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# Comparison between Ropivacaine 0.5% And Levobupivacaine 0.5% in Axillary Brachial Plexus Block for Upperlimb Surgeries in a Tertiary Health Care Centre of Tripura- A Observational Study

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#### ARTICLE INFO

#### **ABSTRACT**

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#### Introduction

Axillary brachial plexus block is an anesthetic option used for surgeries of arm, forearm and elbow. The use of USG has significantly improved the quality of nerve blocks by direct visualization of nerves and related anatomical structures, needle trajectory and spread of local anesthesia during injection and significantly increases the success rate. Various drugs like Ropivacaine, bupivacine and Levobupivacaine are preferred due to greater margin of safety. However the efficacy of Levobupivacaine in Axillary Brachial Plexus Block has not been studied much. Therefore ,we designed this study to compare the clinical effect of Ropivacaine with Levobupivacaine for ABPB using USG technique.

#### Methodology

It's a hospital- based Comparative study done in Department of Anesthesiology, AGMC & GBPH from July 2016 to June 2018 (2 years) where 60 patients aged between 18-55 years with ASA grade 1 & 2 who are posted for upper limb surgery were randomly allocated into two groups of 30 each. Patients belonging to ASA grade (3& 4) and age <18 years,>55 years, Patient with h/o bleeding diathesis, neuromuscular disorder, morbid obesity, prolonged drug therapy & local site infection were excluded from the study.

USG guided axillary nerve block performed under aseptic condition. Sensory and motor blocks were assessed in each nerve territory at 2, 5, 10, 15, 20, 25, and 30 mins, 6 hr, 12 hr, 18hr, and 24 hrs after LA injection after LA injection. Onset of block, duration block and quality of analgesia has been compared. For the duration of the study, the presence of hypotension, bradycardia, hypoxia or nausea and vomiting was recorded and treated according to standard clinical practice.

#### Result

Among 60 study subjects Mean age was 34.7±12.6 years and majority of the study subjects were males (73.3%) and 26.7% were females. The pre-operative parameters e.g. age, sex, body weight etc. were compared between two groups but there was no statistically significant difference between the two groups (p>0.05). The onset of motor blockade among patients of Ropivacaine group was also shorter than patients received Levobupivacaine which was significant. Duration of sensory blockade was shorter in Ropivacaine group & duration of motor blockade was also shorter in Ropivacaine group than Levobupivacaine group and these difference were found to be statistically significant (p<0.05). There was no significant change in vital parameters after administration of both the drugs when observed at specific time intervals. VAS Scores were comparable in both the groups.

#### Conclusion

The following conclusion can be made from the present study

- Ropivacaine has faster onset of sensory and motor blockade when compared with Levobupivacaine.
- But duration of both sensory and motor blockade was lesser than Levobupivacaine
- Ropivacaine provides stable haemodynamic profile similar to Levobupivacaine.
- It provides satisfactory intra-operative &post-operative analgesia comparable to Levo

**KEYWORDS:** ABPB(axillary brachial plexus block), USG(utra sonography), Brachial Plexus, Ropivacine, Levobupivacine, sensory block, motor block.

#### INTRODUCTION

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Axillary brachial plexus block is an anesthetic option used for surgeries of arm, forearm and elbow. The conventional transarterial technique has potential problems such as nerve injury due to needle trauma and intraneural injection, as well as cardiac and CNS toxicity as a result of vascular uptake or accidental intravascular injection. The use of USG has significantly improved the quality of nerve blocks by direct visualization of nerves and related anatomical structures, needle trajectory and spread of local anesthesia during injection. In addition, USG guidance increases success rate, minimize local anesthetic volume needed for effective nerve block and avoids potential complications. 1,2 Bupivacaine, Ropivacaine and Levobupivacaine are the commercially available intermediate acting local anesthetics. They have some difference in risk of cardiovascular and CNS toxicity but they are more or less similar in analgesic and anesthetic potency. In Axillary Brachial plexus block in which relatively large dose of (~30-40ml) LAs are administered, Ropivacaine and Levobupivacaine are preferred due to greater margin of safety.3,4 There have been some studies on the efficacy of Bupivacaine and Ropivacaine in Axillary Brachial Plexus Block, and studies comparing Levobupivaine with Bupivacaine or Ropivacaine for neuraxial peripheral nerve blocks.<sup>5,6</sup> However the efficacy of Levobupivacaine in Axillary Brachial Plexus Block has not been studied much.<sup>7,8</sup> Therefore ,we designed this study to compare the clinical effect of Ropivacaine with Levobupivacaine for ABPB using USG technique.

### **METHODOLOGY**

It's a hospital- based Comparative study done in Department of Anesthesiology, AGMC & GBPH from July 2017 to June 2019 (2 years) where 60 patients aged between 18-55 years with ASA grade 1 & 2 who are posted for upper limb surgery were randomly allocated into two groups of 30 each. Patients belonging to ASA grade (3& 4) and age <18 years,>55 years, Patient with h/o bleeding diathesis, neuromuscular disorder, morbid obesity, prolonged drug therapy & local site infection were excluded from the study.

USG guided axillary nerve block performed under aseptic condition.. A short-bevelled 5 cm needle inserted either inplane or out-of-plane, relative to the probe, towards the four nerves and after careful positioning of needle tip, gentle negative aspiration, and an asymptomatic initial 0.5–1 ml perineural injection, further local anaesthetic is injected in 2 mL aliquots to surround each nerve.

The evaluation of the block of the different nerve territories was performed in an analogous sequence: median, ulnar, radial, brachial cutaneous, musculocutaneous, and intercostobrachial nerve. The characteristics of the sensory block were evaluated by pinprick in the cutaneous areas innervated by all nerves in the upper limb. Sensory block was evaluated by pinprick as follows:

0 (no block): normal sensitivity

- 1 (onset): reduced sensitivity compared with the same territory in the contralateral upper limb
- 2 (partial): analgesia or loss of the sharp sensation of the pinprick
- 3 (complete): anesthesia or loss of sensation to touch

Motor block was evaluated by thumb opposition for the median nerve, thumb adduction for the ulnar nerve, thumb abduction for the radial nerve, and flexion of the elbow for the musculocutaneous nerve.

Motor block was assessed according to the following scale:

0: no block

- 1 (onset): decreased movement with loss of strength
- 2 (partial): decreased movement with inability to perform movement against resistance
- 3 (complete): paralysis

Sensory and motor blocks were assessed in each nerve territory at 2, 5, 10, 15, 20, 25, and 30 mins after LA injection. Patients were considered to be ready for surgery when scores were 2 (partial sensory and motor block).

In the preoperative period, we recorded the following:

- onset of sensory and motor blocks in each nerve (time to reach scores of 1)
- percentage of patients who presented partial or complete sensory and motor block, 30 mins after injection of LA
- time when the patients were ready for surgery (scores of 2 in the nerves involved in the surgical area)

Intraoperatively, quality of anesthesia, fentanyl requirements, and signs of LA toxicity were assessed. In all instances,

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surgery was initiated 45 mins after LA injection. The quality of anesthesia was estimated to be (a) deficient, if the block failed and the patient had to receive general anesthesia; (b) partial, if the patient required more than 0.10 mg of IV fentanyl, skin infiltration, or an additional block; (c) sufficient, when the dose of IV fentanyl was 0.10 mg or less; or (d) complete, if no intraoperative analgesia was required. In the postoperative period, pain intensity was assessed with a visual analogue scale (VAS:1-10) at the time of analgesia request and 30 mins after the administration of analgesia. At the time of analgesia request, patients received a dose of 30 mg of IV ketorolac, which was repeated every 8 hrs, for the first 24 hrs postoperatively. Intravenous metamizol (2 g) was administered if the visual analog scale score 30 mins after ketorolac administration was greater than 3. The quality of analgesia was evaluated by the patients 24 hrs after injection of LA: 0 = poor, 1 = adequate, or 2 = excellent.

Sensory and motor blocks were assessed at 6, 12, 18, and 24 hrs after LA injection. For the duration of the study, the presence of hypotension, bradycardia, hypoxia or nausea and vomiting was recorded and treated according to standard clinical practice.

The sample size was determined on the basis of the variable, time ready to surgery, according to the results obtained by other investigators using the same LA in axillary plexus block. The parameters for the sample size calculation were as follows:

- (1) A 3.5-min difference between groups for clinically relevant targets (\$);
- (2) the > error level was 0.05;
- (3) the statistical power (A) was 0.80;
- (4) an SD of the measurements (R) of 4.8 mins.

Data are expressed as mean values (SD). For comparison between groups, we used the Student t and W2 tests, with P < 0.05 considered statistically significant. Analysis of the data was done using IBM SPSS version-20. Data was finally presented based on principles of descriptive and inferential statistics. Chi-square test and Independent sample t test was used to test the significance.

#### **RESULTS & DISCUSSION**

Among 60 study subjects Mean age was 34.7±12.6 years and majority of the study subjects were males (73.3%) and 26.7% were females. The pre-operative parameters e.g. age, sex, body weight etc. were compared between two groups but there was no statistically significant difference between the two groups (p>0.05). Similar to our study Kulkarni SB et al, Malav K et al, Vampugalla PS et al, 11 Kaur A et al 12 also found no statistical significant association between mean age, sex & body weight of Ropivacaine group and Levobupivacaine group.

The onset of sensory blockade among patients received Ropivacaine was shorter (10.83±4.1mins) than patients

received Levobupivacaine (14.00±3.5mins) & this relation was significant statistically (p<.05). Consistent with our study Kaur et al<sup>8</sup> in their study reported that onset of sensory block was observed from 5 min itself in Ropivacaine group as compared to bupivacaine group (10 min). But in contrast Jain S et al <sup>13</sup> in 2017 and Kulkarni SB et al <sup>5</sup> reported that onset of sensory blockade (p=0.027) was Significantly earlier in group of patients receiving levobupivacaine compared to ropivacaine. Cappelleri et al <sup>14</sup> and Mankad et al <sup>15</sup> has found sensory onset time was almost similar with that of Levobupivacaine and Ropivacaine group which is in contrast to our results.

The onset of motor blockade among patients of Ropivacaine group was also shorter (15.33±4.5 mins) than patients received Levobupivacaine (18.17±3.8mins). This relation was found to be statistically significant (p<0.05). consistent with our study Mankad et al 15 and Cacciapuoti A et al <sup>14</sup> reported motor onset time was faster in ropivacaine group  $(9.50\pm2.403 \text{ mins and } 14.0 \pm 2.3 \text{ min respectively})$ compared with levobupivacaine (12.33 $\pm$ 2.54 mins and 17  $\pm$  5 min respectively). Similar finding also found in other studies conducted by O Liisantti Luukkonen J et al, 16 Susana et al 17, Kaur et al<sup>12</sup>. In one of the study by Heavner et al.<sup>18</sup> there was a rapid onset time of sensory blockade which is consistent with our study finding but slower motor blockade with ropivacaine than levobupivacaine, in contrast to our study finding. Indumathi T et al 19 also reported that levobupivacine group had significantly earlier onset of sensory and motor block than group R (p<0.001)

Duration of sensory blockade was shorter (6.50±.938hours) in Ropivacaine group (7.53±1.00hours) & duration of motor blockade was also shorter (7.43±.817hours) in Ropivacaine group than Levobupivacaine group (8.73±.907hours) and these difference were found to be statistically significant (p<0.05). Our study finding was in agreement with study by Susana et al, <sup>17</sup> Kulkarni SB et al, <sup>9</sup> SIA et al, <sup>20</sup> Jain S et al, <sup>13</sup> and Gautier P et al <sup>17</sup> who reported longer duration of sensory loss in Levobupivacaine group than Ropivacaine group. Mankad et al, Cline et al <sup>8</sup> along with few other studies also reported finding in agreement with our finding, duration of motor block was shorter with ropivacaine when compared with levobupivacaine.

There was no significant change in vital parameters after administration of both the drugs when observed at specific time intervals. VAS Scores were comparable in both the groups. Similar to our study finding Mankad et al, <sup>15</sup> Indumathi T et al, <sup>11</sup> and Upadhyay et al <sup>22</sup> also reported that no significant changes was found in hemodynamic parameters between both the groups and in terms of hemodynamic stability, both groups were comparable (P > 0.005) which was not significant.

This study has further denoted that post-operative pain was lesser after administration of Ropivacaine than Levobupivacaine immediately after administration of drugs

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(at 10 mins, 15 mins and 20 mins intra operative) but on the long run patients received levobupivacaine experienced less pain than Ropivacaine as evident in further recording of pain

at 4, 5, 6, 7, 8, 17 hrs post-operative period. Similar to our result Mageswaran and Choy et al <sup>23</sup> observed no significant difference in VAS score of pain among both the groups.

Table: Comparison of onset of sensory block in Ropivacaine and Levobupivacaine group

Parameter	Ropivacaine	Levobupivacaine	Test applied	t test value	p value
	n=30	n=30			
Onset of sensory block (In mins)	10.83±4.1	14.00±3.5	Independent sample t test	-3.1 (-5.1,-1.1)	.003

The onset of sensory blockade among patients received Ropivacaine was shorter than patients received Levobupivacaine.

**Table:** Comparison of onset of motor block in Ropivacaine and Levobupivacaine group

Parameter	Ropivacaine	Levobupivacaine	Test applied	t test value	p value
	n=30	n=30			
Onset of motor block (In mins)	15.33±4.5	18.17±3.8	Independent sample t test	-2.6 (-5.0,-66)	.01

The onset of motor blockade among patients received Ropivacaine was shorter than patients received Levobupivacaine.

**Table:** Comparison of duration of sensory block in Ropivacaine and Levobupivacaine group

Parameter	Ropivacaine	Levobupivacaine	Test applied	t test value	p value
	n=30	n=30			
<b>Duration</b> of	6.50±.938	7.53±1.00	ent	-4.1	.000
sensory block			t t	(-1.5,53)	
(In hours)			Independent sample t test		
			nde		

Above result denotes statistical significant shorter duration of sensory blockade in Ropivacaine group than in Levobupivacaine group.

Table: Comparison of duration of motor block in Ropivacaine and Levobupivacaine group

Parameter	Ropivacaine	Levobupivacaine	Test applied	t test value	p value
	n=30	n=30			
Duration of motor block (In hours)	7.43±.817	8.73±.907	Independent sample t test	-5.8 (-1.7,85)	.000

Above result denotes statistical significant shorter duration of motor blockade in Ropivacaine group than in Levobupivacaine group.

#### **CONCLUSION**

The following conclusion can be made from the present study

- Ropivacaine has faster onset of sensory and motor blockade when compared with Levobupivacaine.
- But duration of both sensory and motor blockade was lesser than Levobupivacaine
- Ropivacaine provides stable haemodynamic profile similar to Levobupivacaine.

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 It provides satisfactory intra-operative &postoperative analgesia comparable to Levobupivacaine.

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