



RESEARCH ARTICLE

EFFECT OF DEXAMETHASONE AS AN ADJUVANT TO BUPIVACAINE IN ULTRASOUND GUIDED INTERSCALENE BRACHIAL PLEXUS BLOCK IN UPPER ARM SURGERIES

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Abstract

Background: Ultrasound (US) in regional anaesthesia offers a new standard in nerve location and identification, allows real-time imaging of nerves and direct needle guidance. In present study we have access efficacy and safety of Dexamethasone as adjuvant to US guided Interscalene block for Upper arm surgeries.

Aims: The study was conducted to compare the efficacy of Dexamethasone as an adjuvant to Bupivacaine in ultrasound guided interscalene brachial plexus block to provide intraoperative anaesthesia and postoperative analgesia in patients undergoing upper arm surgeries and to evaluate Hemodynamic Stability

Methods: Present study was conducted at tertiary care hospital on 60 patients posted for elective upper arm surgeries, with ASA grade I and II, in age group of 18-70 years. The patients were randomly divided into 2 groups – Group A and Group B. All patients were given ultrasound guided interscalene block with 15 ml of 0.5% Plain Bupivacaine. Inj. Dexamethasone 8mg was added to local anaesthetic drug in Group A and 0.9% normal saline 2ml was added in Group B. All patients received 17 ml of volume of study drug perineurally. All blocks were successful to provide surgical anaesthesia and analgesia. Both the groups were evaluated for onset and duration of sensory and motor blockade, duration of analgesia, perioperative haemodynamic stability, total number of analgesic requests in 24 hours and any adverse reactions/complications.

Results: Both the groups were comparable with respect to patient characteristics. In our study, Onset of sensory block was faster in Group A (3.866 ± 1.33 mins) compared to Group B (7.566 ± 1.67 mins) (p value <0.0001). Onset of motor block was faster in Group A (7.933mins) compared to Group B (11.733 mins) ($p <0.0001$). Duration of mean sensory block was longer in Group A (904 mins) compared to Group B (305.33mins) ($p <0.0001$). Duration of mean motor block was longer in Group A (861 mins) compared to Group B (264.33mins) ($p <0.0001$). Mean Duration of analgesia was longer in Group A (1455.86

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mins) compared to Group B (806.16 mins) ($p < 0.0001$). Total number of analgesic requests was less in Group A (1.2) compared to Group B (2.37) ($p < 0.0001$). Both groups showed perioperative haemodynamic stability. And no adverse reactions or complications were observed in any patient in Group A, whereas 2 patients (6.67%) in Group B had Postoperative Nausea Vomiting.

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Introduction:-

Peripheral nerve block provides good operating conditions, provides excellent perioperative analgesia. They cause the least interference with vital physiological functions of the body, reduction in stress response, systemic analgesic requirements, avoiding polypharmacy, opioid-related side effects and general anaesthesia requirements¹⁹.

Upper arm anaesthesia. The upper extremities receive their innervations from the brachial plexus. Many approaches and techniques are available to block the brachial plexus. It can be blocked through various anatomical approaches such as interscalene, supraclavicular, infraclavicular and axillary approaches⁶.

The Ultrasound-guided regional anaesthesia (UGRA) is a growing area of both clinical and research interest³¹. Ultrasound (US) in regional anaesthesia offers a new standard in nerve location and identification, allows real-time imaging of nerves and direct needle guidance. It has several benefits including faster onset and reduction in the dose of local anaesthetic. Ultrasound-guided needle placement may reduce the risk of complications and increase the accuracy of the block.^(1,5)

Various adjuvant like, dexamethasone, dexmedetomidine, clonidine, magnesium, midazolam, have been used to prolong duration of analgesia of nerve blocks with varying degree of success.

Dexamethasone is a glucocorticoid which are known for their anti-inflammatory, analgesic, immunosuppressive, and antiemetic properties, they exert their action by inhibiting phospholipase A₂, in addition to changes in cell function induced by glucocorticoid receptor activation. Steroids induce a degree of vasoconstriction, acting like epinephrine by decreasing local anaesthetic absorption. Another hypothesis is that dexamethasone may act locally on nociceptive C-fiber to increase the activity of inhibitory potassium channels, thus decreasing their activity and relieving pain.³

Methods:-

After talking approval of IRB, 60 adult patients of upper arm surgeries enrolled in this prospective randomized observational study, after obtaining approval from hospital International Review Board, 60 adult patients with physical status ASA grade I and II aged between 18–70 years, scheduled for upper arm surgeries were included in this study. Study was conducted at Tertiary Care Hospital, during the year of 2019–2021.

Selection of patients:

Inclusion Criteria

Patient aged 18–70 years
Either gender
Having ASA grade I and II
Posted for elective upper arm surgeries

Exclusion Criteria

ASA grade $\geq$ III
, Obese patients, pregnant patients, Patients with distorted neck anatomy

Patients with Mallampati classification Grade 3, Known or suspected coagulopathy

Patients with known hypersensitivity or contraindication to the study drugs, Neurological deficit involving brachial plexus, Infection at site of block, Patient refusal to participate, Patient with chronic pain and using chronic analgesic medications

Method:-

Thorough pre-anaesthetic evaluation was carried out on day before surgery. Patients were explained about the procedure, Informed written consent for procedure was taken from the patient and his/her closer relative. All patients were explained about Visual Analogue scale (VAS) and were made well conversant with it. All patients were advised nil by mouth as per standard fasting guidelines.

Preparation:

An intravenous line was secured and intravenous fluids started. After taking patient in the operation theatre, Pulse oximeter, Non-invasive blood pressure cuff and ECG electrodes were applied and baseline pulse, blood pressure, oxygen saturation and respiratory rate were recorded

Premedication:

All patients were given injection Midazolam i.v. 0.02 mg/kg 10-15 min prior to surgery. The patients were randomly divided into 2 study groups.

Group A:-

Injection Plain Bupivacaine 0.5% - 15cc
Injection Dexamethasone 2ml (8 mg – i.e., 4mg/ml)

GROUP-B :-

Injection Plain Bupivacaine 0.5% - 15cc
Injection 0.9% Normal saline - 2cc.
All patients received 17ml of volume of study drug perineurally.

Under all aseptic precautions, patients in supine position, arms adducted and head turned to opposite side, ultrasound scanning using high frequency linear probe was done. US Transducer position: Transverse on the neck
There are 2 different approaches:¹²

1. Traceback” approach:

Supraclavicular fossa is scanned first to identify the subclavian artery with the nerve trunks which lie posterolateral to the artery. Then the nerve trunks lie in close proximity and are termed as Bunch of Grapes appearing as hypoechoic (black) halos surrounded by hyperechoic (white) thin rim of connective tissue. Move cephalad and trace the bunch of grapes which become arranged in a linear manner between two scalene muscles.

1. Medial to lateral approach:

- Put high frequency linear probe on cricoid cartilage at C6 level.
- Identify trachea, thyroid gland, - Move laterally identifying carotid artery, internal jugular vein, SCM muscle, scalenus anterior and scalenus medius from medial to lateral. Once the groove is identified, the probe adjustments are made for the better visualization of the oval to round anechoic nerve roots of the brachial plexus showing typical “traffic signal” sign.
- Identify vertebral artery in same image posteriorly and medially. Local anaesthetic infiltration 2-3ml at the site of puncture.

Needle:

A 23G 1 inch and half block needle is used.

Needle approach:

In-plane approach

Target:

Needle between the plexus and anterior border of scalenus medius.

Volume:

Group A: Injection Plain Bupivacaine 0.5% - 15cc and Injection Dexamethasone 2ml (8 mg)

Group B: Injection Plain Bupivacaine 0.5% - 15cc and Injection Normal saline - 2cc

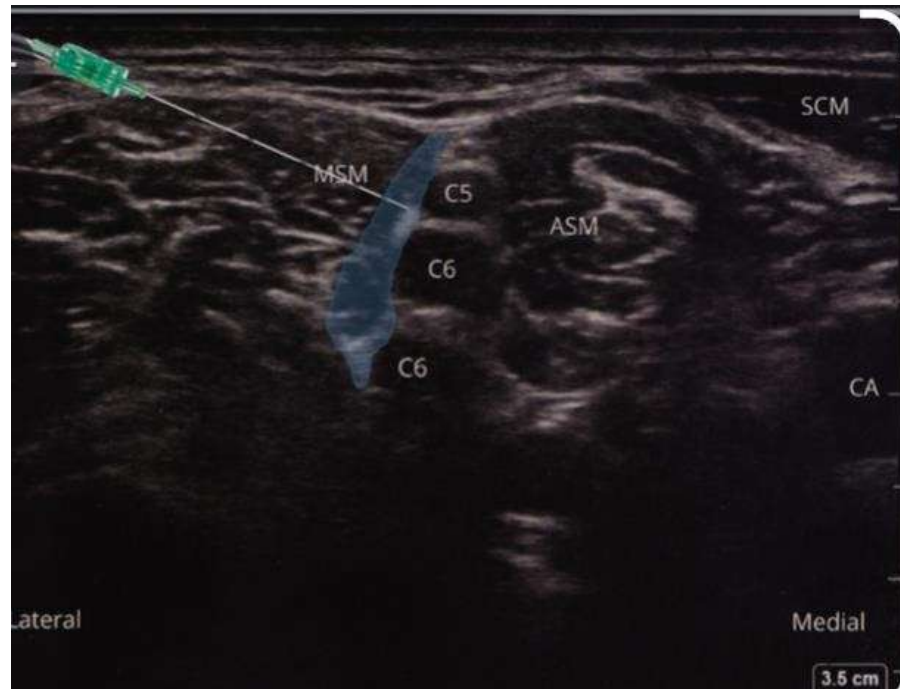


Figure 1:- SCM – Sternocleidomastoid, MSM – Middle Scalene muscle, ASM – Anterior Scalene muscle, CA – Carotid Artery.



Figure 2:- Probe positioning and in-plane approach of needle insertion (22)

Patients were observed for following parameters–

Onset and duration of sensory block
 Onset and duration of motor block
 Perioperative vitals
 Total duration of analgesia
 Complications/Adverse reactions
 Time for first rescue analgesia
 Postoperative analgesic requests in 24 hours

Sensory block:

Sensory block was graded according to Hollmen scale.^(5,40)

- | | | | |
|---------|---|--|------|
| Grade 1 | – | Normal sensation of pinprick | |
| Grade 2 | – | Pinprick felt as sharp pointed but weaker compared | with |
| | | same area in other limbs | |
| Grade 3 | – | Pinprick recognized as touch with blunt object | |
| Grade 4 | – | No perception of pinprick | |

Onset of sensory block was defined as the time between the administration of the drug and establishment of Grade 3 on Hollmen scale.

Motor block:

Motor block was graded according to Hollmen Scale⁽²⁸⁾ Grade 1–Normal muscle function
 Grade 2–Slight weakness in function Grade 3–Very weak muscular action
 Grade 4–Complete loss of muscle action

Onset of motor block was defined as the time between the administration of the drug and establishment of Grade 3 on Hollmen scale. Surgery was allowed to begin after achievement of sensory block grade 3 and motor block grade 3 of Hollmen scale. Vitals at the time of incision were noted.

Success of Block : When Grade 3 of Hollmen Scale was achieved, it was considered a success of block and surgery was allowed to start.

Failure of block was considered if supplementation of any type of general anaesthesia (Inj. Ketamine 2mg/kg/Inj. Propofol 2.5mg/kg) was needed and failure of block was managed by supraglottic airway device insertion (I-gel Insertion).

Intra-operative hemodynamic parameters:

Patients were observed for intra-operative vitals periodically throughout the operation and recorded at the time of block, at the time of incision, then every 5 minutes for 15 minutes after incision, then every 15 minutes for 1 hour and every half an hour thereafter.

Patients received NBM + maintenance solution of Ringer lactate as calculated according to the Holiday Segar formula.

Perioperative blood loss in surgery was managed with crystalloids, blood and blood products. A change of 20% in mean arterial blood pressure from baseline, is defined as Hypertension or Hypotension and corrected pharmacologically. A change of 20% in Heart rate from baseline, is defined as

Tachycardia or Bradycardia (Bradycardia is either $<20\%$ of baseline or <60 beats/min) and corrected pharmacologically.

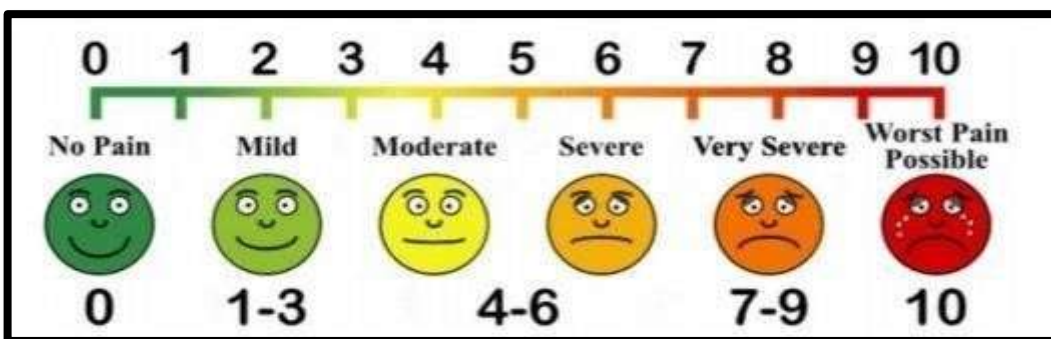
Hypertension and Tachycardia were treated by Inj. Fentanyl 50 mcg intravenously. Hypotension was managed by i.v. fluids and Inj. Mephentermine 6 mg i.v. in all quotes.

Bradycardia was corrected by Inj. Atropine 0.6 mg i.v.

Intraoperative Respiratory depression is defined as $RR < 12/\text{min}$ or $SpO_2 < 90\%$ - Treated with 100% oxygen by mask.

Duration of surgery was noted. It was defined as the time from giving the block till completion of closure of the incision site of surgery. Intraoperatively, patients were observed for adverse reactions of drugs/ complications of procedures and treated according to standard guidelines. Nausea/vomiting: Was assessed by 4 point scale—

VAS(26)



Grade		Treatment
1	No Nausea/vomiting	Ventimask oxygenation and head low position Inj. Ondansetron 4mg i.v.
2	Minor form	
3	Moderate form	
4	Severe form	

For LAST intralipid emulsion was kept ready. If difficulty in breathing occurred, airway assistance was given and oxygenation was provided.

For hoarseness of voice and Horner's Syndrome (ptosis, anhidrosis, miosis, sinking of eyeball into the face, inability to completely close or open the eyelid) patients were reassured and oxygenation with venturimask was given. For Convulsions, Inj. Thiopentone sodium 5-6 mg/kg was kept ready.

After completion of surgery, duration of surgery was noted and patients were shifted to Post Anaesthesia Care Unit (PACU).

Post-operative monitoring:

Postoperative monitoring was done periodically for 24 hours and recorded every hourly till 10 hours then every 2 hourly till 12 hours, then 6 hourly till 24 hours.

Duration of sensory block was defined as the time from onset of sensory block to return of normal sensation from pinprick according to Hollmen scale (Grade 1)

Duration of motor block was defined as the time from onset of motor block to recovery of motor function according to Hollm scale (Grade 1) **Duration of analgesia:** Total duration of analgesia was the summation of

Duration of surgery + time for first rescue analgesia .
Post-operatively patients were assessed according to Visual analogue score

Patients were asked to describe their pain from 0 to 10, where 0 means no pain and 10 means worst pain. Visual Analogue Scale for pain assessment⁵⁰

The duration of analgesia was counted till $VAS \geq 4$. Rescue analgesia was given in form of inj. Tramadol 1 mg/kg intravenously at the VAS score ≥ 4 along with Inj. Ondansetron 4 mg. VAS was observed for 24 hours postoperatively and managed < 4 by analgesic request of Inj. Tramadol 1 mg/kg i.v. Number of analgesic requests in 24 hours were recorded and patients were watched for block related delayed complications like Pneumothorax, Hemidiaphragmatic paralysis and managed by standard protocols. Patients were observed for transient neurological complications in early and late postoperative period for up to 72 hours and also for delay in voiding.

Patients were inquired for any delayed Neurological complications for up to 7 days.

Statistical Analysis

All data were collected in Microsoft Excel Spreadsheet.

Statistical analysis was done by SPSS software.

Descriptive data of both the groups were compared using unpaired 't' tests.

Mean is used to derive the central tendency of the data at a particular time in one study group.

Standard deviation is a measure of dispersion of a set of data from its mean. Categorical data were analysed by Chi Square test.

'P' value < 0.05 was considered statistically Significant (S) and < 0.001 was considered Highly Significant (HS). 'P' value > 0.05 non-significant (NS)

Observation & Results:-

Table 1:- Comparison of Demographic data.

	Group A (N=30)	Group B (N=30)	p value	Inference
Gender				
Male	17 (56.67%)	17 (56.67%)	-	-
Female	13 (43.33%)	13 (43.33%)		
ASA grade (I/II)	10/ 20	17/ 13	-	-
Age (years)	48.46 ± 15.71	43.86 ± 15.69	0.2611	NS
Weight (kgs)	59.3 ± 5.49	60.76 ± 7.09	0.3762	NS
Height (m)	1.71 ± 0.04	1.71 ± 0.03	1.0	NS
BMI (kg/m²)	20.30 ± 2.01	20.61 ± 2.22	0.5729	NS

Duration of surgery (mins)	125.33±17.56	134.33±21.16	0.0782	NS

Table 1 shows various demographic parameters and duration of surgery in each group (P > 0.05)

Table 2:- Types of surgeries.

Type of Surgery	Group A	Group B
Proximal Humerus PHILOS Plating	17	18
Humerus Enders Nail	3	-
Proximal Humerus TENS Nailing	-	1
GT CC Screw	1	1
Distal Humerus CC Screw	-	1
Humerus I/L Nail	1	-
Humerus Shaft Plating	1	2
Distal Humerus Plating	-	2
Humerus MIPPO Plating	-	2
Humerus Ex-fix	3	-
Humerus Enders Nail ROI	1	2
Olecranon TBW + Kwire	1	-
Supraspinatus and Infraspinatus Tear Repair	1	-
Open Supraspinatus Tear Repair	-	1
Shoulder Arthroscopic Repair	1	-

All surgeries in Table 2 were successfully performed under interscalene block without any supplementation of General Anaesthesia.

Table 3:- Mean sensory and motor onset.

Onset (mins) (in minutes)	Group A (N=30)	Group B (N=30)	p value	Inference
Sensory onset (Mean ±SD)	3.866± 1.33	7.566± 1.67	<0.0001	HS
Motor onset (Mean ±SD)	7.933± 1.25	11.733± 2.08	<0.0001	HS

Table 4:- Mean heart rate changes.

Time duration	Group A (N=30)	Group B (N=30)	p value	Inference
	Heart rate (per min) (Mean ±SD)	Heart rate (per min) (mean ±SD)		
Preoperative Mean HR	83.6±11.15	81.3 ± 10.14	0.4067	NS
INTRAOPERATIVE PERIOD				
At the time of induction	81.76 ± 10.89	83.06 ± 11.10	0.6487	NS
At the time of	80.73 ± 10.35	80.2 ± 10.68	0.8459	NS

incision				
15min	80.3 10.43	81.9 11.50	0.5746	NS
30min	80.2 10.57	81.56 10.79	0.6238	NS
45min	80.63 10.23	83.06 11.29	0.3859	NS
60min	80.5 10.28	82.96 10.95	0.3734	NS
75min	80.73 10.35	82.73 11.61	0.4841	NS
90min	80.8 10.37	83.26 11.31	0.3835	NS
105min	81.46 9.51	83.44 11.38	0.4676	NS
120min	82.52 8.80	82.59 11.39	0.9788	NS
135min	82.21 10.31	80.57 10.55	0.5449	NS
150 min	81.42 10.87	79.09 12.93	0.4530	NS
160 min	68.5 0.70	69.25 7.88	0.6056	NS
180 min	68.0 2.82	61.0 1.41	0.0882	NS

Table5:- Variationinmap.

Timeduration	GroupA (N=30)	GroupB (N=30)	pvalue	Inference
	MAP (mmHg) (Mean±SD)	MAP (mmHg) (mean± SD)		
Preoperative MAP	96.04 ± 8.18	95.57 9.62	0.8392	NS
INTRAOPERATIVEPERIOD				
Atthetime of induction	94.46 8.03	96.66 9.02	0.3225	NS
Atthetime of incision	94.0 8.10	94.93 8.59	0.6678	NS
15min	93.0 7.80	95.21 8.53	0.2993	NS
30min	93.84 7.97	94.55 8.53	0.7402	NS
45min	94.08 7.82	94.43 8.10	0.8654	NS
60min	95.0 8.06	94.8 8.5	0.9258	NS
75min	94.0 8.13	93.27 8.07	0.7283	NS
90min	95.06 8.39	95.32 8.83	0.9073	NS
105min	94.49 8.73	95.50 8.85	0.6580	NS
120min	93.0 8.91	95.61 8.77	0.2575	NS
135min	93.0 8.99	93.59 7.76	0.7865	NS
150 min	89.14 5.90	95.42 9.54	0.1399	NS
160 min	89.0 1.41	93.66 9.13	0.2035	NS
180 min	89.0 1.88	95.16 6.83	0.3007	NS

Table 5 shows that there were no statistically significant changes in mean blood pressure in perioperative periods in each group.

Table6:- Duration of motor and sensory block.

Duration of block (mins)	GroupA (N=30)	GroupB (N=30)	pvalue	Inference
Duration of motor block (Mean±SD)	861±104.01	264.33±57.32	<0.0001	HS
Duration of sensory block (Mean±SD)	904±113.88	305.33±63.17	<0.0001	HS

Table7:- Meanpostoperativevas Score.

	GroupA	GroupB
	Mean±SD	Mean±SD
Immediate	0	0
30mins	0	0
60 mins	0	0
120 mins	0	0
180 mins	0	0
240 mins	0	0
360 mins	0	1±0
480 mins	0	1.533±0.68
600 mins	0	3.1±0.84
720 mins	1.26±0.52	4±0
1080 mins	2.56±1.072	0
1440 mins	4±0	0

Table8:- Durationofanalgesia.

	GroupA (N=30)	GroupB (N=30)	pvalue	Inference
Durationof analgesia (minutes) (Mean±SD)	1455.86± 167.25	806.16± 62.47	<0.0001	HS

FromTable9itisevidentthat durationofanalgesia was prolongedinGroupAcomparedtoGroupB(pvalue <0.0001).

Table9:- TotalAnalgesicrequestsin24hours

	GROUPA	GROUPB	Pvalue	Inference
Total number of analgesic requestsin 24hours (MeanSD)	1.2 0.4068	2.370.4901	<0.0001	HS

Table10:- Postoperativecomplications

Complications	GroupA(N=30)	GroupB(N=30)
Tachycardia	Nil	Nil
fentBradycardia	Nil	Nil
Hypotension	Nil	Nil
LAST	Nil	Nil
Convulsion	Nil	Nil
Coughing	Nil	Nil
Nausea/vomiting	Nil	2
LocalHematomaat siteofblock	Nil	Nil
Hoarsenessofvoice	Nil	Nil
Horner'ssyndrome	Nil	Nil

Difficulty in breathing	Nil	Nil
Pneumothorax	Nil	Nil
Hemidiaphragmatic paralysis	Nil	Nil
Neuropathy	Nil	Nil

As illustrated in Table 10 only 2 patients in Group B had PONV Grade 3 (6.67%) which was managed with Inj. Ondansetron 4mg i.v. No patients needed any additional fentanyl supplementation or airway assistance for oxygenation. There were no transient/delayed neurological complications observed in any patient of each group.

Discussion:-

In recent years, there has been a growing interest in peripheral nerve blocks for surgical anaesthesia and analgesia. The development of focal anaesthetic agents with low toxicity and long duration of action had contributed to this change. It is associated with multiple benefits including reduced morbidity and mortality, superior postoperative analgesia, cost-effectiveness, and a low rate of serious complications. However, it is highly dependent on the accurate delivery of a correct dose of focal anaesthetic to attain success and to avoid rare but potentially devastating nerve damage. One of the principle challenges in regional anaesthesia is the unreliability of conventional modalities like electric stimulation and patient-reported paraesthesia for confirming precise nerve localization.

Demographic data:

In our study, Table 1 shows demographic data were comparable in both groups. Badran et al, (2020)⁴ and Jadon et al, (2015)²³ had comparable demographic data and duration of surgery.

Method of Block:

We used Ultrasound technique for giving Interscalene block as by using this technique, we can give lesser volume of drug at perfect site, so very less chances of focal anaesthetic toxicity and transient neurological complications. Drug is injected into the sheath so less chances of nerve damage.

Jadon et al, 2015, (17) had done their study of PNS guided interscalene block for shoulder arthroscopic repair for surgical anaesthesia and analgesia. Badran et al, 2020, had done their study of USG guided interscalene block with Bupivacaine and perineural Dexamethasone along with general anaesthesia for shoulder surgical procedures.

Drug for Interscalene Block

Estimated time of four surgery was approximately 2-3 hours, so we have taken longer acting local anaesthetic drug— Plain Bupivacaine 0.5% Volume

of Drug: We have used Plain Bupivacaine 0.5% 15ml in our study of USG guided Interscalene

Block. Falcao et al, (2012)¹² in their study estimated effective volume of Bupivacaine 0.5% by using the step-up/step-down sequence according to the formula proposed by Dixon, focusing on the analysis of the minimum anaesthetic volume with a 50% probability of effective nerve block. In their study 15ml of 0.5% bupivacaine with epinephrine (1:200,000) was injected, with a direct view of the injection site. The injection was given in two equal aliquots of 7.5ml: one between the upper and middle trunk and one between the middle and lower trunk of the brachial plexus.

Adjuvant :

We have used Dexamethasone 8mg perineurally as adjuvant in GROUP A.

David et al, (2019)¹⁰ in their study observed that i.v. dexamethasone has a direct effect on a blocked nerve, therefore the concentration of steroid reaching that nerve should be very much less than that achieved by placing the local anaesthetic directly onto the nerve. Ke An, Nabil M. El Kassabany, Jiabin Li et al, (2015)²⁷ in their study of comparison of different dose of Dexamethasone 0.14mg/kg and 0.5 mg/kg combined with Bupivacaine for sciatic nerve block demonstrated that Perineural Dexamethasone prevent axonal degeneration and demyelination and no significant caspase dependent apoptosis process in the mouse sciatic nerve among all study

groups in the first study period. Neurotoxicity of plain bupivacaine identified via histomorphology and S-100 expression patterns was no longer significant with the co-administration of preservative-free dexamethasone perineurally. Perineural Dexamethasone prevents the Bupivacaine-induced reversible neurotoxicity and short-term "rebound hyperalgesia" after the resolution of nerve block. In our study, therefore we have taken low dose Dexamethasone as perineural adjuvant.

Type of surgery:

In our study, Table 2 shows various upper arm and shoulder surgeries conducted under US guided interscalene block. Jadon et al, 2015 (17) and Badran et al (3) had used interscalene block for various upper arm surgeries.

Onset of sensory and motor block:

In our study, in Table 3, we observed that in Group A (Bupivacaine 0.5% + Dexamethasone) mean onset of sensory block was 3.866 ± 1.33 min while in Group B (Bupivacaine 0.5% + Normal Saline) it was 7.566 ± 1.67 min ($p < 0.0001$). Mean onset of motor block was faster in Group A 7.933 ± 1.25 min compared to Group B 11.733 ± 2.08 min, which is also statistically highly significant ($p < 0.0001$). From this it is clear that Group A – Dexamethasone group had faster onset of sensory and motor block than Group B without any adjuvant.

In a study conducted by Daret al⁹, who added dexamethasone to ropivacaine in brachial plexus block and observed that the onset time in dexamethasone group for sensory block was 14.65 ± 3.31 min which was significantly shorter than the control group (17.5 ± 4.2 min).

Khalid Ali Khaleel et al¹⁹, in their study of brachial plexus block, the mean time for the onset of sensory block was 7 min in Bupivacaine + Normal Saline Group and 5 min in Bupivacaine + Dexamethasone Group. The onset of motor block was 17.53 ± 1.70 min in Bupivacaine + Normal Saline Group and 14.63 ± 2.79 min in Bupivacaine + Dexamethasone Group.

In a study of Brachial Plexus Block conducted by El-Barade

Elshmaa¹³, they observed the onset time for motor block with bupivacaine and dexamethasone to be 11.4 ± 3.6 min which is comparable to our study.

Jadon et al, (2015)¹⁷ in their study observed that Dexamethasone when mixed with Ropivacaine had no effect on the onset of sensory and motor block.

Duration of sensory and motor block:

In our study, in Table 6 the duration of sensory block was longer in Group A (904 ± 113.88 mins) compared to Group B (305.33 ± 63.17 mins), $p < 0.0001$. Duration of motor block was longer in Group A (Bupivacaine 0.5%

+ Dexamethasone) 861 mins compared to Group B (Bupivacaine 0.5% + Normal Saline) 264.33 min.

($p < 0.0001$). In study of Vieira et al, (2009)⁴⁸ The median sensory block duration was 1457 versus 833 min ($P < 0.0001$) and motor block duration was 1374 versus 827 min ($P < 0.0001$) when compared with the control. Vasconcelos et al, (2020)³² in their study of interscalene block, mean duration of sensory block was 1440 ± 0 minutes in Dexamethasone + Levobupivacaine with vasoconstrictor Group and 1267 ± 164 minutes in Normal Saline + Levobupivacaine with vasoconstrictor Group.

Khalid Ali Khaleel et al¹⁹, in their study of brachial plexus block, the mean duration of sensory block was 289.50 ± 45.71 min in Group BS (Bupivacaine + Normal saline) and 1160 ± 143.10 min in Group BD. This difference was highly statistically significant with $P < 0.0001$. The mean duration of motor block was 216.27 ± 37.73 min in Group BS and 870.87 ± 101.14 min in Group BD. This too was found to be highly statistically significant with $P < 0.0001$.

Choi et al⁶ who conducted a meta-analysis of RCTs and included

801 patients of brachial plexus block. They observed that dexamethasone prolonged the analgesic duration for long-acting local anesthetic from 730 to 1306 min, while motor block was prolonged from

664 to 1102 min without any observed adverse events. Parveen Setal, (2017)²³ in their study of brachial plexus block, mean duration of sensory block was 1085.23 minutes in Dexamethasone group and 322.37 minutes in control group. The mean duration of motor block was 510 minutes in Dexamethasone group and 352.33 minutes in control group.

Duration of analgesia:

In our study, in Table 8 it was observed that the duration of analgesia was longer in Dexamethasone Group A (1455.86 ± 167.25 mins) compared to Group B (806.16 ± 62.47 mins). ($p < 0.0001$).

K.C. Cumming et al (2011)⁷ in their study of interscalene block observed that dexamethasone significantly prolonged the duration of analgesia of both ropivacaine and bupivacaine. Mean duration of analgesia in Bupivacaine with Dexamethasone group was 1344 minutes and Bupivacaine with Normal Saline group was 888 minutes. This study results are comparable to our study. Jadonet al, (2015)²³ in their study of interscalene block, mean duration of analgesia was 1103.72 $\pm$ 96.02 minutes in the Dexamethasone + Ropivacaine group and 551.54 $\pm$ 166.92 minutes in the control group. Rosenfeld et al, (2016)³⁸ in their study of perineural versus intravenous dexamethasone as adjunct to local anaesthetics in interscalene block, it increases duration of analgesia.

Tandoc et al, (2011)²⁴ in their study of interscalene block, mean duration of analgesia was 1408 $\pm$ 158 minutes in Bupivacaine + Dexamethasone group and 798 $\pm$ 60 minutes in control group. This study results are comparable to our study. Viera et al, (2010)⁴⁸ in their study of interscalene block, mean duration of analgesia was 1457 $\pm$ 434 minutes in Bupivacaine + Dexamethasone group and 833 $\pm$ 267 minutes in control group. This study results are comparable to our study. Desmet et al, (2013)¹¹ in their study of interscalene block, mean duration of analgesia was 1433 $\pm$ 619 minutes in Ropivacaine + Dexamethasone group and 824 $\pm$ 510 in control group. This study results are comparable to our study. D. Holland et al, (2018)⁸ in their study of effect of Dexamethasone route and dose on the duration of interscalene block, mean duration of analgesia was 1524 minutes in Bupivacaine + 4mg Dexamethasone given perineurally, 1632 minutes in Bupivacaine + 8mg Dexamethasone given perineurally, 1440 minutes in Bupivacaine (perineural) + 4mg Dexamethasone given intravenously, 1488 minutes in Bupivacaine (perineural) + 8mg Dexamethasone given intravenously.

Perioperative Haemodynamic Parameters:

In our study, intraoperative haemodynamic parameters were stable and comparable in both the groups. It is evident from Table 4, Table 5. Postoperative haemodynamic parameters were stable and comparable in both the groups. It is evident from Table 4, & 5. This may be due to successful block and adequate intraoperative anaesthesia and analgesia. Also, postoperatively VAS was managed by rescue analgesic requests. Badran et al, 2020³, in their study of interscalene block, as regards hemodynamics, they were stable all through surgery and postoperatively in the two groups. They noticed an increase in the HR and MAP in the Control group higher than in the Dexamethasone group at 12h and 24h postoperatively:

Total Analgesic requests in 24 hours:

In our study, in Table 9 it was observed that total number of analgesic requests in Group A with Dexamethasone was 1.2

0.4068 and in Group B was 2.370.4901. ($p < 0.0001$). Thus, need for postoperative analgesia in Group A with adjuvant Dexamethasone is very less compared to Group B without adjuvant.

Zorilla Vecca et al (26) have proved that perineurally given dexamethasone is better adjuvant than IV Dexamethasone.

Tandoc et al (24) showed that prolonged Analgesia in Dexamethasone as adjuvant for US guided Interscalene Block.

Complications:

As evident from Table 10, no complications or adverse effects were noted in Group A. 2 patients (6.67%) in Group B had postoperative nausea and vomiting as side effects. No PONV was observed in Group A; this could be attributed to the antiemetic properties of Dexamethasone. No patients needed any additional airway assistance or oxygenation. There were no delayed neurological complications observed in any patient of each group in our study. Badran et al, (2020)⁴ in their study of interscalene block, there were no dexamethasone-specific adverse effects; however, they noticed that postoperative nausea and vomiting were more common within the Control group (7 out of 23 patients), when compared to the Dexamethasone group (1 out of 22 patients), which can be attributed to the antiemetic effect of dexamethasone. As regards the interscalene block side effects, they had used 30 ml of 0.5% Bupivacaine along with dexamethasone 8 mg perineurally under US guided block. Horner syndrome was the most common and was observed in 5 patients out of 23 (21.7%) in the Control group and 4 patients out of 22 (18.2%) in the Dexamethasone group. Phrenic nerve affection in the form of diaphragmatic paresis was observed in 2 patients out of 23 (8.7%) in the Control group and in 3 patients out of 22 patients (13.6%) in the Dexamethasone group, which was ipsilateral without breathing difficulty. There were no pneumothorax cases found. Jadon et al, 2015¹⁷ also used 30 ml of Ropivacaine 0.5% and observed Horner syndrome in 11/50 (20.5%) in the dexamethasone-ropivacaine group and 15/50 (23.5%) in the ropivacaine-only group, but diaphragmatic paresis was observed in 10 out of 50 patients (20%) in the dexamethasone group and in 8 out of 50 patients (16%) in the ropivacaine-only group. Two patients (2/50) in the dexamethasone-ropivacaine group and 1 (1/50) in the ropivacaine group had external jugular vein puncture, which was managed with external compression and did not require any intervention.

K.C. Cumming et al, (2011)⁷ in their study of interscalene block, no adverse events were detected in 108 patients given dexamethasone, the

95% CI for neurotoxicity was 0–3%. Desmet et al, (2013)¹¹ observed 20.5% Horner Syndrome in patients who received ropivacaine only and 23.5% in patients who received ropivacaine and dexamethasone mixture in interscalene block.

Khan Alam Noor et al, (2016)²⁰ in their study of interscalene block, one patient suffered from ipsilateral wrist drop involving radial nerve and to some extent that was due to surgical compression of the terminal nerves of the brachial plexus and not due to block.

Conclusion:-

In nutshell, it can be concluded that, perineural dexamethasone 8 mg added to 0.5% Plain Bupivacaine in US guided interscalene block for elective upper arm surgeries, provides faster onset of sensory and motor blockage, prolongs the duration of sensory and motor blockage, duration of analgesia and decreases postoperative analgesic requests without any adverse reactions.

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