Comparision of Bupivacaine and Bupivacaine with Dexamethasone Combination in Brachial plexus Block by Supraclavicular Approach

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Abstract

Introduction: As an alternative to general anesthesia, for upper limb surgeries, brachial plexus block is a popular approach. This type of block avoids the untoward effects of general anesthesia including upper airway instrumentation and thus prevents the consequences as shown by the available literatures. Aim: To compare 0.5% bupivacaine with normal saline and 0.5% bupivacaine with Dexamethasone (8 mg) in brachial plexus block by supraclavicular approach. Materials and Methods: This study was a single blinded, randomized study which was taken up among 70 patients aged between 18 to 65 years of ASA I and II posted for upper limbs. They were randomly divided into two equal groups where first group received 28 ml of Bupivacaine + 2 ml NS and second group received 28 ml of Bupivacaine + 2 ml Dexamethasone (8 mg) by supra clavicular approach. Results: Between the mean ages of two groups, there was no statistically significant difference. In Bupivacaine group, the mean time for onset of sensory block was 17.4 (± 3.5) min and in Bupivacaine dexamethasone group was 11.8 (\pm 2.7) min (p < 0.05). The mean time for onset of motor block in Bupivacaine group was 8.5 (\pm 4.4) min and in Bupivacaine dexamethasone group was 6.4 (\pm 1.8) min (p < 0.05). Both the differences were statistically significant (p < 0.05). The mean duration of sensory block in Bupivacaine group was 2.03 (\pm 1.4) hours and in Bupivacaine dexamethasone group was 5.85 (\pm 0.84) hours (p < 0.05). The mean duration of motor block in Bupivacaine group was 2.48 (± 0.59) hours and in Bupivacaine dexamethasone group was 6.97 (\pm 0.47) hours (p < 0.05). In Bupivacaine dexamethasone group patients required only 1 rescue analgesic dose. Rescue analgesic requirement in Bupivacaine group was higher (p < 0.05). No significant difference in hemodynamic variables i.e., pulse rate, systolic BP, diastolic BP and O, saturation. Conclusion: In brachial plexus block, addition of dexamethasone to bupivacaine produced faster and longer duration of block and less number of rescue analgesics in post-op 24 hours.

Keywords: Bupivacaine; Dexamethasone; Brachial plexus Block.

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Introduction

A popular approach for upper limb surgeries is brachial plexus block as an alternative to general anesthesia. To achieve ideal operating conditions by producing muscular relaxation, maintaining stable intra-operative hemodynamic condition and sympathetic block which reduces post-operative pain, vasospasm and edema, this type of anesthesia is helpful.1 The untoward effects of general anesthesia including upper airway instrumentation is avoided by this type of block and thus prevents

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the consequences. It is effective in terms of cost and performance, margin of safety, along with good post-operative analgesia. Consistent and easiest method for anesthesia and post-operative pain management is supraclavicular block.² Bupivacaine is the local anesthetic used most frequently as it has a duration of action which is longer varying from 3 to 8 hours. Delayed onset, patchy or incomplete analgesia are the limiting factors. Many drugs like Neostigmine, Opioids, Hyaluronidase, Midazolam, Clonidine, Dexamethasone etc., have been added to local anesthetics to improve the quality and duration of action and post-operative analgesia and to minimize these drawbacks.³ The steroids have shown to decrease the inflammation and also have shown analgesic effects. Reduction of inflammation by inhibition of Phospholipase A2 is caused after administration of steroids and also blocks the transmission in nociceptive C-fibers to reduce the pain. Membrane injury and edema by generating inflammatory mediators was induced by phospholipase A². For liberation of arachidonic acid leading to the production of prostaglandins and leukotrienes, phospholipase A2 enzyme is used. Small neurons are sensitized by these enzymes and they enhance pain generation by abnormal conduction and intraneural edema. Dexamethasone is a very potent and selective glucocorticoid. It is used as anti-inflammatory and immunosuppressant. Potency of dexamethasone is about 40 times that of Hydrocortisone. For treatment of many inflammatory and autoimmune conditions, clinical uses of dexamethasone was used but to treat patients suffering from neuropathic pain, Complex Regional Pain Syndromes (CRPS), glucocorticoid are also used. So, steroids have anti-inflammatory as well as analgesic effects.4 Hence, this study was taken up to assess the efficacy of Dexamethasone as an analgesic especially for upper limb surgeries.

Materials and Methods

This study was a single blinded study which was taken up among 70 patients aged between 20 and 68 years undergoing upper limb surgeries in Mamata General and Super Specialty Hospital, Khammam, during October 2013 to September 2015. Ethical clearance was obtained by Institutional Ethical review committee. An informed, written consent was obtained from all the patients.

Inclusion Criteria was that all patients with ASA class I and II, aged between 20 and 68 years posted for upper limb surgeries.

Exclusion Criteria was known case of

hypersensitive reaction to local anesthetic, with abnormal BT, CT or on anticoagulation therapy, severe anemia, hypovolemia, shock, septicemia and h/o seizures, for supraclavicular block, local infection at the site of proposed puncture.

Routine blood and urine investigations done along with Chest X-ray, ECG, HIV, HBsAg screening. Written informed consent was obtained. Intravenous access with a 20 gauge IV cannula on the contralateral upper limb under aseptic techniques. Study consisted of control group (Group I) received 28 ml, bupivacaine 0.5% + 2 ml NS. Study group (Group II) received 28 ml bupivacaine 0.5% + 2 ml Dexamethasone (8 mg). Pre-anesthetic checkup would be done and all the patients were informed about the procedure. IV line was secured and patients would be connected to monitors to record pulse, O₂ saturation, NIBP and ECG. Before the procedure, pre-medication with inj Midazolam 0.05 mg/kg body weight was administered. Patient lies supine, arms by the side and head turned to other side. At a point 1.5 to 2.0 cm posterior, after aseptic preparation of the area, and cephalad to midpoint of clavicle, pulsations of subclavian artery is felt. Cephalo-posterior to the pulsations, a skin wheel is raised with local anesthetic. Until either paresthesia is elicited or first rib is encountered, introduction of a 22 guage, 1.5 inches short beveled needle was done through the same point, parallel to head and neck, in a caudal, slightly medial and posterior direction. The needle would be moved over the first rib until paresthesia is elicited in the arm or hand, if the rib is encountered. The study medication would be injected slowly ruling out intravascular injection intermittently, after eliciting paresthesia and negative aspiration of blood, keeping the needle in the same position. By pin prick method with a 23-gauge needle, sensory block is evaluated. The time between injection and complete loss of pin prick sensation in C₂ and T, dermatome and temperature testing using spirit soaked cotton on skin dermatomes C₂ to T₂ is called as onset time. The time when complete sensory blockade achieved would be noted. The motor block was assessed by using bromage three point score which is 0 = normal motor functionwith full flexion and extension of elbow, wrist and fingers, 1 = decreased motor strength with ability to move fingers and/or wrist only, 2 = complete motor blockade with inability to move fingers. The time of motor blockade was noted. The time elapsed between the injection of drug and complete loss of cold perception of the hand, while onset of the motor blockade was defined as the time elapsed from injection of drug to complete motor block is called the time of onset of sensory block. During the surgery, heart rate, non-invasive blood pressure and oxygen saturation were monitored. The time elapsed between the injection of drug and appearance of pain requiring analgesia is called duration of sensory block and duration of motor block was also recorded.

The data thus obtained was compiled and analyzed using Statistical Package for Social services. (SPSS vs 18). Quantitative data was analyzed by using student 't' test. Qualitative data was analyzed using Chi–square test. A p - value of less than 0.01 was considered as statistically significant.

Results

About 70 patients of ASA I and II who had upper limb surgeries were considered in this study as study subjects. They were randomly divided into two equal groups where first group received 28 ml of Bupivacaine + 2 ml NS and second group received 28 ml of Bupivacaine + 2 ml Dexamethasone (8 mg) by supraclavicular approach, shows in Table 1.

Table 1: Age distribution of the study group

Age groups	Bupivacaine group n (%)	Bupivacaine Dexamethasone group n (%)	Total n (%)
Less than 30 years	10 (28.6)	9 (25.7)	19 (27.1)
31-40 years	13 (37.1)	21 (60.0)	34 (48.6)
41-50 years	8 (22.9)	0	8 (11.4)
51 years and above	4 (11.4)	5 (14.3)	9 (12.9)
Total	35 (100)	35 (100)	70 (100)
Mean ± SD	37.7 ± 12.8	35.6 ± 8.8	37.8 ± 9.7

T value = 0.988 p = 0.419 students unpaired t test NS-not significant

Age groups were similar the groups were comparable in age.

The mean time of onset of sensory block in Bupivacaine group was 17.4 minutes and 11.8 minutes in Bupivacaine – Dexamethasone group. This difference in onset of sensory block was statistically significant between the two groups. The mean time of onset of motor block in this study in Bupivacaine group was $8.5~(\pm 4.4)$ minutes and the mean time of onset of motor block in Bupivacaine – Dexamethasone group was $6.4~(\pm 1.8)$ minutes. This difference in onset of motor block was statistically significant between the two groups, shown in Table 2.

The mean duration of sensory block in Bupivacaine group was 2.03 (± 1.4) hours and in Bupivacaine - Dexamethasone group was 5.85 (± 0.84) hours. There was statistically significant difference in duration of sensory block between Bupivacaine and Bupivacaine - Dexamethasone groups. The mean duration of motor block in Bupivacaine group was 2.48 (± 0.59) hours and the mean duration of motor block in Bupivacaine - Dexamethasone group was 6.97 (± 0.47) hours. There was statistically significant difference in duration of motor block between Bupivacaine and Bupivacaine – Dexamethasone groups. The patients of Bupivacaine group had received 2.5 (± 0.5) doses and the patients of Bupivacaine - Dexamethasone group received 1.4 mean doses of rescue analgesic. The difference in receiving the mean doses of rescue analgesic was statistically significant between the Bupivacaine and Bupivacaine - Dexamethasone groups. The mean heart rate in Bupivacaine group was around 77 to 79 beats per minute. The mean heart rate in Bupivacaine - Dexamethasone group was around 79 to 80 beats per minute. There was no statistically significant difference between Bupivacaine and Bupivacaine - Dexamethasone groups in Heart rate at different time intervals. The mean systolic blood pressure in Bupivacaine

Table 2: Onset of sensory and motor block between the study groups

Groups	Bupivacaine group	Bupivacaine – Dexamethasone group	t value*	p - value, Sig
Onset of sensory block in min Mean ± SD	17.4 (± 3.5)	11.8 (± 2.7)	14.897	0.001, Sig
Onset of motor block in min Mean ± SD	8.5 (± 4.4)	6.4 (± 1.8)	12.669	0.001, Sig

Table 3: Mean time of onset of sensory and motor block in the study groups

Groups	Bupivacaine group	Bupivacaine – Dexamethasone group	t value*	p - value, Sig
Duration of sensory block in hrs Mean ± SD	2.03 (± 1.4)	5.85 (± 0.84)	13.26	0.001
Duration of motor block in hrs Mean ± SD	2.48 (± 0.59)	6.97 (± 0.47)	28.9	0.0001

group ranged from 115.4 ± 6.4 mm of Hg to 115.6 ± 7.1 mm of Hg. The mean systolic blood pressure in Bupivacaine - Dexamethasone group was ranging from $119.8 \pm 10.7 \, mm$ of Hg to $120.3 \pm 12.1 \, mm$ of Hg at different time intervals. There was no statistically significant difference in systolic blood pressure between Bupivacaine and Bupivacaine -Dexamethasone groups at different time intervals. The mean diastolic pressure in Bupivacaine group was ranging from 74.2 \pm 5.8 mm of Hg to 76.6 \pm 7.7 mm of Hg. It was ranging from 77.0 \pm 6.9 mm of Hg to 79.2 ± 7.7 in Bupivacaine – Dexamethasone group at different time intervals. There was no statistically significant difference in diastolic blood pressure between Bupivacaine and Biscayne -Dexamethasone groups at different time intervals.

The oxygen saturation was ranging from 98.7 ± 0.8 percent to 99.9 Percent in Bupivacaine group and it was ranging from 98.6 ± 0.7 percent to 98.9 ± 0.6 percent in Bupivacaine – Dexamethasone groups. The difference between the oxygen saturation was not statistically significant between Bupivacaine and Bupivacaine – Dexamethasone groups, shown in Table 3.

Discussion

For prolongation of an algesic effects, dexame thas one has emerged as a potent corticosteroid when used along with Bupivacaine. However, the studies are scanty to evaluate the efficacy of Bupivacaine alone and when used in combination with corticosteroids like Dexamethasone.⁴ Hence, this study was undertaken to evaluate the efficacy of Bupivacaine with Dexamethasone.

About 37.1% of the patients in Group I and 60.0% of the patients in Group II belonged to 31–40 years age group. The mean age of patients posted was 37.7 ± 12.8 years in Bupivacaine and 35.6 ± 8.8 years in Bupivacaine - Dexamethasone groups. There was no statistically significant difference in age between the two groups. Majority of the patients in this study belonged to 31-40 years in both the groups. In a study by Sheshtha et al. in Nepal in 2003, the mean age was 25.5 ± 12.02 years in local anesthetic group and 28.05 ± 16.1 in Dexamethasone groups.5 In a similar study, the mean age in local anesthetic group was 33.8 years and in Dexamethasone group was 30.3 years.6 In our study, the mean time of onset of sensory block in Bupivacaine group was 17.4 minutes where as it was 11.8 minutes in Bupivacaine - Dexamethasone (study) group. The mean time of onset of sensory block was late in Bupivacaine group compared to

Bupivacaine - Dexamethasone group. Our study is comparable to the study done by Shreshtha et al., in which the mean onset of action was $18.15 \pm$ 4.25 minutes in group which received local anesthetic with adrenaline while it was 14.5 ± 2.1 minutes in the group which received local anesthetic with dexamethasone.5 However, the mean onset of sensory anesthesia was slightly lesser in this study in contrary to findings of Shrestha et al.⁵ In another study, Yadav et al. compared the effect of adrenaline, neostigmine and dexamethasone (three different drugs) by supraclavicular brachial plexus block.3 However, the onset of anesthesia in Dexamethasone group was faster than other two groups of drugs which is comparable to our study. In a study by Islam et al., the onset of sensory block was also lesser in Dexamethasone group than the plain local anesthetic group.⁷ Thus, the observation of our study shows that addition of dexamethasone to bupivacaine decreases the onset time of analgesia. In our study, the mean duration of sensory block in Bupivacaine group was 2.03 (± 1.4) hours and 5.85 (± 0.84) hours in Bupivacaine-Dexamethasone group. A similar study in Nepal found that the duration of action of the local anesthetic as 3.16 hours in local anesthetic group and 12.75 hours in steroid group.8 In a study by Shreshtha et al., the mean duration of post-operative analgesia was around 16 hours in a group who received Bupivacaine with Dexamethasone and its was around 8 hours in Bupivacaine-Tramadol group.5 This shows that the addition of steroid to certainly prolongs the duration of anesthesia and also produces earlier onset of action. This might be due anti-inflammatory effect of Dexamethasone. It has also been proved in many studies that the addition of Dexamethasone to local anesthetic prolongs the duration of action. our study shows similar results when compared with other studies done previously which again proves that dexamethasone prolongs the duration of analgesia. In our study, the mean time of onset of motor block was lesser in Group II *i.e.*, Bupivacaine – Dexamethasone group (6.4 mins) than in Group I *i.e.*, bupivacaine group (8.5 mins). This difference was statistically significant between the two groups. Yadav RK et al. reported a little early onset of motor blockade with dexamethasone. Pathak RG et al. did not find significant reduction of motor blockade time.3 This might be due to usage of adrenalized xylocaine along with bupivacaine and dexamethasone. But they found a significant prolongation of motor blockade with dexamethasone. Shrestha BR et al. (2007) in their study found a significant early onset in motor blockade when dexamethasone was added to a

mixture of 2% lignocaine and 0.5% bupivacaine.5 Our study correlates with the study done by Yadav RK and Shrestha BR.35 Birdar PA et al. (2013) studied the effect of admixture of lidocaine 2% and dexamethasone on onset and duration of sensory and motor blockade.9 They concluded that onset of sensory and motor blockade was more rapid in dexamethasone group. It correlates our study but in which bupivacaine was used instead of lidocaine with dexamethasone. Estebe JP, Le Corre et al. in 2003 stated in their study that dexamethasone alone did not produce any motor block.10 When added to plain bupivacaine without pH correction, complete motor block could not be obtained. With corrected pH, addition of dexamethasone to plain bupivacaine seemed to delay the onset of motor block and did not prolong its duration. In our study correction of pH was not done in both the groups. But the results were encouraging with addition of 8 mg dexamethasone to 28 ml of 0.5% bupivacaine. The mean duration of motor block in Bupivacaine group was 2.48 (± 0.59) hours and in Bupivacaine - Dexamethasone group was 6.97 (± 0.47) hours. There was statistically significant difference in duration of action between Bupivacaine and Bupivacaine - Dexamethasone groups. Tandoc MN et al. did a study in 2011 at University of Buffalo.¹¹ They compared 0.5% bupivacaine with dexamethasone 4 mg and 8 mg as an additive to bupivacaine. Their results showed that duration of motor blockade was significantly prolonged in patients receiving dexamethasone. Pathak RG et al. in 2012 conducted a study with the aim to compare effect of addition of dexamethasone to local anesthetic in brachial plexus block.12 They concluded addition of dexamethasone significantly prolongs the duration of analgesia and motor block. This observation is comparable to our study. Vieira PA et al. (2010) also concluded that addition of dexamethasone prolonged both sensory and motor blockade in interscalene block with bupivacaine.¹³ In our study, the mean numbers of rescue analgesic doses were lesser in Dexamethasone group than Bupivacaine alone group significantly. Our study correlates the study done by Yadav et al. 3 where the mean number of rescue analgesic doses was also lesser in Dexamethasone group than other two groups. Tandoc MN et al. also reported that postoperative analgesic consumption was significantly low in patients who received dexamethasone along with bupivacaine.11 Vieira PA et al. concludes in their study that addition of dexamethasone to bupivacaine-epinephrine-clonidine combination in interscalene block significantly reduces the use of opioids post-operatively.¹³ In our study, the

mean heart rate in Bupivacaine group was around slightly higher in Dexamethasone group than the local anesthetic group. There was no statistically significant difference between the pulse rates of the Dexamethasone group than local anesthetic group. The mean systolic and diastolic pressure was also almost similar in both the groups within normal limits. The mean oxygen saturation also not varied much in both the groups. In summary, the hemodynamic responses are crucial in maintenance of patient during anesthesia. However, the Bupivacaine has already proved its safety especially when used as local anesthetic in supraclavicular block. Since, the hemodynamic responses were similar, the study concludes that the Bupivacaine -Dexamethasone combination is also safer to use in supraclavicular block. The adverse effects were not reported in both the groups in this study.

Conclusion

In brachial plexus block, addition of dexamethasone to bupivacaine produced faster onset of sensory block and motorblock, longer duration of sensory block and motor block. Dexamethasone to bupivacaine has less number of rescue analgesics in post-op 24 hours. No significant difference in hemodynamic variables *i.e.*, pulse rate, systolic BP, diastolic BP and O₂ saturation.

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