

Comparison of 0.5% Ropivacaine and 0.2% Ropivacaine in Bilateral Transversus Abdominis Plane Block in Laparoscopic Abdominal Surgeries

Research Article

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Abstract

Background and Aims: Transversus abdominis plane (TAP) block is a popular regional anaesthesia technique for post-operative analgesia after abdominal surgeries. The aim of the study was to evaluate the relative efficacy of 2 concentrations of ropivacaine for post-operative analgesia using ultrasound-guided TAP block in laparoscopic abdominal surgeries.

Methods: Sixty adults undergoing elective laparoscopic abdominal surgeries were randomised to receive ultrasound-guided TAP block at the end of the surgical procedure before extubation with either 0.5% ropivacaine (Group A, n = 30) or 0.2% ropivacaine (Group B, n = 30). All patients were assessed for post-operative pain and rescue analgesic consumption at 0, 2 h, 4 h, 8 h, 12 h and 24 h time points. Means for normally distributed data were compared using unpaired t-test, and proportions were compared using Chi-square or Fisher's exact test whichever was applicable.

Results: Patients receiving ultrasound-guided TAP block with 0.5% ropivacaine (Group A) had significantly lower pain scores when compared to patients who received the block with 0.2% ropivacaine (Group B) at 2h,4h,8h,12h and 24 h. The mean time required for the first rescue analgesia (after administration of block) in hours was significantly higher in Group A as compared to Group B. The mean 24 hours Opioid requirement (No. of doses of inj. Tramadol 50 mg IV) was significantly lower in group A as compared to Group B.

Conclusion: Ultrasound-guided TAP block with 0.5% ropivacaine provides effective and longer analgesia in the post-operative period as compared to 0.2% ropivacaine. And lesser 24 hour opioid requirement.

Keywords: Laparoscopic Surgeries; Ropivacaine; Transversus Abdominis Plane Block.

Introduction

The abdominal wall is significant source of pain after abdominal surgery. Laparoscopic surgery is a popular method of surgery with many advantages. Despite the minimally invasive nature, pain can be moderate to severe in immediate post-operative period [1].

The pain experienced by patients after abdominal surgery is derived from the anterior abdominal wall incision. The anterior abdominal wall is innervated by nerve afferents that course through the transversus abdominis neurovascular fascial plane [2].

The usual trend is to prescribe an opioid or non-steroid anti-inflammatory drugs (NSAIDs) for post-operative analgesia. Regional anaesthesia technique has gained widespread popularity as an important component of post-operative analgesia regimen.

The transversus abdominis plane block is a regional anaesthesia technique that provides analgesia to the parietal peritoneum as well as skin and muscles of the anterior abdominal wall. It has been shown to be a safe and effective post-operative adjunct analgesia method in a variety of surgical procedures and it is suggested as part of the multimodal anaesthetic approach to enhance recovery after abdominal surgeries [3]. It allows sensory blockade of plexus of nerves supplying abdominal wall, skin and muscles via local anaesthetic drug deposition.

The present study is to compare the duration of post-operative analgesia and 24-hour opioid requirement conferred by 0.5% ropivacaine and 0.2% ropivacaine used in transversus abdominis plane block for laparoscopic abdominal surgeries [4, 5].

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Methods

After the Institutional Ethics Committee approval, sixty American Society of Anaesthesiologists Physical Status I/II patients of either sex, aged 18-60 years, scheduled to undergo elective laparoscopic surgery were enrolled in this study. Patients with a history of allergy to test drugs, drug abuse, coagulation disorders or patients on anticoagulants and patients with abdominal infections were excluded from the study. The patients were randomly allocated into two groups to receive TAP block with either 0.5% ropivacaine and 0.2% ropivacaine solution.

Patients were randomly allocated into two groups, one group to undergo ultrasound-guided TAP block with 0.5% ropivacaine (Group A, n= 30) and other group to undergo ultrasound-guided TAP block with 0.2% ropivacaine (Group B, n= 30).

After pre anaesthetic evaluation valid, written, informed consent was obtained from all patients both for conduct of study as well as for administration of general anaesthesia.

All patients were kept nil by mouth from midnight before surgery and tablet Alprazolam (0.01mg/kg) was administered at bed time, the day before surgery.

In the operating room, routine monitors were applied and venous access was secured. Patients were premedicated with inj ondansetron 0.1mg/kg, inj glycopyrrolate 0.005mg/kg, inj midazolam 0.05mg/kg, following pre-oxygenation, the patients received IV fentanyl (2 µg/kg). Anaesthesia was induced with IV propofol 2 mg/kg. Vecuronium bromide (0.1 mg/kg) IV was utilised to facilitate tracheal intubation. Anaesthesia was maintained with nitrous oxide (60%) and isoflurane (0.5-1%) in oxygen.

At the end of surgery with Patient in supine position under all aseptic precautions ultrasound is kept midway between the coastal margin and iliac crest in the midaxillary line. The muscles are identified. The needle is gradually passed through skin, subcutaneous tissue, external oblique and internal oblique until the needle tip is placed between internal oblique and transversus abdominis muscle, after aspiration 2-3ml of local anaesthetic drug is given and spread of drug is seen. After confirmation 20ml of local anaesthetic is given on both sides.

Anaesthesiologist who observed the patient in post operative period is blinded to the drug injected in TAP block. Patients are monitored at 2, 4, 6, 12, 24 hrs post operatively for heart rate, blood pressure, saturation, pain, no of episodes of nausea and vomiting and complications if any. Duration of analgesia will be from the time of administration of transversus abdominis plane

block to administration of first rescue analgesia. All patients will be educated regarding the use of VAS scoring [6] from scale 1 to 10 in which 0 means no pain 10 means worst pain as depicted in figure no 1. Rescue analgesia (inj tramadol 50mg diluted IV slowly) will be given when VAS score is 4 or more.

The number of episodes of retching, nausea and vomiting in 24 hrs post operative period, severity is graded as:

- Mild: 1
- Moderate: 2
- Severe: 3

Statistical analysis

Statistical data was analysed by SPSS 20.0 version software. Collected data were spread on excel sheet and prepared master chart. Through the master chart tables, graphs and diagrams were prepared. For qualitative data analysis chi-square test was applied; for quantitative data analysis unpaired t test was applied for statistical significance. If P-value was less than 0.05 considered as significant.

Sample size calculation for 2 different groups of equal sizes for a continuous outcome measure.

n = sample size per group

α = 0.05 The probability of rejecting the null hypothesis when it is true. A level of 0.05 or 95% is most commonly used. the value was 1.96.

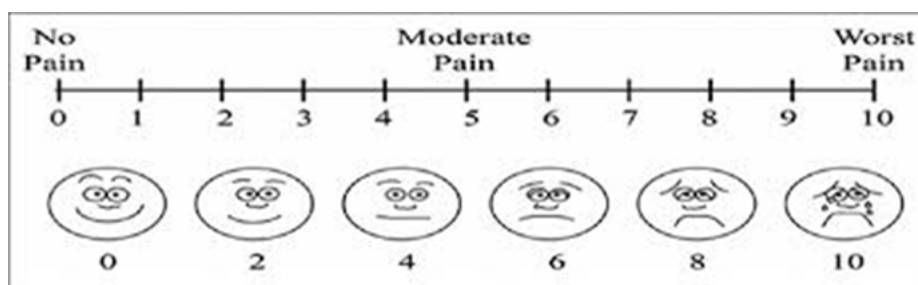
β = 0.2 The probability of failing to reject the null hypothesis if it is false. A level of 0.2 is most commonly used. This corresponds to a study power of 0.8 or 80%.

σ^2 = population variance in mean time to union (standard deviation²) the SD of the study was 18 in the reference study (The sample size was estimated based on the 24-hour morphine requirement in a previous study by Niraj et al. [7]) in Acta Anaesthesiologica Taiwanica.

μ_1 = Mean of group A
 μ_2 = Mean of group B
 $\mu_2 - \mu_1$ = minimum difference was 12.9

$$\begin{aligned} \text{Sample size} &= (Z_{\alpha/2} + Z_{\beta})^2 \times 2\sigma^2 / \mu_2 - \mu_1 \\ &= (1.96 + 0.82)^2 \times 2 \times (18)^2 / (12.9)^2 \\ &= 30 \text{ samples in each group} \end{aligned}$$

Figure 1.



Sample size = Total samples 60 (30 samples were taken in each group).

There was no difference in the demographic data (age, gender, body weight and height). There was no statistical significant difference of mean VAS pain score between Group A and Group B at 0 hours (P>0.05).

There was statistically highly significant difference of mean VAS pain score between Group A and Group B at 2 hours (P<0.002).

There was statistically very highly significant difference of mean VAS pain score between Group A and Group B at 4 hours, 8 hours, 12 hours and 24 hours of (P<0.001).

The mean VAS pain score were significantly higher in Group B as compare to Group A At 2 hours, 4 hours, 8 hours, 12 hours and 24 hours.

There was statistically very high significant difference of mean Time required for the first rescue analgesia (after administration of block) in hours between Group A and Group B (P<0.001).

The mean time required for the first rescue analgesia (after administration of block) in hours was significantly higher in Group A (Inj. 0.5% Ropivacaine) as compare to Group B (Inj.0.2% Ropivacaine).

There was statistically very high significant difference of mean 24 hours opioid requirement (No. of doses of inj. Tramadol 50 mg IV) between Group A and Group B (P<0.001).

The mean 24 hours OPIOD requirement (No. of doses of inj. Tramadol 50 mg IV) was significantly lower in group A as compare to Group B.

Comparison of Time required for the first rescue analgesia, Incidence of post operative nausea and vomiting and 24 hours OPIOD requirement cases between Group A and Group B.

Results

Group A that is Inj.0.5% Ropivacaine was significantly better as compare to Group B Inj. 0.2% Ropivacaine relieving pain. The mean time required for the first rescue analgesia (after administration of block) in hours was significantly higher in Group A (mean time 24.23 hours)as compared to Group B (mean time 5.57 hours). The mean 24 hours opioid requirement (No. of doses of inj. Tramadol 50 mg IV) was significantly lower in group A as compare to Group B.

Discussion

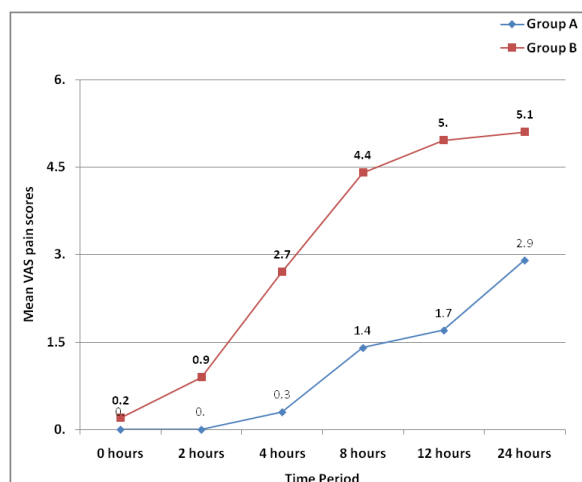
The benefits of adequate postoperative analgesia are clear and include a reduction in the postoperative stress response, reduction in postoperative morbidity, in certain types of surgery, there is

Table 1. Group A and Group B.

Time Period	VAS score				
	Group A	Group B	Std. Error	t –test value	P- value & Significance
	Mean ± SD	Mean ± SD			
0 hours	0.00 ± 0.00	0.17 ± 0.53	0.096	t = 1.723	P = 0.091, NS
2 hours	0.00 ± 0.00	0.89 ± 1.39	0.245	t = 3.515	P = 0.002, HS
4 hours	0.33 ± 0.60	2.67 ± 1.24	0.253	t = 9.252	P = 0.000, VHS
8 hours	1.43 ± 0.50	4.43 ± 0.56	0.132	t = 21.632	P = 0.000, VHS
12 hours	1.73 ± 0.69	4.96 ± 0.31	0.135	t = 23.241	P = 0.000, VHS
24 hours	2.86 ± 1.27	5.03 ± 0.80	0.274	t = 7.843	P = 0.000, VHS

NS = not significant, S = significant, HS = highly significant, VHS = very highly significant

Figure 2. Line Diagram Represents mean VAS pain Score of the Cases.



improved surgical outcome. Effective pain control also facilitates rehabilitation and accelerates recovery from surgery. Other benefits of effective regional analgesic techniques include reduced pain intensity, decreased incidence of side effects from analgesics and improved patient comfort [8], and early discharge from the hospital.

Using local anaesthetic agents in transversus abdominis plane block (TAPB) is a simple and effective analgesic technique, appropriate for surgical procedures where parietal pain is a significant component of postoperative pain. The local anaesthetic agents in TAP block have been demonstrated to provide excellent analgesia to the skin and musculature of the anterior abdominal wall in patients undergoing colonic resection surgery involving a midline abdominal wall incision.

The TAP block is an ideal and novel intervention in postoperative

pain management for lower abdominal surgeries. As compared to the traditional epidural analgesia and parenteral analgesics, the TAP block offers innumerable advantages such as decreased hemodynamic perturbations, decreased respiratory depression, nausea, vomiting, decreased cost, and overall patient satisfaction [9].

In this study local anaesthetic agents like 0.5% ropivacaine and 0.2% ropivacaine used in TAPB produced effective and prolonged postoperative analgesia.

Patients in group A had pain relief of upto 24 hours while patients in group B had pain relief only upto 5 hours postoperatively. 24 hour opioid requirement was less in group A compared to group B.

Though there was no significant difference between the 2 groups

Table 2.

Variables	Group A	Group B	Std. Error	t - test value	P- value & Significance
	Mean ± SD	Mean ± SD			
TIME OF FIRST RESCUE ANALGESIA (AFTER ADMINISTRATION OF BLOCK) in hours	24.23 ± 1.38	5.57 ± 1.30	0.342	t = 53.80	P = 0.000, VHS
24 HOUR OPIOID REQUIREMENT (no. of doses of injection tramadol 50 mg IV)	0.40 ± 0.49	3.23 ± 1.24	0.152	t = 18.43	P = 0.000, VHS

NS = not significant, S = significant, HS = highly significant, VHS = very highly significant.

Figure 3. Simple bar diagram represents comparison of mean Time required for the first rescue analgesia among Group A and Group B.

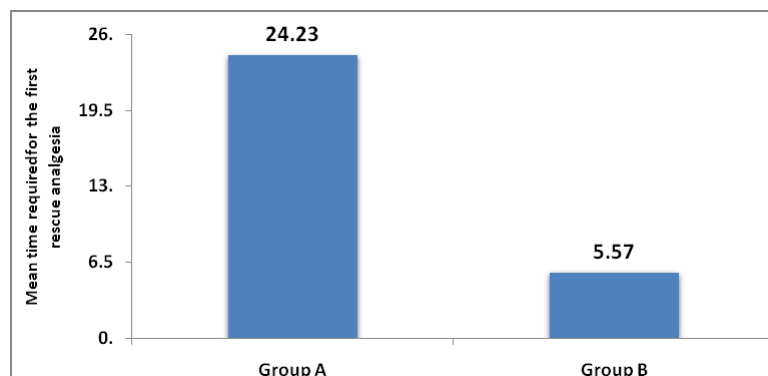
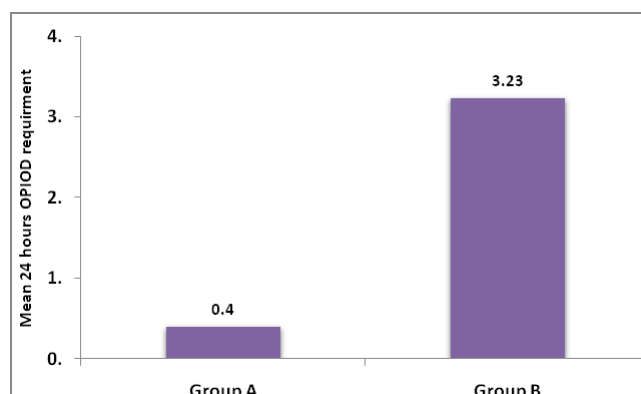


Figure 4. Simple bar diagram represents comparison of mean 24 hours OPIOID requirement among Group A and Group B.



in terms of post operative nausea vomiting but it was less in group A as compared to group B. Patients in group A were more comfortable in post operative period.

Patient receiving 0.5% ropivacaine had longer post operative analgesia as compared to 0.2 % ropivacaine even when laparoscopic surgeries were converted to open surgeries.

Some of the similar studies have been conducted in past few years Chen and Phui [10] found that injection of 20 ml of ropivacaine 0.375% for TAP block was effective as intraoperative and post-operative analgesic in 10 cases of laparoscopic cholecystectomy even after they were diverted to open cholecystectomy. Abdul Jalil et al. [11] in their prospective, randomized, double blind study on 56 patients scheduled for appendectomy under general anesthesia and they received TAP block with ropivacaine 0.2% in one group and ropivacaine 0.5% in the other group at the end of the surgery and they found that both concentrations provided comparable postoperative analgesia.

Oliviera et al. [12] in their prospective randomized, double blind-ed, placebo controlled study on 98 females scheduled for outpatient gynecological laparoscopy in which they compared between ropivacaine 0.25 % and ropivacaine 0.5% in TAP block to provide postoperative analgesia and they found that there was no significant difference between the two concentrations in their analgesic potency and no significant difference in parturients satisfaction score between the two concentrations. The present study has certain limitations. The pain scores at movement have not been taken into account despite the fact that laparoscopic surgeries are aimed to facilitate early ambulation. The serum concentrations of the drugs administered in the TAP were not estimated.

Conclusion

We conclude with our study that 0.5% Ropivacaine as compared to 0.2% ropivacaine used in Transversus abdominis plane block for laparoscopic surgeries of abdomen provided longer duration of post operative analgesia and 24 hour opioid requirement was also less in 0.5% ropivacaine group.

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