

## COMPARATIVE EFFICACY OF ANALGESIC AND ANAESTHETIC DRUGS FOR HIGH EPIDURAL ANALGESIA IN BLACK BENGAL GOATS

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### ABSTRACT

The effects of analgesic and anaesthetic drugs on high epidural analgesia in Black Bengal goats were investigated. The animals were divided into five groups (n=8) and a replication of 8 trials was performed in each group at least one week interval. Two percent (2%) lidocaine hydrochloride, 0.5% bupivacaine hydrochloride, 2% lidocaine hydrochloride with adrenaline, ketamine hydrochloride and diazepam were administered into lumbo-sacral (high epidural) space for high epidural analgesia. Bupivacaine hydrochloride significantly (P<0.05) decreased respiration rates, rectal temperature, however, increased heart rates during high epidural analgesia with prolonged analgesia compared to other drugs. Lidocaine hydrochloride showed a rapid onset and excellent analgesia and did not produce any side effects whereas, diazepam exhibited delay onset with short duration of analgesia. It seems that 2% lidocaine hydrochloride is effective for high epidural analgesia in goats.

**Key words:** Analgesics, anaesthetics, high epidural analgesia, goats

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### INTRODUCTION

Epidural analgesia, a regional analgesia, is needed to perform major surgical operations for correction of goat diseases as well local and regional analgesia have long being practiced for minor and major surgical affections. Among different types of local and regional analgesia, epidural analgesia technique is representing as an alternative to general analgesia for the patients at high risk. It has also been used in veterinary practice for diagnosis and treatment of obstetrical and surgical procedures at the caudal aspect of body; lower abdomen, flank, perineal region and in tail. Lidocaine has a relatively rapid onset and duration of action about 1 to 2 hours (Carpenter *et al.*, 2004), however, bupivacaine is most commonly used for epidural nerve block (Eugene and Nicholas, 1995). Addition of adrenaline is a common practice for prolongation of the action of local analgesic drugs. Ketamine hydrochloride (HCl), a dissociative agent has been used epidurally to relieve pre and post operative pain in both man and animals (Brander *et al.*, 1991). Diazepam, a sedative agent has been used routinely for sedation in animals, though there is lack of evidence of its effects during high epidural analgesia. The present experiment was carried out to find out the analgesic effects of different analgesic and anaesthetic drugs during high epidural analgesia.

### MATERIALS AND METHODS

A total of 8 (3 castrated males and 5 non pregnant females) goats were taken for the experiment in the operation theatre of the Department of Surgery and Obstetrics, Faculty of Veterinary Science and Veterinary Clinic, Bangladesh Agricultural University (BAU), Mymensingh. The body weight of these animals ranged from 7 to 10 kg and age ranged from 8 to 15 months. They were allowed to graze in the open field for 5 to 6 hours daily and water *ad libitum*. The animals were restrained on the operation table in right lateral recumbency. The site of injection was clipped, cleaned and disinfected and analgesic solution was inserted at the lumbo-sacral space. The animals were divided into 5 (five) different groups and a replication of 8 trials were performed in each group at one week interval. Two percent (2%) lidocaine hydrochloride (Jasocaine<sup>®</sup>, Jayson Pharmaceuticals Ltd.) 3ml, 0.5% bupivacaine hydrochloride (Ultracaine<sup>®</sup>, Jayson Pharmaceuticals Ltd.) 3ml, 2% lidocaine hydrochloride with adrenaline (Jasocaine A<sup>®</sup>, Jayson Pharmaceuticals Ltd.) 3ml, ketamine hydrochloride (Calypsol<sup>®</sup>, Gedeon Richter Ltd., Budapest, Hungary) 1ml and diazepam (Sedil<sup>®</sup>, Square Pharmaceuticals Ltd.) 2ml were injected into lumbo-sacral space in groups A, B, C, D and E respectively.

Respiration rate, heart rate and rectal temperature were recorded before, during and after recovery of analgesia. The state of analgesia was determined by needle pricking in the lumbar region in every 5 minutes. Analgesia was assessed as “+++” (no response), “++” (slight movement or reflex response), “+” (avoidance response) by scoring excellent, adequate and poor respectively. Onset, peak point and the duration of analgesia were recorded and measured by a scale. Tail movement, leg movement and any side effects were closely observed and recorded during the course of analgesia. The data were analyzed statistically by Students’ Paired “*t*” test and ANOVA in completely randomized design.

## RESULTS AND DISCUSSION

The effect of various analgesic and anaesthetic drugs on respiration rate, heart rate and rectal temperature in different groups of Black Bengal goats in high epidural analgesia are presented in Table 1.

Table1. Effects of analgesic and anaesthetic drugs on respiration rate, heart rate and rectal temperature during high epidural analgesia in Black Bengal goats

Drugs used	Time of recording analgesia (interval = 10mins)	Respiratory rate (min) <sup>a</sup>	Heart rate (min) <sup>a</sup>	Rectal temperature <sup>a</sup> (°F)
2% lidocaine HCL	Before	32.50±5.53	61.25±5.01	102.71±0.67
	During	30.50±7.76	65.25±8.00	102.25±0.96**
	After	33.13±6.17	62.38 ± 4.60	102.38±0.44
0.5% bupivacaine HCL	Before	31.75±6.54	63.25±8.21	102.38±1.24
	During	28.00±4.28*	67.75±10.71*	102.09±1.32*
	After	31.25±4.65	65.00 ± 8.21	101.96±1.55
2% lidocaine HCL with adrenaline	Before	32.25±4.83	66.25±5.50	101.95±0.99
	During	28.00± 3.85**	68.00±9.07	101.61±1.15*
	After	32.25±4.17	66.50 ± 6.21	101.63±1.62
Ketamine HCL	Before	31.50±4.11	57.75±8.24	103.18±1.09
	During	35.00±10.25	64.25±7.21**	102.43±0.82**
	After	32.25±4.20	60.75 ±7.25	102.43±0.66*
Diazepam	Before	33.00±7.41	62.00±6.14	102.28±0.80
	During	30.25±7.74*	66.25±5.39**	102.11±0.75
	After	32.13±7.64	63.50±5.42	102.03±0.72

<sup>a</sup>Mean±SD, \* = Significant (P<0.05), \*\* = Highly significant (P<0.01).

Two percent (2%) lidocaine hydrochloride insignificantly decreased respiration rates and increased heart rates in all the experimental animals. Bupivacaine hydrochloride (0.5%) significantly (P<0.05) decreased respiration rates and rectal temperature. Singh *et al.* (2005) also found significant fall in respiration rate after epidural administration of medetomidine or bupivacaine in goats. Decreased respiratory rate might result from their depressing action on respiratory center in central nervous system (Hall and Clarke, 1989). When 2% lidocaine hydrochloride with adrenaline was used respiration rates significantly (P<0.01) decreased and heart rates insignificantly increased. Ketamine hydrochloride increased heart rates significantly (P<0.01) but respiration rates insignificantly. This result corresponds with the study reported by Singh *et al.* (2002). The initial bradycardia followed by increase in heart rate might be due to cardiovascular stimulant effect of ketamine (Kumar and Singh, 1990). Diazepam decreased respiration rates significantly (P<0.05) whereas increased heart rate significantly (P<0.01). Sanhouri *et al.* (1991) reported the similar suppressive effect on respiration in goat by diazepam injected through lumbo-sacral space. This reduction in respiration rate resulted from direct depressing effect of diazepam on central nervous system. In this study, temperature in goats of most of the groups decreased. There is no report on the effect of local analgesic agents on body temperature during epidural analgesia in goats.

### Epidural analgesia in goats

The analgesic effects were assessed in different goats by using scoring method. Two percent (2%) lidocaine hydrochloride produced excellent analgesia compared to other drugs. There were significant differences in different variance after epidural injection of same drug with same dose in different goats (Table 2). The onset was rapid in case of 2% lidocaine hydrochloride. On the contrary, the onset was slower in case of 0.5% bupivacaine hydrochloride and diazepam. Mahale and Wakanker (1992) observed the similar findings in goat. There was no report about onset of analgesia after epidural administration of diazepam in goats.

Table 2. Effects of analgesic and anaesthetic drugs during high epidural analgesia in Black Bengal goats

Drugs used	Onset of analgesia (min)	Peak point of analgesia (min)	Desensitized area (cm)	Duration (min)
2% lidocaine HCL	1.88±0.64 <sup>c</sup>	15.50±4.04 <sup>a</sup>	22.63±9.00 <sup>a</sup>	63.50±14.91 <sup>a</sup>
0.5% bupivacaine HCL	5.13±2.10 <sup>b</sup>	15.00±3.63 <sup>a</sup>	25.21±10.30 <sup>a</sup>	81.75±36.64 <sup>a</sup>
2% lidocaine HCL with adrenaline	4.88±1.73 <sup>b</sup>	11.25±3.85 <sup>ab</sup>	21.81±7.73 <sup>a</sup>	81.50±23.14 <sup>a</sup>
Ketamine HCL	5.00±1.60 <sup>b</sup>	8.38±2.50 <sup>b</sup>	18.18±10.55 <sup>a</sup>	27.63±7.63 <sup>b</sup>
Diazepam	7.88±1.36 <sup>a</sup>	10.38±1.69 <sup>b</sup>	18.88±7.09 <sup>a</sup>	25.50±7.03 <sup>b</sup>

Values with different superscript letters in the same column differ significantly (P<0.01).

The high peak time of analgesia was found with 2% lidocaine hydrochloride among different local analgesic agents. This observation corresponds with the previous findings (DeRossi, 2005). The inherent spreading power of lidocaine might be responsible for its wider spread blockade and intense action (Hall and Clarke, 1989). The duration of analgesia with 0.5% bupivacaine hydrochloride was found longer than 2% lignocaine hydrochloride in this experiment. The protein binding characteristics of local analgesic agents influence the duration of action (Gissen *et al.*, 1980). Lidocaine showed intermediate duration of action among other agents used here. This observation was supported by Lemke and Dawson (2000). The duration of analgesia with diazepam was very short among all the groups of the experiment. Drowsiness and shivering were found during epidural analgesia with 0.5% bupivacaine hydrochloride. These findings are in agreement with the study reported by Laishley *et al.* (1988). In case of diazepam more drowsiness and salivation were observed in all the animals. It is suggested that 2% lidocaine hydrochloride is the effective analgesic agent for high epidural analgesia comparing to other agents.

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