**KEYWORDS** 



clonidine, Morphine, bupivacaine, lower limb surgeries, post operative analgesia.

# Dr. P.Harish Gautam

Assistant Professor, Department of Anaesthesiology, Alluri SitaRamaraju Academy of Medical Sciences (ASRAM), Eluru, AP, India.

**ABSTRACT** Background: Severe pain is common after surgeries like TKR (Total Knee Replacement) and can delay early commencement of physiotherapy which is the most important determinant of successful postoperative knee rehabilitation. Attainment of adequate postoperative analgesia in patients who undergo lower limb surgeries is often a challenging task. The current and the most common approach in post operative pain management is multi-modal analgesia. Various combinations of epidural clonidine, opioids and local anaesthetics have improved post operative analgesia after lower limb surgeries. This study is done to determine the optimal epidural bolus dose of clonidine, which provides the best analgesia and least side effects.

Aim of the study: This study was designed to evaluate the effects of the addition of two different doses of clonidine to epidural morphine and bupivacaine on duration and efficacy of post operative analgesia and side effects after lower limb surgeries

Materials and methods: The study included 60 patients, ASA I and II scheduled for lower limb surgeries at ASRAM Medical College and Hospital, Eluru. These patients were randomly allocated into three groups, 20 patients in each group. Group A where patients received 2-mg morphine + 12.5mg Bupivacaine, Group B where patients were given 2-mg morphine + 12.5mg Bupivacaine + 10.5mg Bupivacaine + 100 g of clonidine. Postoperative pain was assessed using Visual Analog Scale (VAS). Demographics, time intervals, and continuous variables (SBP, DBP, heart rate, VAS) were analyzed using the one-way analysis of variance (ANOVA) test. Categorical data (ASA, sex) were analyzed using chi-square test. In all cases, P < 0.001 was considered statistically significant.

Results: Demographic characteristics as well as intraoperative SBP, heart rate and were similar among groups. Epidural administration of combinations of local anesthetic bupivacaine(1mg/ml), opioid morphine(0.2mg/ml), and clonidine(7.5µg/ml in group B and 100µg/ml in group C) resulted in significant improvement of analgesia after lower limb surgeries The total amount of drug consumed in clonidine groups was significantly lower compared with control group (group-A) p<0.001. The clonidine groups (group-B, group-C) compared with control group (group-A) had significantly less pain at 4hr, 12hr and 16hrs as p<0.001. Systolic blood pressure and diastolic blood pressure were low for the clonidine groups (group-B, group-C) compared to control group (group-A) at all sampling intervals. Relative hypotension in group-C who received higher dose of clonidine (435µg over 24 hr period) for diastolic blood pressure were recorded which were statistically significant at 16hr, 20hr and 24hrs though did not require any therapeutic intervention. In this study 10 patients in group-C and 5 patients in group-B have higher (score 2) sedation scores at 4hr and 16hr, scores were statistically significant between clonidine groups and control group at 4hr

Conclusion: When added to lumbar epidural mixture of bupivacaine, morphine and clonidine augmented analgesia both in quality and duration after lower limb surgeries in comparison with patients who received bupivacaine and morphine drug combination only.

Group with lower concentration of clonidine group-B (7.5 $\mu$ g/ml) produced significant analgesia compared to control group (group-A with bupivacaine(0.125%) and morphine 0.2mg/ml only) and experienced significantly less side effects of clonidine compared to group-C( with 10 $\mu$ g/ml of clonidine).

#### Introduction

Pain is one of the most unpleasant experiences for mankind. International association for the study of pain defines the term pain thus: "Pain is an unpleasant sensory and emotional experience associated with actual or potential tissue damage or described in terms of such pain."

Surgical pain is a kind of acute pain, which is always a challenge to the anaesthesiologist. However, the patients' problems of pain do not end with the surgical procedure. Pain during post-operative period is also of concern to the anaesthesiologist.

Post-operative pain control is generally best managed by anesthesiologist, because they offer regional techniques of anesthesia as well as pharmacological expertise in analgesics Adequate postoperative analgesia after total knee Replacement (TKR) is often a challenging task. Pain is rated as severe in the majority of patients who undergo TKR<sup>2</sup>. This can easily hinder early physical therapy, which is the most important factor for successful postoperative knee rehabilitation<sup>3</sup>. Postoperative pain control is complicated by rehabilitation and the recovery goal of continuous movement of the joint.

The current trend in postoperative pain management is multimodal analgesia  $\!\!\!^4$ 

Effective postoperative analgesia can be provided by neuraxially applied local anesthetics<sup>5</sup> or opioids, which may be accompanied by unwanted side effects, such as motor block, hypotension, or respiratory depression.<sup>6</sup>

The analgesic neuraxial effect of the alpha\_2-adrenergic agonist clonidine is well described<sup>7-10</sup>

The alpha\_2-adrenergic agonist clonidine produces analgesia by a different mechanism that mimics the effect of endogenously released norepinephrine to stimulate postsynaptic alpha\_2 receptors in the spinal cord. Intrathecal (IT) and epidural clonidine provide effective analgesia in volunteers<sup>11</sup>, in patients in labor<sup>12</sup>, and in those with postoperative pain<sup>13</sup>, without inducing respiratory depression or motor block<sup>14</sup>

intrathecal morphine is a well described modality for providing prolonged postoperative analgesia. Unfortunately, both drugs are limited by their unique side effect profiles  $^{15\cdot16}$ 

First, we hypothesized that the combined administration of epidural morphine and clonidine would provide superior postoperative analgesia in patients undergoing lower limb surgeries with comparison of two doses of clonidine. Second in addition taking advantage of the potential synergy between neuraxial morphine and clonidine, we hypothesized that the doses of these drugs could be reduced, therefore minimizing their side effect profiles.

#### Aim and objectives:

To evaluate the effects of the addition of two different doses of clonidine to epidural morphine and bupivacaine on duration and efficacy of post operative analgesia and side effects after lower limb surgeries.

#### **Review of literature**

From time to time man has resorted to many means in his search for relief of pain. Painless surgery and painless post operative period is probably the greatest achievement that has been granted to the patients. There are various methods used, each one carrying its own merits and demerits.

During the past two decades, epidural and intrathecal administration of drugs have been increasingly used for relief of pain.

Motsch J, Graber,Ludwig et al in 1990 compared the analgesic effect of the combination of epidural morphine and clonidine verses epidural morphine alone in patients with postoperative pain. They reported pain scores were lower in clonidine and morphine group compared to epidural morphine only group<sup>17</sup>.

Mogensen T, Eliasen K, Ejlersen et al in 1992 compared the value of adding clonidine to a low-dose epidural regimen for postoperative pain by a randomized, doubleblind, placebo-controlled trial .They concluded that a continuous low-dose epidural clonidine infusion enhances analgesia from a combined low-dose epidural bupivacaine and morphine regimen after hysterectomy.<sup>18</sup>

Marjan Jahangiri, J W P Bradley, Dark et al in 1994 studied combined efficiency of epidural infusion containing diamorphine, bupivacaine and clonidine in preventing phantom limb pain in a prospective controlled study undergoing lower limb amputation. They concluded that perioperative epidural infusion of diamorphine, clonidine and bupivacaine is safe and effective in reducing the incidence of phantom pain after amputation<sup>19.</sup>

Michael J. Paech, J. G. Pavy, E. P. Orlikowski, et al 1n 1997 studied the effect of adding clonidine in different doses to

#### Volume : 6 | Issue : 3 | March 2016 | ISSN - 2249-555X | IF : 3.919 | IC Value : 74.50

epidural Bupivacaine and fentanyl infusion on postoperative analgesia in patients undergoing abdominal gynecological surgeries. The study was randomized and double blinded. They also studied to determine an effective dose rate that would minimize the sedative and hemodynamic effects. They concluded that high-quality analgesia after abdominal surgery produced by postoperative epidural bupivacaine and fentanyl was further enhanced by the addition of epidural clonidine<sup>20</sup>

Brian D. Sites, Michael Beach, Russell Biggs et al in 2003 compared the effect of intrathecal Clonidine with morphine versus morphine in a prospective, randomized placebo-controlled study on postoperative analgesia after total knee arthroplasty. They concluded that intrathecal morphine, the combined administration of clonidine and morphine reduces postoperative IV morphine use and improves the VAS score at 24 hours. There was an increase in relative hypotension which did necessitate intervention<sup>21</sup>

Marjan Jahangiri, C H Dark, A P Jayatunga et al studied the effect of adding clonidine to epidural bupivacaine and diamorphine on post operative analgesia in patients undergoing lower limb amputation. They concluded that perioperative epidural infusion of diamorphine, clonidine and bupivacaine is safe and effective in reducing the incidence of phantom pain after amputation.<sup>22</sup>

B.S.Sethy, Mary Samuel, Deepak et al in 2007 studied the effect on post operative analgesia provided by low dose intrathecal clonidine admixed with bupivacaine as compared to bupivacaine alone. They concluded that addition of clonidine to bupivacaine in the dose  $1\mu g/kg$  intrathecally significantly increased the duration of analgesia compared to bupivacaine alone without significant side effects<sup>23</sup>.

Yuan-Shiou Huang, Liu-Chi Lin, Billy K et al in 2007 studied the effect of adding clonidine to epidural ropivacaine and morphine on post operative analgesia in patients undergoing Total knee Arthroplasty. In that double blind study they also sought to determine the optimal epidural bolus dose of clonidine which provides the best analgesia and fewest side effects. They concluded that by adding clonidine to lumbar epidural mixture of ropivacaine and morphine augmented analgesia after TKA surgery without significant adverse effects in lower concentrations of clonidine. They concluded that by adding clonidine to a lumbar epidural mixture of ropivacaine.<sup>24</sup>

#### Materials and methods

This randomized study after approval from ethical committee was conducted in 60 patients. Informed consent was obtained from each patient under study.

#### Inclusion Criteria

- Age group 40 80 years.
- Both sexes.
- ASA grade I and II.

#### **Exclusion Criteria**

- H/O allergic reaction and CI to any of the study drugs.
- CI to epidural catheter placement.
- CI to use of opioids, NSAIDS, corticosteroids
- Inability to understand VAS (visual analogue scale)
- DBP > 100 mm of Hg

After the pre anaesthetic checkup patients were recruited for the study as per the inclusion and exclusion criteria.

Volume : 6 | Issue : 3 | March 2016 | ISSN - 2249-555X | IF : 3.919 | IC Value : 74.50

Patients were familiarized with visual analogue scale at the time of pre anaesthetic checkup.

60 patients undergoing lower limb surgeries were randomly divided into 3 groups of 20 patients each to receive.

- Group –A-2-mg morphine+12.5mg (2.5ml of 0.5%) Bupivacaine and Normal saline to make 10ml.
- Group -B--2-mg morphine+75 μg Clonidine+12.5mg (2.5ml of 0.5%) Bupivacaine and Normal saline to make 10ml.
- Group-C---2-mg morphine+100 µg Clonidine+12.5mg (2.5ml of 0.5%) Bupivacaine and Normal saline to make 10ml.

#### Pre Medication:

All the patients were premedicated the night before and on the day of surgery with Tab. Omeprazole-20mg, Tab. Alprazolam- 0.5 mg. Inj. Tramadol-1mg/kg was given IM one hour before surgery.

Pre operatively lumbar epidural space was identified between L2-L3/L3-L4 and catheter inserted and cephalically advanced up to 3-4 cm in the epidural space.

Epidural catheter will be activated at the end of the surgery by injecting the epidural analgesic mixture of volume 10ml.

Top up dose was given whenever VAS > 4.

Top up dose of 10ml mixture consist of 2mg of morphine+0.125% bupivacaine.

Visual analogue scale, Heart rate, Systolic blood pressure, Diastolic blood pressure, Sedation score were measured every 4hr up to 24hrs post operatively.

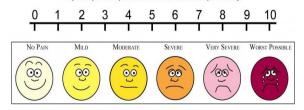
Duration between activation of epidural catheter (first epidural analgesic dose) and first top up dose and total number of top up doses given up to 24hrs in each group were recorded.

**Pain evaluation:** A 0 - 10 scale VAS with end points labeled no pain and worst possible pain

Pain intensity was measured every two hours up to 24hrs post operatively

Universal Pain Assessment Tool

This pain assessment tool is intended to help patient care providers access pain according to individual patient needs. Explain and use 0-10 Scale for patient self-assessment. Use the faces or behavorial observations to interpret expersed pain when patient cannot communicate his/her pain intensity.



## HEMODYNAMIC EFFECTS

SBP & DBP were measured for every two hours upto 24hrs postoperatively.

Hypotention defined as decrease by 20% of basal value.

Bradycardia defined as HR decrease by 20% of basal val-

**Sedation:** Score noted every two hours up to 24hrs post operatively as per the following scale.

#### Scale:

ue

- 1. Awake and alert
- 2. Awake but drowsy responsive to verbal stimulus
- 3. Drowsy but arousable responsive to physical stimulus
- 4. Unarousable not responsive to physical stimulus

## Sensory level: Assessed by pinprick

## Data analysis:

The collected data was summarized by calculating the mean and standard deviation and presented in the form of table and diagrams. Paired "t" test and analysis of variance for repeated measures were used for the analysis of significance. Chi Square test was used to obtain other possible associations.

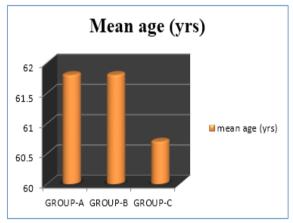
## **Observation and Results**

Demographic data like age, sex and ASA physical status are presented as mean  $\pm$  SD where appropriate. There were twenty patients in each study group, and the groups were demographically similar.

#### TABLE-I: AGE DISTRIBUTION (MEAN±SD):

	GROUP-A	GROUP-B	GROUP-C
MEAN±SD	61.8 ± 4.3	61.8 ± 5.88	60.7 ± 6.86





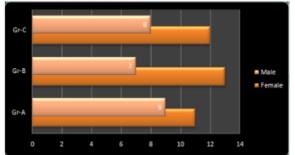
The table-I and graph-I shows age distribution of the three Groups. Group A is the bupivacaine 0.125% + morphine 2mg group, Group B is the bupivacaine 0.125% + morphine 2mg + clonidine 75µg group and group C is the bupivacaine 0.125% + morphine 2mg + clonidine 100µg group. The range of age is comparable in all the groups' i.e Group A 61.8 ± 4.3, Group B 61.8 ± 5.88 and Group C 60.7±6.86.

## TABLE II: SEX DISTRIBUTION (MEAN±SD):

SEX	Group-A	Group-B	Group-C
Female	11	13	12
Male	9	7	8

p = NS (chi-square)

## Graph -2: Comparison of Sex Distribution



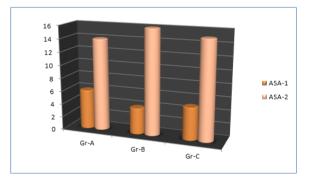
The three Groups had comparable patients in terms of male and female distribution. Group-A (bupivacaine 0.125% + morphine 2mg) had 9 male and 11 female patients. Group-B (bupivacaine 0.125% + morphine 2mg + clonidine 75µg) had 7 male and 13 female patients. Group-C (bupivacaine 0.125% + morphine 2mg + clonidine 100µg ) had 8 male and 12 female patients.

#### TABLE III: ASA PHYSICAL STATUS (MEAN±SD)::

ASA STATUS	ASA-1	ASA-2
Group-A	6	14
Group-B	4	16
Group-C	5	15

P = NS (chi-square)

## Graph-3: Comparison of ASA physical status



All the patients in our study belonged to ASA physical status I and II and the distribution of these patients randomly into the three study groups were comparable.

#### ANALGESIC EFFECTS AND TOTAL DRUG CONSUMP-TION

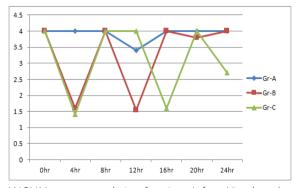
Means for demographic variables did not differ significantly among the three groups (Table 1, 2 and 3). No patients required rescue analgesic (inj. Tramadol). No patients were excluded from the study. Patients in the clonidine groups experienced significantly less pain than those in the control group during the 24 hr period after the surgery.

Table-IV -Visua	al analog	scale	(VAS)	pain	scores	(means)
-----------------	-----------	-------	-------	------	--------	---------

VAS	0hr	4hr	8hr	12hr	16hr	20hr	24hr
Group-A	4	4	4	3.4	4	4	4
Group-B	4	1.6	4	1.55	4	3.8	4
Group-C	4	1.4	4	4	1.6	4	2.7

Volume : 6 | Issue : 3 | March 2016 | ISSN - 2249-555X | IF : 3.919 | IC Value : 74.50

Graph-4: Comparison of Visual analog scale (VAS)pain scores



ANOVA(one way analysis of variance) for Visual analog scale pain scores

#### VAS at 4hr

if the value of t is greater that than 0.05.	n 2.467 then	the P val	lue is less
	Mean		P value
Connariano			
Comparison	Difference		
Comparison Column & vs Column B	2.400		*** Pc0.001
		12.029	

	Group A	Group B	Group C
Col. title			
Mean	4	1.6	1.4
Standard deviation (SD)	0.000	0.9403	0.5026
Sample size (N)	20	20	20
Std. error of mean(SEM)	0.000	0.2103	0.1124
Lower 95% conf. limit	4.000	1.160	1.165
Upper 95% conf. limit	4.000	2.040	1.635
Mnimum	4.000	1.000	1.000
Median (50th percentile)	4.000	1.000	1.000
Maximum	4.000	4.000	2.000
Normality test KS	-	0.3383	0.3869
Normality test P value	1	<0.0001	<0.0001
Passed normality test?		No	No

## VAS at 12hr

			Group A	Group B	Group C
		Col. title			
		Mean	4	4	1.6
Bonferroni Multiple Comparison	a Test	Standard deviation (SD)	0.000	0.000	0.5026
	than 2.467 then the P value is 1	Sample size (N)	20	20	20
	UND CLART POST POST A VALUE TO T				
than 0.05.		Std. error of mean(SEM)	0.000	0.000	0.1124
		Lower 95% conf. limit	4.000	4.000	1.365
		Upper 95% conf. limit	4.000	4.000	1.835
	Nean				
	P96.8.0	Minimum	4.000	4.000	1.000
Comparison	Difference t P val	Median (50th percentile)	4.000	4.000	2.000
		Maximum	4.000	4.000	2.000
Column & vg Column B	1.850 9.471 *** Pc0	.001 Normality test KS			
Column & vs Column C	-0.6000 3.072 ** Pd				0.3869
		rearry cover to be			<0.0001
Column B vs Column C	-2.450 12.542 *** Pc0	Passed normality test?			No
1					

## VAS at 16hr

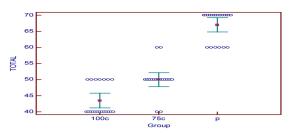
			Group A	Group B	Group C
		Col. title			
		Mean	3.4	1.55	4
Read-annel Malainta Anneala		Standard deviation (SD)	0.9403	0.5104	0.000
Bonferroni Nultiple Compariso If the value of t is greater	than 2.467 then the P value is less	Sample size (N)	20	20	20
than 0.05.		Std. error of mean(SEM)	0.2103	0.1141	0.000
		Lower 95% conf. limit	2.960	1.311	4.000
		Upper 95% conf. limit	3.840	1.789	4.000
	Mean	Minimum	2.000	1.000	4.000
Conparison	Difference t P value	Median (50th percentile)	4.000	2.000	4.000
		Maximum	4.000	2.000	4.000
Column & vs Column B	0.000 0.000 ns 950.05	Normality test KS	0.4383	0.3610	
Column & vs Column C	2.400 26.153 *** Pc0.001	Normality test P value	<0.0001	<0.0001	
Column B vs Column C	2.400 26.153 *** Pc0.001	Passed normality test?	No	No	

Visual analog scale (VAS) pain scores for the control Group (Group-A) were significantly higher at 4hr,12hr and 16hr than those for the 2 clonidine Groups(Group-B, Group-C)

Visual analog scale (VAS) pain scores for the Group-B with lower dose of clonidine (75µg) were higher at 4hr, 16hr and 24hr.VAS score was lower at 12 hr .

The total volumes of analgesic solution consumed by the different Groups during the study period were Group-A 67  $\pm$  4.58ml; Group-B 50  $\pm$  4.47ml; Group-C 43.5  $\pm$  4.769ml.

Graph-5: Comparison of total amount of drug mixture consumed



## ANOVA

cance level

Source of vari- ation	Sum of squares	DF	Mean square
Be- tween Groups (influence fac- tor)	5890.0000	2	2945.0000
Within Groups (other fluctua- tions)	1275.0000	57	22.3684
Total	7165.0000	59	
F-ratio	131.659		
Signifi-	P < 0.001		

## Student-Newman-Keuls test for all pair wise comparisons

P < 0.001

Factor	n	Mean	Different (P<0.05) from factor nr
(1) 100C	20	43.5000	(2)(3)
(2) 75C	20	50.0000	(1)(3)
(3) P	20	67.0000	(1)(2)

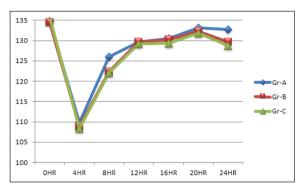
Group-C used a significantly lower volume of analgesic solution (p<0.001) than the other Groups during first 24hrs after surgery.

Groups with clonidine (Group-B and Group-C) used a significantly lower volume of analgesic solution (p<0.001) than the control Group(Group-A)

## **HEMODYNAMIC EFFECTS** Table-V - SYSTOLIC BLOOD PRESSURE (MEAN)

SBP	OHR	4HR	8HR	12HR	16HR	20HR	24HR
Group- A	134.8	109.8	126	129.7	130.5	133.1	132.7
Group- B	134.4	108.7	122.4	129.6	130.25	132.2	129.65
Group- C	135.8	108.3	122.1	129.2	129.3	131.8	128.7

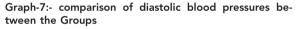
Graph-6:- comparison of systolic blood pressures between the Groups

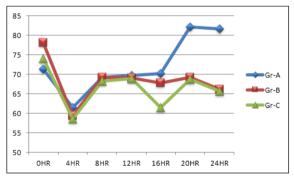


The systolic blood pressures were lower in clonidine Groups (Group-B, Group-C) than in the control Group (Group-A) but the difference was not statistically significant.

## Table-VI - DIASTOLIC BLOOD PRESSURE

DBP	OHR	4HR	8HR	12HR	16HR	20HR	24HR
Group- A	71.4	61.5	69.2	69.75	70.15	82.1	81.6
Group- B	78.1	59.4	69.15	69.1	67.7	69.2	66.1
Group- C	74	58.6	68.2	68.9	61.5	68.7	65.7





#### ANOVA DBP at 16hr

	Group A	Group B	Group C	Bonferroni Hultiple Comparisons Te			
Col. title				If the value of t is greater than		te P val	ue is less
Mean	70.15	67.7	61.5	than 0.05.			
Standard deviation (SD)	3.150	1.838	3.171				
Sample size (N)	20	20	20				
	1				Mean		
Std. error of mean(SEM)	0.7044	0.4110	0.7090	Comparison	Difference	t	P value
Lower 95% conf. limit	68.676	66.840	60.016				********
Upper 95% conf. limit	71.624	68.560	62.984	Column & vs Column B	2,450	2,777	
				Column & vs Column C	8,650		*** Pc0.001
Minimum	62.000	64.000	58.000	Column 5 va Column C	6,200	7,027	*** P(0.001
Median (50th percentile)	70.500	68.000	60.000		Mean	101 Card	Lidence Inter
Maximum	74.000	70.000	70.000	Difference	Difference	Free	To
Normality test KS	0.1715	0.1945	0.2819	Column A - Column B	2,450	0.1735	4.626
Normality test P value	>0.10	0.0458	0.0002	Column & - Column C	8,680	6.474	10.826
Passed normality test?	Yes	No	No	Column B - Column C	6.200	4.024	0.376

## DBP at 20hr

	Group A	Group B	Group C	Bonferroni Multiple Comparisons Test
Col. title				If the value of t is greater than 2.667 then the P value is less
Mean	82.1	69.2	68.7	than 0.05.
Standard deviation (SD)	6.703	2.093	4.014	
Sample size (N)	20	20	20	
	1			Hean
Std. error of mean(SEM)	1.499	0.4679	0.8977	Comparison Difference U P value
Lower 95% conf. limit	78.963	68.221	65.821	
Upper 95% conf. limit	85.237	70.179	70.579	Column & vs Column 5 12,900 8.735 *** P<0.00
	1			Column & vs Column C 19.400 9.073 *** Pc0.001
Minimum	68.000	64.000	60.000	Column B vs Column C 0.5000 0.3386 ms P>0.0
Median (50th percentile)	82.000	70.000	70.000	Nean 93% Confidence Inte
Maximum	92.000	72.000	76.000	Difference Difference From To
Normality test KS	0.1770	0.2989	0.1770	Column & - Column B 12,900 9.257 16.543
Normality test P value	×0.10	<0.0001	>0.10	Column & - Column C 13.400 9.757 17.043
Passed normality test?	Yes	No	Yes	Column B - Column C 0.5000 -3.143 4.143

#### DBP at 24hr

	Group A	Group B	Group C	Bonferroni Haltiple Comparisons Test
Col. title				If the value of t is greater than 2.467 then the P value is less
Mean	81.6	66.1	65.7	than 0.05.
Standard deviation (SD)	5.642	2.198	3.197	
Sample size (N)	20	20	20	liten
				Comparison Difference t P value
Std. error of mean(SEM)	1.262	0.4915	0.7149	
Lower 95% conf. limit	78.960	65.071	64.204	Column & vs Column B 15,500 12,399 *** Pc0.001
Upper 95% conf. limit	84.240	67.129	67.196	Column & vs Column C 15,900 12,719 *** Pc0.001
				Column B vs Column C 0.4000 0.3200 ns \$50.05
Minimum	70.000	62.000	60.000	
Median (50th percentile)	81.000	66.000	65.000	Nean 95% Confidence Inter
Maximum	90.000	70.000	72.000	Difference Difference From To
Normality test KS	0.1884	0.2819	0.2374	Column A - Column B 15.500 12.416 18.584
Normality test P value	0.0611	0.0002	0.0044	Column A - Column C 16.900 12.816 18.984
Passed normality test?	Yes	No	No	Column B - Column C 0.4000 -2.654 3.454

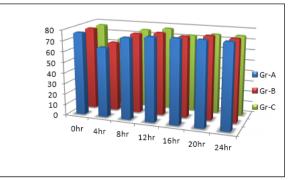
The diastolic blood pressure in clonidine Groups (Group-B, Group-C) were lower compared to control Group (Group-A) at all sampling intervals during post operative period. The difference was statistically significant at 16hr, 20hr and 24hrs.

The difference of diastolic blood pressure between clonidine Groups (Group-B and Group-C) were not statistically significant.

TABLE – VII - HEART RA
------------------------

HR	0hr	4hr	8hr	12hr	16hr	20hr	24hr
Group- A	77.25	65	75.5	77.65	77.9	78.1	78
Group- B	77.45	65.3	75.2	77.3	75.5	77.6	76.45
Group- C	77.2	65.25	75	77.55	72.4	74.7	74.9





The heart rate in clonidine Groups (Group-B, Group-C) were lower compared to control Group (Group-A) at all sampling intervals during post operative period. The difference of heart rate between clonidine Groups (Group-B and Group-C), between clonidine and control

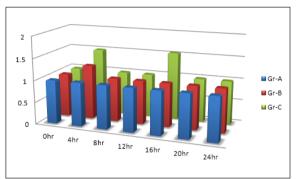
#### Volume : 6 | Issue : 3 | March 2016 | ISSN - 2249-555X | IF : 3.919 | IC Value : 74.50

Groups were not statistically significant.

## SEDATION Table- VIII (mean of sedation scores)

SS	0hr	4hr	8hr	12hr	16hr	20hr	24hr
Group-A	1	1	1	1	1	1	1
Group-B	1	1.25	1	1	1	1	1
Group-C	1	1.5	1	1	1.55	1	1

Graph-9:- comparison of sedation scores between the Groups



10 patients in Group-C and 5 patients in Group-B have higher (score 2) sedation scores at 4hr and 16hr, but the sedation scores were not statistically significant between clonidine groups . Scores were statistically significant between clonidine groups and control group at 4hr. Group-C experienced a higher incidence of clonidine related side effects than groups A &B (p<0.05).

Groups	Count	Sum	Average	Variance		
Column 1	20	20	1	0		
Column 2	20	25	1.25	0.197368		
Column 3	20	30	1.5	0.263158		
Source of Variation	SS	df	MS	F	P-value	F crit
Between Groups	2.5	2	1.25	8.142857	0.000775	3.158843
Within Groups	8.75	57	0.153509			
Total	11.25	59				

# ANOVA (one way analysis of variance) for SEDATION at 4hr

#### Conclusions:-:

All the three groups A, B and C were comparable with respect to age, sex, weight and ASA physical status.

Base line pulse rate, systolic blood pressure and diastolic blood pressure were comparable in all the three groups.

Intraoperative and postoperative hemodynamics in clonidine groups (groups B and C) are more stable when compared to group A.

Visual analog scale (VAS) pain scores for the group A were significantly higher at 4 hr, 12 hr and 16 hr than those for

#### Volume : 6 | Issue : 3 | March 2016 | ISSN - 2249-555X | IF : 3.919 | IC Value : 74.50

the clonidine groups (group B and group C).

Total volume of analgesic solution consumed by the group A (67  $\pm$  4.58 ml) was significantly higher than the clonidine groups (group B 50  $\pm$  4.47ml and Group-C 43.5  $\pm$  4.769 ml)

**Recommendations:-** Clonidine at a dose of 75  $\mu$ g or 100  $\mu$ g is a efficient adjuvant to epidural bupivacaine and morphine for the management of post operative pain after lower limb surgeries because of its ability to produce hemodynamic stability and less complications.

#### Acknowledgement

I thank all my seniors, colleagues and students for helping me throughout the study. I thank my orthopedic consultants for supporting me and definitely the patients for supporting me to continue with the study.

#### References

- Albert Schweitzer .On The Edge Of The Primeval Forest and More From the Primeval Forest 1948.
- Bonica J. Postoperative pain. In: Bonica J, ed. The management of pain. 2nd ed. Philadelphia: Lea & Febiger, 1990:461–80.
- Shoji H, Solomonow M, Yoshino S, et al. Factors affecting postoperative flexion in total
- 4. knee arthroplasty. Orthopedics 1990;13:643-9.
- 5. Kissin I. Preemptive analgesia. Anesthesiology 2000;93:1138-43.
- Buggy DJ, Hall NA, Shah J, et al. Motor block during patient controlled epidural analgesia with ropivacaine or ropivacaine/ fentanyl after intrathecal bupivacaine for caesarean section. Br JAnaesth 2000;85:468–70.
- Standl TG, Horn E, Luckmann M, et al. Subarachnoid sufentanil for early postoperative pain management in orthopedic patients: a placebo-controlled, double-blind study using spinal microcatheters. Anesthesiology 2001;94:230–8
- Eisenach J, Detweiler D, Hood DD. Hemodynamic and analgesic actions of epidurally administered clonidine. Anesthesiology 1993;78:277–87
- Eisenach J, Hood DD, Tuttle R, et al. Computer controlled epidural infusion to targeted CSF concentrations in humans: clonidine. Anesthesiology 1995; 83:33–47.
- Eisenach J, Dupen S, Dubois M, et al. Epidural Clonidine Study Group: epidural clonidine analgesia for intractable cancer pain. Pain 1995; 61:391–9.
- Eisenach J, De kock M, Klimscha W. alpha2-Adrengeric agonists for regional anesthesia: a clinical review of clonidine (1984 –1995). Anesthesiology 1996;85:655–74
- Eisenach JC, Hood DD, Curry R. Intrathecal, but not intravenous, clonidine reduces experimental thermal or capsaicin induced pain and hyperalgesia in normal volunteers. Anesth Analg 1998; 87:591–6.
- Chiari A, Lorber C, Eisenach JC, et al. Analgesic and hemodynamic effects of intrathecal clonidine as the sole analgesic agent during first stage of labor: a dose-response study. Anesthesiology 1999;91:388–96.
- Filos KS, Goudas LC, Patroni O, Polyzou V. Hemodynamic and analgesic profile after intrathecal clonidine in humans: a dose response study. Anesthesiology 1994;81:591–601
- Eisenach JC, De Kock M, Klimscha W. Alpha (2)-adrenergic agonists for regional anesthesia: a clinical review of clonidine (1984–1995). Anesthesiology 1996; 85:655–74.
- 15. Filos K, Goudas L, Patroni O, Polyzou V. Hemodynamic and analgesic profile after intrathecal clonidine in humans. Anesthesiology 1994; 81:591–601
- Bailey P, Rhondeau S, Schafer P, et al. Dose response pharmacology of intrathecal morphine in human volunteers. Anesthesiology1993;79:49–59
- Motsch J, Graber, Ludwig k Addition of clonidine enhances postoperative analgesia from epidural morphine: a double blind study Anesthesiology 1990;73:1067-73.
- Mogensen T, Eliasen K, Ejlersen E, et al. Epidural clonidine enhances postoperative analgesia from a combined low-dose epidural bupivacaine and morphine regimen. Anesth Analg1992; 75:607-610.

- Marjan Jahangiri, JW P Bradley, Dark et al prevention of phantom pain after major lower limb amputation by epidural infusion of diamorphine, clonidine and bupivacaine. Ann R Coll Surg Engl 1994; 76: 324-326
- Michael J. Paech, J. G. Pavy, E. P. Orlikowski Postoperative epidural infusion :A randomized, double-blind, dose-finding trail of clonidine in combination with Bupivacaine and fentanyl (Anesth Analg 1997;84:1323-8)
- Brian D. Sites, Michael Beach, Russell intrathecal clonidine added to Bupivacaine-Morphine Spinal anesthetic improves postoperative analgesia for Total knee arthroplasty (Anesth Analg 2003;96:1083–8)
- Marjan Jahangiri,C H Dark,A P Jayatunga . Prevention of phantom pain after major lower limb amputation by epidural infusion of diamorphine, clonidine and bupivacaine. Ann R Coll Surg Engl 1994; 76: 324-326
- B.S.Sethy, Mary Samuel, Deepak .Efficacy of analgesic effects of low dose intrathecal clonidine as adjuvant to bupivacaine. Indian journal of anaesthesia 2007;51(5):415-419
- Yuan –shiou Huang, Liu-Chi Lin,Michael J. Epidural clonidine for postoperative pain after total knee arthroplasty: A dose response study Anesth Analg 2007;104:1230-5
- Gutierrez A: Anaesthesia extra dural. Buenos Aires Rev. Cirug; 18 : 52, 1939.
- 27. Harrison GR and Clowes N : The depth of the lumbar epidural space from the skin. Anesthesia, 40 : 685, 1985.
- Palmer SK, Abram SE and Maitra AM et al : Distance from the skin to the lumbar epidural space in an obstetric population. *Anaesth. Analg*, 62 : 944, 1983.
- Gossch M (ed): Gray's Anatomy of the Human body, 29<sup>th</sup> ed. Philadelphia, Lea and febiger, 23: 880, 1973.
- Cheng PA: The anatomical and clinical aspects of epidural anesthesia. Anaesth. Analg, 42: 398, 1963.
- Lund PC, Cwik JC and Quinn JR: Experiences with epidural anesthesia. Anaesth Analg, 40: 164, 1961.
- Dawkins M: Identification of anaesthesia in etarmerica peridural. Rev. Cir Buenos Aires, 11: 665, 1932
- Gutierrez A: Anaesthesia in etarmerica peridural. Rev. Cir. Buenos Aires 11: 665, 1932.
- Gutierrez A: Valor de la aspiracion liquida en el espacio peridural en la anaestesia peri dural *Rev. de. Cir. de Buenos Aires*, 12: 225, 1933.
- Odom CB : A review of pages epidural anaesthesia with a report of one hundred cases. New Orleans Med. Surg. J, 88: 618, 1936.
- Brooks AM and Debour SB: Anaesthesia in local, regional and spinal. Anesthesiology, 40: 79-81, 1939.
- Zelenka L: U-Tube and balloon indicator. A new indicator for spinal epidural analgesia. Anesthesiology, 17: 210, 1956.
- Dogliotti AM and Debour SB: Narcosis in local, regional and spinal. Anaesthesia, 42: 80-85, 1939.
- Lund PC, Cwik JC and Magoswer R: Peridural Anaesthesia in general surgery Modified pressure technic . Anesthesiology, 17: 605, 1956.
- Brunner C, and Ikle A : Beitrag Zur peridural anesthesia schweir med. wochensche, 79 : 799, 1949.
- Ikle A : Die peridural anesthesia inder Geburtshilfe . Anesthetist, 2 : 29, 1953.
- 42. De souse E : Contribuicao a technical de puncae extradural. *Rev Bras. Cirur*, 12: 363, **1943**.
- Castonos CC: Origin of the balloon technique as epidural indicator. Anesthesiology, 52: 193, 1980.
- 44. Macintosh RR : Extradural space indicator. Anesthesia, 5: 98, 1950.
- 45. Brooks W: An epidural indicator. Anesthesia, 12: 227, **1957**.
- 46. Dawkins CJM: A drip epidural indicator. Anesthesia, 16: 102, 1961.
- 47. Heldt TJ and Moloney J C : Negative pressure in the Epidural space. A.M.J. Med Sci, 175: 371, **1928**.
- Eaton LM: Observation on negative pressure in epidural space. Proc. staff meets. Mayo Clinic, 14: 566, 1939.
- 49. Lawrence ED : Spinal epidural block. Anesthesiology, 9: 601, 1948.
- Johnston GM, Rodgers RC and Tunstall ME: Alteration of maternal posture and its immediate effect on epidural pressure. *Anesthesia*, 44: 750, 1989.
- 51. Telford RJ and Hollway TE: Observations on deliberate dural puncture

#### Volume : 6 | Issue : 3 | March 2016 | ISSN - 2249-555X | IF : 3.919 | IC Value : 74.50

with a tuohy needle, Pressure measurements. Anesthesia, 46: 725, **1991**.

- Macintosh RR and Mushin WW: Observation on the epidural space. Anesthesia, 2: 100, 1947.
- 53. Brice Smith R: Pressures in the extra dural space. *Anesthesia*, 5: 213, **1950**.
- Foldes FF, Colavincenzo JW and Birch JH: Epidural Anesthesia: A reappraisal. Anesth. Analg (Cleve), 35: 33, 1956.
- Bromage PR: The physiology and pharmacology of epidural blockade. Chapter 3 in regional anesthesia, Bonica JS(ed) et al. Clin Anaes, Philadelphia. F.A. Davis 2: 45, 1969.
- Macintosh RR and Mushin WW: Observation on the epidural space. Anesthesia, 2: 100, 1947.
- Bromage PR: The physiology and pharmacology of epidural blockade. Chapter 3 in regional anesthesia. Bonica. J S (ed) et al. Clin Anaes, 2: 45, 1978.
- Mather LE, Tucker GT and Murphy TM et al: The effects of adding adrenaline to etomidate and lignocaine in extradural anesthesia. ii) pharmaco kinetics Br. J. Anesth, 48: 989, 1976.
- Cheng P: Epidural space, anatomical and clinical aspects. Anesth. Anal (cleve) 42; 407, 1963.
- Bromage PR: Spread of analgesic solutions in the epidural space and site of action. Br. J. Anaesth, 34: 161, 1962.
- Strichartz GR: The inhibition of sodium currents in myelinated nerve by quarternary derivatives of lidocaine. J. Gen. Physiol, 62: 37 -57, 1973.
- Corke BC, Datta S, Ostheimer GW, Wiess JB and Alper M H : Spinal anaesthesia for cesarian section, the influence of hypotension on neonatal out come. Anesthesia, 37: 658, 1982.
- Snider SM, Wright RG and Levinson G et al: Uterine blood flow and plasma norepinephrine changes during maternal stress in the pregnant ewe. Anesthesiology, 50: 524-527, 1979.
- Farag E, Dilger J, Brooks P, Tetzlaff JE. Epidural analgesia improves early rehabilitation after total knee replacement. J Clin Anesth 2005;17:281–5
- Forster JG, Rosenberg PH. Small dose of clonidine mixed with low-dose ropivacaine and fentanyl for epidural analgesia after total knee arthroplasty. Br J Anaesth 2004; 93:670 –7.
- Solomon RE, Gebhart GF. Synergistic antinociceptive interactions among drugs administered to the spinal cord. Anesth Analg 1994; 78:1164 –72.
- Bernard JM, Hommeril JL, Passuti N, Pinaud M. Postoperative analgesia by intravenous clonidine. Anesthesiology 1991;75:577–82
- Duggan AW, Morton CR. Tonic descending inhibition and spinal nociceptive transmission. Prog Brain Res 1988;77:193–211
- Ossipov MH, Suarez LJ, Spaulding TC. Antinociceptive interactions between alpha 2-adrenergic and opiate agonists at the spinal level in rodents. Anesth Analg 1989; 68:194 –200.
- Mogensen T, Eliasen K, Ejlersen E, et al. Epidural clonidine enhances postoperative analgesia from a combined low-dose epidural bupivacaine and morphine regimen. Anesth Analg 1992; 75:607–10.
- Paech MJ, Pavy TJ, Orlikowski CE, et al. Postoperative epidural infusion: a randomized, double-blind, dose-finding trial of clonidine in combination with bupivacaine and fentanyl. Anesth Analg 1997; 84:1323–8.
- Eisenach JC, De Kock M, Klimscha W. Alpha (2)-adrenergic agonists for regional anesthesia. A clinical review of clonidine (1984 –1995). Anesthesiology 1996; 85:655–74.
- Armand S, Langlade A, Boutros A, et al. Meta-analysis of the efficacy of extradural clonidine to relieve postoperative pain: an impossible task. Br J Anaesth 1998; 81:126 –34.
- Sveticic G, Gentilini A, Eichenberger U, et al. Combinations of bupivacaine, fentanyl, and clonidine for lumbar epidural postoperative analgesia: a novel optimization procedure. Anesthesiology 2004; 101:1381– 93.
- Moss J, Glick D. The Autonomic Nervous System.In:Miller RD Editor. Miller's Anesthesia. 6th Ed.Philadelphia:Elsevier Churchill Livingstone 2005:617-77.
- Stoelting RK, Hillier SC. Editors. Antihypertensive Drugs. In: Pharmacology& Physiology in Anesthetic Practice, <sub>4</sub>11 Ed. Philadelphia: Lippincott Williams& Wilkins 2006: 338-51

- Davis M, Gendelman DS, Tischler MD, Gendelman PMI. A primary acoustic startle circuit: Lesion and stimulation studies. Journal of Neuroscience 1982; 6:791-05.
- De Vos H. Bricca G. De Keyser J. Dc Backer J P Bousquet P. Vauquclin G Imidazoline receptors, non-adrenergic idazoxan bindin\_g sites and alpha 2-adrenoceptors in the human central nervous system. Neuroscience 1994:59:589-98
- Hamilton C A. The role of imidazoline receptors in blood pressure regulation. Pharmacol Ther 1992; 54:231
- Guyenet P G, Cabot G B. Inhibition of sympathetic preganglionic neurons by catecholamines and clonidine: Mediation by an alpha adrenergic receptor. Neurosci 1981:1:908
- Eisenach JC, DeKock M, Klimscha W. Alpha-2 adrenergic agonist for regional anesthesia. A clinical review of clonidine (1984-1995). Anesthesiology 1996;85:655-74
- Aho M. Erkola O. Korttila K. Alpha-2 Adrenergic agonists in anaesthesia. Curr Opin Anesthesiol 1992;5:481
- Metz SA, Halter JB. Robertson RP. Induction of defective insulin secretion and impaired glucose tolerance by clonidine. Selective stimulation of metabolic alpha adrenergic pathways. Diabetes 1978;27:554
- Ghignone M, Calvillo O, Quintin L. Anesthesia and hypertension: the effect of clonidine on perioperative hemodynamics and isoflurane requirements. Anesthesiology 1987:67:3.