at the Indian Veterinary Research Institute (I.V.R.I.), Izatnagar farm were also included in this study. Most of the ewes of different breeds were recorded for single inter-lambing period. However, two or more than two interlambing periods per animal were also recorded in few animals. The climate at Izatnagar (U.P.) and Hissar (Haryana) was divided into four seasons on the basis of humidity and temperature as follows:

Izatnagar Climate

- 1. Cold season (second half of November to February)
- ⁹ 2. Temperate season (second half of September to first half of November and March to first half of April)
 - 3. Hot dry season (second half of April to first half of June.

4. Hot humid season (second half of June 1 to first half of September).

Hissar Climate

1. Cold season (November to January)

2. Temperate season (second half of October, February and March)

- 3. Hot dry season (April to June)
- 4. Hot humid season (July to first half of October)

 TABLE 1: Interlambing period in different breeds. (days).

Bikaneri	Nali	Hissar Dale	Muzaffarnagri
357.734	341.314	434.706	330,48a
±7.17	±4.62	±13.92	±8.34
(332)	(960)	(243)	(264)

Averages with the same superscript are not significantly different at 5% probability level from each other. Figures in parentheses are number of observations.

For seasonal calculation the interlambing period was considered as the lambing in that particular season from there the time taken by lambs again. Standard statistical procedures were used for analysis.

Obeservation on inter-lambing periods available for study in different breeds were as follows:

Bikaneri 332, Hissar Dale 245, Nali 960 and Muzarffanagri 264.

Results and Discussion

INTER-LAMBING PERIOD

The inter-lambing period averaged 335.73±7.17 days in Bikaneri, 341.31

 ± 4.62 days in Nali, 434.70 ± 13.92 days in Hissar Dale and 330.48 ± 8.34 days in Muzaffarnagri breeds of sheep, respectively (Table 1). It was longest in Hissar Dale which differed significantly from other three breeds. However, the difference in the inter-lambing periods of Bikaneri, Muzaffarnagri and Nali was not significant (Table 1). Incidentally, Hissar Dale breed had the maximum age at first lambing which also indicated sexual maturity in this breed at a slower rate as compared to other breeds of sheep studied.

Analysis of variance (Table 2) revealed that the seasons had a significant (P< 0.01) effect on inter-lambing period in Bikaneri and Nali breeds. Season 2 appeared to have favoured the interlambing period while season 1 adversely affected this trait and it differed significantly from the other three seasons in Bikaneri breed (Table 3). Seasons 1, 3 and 4 did not vary significantly from each other in Bikaneri breeds. The interlambing period during season 2 was significantly shorter than that during seasons 1, 3 and 4 in Nali breed (Table 3). Averages of iner-lambing period for seasons 1, 3 and 4 did not differ significantly in Nali breed. The inter-lambing period was shortest during season 2 in both the breeds i.e. Bikaneri and Nali. Seasonality did not seem to have significant effect on inter-lambing period in Hissar Dale and Muzaffarnagri breeds (Table 3),. Breed differences were highly

TABLE 2: Inter-lambing period for different seasons within breeds.

	-			ANOV	A			
Source of	Bika	neri		Nali	Hisse	r Dale	Muza	farnagri
variation	df	MS	df	MS	df	MS	df	MS
Seasons Within seasons	3. 328,	108323.8** 16276.8	3 956	291771.9** 19674.9	3 239	55503.3 46969.9	3 260	12387.7 18436.0
	1	Y Har to	** P	< 0.01			2.7	1

70

ent S a	Seasons		Breeds	ente estand Rectante d	lat i a
	1000 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	Bikaneri	Nali	Hissar Dale	Muzaffarnagri
	I. Cold	373.364	360.66 ^b	459.79	3 321.75 2 1g /
		±11.96	± 6.96	± 18.53	± 19.01
· · ·		(125)	(323)	(104)	(66)
	2. Temperate	300.28b	298.32a	429.80	311.26
		± 16.36	±7.81	± 24.35	± 24.85
		(82)	(301)	(86)	(35)
	3. Hot dry	329.826	368.416	371.00	329.82
	I HAR - Is - Sec	±11.98	±9.02	± 72.86	± 17.35
		(85)	(265)	(13)	(75)
	4. Hot humid	300.356	334.480	400.70	\$45.23
		± 14.28	± 22.62	± 39.09	±10.46
		(40)	(71)	(40)	(88)

TABLE 3. Inter-lambing period (days) for different seasons within breeds.

Seasons averages with the same superscript are not significantly different at 5% probability level. Figures in parentheses are number of observations.

significant (P < 0.01) for inter-lambing period studied for Hissar Dale, Nali and Bikaneri breeds which were located at the same farm (Hissar) (Table 4).

The ewes in the present study were at different stages of parity. It was not possible to determine the intervals between first and second lambing, second and third lambing and so on. Data, therefore, where analysed for intervals between lambing and those for subsequent ones.

The results revealed that in the Hissar Dale breed, inter-lambing period was the longest and the differences were significant (P < 0.05) as compared to other three breeds. Differences among the other two breeds at Hissar farm and Muzaffarnagri breed at Izatnagar farm were not significant.

Narayanswamy et al. (1976) reported lambing interval in Bannur (Mandya) sheep which averaged 304.42 ± 11 days. This value is quite lower in comparison to the values observed in the present study. This night be due to breed difference or by virtue of its being controlled and influenced by the factors other than genetic ones and can be reduced through improved managemental practices.

Sex of the lamb had no effect on any of the inter-lambing periods, which is not in agreement with the findings of Narayanswamy *et el.* (1976) who reported that

TABLE 4:	Inter-lambing	period	for	Bikanei,	Nali	and	
	Hissar Dale br	eeds					

Source of variation	df	MS
Breeds	2	923274.9**
Within breeds	1532	23982.1

ewes carrying male lambs had longer lambing interval by 4.3 days than the ewes carrying female lambs and thids difference due to sex was highly significant. Acknowledgement The first author is thankful to the Director, Indian Veterinary Research Institute for research facilities and to Indian Council of Agricultural Research for financial assistance.

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Studies on the Regeneration in Macrostomum orthostylum (BRAUN) (Turbellaria: Macrostomida)

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ABSTRACT

The regeneration process was observed in the laboratory cultured macrostomid turbellarian, *Macrostomum orthostylum*. Recovery of cut portions was very fast at the anterior region as compared to the posterior region. The average regeneration rate observed was 0.069 mm/day. However, regeneration was not observed in the region anterior to the cerebral ganglion.

*

Information available on the regeneration in free-living Turbellaria indicates that acoelid turbellarians have limited power of regeneration (Ghild, 1907; Peebles 1913 and Keil 1929). In the case of alloeocoels and macrostomid, Ruhl (1927) has reported that the former are incapable of any regeneration and the latter whilst incapable of a sequal reproduction, can regenerate, particularly towards the tail portion.

Keeping in view the paucity of information on the regeneration in M. orthostylum (BRAUN), the present studies was undertaken and results are reported here.

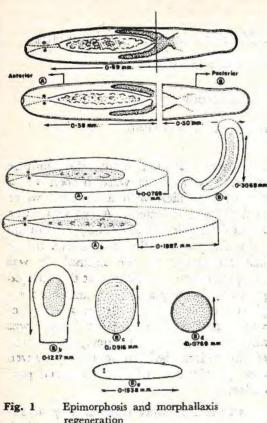
MATERIALS AND METHODS

A culture of *Macrostomum orthostylum* was maintained in the laboratory in glass beakers following the method described by Shirgur (Shirgur 1980). For the present study, healthy adult specimens of uniform size, were isolated from the culture stock. The animals were placed on a microscopic slide with a drop of water and allowed to extend. A cut was made at desired part of the body of animal by a sharp razor. Sometimes due to the fast movement of the animal, it was not possible to make a cut at the desired part of the body and thus, animals were placed on a cavity slide in a drop of water, and then a drop of PHCl₃ was added by a fine dropper. Movement of the animal was subsequently slowed down, making a cut at the desired part easier.

Cut portions were immediately transferred into separate glass cavity block (2 ml capacity) having culture media of 9% salinity. Observations on the cut portion were carried out every day under a low power disecting microscope. The figures, of different developmental stages of the cut portion, were made using a camera lucida, while all measurements were taken by a micrometer.

RESULTS AND DISCUSSION

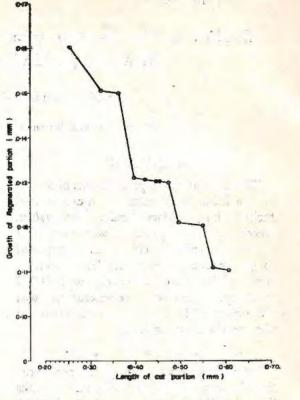
Macrostomum arthostylum, an easyhaline turbellarian, recorded first time from Indian water (Shirgur 1980) and cultured in the laboratory for nutritional requirements of hatchery level prawn seed production (Shirgur and Ingole 1983). It has been observed in the mass culture of *M orthostylum* that sometimes the tail portion of the animal gets detached in



- regeneration 1.4 14
 - Anterior cut portion
 - 1Aa Anterior portion regenerating its lost portion
 - 1AB Anterior portion completely regenerated
- Poeristor cut portion Fig. 1B
 - 1Ba Posterior cut portion changed into curve shape
 - 1Bb Curve shape changed into cap like structure
 - Cut portion transformed into spherical 1Bc shape
 - 1Bd Egg-shaped cut portion
 - 1Be Newly hatched youngone

1-1-1-1

the course of laying of subitaneous eggs and regenerates to a new young animal. Specimens below 0.4 mm length were immature and did not show morphallaxis type of regeneration. Therefore, only the adult specimens measuring more than 0.4 mm, which were found to undergo



Growth of regenerated portion Fig. 2

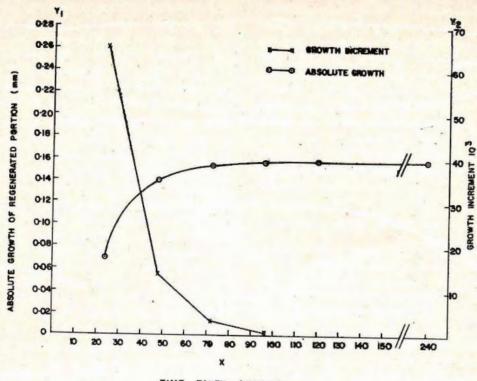
epimorphosis and morphallaxis regeneration, selected in the present study.

Process of Regeneration

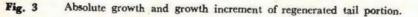
The animal measuring 0.69 mm (Fig. 1) was cut into two portions — an anterior portion of 0.38 mm and a posterior portion of 0.30 mm (Fig. 1, A & B.)

The regeneration of the lost portion at the anterior end was epimorphosis type. Growth in tissue mass was very fast resulting in the formation of complete animal after five days (Fig. 1A a; and 1A b).

To study the regeneration of posterior portion, the cut part was kept in a separate cavity block in fertilized culture media, where it showed circular movements



TIME TAKEN (HOURS)



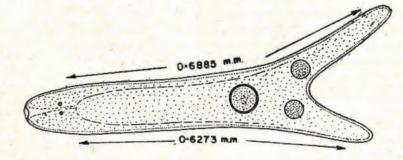


Fig. 4

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Animal with bifurcated tail portion

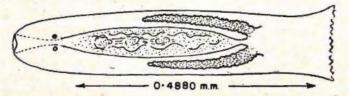


Fig. 5 Animal with severed tail portion

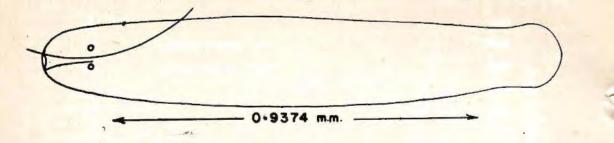
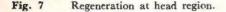


Fig. 6 Cut showing removal of eye.





at first and then finally settled at the bottom of the cavity block. One end of the portion was attached to the bottom and undulation of the body occurred. The type of regeneration in this portion was totally morphallaxis type. At first, the cut portion drummed to a curveshaped structure measuring 0.3068 mm in length (Fig. 1B a). At the end of 4th day of observations, the curve-shaped portion changed to a cap like structure at the distal end with a length of 0.1227 mm (Fig. 1B b). However, no change was observed on the 5th and 6th days. The entire portion was transformed into a egg shaped structure having diameter of 0.0916 mm, at the end of 7th day (Fig. 1B c). At the end of 8th day, the shape of the cut portion became exactly similar to that of a subitaneous egg with a diameter of 0.0765 mm (Fig. 1B d) and then a single young animal was hatched out at the end of 11th day of observation (Fig. 1B c.) Discussing the similar type of regeneration in other species Hyman (1951) and Pascolini (1968) have reported that a piece of moderate size from any level forms a complete new worm. While discussing the process of cell totipotency in *Digesia* gonocephalla, Gremigni (1972) has reported that a single cell of somatic tissue develops a complete organisms.

Absolute growth and growth increment of the regenerated tail portion.

It was observed that an anterior portion normally regenerated its lost portion within five days some observations were made making cuts at different body line towards the anterior end. The pattern of regeneration was observed in twelve successive experiments and the average rate of regeneration calculate. In the case of an anterior cut portion of 0.25 mm in length, the length recovery

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of the regenerated portion was 0.16 mm at the end of 120 hours (Fig. 2). In the third experiment of the series, in a cut portion of 0.45 mm length; the regenerated part was 0.13 mm, while 0.11 mm in the case of a 0..60 mm cut portion, at the end of 120 hours.

Absolute growth was 0.07 mm after 24-hour interval and 0.14 mm after a 48-hour interval. At the intervals of 72 and 96 hours absolute growth were 0.15 mm and 0.16 mm respectively, while at the end of 120 hours an average growth of 0.16 mm was observed which remained constant subsequently. Average growth rate was 0.069 mm/day, whereas the growth increments calculated from 12 experiments were 0.0681 mm, 0.0139 mm, 0.0037 mm and 0.0035 mm at the end of 24, 48, 72 and 96 hours, respectively (Fig. 3). It was found that the growth rate of the regenerated portion was faster at first (0.24 hours) in the anterior region. Fast growth rate of this portion may be due to the presence of extraordinary high number of undifferentiated cells in the parenchyma. Germigni and Banchetti (1972) have also reported the presence of extraordinary cells in the parenchyma of Dugesia gonocephala. Fast growth increment indicates the high activity of regenerating cells within a determined period. It is also reported in Digesia mediterranea and Dugesia tigrina that the rapid accumulation of neoblast cells at the beginning of regeneration and a decrease after 2-3 days (Baguna and Romero, 1981).

Bifurcation at tail portion (Fig 4)

The bifurcation of a tail portion was

usually noticed in the case of an individual which sustained a severed tail portion during the release of large subitaneous egg (Fig. 5). Further observations on such animals show that they reproduce normal hatchings (without biaxial tail). However, some observations were also made in case of the selected adult specimens by making cuts in the tail region. It was observed that their tails regenerated into normal form, without bifurcation.

Observations at the head region

Cuts were made at the anterior end to observe the power and pattern of regeneration of structures in the head region. A cut just posterior to the eyes, was observed to replace its lost portion within few days. However, when a portion of head with one eye of the animal was detached (Fig. 6), it was observed that the animal grew to a normal size without further replacement of detached eye (Fig. 7). This is further supported by Peebles (1913) finding, where he has reported that the portion removed anterior to the cerebral ganglie replaced perfectly but without replacing the cerebral ganglia and stytocyst.

Acknowledgement

The authors are thankful to late Dr. M.R. Ranade, Associate Dean, Marine Biological Research Station, K.K.V. Ratnagiri, for facilities. Thanks are extended to Dr. A.H. Parulekar and Dr. Anil Chatterji, Scientists, NIO, Goa, for going through the manuscript.

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SHORT COMMUNICATIONS

Phosphatases In Seminal Plasma Of Buffalo Bulls

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Bovine semen contains high concentration of Acid phosphatase but comparatively low concentration of Alkaline phosphatase. The acid phosphatase in mammalian semen is produced by the prostate gland. (Mann, 1964).

Although the enzyme Alkaline phosphatase and Acid phosphatase in buffalo semen have been studied extensively by various workers (Chauhan and Srivastava, 1973; Mishra et al., 1969 and Sen Gupta etal, 1963), there are no reports on the activity and kinetics of Fructose 1:6 bisphosphatase (FruP2ase) and Glucose 6 Phosphatase (G6Pase) in buffalo semenal plasma. Sheth and Panse (1979) have reported the effect of antiandrogens on FruP2ase in rat caudal spermatozoa.

The present communication reports the activity and kinetic properties of these enzymes.

Buffalo seminal plasma was obtained from four bulls by centrifuging the semen samples at 1400xg and the experiments were replicated. Acid and Alkaline Phosphatase was assayed with the beta glycerophosphate substrate by the method of Bodansky (1932) Glucose 6 Phosphatase and Fructose 1:6 bisphosphatase were assayed by the method of Swanson (1950) and Pogell and McGilvery (1952) respectively, Pi liberated was estimated by the method of Chen et al (1956) and Protein by the method of Lowry et al. (1951). The specific activity of the enzyme is expressed as micromoles of piliberted per mg of protein per hour at 37°^c. No experiment was done on the pH optima of G6Pase.

The activity and kinetic properties of the phosphatase in the seminal plasma are presented in Table 1.

It is clear from the table that acid phosphatase activity was very high as compared to alkaline phosphatase.

The FruP2ase activity was considerably low as compared to acid and alkaline phosphatase. G6Pase, was comparatively extremely low. The lowest Km was observed in case of G6Pase and FruP2ase. The pH optima of the Fru-P2ase and alkaline phosphatase were similar whereas acid phosphatase showed a pH optima of 5.0.

In order to verify whether the enzyme present in the seminal plasma is specific for the catalytic hydrolysis of FruP2ase does it represent conventional alkaline phosphatase activity the Fru2P in the reaction mixture was replaced by F6P and the rate of hydrolysis was measured. It was observed that the rate of hydrolytic rate of FruP2.

That the specific phosphatase FruP2ase and G6Pase do not represent the generalized alkaline phosphatase is also confirmed by the low Km value as compared to the high Km of acid and alkaline phosphatase.

A high level of alkaline and acid phosphatase has been reported in buffalo

Enzyme	Activity	Km	pH
Acid Phosphatase	68.0	4.0 mM	5.0
Alkaline phosphatase	18.5	6.4 mM	10.0
FruP2ase	3.78	9.9 UM	9.5
G6Pase	0.40	6.6 UM	

TABLE 1 Activity and kinetic properties of phosphatases in Buffalo seminal plasma

semen. (Chauhan and Srivastava, 1973). It is well known that high level of inorganic phosphorus inhibited the respiration and viability of the spermatozoa which may be one of the factor for the poor preservation and freezability of buffalo semen. (Srivastava 1979)

The enzyme FruP2ase is a pacemaker in gluconeogenesis and involved in second stage to overcome energy barier in formation of carbohydrate from non carbohydrate precursors, suggests the operation of gluconeogenesis. Further, Sheth and Panse (1979) have shown the enhancement of FruP2ase activity in rat caudal spermatozoa by administration of antiandregens, thereby inhibiting glycolysis. Mohri et al. (1975) have reported that the flux of carbon ceases due to inhibition of some of glycolytic enzymes by male antifertility agent alpha chlorohydrin in ram spermatozoa. Due to the lack of sufficient energy gained from the carbohydrates, the ram is therefore unable to ejaculate spermatozoa for its essential activities, such as motility in the femal tract. This may render them infertile. It appears that the glycolytic-gluconeogenic enzyme play a significant role in the fertility of spermatozoa.

Acknowledgement

Our thanks are due to Dr. M.R. Patel the ex. Dean of veterinary college, Jabalpur for providing the facilities and keen interest in the work.

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Anoestrus In Buffaloes-Treatment With "Estrona"

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ABSTRACT

The present trials were undertaken to study the efficacy of "Estrona" Capsules in induction of oestrus and fertile oestrus in buffaloes. Out of a total of 40 animals, oestrus was induced in 27 (67.50%) animals of which 18 (66.6%) animals conceived. The mean time interval for oestrus induction was 15.14 ± 2.68 days and for conception it was 31.9±5.37 days. Among control group of 10 animals, 3 (30.0%) came in oestrus and only One (10.0%) conceived The mean time interval for exhibition of oestrus was 36.5+7.83 days, and the only animal conceived took more than 90 days for conception.

Both the doses (2 capsules/day for 5 days and 3 capsules/day for 3 days) tried were found effective for induction of oestrus and fertile ocstruses in buffaloes.

Anocstrus is one of the most common problem encountered in buffaloes which affects the farmers economy by reducing calf crop and milk yield. several indigenous drugs have been tried to induce oestrus and fertile oestrus in anoestrus animals (Galhotra *et el.*, 1970; Porwal *et al.*, 1976; Kodagali *et al.*, 1981; Shah and Kodagali, 1984) but literature on the use of ayurvedic product like "Estrona" is very scanty. Hence the present study was undertaken to assess the efficacy of "Estrona" in induction of oestrus and conception in anoestrus buffaloes and buffalo heifers.

Materials and Methods

A total of 50 true anoestrus buffaloes maintained by farmers of Vatva Village of Ahmedabad district were included in this study, out of which 40 animals were involved in treatment trials (with Estrona capsules) and 10 animals were kept as control group. The buffaloes were considered to be true anoestrus based on two gynaecological examination at 8 days interval confirming the absence or lack of Graafian follicles or corpus luteum on the ovaries.

The composition of Estrona capsule is as follows:

1.	Loha Bhasma	50 mg,
3.	Rubia Cardifolia	100 mg,
2.	Myrsh	50 mg
4.	Aloes indica	50 mg
5.	Harmal	50 mg,
6.	Magella Sativa	50 mg
	Gogul	50 mg.

This is an ayurvedic product of Rakesh Pharmuceuticals, Ahmedabad.

CLINICAL TRIALS:

Estrona capsules were administered orally at random in 40 animals confirmed to be true anoestrus.

DOSE EFFECT:

Two different doses were tried, 2 capsules/day for 5 days and 3 capsules day for 3 days. Vaginal inspection and rectal palpations were made at an interval of 8-10 days and findings were recorded.

TABLE 1	Results of Estrona for oestrus and fertile oestrus induction in treatment vs control group	Ł
	of buffaloes.	

Sr. No. ITEM		Group				
		Treatment			Control	
		No.	Percent	No.	Percent	
1. No. of	total Animlas	40	100	10	100	
2. Oestru	induced/occured	27	67.5	3	30.00	
3. Pregna	ncy	18	45.00	1	10.00	
f. Pregna	ncy rate among served	_	66.60		33.30	
5. Interva	1 in days for oestrus induction	15.14	±2.68	36.50	±7.83	
6. Interva	l in days for fertile oestrus induction	31.9	±5.37	>90		

TABLE 2 Comparison of results with two different doses of Estrona capsules

Sr.	No. It	em		Group			
			2	Cap/day	3 (cap/day	
			No.	Percent	No.	Percent	
1.	No. of animals in trials		18	100	22	100	
2.	Oestrus induced		12	66.6	16	72.7	
3.	Pregnancy		7	37.0	11	50.0	
4.	Pregnancy rate among served			58.33	-	68.78	
5.	Interval in days for oestrus Indu	uction	17.7	3±2.54	14.50	± 2.32	
6.	Interval in days for fertile oestr	us	40.14	±±3.71	28.50	+5.26	

The experimental work was carried out for the period of 6 months. On expression of oestrus, the animals were served either by natural services or by A.I. and regular rectal examinations were made for pregnancy after 6-8 weeks of service.

Results and Discussion

The details of results regarding oestrus induction and pregnancy have been furnished in Table 1. Out of total 40 animals under treatment trials. in 27 (67.50%) animals oestruses were induced and 18 (45.00%) animals conceived. The mean time interval for oestrus induction was 15.14 ± 2.68 days and for conception it was 31.9 ± 5.37 days.

Among control group of 10 animals, 3 (30.00%) animals came into oestrus and only one (19.00%) conceived. The mean time interval for exhibition of oestrus was 36.5 ± 7.83 days and conception took more than 90 days.

Chi Square test revealed significant difference (P < 0.05) between treatment and control group as regards oestrus and fertile oestrus induction.

The results of oestrus and fertile oestrus induction nearly agree with the results reported by Galhotra et al. (1970), Kodagali et al. (1973) and Desphande and Sane (1977) who tried the indigenous drugs.

The details of results on dose effect have been illustrated in table 2.

It can be seen from the above figures that apparently better results were observed with the dose of 3 cap/day bat statistically there was no significant difference. Thus both the doses (2 cap/day and 3 cap/day) are effective in induction of oestrus and pregnancies.

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Reproductive-Performance of $F_1 \& F_2$ Crossbred Buffaloes (Swamp \times Murrab) in Vietnam

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ABSTRACT

The present preliminary study revealed values of certain reproductive-parameters of Swamp \times River Crossbreds (F₁ & F₂) in VietNam and suggested for further detailed study in this direction.

*

Swamp buffaloes are often slow in coming into heat and that their reproductive-attributes, like gestation-period, post partum oestrus period and calving interval are long with little milk leading to low return. Hence, the introduction of exotic (river buffalo, Murrah) germ plasm in "VietNamesetrau" (Swamp) is accepted as a tool to bring about rapid improvement in milk-production and also to maintain its superior draughtability in rich paddy-area of SouthEast Asia. However, no report is seem to be available on the said reproductiveattributes of F1 and F2 Swamp Crosses in VietNam. The present paper is an an attempt to put on record in this direction.

Materials and Methods

The breeding data in two different genetic groups of Swamp Crosses namely Group-:1/2 Swamp X 1/2 Murrah (F_1) ; and Group-II:1/4 Swamp X 3/4 Murrah (F_2) were collected from Buffalo Breeding and Research Centre, Song Be (BenCat). The overall macro-environment of BenCat is "Monsoon-Tropical" climate due to its heavy rain (Yearly Rainfall 2028.96 mm) alongwith high ambient temp. (Yearly Average Temp. 28.58°C) as reported earlier by Verma et al. (1984). The ability to adopt the "Monsoon-Tropical Climate" of VietNam was evaluated on the reproductive-attributes like gestation-period, postpartum oestrus interval and calving-interval. The results were presented in Table 1.

Gestation-Period:- Table 1 indicated that the overall average gestation-length of crossbred-buffaloes irrespectively of grade was 308.0 ± 1.2 days in VietNam. In Philippines, Villegas (1959) has reported a little higher values (316.2 days) of the average gestation-period in river \times swamp type crosses (Philippine X Murrah Crosses, their various grades), which probably might be due to variation of the strain of swamp from location to location.

During the present study, it was observed that the average gestationlength of F_1 (308.1 \pm 1.6 days) was found to be slightly higher than F_2 Crosses (307.8. \pm 1.8 days), which might be due to higher blood level of Swamp in F_1 than F_2 Cross. Because the gestation of Swamp is more than the gestation period of Murrah (Fahimuddin, 1975).

Postpartum-oestrus interval: The mean postpartum oestrus interval of F_2 (60.5 ± 8.3 days) was comparatively less than

F1	Fa	Overall
308.1±1.6	307.8±1.8	308.0±1.2
(10) 78.8±10.0	(4) 60.5±8.3	(14) 70.7±7.0
(5)	(4)	(9) 520.5+85.1
574.0±93.6 (3)	300.0±0.0 (1)	(4)
	$\begin{array}{c} 308.1 \pm 1.6 \\ (10) \\ 78.8 \pm 10.0 \\ (5) \\ 574.0 \pm 93.6 \end{array}$	$\begin{array}{ccccc} 308.1 \pm 1.6 & 307.8 \pm 1.8 \\ (10) & (4) \\ 78.8 \pm 10.0 & 60.5 \pm 8.3 \\ (5) & (4) \\ 574.0 \pm 93.6 & 360.0 \pm 0.0 \end{array}$

TABLE 1. Reproductive-estimates of F1 and F2 Crossbred-buffalo at BenCat (VietNam)

Note:Figures in parentheses indicate number of observations.

 F_1 (78.8±10.0 days). The overall average postpartum oestrus of crossbred buffalo was 70.7±7.0 days in VietNam. In Philippine the postpartum oestrus in Swamp X River Type Crosses (Philippine X Murrah Crosses) were reported to be 44 (Ocampo, 1939) and 45.8 days (Villegas, 1959). The variations might be due to variations in the strain of Swamp, blood-level of swamp in Cross etc.

Calving Interval: The calving interval was found to be minimum in F_2 Crossbreds (360.0 \pm 0.00 days) against F_1 Crossbreds (574.0 \pm 93.6 days). However, overall calving interval was 520.5 \pm 85.1 days in River X Swamp Crossbreds in VietNam. In Philippine, Villegas (1959) has reported 429.2 days for calving interval in Swamp X River Type Crosses (Philippine Carbao X Murrah Crosses). The differences of finding probably might be due to difference of strain of Swamp, blood level of Swamp in its Crossbreds, location, number of observations etc.

Acknowledgement

Authors are thankful to the Govt. of Socialist Republic of VietNam and Govt. of India for taking their interest in dairyindustry through development of Murrah buffaloes in VietNam. Authors are thankful to Dr. N. V. Thuong, Director, Dr. N. D. Thac, Head, Buffalo Breeding Research, AHRI, Dr. N. V. Vuc, Director, and all staff members of SongBe-Centre, Mr. N. M. Dung, Mrs. L. N. Van, interpreters for their help.

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Gynaeco-clinical Investigation of Repeat Breeder Cows and Buffaloes

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ABSTRACT

The detailed gynaeco-clinical examination of 318 repeat breeder buffaloes and 81 repeat breeder cows revealed detectable problems in the genital organs to the extent of 40.67 per cent and 49.15 per cent, respectively as a possible actiology. The over all incidence of vaginitis (1.88% and 28.39%), kinked cervix (19.38% and 25.92%), uterine affections (5.01% and 4.93%), salpingitis (1.88%) and 1.23%), ovario-bursal adhesions (4.08% and 4.93%) and cystic ovaries (0.94% and 3.70%) was noted in buffaloes and cows respectively. It was revealed that with critical and careful clinical examination of repeat breeders, major disorders of genital tract were detectable. The disorders like vaginitis, mild kinked cervix and low grade infections could be managed, whereas severe fibrosed cervix, salpingitis, ovario-bursal adhesions and chronic cases of cystic ovaries were the extreme conditions found in chronic repeat breeders in which chances of recoveries were remote.

Detailed clinical examination of the reproductive system is possible in most

bovines and this provides the valuable information about reproductive status of the individual animal. With the emphasis to ensure optimum reproductive efficiency, especially in dairy herds, routine examinations are frequently performed for which an appreciation of the normal genital organs is important.

The cyclic, functional or pathological changes and congenital or hereditary defects can be detected relatively easily by gynaecological examination. For this a clinician must be familiar with the anatomy, physiology and pathology of reproductive organs as well as he should possess sufficient clinical experience. Many of the changes can be ascertained if daily gynaeco-clinical investigations are made. However, following a single examination accurate diagnosis is not always possible.

Materials and Methods

The Gynaeco-clinical investigation was carried out on 318 repeat breeder buffaloes (273 buffaloes and 45 heifers) which attended the college A.I. clinic during the period of three years (January 1980 to December, 1982). Eighty-one repeat breeder cows (58 cows and 23 heifers) were also investigated for genital abnormalities.

Animals which had completed 4 or more inseminations or natural services without pregnancy were investigated clinically. Breeding records pertaining to these animals were studied. After restraining and cleaning the vulva and perinium region, careful examination of genital organs was performed. Vaginal

1	Abnormalities detected									
No. of	Vagina		Cervix		Uterus		Fallopian tube	Ovary and Bursa		
animals	Vaginitis (G.V.)	Hypo- plastic	Enlarged and Hard		External cysts	Hard ute- rus with watery discharge		Ovario- Bursal adhesions	Ovarian cysts	Para- ovarian cysts
Cows	28.39	1.23	43.20	25.92	1.23	3.70	1.23	4.93	3.70	1.23
(81)	(23)	(1)	(35)	(21)	(1)	(3)	(1)	(4)	(3)	(1)
Buffaloes	1.88		5.34	19.18	1.25	3.14	1.88	4.08	0.94	2.93
(318)	(6)		(17)	(61)	(4)	(10)	(6)	(13)	(3)	(9)

TABLE 1. Results of Clinical Examination of Repeat Breeder Animals

inspection and rectal palpation for detecting and visual and palpable disorders of vagina, cervix, uterine horns, fallopian tubes and ovaries were made. When detailed inspection was necessary speculum examination was carried out for knowing the changes in vaginal mucus membrane, external cervical os and the nature of the discharge if any.

Results and Discussion

During the course of study 318 repeat breeder buffaloes and 81 repeat breeder cows were investigated. However, out of 318 buffaloes and 81 cows in 189 (59.43%) buffaloes and 42 (51.85%) cows respectively had no reproductive disorders which could be clinically detected. The abnormalities detected in repeating buffaloes were in the order of 1.88 per cent vaginal, 25.46 per cent cervical, 5.01 per cent uterine, 1.88 per cent tubal and 7.85 per cent ovario-bursal lesions. The details of reproductive disorders detected on clinical examination of repeating buffaloes and cows have been presented in Table - 1.

The frequency of occurrence of different detectable disorders in the reproductive tracts of repeat breeder buffaloes and cows varied considerably. The granular lesions in the vagina were more marked in cows, whereas the incidence of kinked cervix was high in cows and buffaloes. A very high incidence of hard and fibrous cervix noted in cows was in agreement with the findings of Bhosrekar (1973). This might be due to repeated breedings in these animals.

It can be observed from results (Table 1) that on clinical examination of tubular genitalia (uterus) in the present study in repeat breeders did not reveal any gross changes. Hard uterus with watery mucus discharge was observed in 10 (3.14%) buffaloes and 3 (3.7%) cows, respectively. In all these cases oestrous period was longer than normal. The uterus was slightly enlarged and hard. This may be due to either high estrogenic effect for longer duration or chronic lesions in the uterus. The ovario-bursal adhesions were detected in very low frequency, and this finding is in agreement with the observations made by Bhosrekar (1973), but earlier report of Dawson (1956) showed bursal adhesions in larger percentage of repeat breeder cows.

It is suggested that vaginitis, mild kinked cervix, and low grade infections were the conditions amneable to treatment, whereas severe ovario-bursal adhesions, cystic ovaries and fibrosed hard cervix were the extreme conditions of chronic repeat breeders in which the chances of restoration of fertility are remote.

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CASE REPORTS

IJAR 6: 1: 89-90 1985

Persistant Hymen In A Camel

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Imperforate or persistant hymen—a developmental defect of the tubular genital tract — has been frequently described in cows and marcs. The condition may be frequently associated either with segmental aplasia in cows or with an otherwis normal set of genitalia in cows and marcs. The present paper records a case of persistant hymen in a camel.

CASE REPORT:

A four and a half year old nulliparous, she camel was presented to our clinic with the history of frequent appearance of a distended, glistening mass between the vulval lips, particularly when the animal lied down. The condition had been existing for at least two months and the frequency of its appearance between the vulval lips as well as the degree of distension had progressively increased over this period.

The animal was cast on its abdomen and the fore and hind legs were secured separately. Immediately after casting, a cone shaped, glistening, pink coloured mass appeared between the vulval lips (Fig. 1). The distended mass was soft, firm and fluctuating and easily replaceable into the vagina., but reappeared again with each abdominal contraction. Further examination of the vaginal passage revealed that the vaginal canal was completely obliterated from immediately anterior to the urethral orifice by a tough membranous partition. Rectal examination revealed a marked distension of the vagina immediately caudal to the cervix and anterior to the membranous parition. No other palpable abnormalities of the cervix, uterus or the ovaries were detected.

TREATMENT:

The animal was tranquilized with 60 mg. of Triflupromazine hydrochleride (Siguil) Sarabhai Chemicals, Baroda, India administered intravenous. The tail was secured and the perenial region thoroughly cleansed with weak solution of potassium permanganate. A 2 cm. incision was made over the point of the greatest bulge and about two litres of mucus was seen to escape. A sample of the mucus, collected by aspiration through a sterile needle, was found to be negative for microorganisms on subsequent cultural examination. Complete draining of the genital passage was followed by enlargement of the incision at right angles in all four directions and the membrane was removed in pieces. The vaginal canal was irrigated with antiseptic solution and liberal quantities of Hostacycline Hoechst Pharmaceuticals, Bombay, India. powder (Tetracycline hydrochloride) was applied over the site.

Post operatively the animal received 4 million units of penicillin, intramuscular, for five days. The vaginal canal was examined every day with a vaginal speculum, in an attempt to dilate the



Fig. 1. Completely imperforate hymen in camel. The bulging is due to the accumulated fluids.

vulvo-vaginal ring. The animal was discharged after an uneventful recovery.

Discussion

Imperforate or persistant hymen has been rarely described in camels. The animal presented, to us, had a completely imperforate hymen with an otherwise normal set of genitalia. The vaginal distension immediately anterior to the hymenal partition indicated that the animal had probably cycled repeatedly with the consequent accumulation of the discharges posterior to the cervix. The age of puberty in female camels, about 31 years., and a relatively short breeding season, would perhaps mean that the animal should have been cycling for about four months, during which period the fluid accumulation occurred. The appearance of the bulging hymen between the vulval lips was due to the great distension of the vagina resulting into frequent straining. That the fluid inside the genitalia was sterile indicated that there was no chance of infection setting is due to the complete blockage by the hymenal membrane.

Persistant hymen has been recognised as a development defect of the vagina and may be of hereditary origion. Cases of complete imperforate hymen often require surgical intervention. Although some veterinarians prefer removing the imperforate hymen by circular incision along the hymenal attachment with the vaginal walls., enlarging the hymen by incision at right angles in all four directions, as used in the present case, has been advocated more frequently.

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A Cephalothoracopagus Monster In An Indian Water Buffalo (Bubalus Bubalis)

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Different types of foetal anomalies and monstrosities have been rarely recorded and cited in cattle (Jubb & Kennedy, 1970; Roberts, 1971). Reports on incidence of monstrosities in buffaloes are scarce (Mahalingam, 1968; Sahu, 1968; Bhaskar Singh, 1973 and 1976; Rai et al., 1975; Bugalia et al., 1980; Sreemannaryana et al., 1980; Nowshahri, et al., 1984). The present report records a case of cephalothoracopagus monster in an In lian water buffalo (Bubalus Bubalis). Case history, clinical examination and Obstetrical

Case history, clinical examination and Obstetrical operation

A pluriparous buffalo at partum after the local quacks had failed to deliver the calf even after amputation of three limbs, was presented to Veterinary College pervaginal Clinics, Ludhiana. On more limbs and examination two a head of quite a large size were discernible. It was diagnosed to be a case of foetal monster. Fetotomy was considered unsuitable and hence caesaritomy was performed and a conjoint twin monster was delivered.

General description

The conjoint twin monster delivered possessed a pair of cranie with complete spinal column's, Four fore limbs (Three of which had already been amputated), Four hind limbs, two vulva and two tails. Anus was discernible in one twin while there was atresia ani in the other. There was a fusion of the foetus ventrally. The fusion started from the head (Lower jaws) and extended to neck and thorax (Sternum to sternum). Emphysema of the fetus had already commenced.

Internal organs

The Lower Jaw of one twin was complete and was adhered to a thin sheath of tissue and skin representing the lower jaw of the other twin. The buccal cavity of one twin had tongue while on ly remnant of tongue was present in the other buccal cavity. This buccal cavity ended in a blunt pouch. The trachea and oesophagus opened into the common thoracic cavity of the twins formed due to sternum to sternum fusion. The thoracic cavity also enclosed one heart, two lungs, one oesophagus and a common intact diaphragm. The abdominal cavity comprised of single liver, spleen, pancreas, gall bladder and intestines. Single Female genito urinary tract was present. It was comprised of two kidneys, two ovaries, two oviducts and two uterine horns with cervix and vagina.

Discussion

The monster confirmed typical morphology of cephalothoracopagus as per



Fig. 1. Ventral opposition of crania due to fusion of lower jaws.

the classification of Potter (1961) cited by Roberts (1971). Roberts (1971) mentioned duplication of internal organs in such monstrosities as have also been observed by Dhingra *et al.* (1984). However, Arey (1966) stated that duplication of internal organs varies with intimacy



Fig. 2. Conjoint twin monster showing fusion of Lower Jaws, neck and thoracic region (Sternum to Sternum).

of fusion in twins.

Conjoined twins arise from single ovum and are monozygotic. The present case seems to be a teratologic defect of development arising from interruption of specific developmental stages that lead to incomplete twining.

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Uterine Prolapse In A Mare

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Uterine prolapse, a common postparturient complication in bovines and ovines, is considered to be rather infrequent in equines¹. Its occurrence in this species is usually as a sequel to the forced extraction of the foetus or retained placenta². The present report records a case of complete uterine prolapse following mid-term abortion in non-descript draft mare.

CASE REPORT:

A six and a half year old pluriparous, non-descript draft mare was presented with complete uterine prolapse. The animal had aborted the same morning, a six and a half month old foal, and although the placental membranes were reported to have been expelled immediately after abortion., eversion and prolapse of the uterus occurrred which was preceeded by mild to moderate straining. The prolapsed organ was fresh and moist (Fig. 1) and no attempts had been made to replace the same. Examination of the prolapsed mass revealed no signs of injury, lacerations or haemorrhage Inspite of the mild recurrent straining the animal appeared alert with normal body temperature and respiration.

TREATMENT:

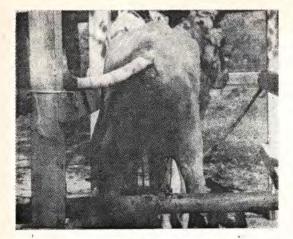
The mare was restrained in a chute and tranquilized with Siquil Sarabhai Chemicals, Baroda, India. (Triflopromazine Hydrochloride) 60 mg. I.M. The tail was bandaged and cpidural anaesthesia induced with 5 ml. of 2 percent Gesicain*. SG Pharmaceuticals, Baroda, India. (Xylocain Hydrochloride) . The perineal region was cleaned and the prolapsed mass was thoroughly washed with ice cold antiseptic solution. Antiseptic cream was liberally applied on the prolapsed mass, which was subsequently replaced by lifting and pushing with palm-pressure as in the bovines. Steclin Sarabhai Chemicals, Baroda, India. (Tetra-cycline Hydrochloride) bolus-2 g. was inserted into the uterine cavity and the vulvar lips were sutured with three zero silk using horizontal matress pattern, closing 2/3ds of the vulvar opening.

After replacement of the prolapsed uterus, the mare was given 60 I.U. of posterior pituitary extract, three million units of procain penicillin and 1500 uits of tetanus antitoxin, intramuscular.

Twenty four hours later, the mare was reported to be eating and drinking normally with no evidence of straining. The animal had also passed a small quantity of blood tinged uterine discharge. The antibiotic therapy and the local dressing of the vulvar sutures was continued till day 5 and the animal was discharged on day 6 after removal of the vulvar sutures and an uneventful recovery (Fig. 2).

Discussion

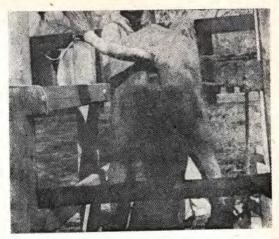
Uterine prolapse in equines is considered to be rather infrequent, with most



 Prolapsed uterus immediately after presentation at the clinic.

cf the isolated cases reported to occur during the immediate post-partum³,⁴ period. A long mesometrium and flaccidity of the pelvic viscera and the perineum associated with parturition probably predispose certain marcs to prolapse⁵. It has also been suggested that the free portion of the placental membranes, which remains hanging from the external genital orifice, exerts traction on the uterus and invariably leads to the uterine prolapse².

However, in the present case the mare aborted at approximately six and a half



The prolapsed organ has been replaced and the vulva sutured.

months of gestation and there was no evidence either of forced extraction of the foetus or of the foetal membranes. Therefore, none of the previously reported factors appear to be responsible in the present condition. However, in view of the general condition of the animal, the type of work for which it was used and the relative case with which the prolapsed mass was replaced, it seems highly probable that the abortion followed by mild to moderate straining, which coupled with the uterine inertia resulted into eversion and prolapse of the uterus.

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Fibro-Adenoma Of Cervix In A Kankrej Cow, A Case Report

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ABSTRACT

A case of fibro-adenoma of cervix in a Kankrej cow is reported.

The incidence of pedunculated tumours in the bovine genital system is rare while as the cervical tumours are still rarer. Wadsworth (1952), Cotchin (1956) and Moulton (1961) described fibromas, fibrosarcomas, leiomyomas and carcinomas affecting bovine cervix. Jubb and Kennedy (1970) have reported carcinomas of uterus and cervix. Chenna Reddy (1980) reported a voluminous fibroma of vagina weighing 7 kg in a non-descript cow. Cervical tumour has been reported as a cause of infertility in cow by Chatterjee (1971). Arthur (1975) commented that such tumours of cattle reported were of benign type. Sharma et al. (1977) described tumours of cervix in buffaloes, during routine examination of slaughter house material. Kohli and Bishnoi (1980) reported similar tumour in a pregnant Rathi cow while Sindhaye (1982) reported in a cross-bred lactating cow. The authors present here a case of long pedunculated cervical tumour in a Kankrej cow, which was removed successfully by surgical intervention.

CASE HISTORY AND CLINICAL EXAMINATION

An old pleuriparous dry Kankrej cow aged about 10 years (4th lactation) was presented to the Veterinary College, AI Clinic, Anand (case No. 7987) on 13th of September, 1983 with the complaint that there was a cervico-vaginal prolapse and the animal was straining intermittently since last two days, hence brought for the treatment. The animal had calved two years ago and was served six months previously. There was no estrous exhibited afterwards.

The general body condition of the animal was poor. Visual examination revealed a fleshy mass soiled with dung and having bruised wounds with necrotic shreds hanging out from the vulvar lips (fig, 1). Rectal examination revealed that the cow was non-pregnant, uterine horns were flacid, cervix was enlarged and somewhat hard, and both ovaries were smooth, soft and flabby indicating typical anoestrus condition. A thick long peduncle was felt on ventral aspect of os uteri externus. Cleaning of the hanging mass with saylon solution confirmed that it was a irregular hard tumourous growth, weighing about 2 kg with a long thick peduncular stalk originating from the inner aspect of cervix.

SURGICAL MANAGEMENT

The cow was restrained in the travis and the operation was performed in standing position. The tumourous mass was cleaned thoroughly with plenty of savion solution up to the cervix and was dried. Caudal epidural anaesthesia was induced with Novocain 7 ml injection

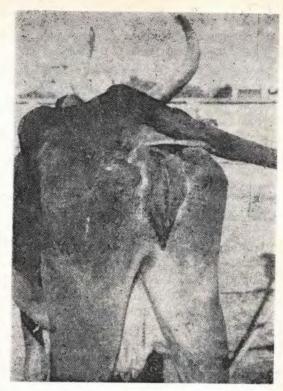


Fig. 1



Fig. 2

Fig. 3



Fig. 4. Microphotograph of tumour tissue showing extensive proliferation of fibrous connective tissue with cystic dilatations of the gland. 120×H.E. stain.

in the first intercoccygeal space, after giving Siquil 3 ml I/M. After setting of anaesthesia, cervix was everted through vaginal passage along with the tumour (fig, 2). The tumour was found to be supplied with plenty of blood vessels. The prominent blood vessels in the stalk of tumour were ligated with catgut (chromic No. 2/0) at the base of stalk; then stalk was clamped with two artery forceps from both ends of the stalk close to the base. A bold incision was made close to the artery forceps and the pedunculated tumour with stalk was removed. Then the surgical wound was sutured with catgut (chromic size No. 2) to check the bleeding and then artery forceps were removed. The wound was dressed with dusting of 2.5 gm Dicrysticin powder and the everted cervix was placed back in position (fig, 3). Dicrysticin 2.5 gm daily in 20 ml of distilled water was infused in the cervix for consecutive two days. On inquiry after 15 days of operation the animal had made an uneventful recovery.

HISTO-PATHOLOGICAL EXAMINA-TION

Grossly the tumourous tissue when cut showed few cystic spaces of varying sizes filled with mucous like substance. Parafin sections of tumour tissue were taken and were stained with routine H & E stain (Culling, 1963). On histopathological examination the tumour was identified as fibro-ademoma. Scattered focal areas of glandular acini lined by columnar epithelium were seen amidst extensively proliferated fibrous connective tissue (fig, 4). At places the acini were cystic and filled with mucous. No malignant changes could be noticed.

Acknowledgement

The authors are grateful to Dr. M. R. Patel, Principal, College of Veterinary Science and Animal Husbandry, GAU, Anand for the facilities provided.

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Campylorrachis Scoliosa—A Fetal Monster Causing Dystocia In Buffalo—A Case Report

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ABSTRACT

Campylorrachis scoliosa—a fetal monster seen very rarely in cattle and swine is reported for the first time in a buffalo. The 12 kg male dead monster, removed by cesarotomy in a buffalo suffering from dystocia, had laterally bent vertebral column, deformed fore limbs and complete ankylosis of all joints. Rest of the body was apparently complete except that testis were retained in abdomen, penis was missing and prepuce closed.

*

Fetal monstrosities are quite common in domestic animals. A large number of fetal anomalies and monstrosities of cattle are enlisted by Roberts (1971) and Schistosomus reflexus, double monsters and Persosomus elumbis comprise the commoner gross abnormalities of obstetrical interest (Arthur et al, 1982). Various types of monstosities of obstetrical interest that are on record in buffaloes include Schistosomus reflexus (Rai et al, 1975; Rao and Sreen, 1984), double monsters (Dingra et al, 1984), and Diprospus (Sreemannaryana et al, 1980). However, the literature available lacks in reports on campylorrchis scoliosa in buffaloes-a monster that very rarely occurs in cattle and swine. The present report puts on record the occurrence of this fetal monster causing dystocia in a buffalo.

Materials and Methods

A five years old primiparcus buffalo which had completed gestation period was presented to Veterinary clinics of Punjab Agricultural University for the treatment of dystocia. The water bags were ruptured and partial fetotomy had been done on limbs presented in birth canal by the field veterinarian a day before refering it to our clinics. According to history, this was the fourth such monster from four different buffaloes served by the same bull.

Per-vaginal examination of the case revealed a monstrous fetus with ankylosed limbs. The handling that had already been done caused lacerations and oedema of vulva and vagina. The stenosis of birth canal thus occurred rendered the case unfit for fetotomy. Cesarean was done and small dead monster was removed. The monster was weighed and preserved in 10% formaline for gross dissection. The dam was treated as routine and discharged on recovery.

Results and Discussion

Clinical Examination and treatment of dam: Simultaneous presentation of head and hind quarters of a totally anakylosed fetus in an cedematous birth caral made the handling of the case more tedious. Eventually, cesarean section was done

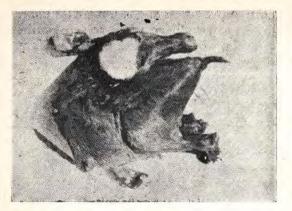
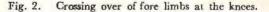


Fig. 1. Laterally bent vertebral column in Campy lorrachis scoliosa — a buffalo fetal monster.

and the monster removed. Ankylosed monsters often cause dystocia where mutuation, force traction and fetotomy seldom succeed. Bugalia et al (1982) and Nanda et al (1983) also preferred cesarotomy to remove Schistosomus reflexus—a monster of similar contour, while Roberts (loc.-cit-) however, could deliver them by force traction and he fetotomy.

Gross Examination of Monster: The monster was a male fetus weighing only 12 kgs. as against 29 kgs. average weight of a buffalo calf (Gururg and Johar, 1983). The vertebral column was bent laterally over the thorax region making a 'U' shaped curve in a way that head was resting against lumber region and nose was approximating the pelvis (Fig. 1). The pelvis was tapering posteriorly forming a cone and was notably small. The hind limbs were already amputated just below the hocks. The hocks and the rest of the limbs and vertebral column were ankylosed. The fore limbs were crossing over each other over the knees (Fig. 2). The rudimentary scrotum was present but without testis. The prepuce was closed. The rest of the body was apparently complete.





Detailed dissection were performed on the monster. It was found that all internal organs of skull, buccal cavity, thorax, abdomen and pelvis were present and size developed normally appropriate to the fetal size. However, both testicles along with respective epididymii were retained in abdomen. The penis was missing and prepuce was closed. Unlike Schistosomus reflexus, where liver is usually oversized and/or cystic, the liver was grossly normal with gall bladder.

Owing to the bend in the vertebral column and ankylosis of limb, this monster ressembled with Schistosomus reflexus where the vertebral column is sharply bent dorsally and the abdomen is open at linea alba exposing the visceral organs. Unlike Schistosomus reflexus, this monster had a laterally bent vertebral column, fore limbs were deformed and visceras were enclosed in a complete abdomen and was confirming the characteristics of Campylorrachis scoliosa as described by Roberts (loc.-cit.).

Cryptorchidism as seen in this case and ankylosis and death of fetus in last month of pregnancy is transmitted genetically (Thompson et al, 1957). Nothing is reported about Campylorrachis scoliosa being genetic or because of some environmental factors However, the fact that four similar monsters were born to two sisters and two more buffaloes in the same village when served by the same bull over a period of two years is suggestive of its being genetic.

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Campylorrachis Scoliosa Monster And Spontaneous Vaginal Rupture In A Goat

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ABSTRACT

A case of spontaneous vaginal rupture at parturition associated with Campylorrachis Scoliosa monstrosity with abnormally long and contracted limbs in a goat has, been reported.

* *

Campylorrachis scoliosa is a foetal monster, rarely seen in cattle and swine, characterised by a lateral curvature of the spine and the limbs are usually deformed and ankylosed (Roberts, 1971). The spontaneous vaginal rupture at parturition due to such monster is uncommon. However, uterine rupture is seen in cases with malpositions which are difficult to correct. Verma and khar (1972) reported ankylosis of fetal legs as a cause of dystocia and uterine rupture in a cow. Williams (1951) and Roberts (1971) described rupture of dorsal wall of vagina and uterus as a sequelae to foot nape posture, ventro-transverse presentation and dog sitting posture of the foetus. Singh and Purbey (1984) observed 3 cases of curved vertabral spine associated with prolonged deformed limbs in Muzzaffernagari sheep. As there were no reports on Campylorachis scoliosa monstrosity in the goat in the literature reveiwed, the present one is reported.

Case History

A she-goat, aged about 5½ years having delivered normal kids twice previously,

100-1

was presented to the Veterinary College, Hospital, Anand on 14th of February, 1985 (Case No. 12371) in the evening at 6.00 P.M. with the complaint that the animal was in full term pregnancy showing labour pain, severe straining and bleating very frequently since last evening without any progress in the process of parturition. The water bag had appeared and ruptured earlier. No efforts or manipulations were made to deliver the foetus.

CLINICAL EXAMINATION- DIAG-NOSIS & TREATMENT

The goat was dull, depressed, anxious and was straining and bleating frequently. Per-vaginal examination, after applying sufficent quantity of Fairginol obstetrical cream on the gloved hand, revealed a transverse tear of about 4-5 inches long on the dorsal wall of vagina just behind the os through which all the four limbs had gone below the rectum, the foetal head could not be located. Cervix was fully dilated and scanty fluid was present in the birth canal mixed with blood. Due to the presence of four limbs simultaneously into the birth canal a double monster or twin pregnancy was suspected.

Further examination by repulsion of the foetus showed that there was a single foetus which was presented in anterior longitudinal presentation, dorso-sacral position and both the hind limbs were extended simultaneously beneath the



foetal body and into the birth canal along with extended forelimbs. The head was severely deviated downward between the forelimbs reaching the sternum (Fig. 1). The spinal column of the foetus was found to have a "S" shaped dorso-lateral cnrvature. All the joints and spinal column were very rigid. The limbs were found to be excessively long and rigid. The correction of the deviated head and limbs was tried through birth canal. The head deviation could be corrected with great difficulty but the limbs could not be corrected because of rigidity. The deformed, dead foetus was then delivered by gentle traction on the forelimbs with a hand pressing the hind limbs inside to prevent their simultaneous progression as the repulsion and correction was not possible. The part of the placenta came out immediately behind the foetus but rest of the placenta was not completely separated and a little amount of bleeding was observed. The tear of dorsal wall of vagina was sutured by everting the cervix and vagina through vulvar canal. Injection Caldee-12 10 ml S/C, Oxytocin 25 i.u. I/M and Oxysteclin 5 ml I/M were given to hasten the involution of uterus and to combact infection. Two Furea boluses were inserted into the

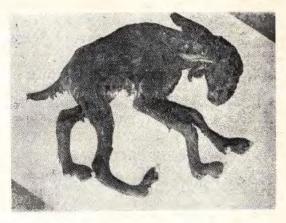


Fig. 2

uterus. As there was no bleeding and the animal stopped straining, the animal was not hospitalized but the owner was advised to bring the animal for further treatment.

GROSS EXAMINATION OF MONS-TER KID

The monster was a male kid with wattles and confirmed the description of Campylorrachis scoliosa described by Roberts (1971). The kinking and ankylosis of vertabral column with dorsolateral deviation involving posterior half of thoracic and first two lumber vertabrae (Fig. 2) was distinctly visible. There was dorsal flexon of fetlock joints due to contracted digital extensor tendons. The radiological findings were confirmed by detailed dissection. The right testis was located in the inguinal canal while left one was fully descended into the scrotum. The head, tail and the other viscera were almost normal. However, due to the vertebral deformity the tophography of various thoracic and abdominal viscera, was slightly altered.

Acknowledgement

The Authors are greatly indebted to Dr. M. R. Patel, Principal, Gujarat Veterinary College, Anand for the facilities provided.

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Duplication of Right Hind Limb In a Cross Bred Jersey Calf

M.R.K. IYER

Superintendent, District Livestock Farm, Kodappanakunnu, Trivandrum

Developmental anomalies have been reported by Jubb and Kennedy 1963 and Christopher 1971. Since it is a facinating subject a case of duplication of right hind limb in a cross bred Jersey calf is reported.

A Jersey cross bred cow aged 6 years had normal calvings thrice before and with a gestation period of 290 days was presented with the history that calving did not take place inspite of prolonged straining. The animal was off feed since morning and the temperature was 38°c. Per vaginal examination revealed unruptured bag and breech presentation. On rupturing the water bag the presence of additional hind limb could be felt. Both the well developed hind limbs were corrected and by judicious traction the calf was removed.

The calf was female, fully grown but weak and weighed 28 Kgs. Except for an accessory right hind limb and horny . growth measuring 4×1.5 c.m. on the internal aspect of left metatarsal no other defect were visible. The accessory limb was placed posterior to the right hind and the hoof tip was facing forwards. Detailed examination revealed the presence of extra acetabulam, all bones and joints. All the bones compared with the healthy right hind were proportionately shorter and the hoof was placed 8 c.m. above the ground level. All joints except the hock and stifle were well formed. The hock joint was less angular and tubercalcis very short. The two



stifle joints were placed in a line and the distance between the two were 9 c.m. (Photograph of the calf is enclosed).

According to the owner, the calf succumbed suddenly on the fifth day due to convulsions and planned amputation of additional limb at the stifle region could not be carried out.

According to Jubb and Kennedy (1963) the skeleton develops by a complex process from the mesenchyma and hence there is ample opportunity for error. A calf with a parasitic limb was illustrated by Roberts (1956) and an extraneous limb like appendage with a horny clubbed extremity was reported by Misra et al (1984) in a murrha buffalo calf. This case differs significantly from others since the accessory limb has all the bones and joints.

A case of duplication of right hind limb in a cross bred Jersey Calf is reported.

Acknowledgement

The author is thankful to Dr. N. K. Unni, retired Director of Animal Husbandry and Dr. N. Madhavan Nair, Deputy Director for the encouragement.

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Unilateral Orchitis In Bucks

JOSEPH MATHEW, E. MADHAVAN and C.P.N. IYER

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ABSTRACT

Acute unilateral orchitis noticed in two Alpine bucks maintained in the AICRP on Goats, Kerala is described. In one case unilateral removal of the affected organ was done and in the other only broad spectrum antibiotics and allied treatments were tried. In both the cases the animals died within a short span due to affection of the vital organs like lungs and kidneys.

*

Two cases of acute unilateral orchitis noticed in Alpine Bucks belonging to the All India Co-ordinated research project on Goats, Trichur, Kerala are presented in this paper. The therapeutic measures adopted and the gross and histopathological lesions of the condition are also documented.

Case 1.

The Buck No. A 58 showed hot and painful swelling on the left side of scrotum (Fig. 1), and was diagnosed as left lateral orchitis. A course of broad spectrum Antibiotics was started for treating the acute inflammatory reaction. On the second day the inflammed left testicle was removed by open method of castration. Antibiotic treatment was continued but the condition of the animal became poor and the animal died three days later.

The affected left testicle together with its tunics removed during the surgical

operation was about 2 to 3 times larger than the right testis. The organ when cut into two longitudinal halves revealed thickening of the parietal layer of tunica vaginalis and a thick gelatinous material between the parietal and visceral layers (Fig. 2). The thickness of this gelatinous material increased to about 3 cm. ventrally near the cauda but it reduced gradually and was almost completely absent on the posterior aspect of the testis. The cut surface of the testicular parenchyma revealed slightly reddish and congested appearance with only poor bulging of the cut surface. While incising the caput epididymis, a cavity, 1.5 cm. in diameter, containing yellowish creamy material was noticed on its lateral side. From the cut surface of the cauda, blood-tinged, thick whitish material oozed out. There was severe congestion and reddening of the cauda.

Histopathological examination of the testis revealed severe degenerative changes characterized by almost complete desquamation of the epithelial lining of the seminiferous tubules. The desquamated cells had highly condensed, hyperchromatic nucleus and vacuolated cytoplasm. Sections of the caput epididymis near the cavity revealed typical granulomatous reaction characterized by the presence of a large number of macrophages and few giant cells, as described by cohrs (1967). Sections of the cauda revealed



Fig. 1 Left lateral orchitus

severe congestion and infiltration of neutrophils and few mono-nuclear cells.

The failure to save the life and reproductive capacity of the animal by unilateral removal of the affected testis (Lagerlof, 1934) might be due to primary infection already present in other vital organs as revealed by the presence of nephritis and multiple abscess in the spleen and lungs on post mortem examination.

Case 2.

The Buck No. A. 62 developed sudden hot swelling of the right testis. The animal was treated with broad spectrum antibiotics parenterally and application of Mag. Sulp. glycerine externally on the scrotum. Although the swelling was reduced considerably the semen picture revealed continuous deterioration in quality, suggestive of severe degenerative changes in the testis. The animal died 14 months after the onset of the condition.

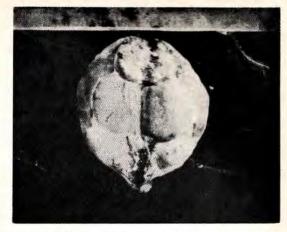


Fig. 2 Thick gelatirous nateral leturow the layers of tunica Vagirales.

The cause of death was due to acute congestion and oedema of the lungs together with multiple abscesses in the spleen.

The testicles were very hard to feel. both testes revealed On sectioning, numerous whitish gritty areas in the parenchyma. Histopathologicalexamination of the testicles revealed severe degeneration of the seminiferous tubules. The tubules were lined only by a single layer of sertoli cells. Prominent calcification of the desquamated contents was also noticed at many areas. The epididymis revealed chronic inflammatory changes characterized by prominent inter tubular fibrosis with infiltration especially of mononuclear cells around the epididymal tubules. The failure of saving the animals reproductive capacity with antibiotics and allied therapy might be due to the reversion of acute inflammation into a chronic form and extending into both the gonads.

Both the animals in the present study were checked and were found free of Brucella infection. It could be presumed from the course of events of the disease and the post mortem findings that the affection of testicles in both the case could he due to a sequale of generalised infection.

Acknowledgement

Grateful acknowledgement is made to Dr. M. Krishnan Nair, Dean, for the facilities provided.

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A Case Of Bifid Scrotum In A Young Rathi Bull

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Though on the whole quite a voluminous literature is available on the malformation of the genital organs in a bull but not much has been reported specifically on the scrotum. A case of bifid scrotum in a buffaloe bull had been reported by Kohli and Rajwanshi (1967).

Case Report

A young Rathi cow bull, aged about $1\frac{1}{2}$ years was brought to the Gynaecology outdoor clinic of the College along with two cows which the owner wanted to get treated for purperial infections. The young bull was dark brown in colour and its scrotum was bifid (Fig. 1). Both the testicles were well developed and were freely movable in the scrotum.

In this case there was no rotation of the scrotum on its longitudinal axis while the case of bifid scrotum in a buffalo bull reported by Kohli and Rajwanshi (1967) had a rotation of 90° to its left.



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FARM NEWS

Timing Of Parturition In Surati Buffalo

LALITA V DESHPANDE and K JANAKIRAMAN

Reproductive Biology Research Unit Faculty of Veterinary Science and Animal Husbandry Gujarat Agricultural University Anand

At the time of parturition the farm animals deliver a vulnerable young one. Under natural conditions the timing of parturition are so arranged, as to limit the vulnerability of the newborn, hence more births occur during night than during the day (Fraser, 1968). In cattle higher percentage of calvings have been reported during night hours by McDonald (1971). But Edwards (1979) reported equal distribution of calvings throughout the 24 hours.

In buffalo the frequency of calving during various parts of the day has received little attention. The knowledge of the peak hours of calving would help the farm personnel, in giving proper care to the newborn (e.g. timely feeding of colostrum) and assistance required if any, to the dam at the time of calving.

Bhuller and Tiwana (1984) have reported that buffalo calves born during day time are provided with better care by animal house staff, which is reflected by the lower mortality rate in the calves born during day time as compared to the higher mortality rate in the calves born during evening and night hours.

Further weaning of the newborn calf from the dam is being regularly preticed at most of the organized buffalo farms. If the calf suckles the dam, weaning becomes difficult later on. Knowledge of the timing of parturition especially the peak hours will help personal attention at the required time to wean the calf.

In the present study the data on 94 normal calving with the birth of viable calf in Surati buffalo are reported. The animals included, ranged from primipara to those with 6 calvings. They were maintained under standard managemental conditions. The buffaloes approaching calving were isolated to facilitate better observations. The time of calving was recorded for each birth. All the parturitions were single births and occurred with anterior, normal presentations, except in one case where the presentation was posterior.

When the frequency distribution for calvings during 24 hours of the day was worked out, it pointed out that though the calving frequency is located almost at all hours of the day, the clustering occurred between 2 to 3 (10 calvings) and 20 to 21 (7 calvings) hours.

The data was further grouped and for each 6 hour period, the percentage of calvings was worked out as detailed in Table-1.

In this particular study out of 94 observations maximum (29) number of calvings occurred between 6 to 12 hours, followed by 27 calvings during 0 to 6 hours. Almost equal number of delive-

TABLE 1: Frequency of calving

Sr. No.	Hours of the day		Percentage of calving
1.	0-6	27	29.0
2.	6 - 12	29	31.0
3.	12 - 18	18	19.0
4.	18 - 24	20	21.0
	Total	94	100.0

ries occurred between 18 to 24 and 12 to 18 hours.

The day and night frequencies of calving were equal (50% each). However it appears that the period between 0 to 12 hours is preferred by buffaloes, when 60% of the total calvings occurred.

Interestingly, out of the total 94 calvings, 25 parturitions (27%) occurred on weekends (Saturdays and Sundays), which indicates the need to attend to the animals on odd days also.

Finally, the data makes it evident that in buffaloes the calvings are spread throughout the 24 hours of the day with maximum calvings between 2 to 3 a.m. probably due to least disturbances. And although the frequency of calving is equal during day and night, the period from midnight to midnoon is more favoured making it evident that arrangements must be made, especially during odd hours to attend the prospective calvers, to help proper weaning, reduce complications and calf mortality.

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ISSAR NEWS

1. ANNOUNCEMENT OF THE ASIAN CONGRESS NOTIFICATION

ISSAR is pleased to announce that the FIRST ASIAN CONGRESS ON ANIMAL REPRODUCTION will be organised in association with I.C.A.R. and Konkan Agricultural University, Dapoli at Bombay on 11th., 12th., & 13th. December, 1985. Last date for receiving the abstracts of Scientific articles is 31st. August, 1985 and for full Text of the paper 30th. Sept. 1985. Delegation fee is Rs. 200/- upto 31-8-85 and Rs. 225/upto 30-9-85 and Rs. 250/- after 1-10-85 (including spot registration).

Scientists from Asian Countries are expected to participate in the congress.

For further details, please contact the undersigned. Venue of the congress:-New Complex of the Bombay Veterinary College, at Aarey Colony, Bombay-400 065.

(Dr. D. P. Velhankar) Hon. Sec. I.S.S.A.R. & Organising Secretary

2. It is experienced that the journal issues are returned back undelivered. The members are earnestly requested to communicate change in their address to avoid necessary postal expenditure and delay.

Dr. D. P. Velhankar) Hon. Secretary ISSAR

3. Prof. C. R. Sane and Dr. B. R. Deshpande were invited for the All India Symposium on Cross Breeding and Embryo Transfer Technology organised by the Rajasthan Cooperative Dairy Federation held at Jaipur on 16th. & 17th. NOV-1984. Prof. Sane delivered the key note address and Dr. B. R. Deshpande chaired the sessions on Panel discussion on A.I. and Embryo Transfer Technology.

It is to the credit of ISSAR that the proposal submitted to the Govt. of Rajasthan for opening one more Veterinary College, has been accepted by the Rajasthan Govt. A communication to this effect has been received from the Director of A.H. Rajasthan State.

4. ISSAR NEWS:

2.4 2

Congratulations to Dr. A. Ramamohana Rao:

ISSAR feels happy that Dr. A. Ramamohana Rao has assumed the charge of the Dean, Post-graduate Studies, Andhra Pradesh Agricultural University, Rajendra Nagar, Hyderabad. On behalf of the members of ISSAR we convey him Hearty Congratulations and Best Wishes.

> Dr. D. P. Velhankar Hon. Sec. ISSAR

5. LIST OF PARTICIPANTS OF THE 16TH SWEDISH INTERNATIONAL POSTGRADUATE COURSE ON ANIMAL REPRODUCTION- 1985

Otavio Mitio Ohashi Li Chih Fu Eulogio Padron Moreira Merga Bekana Gonfa Tapan Kumar Barthakur Ashok Wasudeorao Deshmukh Gurdial Singh Randhawa Logeswaran s/o Vannyasingam Pedro Henrique Antonio Halar Angel Alberto Tornamira Romero Hussein Mohamed Nur Duwearachchige Ajantha Melani Hewakopara Sunanda Sirimathie Weerathunga Wakista Elamin Dafalla Gasm Elseed Nussara Vadhanakul Brasil China Cuba Ethiopia India India India Malaysia Mozambique Peru Somalia Sri Lanka Sri Lanka Sri Lanka Sudan Thailand

Obituary

Biodata of Dr. Sundaresan

It is with great measure of shock to learn the untimely passing away on 1st March 1985 of Late Dr. Devadasan Sundaresan who was Director of NDRI, Karnal up to 30th June 1981.

Dr. Sundaresan was born on 10th March 1925. He graduated from the Allahabad Agricultural Institute in 1946. He obtained his master's Degree from Kansas and Ph.D. in 1959 from Iowa State University.

Dr. Sundaresan served as Research officer, Animal Breeding at NDRI, Karnal till 1964. Afterwards he joined the PAU and HAU where he served as Dean and Director of Research between 1964-70. From Nov. 1970 he served as Director, NDRI, Karnal till 30th June-1981. Dr. Sundaresan had number of FAO assignments.

After his retirement from NDRI he had joined Allahabad Agricultural Institute as Director in 1982.

In his passing away the NDRI has lost one of its most distinguished builders. His dedication will long be remembered in the cause of Agricultural Research and and education in India and abroad.

The members of ISSAR share the sorrow and pay their Homage.

May his Soul rest in peace.

DECLARATION

Statement about ownership and other particulars about THE INDIAN JOURNAL OF ANIMAL REPRODUCTION as required under Rules No 8 of the Registration of News papers (Central) Rules 1956.

FORM NO. IV (Rule No. 8)

1. Place of Publication

Editorial Office: Dept. of Gynaecology & Obstetrics Gujarat Veterinary College, Anand

2. Periodicity of Publication

3. Printer's Name Nationality Address

- 4. Publishers Name Nationality Address
- 5. Editor's Name Nationality Address

Names and addresses of individual who own the news-paper and partners, share-holdersholding more than 1 per cent of the total capital Bi-annual (JUNE & DECEMBER)

Anand Press, Gamdi Anand-388 001

Dr. D. P. Velhankar Indian Bombay Veterinary College, Bombay 400 012

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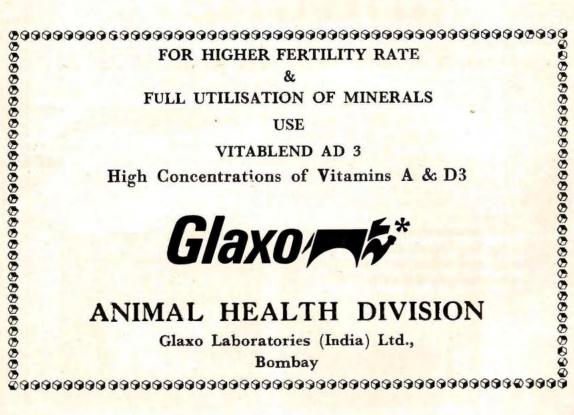
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I Prof Dr SB Kodagali, Editor of THE INDIAN JOURNAL OF ANIMAL REPRODUCTION hereby declare that the particulars given above are true to the best of my knowledge and belief.

> PROF DR SB KODAGALI Editor

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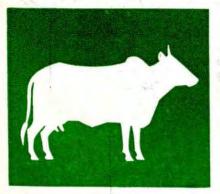


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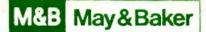
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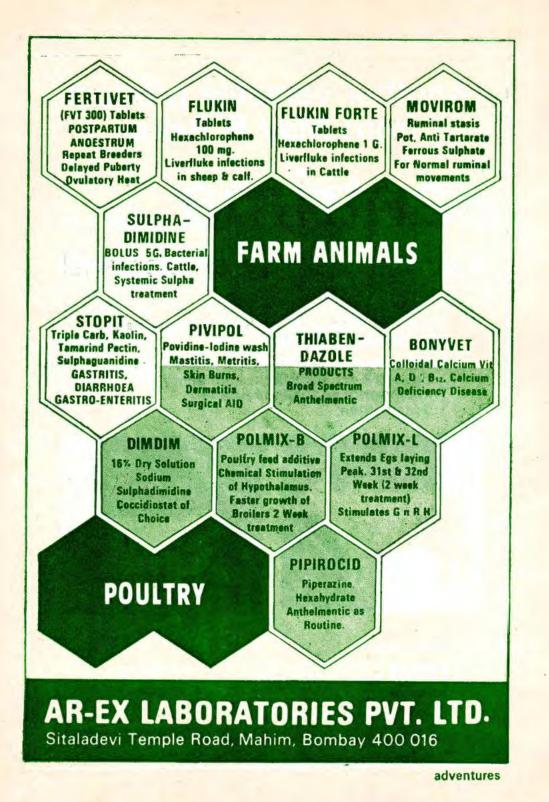


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This Video Cassette was even shown to about 1000 Gynaecologists coming from all over the World at the "Fourth World Congress on Human reproduction," which was held at Bombay from November 26th to December 2nd 1983.

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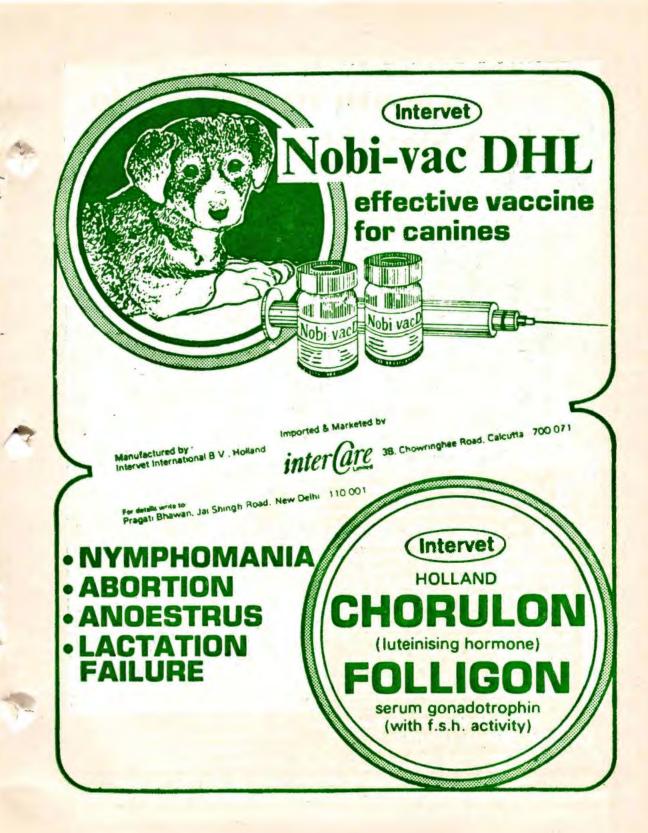
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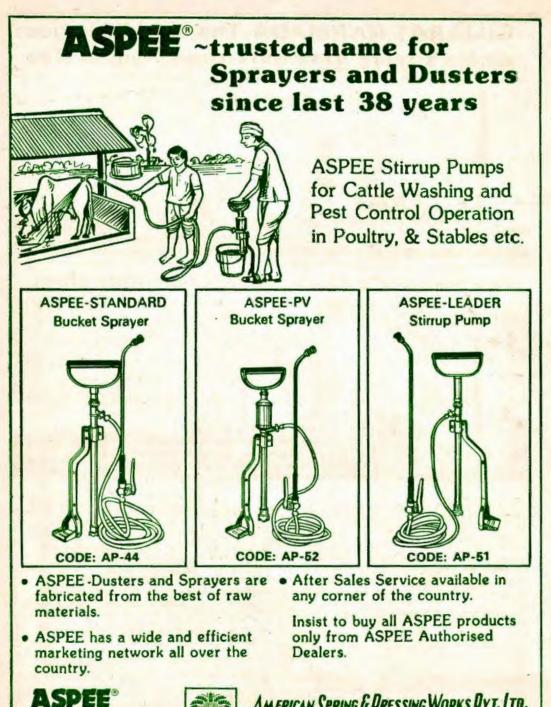
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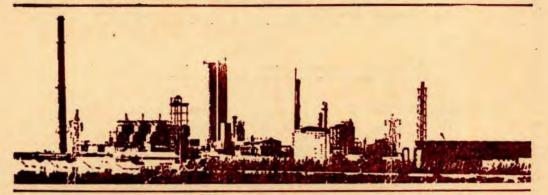


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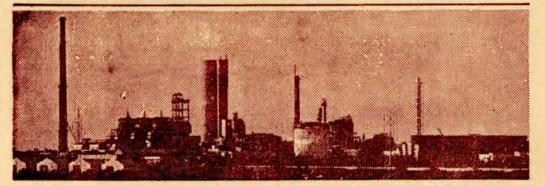
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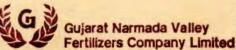


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