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## INDIAN JOURNAL OF VETERINARY AND ANIMAL SCIENCES RESEARCH (Formerly Tamil Nadu Journal of Veterinary and Animal Sciences)

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# Type of whelping and its infleunce on Apgar score

## A. Subramani<sup>1</sup>, N. Arunmozhi<sup>\*2</sup>, P. Sridevi<sup>3</sup>, M. Shafiuzama<sup>4</sup>, Cecilia Joseph<sup>5</sup> and K. Krishnakumar<sup>6</sup>

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#### ABSTRACT

The modified Apgar score for puppies was evolved based on the reference range of heart rate, respiration, irritability reflex, motility and membrane mucous colour according to the physiology of canine neonate. A total of 100 puppies born to the bitches that underwent Spontaneous Whelping (SW), Assisted Whelping (AW) and Caesarean section (CS) were selected and were divided into III groups viz. SW (n=30), AW (n=35) and CS (n=35). A modified Apgar score model was used to assess neonatal viability of the puppies. Among the total 100 pups evaluated 36, 13 and 51 pups were having Apgar scores of 0 to 4, 5 to 9 and 10 to 14, respectively. Out of 36 pups having Apgar scores of 0 to 4 at birth, 5.56, 19.44 and 75.00 per cent of pups were born through SW, AW and CS, respectively. Among 13 pups born with Apgar scores of 5 to 9 at birth, 0, 38.46 and 61.54 per cent of pups were born through SW, AW and CS, respectively. Among the viable pups at birth, the pups born through CS were having low and medium Apgar scores of 0 to 4 and 5 to 9, respectively. All the pups delivered through SW had highest Apgar scores at birth while the pups delivered through AW had shown Apgar scores between 5 to 9 and 10 to 14.

Keywords: Apgarscoringsystem, Canine, Dystocia, Puppies, Mortality

#### INTRODUCTION

The incidence of neonatal mortality following normal delivery is reported to be as high at 9 to 26 per cent (Davidson, 2014) and 30 to 40 % (Mosier, 1986) following complicated whelping in dogs. Till date, there are very few reports on the use of Apgar scoring system to evaluate newborn puppies, mainly due to the polytocous condition of the dog. The modified Apgar score for puppies was

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evolved based on the reference range of heart rate, respiration, irritability reflex, motility ad membrane mucous colour according to the physiology of the canine neonate (Veronesiet al., 2009).

## MATERIALS AND METHODS

The present study was conducted in the puppies born to bitches of different breeds brought to the Small Animal Gynaecology and Obstetrics outpatient ward (SAP-OP-OG) of Madras Veterinary College Teaching Hospital with the history of progressive whelping and / or dystocia. A total of 100 puppies born to the bitches that underwent Spontaneous Whelping (SW), Assisted Whelping (AW) and Caesarean section (CS) were selected and were divided into III groups as follows.

Group I (n=30) comprises of Spontaneous Whelping in which puppies which were born to five bitches through spontaneous whelping without any assistance. No medical/manual or surgical assistance were given to the bitches for delivering the puppies. Group II (n=35) comprises of assisted whelpingin which puppieswere born to 10 bitches that delivered through either manual or medical assistance. Manual assistance was attempted to the puppies which were partly expelled from the vagina and/or the puppies whose parts were within reach on vaginal examination but not progressed further. Medical assistance was supported with induction protocol with slow intravenous injection of oxytocin @ 1.1 to 2.2 IU/kg IM or SC with a dose range between 5 and 20 IU every 30 min and concurrent administration of 10% calcium gluconate @ 0.5 to 1.5 ml/kg to augment oxytocin's effect on myometrial contraction and intravenous fluids to correct hydration, electrolyte, and blood glucose abnormalities.Puppies born to 9 bitches that underwent caesarean section were included in group III. Under general anaesthesia using propofol @ 3 mg/kg and diazepam @ 0.5mg/kg for induction and 2 per cent isoflurane for maintenance, caesarean section was performed through midventral approach by adopting standard surgical procedures The Apgar scores were evaluated at birth, 30 min, 2 and 24h after birth.

A modified Apgar score model developed by Groppetti et al. (2010) (Table 1) was used to objectively assess neonatal viability of the puppies. The following parameters were evaluated: mucous membrane colour, heart and respiratory rate, reflex irritability, mobility, suckling, and vocalization, as described in Table 1. Each parameter was assigned a value 0, 1, or 2. Apgar score for each pup is the summation of the value of each parameter.

The number of animals in each category is expressed as percentage of total number of animals in that category.

## **RESULTS AND DISCUSSION**

Percentage of pups with different Apgar scores at birth in relation to type of delivery is presented in Table 2. Among the total 100 pups evaluated, 36, 13 and 51 pups were having Apgar scores of 0 to 4, 5 to 9 and 10 to 14, respectively. Out of 36 pups having Apgar scores of 0 to 4 at birth, 5.56 (2/36), 19.44 (7/36) and 75.00 (27/36) per cent of pups were born through SW, AW and CS, respectively. Among 13 pups born with Apgar scores of 5 to 9 at birth, 0, 38.46 (5/13) and 61.54 (8/13) per cent of pups were born through SW, AW and CS, respectively. Out of 51 pups with Apgar scores of 10 to 14 at birth, 54.90 (28/51), 45.10 (23/51) and 0 per cent of pups were born through SW, AW and CS, respectively.

The survival rate of the puppies at birth, 30 min, 2 h and 24 h in relation to type of delivery and Apgar scores is presented in Table 3. Out of 75 viable pups at birth, 14.67 (11/75), 17.33 (13/75) and 68.00 (51/75) per cent pups were showing an Apgar scores of 0 to 4, 5 to 9 and 10 to 14, respectively.

Among 67 viable pups at 30 min, 5.97 (4/67), 11.94 (8/67) and 82.09 (55/67) per cent pups were showing Apgar scores of 0 to 4, 5 to 9 and 10 to 14, respectively. Out of 11 pups born through CS that had low Apgar scores of 0 to 4 at birth, 6 pups died within 30 min. Of the remaining 5 pups, one pup had an improved Apgar scores of 5 to 9 at 30 min while the remaining 4 pups continued to show low Apgar scores. Out of 13 live pups that had Apgar scores of 5 to 9 at birth, 2 died within 30 min and 4 had improved Apgar score of 10-14. The remaining 7 pups continued to show low Apgar scores of 5 to 9 at 30 min. (Totally, 8 pups showing 5 to 9 Apgar score at 30 minutes is because one pup had improved Apgar score from 0-4 at birth to 5-9 at 30 min)

Out of 61 viable pups at 2 h, 1.64 (1/61), 4.92 (3/61) and 93.44 (57/61) per cent of pups were having Apgar scores of 0 to 4, 5 to 9 and 10 to 14, respectively. In CS group, out of 4 pups which had low Apgar scores of 0 to 4 at 30 min, 3 pups died and one pup remained in the same group at the end of 2 h and subsequently died at 24 h. Out of 8 pups which had Apgar scores of 5 to 9 at 30 min, 3 died and 2 had improved Apgar scores of 10 to 14 at the end of 2 h. The remaining 3 pups failed to survive through 24 h.

All the 57 live pups at the end of 24 h showed Apgar scores of 10 to 14. Out of 28 pups live at birth born through spontaneous whelping all had good Apgar scores of 10 - 14 and all survived after 24 h. In the AW group, of the 13 pups, 5 had scores of 5-9 at birth, of which 2 died and the remaining 3 showed improved Apgar scores at 2 h and further survived after 24 h. All the remaining 23 pups of this group with Apgar scores of 10 - 14 survived after 24 h.

Among the viable pups at birth, those with low and medium Apgar scores of 0 to 4 and 5 to 9, respectively, were more in the CS group. All the pups delivered through SW had highest Apgar scores at birth while the pups delivered through AW had shown Apgar scores between 5 to 9 and 10 to 14. This is in accordance with the results of Groppettiet al. (2010) who have reported 92 per cent pups delivered through CS, had shown poor Apgar. This reduction in the Apgar scores in CS group might be due to the effect of anaesthesia used in the CS. Propofol is a short acting hypnotic agent and provides rapid induction of recovery from anaesthesia and it crosses placental barrier. Propofol is also reported to cause more cardio respiratory depression and increased PCO2 which might be the reason for reduced Apgar scores in the pups delivered through the CS.

Similar results of the pups delivered through CS having lowest Apgar scores and that of SW with highest Apgar scores and those delivered through AW with intermediate and high Apgar scores were found at 30 min, 2 and 24 h. These results were in accordance with that of Veronesi et al. (2009), Groppettiet al. (2010) and Jayakumar et al. (2015). Bharathidasan et al. (2016), who reported that low Apgar scores in the pups delivered through CS beyond 4 and a half hour after the onset of labour and have added that the duration of delivery had an indirect relationship with the Apgar scores. They have also added that CS should be performed within 4.5 h after the onset of second stage of labour and the number of inductions should be limited to one.

All the 28 live pups born through the SW at birth were having Apgar scores of 10 to 14 and all of them survived beyond 24 h. This result is similar to that of Jayakumaret al. (2015) whoreported that all the live pups born through SW have survived beyond 60 min, the numbers of which are slightly higher than Veronesi et al. (2009) who have reported that 92.6 per cent pups born through SW survived beyond 24 h.

Among the 28 live pups born through AW at birth, 23 were having Apgar scores of 10

to 14 and all survived beyond 24 h. Of the 5 pups having moderate Apgar scores of 5 to 9 at birth, 2 died and 3 pups had improved Apgar scores of 10 to 14 at the end of 24 h. Significant improvement in survival rates were observed when Apgar scores improved from medium to high grade at 2 h of evaluation after resuscitation,. The beneficial effects of resuscitation procedures were more apparent in improving the Apgar scores of puppies born with medium Apgar scores. Though dystocia promoted a long lasting bradycardia and slowed down Apgar scores progression in pups (Lucio et al., 2009), resuscitation measures helped to attain satisfactory improvement.

Among 19 live pups born through CS, 11 were having low Apgar scores of 0 to 4 and died at 30 min. 8 were having moderate Apgar scores of 5 to 9, among which 3 had improved to high Apgar scores of 10 to 14 at 2 h and remaining 5 pups died even after adopting resuscitation procedure. Systemic evaluation of puppies in this study allowed the timely detection of puppies with poor Apgar scores that might otherwise have been overlooked at birth. Three out of 8 pups with Apgar scores of 5 to 9 recovered completely which proved that prompt detection of less viable newborns, followed by attempts to resuscitate, could improve neonatal survival.

To conclude, the type of delivery had a significant impact on Apgar scores with survival rate being lowest for pups born of CS and adequate prompt adoption of resuscitation measures would help in reducing the neonatal mortality in pups.

## ACKNOWLEDGEMENTS

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## REFERENCES

- Bharathidasan, M., George, R.S., Shafiuzama, M., Kannan, T.A and Ramesh, S. (2016). Modified Apgar score system for prediction of neonatal survival puppies delivered through cesarean section. International Journal of Developmental Research, 6(11):10281-10284.
- Davidson, A.P. (2014). Neonatal resuscitation improving the outcome. Veterinary Medicine, Small Animal Clinician, 44:191-204.
- Groppetti, D., Pecile, A., Del Carro, A.P., Copley, K., Minero, M and Cremonesi, F. (2010). Evaluation of newborn canine viability by means of umbilical vein lactate measurement, Apgar score and uterine tocodynamometry. Theriogenology, 74 (7): 1187-1196.
- Jayakumar, C., Krishnaswamy, A., Sudha and Gand Honnappa, T.G. (2015). Assessment of canine neonate by Apgar scoring system and its value as an index of short term neonatal survival. Indian Journal of Animal Reproduction, 36 (2):14-20.
- Lucio, C. F., Silva, L.C.G., Rodrigues, J.A., Veiga, G.A.L and Vannucchi, C. I. (2009). Acid-base changes in canine neonates following normal birth or dystocia. Reproduction in Domestic Animals, 44(2):208–210.
- Mosier, J.E. (1986). Small animal reproduction and infertility. A clinical approach to diagnosis and treatment. Ed. T.J. Burke, Lea and Febiger, Philadelphia. Pp. 335-345.
- Veronesi, M.C., Panzani, S., Faustini, M and Rota, A. (2009). An Apgar scoring system for routine assessment of newborn puppy viability and short-term survival prognosis. Theriogenology, 72(3):401-407.

Parameters	Score 0	Score 1	Score 2
Mucous colour (Appearance)	Cyanotic, pale	Pink	Reddish
Heart rate (BPM)	<120	120-180	>180
Respiratory rate (bpm)	<15	15-30	>30
Reflex irritability	None	Feeble reaction	Active reaction
Mobility	None	Hypo-mobility	Active mobility
Suckling	None	Weak	Energetic
Vocalization	None	Mild	Vigorous

Table 1.Apgar score model used in this study

BPM: beats/min; bpm: breaths/min

# Table 2. Percentage of pups with different Apgar scores at birth in relation to the type of delivery

	Total no. of	% of pups with different Apgar scores at birth					
Type of delivery	pups (n)	0 to 4	5 to 9	10 to 14			
Spontaneous whelping	30	5.56 (2)	0.00	54.90 (28)			
Assisted whelping	35	19.44 (7)	38.46 (5)	45.10 (23)			
Caesarean section	35	75.00 (27)	61.54 (8)	0.00			
Total	100	36	13	51			

Values within parenthesis indicate numbers.

Type of				Percel	ntage of vi	iable pups v	vith differ	ent scores				
Whelping		At birth			At 30 min			At 2 hrs			At 24 h	s
	0 to 4	5 to 9	10 to 14	0 to 4	5 to 9	10 to 14	0 to 4	5 to 9	10 to 14	0 to 4	5 to 9	10 to 14
Spontaneous whelping	0	0	54.90 (28)	0	0	50.91 (28)	0	0	49.12 (28)	0	0	49.12 (28)
Assisted whelping	0	38.46 (5)	45.10 (23)	0	37.50 (3)	41.82 (23)	0	0	45.62 (26)	0	0	45.62 (26)
<b>Caesarean</b> section	100.00 (11)	61.54 (8)	0	100.00 (4)	62.50 (5)	7.27 (4)	100.00 (1)	100.00 (3)	5026 (3)	0	0	5.26 (3)
Total	14.67 (11)	17.33 (13)	68.00 (51)	5.97 (4)	11.94 (8)	82.09 (55)	<b>1.64</b> (1)	4.92 (3)	93.44 (57)	0	0	100.00 (57)
Values within p	arenthesis	indicate nu	umbers.									

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# Lifetime semen production performance of cattle and buffalo bulls

# K.G. Bhave<sup>\*1</sup>, K. Thilak Pon Jawahar<sup>2</sup>, P. Kumarasamy<sup>3</sup>, T. Sivakumar<sup>4</sup>, C. Joseph<sup>5</sup>, S. Sontakke<sup>6</sup>, J. Khadse<sup>7</sup> and R.Venkataramanan<sup>8</sup>

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#### ABSTRACT

Semen production performance data of 88 buffalo and 179 cattle bulls of different breeds from 1975 to 2018, were collected from two frozen semen stations of BAIF Research Development Foundation. The objective of the present study was to document the lifetime semen producing performanceof cattle and buffalo bulls. Traits studied were lifetime frozen semen production doses (LFSP),lifetime production period (LPP),age at first semen production (AFSP) and age at last semen production (ALSP). The factors tested for influence on these traits were breed, period and season of birth.Overall means of AFSP, ALSP, LPP and LFSP were 1859.6, 4257.2, 2581.7 days and 183631.9 semen doses in buffalo bulls, while 1386.7, 4029.4, 2076.16 days and 264566.3 semen doses in cattle bulls.Production performance for results obtained for various breeds could be useful in devising important culling and disposal policies for AI centres in the country.

Key words: Semen Production, Lifetime Production, Breed

#### INTRODUCTION

Artificial insemination (AI), one of the widely used biotechnological tools is used to coverthe large population of cattle and buffalo in India. AI is not only used for the genetic improvement of livestock but also for the conservation purpose.With the consistent growing population of cattle and buffalo (PKR, 2019), mostly consisting of non-descript population, the demand for good quality semen is on rise to <u>upgrade the non-descript population</u>.

The information on semen production capacity of bulls of different breeds would prove useful inplanning to covermaximum breedable population. The requirement of semen doses of a

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particular breed over a period of time would help decide a minimumproduction target. A knowledge of the average first and last collection of freezable semen would help serve in the selection, culling and replacement decisions as the AI centersneed adult bulls and testing bulls to replace the adult stock. With this objective, the present study was designed to document the lifetime semen producing performance of cattle and buffalo bull breeds reared in the AI Stud Farm.

## MATERIALS AND METHODS

Semen production performance data of 88 buffalo and 179 cattle bulls of different breeds were collected from two frozen semen stations of BAIF Research Development Foundation, viz. Uruli Kanchan, Pune, Maharashtra for the years between 1975-2018, and at Dharouli, Jind, Haryanafor the years between 2011-2018. The buffalo breeds included in the study were Banni (2), Bhadawari (4), Murrah (66), Jaffarabadi (5), and Surti(11) while the cattle breeds included in the study were Gir (16), Hallikar (4), Holstein Friesian (HF)100% (30), HF50% (31), HF62.5% (4), HF75% (27), Jersey 100% (25), Jersey 50% (12), Jersey 62.5% (7), Jersey 75% (9), Khillar, (2) and Sahiwal (12). The date of birth, first and last collection records and frozen semen doses produced per ejaculate were utilized for the analysis. The traits considered for the analysis were lifetime frozen semen doses(LFSP),lifetime production production period (LPP), age at first semen production (AFSP) and age at last semen production (ALSP). The period of birth was divided into five groups viz. <1980, 1981-1990, 1991-2000, 2001-2010 and 2011-2018. Season of birth were divided into three categories viz. summer (March to May), monsoon (June to October) and winter (November to February). Until 2004, the sperm concentration per frozen semen dose was 30 million. From 2005 onwards, the sperm concentration per semen dose was revised from 30 million to 20 million per dose. Considering this fact, the LFSP was studied only for the period after 2005. The records were subjected to statistical analysis using least square analysis using "*lm*" function in R statistical software (RCT, 2019 version 4.0.1.). Along with "*lm*" function, "*car*" and "*agricolae*" packages were used for computing ANOVA (Type III sum of square) and Duncan's multiple range test. The model for the analysis is given below:

$$\mathbf{Y}_{iikl} = \boldsymbol{\mu} + \mathbf{B}_i + \mathbf{S}_i + \mathbf{A}_k + \mathbf{e}_{iikl}$$

Where,

- $Y_{ijkl}$  = Lifetime semen production performance trait
- $\mu$  = Overall mean
- $B_i = Effect of m^{th} breed (m = 1 to 5 or 12)$
- $S_i$  = Effect of n<sup>th</sup> season of birth (n = 1 to 3)
- $A_k$  = Effect of k<sup>th</sup> period of birth (k = 1 to 5)

Random error associated with Y<sup>ijkl</sup> which is assumed to be normally and independently

e<sub>ijkl</sub> = distributed with mean zero and constant variance

## **RESULTS AND DISCUSSION**

# Lifetime semen production performances of Buffalo and Cattle bulls

The least square means of lifetime frozen semen production traits like age at firstsemen production (AFSP) and last semen production (ALSP), lifetime frozen semen production doses (LFSP) and lifetime semen production period (LPP) of buffalo and cattle bulls are presented in Tables 1 and 2, respectively.

Breed had significant effect on AFSP. Banni and Bhadawari breeds had a higher AFSP in comparison with the other breeds while Murrah had the lowest AFSP. LFSP was highest in Murrah and lowest in Bhadawari buffalo breed. Banni and Jaffarabadi bulls did notdifferfrom each other in these traits. Breed showed no significance for the ALSP, LFSP, and LPP traits.

The breed differences in the age at first collection of buffalo bull semen were due to difference in the age of the bulls purchased. Murrah bulls were purchased at a very young age of 18 to 24 months, while Banni and Bhadawari bulls were purchased at a comparatively older age of 3 years, subsequently thus delaying the collection of their first ejaculate. Mukhopadhyay *et al.* (2010), Khatun *et al.* (2013) and Ramajayan (2016) had reported a still lower age at first semen collection in Murrah buffaloes. Early sexual maturity leads to collection of semen at an early age resulting in an overall production of semen doses, reduction in generation intervaland subsequentlyresulting inincreased genetic gain.

Breed effect on lifetime production of semen doses was particularly due to preferential treatment at the semen station over the bulls. Murrah breeds are in huge demand in the field unlike the Banni and Bhadawari which have a very small breeding tract. They are also not been used for grading up purposes. Thus, the cattle breeds with less demand were subjected to fewer collections of semen and which resulted in the production of lesser number of doses.

From the averages presented for age at last semen collection and lifetime production period, it was also observed that on an average, the semen collection was carried out for approximately 8-9 years of productive life and which was found similar for all the breeds and may perhaps explain the reason for the non-significant effect due to breed.

All the lifetime production traits except LPP was found to be significantly affected by breed factor in cattle. AFSP was found to be the lowest in HF pure bulls. However, HF pure bulls did not differ significantly from other HF crossbreds and Jersey and their crossbred bulls except Jersey 62.5%

bulls which had the highest AFSP. Indigenous bulls showed comparatively a higher AFSP and ALSP than the exotic and crossbred bulls. HF 100% bulls had the highest LFSP means in comparison to other breeds. Apart from Gir and Hallikar bulls, all the other breeds did not differ statistically from each other.

The different genotypic groups present, the number of bulls available in a particular genotype, the demand in field and the age of puberty was found to have a significant effect on all the lifetime semen production traits except lifetime production period in cattle. The exotic breeds HF and Jersey had a lower age of semen production and thus produced higher semen doses in comparison with the other breeds. This was followed by the crossbred bulls which had a lesser age at puberty in comparison with the indigenous breeds of cattleand which had a huge demand in the field for crossbreeding purposes. Thus the semen production from each pure and crossbred bull was observed to be higher than that produced by the indigenous bulls. The Bos taurus bulls attain sexual maturity at the age of 8 to 10 months of age while the Bos indicus bulls take a little longer to attain sexual maturity viz., 16-18 months (Wolf et al., 1965; Almquist et al., 1976; Lunstra et al., 1978; Chenoweth et al., 1996 and Vale Filho et al., 1997). Brito et al. (2002) reported a slightly higher age at puberty in crossbred bulls (13.8 months; 421 days). Age at puberty depends on the early release of gonadotropin secretion and which usually increases between 2 to 5 months after birth. It had been observed that the pattern of secretion of gonadotropin differedbetween the early and late maturing bulls. A higher concentration of luteinizing hormone was found in early maturing bulls than in late maturing bulls(Evans et al., 1995 and Aravindakshan et al., 2000). Age at last semen production in cattle bulls showed little variation among the breeds (Table 2). It was also observed that the average lifetime production period among the cattle breedsin the present study was 6 to 7 years. This may perhaps explain the reason why there was no difference among the breeds for lifetime production period.

Age at first and last semen collection among the HF and Jersey crossbred and Sahiwal breed cited by different researchers (Rao, 1984; Suryaprakasam and Rao, 1993; Chauhan*et al.*, 2010; Mukhopadhyay *et al.*, 2010; Gopinathan, 2014; Khatun *et al.*, 2013 and Thippesamy *et al.*, 2014) reveal slightly lower means than the present study. Similarly, the lifetime production period and semen doses reported by Mukhopadhyay *et al.* (2010) and Gopinathan (2014) in Jersey and HF crossbred bulls were found to be lower than observed in the present study. The difference among the earlier reports and present study may perhaps be attributed to management measures practicedin the semen stations.

Period of birth had a significant effect on all the lifetime semen production traitsin cattle and buffalo bulls (except for lifetime frozen semen production in which the period of birth effect was not tested). The significant difference among the traits due to periodmay be due to the number of bulls studied and the increase in demand in the last two decades. Season of birth had no statistically significant effect on any lifetime production traits of both cattle and buffalo bulls.

## SUMMARY

Production performance for results obtained for various breeds could prove useful in culling decisions and disposal policies for AI centres in the country.

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## REFERENCES

- Almquist, J.O and Amann, R.P. (1976). Reproductive capacity of dairy bulls. XI. Pubertal characteristics and post-pubertal changes in production of semen and sexual activity of Holstein bulls ejaculated frequently. Journal of Dairy Science, 59:986–991.
- Aravindakshan, J.P., Honaramooz, A., Bartlewsliki, P.M., Beard, A.P., Pierson, R.A and Rawlings, N.C. (2000). Patterns of gonadotrophin secretion and ultra-sonographic evaluation of developmental changes in the testes of early and late maturing bull calves. Theriogenology, 54:339–354.
- Brito, L.F., Silva, A.E., Unanian, M.M., Dode, M.A., Barbosa, R.T and Kastelic, J.P. (2004). Sexual development in early and late maturing Bos indicus and Bos indicus× Bos taurus crossbred bulls in Brazil. Theriogenology, 62(7): 1198-1217.
- Chauhan, I.S., Gupta, A.K., Khate, K., Chauhan, A., Rao, T.K.S., Pathak, S., Hazra, R and Singh, M. (2010). Genetic and non-genetic factors affecting semen production traits in Karan Fries crossbred bulls. Tropical Animal Health and Production, 42:1809-1815.
- Chenoweth, P.J., Chase Jr, C.C., Thatcher, M.J.D., Wilcox, C.J and Larsen, R.E. (1996). Breed and other effects on reproductive traits and breeding soundness categorization in young beef bulls in Florida. Theriogenology, 46:1159–1170.
- Evans, A.C.O., Davies, F.J., Nasser, L.F., Bowman, P and Rawlings, N.C. (1995). Differences in early patterns of gonadotrophin secretion

between early and late maturing bulls, and changes in semen characteristics at puberty. Theriogenology, 43:569–78.

- Gopinathan, A. (2014). Genetic studies on semen production in Jersey crossbred and Holstein Friesian crossbred bulls. M.V.Sc dissertation submitted to Tamil Nadu Veterinary and Animal Sciences University, Chennai, Tamil Nadu, India.
- Khatun, M., Kaur, S and Kanchan, C. S. (2013). Subfertility problems leading to disposal of breeding bulls. Asian-Australasian Journal of Animal Sciences, 26(3): 303.
- Lunstra, D.D., Ford, J.J and Echternkamp, S.E. (1978). Puberty in beef bulls: hormone concentrations, growth, testicular development, sperm production, and sexual aggressiveness in bulls of different breeds. Journal of Animal Science, 46:1054–1062.
- Mukhopadhyay, C.S., Gupta, A.K., Yadav, B.R., Khate, K., Raina, V.S., Mohanty, T.K and Dubey, P.P. (2010). Subfertility in males: an important cause of bull disposal in bovines. Asian-Australian Journal of Animal Science, 23:450-455.
- PKR. (2019). Provisional Key Results of 20<sup>th</sup> Livestock Census. (2019). Department of Animal Husbandry Dairying, Government of India Accessed at http://dahd.nic.in/division/ provisional-key-results-20th-livestock-census.

- RCT. (2019). R: A language and environment for statistical computing. R Foundation for Statistical Computing, Vienna, Austria. Version: 3.6.1.
- Ramajayan, P. (2016). Genetic studies on semen production in Murrah buffalo bulls reared in Tamil Nadu. M.V.Sc dissertation. Tamil Nadu Veterinary and Animal Sciences University, Chennai, Tamil Nadu, India.
- Rao, A.V.N. (1984). Reproductive efficiency of exotic and crossbred AI bulls in Andhra Pradesh. Indian Veterinary Journal, 61:431-433.
- Suryaprakasam, T.B and Rao, A.V.N. (1993). Studies on breeding and disposal pattern of A.I. sires in Andhra Pradesh. Indian Veterinary Journal, 70:1022-1024.
- Thippeswamy, V.B., Layek, S.S., Kumaresan, A., Mohanty, T.K., Gupta, A.K., Chakravarty, A. K and Prasad, S. (2014). Effects of pedigree and exotic genetic inheritance on semen production traits of dairy bulls. Asian Pacific Journal of Reproduction, 3(1):13-17.
- Vale Filho, V.R., Bergmann, J.A.G., Andrade, V.J., Quirino, C.R., Reis, S.R and Mendonca, R.M.A. (1997). Andrologic characterization of Nellore bulls selected for the first breed season. Revista Brasileira de Reproduo Animal, 21:42–44.
- Wolf, F.R., Almquist, J.O and Hale, E.B. (1965). Prepuberal behaviour and puberal characteristics of beef bulls on high nutrient allowance. Journal of Animal Science, 24:761–765.

## Table 1.

Levels	AFSP	ALSP	LFSP	LPP
Overall mean	1859.6±110.52	4257.2±229.8	183631.9±25077	2581.7±235.2
Breed	**	NS	NS	NS
Banni	2056±234.1ª	5175±710	171812±65962	2765±736
Bhadawari	2270±188.4ª	4548±524	100053±65300	1771±543
Jaffarabadi	1782±120.3 <sup>ab</sup>	4538±480	209573±47807	2644±498
Murrah	1585±78.1 <sup>b</sup>	4835±170	242330±21673	3166±176
Surti	1729±110.1 <sup>ab</sup>	5039±301	-	2991±312
Period of birth	**	**	-	**
≤1980	1416±269ª	6477±536ª	-	2975±555 <sup>ab</sup>
1981 to 1990	1315±137°	4916±291 <sup>b</sup>	-	3441±301ª
1991 to 2000	1712±129 <sup>b</sup>	3850±320°	-	1725±332 <sup>b</sup>
2001 to 2010	1288±109°	4065±208°	-	2528±216 <sup>b</sup>
2011 to 2018	1567±226 <sup>bc</sup>	-	-	-
Season of birth	NS	NS	NS	NS
Summer	1879±133	4781±296	175205±33227	2629±307
Monsoon	1849±117	5017±275	166817±44333	2816±286
Winter	1851±116	4684±284	200804±34978	2557±294

## Least square mean of lifetime semen production traits in buffalo bulls

(LFSP: Lifetime frozen semen production doses; LPP: Lifetime production period; AFSP: Age at first semen production; ALSP: Age at last semen production)

## Table 2.

Levels	AFSP	ALSP	LFSP	LPP
Overall mean	1386.71±67.8	4029.4±109.08	264566.3±23939.1	2076.16±148.6
Breed	**	*	*	NS
Gir	1495±148.4 <sup>b</sup>	4162±270 <sup>b</sup>	151015±50337 <sup>b</sup>	1844±291
Hallikar	1741±277.5 <sup>ab</sup>	4263±504 <sup>b</sup>	122239±87186 <sup>b</sup>	1502±543
HF100%	757±52.6 <sup>b</sup>	3936±176 <sup>b</sup>	391348±45254ª	2295±189
HF50%	1259±99.5 <sup>b</sup>	4247±197 <sup>ab</sup>	298543±62184 <sup>ab</sup>	2382±212
HF62.5%	1203±163.2 <sup>b</sup>	5131±511 <sup>ab</sup>	-	2293±551
HF75%	1135±74 <sup>b</sup>	4395±209 <sup>ab</sup>	347515±49024 <sup>ab</sup>	2480±225
JR100%	941±59 <sup>b</sup>	3910±178b ab	236561±53524 <sup>ab</sup>	2176±192
JR50%	1211±134.3 <sup>b</sup>	4731±297 <sup>ab</sup>	354490±67826 <sup>ab</sup>	2711±320
JR62.5%	2256±189.1ª	5442±377ª	-	1660±407
JR75%	1218±157.3 <sup>b</sup>	4332±336 <sup>ab</sup>	313282±107676 <sup>ab</sup>	2175±362
Khillar	1315±366 <sup>b</sup>	3996±699 <sup>b</sup>	187285±111309 <sup>ab</sup>	1742±754
Sahiwal	1663±150.8 <sup>ab</sup>	4392±298 <sup>ab</sup>	243387±89101 ab	1648±321
Period of birth	**	**	-	**
≤1980	1547±104.6 <sup>b</sup>	5049±378ª	-	1089±408ª
1981 to 1990	1762±104.1ª	4836±192 <sup>ab</sup>	-	2154±207 <sup>b</sup>
1991 to 2000	1290±86.2 <sup>b</sup>	3952±147 <sup>b</sup>	-	2530±159 <sup>b</sup>
2001 to 2010	1193±69.9 <sup>b</sup>	3809±133 <sup>b</sup>	-	2530±143 <sup>b</sup>
2011 to 2018	1268±93.6b	-	-	-
Season of birth	NS	NS	NS	NS
Summer	1463±82	4332±174	239561±41027	2047±187
Monsoon	1383±76.8	4536±184	241412±36884	2079±199
Winter	1390±77	4367±158	312726±36699	2100±170

## Least square mean of lifetime semen production traits in cattle bulls

(LFSP: Lifetime frozen semen production doses; LPP: Lifetime production period; AFSP: Age at first semen production; ALSP: Age at last semen production)

# **Expression of Growth Differentiation Factor-9 from buffalo follicular fluid – a marker gene for fecundity**

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#### ABSTRACT

Growth factors synthesized from mammalian oocytes popularly known as Oocyte Secreted Factors (OSFs) play numerous role in ovarian folliculogenesis. Growth differentiation factor-9 (GDF-9) produced within the ovary plays an essential role during follicle maturation through their actions on granulosa cells, but their expression in follicular fluid has not been studied. The purpose of this study was to detect the temporal expression pattern of GDF-9 from the follicular fluid of buffalo ovary, which may be correlated with the oocytes and embryo quality. The sensitive messenger ribonucleic acid detection of GDF-9 from follicular fluid was determined by Reverse Transcriptase-Polymerase Chain Reaction using bovine oligonucleotide primers. Our finding showed the qualitative detection of GDF-9 mRNA transcripts from follicular fluid of buffalo ovary.

Key words: Buffalo, Follicular fluid, GDF-9 expression, RT-PCR

## INTRODUCTION

Growth Differentiation Factor-9 (GDF-9), a prominent member of the Transforming Growth Factor- $\beta$ (TGF- $\beta$ ) superfamily functions as paracrine factor in the regulation of granulosa cell proliferation and differentiation (Gilchrist *et al.*, 2006). Furthermore, GDF-9 is the first oocyte derived growth factor found to be indispensable in ovarian folliculogenesis and expressed throughout the development of the maturing follicle. Besides its activities in cumulus expansion, oocyte maturation and ovulation (Gui and Joyce, 2005; Yoshino *et*  *al.*, 2006), italso play important roles during many steps of ovarian follicular development. GDF-9 involved in cell survival signaling (Orisaka *et al.*, 2006) and act as modulators of many other growth factors and endocrine hormones (Juengel *et al.*, 2004).

Given these results, GDF-9may be considered as a candidate gene for controlling ovulation rate in buffalo. Several studies have reported that supplementation of *in vitro* maturation media with FSH enhanced the cumulus

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expansion, nuclear maturation and cleavage rate of buffalo oocytes (Chauhan *et al.*,1996; Abdoon *et al.*, 2001). Few other studies have reported the beneficial effects of including follicular fluid replacing hormones during *in vitro* maturation of buffalo oocytes (Chauhan *et al.*, 1997).

In the ovaries, GDF-9 has been expressed in the oocytes of primordial follicles of sheep and cows (Bodensteiner et al., 1999). The expression begins at the primary follicle stage in women (Aaltonen et al., 1999), mice (Elvin et al., 1999) and also in rats (Jaatinen et al., 1999). GDF-9 mRNA expression has also been identified in extraovarian sites such as the pituitary, hypothalamus, placenta, testis and adrenal gland (Pennetieret al., 2004; Faure et al., 2005; Farnworth et al., 2006). Follicular fluid contains proteins, anticoagulants, enzymes, etc., thereby providing the oocytes with a microenvironment which contains the necessary regulatory factors (Fortune, 1994). Since GDF-9 affect oocyte development during folliculogenesis, follicular fluid level of this oocyte secreted paracrine factor may be important for oocyte function and subsequent embryo quality. Although GDF-9 and its functions in mammals have captured increased attention in recent years, much remains to be learnt about this molecule in buffalo. There is a lack of information on the availability of this growth factor from follicular fluid. Therefore, this study was designed to investigate the expression of GDF-9 from follicular fluid of buffalo ovary.

## MATERIALS AND METHODS

Buffalo ovaries were collected from a local abattoir at Perambur, Chennai within two hours of slaughter and washed five times in phosphate buffered saline (PBS) containing 100 IU/ml penicillin and 100  $\mu$ g/ml streptomycin. The aspiration of non-atreticantral follicles (3-8 mm) was carried out using 18-gauge needle connected to a 10 ml disposable syringe (Fig. 1). The aspirated contents were centrifuged at 500g for 30 minutes in a cooling centrifuge at 4°C. After settling of follicular cells, supernatant part of follicular fluid was slowly aspirated and transferred into a fresh 1.5 ml microcentrifuge tube and then frozen at -80°C for RNA purification.

Total RNA was purified under RNase-free conditions at room temperature for each sample using Trizol Reagent (Invitrogen) according to the protocol of the manufacturer. In microcentrifuge tube, 250µl of cells free follicular fluid was lysed by the addition of 750µl Trizol followed by incubation at room temperature for 5 minutes. Then, 200µl of chloroform was added, vortexed for 5 seconds and incubated at room temperature for 5 minutes. The centrifugation at 12,000 rpm for 15 minutes at 4°C in cooling centrifuge separated the solution into an upper aqueous phase containing RNA and a lower phenol-containing organic phase. The upper aqueous phase was transferred to a clean, RNase-free eppendorf tube and again centrifuged at 12,000 rpm for 10 minutes at 4°C. Then, the pellet was washed with 1ml of 75% ice cold ethanol, subsequently centrifuged at 7500 rpm for 5 minutes at 4°C and ethanol was discarded. The microcentrifuge tube was air dried and suspended with 10ul of DEPC water. The purified total RNA was stored at 80°C. The quantity and purity of the RNA were determined by spectrophotometry.

The cDNA synthesis was carried out using Revert Aid M-MuLV Reverse Transcriptase Kit (Fermantas, USA)in a standard  $20\mu$ l reaction mixture. One microgram oftotal RNA was used as template RNA for reverse transcription as per manufacturer's instructions. RT-PCR for GDF-9 was carried out by using the published (NCBI accession No: AB058416) forward (5' AGAAGCTGCTGAGGGTGTAAGATT3') and thereverse(5'AAGCAATTGAGCCATCAGGC3') primer sequences (Hosoe*et al.*, 2011) from bovine. These oligonucleotide primers amplified a fragment size of 401 base pair. The  $\beta$ -actin forward primer5'-GATGAGGCTCAGAGCAAGAGA-3' and reverse Expression of growth differentiation factor-9 from buffalo follicular fluid – a marker gene for fecundity

primer 5'-TCGTCCCAGTTGGTGACGAT-3'(amplicon size 596 was used as the internal control for Polymerase Chain Reaction). The PCR programme comprised an initial denaturation at 95°C for 5 minutes, followed by denaturation at 95°C for 30 seconds, annealing at 52°C for 30 seconds, and extension at 72 °C for 1 minutes and final extension at 72 °C for 5 minutes. Each PCR was performed for 30 cycles for each sample. Products of the RT-PCR were separated by electrophoresis on 1.5% agarose gel and visualized by ethidium bromide staining.

## **RESULTS AND DISCUSSION**

In this PCR study, the existence of GDF-9 mRNA expression in follicular fluid of buffalo ovary was observed. The presence of this fecundity gene from follicular fluid of buffalo was detected by using the reverse transcriptase PCR. Fig. 2 shows the PCR ampliconsof GDF-9 (NCBI accession No: AB058416) along with  $\beta$ -actin gene (NCBI accession No: BG689033) electrophoresed through 1.5% agarose gel.



Fig.1. Picture showing ovaries of buffalo with follicles. Arrow mark showing fully grown good quality follicle from which the follicular fluid was collected.



Fig. 2.Agarose gel showing PCR amplicons of GDF-9 (401 bp) and  $\beta$ -actin (endogenous control) from follicular fluid of buffalo ovary.Lane 1: 100bp Marker, Lane II: GDF-9 (401bp), Lane III:  $\beta$ -actin (596bp).

Folliculogenesis is a complex process dependent upon the intricate interplay of various growth factors and hormones. It is now well established that during ovarian folliculogenesis there is a dynamic interplay between the oocyte and the surrounding somatic cells that mutually influences growth and differentiation of the somatic granulosa cells and the oocyte (Gilchrist et al., 2004). The members of the TGFB superfamily influence a wide variety of growth and differentiation processes in a number of tissues and species. The newest member of this superfamily, GDF-9 appears to be no exception from this important function. Presumably, GDF-9 is the candidate gene for increasing the ovulation rate in sheep (McNatty et al., 2003). The essential role of GDF-9 in fertility was demonstrated by targeted deletion of the GDF-9 gene in mice (Dong et al., 1996). Long term immunization against GDF-9 in sheep disrupts early folliculogenesis and leads to the absence of normal follicles beyond the primary stage of development (Juengel et al., 2002). In addition, a naturally occurring mutation in this gene causes infertility in sheep (Hanrahanet al., 2004). Positive staining for the GDF9 protein was identified in oocytes and granulosa cells in human ovarian samples (Oronet al., 2010) and in buffalo oocytes (Kathirvel et al., 2013).

In the present study, we have reported the expression of GDF-9 from the follicular fluid of buffalo ovary. There are no expression studies available to support the detection of GDF-9 gene from ovarian follicular fluid of buffalo. From previous ovarian expression studies, GDF-9 transcripts have been found to be localized in the oocytes from primordial follicles in cows and sheep (Bodensteiner et al., 1999), whereas it is not found in primordial follicles of mice. These investigations suggest that expression pattern of GDF-9 differs between species. In goats, the polymerase chain reaction technique revealed the presence of GDF-9 mRNA in primordial, primary and secondary follicles, oocytes and granulosa cells of antral follicles (Silva et al., 2004). Moreover it is reported earlier that the numbers of antral follicles per unit of ovarian mass and expression GDF9 serve as an important clue for higher prolificacy (Pramodet al., 2013). Although results of our RT-PCR revealed the availability of GDF-9 expression from follicular fluid, the relative amount of mRNA transcripts of this gene need to be analysed through quantitative real time-PCR for further understanding the exact level of GDF-9 in follicular fluid of different category follicles. Additional studies on preantral follicles are needed to clarify the role of GDF9 as well as other growth factors.

In summary, the present study has demonstrated for the first time, GDF-9 expression from follicular fluid of buffalo ovary. This evidence of expression of GDF-9 in buffalo ovarian follicular fluid supports the effect of supplementation of buffalo follicular fluid during *in vitro* maturation of buffalo oocytes in improving the maturation rate. Moreover, the precise physiologic role of GDF-9 protein from follicular fluid in influencing oocyte needs further study.

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## REFERENCES

- Abdoon, A.S.S., Kandil, O.M., Otoi, T and Suzuki, T.(2001) Influence of oocyte quality, culture media and gonadotrophins on cleavage rate and development of *in vitro* fertilized buffalo embryos. *Animal Reproduction Science*, **65**:215–223.
- Aaltonen. J., Laitinen, M.P., Vuojolainen, K.,Jaatinen, R.,Horelli-Kuitunen, N.,Seppä, L.,Louhio, H.,Tuuri, T.,Sjöberg, J.,Bützow, R. Hovata. O. Dale.L and Ritvos.O., (1999). Human growth differentiation factor 9 (GDF- 9) and its novel homolog expressed GDF-9B are in oocytes during early folliculogenesis. Journal ofClinicalEndocrinology&Metabolism, 84:2744-2750.
- Bodensteiner, K.J., Clay, C.M.,Moeller,C.L and Sawyer,H.R. (1999). Molecular cloning of the ovine growth/differentiation factor-9 gene and expression of growth/ differentiation factor-9 in ovine and bovine ovaries.*Biology ofReproduction*, **60**:381– 386.
- Chauhan, M.S., Katiyar, P.K., Madan, M.L., Singla, S.K and Manik, R.S. (1996). Influence of follicle stimulating hormone on *in vitro* maturation and cleavage of buffalo (*Bubalusbubalis*) oocytes after *in vitro* fertilization. *Theriogenology*, **45**:243.
- Chauhan, M.S., Palta, P., Das, S.K., Katiyar, P.K and Madan, M.L. (1997). Replacement of serum and hormone additives with follicular fluid in the IVM medium: effects

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of maturation, fertilization and subsequent development of buffalo oocytes *in vitro*. *Theriogenology*, **48**:461–469.

- Dong, J., Albertini, D.F., Nishimori, K., Kumar, T.R., Lu, N and Matzuk, M.M. (1996). Growth differentiation factor-9 is required during early ovarian folliculogenesis. *Nature*, 383:531–535.
- Elvin, J.A., Clark, A.T., Wang, P., Wolfman, N.M and Matzuk, M.M. (1999). Paracrine actions of growth differentiation factor-9 in the mammalian ovary. *MolecularEndocrinology*, **13**:1035–1048.
- Farnworth. P.G., Wang, Y.,Leembruggen, P.Ooi. G.T.,Harrison, С., Robertson D.M and Findlay, J.K. (2006). Rodent adrenocortical cells display high affinity binding sites and proteins for inhibin A, and express components required autocrinesignallingby for activins and bone morphogenetic proteins. JournalofEndocrinology, 188:451-465.
- Faure, M.O., Nicol, L.,Fabre, S.,Fontaine, J.,Mohoric, N.,McNeilly,A and Taragnat,C. (2005). BMP-4 inhibits follicle-stimulating hormone secretion in ewe pituitary. *Journal ofEndocrinology*,**186**:109–121.
- Fortune, J.E. (1994).Ovarian follicular growth and development in mammals.*Biology of Reproduction*. **50**:225–232.
- Gilchrist, R.B., Ritter,L.J and Armstrong,D.T. (2004). Oocyte-somatic cell interactions during follicle development in mammals. *Animal Reproduction Science*, **82–83**:431– 446.
- Gilchrist, R.B., Ritter, L.J.,Myllymaa, S.,Kaivo-Oja, N.,Dragovic, R.A.,Hickey, T.E.,Ritvos,O and Mottershead,D.G. (2006). Molecular basis of oocyte-paracrine

signaling that promotes granulosa cell proliferation. *Journal ofCell Science*, **119**: 3811–3821.

- Gui, L.M and Joyce I.M. (2005). RNA interference evidence that growth differentiation factor-9 mediates oocyte regulation of cumulus expansion in mice. *Biology of Reproduction*, **72**:195–199.
- Hanrahan, J.P., Gregan, S.M., Mulsant, P., Mullen, M., Davis, G.H and Powell.R. (2004). Mutations in the genes for oocyte-derived growth factors GDF-9 and BMP-15 are associated with both increased ovulation rate and sterility in Cambridge and Belclare sheep (*Ovis Aries*). *Biology of Reproduction*, **70**:900–909.
- Hosoe, M., Kaneyama, K., Ushizawa, K., Hayashi, K.G and Takahashi,T.(2011). Quantitative analysis of bone morphogenetic protein 15 (BMP15) and growth differentiation factor 9 (GDF9) gene expression in calf and adult bovine ovaries.*Reproductive Biology and Endocrinology*, **9**:33.
- Jaatinen, R., Laitinen, M.P., Vuojolainen, K., Aaltonen, J., Louhio, H., Heikinheimo, K., Lehtonen, E and Ritvos, O. (1999). Localization of growth differentiation factor-9 (GDF-9) mRNA and protein in rat ovaries and cDNA cloning of rat GDF-9 and its novel homolog GDF-9B.*Molecular and Cellular Endocrinology*, **156**:189–193.
- Juengel, J.L., Hudson, N.L.,Heath, D.A.,Smith, P.,ReaderK.L and Lawrence,S.B. (2002). Growth differentiation factor 9 and bone morphogenetic protein 15 are essential for ovarian follicular development in sheep. *Biology of Reproduction*, 67:1777–1789.
- Juengel, J.L., Bodensteiner, K.J.,Heath, D.A.,Hudson, N.L.,Moeller, C.L.,Smith,

P.,Galloway, S.M.,Davis, G.H.,Sawyer, H.R and McNatty,K.P. (2004). Physiology of GDF9 and BMP15 signalling molecules. *AnimalReproduction Science*, **82–83**:447– 460.

- Kathirvel, M., Eswari, S., and Kumanan, V.(2013).
  Differential expression dynamics of Growth differentiation factor9 (GDF9) and Bone morphogenetic factor15 (BMP15) mRNA transcripts during in vitro maturation of buffalo (*Bubalusbubalis*) cumulus–oocyte complexes. *SpringerPlus*, 2:206-211.
- McNatty, K.P., Juengel, J.L.,Wilson, T.,Galloway, S.M.,Davis, G.H.,Hudson, N.L.,Moeller,C.L.,Cranfield, M.,Reader, K.L.,Laitinen, M.P.,Groome, N.P.,Sawyer,H.R and Ritvos,O. (2003). Oocyte- derived growth factors and ovulation rate in sheep. *Reproduction*, **61**(Suppl):339-351.
- Orisaka, M., Orisaka, S.,Jiang, J.Y.,Craig, J.,Wang, Y.,Kotsuji,F And Tsang,B.K. (2006). Growth differentiation factor 9 is antiapoptotic during follicular development from preantral to early antral stage. *Molecular Endocrinolology*, **20**:2456–2468.
- Oron, G., Fisch, B., Ao, A., Zhang, X.Y., Farhi, J., Haroush, A.B., Icekson, G.K and Abir, R. (2010). Expression of growth-differentiating

factor 9 and its type 1 receptor in human ovaries. *ReproductiveBioMedicine Online*, **21**:109–117.

- Pennetier, S., Uzbekova, S., Perreau, C., Papillier, P., Mermillod, P and Dalbies-Tran, R. (2004). Spatio-temporal expression of the germ cell marker genes MATER, ZAR1, GDF9, BMP15, and VASA in adult bovine tissues, oocytes, and preimplantation embryos. *Biology of Reproduction*, **71**:1359–1366.
- Pramod, R.K., Sharma, S.K., Singhi, A., Pan, S and Mitra, A. (2013). Differential ovarian morphometry and follicular expression of BMP15, GDF9 and BMPR1B influence the prolificacy in goat. *Reproduction in Domestic Animimals*,**48**: 803–809.
- Silva, J.R.V., Van den Hurk, R.,VanTol, H.T.A.,Roelen,B.A.J. and Fiueiredo,J.R. (2004). Expression of growth differentiation factor 9 (GDF9), bone morphogenetic protein 15 (BMP15), and BMP receptors in the ovaries of goats.*Molecular Reproduction* and Development, **70**:11–19.
- Yoshino, O., McMahon, H.E., Sharma, S. and Shimasaki, S. (2006). A unique preovulatory expression pattern plays a key role in the physiological functions of BMP-15 in the mouse. *Proceedings of the National Academy of Sciences of the United States of America*, **103**:10678–10683.

## Performance of farmbred Jersey cattle under tropical climatic conditions of Tamil Nadu

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## ABSTRACT

Data on production and reproduction performances of 373 Jersey cows born and reared at the Exotic Cattle Breeding Farm, Eachenkottai, Tamil Nadu, India, pertaining to 27 years were collected. They were analysed to study the effects of various non-genetic factors on milk production and reproduction traits. The least-squares means ( $\pm$  SE) for 305-day and total lactation milk yields for all the parities were 1491.6  $\pm$  25.9 and 1560.9  $\pm$  29.9 kg respectively. The averages for lactation length, service period, calving interval and dry period for all lactations were 303.1  $\pm$  4.0, 177.0  $\pm$  10.0, 461.0  $\pm$  9.7 and 160.2  $\pm$  9.7 days respectively. Years grouped into five periods had significant (P<0.01) influence on all the traits studied. Season of calving showed a significant (P<0.01) source of variation in milk yield traits and calving interval. Parity was found to influence 305-day milk yield, service period, dry period and calving interval significantly (P<0.01). The study revealed that the performance of the Jersey cattle was much lower than those maintained under high altitude conditions in Tamil Nadu and other places in India.

Key words: calving interval, dry period, heritability, milk yield, non-genetic factors

## INTRODUCTION

As a policy the system, the cross breeding programme has been adopted to alter the genetic makeup of native zebu stock in the tropics. This tool has had India, transverse a long way in the global scenario of milk production. For cross breeding, bulls of Holstein Friesian, Jersey and other European milch breeds were introduced into India. Among them, the Jersey breed was preferred because of its perceived better adaptability to tropical climate. Importation of either bulls or frozen semen alone could not fulfill the needs of cross breeding programme in India. Hence, a programme of raising bulls within the country was made by establishing bull mother farms at different places. For this, the Government of Tamil Nadu, India imported a herd of 150 pregnant Jersey heifers in 1978 and 1979 in two batches from Australia to produce bulls of high genetic merit and to ensure regular supply of good quality semen. Although investigations have been carried out on Jersey cattle maintained in bull mother farms (Singh and Mishra, 1980; Sadana and Tripathi, 1986; Sreemannarayana and Rao, 1994; Pal *et al.*, 2002; Venkataramanan*et al.*, 2007), a detailed study on Jersey cattle maintained under hot and humid climatic conditions of Tamil Nadu is lacking. Hence, an investigation on production and reproduction performances of the Jersey cattle

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born and bred on the farm has been made. This will be useful to understand the performance and the influence of various non-genetic factors affecting the economic traits of Jersey cattle.

#### MATERIALS AND METHODS

This study was based on the data pertaining to the progeny of imported Australian Jersey heifers and cows born and bred at Exotic Cattle Breeding Farm, Eachenkottai, Tamil Nadu, India from 1980 to 2006 (27 years). This farm is located in the east-coastal region of Tamil Nadu and the climate is generally hot, humid and tropical in nature. The Jersey cattle were housed in permanent sheds with open type ventilation and maintained under stall-fed conditions. Roughages in the farm of green fodder and paddy straw were provided. In addition, concentrate mixture was provided to all age groups as per the standard requirements. Cows were hand-milked twice daily in the morning and evening. Data on production and reproduction performances of farmbred Jersey cattle (1172 lactation records form 373 Jersey cows) were extracted from history and pedigree sheets. The traits studied were 305-day milk yield, lactation length, lactation milk yield, service period, calving interval and dry period. Period and season were the fixed environmental effects considered for all the traits. As the calvings were less in a year, year-season analysis was not done. To utilise all available data, the entire duration of the study was grouped into five periods and each year was classified into four seasons viz. winter (January to February), summer (March to May), southwest monsoon (June to September) and north-east monsoon (October to December). In addition, parity effect was also considered. First six parities were considered and parties six and above were lumped together as sixth parity. LSMLMW and MIXMDL PC-2 VERSION computer programme of Harvey (1990) was used to study the effect of various non-genetic factors. The model used for analysis was  $Y_{ijkl} = \mu + P_i + S_i + O_k + e_{ijkl}$ . Where,  $Y_{ijkl}$  = the l<sup>th</sup> observation in i<sup>th</sup> period, j<sup>th</sup> season and k<sup>th</sup> parity,  $\mu$ = overall mean when equal subclass frequencies exist,  $P_i$  = effect of i<sup>th</sup> period (<sub>i</sub> =1 to 5),  $S_j$  = effect of j<sup>th</sup> season (<sub>j</sub> =1 to 4),  $O_k$  = effect of k<sup>th</sup> parity (<sub>k</sub>=1 to 6) and e<sub>ijkl</sub> = random errors NID (0,  $\sigma_e^2$ ). The least-squares means were compared by Duncan's multiple range test.

### **RESULTS AND DISCUSSION**

The least-squares means  $(\pm SE)$  for production traits are presented in Table 1. The milk yield in period 1 (1980-84) was significantly (P<0.01) more when compared to other periods. North-east monsoon and winter calvers were found to give the highest milk yield and they were different (P<0.01) from summer and southwest monsoon season calvers. Most of the 305day milk yields reported for locally bred Jersey (Sreemannarayana and Rao, 1994; Pal et al., 2002; Ahmed et al., 2004; Venkataramanan, et al., 2007) was higher than the values observed in the present study. However, Singh and Mishra (1980) reported much lower value of 1176.2  $\pm$ 87.49 kg (n=17) for Jersey cattle maintained in Orissa, also a hot and humid region in India. A comparable value of 1399.10 kg was also reported by Sadana and Tripathi (1986) for Jersey cattle maintained at Palampur, Himachal Pradesh, a high altitude region in India. The significant influence of period and season of calving on 305-day milk yield and lactation milk yield in the present study corroborated with the findings of Chauhan (1990) and Murdia and Tripathi (1992). Parity had highly significant effect on 305-day milk yield and significant effect on lactation milk yield. The milk yields were lowest in first lactation and increased with advancement of lactation. Similar results were also reported for Jersey cows maintained in different places in India and Pakistan (Chauhan, 1990; Murdia and Tripathi, 1992; Ahmed et al., 2004; Venkataramananet al., 2007). In general, the low productivity of *Bostaurus* in the tropics is attributable to the interaction of a complex group of factors. Among these are the high atmospheric temperature and relative humidity, low content of essential nutrients in feeds, wide seasonal fluctuations in quantity and quality of feed and high incidence of disease (Mahadevan, 1966)

The least-squares means (± SE) for service period, calving interval and dry period are presented in Table 2. The service period observed was higher than the averages reported for Jersey cows by Jain et al. (1999) in India and Sattaret al. (2004) in Pakistan. Venkataramananet al. (2007) reported that the calving interval of Jersey cows maintained at Nucleus Jersey and Stud Farm, located in highlands in Tamil Nadu as 416.5 ± 7.2 days and many other studies (Chauhan, 1990; Das et al., 1990; Sattaret al., 2004) also indicated lower calving interval than that observed in the present study. However, comparable values were reported in India by Karet al. (1987) and Jain et al. (2001). Gogoiet al. (1993) reported higher value of more than 530 days for Jersey cattle maintained in Assam, India and indicated the possible cause as the hot and humid climatic conditions. The dry period pooled over parities observed was higher than the value reported by Duc and Taneja (1984). The highly significant effect of period of calving on calving interval observed in the present study corroborated with the reports of Chauhan (1990) and Venkataramananet al. (2007). Some of the studies (Das et al., 1990; Venkataramananet al., 2007) including the present one revealed that the mean calving interval for the first parity was higher than the records pooled over all parities.

The production performance of farmbred Jersey at Eachenkottai was generally distinctly lower than those observed in other parts of India. But the reproduction performance observed is comparable to earlier studies made in bull mother farms located at different places in India. Period and season of calving had highly significant effect on all the production traits studied, which indicated that the animals exposed to different environments during different years and seasons and this might be one of the reasons for poor production performance of the herd.

## REFERENCES

- Ahmed, B., Khan, S., Manna, Aand Abdullah. (2007). Production and reproduction performance of Jersey cattle at Cattle Breeding and Dairy Farm, HarichandCharsaddaNWFP. *Journal* of Agriculture Biological Science, **2**:1-5.
- Chauhan, V.P.S. (1990). Production and reproduction performance of Jersey cows in India. *Indian Journal of Animal Genetics and Breeding*, **12**:18-20.
- Das, G.C., Das, D and Aziz, A. (1990).Genetic and non-genetic factors affecting some of the reproductive traits of Jersey cows.*Dairy Guide*, **12**:25-28.
- Duc, N.V and Taneja, V.K. (1984). Performance of purebred and *crossbred grades for some reproductive traits in India.Indian* Journal of Animal Sciences, **54**:141-144.
- Gogoi, D.N., Goswami, R.N and Das, D. (1993). First lactation performance of Jersey, Red Sindhi and their Fl cross under the farm conditions of Assam. *Indian Journal of Animal Sciences*, **63**:569-572.
- Harvey, W.R. (1990). User's Guide for LSMLMW and MIXMDL PC-2 Version.Mixed model least-squares and maximum likelihood computer program.Ohio State University, Columbus, Ohio, U.S.A.
- Jain, J.K., Khan, F.H and Singh, A. (2001).Factors affecting calving interval in Jersey.*Indian Veterinary Journal*, **78**:444.
- Jain, J.K., Khan, H and Saha, D.N. (1999). Studies on first service period in Jersey cows.*Indian Veterinary Journal*, **76**:230-232.

- Kar, B.K., Mohanty, A and Mishra, M. (1987). Relative economic efficiency of Jersey, its crosses with Red Sindhi and Hariana and Red Sindhi cows and Murrah buffaloes. *Indian Journal of Animal Production and Management*, 3:72-77.
- Mahadevan, P. (1966). Breeding for Milk Production in Tropical Cattle. Technical Communication No.17 of the Commonwealth Bureau of Animal Breeding and Genetics, Edinburgh, Commonwealth Agricultural Bureaux, England. 154 pp
- Murdia, C.K and Tripathi, V.N. (1992).Effect of farm, period, season and parity on performance traits of Jersey cattle.*Indian Journal of Animal Sciences*, **62**: 77-180.
- Pal, V., Barhat, N.K and Beniwal, B.K. (2002). Performance of imported Danish and Australian Jersey cattle and their farm bred progenies in Rajasthan. SARAS Journal of Livestock and Poultry Production, 18:34-40.

- Sadana, D.K and Tripathi, V.N. (1986).Comparative performance of the exotic and locally born Jersey cattle.*Indian Dairyman*, **38**:360.
- Sattar, A., Mirza, R.H and Ahmed, I. (2004). Reproductive efficiency of Jersey cows under subtropical conditions of Punjab. *Pakistan Veterinary Journal*,**24**:129-133.
- Singh, A.S and Mishra, M. (1980). Physiological responses and economic traits of Holstein, Jersey, crossbred and Hariana cows in hot and humid environment. *Indian Journal of Dairy Science*, **33**:174-181.
- Sreemannarayana, O and Rao, A.V.N. (1994). Effect of weight at first calving on productive and reproductive traits in Jersey cows. *Indian Veterinary Journal*, 71:514-515.
- Venkataramanan, R., Panneerselvam, S and Kandasamy, N. (2007).Non-genetic factors affecting production and reproduction traits of Jersey cattle at highlands in South India. *Indian Journal of Dairy Science*,60:295-299.

## Table 1.

T.C. at		305-day milk yield	Lactation length	Lactation milk yield	
Effect	n	( <b>kg</b> )	(days)	( <b>kg</b> )	
Overall mean (µ)	1172	$1491.6 \pm 25.9$	$303.1 \pm 4.0$	$1560.9 \pm 29.9$	
Period of calving		**	**	**	
P <sub>1</sub> (1980-84)	197	1729.5 <sup>b</sup> ± 35.7	331.8°± 5.5	1890.7 <sup>b</sup> ± 41.2	
P, (1985-89)	468	1451.7ª± 22.3	$315.4^{bcd} \pm 3.5$	1494.9ª± 25.7	
P <sub>3</sub> (1990-94)	418	1538.1ª± 23.0	$304.2^{bd} \pm 3.6$	$1579.2^{a} \pm 26.5$	
$P_{4}(1995-99)$	69	$1400.6^{a} \pm 54.5$	$264.2^{a} \pm 8.5$	1415.8ª± 63.0	
P <sub>5</sub> (2000-04)	20	1338.3 <sup>a</sup> ± 98.3	300.4 <sup>ad</sup> ± 15.3	$1423.8^{a} \pm 113.5$	
Season of calving		**	**	**	
Winter (Jan. – Feb.)	156	1566.5 <sup>b</sup> ± 42.6	313.2 <sup>b</sup> ± 6.6	1629.8 <sup>bc</sup> ± 49.2	
Summer (Mar May)	305	1383.2ª± 32.3	289.7ª± 5.0	1453.7ª± 37.3	
South-west monsoon	329	$1447.0^{a} \pm 32.7$	$301.6^{ab} \pm 5.1$	$1523.2^{ac} \pm 37.8$	
(Jun. – Sep.)					
North-east monsoon	382	1570.0 <sup>b</sup> ± 32.0	$308.2^{b} \pm 5.0$	1636.9 <sup>b</sup> ± 36.9	
(OctDec.)					
Parity		**		*	
First	373	1396.8 <sup>a</sup> ± 30.0	$299.1 \pm 4.7$	1469.5 <sup>a</sup> ± 35.0	
Second	284	1448.7 <sup>a</sup> ± 33.5	$304.0 \pm 5.2$	1511.2 <sup>ab</sup> ± 38.7	
Third	210	1496.5 <sup>a</sup> ± 36.7	$311.4 \pm 5.7$	1563.6 <sup>b</sup> ± 42.4	
Fourth	144	$1511.3^{a} \pm 42.3$	$303.6 \pm 6.6$	1597.2 <sup>b</sup> ± 48.8	
Fifth	87	$1537.7^{ab} \pm 52.4$	$298.2 \pm 8.1$	1584.5 <sup>b</sup> ± 60.5	
Sixth and above	74	1558.8 <sup>b</sup> ± 55.5	$302.9 \pm 8.6$	1639.3 <sup>b</sup> ± 64.1	

## Least-squares means $(\pm SE)$ for different milk production traits of farmbred Jersey cows

n= number of observations. \* P<0.05, \*\* P<0.01.

Means bearing same superscript do not differ significantly.

## Table 2.

Effect	n	Service period	n	Calving interval	n	Dry period
Overall mean (µ)	829	177.0 ± 10.0	878	461.0 ± 9.7	861	$160.2 \pm 9.7$
Period of calving		**		**		**
P <sub>1</sub> (1980-84)	156	170.3 <sup>ab</sup> ± 11.4	178	459.8 <sup>bc</sup> ± 11.2	168	125.9ª± 11.5
P <sub>2</sub> (1985-89)	355	169.1ª± 7.3	372	$444.6^{ac} \pm 7.4$	366	129.8ª± 7.5
P <sub>3</sub> (1990-94)	278	145.3ª± 7.9	286	422.2ª± 8.1	283	117.7ª± 8.2
P <sub>4</sub> (1995-99) 30		235.9 <sup>b</sup> ± 22.8	30	515.2 <sup>bd</sup> ± 23.5	32	261.7 <sup>bc</sup> ± 22.9
P <sub>5</sub> (2000-04)	(2000-04) 10 164.4 <sup>a</sup> ± 38.9 12 463.4 <sup>ac</sup>		$463.4^{acd} \pm 36.6$	12	$165.8^{ac} \pm 36.8$	
Season of calving				*		
Winter (Jan. – Feb.)	106	$199.3 \pm 15.3$	111	487.2 <sup>b</sup> ± 15.1	113	177.7 ± 15.1
Summer (Mar May)	208	$174.8 \pm 11.8$	218	$460.4^{ab} \pm 11.7$	216	167.1 ± 11.7
South-west monsoon (Jun. – Sep.)	235	170.1 ± 11.9	253	451.1ª± 11.7	243	151.9 ± 11.8
North-east monsoon (Oct Dec.)	280	163.8 ± 11.9	296	$445.5^{a} \pm 11.6$	289	144.1 ± 11.7
Parity		**		**		**
First	260	$208.0^{bd} \pm 11.4$	288	496.5 <sup>b</sup> ± 11.0	283	$201.6^{b} \pm 11.0$
Second	205	$203.7^{\rm bc} \pm 12.0$	215	$486.0^{ab} \pm 11.8$	210	$184.6^{bc} \pm 11.9$
Third	155	$183.8^{ab} \pm 13.3$	159	$466.9^{ab} \pm 13.1$	157	$157.6^{ac} \pm 13.2$
Fourth	102	$164.2^{ac} \pm 15.1$	106	450.2ª± 15.0	104	143.5ª± 15.2
Fifth	60	157.3 <sup>acd</sup> ± 17.8	61	437.9ª± 18.0	60	136.4ª± 18.2
Sixth and above	47	145.0ª± 20.0	49	428.7ª± 20.1	47	137.9ª± 20.5

## Least-squares means $(\pm SE)$ for different reproduction traits (days) of farmbred Jersey cows

n= number of observations. \* P<0.05, \*\* P<0.01.

Means bearing same superscript do not differ significantly

## Adoption and Perceived Effectiveness of Traditional practices to mitigate human-wild pig conflict situations

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## ABSTRACT

Human-wild pig conflict is one of the main threats to farmers as wild pig destroys the growing crops completely. Krishnagiri district of Tamil Nadu was purposefully selected for the study as it ranked first in human-wild pig conflict incidents in Tamil Nadu. Farmers who had at least one wildlife conflict incidence in their lifetime were selected for this study during 2015-16 on adoption and effectiveness of traditional practices to manage human-wild pig conflict situations. Sixty participants were selected using the snow ball sampling and data were collected using semistructured interviews, complemented by free listing techniques, nonspecific prompting and reading back. A total number of nine traditional practices were identified and found that they were adopted at various levels. Further, boundary clearing, using metal cow bells and using shining tapes (100 %) were found to be most effective traditional method followed by fireworks/ crackers (75 %), noise making (72.2 %) and scarecrows (69.6 %). Although encouraging, these results require more widespread testing and demonstration to ensure their effectiveness at broader scales.

Keywords:Human-pig conflict, traditional practices, metal cow bells, shining tapes, scarecrows, fireworks

## INTRODUCTION

Wild pig (*Susscrofa*) is the most widely distributed large mammal and distributed in North Africa, Europe and Asia (Rao*et al.*, 2015). The wild pig population is increasing drastically not only due to its prolificacy in breeding but also depletion in the head counts of its predators such as

tiger, leopard, wild dogs, wolf and jackals (Khan *et al.*, 2019). Overabundances of wild pig population, scarcity of food and shrinking natural habitats have compelled the wild pigs to intrude the forest fringe villages and farms. Cutting of forest trees for

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farming and industrial activities often accelerate the humans and wild animals conflicts.

The farmers in such areas have to bearconsiderable economic loss (Fig.1).Therefore, the farmers resort to different traditional methods to mitigate the intrusion of wild pigs into their farming land. Hence a study was carried outto identify the traditional practices followed by the farmers and also find adoption and perceived effectiveness.

#### MATERIALS AND METHODS

Among the 38 districts of Tamil Nadu state, this study on adoption and perceived effectiveness of traditional practices to mitigate human-wild pig conflict situations was purposively carried out in Krishnagiri district of Tamil Nadu state during 2015-2016 due to the high incidence of human-wild pig (210 incidents during 2014-15) conflict on the basis of data from Tamil Nadu Forest Department. From the four forest zone blocks of Krishnagiri district, two blocks viz., Denkanikottai and Thally were randomly selected. From these two blocks, six villages were also chosen randomly. Sixty respondents (ten farmers per village) were selected using the snowball technique and data were collected using semi structured interview schedule, complemented by free listing techniques, nonspecific prompting, and reading back. The data was analysed using average and percentage.

## **RESULTS AND DISCUSSION**

A total number of nine traditional practices were identified that include noise-making, fire/ smoke producing, manual guarding, traditional fences, scarecrows (Fig.2), boundary clearing, fireworks/crackers, metal cow bells and shining tapes. The results in Table 1 revealed that among the nine chosen traditional methods, fireworks/ crackers (66.7 %) ranked first in the adoption of traditional practice to mitigate human-wild pig conflict.

About 63 per cent of the respondents of the study area were manually guarding the crop fields during the harvest time to drive away the intruding wild pig. Similar strategy was followed by the farmers of Mazowe District of Mashonal and Central Province, Zimbabwe (Nyika, 2017). During certain period of high crop vulnerability, mostly male members of the family would take the turns to guard the field crops. They used the different methods to cope with human-wild pig conflict. Farmers tied the metal cow bells and hung them at fence to act as an alarm if a wild pig tried to break through the perimeter fencing. Similar technique was recorded by Graham and Ochieng (2008) to mitigate the human-elephant conflict by the farmers in Laikipia District, Kenya.

# Table 1. Adoption and perceived effectiveness of traditional practices to mitigate human-wild pig conflict situations

n=60

S.	Traditional	Ado	pters	Non-ad	opters	Effe	ctive	Inef	fective
No.	practices	Frequency	Percentage	Frequency	Percentage	Frequency	Percentage	Frequency	Percentage
1	Noise- making	36	60.0	24	40.0	26	72.2	10	27.8
2	Fire/Smoke	22	36.7	38	63.3	9	40.9	13	59.1

Adoption and perceived effectiveness of traditional practices to mitigate human-wild pig conflict situations

3	Manual Guarding	38	63.3	22	36.7	22	57.9	16	42.1
4	Traditional fences	4	6.7	56	93.3	-	-	4	100.0
5	Scarecrows	23	38.3	37	61.7	16	69.6	7	30.4
6	Boundary Clearing	4	6.7	56	93.3	4	100.0	-	-
7	Fireworks/ Crackers	40	66.7	20	33.3	30	75.0	10	25.0
8	Metal cow bells	4	6.7	56	93.3	4	100.0	-	-
9	Shining tapes	6	10.0	54	90.0	6	100.0	-	-

Use of shining tapes like video/audio tapes had been observed as well established fact that wild pigs were afraid of shining materials which reflect light from far away. Due to this reason, farmers tie audio and video tapes roll all around the crop fields with wooden sticks. During daytime, the tapes glare due to scorching sunlight and in night, they reflect in dark that made the wild pigs frightened to keep them away from the agricultural fields. This is in accordance with the findings of Meena et al. (2014) who proved the shining tapes were very successful technique to frighten the blue bulls in Rajsamand district, Rajasthan, India. Moreover, there is a need to develop appropriate extension mix for disseminating the locally relevant ITK through vernacular language targeting the resource poor farm families (Ponnusamy et al., 2017).

Even though using fireworks/crackers ranked first, only 75 per cent of the farmers perceived it to be effective. But cent per cent efficacy was recorded by the farmers practicing boundary clearing, metal cow bells, and shining tapes though these were adopted by 6.7 per cent and 10 per cent farmers respectively. The farmers created noises by various means to frighten the intruding wild pig herd. Some of the farmers also released dogs during encounter with wild animals. Raoet al. (2015) used castor (*Ricinuscommunis*) as a barrier to reduce the damage caused by wild pig conducted at Agricultural Research Station, Tandur (Telangana state) in the Maize (*Zea mays*) crop. In addition to this, Naik and Basavadarshan (2020) used color sarees to drive the intruding wild pigs and found to be effective at Arjunahalli and surrounding villages which were in the vicinity of the forest, KanakapuraTaluk, and Ramanagara District of Karnataka State.

various The losses created by wild animals to the farmers are given in the table 2. This table clearly depicted that majority (48.30 %) of the elephant conflict respondents experienced losses which was below Rs.10,000 followed by 23.30 per cent of them suffered losses between Rs.25,001- 50,000. The major losses created by the wild pigs on the farmers were recorded in Paddy fields followed by Ragi(finger millet). McKeeet al. (2020) estimated crop loss due to feral swine of \$272 million for the surveyed set of crops in 12 states in United States.

S. No.	Losses to crops (Rs.)	Number of farmers (Percentage)
1	Below 10000	29 (48.30%)
2	10001-25000	13 (21.70%)
3	25001-50000	14 (23.30%)
4	Above 50000	4(6.70%)

 Table 2. Distribution of losses to crops of human wild pig conflict affected farmers

(N=60)

## CONCLUSION

Crop depredation by wild pigs is enormous. Hence the farmers use different traditional technologies to mitigate the human-wild pig conflict. The results of the present study indicate that among the different traditional methods used to mitigate Human wild pig conflict, fireworks/ cracker, manual guarding and noise making were highly adopted than rest of the traditional methods. But boundary clearing, metal cow bells and shining tapes were found to be highly effective in reducing the wild boar followed by noise making, fireworks/ crackers and scarecrowsaround the crop was promising. Hence proper educational programme may be conducted among the farmers of affected areato popularize the effective traditional methods to mitigate human wild pig conflict through training. Although encouraging, these results require more widespread testing and demonstration to ensure their effectiveness at broader scales.

## REFERENCES

- Khan, S., Gupta, S., Ilyas, O., Roy, A and Haleem,
  A. (2019).Evaluation of suitable habitat for wild boar (*Susscorfa*) in Pench Tiger Reserve,
  Madhya Pradesh, India.*International Journal* of Ecology and EnvironmentalSciences,
  45(2):157-164.
- Graham, M. D and Ochieng, T. (2008). Uptake and performance of farm-based measures for reducing crop raiding by elephants *Loxodontaafricana* among smallholder farms in Laikipia District, Kenya. *Oryx*, 42(1):76-82.

- McKee, S., Anderson, A., Carlisle, K and Shwiff, S. A. (2020). Economic estimates of invasive wild pig damage to crops in 12 US states. *Crop Protection*, **132**(6) 105105.
- Meena, R. P., Meena, B. L., Nandal, U and Meena, C. L. (2014) Indigenous measures developed by farmers to curb the menace of blue bull (*Boselaphustragocamelus*) in district Rajsamand, Rajasthan, India. *Indian Journal of Traditional Knowledge*, **13**(1):208-215.
- Naik, M. I and Basavadarshan, A. V. (2020). Incidence and efficacy of crop protection measures against wild boar (Susscrofa L.) in groundnut (Arachishypogaea L.).Journal of Entomology and Zoology Studies, 8(3):1616-1620
- Nyika, E. (2017). The human-wildlife conflict: a survey of attitudes of local farmers towards crop raiding baboons in concession, Mazowe District of Mashonaland Central Province, Zimbabwe (Doctoral dissertation, BUSE).
- Ponnusamy, K., Kale, R.B., Ravi, K.N., Devi.M.C.A and Sharma, P. (2017).Cross-regional analysis on usage of Indigenous Technical Knowledge in dairy farming.*Indian Journal of animal Research*, **51**(3):549-556.
- Rao, V. V., Naresh, B., Reddy, V. R., Sudhakar, C., Venkateswarlu, P and Rao, D. R. (2015). Traditional management methods used to minimize wild boar (*Susscrofa*) damage in different agricultural crops at Telangana state, India. *International Journal of Multidisciplinary Research and Development*, 2(2):32-36.

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Adoption and perceived effectiveness of traditional practices to mitigate human-wild pig conflict situations



Fig.1 Wild pig Trampled field



Fig.2 Scarecrows to drive out intruding wild pigs

# Energy consumption pattern in manufacturing of different types of broiler feeds

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#### ABSTRACT

The present study was carried out to determine the specific energy consumption pattern in the manufacturing of different type of broiler feeds with different process conditions and thereby analyze the gaps between the benchmarking and actual energy usage pattern. The study was conducted in three commercial feed plants located at different locations. This was achieved by conducting energy analysis for the specific energy consumption for different type of the broiler feeds. The actual consumption was recorded from the electricity board department's energy meter readings. The results indicated that the gap between actual consumption and bench marking was lesser at one plant whilst, it was considerably higher at other two plants. The gap was observed to be higher in the pre-starter feed and lesser in the finisher pellet feed. The gap between actual power consumption and benchmarking observed for pre-starter feeds at different locations viz., Muzaffarpur, Kanpur and Ambala was 2.70, 2.43 and 0.85 kwh units respectively. The gap analysis for starter feed was 1.71, 1.46 and 0.59 kwh units and for finisher feed 1.37, 1.81 and 0.92 kwh units respectively. The cost saving per metric tonne of feed ranged from Rs. 16.00 to Rs. 22.00. Among the three feed manufacturing plants, the gap between bench marking and actual power consumption was very minimal at Ambala plant due to better production efficiency. From the boiler's fuel cost point of view, coal was found to be cheaper than wood due to its high calorific value and hence it was suggested to replace fire wood with coal at Ambala plant for further improvement of fuel efficiency. The present findings also revealed that the timely replacement of press roller of pellet mill and reduction in pellet fines recycling leads to better efficiency and cost control.

Key words: energy efficiency, boiler fuel, coal, wood, fuel efficiency

#### **INTRODUCTION**

Feed milling industry is one of the most energy consuming industries in the world. Like capital, labor and materials, energy is also one of the major production factors which are used to produce final product. In economical terms, energy is demand-derived goods and can be regarded as intermediate good whose demand depends on the demand of final product. Electricity is the main energy source for these feed mills which is imported from the state electricity board grids. Electricity is

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used to run motors, pumps, blowers, conveyors, fans, lights, etc. The variations in the consumption rate of energy through the use of utilities during processing must also be accounted for final cost of the finished product. The feed milling consumes significant quantity of fuels and electricity. The major energy consuming equipments in the feed milling units are; pellet mill, hammer mill, boiler and steam distribution units, blowers, pumps, conveyers, elevators. motors. transmission systems, weighing, etc. Energy, its utilization and conservation, may be the greatest challenge facing management in the feed industry today and in the years ahead. All forms of energy are increasing in cost and diminishing in supply; yet effective energy management, or conservation, can reduce the cost impact and make better use of available energy supplies. As energy is used more effectively, product costs can be reduced and profits improved.

Energy is one of the most critical input resources in the manufacturing industries. In most cases, energy cost outweighs the costs of other resources such as raw material, labors, depreciation and maintenance (Fadare, 2003). Energy is one of the most important material bases for the economic growth and social development of a country or region. Scientific forecasts and analysis of energy consumption will be of great importance for the planning of energy strategies and policies. Nowadays, energy usage in agriculture has been intensified in response to continued growth of human population, tendency for an overall improved standard of living and limited supply of arable land; thus, the farmers use their inputs in excess and inefficiently, particularly when the inputs have low price or are available in plenty. The enhancement of energy efficiency not only helps in improving competitiveness through cost reduction, but also results in minimized energyrelated environmental pollution, thus positively contributing towards sustainable development (Kizilaslan, 2009 and Ghorbaniet, al., 2010).

Improvement of energy efficiency in our facility improves the bottom line; increases productivity and market competitiveness; lessens the impact and protects the business from fluctuations in energy prices and reduces carbon emission and stay ahead of government regulations.

Though, wide variety of technologies has been evolved for efficient use of energy for various equipments of feed mills, so far, only a few have improved their energy efficiency levels. Most of the feed mills use old and locally available technologies and are also completely dependent on locally available technical personnel. In this pretext the present investigation was carried out at three different commercial feed mills located at Muzaffarpur (Bihar), Ambala (Harvana) and Kanpur (UP) to determine energy consumption pattern during different stages of feed manufacturing process for each type of feed viz., pre-starter, starter and finisher feed with different type boilers and work out the economic impact of energy utilization on feed production.

## MATERIALS AND METHODS

The determination of the energy consumption in manufacturing of different types of broiler feeds (crumbs and pellets), was the main target of this study. This was achieved by determining the energy consumed in each stage of processing to assess the most consumable stage in the different types of feed. The bench marking study is done for specific energy consumptions of major sections in the feed milling process *viz.*, intake, batching & grinding, pelleting and boiler and other utilities.

Energy benchmarking for industry is a process in which the energy performance of an individual plant or an entire sector of similar plants is compared against a common metric that represents 'standard' or 'optimal' performance. As a benchmarking standard "Specific Energy Consumption" energy analysis was conducted by measuring running amps of the individual motor load. Load factor of each motor is calculated. From the input Volts and measured current (amps), individual motor power consumption unit has been calculated. The benchmarking energy analysis was studied for all of three plants for different types of feeds of prestarter, starter and finisher feeds. Actual energy consumption data was taken from the electricity meter installed in the feed mill.

## Methodology Used

Electrical power consumption was estimated from the measured electric current and voltage values and estimated according to Kurt (1979)as follows from equation:

The amount electricity (kwh) used to produce one tonne of feed (kwh/T) is calculated as follows as specific energy consumption of the individual motor by using the following formula :

Downer (lowb)	_	A x voltage x $\sqrt{3}$ x power factor	MT/ Hour
Power (Kwii)	= -	1000	MIT/ Hour
	(p	ower factor has been taken as 0.95)	

Specific energy consumption in terms of kwh  $\setminus$  MT of feed was calculated for Pre-Starter, Starter and Finisher feed and the benchmarking data has been arrived under standard process conditions. The specific energy consumption bench mark data was compared with the actual energy consumption data which were being recorded from energy meter installed by the electricity department.

## Method of Motor Load measurement

To estimate the motor load factor power measurement was carried out in induction motor as a part of work during the detailed energy studies. The power measurements were used to calculate the actual load factor of induction motor at site conditions. The power measurements were done with the help of a Class 1 accuracy clamp-on power meter.

Actual power consumption was arrived from the data collected from the EB meter installed at the respective plants and it was calculated as per MT of prestarter, starter and finisher feed.

### **Boiler Fuel Consumption**

Coal is being used for the boilers located at Muzaffarpur and Kanpur and wood is used at the Ambala plant's boiler unit. Fuel usage per MT of feed was calculated from the daily consumption record.

		Total Consumption in
Fuel Consumption	_	Kgs
per MT of feed	=	Qty of feed produced
1		in MT

### Table 1. Technical specifications of boiler unit at three feed plants

S No	Dontioulon	Details							
5.110.	rarucular	Muzaffarpur	Kanpur	Ambala					
1	Firm Name	Ugraya foods and feeds	Prime avian feeds pvt.	Ugraya foods and					
		pvt. Ltd.	Ltd.	feeds pvt. Ltd.					
2	Type of Boiler	Horizontal Multi Tubular	Horizontal Multi Tubular	Horizontal Multi					
		Fully Wet back Three pass	Fully Wet back Three	Tubular Fully Wet					
		smoke tube Boiler	pass smoke tube Boiler	back Three pass					
				smoke tube Boiler					

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3	Design,	IBR 1950 with latest	IBR 1950 with latest	IBR 1950 with
	Fabrication	Amendments. inspection	Amendments. inspection	latest Amendments.
		& testing code	& testing code	inspection & testing
				code
4	Model	IFB-20D -THERMAX	IFB-10D -THERMAX	IFB-10D
		MAKE	MAKE	-THERMAX MAKE
5	Evaporation	2000 Kg/hr (F&A 100°C)	1000 Kg/hr (F&A 100°C)	1000 Kg/hr (F&A
	Capacity			100°C)
6	Max. Working	10.54 Kg/cm2(g) (Safety	10.54 Kg/cm2(g) (Safety	10.54 Kg/cm2 (g)
	Pressure	valve set off)	valve set off)	(Safety valve set off)
7	Efficiency	72%	72%	72%
8	Fuel	Coal	Coal	wood
9	Gross Calorific	3500 kCal/kg	4500 kcal/kg	4500 kCal/kg
	Value			
10	Mode of	Mechanical Draught	Mechanical Draught	Mechanical Draught
	combustion			
	Type of feeding	Manually through fire	Manually through fire	Manually through
		door	door	fire door

## **RESULTS AND DISCUSSION**

The process condition under which energy consumption study was conducted is given in Table 2. The pelleting process conditions are different from one another at each plant. Conditioner design is different from one another at all of the three plants and the moisture addition is maintained in the range of 2.0- 2.5% by adjusting steam flow valves. Different hole diameter and thickness of the die has been used at all the three plants. It has been observed that there is variance in the electrical consumption at all the three plants.

The comparative details of benchmarking of specific energy consumption analysis per MT of feed of pre-starter crumbs (PSC), starter crumbs (SC) and finisher pellet (FP), produced at muzaffarpur, kanpur and ambala respectively are depicted in Table 3. The specific energy

consumption benchmarking analysis was done for the different processes such as intake, batching and grinding, pelleting, boiler and other utilities. From the individual process study values, total power consumption in kwh units were calculated. The actual power consumption units were calculated from the energy meter's initial and final readings. From the total benchmarking and actual power consumption data, the gaps were identified. The gap between benchmarking and actual observed was more at muzaffarpur plant and lesser gap was observed at ambala plant. The electrical energy utilization efficiency was found to be better at ambala plant and less efficient at muzaffarpur plant, followed by kanpur plant. Overall, high cost impact per MT of feed was observed at Kanpur due to high power cost. The potential cost saving per month by considering average production volume at respective plants has been calculated as Rs. 79,459 at muzaffarpur, Rs. 21,091 at kanpur and Rs. 17,158 at ambala.

C No	Dellating Verichlag	Plant				
5.INO	reneting variables	Ambala	Kanpur	Muzaffarpur		
1	Ground Particle Size passing on 0.71mm (%)	65-70	65-70	65-70		
2	Moisture Addition at Conditioner (%)	2.0	2.5	2.0		
3	Added Fat (%)	0.5 -0.9	0.5 -0.9	0.5 -0.9		
4	Steam Pipe Size from Boiler (NB)	40	40	50		
5	Die Hole diameter (mm)	3.2	2.8	3		
5	Pellet Die L/D Ratio	01:10.5	1:12	1:13		
6	Die Material	Chrome	Chrome	Chrome		
7	Retention Time at Conditioner ( Seconds )	10	15	30		
8	Conditioner Type	Two Stage	Single	Two Stage		
9	Moisture Addition at the Conditioner (%)	2.00-2.5	1.8-2.2	2.0- 2.5		
10	Conditioner Length (mt)	2.0	3.0	2.5		
11	Retention time (Seconds)	20	10	30		
12	Steam Pressure (Kg/cm2) at PRV	2.5	2.5	2.5		

## Table 2. Process design Variables

The boiler fuel consumption pattern is presented in Table 4. Fuel consumption quantity per MT of feed was calculated from the boiler operators log book. The fuel consumption usage per MT of feed was arrived. Coal was used as a fuel at two boilers at Muzaffarpur and Kanpur and fire wood was used at Ambala. From the fuel consumption volume and actual fuel price, the fuel cost per MT of feed was calculated as Rs 94.15 at Ambala, Rs 73.98 at Kanpur and Rs 88.57 at Muzaffarpur. Coal has been identified as cheaper source of fuel as compared to firewood and up to Rs 20 can be saved per MT of feed, if coal is used as a fuel.

Table 3	3. Data	on	boiler	design	and	operation	for	three	different	feed	nlants
Table .	. Data	on	DOLLET	ucoign	anu	operation	101	unice	uniciciit	iccu	plants

C No.	Description	Plant Locations				
5.110	Description	Ambala	Kanpur	Muzaffarpur		
1	Make	Thermax	Thermax	Thermax		
2	Model. No	IFB-10D	IFB-10D	IFB-20D		
3	Efficiency (%)	73	73	78		
3	No of Passes	Three	Three	Three		
4	Steam Generation Capacity (Kgs/Hr)	1000	1000	2000		
5	Year of installation	2012	2014	2013		
6	Design Pressure (Kg/cm2)	10.54	10.54	10.54		
7	Max Working Steam Pressure (Kg/cm2)	10.5	10.5	10.5		

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8	Operational Pressure (Kgs/cm2)	8.5	9	8.5
6	ID (Induced Draft Fan)	Available	Available	Available
7	FD (Forced Draft Fan	Available	Available	Available
8	Water Pre heater	Available	Available	Available
10	Fuel Type Used	Wood	Coal	Coal
11	Net Calorific Value (Kcal /kg)	3500	4500	4500
12	Production per hour (MT)	10	10	20
13	Fuel Consumption (Kgs) /Hr	207.13	82.2	208.4
14	Fuel Consumption (Kgs) / MT of feed	18.83	8.22	10.42
15	Fuel Cost per kg ( Rs )	5	9	8.5
16	Fuel cost per MT of feed (Rs)	94.15	73.98	88.57

# Table 4. Data on Electricity Consumption Analysis - Benchmarking of specific energy consumption Vs Actual Consumption Units

S NO	Dontioulous	Muzaffarpur			KANPUR			Ambala		
5. NU	raruculars	PSC	SC	FP	PSC	SC	FP	PSC	SC	FP
	Intake Bench Marking Units									
1	(kwh)	1.75	1.75	1.75	1.36	1.36	1.36	1.11	1.11	1.11
	Batching & Grinding Bench									
2	marking units (kwh)	9.25	9.25	9.25	7.23	7.23	7.23	7.68	7.68	7.68
	Pelleting bench marking units									
3	(kwh)	20.14	18.13	15.47	20.54	20.5	17.46	18.2	16.5	15.17
	Boiler & other utilities									
4	benchmarking units (kwh)	1.16	1.16	1.16	1.44	1.44	1.44	1.12	1.12	1.12
	Total Benchmarking Units									
5	(kwh)	32.3	30.29	27.63	30.57	30.5	27.49	28.2	26.4	25.1
	Actual Power Consumption									
6	Units (kwh)	35	32	29	33	32	29.3	29	27	26
	Gap between standard and									
7	Actual (kwh)	2.7	1.71	1.37	2.43	1.46	1.81	0.85	0.59	0.92
8	KWH Cost (Rs) /Unit	6.15	6.15	6.15	9.06	9.06	9.06	8.37	8.37	8.37
	Cost impact per MT of feed									
9	(Rs)	16.61	10.52	8.43	22.02	13.23	16.4	7.11	4.94	7.7
	Average Production Qty									
10	.Month (MTs)	1090	4048	2230	162	762	454	296	946	1348
11	Cost Impact per month (Rs)	18099	42571	18789	3567.2	10081	7445	2106	4672	10380
	Total potential Cost Savings /									
12	Month (Rs)		79459			21091			17158	3

PSC - Prestarter crumbles; SC - Starter crumbles; FP - Finisher pellets

The actual power consumption units (kwh) for pre-starter feed were 35, 33 and 29 at Muzaffarpur, Kanpur and Ambala plants respectively which is of 2.7, 2.45 and 0.85 more than bench marked specific energy consumption units. For pre-starter feed, power consumption was high at muzaffarpur plant and lower at ambala plant. For Starter feed, the actual power consumption were 32,32 and 27kwh, which is of 1.71, 1.46 and 0.59 units more than the bench marked specific energy consumption units at muzaffarpur, kanpur and ambala plants respectively. For finisher feed, the actual power consumption were 29, 29.3 and 26kwh, which is of 1.37, 1.81 and 0.92 units more than the bench marked specific energy consumption units at muzaffarpur, kanpur and ambala plants respectively. From the boiler's fuel consumption study, coal consumption was 8.22 Kg and 10.42 kg/MT of feed and fuel cost was Rs 73.98 and Rs 88.57/MT for Kanpur and Muzaffarpur boilers respectively. Fire wood was used at Ambala boiler plant. The fuel consumption per MT of feed was 18.83 kg and cost per MT of feed was Rs. 94.15.

Kilborn et. al.(1982) found that the total specific milling energy ranged from 46 kJ·kg<sup>-1</sup> for soft wheat cultivars to 124 kJ·kg<sup>-1</sup> for durum wheat. Dziki, (2008)reported that the specific grinding energy of uncrushed kernels ranged from 72.3 to 146.7 kJ·kg<sup>-1</sup> and from 67.0 to 114.4 kJ·kg<sup>-1</sup> for Turnia and Slade, respectively. The crushing caused a decrease of specific grinding energy in both cultivars. The total specific grinding energy of crushed kernels (the sum of crushing energy and grinding energy) ranged from 47.6 to 100.5 kJ·kg<sup>-1</sup> and from 44.6 to 85.3 kJ·kg<sup>-1</sup> for hard and soft wheat, respectively. Dziki, (2008)presented the results concerning the influence of grain mechanical properties on wheat grinding energy requirements. The investigations were carried out on 10 wheat varieties (grain moisture was 15%). The results showed that the specific grinding energy ranged from 22 to 37 kJ.kg<sup>-1</sup>. The grinding efficiency index ranged from 0.215 to 0.342 m<sup>2</sup>.kg<sup>-1</sup>.

Kulig and Laskowski (2005) reported that an increase in fat concentration in feed material from 2 to 5.5% reduces energy consumption during pelleting by 30%. In general, the specific energy required for pelleting (i.e., energy consumed by the pellet mill motor) may range from 4 to 40 kwh/t (Stevens, 1987; Israelsenet. al., 1981 and Tabil et. al., 1997). In addition, steam conditioning/ preheating the feed may require considerable energy. For example, Skoch et. al. (1981) estimated that steam conditioning to increase the temperature from 27 to 80°C consumed about 26 kwh/t. Steam addition in pelleting operations improves pellet durability. Added steam provides heat and moisture and it also helps to reduce energy consumption during pelleting.

There is no known report in the literature on the energy requirements of feed processing operations. Such information is vital so as to enable the management of this industry to develop strategies for better control of their production operations and modify areas of waste. It will also enable the management to properly appraise their energy consumption for effective planning of production network.

From the electric energy consumption study, it has been observed that ambala plant was more energy efficient as the gap between actual power consumption and bench marked data is lower which was highest at muzaffarpur followed by kanpur plant. The present study revealed that the power consumption was higher at muzaffarpur due to lack of efficient pelleting output followed by kanpur plant due to installation of high capacity motor. If the actual energy efficiency is achieved to benchmarking data, the power cost per MT of feed can be saved to the tune of Rs 16.61, 10.52 and 8.43 for Pre-starter, starter and finisher feed respectively at muzaffarpur, Rs 22.02, 13.23 and 4.94 at kanpur and Rs 8.43, 7.11 and 7.7 at ambala plant .The overall cost saving per month can be Rs 79,459, Rs 21,091 and Rs 17,158 at muzaffarpur, kanpur and ambala plants respectively. If energy is used more effectively, product costs can be reduced and profits improved.

#### REFERENCES

- Dabbour, A and Bahnasawy, S Ali. (2014). Energy consumption in manufacturing of different types of feeds . (2nd International Conference on Biotechnology Applications in Agriculture (ICBAA), Benha University, Moshtohor and Hurghada, 8-12, April 2014, Egyp; http://www.bu.edu.eg/portal/uploads/ Agriculture)
- Dziki, D. (2008). The crushing of wheat kernels and its consequences on the grinding process.Powder Technology, 185: 181–186. (https://www.researchgate.net/)
- Fadare, D.A. (2003). Development of an organomineral fertilizer processing plant.Ph.D. thesis of Department of Mechanical Engineering, University of Ibadan. Ibadan, Nigeria. (https://www.academia.edu )
- François Lucas FabricePutier. (2012). Energy benchmarking for the animal feeds sector in France.(https://www.eceee.org/library/ conference\_proceedings/eceee\_Industrial\_ Summer\_Study)
- Ghorbani, R., Mondani, F., Amirmoradi, S., Feizi, H., Khorramdel, S and Teimouri, M. (2010). A case study of energy use and economical analysis of irrigated and dryland wheat production systems. Applied Energy, 88:283–288.
- Gupta, S.,Handa, K.,Saxena, S. K and Bandyopadhya, S. (2000).Assessment of energy conservation potential in petroleum refineries through benchmarking and targeting techniques (http://www.ese.iitb. ac.in/~santanu/Ref\_Benchmark\_152.pdf)

- Indian Animal Feed (Poultry, Cattle & Aquaculture) Market. (2017). Industry Analysis, Size, Share, Growth, Trends and Forecast by 2022 (http://www.digitaljournal.com/ pr/3584516)
- Israelsen, M., Busk, J and Jensen, J. (1981). Pelleting properties of dairy compounds with molasses, alkali-treated straw and other byproducts.Feedstuffs, 7:26–28.
- Jahangeer, K Abdul Halim. (2015). Energy efficiency benchmarking study of food manufacturing plants in Singapore. (http:// www.e2singapore.gov.sg)
- Kilborn, R. H., Black, H. C., Dexter, J. E and Martin D. G. (1982).Energy consumption during flour milling.Description of two measuring systems and influence of wheat hardness on the energy requirements. Cereal Chemistry, 59:284–288.
- Kizilaslan, H. (2009). Input–output energy analysis of cherries production in Tokat Province of Turkey.Applied Energy, 86:1354–8.
- Kulig, R and Laskowski.(2005). Wpływzawartościtłuszczuna process granulowania material łówpaszowych. InżynieriaRolnicza, 7(67):59-68.
- Kurt, G. (1979). Engineering formulas. 3<sup>rd</sup>. Ed. McGraw Hill book Co.
- McEllhiney, R.R. (1980). "The Economics of Feed Manufacturing Processes," In: proceedings of thirty-fifth Kansas Formula Feed Conference, Kansas State University, Manhattan, KS, Jan.
- Metwally, K. A. (2010). Study the effect of some operational Factors on hammer mill. AMSC thesis of Department of Agricultural Engineering, Faculty of Agriculture,

Zagazig University. Egypt. (http://www. mjae.eg.net/pdf/jan2010/4.pdf )

- Mokhtar Ibrahim Dabbour. (2014). Role of feeding rate in energy consumption and mechanical properties for different types of feed pellet (www.researchgate.net/publication /275518848)
- Odigboh, E. U. (1997). Machines for crop production. In: Stout BA, editor. Handbook of agricultural engineering-plant production engineering.American Society of Agricultural Engineers. (http://www.cigr. org/documents/CIGRHandbookVol3.pdf)
- Robinson, Roy, "Pelleting Introduction and General Definitions." In Feed Manufacturing Technology, American Feed Manufacturers Assn., Arlington, VA 1976 (book Feed

manufacturing technology IV Robert R McEllhiney) Published in 1994 in Arlington (Va.) by American feed industry association

- Skoch, E. R., Behnke, K. C., Deyoe, C. W and Binder, S. F. (1981).The effect of steamconditioning rate on the pelleting process. Animal Feed Science and Technology, 6:83–90. (https://www.sciencedirect.com/ science/article/pii/037784018190033X
- Stevens, C. A. (1987). Starch gelatinization and the influence of particle size, steam pressure and die speed on the pelleting process. Ph.D. Dissertation.Kansas State University, Manhattan, KS.
- Tabil, L.G., Sokhansanj, S and Tyler, R.T. (1997). Performance of different binders during alfalfa pelleting. Canadian Agricultural Engineering, 39(1):17-23.

## Fatal secondary septic peritonitis associated with multiple renal and splenic infarcts in Lhasa Apso dog

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#### ABSTRACT

An eight year old male Lhasa Apso dog was presented for necropsy with the history of persistent vomition, anorexia, lethargy, dehydration and abdominal distension. Blood picture revealed marked neutrophilic leukocytosis along with left shift. Necropsy examination revealed the presence of secondary septic peritonitis and multiple renal / splenic infarcts. Histologically, perforative/haemorrhagic enteritis, renal infarcts with diffuse glomerular / tubular necrosis of coagulative type and splenic infarction with occluded artery containing bacterial thrombo-emboli were confirmed. Secondary peritonitis has been observed to be a frequent complication associated with perforation of the intestine. These fatal complications of septic peritonitis present a therapeutic challenge and needs prompt veterinary care and treatment.

key words: Dog, Renal infarcts, splenic infarcts, septic peritonitis

Peritonitis in animals may occur as a primary disorder or may be secondary to other pathophysiological conditions. A variety of agents can cause peritonitis resulting in several clinical symptoms, disease progression and adverse effects. The etiologiesrange from viral to bacterial to parasitic to physical agents and organrupture (Zachary, 2012). Gastrointestinal ulcers account for nearly 24% to 35% of gastro-intestinal tract associated peritonitis (Greenfield and Walshaw, 1987).Peritonitis can be classified as acute or chronic depending on the duration; septic or non-septic depending on the infectious nature of

the causative agent; local or diffuse depending on the area involved and adhesive or exudative depending on the type of inflammatory product. The incidence of primary peritonitis is lesser than that of secondary peritonitis and can be infectious or idiopathic. In case of primary peritonitis caused by pathogenic organisms, the organisms spread *via* blood to the peritoneal cavity.

Primary peritonitis is rare in healthy dogs and is often observed in immune compromised animals. In case of secondary peritonitis, the peritoneal cavity becomes contaminated

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with infectious agents via translocation or penetration through the wound or leakage from the gastrointestinal tract (most common cause) or through leakage from other infected visceral organs (eg, hepatic/ splenic abscess, inflamed bladder/ uterus, etc.). Pathogenic organisms frequently isolated are Nocardia, Mycobacterium, Haemophilusparasuis, etc. Perforations in the abdomen or wound dehiscence can cause laceration of the visceral organs, followed by translocation of extraneous substances and infectious agents into the peritoneal cavity. Migration of endoparasites can also cause leakage of the ingesta leading to septic peritonitis. The gross and microscopic features of a case of fatal secondary septic peritonitis in a LhasoApso dog along with splenic and renal infarcts as its complicationare described in the present communication.

An eight year old male Lhasa Apsodog was admitted with the history of persistent vomition, anorexia and lethargy for one week. The dog had injected mucous membranes, sunken eyes with marked dehydration and exhibited abdominal distension and localised pain on cranial abdominal palpation. Blood picture revealed marked leukocytosis with neutrophilia and shift to left. The animal collapsed during clinical examination.

A detailed post-mortem examination was conducted and representative tissue samples from liver, lungs, kidneys, spleen, stomach and intestines were collected in 10% neutral buffered formalin. The tissue samples were processed for histopathological studies employing paraffin embedding technique. Sections of 3-5mm thick were prepared and stained with hematoxylin and eosin (H&E) following standard procedures.

At necropsy, the abdominal cavity contained abundance of sero-sanguinous fluid, fibrin deposits and undigested feed contents (Fig.1). The serosal surface of the intestine and the mesentery showed multifocal, randomly distributed

areas of marked congestion and haemorrhage. Three perforations measuring around 1-2 cm diameter with irregular haemorrhagic borders were present in the duodenal segment (Fig.2). Kidneys were moderately shrunken and had multiple irregular depressed areas with fibrin deposits. The cut surface appeared diffusely congested and had multiple, well demarcated, yellowish white wedge shaped areas of necrosis surrounded by thin zones of congestion (Fig.3). The broader end of spleen appeared markedly enlarged, oedematous, congested and haemorrhagic with discrete red black wedge shaped foci filled with blood and adherent fibrin deposits (Fig.4). Cut surface of spleen showed well demarcated zones of necrosis and haemorrhage.

histopathological On examination, intestinal segment adjacent to the site of perforation showed marked necrosis and extensive areas of congestion and haemorrhages in the mucosa/ sub-mucosa. The renalinfarcts were characterized by diffuse glomerular/ tubular necrosis with the preservation of architecture (coagulative necrosis) along with the presence of moderateoedema/ infiltrates in the interstitium and the adjacent cortex (Fig. 5).Splenic artery appeared to be occluded with a bacterial embolic thrombus firmly attached to the lumen of the vessel consisting of concentric layers of fibrin, numerous inflammatory cells and RBC's and had clusters of basophilic bacterial colonies (Fig. 6).

Septic peritonitis observed in the present case was due to the presence of 3 perforations in the duodenal segment.Septic peritonitis has been observed to be a frequent complication associated with injuries that affect the gastrointestinal integrity (Ragetly*et al.*, 2011). Perforation of the stomach / intestine leading to leakage of the partially digested feed material as well as rupture of the infected uterus can result in acute secondary septic peritonitis. Agents frequently associated with secondary peritonitis are intestinal foreign bodies (Penninck and Mitchell, 2003); perforations of the stomach, intestines, rectum, urinary bladder, or uterus; gastrotomy or enterotomy; gastric/intestinal neoplasms (eg, mesothelioma); inflammatory / traumatic injuries of liver, gall bladder, pancreas, dilatation / volvulus of gastrointestinal segments, etc.

The dog had persistent vomition, was anorectic and lethargic, prior to death. It had injected mucous membranes, sunken eyes with marked dehydration, abdominal distension and localised pain on cranial abdominal palpation. Animals with secondary peritonitis may also show clinical signs associated with the primary disease. The classical observation through radiography is the absence of abdominal details and there is a focal or generalized "ground-glass" appearance. The GI tract can be dilated with air and/or fluid. Free air inside the abdomen can be present when there is rupture of any luminal organ or at times may be due to gas-producing anaerobic organisms in the absence of organ rupture (Fossum*et al.*, 2009).

The chemical mediators of inflammation produced such as histamine, serotonin, proteases and endotoxinsas a result of the leaked contents result in enhanced capillary permeability and outpouring of plasma proteins, solutes and water into the peritoneal cavity. The resulting inflammation enhances fibrin deposition, particularly in and around the site of injury and peritoneum, as well as the loss of isotonic fluid into the peritoneal cavity with concurrent haemo-concentration (Culpet al., 2009). The fluid produced is clear and transparent (transudate) during the early stages, but becomes turbid as a result of exuded protein, macrophages and polymorphonuclear leukocytes (Kirby., 2003, Culp et al., 2009). In the present case, the abdominal cavity contained abundance of sero-sanguinous fluid and fibrin deposits. The resulting exudate further inhibits microbial clearance and enhances microbial multiplication (Kirby, 2003).

Infarction of different organs has been frequently reported during peritonitis as a result of ischemic injury caused by the vascular obstruction and thromboembolism. Owing to the greater circulating blood supply (20% to 25%), kidneys and spleen are the most frequent targets of thromboembolism. In the present case also, we observed renal and splenic infarcts. The small blood vessels are frequently occluded, i.e., the interlobular arteries and therefore, infarction occurs mostly in the cortical region (Zachary, 2012). In the present case, renal infarcts were seen in the cortex concurring with the previous reports.

Several agents including gastric mucin, bile salts, haemoglobin, etc., are regarded as adjuvants of the inflammation of peritoneum and they accentuate the localized as well as the systemic inflammation (Kirby, 2003). Opsonins, immunoglobulins and complement components get triggered with increasing severity of the inflammation (Schein and Paladugu2000, Culp et al., 2009). In addition, the levels of several cytokines including TNF, IL-1 and IL-6, PGE, and platelet aggregation factor are also increased (Culpet al., 2009). As a result of the production of these chemical mediators, there is a marked decrease in cardiac output, vasodilation and decreased venous return, which further intensify the systemic hypotension and enhance production of endotoxins. The presence of bacteria, the endotoxins, inflammatory cells and their cytokines lead to endothelial damage and tissue factor expression that result in generalized activation of the coagulation cascade, causing thrombosis and fibrinolysis. This results in disseminated intravascular coagulation with the loss of the anti-thrombin III protein into the abdomen. Micro and macro vascular thromboses may lead to tissue hypoxia and organ damage involving the myocardium, lungs, and gastrointestinal tract (DeLaforcadeet al., 2003;Estrinet al., 2006).These pathological changes that occur during septic peritonitis result in systemic vasodilation, enhanced capillary permeability, reduced cardiac function and multiple organ failure, the characteristic signs of systemic inflammatory response syndrome and septic shock (Schein and Paladugu, 2000). In spite of the advanced treatment strategies available, the prognosis of peritonitis is poor.

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#### REFERENCES

Culp, W.T., Zeldis, T.E., Reese, M.S and Drobatz, K.J.(2009). Primary bacterial peritonitis in dogs and cats: 24 cases (1990-2006). *Journal* of American Veterinary Medical Association, **234**(7):906-913.

De Laforcade, A.M., Freeman, L.M and Shaw, S.P. (2003). Hemostatic changes in dogs with naturally occurring sepsis. *Journal of Veterinary Internal Medicine*, **17**(5):674-679.

Estrin, M.A., Wehausen, C.E., Jessen, C.R and Lee, J.A.(2006). Disseminated intravascular coagulation in cats.*Journal of Veterinary Internal Medicine*, **20**(6):1334-1339.

Fossum, O.,Jansson, D.S., Etterlin, P.E.and Vågsholm, I. (2009).Causes of mortality in

laying hens in different housing systems in 2001 to 2004. *ActaVeterinariaScandinavica*, 51 (3), doi:10.1186/1751-0147-51-3.

Greenfield, C.L and Walshaw, R. (1987). Open peritoneal drainage for treatment of contaminated peritoneal cavity and septic peritonitis in dogs and cats:24 cases (1980-1986). *Journal of the American Veterinary Medical Association*, **191**:100-105.

Kirby, B.M. (2003). Peritoneum and peritoneal cavity. In: Slatter D, ed. Textbook of Small Animal Surgery.3<sup>rd</sup>Edn. Elsevier Science, Philadelphia, PA.414-445.

Penninck, D and Mitchell, S.L. (2003). Ultrasonographic detection of ingested and perforating wooden foreign bodies in four dogs. *Journal of American Veterinary Medical Association*, **223**(2):206-209.

Ragetly, G.R., Bennett, A. V and Ragetly, C.A. (2011). Septic Peritonitis: Etiology, Pathophysiology, and Diagnosis. *Compendium: Continuing Education for Veterinarians*, **33** (10):E1-6.

Schein, M and Paladugu R.(2000). What's new in pathophysiology of peritonitis. *Acta Chir Austriaca*, **32**:162-166.

Zachary F. J. (2012). Peritonitis.Pathological basis of Veterinary Diseases. 6<sup>th</sup>Edn. Saunders Elsevier, pp.338.

Fatal secondary septic peritonitis associated with multiple renal and splenic infarcts in lhasa apso dog



Figure 1: Gross changes in the abdominal cavity



Figure 2: Intestine showingperforations



Figure 3: Cut surface of Kidney showing gross lesions



Figure 4. Spleen showing gross lesions



Figure 5: Kidney showing infarct and diffuse coagulative necrosis of the glomeruli (G)and the tubules (T), interstitial oedema / congestion H&E,100×.



Figure 6: Section of splenic artery occluded with a bacterial embolic thrombus, H & E, 40  $\times$ 

# **Comparative advantage of livestock component in reduction of poverty – logistic regression approach\***

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#### ABSTRACT

A logistic model was fitted for the sample respondents to explore the determinant factors of poverty and comparative advantage of livestock component in reduction of poverty was ascertained. The primary data were collected through personal interview from randomly selected 540 sample households from six poverty prone districts using pre-tested interview schedules. Among various variables presumed to be the determinants of poverty, the variables viz., Ariyalur district dummy, cattle holding, sheep holding, family size, family dependency ratio were found to be statistically significant and rest were non-significant. The number of cattle and sheep were found to reduce the probability of fell down below the poverty line. Thus, these components may be considered while framing any poverty alleviation programmes in rural India.

Key words: Poverty - Livestock - Determinants - Logistic regression

Indian economy is a clear example of dualism. India excels in food production on one side, however on the other hand nearly 363 million people live in extreme poverty in India and face deprivation in terms of nutrient intake and access to basic services. These facts revealed that despite the country's meteoric GDP growth rate, poverty in India is still pervasive, especially in rural areas. Hence, the study of poverty is an important issue in the field of development. No development can even be thought of if any household / person in any country lives below the poverty line (Chatterjee, 2009). In this context, the present study was undertaken to fit a logistic model to explore the determinant factors of poverty and comparative advantage of livestock component in reduction of poverty.

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<sup>&</sup>lt;sup>#</sup>Part of the Ph.D. thesis of the first author submitted to the Tamil Nadu Veterinary and Animal Sciences University, Chennai

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Six sample districts of Tamil Nadu *viz.*,Thiruvannamalai, Villupuram, Dharmapuri, Pudukottai, Ariyalur and Ramanathapuram located in southern part of India were selected based on the composite poverty indices. From each selected district, 90 respondents comprising different occupational groups were selected through multi-stage random sampling leading to the sample size of 540 sample households. The data pertaining to the objectives of the study were collected through personal interview using structured pre-tested interview schedule. The period of data collection was from August 2013 to October 2014.

A logistic model was fitted for the sample respondents to explore the determinant factors of poverty. In explaining a dichotomous dependent variable ( $Y_i$ ), where '1' represents person falling below poverty line, and '0' represents above poverty line. The relationship between dependent and independent variables is non-linear, a logistic function was used to estimate the association between binary dependent variable  $Y_i$  and the independent variables ( $X_{ij}$ s). The following mathematical form of the model was used in the present study.

$$\ln\left(\frac{p_i}{1-p_i}\right) = \alpha + \sum_{j=1}^k \beta_j X_{ij} + \mu$$

Where,  $p_i$  is the probabilityofi<sup>th</sup> household to fall below poverty line and  $X_{ij}$  is the j<sup>th</sup> explanatory variable of i<sup>th</sup> household. The dependent variable ln  $\left(\frac{p_i}{1-p_i}\right)$  is the log-odds ratio in favour of sample households to fall below poverty line.

The results of the logit model to assess the determinant factors of poverty are given in Table 1. On perusal of table, it could be noted that the model Chi square was 411.018, which implied that the model was statistically significant. Among 25 variables presumed to be the determinants of poverty, the variables *viz*., Ariyalur district dummy, cattle holding, sheep holding, family size, family dependency ratio were found to be statistically significant and rest were non-significant (P>0.05).

Among the significant variables, the variables cattle holding and sheep holding alone were the negative determinant factors of poverty, which indicated the negative relationship between cattle and sheep holding with the existence of poverty.

The variables namely Ariyalur district dummy, family size and family dependency ratio were found to be positive determinant factors of poverty similar to the findings of Hashmi et al. (2008). As the family size especially number of dependents increases, the total family income has to be shared for more number of family members which lead to decrease in per capita income. Thus the individual household might have more chance to fall below the poverty line, if the family size and number of dependents increases. In contrast, as the cattle and sheep holding provides regular and more income for the livestock farmers, these variables had negative relationship on the determinants of poverty. The results are in line with the findings of Ambika (2003).

The reliability of the usage of the logit model for the correct prediction of below poverty line and above poverty line was tested by comparing the observed and predicted values. The percentage of correct prediction by the logit model was 87.40 per cent. The model was successful in predicting the above poverty line (88.10 per cent) correctly than the below poverty line (86.60 per cent).

Thus, it could be concluded that special attention must be given to Ariyalur district so as to uplift the people from clutches of poverty. As the number of cattle and sheep were found to reduce the probability of fell down below the poverty line, these components might be considered while framing any poverty alleviation programmes in rural India.

## ACKNOWLEDGEMENT

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## REFERENCES

Ambika, L. (2003). Livestock as a source of improving rural equity and poverty: A study in Karur district of Tamil Nadu. Unpublished MVSc. thesis submitted to the *Tamil Nadu Veterinary and Animal Sciences University, Chennai.* 

- Hashmi, A.A., Sial, M.H and Hashmi, M.H. (2008). Trends and determinants of rural poverty; A logistic regression analysis of selected districts of Punjab. *The Pakistan Development Review*, **47**(4):909-923.
- Chatterjee, S. (2009). Estimation of rural poverty; A discussion with reference to India. WYE City group on statistics on rural development and agriculture household income.Second meeting, Italy, Rome.11-12 June.

v	Evolanatory variables	Co-officients	Standard error	t volue	P voluo
	Villement district				
	Villupuram district	-0.185	0.484	0.146	0.703
	Dharmapuri district	0.096	0.515	0.035	0.851
X	Pudukottai district	1.017	0.565	3.241	0.072
	Ariyalur district	1.207*	0.571	4.462	0.035
X <sub>5</sub>	Ramanathapuram district	0.240	0.681	0.124	0.725
X <sub>6</sub>	Cropping occupation	-0.140	0.505	0.077	0.781
X <sub>7</sub>	Livestock farming occupation	-0.107	0.444	0.058	0.809
X <sub>8</sub>	Fishing occupation	-0.164	1.053	0.024	0.877
X <sub>9</sub>	AgriculturalLabourer occupation	19.873	5939.621	0.000	0.997
X <sub>10</sub>	Non-farm occupation	0.193	0.804	0.058	0.810
X	Number of cattle	-1.453**	0.343	17.920	0.000
X <sub>12</sub>	Number of buffalo	0.220	0.682	0.104	0.748
X <sub>13</sub>	Number of sheep	-0.302*	0.129	5.494	0.019
X <sub>14</sub>	Number of goat	-0.243	0.359	0.459	0.498
X <sub>15</sub>	Age of the head of household	0.000	0.015	0.002	0.968
X <sub>16</sub>	Gender of the head of household	0.249	0.404	0.379	0.538
X <sub>17</sub>	Illiterate dummy	0.540	0.340	2.516	0.113
X <sub>18</sub>	Family size	0.237*	0.100	5.640	0.018
X <sub>19</sub>	Family dependency ratio	14.249**	1.433	98.903	0.000
X <sub>20</sub>	Hindu Religion dummy	0.759	0.781	0.944	0.331
X <sub>21</sub>	Christian Religion dummy	-0.072	1.000	0.005	0.943
X <sub>22</sub>	Scheduled Castes (SC) dummy	0.523	0.420	1.551	0.213
X <sub>23</sub>	Scheduled Tribes (ST) dummy	0.667	0.458	2.122	0.145
X.24	Landholdings	-0.032	0.048	0.435	0.510
X25	Value of assets owned	0.000	0.000	2.021	0.155

 Table 1

 Estimates of binary logistic model to explore determinants of poverty

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## Table 1

## Estimates of binary logistic model to explore determinants of poverty

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X,	Dharmapuri district	0.096	0.515	0.035	0.851
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X <sub>4</sub>	Ariyalur district	1.207*	0.571	4.462	0.035
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X <sub>23</sub>	Scheduled Tribes (ST) dummy	0.667	0.458	2.122	0.145
X <sub>24</sub>	Landholdings	-0.032	0.048	0.435	0.510
X <sub>25</sub>	Value of assets owned	0.000	0.000	2.021	0.155
	Constant	-9.895	1.530	41.843	0.000
	Model Chi square	411.018**			
	-2 Log likelihood	333.658			
	Nagelkerke R square	0.712			
	Ν	540			

\*\* Significant at one per cent level and \* Significant at five per cent level

Comparative advantage of livestock component in reduction of poverty – logistic regression approach

## **Classification Table**

		Predicted			
Observ	ved	Below p	Percentage correct		
		Yes	No		
Dolom noverte line	Yes	214	33	86.6	
Below poverty line	No	35	258	88.1	
	87.4				

## Dystocia due to dizygotic twins in a crossbreed cow - a case report

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#### ABSTRACT

Twining incidence is influenced by a wide variety of genetic and environmental factors ranging from 0.5 to 4%. This case report places on record the successful management of dystocia due to dizygotic twins in a pluriparous cross-breed cow of four and a half year of age in her third parity. Both fetuses were non identical and had separate placenta which suggest 'Dizygotic twins'.

Key words: dystocia, dizygotic twins, cow

In uniparous species, twining is a highly undesirable trait. The incidence of twinning in sheep and goat is very high (60-70%), whereas low in dairy cattle (1.04%) and mare (0.5-1.05%)(Roberts, 1971). Bovine twins are of the two types 1) monozygotic twins, genetically and phenotypically identical, since they are formed from one fertilized egg, which splits into two identical halves during early embryonic developmental stages and thereby both individuals are always of the same sex. 2) Dizygotic or fraternal twins are not identical genetically or phenotypically as monozygotic twins, since they are formed from two different sperms fertilizing with two completely different ova at the same time. Dizygotic twins are the most common typeand not necessarily of the same sex because they are the result of ovulation and fertilization of two different oocytes. They can be also as similar or different as any two siblings born from the same parents during different

gestations (Wakchaure and Ganguly, 2016).

A pluriparous cross-breed cow of four and a half year of age in her third parity was brought to Government Veterinary Hospital, Prahladpur, District Kurukshetra (Haryana) with a history of unproductive straining with all the four limbs of the calf partially visible near the vulval lips. The general condition of the animal was fair and the pelvic ligaments were totally relaxed. Rectal temperature, respiratory rate and pulse rate of the animal were within the normal range.

The birth passage was completely relaxed and sufficient fetal fluids were present. Per vaginal examination of all the four limbs revealed that they were all forelimbs. Fetal head was also deviated to right side and fetal reflexes were also present. Further exploration revealed another head deep inside the birth canal and it was diagnosed as a case of dystocia due to twin fetus.

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**Fig.-** Twin live female fetus

After determining the relation of appendages to each fetus, mutational operations like retropulsion of two limbs, correction of lateral deviation of head and neck and traction of one fetus followed by another was performed and two live female fetus were delivered. Both fetuses were non identical and had separate placenta which suggest 'Dizygotic twins'.Careful examination of the birth canal revealed no injury or laceration subsequent to mutational operation. Parenteral administration of antibiotics [Ceftrioxone+Tazobactum-4.5gm (Intaceftazo-Intas Pharmaceuticals)] and analgesics [Meloxicam 30 ml (Melonex-Intas Pharmaceuticals)] intramuscularly daily for three days to prevent secondary bacterial infection due to external contamination during manipulation was followed and the cattle recovered uneventfully.

Twining incidence is influenced by a wide variety of genetic and environmental factors ranging from 0.5 to 4% (Fricke,2015). There is higher incidence of twins in dairy cattle as

opposed to beef cattle (Rutledge, 1975). Increased nutritional intake to improve productivity also increases the twinning incidences. Economic loss due to twining in cows often results from higher incidence of abortions, subclinical or clinical ketosis, reduced birth weight or stillbirths, mastitis, and problems related to dystocia.

### REFERENCES

- Fricke PM.(2015). Double Vision: Management of Twinning in Dairy Cows. Madison: Universityof Wisconsin-Madison;
- Roberts, S.J. (1971). Veterinary Obstetrics and Genital Diseases. (2<sup>nd</sup>edn.), CBS Publishers, New Delhi.
- Rutledge J. (1975). Twinning in cattle. *Journal of Animal Science*, **40**:803
- Wakchaure, R and Ganguly S. (2016). Twinning in Cattle: A Review. Academicians Research Center Journal of Gynecology and Obstetrics, 1(4):1-3.

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## **INSTRUCTIONS TO AUTHORS**

## Scope of the Journal

**"Indian Journal of Veterinary and Animal Sciences Research"** published six times in a year will consider original papers for publication on all aspects of animal and fisheries sciences. The scope of the journal includes animal and fisheries health, management, production and marketing of products. Acceptance of manuscript will be based on scientific merit as judged by referees and Editorial Board.

## Submission of manuscripts

Manuscripts should be written in English and the spelling should follow the Oxford

English Dictionary. Manuscripts should be submitted in triplicate along with Rs. 500/- as Demand Draft drawn in favour of "The Editor, IJVASR & Director of Research, TANUVAS, Chennai – 600 051" as processing fee to the Editor, "Indian Journal of Veterinary and Animal Sciences Research", Directorate of Research, Madhavaram Milk Colony, Chennai – 600 051, INDIA. Manuscripts can also be submitted by email to the email id: ijvasr@tanuvas.org.in. Payment can also be made online to the following account.

Account Name: **The Editor, IJVASR & Director of Research, TANUVAS, Chennai** Account Number: **332902010721641** IFSC Code: **UBINO533297** Reg. No.: / Transaction I.D./NEFT :

The authors should give a statement to the effect that the "Articles sent to IJVASR have not been sent elsewhere for publication". The statement should also be signed by all the authors and certified that the work is done as per the mandate of the respective institute. Email id and contact phone of the first or corresponding author should be provided whereas the authors should submit the copy of the IAEC approval if experimental animals are used.

## **Preparation of manuscripts**

All manuscripts should be typed on one side of the A4 paper, double-spaced throughout, with margins of at least 25mm all around. All contributions will be subjected to editorial revision.

**Major headings** are centered, all capitals except for scientific names and the full length papers should consist of Abstract, Introduction, Materials and Methods, Results and Discussion, Acknowledgement (optional) and References. First subheadings begin at the left margin and the text that follows a first subheading should be in a new paragraph.

**Full length papers** should normally not exceed 3000 words in length including tables and illustrations i.e. approximately five journal pages and should contain the following section, each written concisely:

A **Title** page containing (a) the title of the paper in capital letters in exception for scientific names, (b) the names of authors in full with initials at the beginning, (c) the authors' department and complete postal address. Superscript numbers should be used to link authors with other institution. Provide maximum of five key words for full length paper and three for short communication for subject indexing. The author wise contribution should also be mentioned in nutshell.

An **Abstract** will be printed at the beginning of the paper. Abstract should not be more than 150 words emphasizing objectives, experimental procedure, results and conclusions. Use complete sentences and limit the use of abbreviations. It should be in a form suitable for abstracting journals to use.

A brief **introduction** with specific emphasis on the necessity for such a kind of research may be given.

**Materials and methods** section may refer to previous description of methods whenever possible. This section should include experimental designs and methods of statistical analysis.

**Results and Discussion** may contain subheading if appropriate. This part should be brief and to the point, without repetition of results.

An Acknowledgement section, if required, may be given.

**References** section should contain only essential references which should be listed alphabetically and written as indicated below. In the text, give the author's name followed by the year in parenthesis: Suresh (2009). If there are two authors, use 'and': Suresh and Mani (2015); but if cited within parenthesis: (Suresh and Mani, 2015). When reference is made to a work by three or more authors, the first name followed by et.al. should be used: Rama et.al. (2015); but if cited within parenthesis: (Rama et.al., 2015). Reference to unpublished data and personal communications should not appear in the list but should be cited in the text only (e.g. Amutha T, 2015. Unpublished data).

## Journal articles and abstracts

- Bardbury, J.M., Mc Carthy, J.D and Metwali, A.Z. (1990). Micro immunofluorescence for the serological diagnosis of avian Mycoplasma infection. *Avian Pathology*, **19**:213-222.
- Raja, S., Rani, A., Ravi, M and Kumar. K. (2007). Histopathology of CPV infection. Page no. 120-122....Venue...Date...Place...

### Books and articles within edited books

Rundall, C.J. (1991). A colour Atlas of Diseases of the Domestic Fowl and Turkey. 2<sup>nd</sup> ed. London. Wolf Publishing Ltd. 175 p.

## Handbooks, Technical bulletins, Thesis and Dissertations

Callow, L.L and Dalgliesh, R.J. (1982). Immunity and Immunopathology in Babesiosis. In: S. Choen and K.S. Warren (Ed) Immunology of Parasitic Infections. Blackwell, Oxford. pp 475-526.nded.

## **Electronic publications**

**Tables** should be typed on separate sheets, numbered consecutively in Arabic Numerals and have a short descriptive heading. Units of measure for each variable measured should be indicated. Appropriate estimates of variation (Standard error, standard deviation) must be provided with means. Use superscript letters for the separation of means in the body of the table and explain these in footnotes.

**Illustrations,** referred to as "figures" (Fig. 1etc.) should be on separate sheets and submitted Larger than the size desired for reproduction. Information in tables must not be duplicated in figures and vice versa. Legends, should be provided for each illustration. Line drawings should be either in black ink on smooth white paper or thin board or a good quality laser printout. Photographs and photomicrographs should be printed on glossy paper with good contrast. Magnification For photomicrographs should be indicated. Allillustrations should be indicated. While sending the manuscripts in email, and the figures should be separately sent in JPEG format but for gel pictures it should be in TIFF format with good resolution.

**Short communications and Case Reports** should have a title page as described for full length papers and should comprise approximately 1000 words including tables, illustrations and references. They may contain not more than two tables or illustrations. Methods, results and discussion should be in single section without headings. References should be kept to a minimum and be in the form described above.Review should have a title page as described for full length papers and should contain approximately 4000 words including tables, illustrations and references.

## Units, symbols and abbreviations

Units should conform to the International System of Units (refer Baron, D.N. (1994). Units, Symbols and Abbreviations: A Guide for Biological and Medical Authors. 4th ed. London.Royal Society of Medicine). Abbreviations should not be used in the title, section heading or at the beginning of sentences. As a rule, author-coined abbreviations should be in all capital letters. These should be spelled out in full with the abbreviation following in parentheses the first time they are mentioned.

## Proofs

Proofs will usually be sent to the first or corresponding author. Only typesetter's errors may be corrected; no changes in, or additions to, the edited manuscript will be allowed. It is a condition of acceptance that the Editors reserve the right to proceed to press without submitting the proofs to the author. While reasonable care will be taken to ensure that proof reading is correctly done, neither the Editors nor the Publishers shall be responsible for any errors.

## **Reprints**

It has been decided to discontinue the supply of 25 reprints as the contents of the articles is hosted as PDF in TANUVAS website. (www.tanuvas.ac.in/ijvasr.html).

## **Rejected article**

Hard copy of the rejected articles will not be referred to the authors. The chief editor has the sole rights to either accept or reject the manuscripts based on their merits without reasoning. The first/corresponding authors are requested to inform their email addresses and contact numbers while submitting manuscripts to this journal.

## EDITOR

## **ATTENTION CONTRIBUTORS**

The Editorial Board of Indian Journal of Veterinary and Animal Sciences Research has decided to collect Rs.500/- (Rupees Five hundred only) as processing fee in accordance with the order of Registrar, TANUVAS-(U.S.O.No.500601/G4/2016 Proc.No. 5639/G4/2016 dt 3.5.2016), from the authors at the time of submission of articles for publication in the Journal. This would help the authors to hasten the publication of their articles without any delay.

Hence, the corresponding author is requested to draw a demand draft for Rs.500/- in favour of "The Editor, IJVASR & Director of Research, TANUVAS, Chennai-600051" along with the manuscript during submission. The articles may be addressed to the Editor, IJVASR & Director of Research, TANUVAS, Madhavaram Milk Colony, Chennai-51. The authors are also requested to mention their contact phone number and E-mail address.

## EDITOR

## **REVIEW ARTICLES INVITED FROM EMINENT SCIENTISTS**

The Editorial Board of Indian Journal of Veterinary and Animal Sciences Research invites review articles from eminent research scientists in the field of Veterinary and Fisheries Sciences, on the latest/ current topics of interest for publication in the Journal. The review article (both hard and soft copy) may please be sent to the Editor/Associate Editor, Indian Journal of Veterinary and Animal Sciences Research for publication.

## FORM IV (See Rule 8)

1.	Place of Publication	:	University Publication Division (Printing Press) Tamil Nadu Veterinary and Animal Sciences University, Madhavaram Milk Colony, Mathur Road, Chennai – 51. Ambattur Taluk Thiruvallur District
2.	Periodicity of Publication	:	Bi-Monthly
3.	Printer's Name Whether citizen of India Address	:	<b>Dr. K.N.Selvakumar</b> Yes Director of Distance Education Tamil Nadu Veterinary and Animal Sciences University, Old.No. 327, New No. 485, Anna Salai, Nandanam, Chennai - 600 035
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5.	Chief Editor's Name Whether citizen of India	:	The Vice-Chancellor Yes Vice-Chancellor Tamil Nadu Veterinary and Animal Sciences University, Madhavaram Milk Colony, Chennai – 600 051.
6.	Name and address of individuals who own the newspaper and parents or share holders holding more than one per cent of the total capital	:	<b>The Registrar</b> Tamil Nadu Veterinary and Animal Sciences University, Madhavaram Milk Colony, Chennai – 600 051.

I, Dr. K.N. Selvakumar hereby declare that the particulars given are true to the best of my knowledge and belief.

Dr. K.N. Selvakumar Signature of Publisher All the contributing authors are requested to bestow their personal dattention while submitting the revised manuscripts for spelling mistakes and correctness of language.

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**Chief Editor** 

The Indian Journal of Veterinary and Animal Sciences Research (IJVASR) is indexed in the abstracting journals of the CAB International, Zoological Abstracts of Web of Knowledge published by Thomson Reuters and Indian Science Abstracts published by NISCAIR, India.