COMPARISON OF CLONIDINE AND ADRENALINE AS ADJUVANTS TO BUPIVACAINE IN SUPRACLAVICULAR APPROACH TO BRACHIAL PLEXUS BLOCK

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ABSTRACT

BACKGROUND
Aims and Objectives - In our study we compared the clonidine and adrenaline as adjuvants to bupivacaine in supravacular approach to brachial plexus block to prolong the duration of anaesthesia.

MATERIALS AND METHODS
It was a prospective, randomised, double-blinded study with eighty patients. The randomly allocated Group A [n = 40] received 40 mL of 0.25% Bupivacaine + 200 micrograms of adrenaline and Group B [n = 40] received 40 mL of 0.25% bupivacaine + 150 micrograms of clonidine in supravacular block. The onset of analgesia and motor blockade, duration of surgery, duration of motor blockade and analgesia, VAS score, vital parameters and sedation score were recorded, tabulated and analysed.

RESULTS
No difference in the onset of sensory and motor blockade in two groups. Duration of motor blockade and duration of absolute pain free period and duration of post-operative analgesia significantly prolong in clonidine group. No significant side effects were observed in either of the two groups.

CONCLUSION
Thus, we concluded that addition of clonidine to bupivacaine in supravacular approach to brachial plexus block significantly prolong the sensory and motor blockade with no side effects.

KEYWORDS
Clonidine, Adrenaline, Bupivacaine, Brachial Plexus, Supraclavicular Block


MATERIALS AND METHODS
The present study was conducted at Melmaruvathur Adhiparasakthi Institute of Medical Sciences and Research involving 80 patients. Ethical Committee approval was obtained for this prospective, randomised, controlled study. These patients were divided into two groups; 40 patients were assigned to clonidine group and 40 patients to adrenaline group after proper randomisation.

Inclusion Criteria
1. Age more than 20 years.
2. ASA I and II.
3. Upper limb surgeries.

Exclusion Criteria
1. Age less than 20 years.
2. ASA III and IV.
3. Drug allergy to local anaesthetics.
4. Patients in whom paraesthesia not elicited.
5. Failure of blockade.
6. Unwilling patients.

After obtaining informed consent, no pre-anaesthetic medications were given. IV line secured. Brachial plexus block performed by classic method of supravacular approach by paraesthesia technique.

Clonidine Group
40 mL of 0.25% bupivacaine and 150 µg clonidine was given to the patients.
Adrenaline Group
40 mL of 0.25% bupivacaine and 200 µg adrenaline was given to the patients.

Parameters Observed
Onset of Analgesia
Onset of analgesia was taken as abolition of pin prick pain over the distribution of ulnar and median nerve. Patients were assessed every minute.

Onset of Motor Blockade
Onset of motor block was assessed every two minutes after the block using four point scales -
0 - Normal power.
1 - Able to move arm, but weakness.
2 - Fingers can be moved, but not able to move arm.
3 - Complete motor block.

A score of 2 was considered as onset of motor block.

Surgery Duration
Motor Blockade Duration
The time taken for a score of 3 to become 2 score in a four point scale is considered as motor blockade duration.

Duration of Analgesia
Visual Analogue Scale (VAS) was used for pain assessment. VAS has 10 cms length, numbered from 0 to 10. After explaining clearly about the VAS to the patients as 0 - no pain and 10 - the worst possible pain, the patient themselves was asked to score the pain level according to VAS.

Postoperatively, the patients were observed every half an hour till the motor blockade is reversed. After that every 60 minutes (hourly) for 6 hours, 120 minutes (2 hour) for next 6 hours and 24 hours later.

Duration of Absolute Pain Free Period
If the patient did not have any pain in the post-op period is called as absolute pain free period (VAS - 0).

Duration of Postoperative Analgesia
The time period from the onset of analgesia to requirement of analgesia (VAS > 5).

Vital Parameters
1) Pulse rate, 2) Blood pressure, 3) Respiratory rate, monitored periodically.

Sedation Score
Ramsay sedation score
0 - Fully awake.
1 - Drowsy.
2 - Drowsy, but arousable on touch or call.
3 - Drowsy, but arousable on deep stimuli.
4 - Somnolent.

Side effects noted are hypotension and bradycardia. Patients in whom the block was unsuccessful due to total failure of missed dermatomes, which needed intravenous supplementation or general anaesthesia were excluded from the study.

Statistical Tool
Data analysis was done using Epi Info 2008 version. Means, standard deviations, chi square and “p” values were calculated. Yate’s test was used to find the qualitative variables. Kruskal-Wallis chi-square test was used with “p” less than 0.05 is considered significant for quantitative variable.

RESULTS
In our prospective, randomised, controlled study type of surgery, gender, weight and age were similar in both groups (Table 1).

Onset of Sensory Block
In Adrenaline Group, onset of sensory block varied from 6 to 12 minutes with a mean value of 9.17 minutes with a standard deviation of 1.56. In Clonidine Group, it varied from 7 to 12 minutes with a mean value of 8.77 minutes and standard deviation 1.13 (Table 2).

Onset of Motor Blockade
In Adrenaline Group about 10 to 18 minutes of motor block was seen with mean and standard deviation of 14.13 minutes and 1.67, whereas in clonidine the time ranges from 10 to 16 minutes with a mean and standard deviation of 13.78 minutes and 1.27 (Table 2).

Duration of Surgery
In Adrenaline Group, time varied from 90 to 240 minutes with a mean and standard deviation of 107 minutes and 24.4. In Clonidine Group, the time varied from 90 to 150 minutes with a mean and standard deviation of 113.8 minutes and 14.3 (Table 2).

Duration of Motor Blockade
In Clonidine Group, the duration was found to be prolonged to 480 to 620 minutes with a mean and standard deviation of 550.8 minutes and 28.7, but in Adrenaline Group the time was found to be decreased to 250 to 330 minutes with a mean and standard deviation of 280.75 minutes and 18.7 (Table 2).

Duration of Absolute Pain Free Period
In Clonidine Group, the duration was found to be prolonged to 500 to 720 minutes with a mean and standard deviation of 643.8 minutes and 36.6, but in Adrenaline Group the time was found to be decreased to 330 to 480 minutes with a mean and standard deviation of 371.4 minutes and 27.5 (Table 2).

Duration of Postoperative Analgesia
In Clonidine Group the duration was found to be prolonged to 840 to 1080 minutes with a mean and standard deviation of 959.3 minutes and 38.3, but in Adrenaline Group the time was found to be decreased to 480 to 670 minutes with a mean and standard deviation of 564.8 minutes and 24.2 (Table 2).

Sedation Score
In Adrenaline Group it was mean 0.2 ± 0.1, in Clonidine Group it was mean 1.7 ± 0.51 (Table 2).
DISCUSSION

Alpha-2 agonist like clonidine was introduced in the early 1960s as a nasal decongestant. During its use, a nasal decongestant, the antihypertensive property of drug was found out. Subsequently, more insights into the pharmacological properties have led to its use in clinical anaesthetic practice as well.

Clonidine assumes greater importance as anaesthetic adjuvant and analgesic. Sympatholytic is the primary effect. The main action of the clonidine is the decreases in the level of peripheral norepinephrine release by stimulating the prejunctional inhibitory α2 adrenoreceptors. It inhibits central neural transmission in the dorsal horn by presynaptic and postsynaptic mechanism and directly in spinal preganglionic sympathetic neurons. Traditionally, it was used as an antihypertensive drug, but nowadays the uses is mostly based on its analgesic, sedative and anxiolytic properties being developed. The release of endogenous encephalin-like substances was mediated by the peripheral action of clonidine and was first reported by Nakamura et al in 1988.

In 1991, Maze and Tranquil et al reported that alpha-2 adrenergic agonists has an analgesic activity like a potent opioid, is anxiolytic and sedative as benzodiazepine and sympatholytic and its action is reversible. In peripheral nerve blockade, adjuvants added to the local anaesthetics may increase the duration of intraoperative and postoperative anaesthesia. Clonidine has central analgesic action. In addition to its central action, the peripheral antinociception by an α2 adrenoreceptor-mediated local release of encephalin like substances is also seen.

In peripheral nerve and central neuraxial blockade both sensory and motor blockade of local anaesthetics are prolonged by clonidine. Conduction of C and A gamma fibers was blocked by local anaesthetics, which is intensified by clonidine. Clonidine may modify the action of local anaesthetics in sodium channel, either directly or indirectly.

By statistical analysis the duration of surgery, age and weight in the two groups was not found to be statistically significant with a p value of 0.5502, 0.7508 and 0.0755 (p > 0.05). So both groups were comparable.

Onset of Sensory Blockade

Clonidine group had a mean onset of sensory blockade of 8.77 ±1.13 minutes similar to the adrenaline group with 8.77 ± 1.56 minutes. The two groups were found to be statistically insignificant with a p value of 0.2208 (p > 0.05).

Onset of Motor Blockade

Clonidine group had a mean onset of motor blockade of 13.25 ± 1.37 minutes similar to the adrenaline group with 14.1 ± 1.67 minutes. The two groups were found to be statistically insignificant with a p value of 0.0744 (p > 0.05). Similar to the present study, Eledjam JJ et al9 and Popping Daniel M et al10 also concluded that there was no difference in the onset of sensory and motor blockade when clonidine was added to the local anaesthetic solution.

Duration of Motor Blockade

The mean duration of motor blockade in clonidine group was 550.8±28.7 minutes, whereas in the adrenaline group with 564.75±24.2 minutes. The two groups were found to be statistically significant with a p value of 0.0001. Similar to the present study, Eledjam JJ et al9 (580±38.7 minutes, 290.6±34.5 minutes) and Popping Daniel M et al10 also concluded that clonidine prolongs the duration of motor blockade when added to the local anaesthetic solutions.

Duration of Absolute Pain Free Period

Clonidine group had an absolute pain free period of 643.8±36.6 minutes, whereas in the adrenaline group with 371.4±27.5 minutes. The two groups were found to be statistically significant with a p value of 0.0001 (p < 0.05). Similar to the present study, Eledjam JJ et al9 and Popping Daniel M et al10 also concluded that clonidine prolongs the duration of motor blockade when added to the local anaesthetic solutions.

Duration of Postoperative Analgesia

Clonidine group had a duration of post-operative analgesia of 959.3±38.3 minutes, whereas in the adrenaline group with 564.75±24.2 minutes. The two groups were found to be statistically significant with a p value of 0.0001. Similar to the present study, Eledjam JJ et al9 (in clonidine group it was 994.2±34.2 minutes compared to adrenaline group it was 728.3±35.8 minutes) and Popping Daniel M et al10 also concluded that clonidine prolongs the duration of motor blockade when added to the local anaesthetic solutions.

Sedation Group

The sedation score in Group B it was mean 1.72±0.51, in Group A it was mean 0.2±0.1. In clonidine group since the sedation score was not more than 2, the respiratory function was not compromised, so intraoperative sedation is well observed in clonidine group.
Patients were observed for the side effects, such as hypotension and bradycardia. In both groups, there is no incidence of hypotension and bradycardia. No complications related to brachial plexus block were observed.

CONCLUSION
When compared to the adrenaline, clonidine added to local anaesthetic solution in supraclavicular approach to brachial plexus block prolongs the duration of postoperative analgesia and motor blockade.

REFERENCES