Efficacy of dexamethasone as an adjuvant to bupivacaine in supraclavicular brachial plexus block

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ABSTRACT

Introduction: Supraclavicular brachial plexus block provides the optimal operating conditions for upper arm surgery. The study was conducted to assess the efficacy of dexamethasone as an adjuvant to bupivacaine in supraclavicular brachial plexus block. Methods: An experimental study was performed in Gandaki Medical College and Teaching hospital, Pokhara. Forty patients undergoing surgery of the elbow, forearm, and hand under supraclavicular brachial plexus block, with 0.25% bupivacaine with normal saline (n = 20) in the Group I versus 0.25% bupivacaine with dexamethasone (n = 20) in Group II with an equal volume of 30 ml in each group. Randomization was done and the onset of sensory and motor blockade and duration of analgesia was compared in both groups. Statistical package for the social sciences version 24.0 software was used. Continuous variables like the onset of sensory and motor block and time to first rescue analgesia were assessed with the student’s t-test and the chi-square test was applied to assess categorical variables like visual analogue score score. Results: The time to onset of sensory block in the bupivacaine only group was 504.10±70.24 seconds compared to bupivacaine with dexamethasone group (255.65±75.28) seconds. The time to onset of motor block in the bupivacaine-only group was 813.00±74.57 seconds compared to bupivacaine with dexamethasone group (524.25±80.38) seconds. Time to first rescue analgesia in bupivacaine only group was 542.5±126.33 minutes compared to bupivacaine with dexamethasone group (1245±325.66) minutes. The difference was statistically significant in all the findings (p<0.0001). Conclusions: The study concluded that the addition of dexamethasone to bupivacaine prolongs the duration of analgesia and provides faster onset of sensory and motor blocks.

Keywords: Analgesia, brachial plexus block, bupivacaine, dexamethasone.

INTRODUCTION

Regional anesthesia enables site specific, long lasting, and effective anesthesia and analgesia which is suitable for many surgical patients and can improve analgesia. William Stewart Halsted first reported the use of cocaine to block upper extremity nerve in 1884 and performed the first brachial plexus block in 1885. Brachial plexus block is considered as method of obtaining ideal operating conditions for upper limb surgeries providing extended postoperative analgesia with minimal side effects. It also decreases risk of aspiration due to intact pharyngeal and laryngeal reflexes and maintain haemodynamic stability.

Brachial plexus block can be done through various approaches along their course such as supraclavicular, interscalenous, infraclavicular and axillary route. However, for anesthesia and perioperative pain management below shoulder joint surgeries, supraclavicular approach provides most consistent and easiest method compared to other approaches. However, pneumothorax, hemotherax, Horner’s syndrome, phrenic nerve block are the potential complication of Supraclavicular block. Ultrasonography guided supraclavicular brachial plexus block also allows better visualization of underlying...
structures, movement of the needles and thereby making procedure safe and effective.6

Local anesthetics when used alone for supraclavicular brachial plexus block provide good operative conditions but inevitably, the effects dissipate after several hours unmasking the moderate to-severe pain of the surgical insult.7 So, in order to prolong the effect of local anesthetics various adjunct like morphine, tramadol, fentanyl, clonidine, dexametomidine midazolam were also added.8 Recently steroids particularly dexamethasone has been studied as an adjuvant to local anesthetics in brachial plexus block.9 Dexamethasone is a very potent and long acting (t1/2 >36h) glucocorticoid having potent anti-inflammatory and analgesic effects. It has nerve block prolonging effects and also produce analgesia by blocking transmission of nociceptive non-myelinated c-fibers. They might bring about this effect by altering the function of potassium channels in the excitable cell.10 Dexamethasone also causes degree of vasoconstriction to the tissue and local anesthetic will have a slower uptake and absorption, thus prolonging its duration of action.11 The study was conducted with objective to compare the onset of sensory block and motor block along with duration of analgesia between two groups when dexamethasone is added as an adjuvant to bupivacaine in supraclavicular brachial plexus block.

METHODS

This is a quantitative, experimental study performed in Gandaki Medical College from June 2019 to June 2020. Adult patients of either sex aged 18 to 60 years, American Society of Anesthesiologists (ASA) physical status I or II, patient posted for elective orthopedics surgery of elbow, forearm and hand under supraclavicular brachial plexus block were included. Patients with history of hypersensitivity to drugs in study, peptic ulcer diseases, diabetes mellitus, renal failure, hepatic failure, neuromuscular or psychiatric disease, patient receiving psychotropic drug, chronic analgesic therapy Coagulation disorders, pregnancy and lactation, skin lesion at the site of injection, duration of surgery more than three hours, failure to get sensory motor block or need of other modality of anesthesia were excluded from the study.

Pre-anesthetic evaluation was done day before surgery. Written informed consent was obtained from each patient after explaining about the study & anesthetic procedure. VAS scoring system for the assessment of pain consisting of 10 cm line with 0 = no pain and 10 = worst possible pain was explained to the patient. The patients were randomly allocated into two groups of 20 each in which group using lottery system. Group I received 0.25 % bupivacaine with 2 ml normal saline and group II received 0.25 % bupivacaine with 2ml dexamethasone (8 mg) with equal volume of 30 ml in each group. Study drug was prepared by anesthesiologist not involved in the study and anesthesiologist performing block was blinded about the study drug. Heart rate (HR), systolic and diastolic blood pressures, respiratory rate and oxygen saturation (SpO2) was recorded just before the block and at five minutes intervals thereafter. An intravenous drip was started before undertaking the procedure with 18 G cannula which continued throughout the duration of surgery for maintenance fluid.

Patient was kept in supine position with head turned to the opposite side of the block and ipsilateral forearm adducted. Injection site was prepared by aseptic technique with povidine iodine solution. Block was performed with help of 22 G 2" insulated blunt beveled nerve stimulating needle with peripheral nerve stimulator (Stimuplex®; B Braun) and portable ultrasonography machine (Alpinion E-CUBE 7). Portable ultrasound machine with linear superficial probe (4.2 cm) with frequency of 5 to 10 MHz was used. The USG probe was covered with sterile plastic sheath and sterile ultrasound gel was used for better visualization of image and minimization of interference. After identifying brachial plexus a skin wheal was created with lidocaine 1% 2 ml and stimulating needle was advanced by in-plane approach with respect to the probe. In real time vision and with the help of peripheral nerve stimulator, motor response to the electrical stimulus was observed in corresponding area of the hand. The nerve stimulator was set at frequency of 1 Hz with current strength of 2 mA. After obtaining motor response the strength of the current was gradually decreased and if the motor response persisted even at 0.3 to 0.5 mA, local anesthetic was injected under ultrasound vision. Total volume of 30 ml of 0.25% plain bupivacaine with normal saline or dexamethasone was injected increasing with aspiration after each 5 ml of injection. The inter-costobrachial nerve (T2) was also blocked with 5 ml of 2% lignocaine with 1:200000 adrenalin to avoid tourniquet pain.

Sensory block was tested by using pin prick and assessed by three-point scale:

0 = normal sensation,
1 = loss of sensation to pin prick (analgesia),
2 = loss of sensation to touch (anesthesia).

The palmar surfaces of the index and little finger was used to test the median and ulnar nerve in the hand, respectively.
To test the radial nerve the dorsal surface of the thumb was used.

Motor Score (Modified Bromage Scale)12

0 = able to raise the extended arm to 90° for full two seconds
1 = Able to flex the elbow and move the fingers but unable to raise the extended arm
2 = Unable to flex the elbow but able to move fingers
3 = Unable to move the arm, elbow, or fingers

Both sensory and motor block was assessed every five minutes till acceptable sensory and motor blockade of whole arm was achieved or 30 minutes whichever is shorter.

Score 1 for sensory and score 2 for motor block was considered to be adequate for the procession of surgery. The brachial plexus block was supplemented by intravenous sedation (midazolam 0.015 mg/kg intravenously) and repeated in 30 to 60 minutes as needed. In the event of failure to achieve acceptable sensory and/or acceptable motor blockade, the patient was administrated general anesthesia and was excluded from the study. Pain was assessed every 15 minutes during the first hour, every 30 minutes in the next hour and then hourly. After the operation, patient was asked to fill up the visual analogue scale (VAS). Duration of analgesia was considered from the time of onset till the requirement of first rescue analgesia. Rescue analgesic injection sodium diclofenac 75 mg intramuscularly was given when VAS score is ≥5.

Pain was assessed by using VAS score (0= no pain, 5= moderate pain, 10= worst pain possible) every 15 minutes after the surgery in first hour; every 30 minutes during the second hour and hourly thereafter: The time of onset of acceptable sensory block is defined as the interval between the completion of injection and loss of pain in pinprick sensation (Grade 1) on palmar surfaces of the index and little finger dorsal surface of the thumb and at around the injury site of forearm. The onset time of acceptable motor block defined as the time between the completion of the local anesthetic injection and unable to flex the elbow but able to move the fingers (Grade 2).

Heart rate, systolic blood pressure, diastolic blood pressure, SpO₂, was recorded every 5 minutes till the end of the surgery. Sample size was estimated based on a pilot study of 60 patients with 1.96 (95% confidence interval) and 1.64 (power 95%). Total sample calculated was 40. Statistical analysis was performed with statistical package for social science (SPSS) version 24.0. The observations recorded in both groups were tabulated and statistical analysis of demographic data and comparison of groups was carried out using Student’s t-test (paired for intragroup and unpaired for intergroup comparison) and chi-square test. The p-value of <0.05 was taken to be statistically significant.

RESULTS

The mean onset of sensory block was faster in dexamethasone with bupivacaine group (group I) than in bupivacaine with normal saline group (group II). The mean time in group I was 255.65 seconds and in group II was 504.1070.243 seconds which was statistically significant (p<0.0001). (Figure 1)

Figure 1: Onset of sensory block (sec)

The mean onset of motor block was faster in dexamethasone with bupivacaine group (group I) than in bupivacaine with normal saline group (group II). The mean time in group I was 524.2580.382 seconds and in group II was 81374.576 seconds which was statistically significant (p<0.0001). (Figure 2)

Figure 2: Onset of motor block (sec)

The mean time to first rescue analgesia among the two groups were highly variable. The mean time to first rescue analgesia in bupivacaine with dexamethasone group (Group I) was 1245 ± 325.66531 minutes and in bupivacaine with normal saline group (Group II) was 541.5000 ± 126.33602 minutes which was significant statistically (p<0.001). (Figure 3)
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Figure 3: Time for first rescue analgesia

No statistically and clinically significant changes were observed in hemodynamic and respiratory parameters in either group and demographic variables such as age, weight, gender, ASA.

DISCUSSION

Regional anesthesia because of its simplicity of technique, avoidance of airway manipulation and instrumentation, preservation of consciousness and rapid recovery with adequate post-operative analgesia confers advantage over other choices of anesthesia. The brachial plexus consists of injecting local anesthetic agents in fascial plane surrounding the nerve plexus, thereby blocking autonomic, sensory and motor fibers supplying upper extremity.

In our study the patients characteristics (age, weight, sex, ASA) were similar in two groups and not significant. On comparison of heart rate, systolic and diastolic blood pressure in both groups at different intervals, no significant difference was observed. None of the patient had bradycardia, tachycardia, hypertension or hypotension following administration of dexamethasone along with local anesthetic agents. Our finding was similar to that of Baloda et al.,3 Choi et al.,7 and Shrestha et al.9 who also found no significant difference in hemodynamic parameters on addition of dexamethasone.

Our study is comparable with study conducted by Shrestha et al.9 in which they had used a mixture of lidocaine 2% with 1:200,000 adrenaline and bupivacaine 0.5%, dexamethasone was added to the local anaesthetic solution in the steroid group. They demonstrated that there was significant faster onset of action and prolonged duration of analgesia in dexamethasone group than in local anesthetic group. However, compared to our study the duration of onset of sensory block and motor is higher in this study. This could be because in their study position of needle was considered to be acceptable when distal motor response was observed with output current of less than/equal to 0.6 mA whereas in our study position of needle was considered to be acceptable when distal motor response was observed with output current of less than/equal to 0.5 mA under ultrasound vision. So, deposition of local anaesthetic was closer in vicinity of brachial plexus, as well as under ultrasound guidance could be the explanation for quicker onset of sensory and motor block.

Another study done by Pathak et al.,14 patients were divided into two groups in which one group received 1.5% lidocaine + Bupivacaine 0.5% + dexamethasone 8 mg, while another group received 1.5 % lidocaine + 0.5 % bupivacaine + normal saline. They concluded that dexamethasone did not produce significant difference in onset time of sensory and motor block. However, duration of analgesia was significantly prolonged in dexamethasone group compared to normal saline group, which was comparable with our study.

The study conducted by Gandhi et al.15 divided the patients into two groups where one group received 2% lignocaine with adrenaline with 0.5% bupivacaine and another group received added dexamethasone 4 mg. They concluded that there was significantly faster onset of sensory and motor statistically significant, similar to this in our study duration of analgesia was significant in dexamethasone but was higher in our study. This may be due to use of real time ultrasonography with peripheral nerve stimulator in our study.

Our study was also comparable with Baloda et al.,2 they concluded that onset of sensory and motor block were faster in levobupivacaine with dexamethasone group and duration of analgesia was also significantly higher in dexamethasone group compared to levobupivacaine with normal saline group. The results were similar to our study. However, the prolonged onset of mean sensory block and mean onset of motor in this study than in our study may be due to use of paraesthesia technique in this study vs use of real time ultrasonography combined with peripheral nerve stimulator in our study.
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This study still bears some limitations. The limited number of cases were enrolled in the study. Only perineural use of dexamethasone was compared with control group and effect of dexamethasone via other routes were not observed and only immediate post-operative complications following supraclavicular brachial plexus block were observed.

CONCLUSIONS

From our study, we can conclude that the addition of dexamethasone 8 mg to 0.25% bupivacaine in supraclavicular brachial plexus block fastens the onset of action of sensory and motor block and prolongs the duration of analgesia significantly when compared to use of bupivacaine alone.

CONFLICT OF INTEREST: None declared

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REFERENCES


