

Comparison of analgesic efficiency between wound site infiltration and ultrasound-guided transversus abdominis plane block after cesarean delivery under spinal anaesthesia

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Abstract

Background: Local anesthetic infiltration applied on the wound site or abdominal wall may be used for relieving postoperative pain after delivery by caesarean section. The aim of this study was to compare the analgesic efficiency of ultrasound (USG)-guided transversus abdominis plane (TAP) block with local anesthetic infiltration on a wound site.

Methods: This study was designed as a prospective randomized trial, and consisted of 70 pregnant women of American Society of Anesthesiologists (ASA) class I-II. Patients were randomized into Group I (wound site infiltration, n=35) and Group T (TAP block, n=35). Spinal anaesthesia was administered to all patients. In Group I, wound site infiltration was applied by the surgical team. In Group T, a USG-guided bilateral TAP block was applied. Patients' numeric pain scale (NPS) levels at 2, 6, 12 and 24th hours, after the operation (NPS₀) and during mobilization were assessed. Postoperative complications, time to first analgesic request and patient satisfaction were recorded.

Results: The NPS₀ values of Group T were found to higher and time to first analgesic request longer than those of Group I. The NPS values of Group I at 2, 6, 12, and 24th hours were found to be statistically significantly higher than those of Group T.

Conclusions: According to our results, USG-guided TAP block might be superior to infiltration anaesthesia for postoperative pain management of patients who have had caesarean section and it provided longer-lasting and more efficient analgesia. Hippokratia 2014; 18 (1): 28-31.

Keywords: Ultrasound, transversus abdominis plane block, infiltration anaesthesia, cesarean section

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Introduction

Long acting local anesthetics administered to both sides of the wound site or on/under the skin after surgery have been demonstrated to be effective for postoperative analgesia^{1,2}. It has been reported that, in addition to general or regional anaesthesia, local anesthetic infiltration and abdominal wall blocks are also useful for postoperative analgesia in cases of delivery by caesarean section³.

Transversus abdominis plane (TAP) block is a newly defined block that covers the nerves of the abdominal front wall. It was developed for postoperative pain control after gynecological and abdominal surgeries. Transversus abdominis plane block provides effective analgesia as part of a multimodal analgesic regimen for surgical procedures such as open appendectomy, laparoscopic cholecystectomy, hysterectomy, caesarean section, and large intestine resection performed by midline abdominal incision^{4,5}. Our aim was to compare the efficiency of ultrasound (USG)-guided TAP block with wound site in-

filtration anaesthesia. Our primary endpoints were pain scores at 0, 2, 6, 12 and 24 hours after surgery and patient satisfaction and our secondary endpoints were postoperative complications of the TAP block and wound infiltration approaches for postoperative analgesia and postoperative first analgesic application time.

Methods

After obtaining the approval of the ethics committee and the informed consent of the patients, a prospective randomized, double blinded study was undertaken of patients over 18 years old scheduled to caesarean section in Bagcilar Training and Research Hospital. Seventy pregnant women of American Society of Anesthesiologists (ASA) class I-II were involved in the study, which was completed in a 4-month period.

Elective cases with an eight-hour fast were included in the study; emergency cases were excluded. Patients with spinal anaesthesia contraindications (e.g., coagu-

lopathy, infection at puncture site) and those not wanting spinal anaesthesia were also excluded.

Before initiation, peripheral vascular access was obtained with a 16 or 18-gauge (G) intravenous cannula in all patients and preoperatively 8 mL kg⁻¹ h⁻¹ NaCl 0.9% was infused. General anaesthesia conditions, 0.50 mg atropine sulfate and 10 mg ephedrine were prepared for all patients. Patients' standard monitoring was applied when they were taken to the operating table. Blood pressure (mmHg), heart rate (HR, beat min⁻¹), and peripheral oxygen saturation (SpO₂) values were tracked by noninvasive methods.

Patients were randomized into groups (Group I: Wound site infiltration, n=35 or Group T: TAP block, n=35) with the help of a computer by an anaesthesia nurse according to their arrival time. An experienced anaesthesiologist, expert in TAP block, performed all spinal and TAP blocks on all patients; the spinal technique was performed with the patient in the sitting position. Using a midline approach, the intrathecal space was accessed by traversing the L3–L4 interval, with a 26-G Quincke needle (Exel Int, Los Angeles, CA). After confirmation of clear cerebrospinal fluid flow, 10 mg of hyperbaric bupivacaine (Heavy Marcaine 0.5%; Astra Zeneca, London, UK) plus fentanyl 20 µg were injected intrathecally. Only patients with successful spinal anaesthesia were included in the study.

In Group T, after the surgical procedure was accomplished, sterile skin cleaning was performed with patients lying on their backs, and the linear probe (5–10 mHz) of the USG device (SDU-450 XL, Kyoto, Japan) was prepared under sterile conditions. The probe was placed subcostally between the coastal margin and the iliac crest in the lateral abdominal wall, and the external oblique, internal oblique, and transversus abdominis muscles were identified. In the in-plane technique, a 100 mm 20-G peripheric blockage needle (B. Braun, Melsungen, Germany) was advanced, and local anesthetic medicine was administered over the transversus abdominis muscle. Forty mL (20 mL+20 mL, bilaterally) 100-mg levobupivacaine of 0.25% concentration was applied in Group T patients. All local anesthetics were prepared by the same co-investigator and the same co-investigator assisted during the whole TAP block procedure.

After the completion of the surgical procedure, in total 40 mL (20 mL for each wound site) 100-mg levobupivacaine of 0.25% concentration was used for subcutaneous wound site infiltration of the patients in Group I.

Intraoperative complications (nausea, vomiting, hypotension and bradycardia), postoperative complications (nausea, vomiting, hypotension, bradycardia and headache), need for analgesic, first postoperative analgesic administration time were recorded. Patient satisfaction was determined by asking the patients orally to provide a number between zero and ten (0: not satisfied, 10: satisfied a lot), and the number was recorded. Patient satisfaction evaluation was performed 24 hour after the the block.

Patient pain was evaluated by Numeric Pain Score (NPS), a scale of one to ten. The patients were asked to provide a number, with ten being the most violent pain they had ever had, and zero being no sensation of pain. In Group I NPS after local anaesthetic infiltration and in Group T NPS after TAP block was considered as NPS₀ (zero). Patients were evaluated at the 2, 6, 12, 24th hours and at the first mobilization by a co-investigator, who was blind to the used method and asked for their pain scores and the same co-investigator recorded all pain scores. All patients were routinely mobilized 8 hours after the end of the operation. If the patient suffered from pain (NPS>3) at any hour, intramuscular diclofenac sodium 75 mg was administered and if this was not efficient, 50 mg of tramadol were also given intravenously.

The sample sizes were calculated with the assumption of a possible at least of 35% difference between the two groups. Therefore 35 patients were allocated into each group in order to obtain an alpha error of 5% and statistical power of 80%.

Statistical Analysis

Complementary statistical methods (mean, standard deviation, median, interquartile range) were used to evaluate the data. The following tests were conducted: Friedman test for repetitive measurement of variable groups not showing normal distribution; Dunn's multi-comparison test for comparisons of subgroups; Mann-Whitney U test for comparisons of binary groups; independent t test for comparison of binary groups of variables showing normal distribution; and chi square test for comparisons of qualitative data. Results were considered statistically significant when p value was under 0.05.

Results

All applied TAP blocks were successful. No statistically significant differences were observed in age (p=0.341), body weight (p=0.271), or pregnancy frequency (p=0.912) between the groups (Table 1).

No statistically significant differences were observed in ASA scores (p=0.077), allergy history (p=1), or incidence of complications (p=0.060) between the groups. In both groups, intraoperative complications were hypotension and bradycardia. No postoperative complications were observed in any of the patients (Table 2).

The NPS₀ values of Group T were found to be significantly higher than those of Group I (p=0.012). The NPS values of Group I at 2, 6, 12 and 24th hours were found to be statistically and clinically significant higher than those of Group T (p=0.005, p=0.003, p=0.0001, p=0.0001). No statistically significant differences between the groups were observed in NPS values during the first mobilization (p=0.123) (Table 3).

The first analgesic administration of Group T (p=0.003) was found to be significantly later than that in Group I. No statistically significant difference was observed between the groups in patient satisfaction means (p=0.081) (Table 4).

Table 1: Age, body weight and number of pregnancy in Group I (wound site infiltration, n=35) and Group T (TAP block, n=35).

	Group T (n=35)	Group I (n=35)	p
Age (year)	27.77 ± 4.13	28.89 ± 5.5	0.341
Body Weight (kg)	73.14 ± 10.68	75.97 ± 10.67	0.271
Number of Pregnancy	2.46 ± 1.07	2.49 ± 1.1	0.912

Presented as Mean ± Standard Deviation.

Table 2: ASA score, allergy history and intraoperative complications in Group I (wound site infiltration, n=35) and Group T (TAP block, n=35).

		Group T(n=35)		Group I(n=35)		p
ASA score	I	35	100.00%	32	91.40%	0.077
	II	0	0.00%	3	8.60%	
Allergy History	No	33	94.30%	33	94.30%	1
	Yes	2	5.70%	2	5.70%	
Intraoperative Complication	No	29	82.90%	22	62.90%	0.060
	Yes	6	17.10%	13	37.10%	

ASA: American Society of Anesthesiology.

Table 3. Numeric Pain Scale in Group I (wound site infiltration, n=35) and Group T (TAP block, n=35).

	Group T (n=35)	Group I (n=35)	p
NPS ₀	0 (0-5)	0 (0-0)	0.012*
2 nd hour NPS	4 (2-6)	6 (4-7)	0.005*
6 th hour NPS	4 (3-5)	5 (4-6)	0.003*
First Mobilization NPS	4 (2-5)	4 (4-5)	0.123
12 th hour NPS	2 (1-3)	5 (4-6)	0.0001*
24 th hour NPS	2 (1-3)	4 (4-5)	0.0001*

NPS: Numeric Pain Scale, * p<0.05.

Table 4. Postoperative first analgesic application time and patient satisfaction in groups.

	Group T (n=35)	Group I (n=35)	p
Postoperative first analgesic application time (hour)	6.11 ± 6.2	2.63 ± 1.83	0.003*
Patient Satisfaction	8.89 ± 0.63	8.54 ± 0.82	0.081

Presented as Mean ± Standard Deviation, *p<0.05.

Discussion

Bamigboye et al³ compared wound site infiltration with a placebo in patients who had caesarean sections with regional anaesthesia and reported that NPS at first hour with wound site infiltration was lower. It has also been reported that wound site infiltration applied as a single dose for pain relief after caesarean section is an active, reliable, and simple method for the first four hours postpartum⁶. Similarly, in our study, NPS scores (NPS₀) after surgery were lower in Group I than in Group T. We are concluding that the difference between the two groups was due to rapid application of wound site administration in contrary to USG guided TAP block, which was more

time consuming.

McDonnell et al⁷ compared a placebo with TAP block and reported that TAP block provided superior analgesia until 48 hours. In addition, it has been reported that TAP block not only reduced postoperative opioid need but also extended first analgesia application time⁸. We did not use a control group, as both TAP block and infiltration anaesthesia had already proved their superiority over placebo. Our study is the first study to compare TAP block with infiltration anaesthesia after spinal anaesthesia in deliveries with caesarean section, and our primary goal was to compare patient pain scores and patient satisfaction. In our study, no analgesic was administered to seven

patients in Group T and five patients in Group I in the first 24 hours. However, the first analgesia administration time was longer in Group T than in Group I. In both groups, anti-inflammatory medicine administration was sufficient. No opioid was needed for any patients.

Belavy et al⁹ demonstrated that morphine consumption is lower when USG-guided TAP block is used as a component of a multimodal regimen after spinal anaesthesia. In their retrospective study on patients with or without TAP after caesarean section, Patel et al¹⁰ found that oral narcotic tablet use was low, 24-48 hours postoperatively, in the TAP group. In some newly conducted studies, it has been reported that total morphine need is lower in TAP patient groups and that side effects, such as sedation related to opioid use, postoperative nausea, and vomiting, are also low and that patient satisfaction is better^{4,8}. Scharine¹¹ reported that a long and effective analgesia is obtained by TAP block, and that lower pain score, earlier oral nutrition, and earlier mobilization are seen, and duration of hospital stay is shortened, as when no narcotic analgesic is used. In their study, Tan et al¹² applied USG-guided TAP block after general anaesthesia in caesarean section operations, and they found that morphine need is lowered and mother satisfaction is increased. In our study, there were no differences between groups in terms of patient satisfaction. We believe the reason for this result is that both methods we used provide more effective analgesia than placebo.

Ganta et al¹³ compared patients to whom they applied wound site infiltration and ilioinguinal block after caesarean section with a control group, and they did not find a statistically significant difference. In a study similar to our study, on/under skin local anesthetic administration and USG-guided TAP blocks in total hysterectomy patients were compared with a control group. Pain scores of the group with TAP block were found to be lower than those of the infiltration group in the 6 and 24th hours, and it was reported that TAP block was more effective than surface wound site infiltration in postoperative pain management¹⁴. In our study on caesarean section patients, the pain scores in the TAP block group were lower at 2, 6, 12 and 24 hours postoperatively. However, there were no differences in pain scores during their first mobilization.

Conclusion

According to our results, USG-guided TAP block could be considered superior to infiltration anaesthesia for the postoperative pain management of patients who have had caesarean section under spinal anaesthesia, and it provides longer-lasting and more efficient analgesia than infiltration anaesthesia.

Conflicts of interest

None of the authors have any conflicts of interest to declare.

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