

ORIGINAL RESEARCH

Evaluation of the Analgesic Efficacy of Bilateral Ultrasound-Guided Transversus Thoracic Muscle Plane Block on Post-Sternotomy Pain: A Randomized Controlled Trial

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¹Department of Anesthesiology, Ain Shams University, Cairo, Egypt; ²Department of Cardiothoracic Surgery, Ain Shams University, Cairo, Egypt **Purpose:** Transverse thoracic block is a new perioperative analgesic method for poststernotomy discomfort. This study discusses the efficacy of an ultrasound-guided transversus thoracic muscle plane block (TTPB) in patients undergoing heart surgery, including sternotomy.

Patients and Methods: In this prospective trial, 60 patients were randomly assigned to two groups: transversus (T) or general anesthesia (GA). Patients in the T group received TTPB with GA and 15 mL of 0.25% bupivacaine was administered on either side. Patients in the GA group received TTPB along with GA, but 15 mL saline was injected on either side instead. The patients were followed up postoperatively. The percentage of patients receiving additional opioids, extubation time, intensive care unit (ICU) length, postoperative morphine dose, and adverse events were documented.

Results: The percentage of patients requiring additional morphine was significantly lower in the T group (p<0.015). Pain scores were comparable between the groups at extubation time and 18 h postoperatively. At 8, 12, and 24 h after surgery, pain scores in the T group were significantly lower (P<0.001). The total amount of morphine required after surgery was much lower in the T group (8 \pm 0.74 mg), and the duration to initial analgesic necessity was significantly greater in the GA group (P<0.001). Patients in the GA group had a considerably longer ventilation time (11.07 \pm 0.64) (P<0.001). The ICU stay in the T group was substantially shorter (P<0.001), and postoperative complications were comparable.

Conclusion: For patients undergoing heart surgery, TTPB offers sufficient postoperative pain control, resulting in lower pain scores, lower postoperative analgesic doses, shorter extubation times, and shorter ICU stay lengths.

Keywords: transversus thoracic block, adult cardiac surgery, postoperative analgesia, sternotomy

Introduction

Cardiac surgery with a median sternotomy generates severe discomfort in the sternum, which may lead to delirium, hemodynamic instability, or pulmonary problems.¹ Multimodal analgesic techniques for perioperative pain control in cardiac surgery rely heavily on systemic opioids which cause side effects such as vomiting, respiratory depression and itching.^{2,3} Because of the risk of hematoma formation following full heparinization intraoperatively, regional methods such as

Correspondence: Hoda Shokri Ain Shams University, Cairo, Egypt Tel +20 1211179234 Email Drhoda10@yahoo.com epidural or spinal blocks are not routinely employed in heart surgery but it drew the attention to increased use of plane blocks due to their safety and closeness to the field of surgery.⁴

The anterior branches of the T2–T6 intercostal nerves, which provide sensory supply to the internal mammary area, carry sternotomy-induced discomfort. It has been discovered that inhibiting nerves at the surgery site can relieve sternotomy discomfort while avoiding the hazards of more intrusive procedures.⁵

Ultrasound-guided transversus thoracic muscle plane block (TTPB), introduced by Ueshima in 2015, is a novel technique in which a local anesthetic (LA) is given as a single shot into the fascial plane between the transversus thoracic muscle (TTM) and the internal intercostal muscles (IIM) to block the anterior branches of T2-T6 as part of multimodal analgesic techniques. ^{6,7} The goal of this prospective randomized trial was to determine the effectiveness of TTPB for adult patients undergoing heart surgery including sternotomy incision in terms of the percentage of patients requiring extra doses of morphine, postoperative pain scores, postoperative opioid consumption, intensive care unit (ICU) stay length, and the incidence of local or systemic complications of the technique or the agents injected.

Patients and Methods

The study was approved by the Research Ethics Committee of Ain Shams University (approval number FMASU R 20/2020/2021) done on January 21st, 2021. This study was registered at ClinicalTrials.gov (registration number NCT04872192).

Inclusion and Exclusion Criteria

All patients signed a written informed consent form. This prospective, double-blind, parallel-group clinical investigation included 60 adult patients aged 55–74 years who were ASA (American Association of Anesthesiologists) II and III who underwent elective cardiac surgery with median sternotomy performed by the same surgical team at Ain Shams University Hospital, cardiothoracic academy, from May to August 2021.

Patients who had been intubated for more than 24 h prior to surgery, patients undergoing complex cardiac procedures, patients with severe chronic obstructive pulmonary disease, patients unable to communicate, patients with chronic pain and severe pulmonary hypertension, as

well as any contraindication to regional anesthesia or drug allergy, were all excluded from the study.

Study Design and Participants Induction of Anesthesia and Monitoring

A detailed medical history, including medications used, and full investigations were performed at the night of the surgical procedure. In the pre-induction room, midazolam (1-3 mg IV) was used as a premedication. Fentanyl (5-6 g/kg), propofol (2 mg/kg), and rocuronium (0.6 mg kg-1) were used to induce anaesthesia, which was maintained with 1%-2% isoflurane. The heart rate and blood pressure were maintained within 20% of baseline levels. After intravenous catheterization and endotracheal intubation, arterial and central venous catheters were inserted. Pulse oximetry, esophageal temperature, and invasive arterial blood pressure by cannulating the radial and/or femoral artery to evaluate beat-to-beat changes in blood pressure with a 5-lead electrocardiogram with automatic ST segment analysis, transesophageal echocardiography, and coagulation profile monitoring using activated clotting time.

Nerve Block and Grouping

After induction of anesthesia, patients were assigned to either the transversus group (T group) or general anesthesia (GA) group at random (N group) using a computer-generated random number in various block sizes in a 1:1 ratio. Randomization was performed using the SAS statistical package version 9.3 (SAS Institute, Cary, NC, USA) by a statistician who was not involved in the study.

Patients in the T group (n=30) received a bilateral TTPB with general anesthesia and 15 mL of 0.25% bupivacaine administered on either side. The patient was placed in the supine position and after proper sterilization, a 12 L-RS linear probe of the Sonosite M Turbo ultrasonography system was used prior to surgical incision (National Electrical Manufacturers, USA) and positioned in the sagittal plane under the clavicle lateral to the sternal edge and the ribs were counted by sliding the probe downward till the fifth rib was encountered. The probe was placed horizontally over the fourth intercostal space to identify the internal mammary artery, transverse thoracic (TT) and internal intercostal muscles(II) as shown in Figure 1A. The probe then rotated vertically to identify internal mammary artery in parasagittal view as seen in Figure 1 (B). After that the probe was slid more laterally to identify transverse thoracic plane between II and TT muscles Figure 1C. A 20-gauge Tuohy needle was advanced using inplane technique from caudal Dovepress Shokri et al

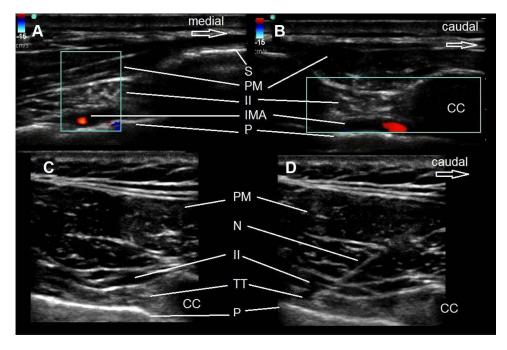


Figure I The block technique. (A) The probe was placed horizontally over the 4th intercostal space to identify the internal mammary artery (IMA) in the horizontal view, (B) then rotated vertically to identify the IMA the in the longitudinal (parasagittal) view. (C) The probe was slide more laterally to identify the transversus thoracic plane, between the internal intercostal (II) and the transversus thoracic (TT) muscles, (D) where the needle (N) tip was adjusted.

Abbreviations: CS, costal cartilage; P, pleura; PM, pectoralis major muscle; S, sternum.

to cranial and its tip was adjusted in the transverse thoracic plane between the IIM and the TTM. To identify this plane, 1 mL of normal saline was injected, followed by 15 mL of 0.25% bupivacaine on each side following intermittent aspiration, monitoring of the LA spread and pressure on the pleura to confirm the accurate injection (Figure 1). To avoid unintended arterial puncture and subsequent bleeding, it is critical to identify the internal mammary artery (IMA), which is a hypoechoic pulsatile structure. Proper visualization of the IMA can also help with the block. Patients in the GA group (n=30) received a bilateral ultrasound-guided TTPB technique combined with general anesthesia, but instead received 15 mL saline (control) on either side. The TTPB approach was used in both groups, along with general anesthesia. A single expert anesthetist performed all the blocks. A pharmacist who was uninformed of the nature of the study produced drug packs prior to the start of the study. Additional 0.5-1 micrograms/kg IV fentanyl dosages were administered if hemodynamic abnormalities, such as elevated blood pressure or severe tachycardia, were observed. All cardiac surgical techniques included medial sternotomy.

Recovery and Postoperative

After the cardiac surgical procedure, all patients were moved to the ICU to maintain hemodynamics, control bleeding, and correct hemoglobin levels, serum electrolytes, and acid-base balance. Standard postoperative analgesia was accomplished using acetaminophen (1 gm/6 h).

All patients received postoperative care according to the protocol, which was implemented by well-trained, qualified bedside nurses who were overseen 1:1 by well-trained ICU specialists. When the patient was able to maintain spontaneous breathing following extubation, all patients were extubated when deemed clinically appropriate according to the local ICU procedure by the ICU staff. The patients were encouraged to sit in a chair and move around with the help of health care workers in the ICU, after which the physiotherapist ensured the enhancement of the patients' mobility and rehabilitation until they were discharged.

Patient demographics, medical status before surgery, left ventricular function, and surgical data (total ischemia duration and number of grafts) were recorded.

Measurements

The percentage of patients requiring additional doses of morphine was the primary outcome of the study. The secondary outcomes included the total dose of postoperative morphine consumed, postoperative visual analogue pain scores obtained after patient extubation for pain assessment at rest (0 = no pain, 10 = maximum unbearable pain) were recorded at the time of extubation, and at 8h, 12 h,18 h and 24 h postoperatively. Pain scores >4 were recorded, and patients were given morphine (0.05 mg/kg) administered by a physician who was blinded to the nature of the study. Time needed for first rescue analgesic, postoperative mean blood pressure, and heart rate were recorded in addition to extubation time, length of ICU stay, and incidence of complications related to the technique such as hemothorax or pneumothorax, arrhythmias, and LA toxicity. The endpoints were unstable medical conditions such as difficult weaning from cardiopulmonary bypass, significant postoperative bleeding that necessitated re-exploration. All difficulties were dealt with in accordance with surgical and medical protocols.

Sample Size Calculation

After accounting for a 20% dropout rate, software for statistics and data science (Stata program version 10; Stata Corp LLC, Texas, USA) revealed that 60 patients (30 per arm) were required (setting an alpha error of 5% and power of 90%). The percentage of patients who required extra analgesia (pentazocine) within 24 h after surgery was much lower in the T group (57%) than in the GA group⁸ (94%).

Statistical Analysis

SPSS Statistics version 23 (IBM Corp., Armonk, NY, USA) was used to analyze the data. The mean and standard deviation were used to represent normally distributed numerical data, whereas the median and interquartile range were used to represent skewed data. The number, percentage, or ratio was used to present qualitative data. An unpaired *t*-test was used to compare normally distributed numerical data. The Mann–Whitney test was used to compare skewed numerical data, and Chi-square test was used to analyze qualitative data. Statistical significance was defined as a P<0.05.

Results

This research was conducted by researchers from Ain Shams University Hospital from May to August 2021. As illustrated in the CONSORT flow diagram, a total of 60 patients were enrolled and analyzed in this study (Figure 2).

No significant difference was observed between the study groups in terms of demographic, surgical, and preoperative data (Table 1). At the extubation time and 18 h postoperatively, pain scores were similar between the study groups; on the other hand, pain scores were significantly lower for the T group than the GA group at 8, 12, and 24 h postoperatively, P<0.001 (Table 2). The total dose of postoperative morphine required was significantly smaller in the T group than in the GA group, and the time to first analgesic requirement was markedly longer in the GA group (P<0.001) (Table 3). The percentage of patients requiring additional doses of morphine was higher in the GA group than in the T group (P<0.015) (Table 3).

Postoperative mean blood pressure and pulse rate were lower in the T group than in the GA group (P<0.001) (Table 4). The ventilation time was significantly longer in the GA group (P<0.001) (Table 4). The need for inotropic support was comparable between groups (Table 4). ICU stay length was significantly shorter in the T group (p<0.001) (Table 5). Postoperative complications such as pneumothorax, arrhythmia, LA toxicity, and skin infection were not significantly different between the study groups. There were no recorded cases of hemothorax or anaphylactic shock (Table 5).

Discussion

This study investigated the effectiveness of TTPB in the management of postoperative pain after cardiac surgery in adults with a median sternotomy.

The results of this prospective study showed that in the first 24 h after surgery, the percentage of patients needing extra morphine doses, postoperative morphine use, pain scores, and first analgesic requirement were significantly lower in the T group than in the GA group. Ventilation time and ICU stay were significantly shorter in the T group. There were no significant differences in postoperative complications between the study groups.

Earlier studies reported that post-sternotomy pain was conducted through the T2-T6 intercostal nerves; thus, blocking these nerves can efficiently control sternotomy-induced pain and reduce ICU and hospital stay length.⁴

In a study by Chaudhary et al, pediatric patients scheduled for cardiac surgery received parasternal block to block the T3-T6 intercostal nerves prior to sternal closure. They found that ventilation time was significantly shorter. In addition, pain scores and postoperative opioid requirements were significantly lower. These results support the findings of the present study.

A retrospective study by Ueshima et al who underwent single-shot TTPB for patients undergoing cardiac surgery through sternotomy, revealed fentanyl doses as rescue

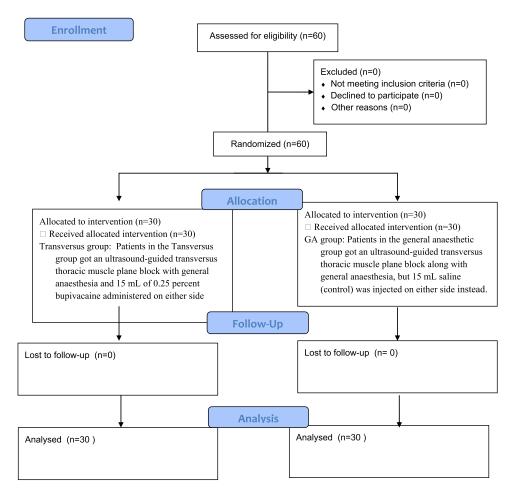


Figure 2 CONSORT flow diagram.

Notes: Adapted from: Schulz KF, Altman DG, Moher D, CONSORT Group. CONSORT 2010 Statement: Updated Guidelines for Reporting Parallel Group Randomised Trials. PLoS Med. 2010;7(3):e1000251. Copyright: © 2010 Schulz et al. Creative Commons Attribution License.

analgesics and pain scores were significantly reduced. These findings are consistent with ours. 6

Kendigelen et al demonstrated that TTPB has a duration of more than 48 h because it is one of the interfascial blocks known for prolonging analgesia. 10

After reviewing the literature, no complications related to TTPB have been reported, such as hemothorax, pneumothorax, wound infection, or hematoma.¹¹

Previous studies reported that a large volume of diluted LA must be injected to ensure adequate diffusion of the drug and increase its efficacy without exceeding the maximum dose.¹²

A retrospective study by Cakmak et al studied the analgesic efficacy of TTPB in pediatric patients undergoing congenital cardiac surgery through median sternotomy. They found that pain scores and postoperative fentanyl doses were significantly lower in the T group than in the non-T group. Extubation time was significantly shorter in the T group (p<0.001). These results are consistent with the findings of the present study.

Severe postoperative pain following cardiac surgery lasts for 1–7 days originally caused by median sternotomy, as mentioned by Mueller et al. In addition, Lahtinen et al reported that 75% of patients experienced postoperative chest pain. 14

The superficial technique of TTPB reduces the incidence of bleeding, especially with heparinization and the presence of coagulopathy.¹⁵

Earlier studies showed that the optimal place of administration of LA was the T4–T5 intercostal space because it resulted in better spread from the manubrium to the fifth intercostal space than injection at the T3-T4 intercostal space.¹⁶

The duration of action of the TTB can last approximately 24 h postoperatively because of the roughness of the TTM and LA spread intramuscularly; thus, its absorption is slower.¹⁷

A study by Ueshima et al studied 299 patients receiving TTPBs and reported that there were only two cases of mild skin infection at the puncture site of the block.¹⁸

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Table I Demographic Data, Surgical Data and Comorbidities

	N Group	T Group (N=30)	Tests	
	(N=30)		t/ X ²	P-value
Sex	•			
Female Male	12(40%) 18(60%)	13(43.3%) 17(56.7%)	0.069	0.793
Age	65.57±5.65	65.13±5.59	0.299	0.766
ASA	•			
II III	22(73.3%) 8(26.7%)	20(66.7%) 10(33.3%)	0.317	0.573
Procedure duration (min)	461.87±1.72	460.97±2.22	1.756	0.084
Cross clamp time (min)	70.27±3.5	71.87±4.05	1.637	0.107
CPB time (min) Hypertension Diabetes mellitus	145.73±0.74 7(23.3%) 5(16.7%)	145.47±0.9 10(33.3%) 3(10%)	1.254 0.739 0.577	0.215 0.390 0.448
Type of cardiac surgeries				I
Aortic valve replacement Mitral valve replacement CABG (3 grafts)	3(10%) 12(40%) 15(50%)	4(13.3%) 14(46.7%) 12(40%)	0.630	0.730
Preoperative EF (%)	55.2±1.19	54.77±1.07	1.484	0.143
Preoperative mean blood pressure	68.7±1.21	69.27±1.23	1.801	0.077
Preoperative mean pulse rate	77.6±3.07	78.67±2.89	1.385	0.171
Re-exploration (no significant bleeding or collection was found)	I (3.3%)	2(6.7%)	0.351	0.554
BMI kg/m2	23.35±0.95	23.23±1.74	0.332	0.741

Abbreviations: ASA, American Association of Anesthesiologists; CPB, cardiopulmonary bypass; EF, ejection fraction; BMI, body mass index.

Table 2 Postoperative Pain Scores

Pain Score	N Group (N=30)		T Group (N=30)			ann– ney Test
					Z	P-value
At extubation time	(0-0)	0	(0-0)	0	0.000	1.000
8 h	(2-6)	4	(0-3)	0	6.188	<0.001*
I2 h	(3-4)	4	(1-4)	2	4.427	<0.001*
18 h	(1-2)	2	(0-2)	2	1.105	0.269
24 h	(0-4)	4	(0-4)	2	3.960	<0.001*

 $\textbf{Notes} \hbox{:}\ \mathsf{Data}\ \mathsf{are}\ \mathsf{median}\ (\mathsf{interquartile}\ \mathsf{range}).\ \mathsf{h,}\ \mathsf{hours.}\ \mathsf{*Mann-Whitney}\ \mathsf{test.}$

Satoru et al studied single-shot TTPB efficacy in adult patients undergoing elective cardiac surgery. They revealed that pain scores were significantly lower in the block group than in the control group. On the other hand, 24-hour total hydromorphine doses were similar in both groups, unlike the current study that showed lower morphine doses required for the block group.¹⁹

Ueshima et al investigated the effectiveness of TTB in female patients undergoing breast surgery and found that the pain scores as well as the number of patients requiring further doses of analgesics were lower in the T group than in the GA group.⁸

A study by Abd Elbaser et al²⁰ performed TTB in pediatric patients undergoing cardiac surgery with median sternotomy, and found that postoperative fentanyl requirements were significantly lower in the T group than in the saline group, and pain scores were significantly lower in the T group at all time points. These findings are consistent with our results.

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Table 3 Total Postoperative Morphine Dose, Time to First Analgesic Dose and Percentage of Patients Receiving Extra Doses of Morphine

	N Group (N=30)	T Group (N=30)	T-to	est	
			t	P-value	
24 h postoperative morphine (mg)	13.9±0.66	8±0.74	32.484	<0.001*	
Time to first analgesic dose (hours)	5.57±0.5	9.07±0.69	22.404	<0.001*	
			Chi-square test	P-value	
	N group (N=30)	T group (N=30)			
Patients receiving extra doses of morphine	15 (50%)	24 (80%)	5.934	<0.015*	

Notes: Postoperative morphine dose and time to first analgesic dose were presented as Mean±SD and compared using Independent *t*-test, patients receiving extra doses of morphine presented as percentage and compared by chi-square test. *Significant.

Table 4 Postoperative Hemodynamic Values, Need for Inotropic Support and Ventilation Time

	N Group (N=30)	T Group (N=30)	Tests	
			t/ X ²	P-value
Postoperative mean blood pressure	80.83±0.75	78.57±1.01	9.908	<0.001*
Postoperative mean pulse rate (beats/min)	96.40±1.00	94.63±0.96	6.953	<0.001*
Need for inotropic support	2(6.7%)	4(13.3%)	0.741	0.389
Ventilation time (hours)	11.07±0.64	7.50±0.51	23.905	<0.001*

Notes: All data were presented as Mean±SD and compared using Independent t-test while need for inotropic support presented as percentage and compared using Chi-square test. *Significant.

Table 5 ICU Stay Length and Postoperative Complications

	N Group (N=30)	T Group (N=30)	Te	ests
			t/ X²	P-value
ICU stay length (days)	2.40±0.50	1.57±0.50	6.440	<0.001*
Pneumothorax	I (3.3%)	0(0.0%)	1.017	0.313
Hemothorax	0(0.0%)	0(0.0%)	0.000	1.000
Postoperative arrythmia	5(16.7%)	3(10.0%)	0.577	0.448
Local anesthetic toxicity	0(0.0%)	0(0.0%)	0.000	1.000
Anaphylactic shock	0(0.0%)	0(0.0%)	0.000	1.000
Skin infection	0(0.0%)	3(10.0%)	3.158	0.076

Notes: All data were presented as percentage and compared by Chi-square test except for ICU stay length presented as Mean±SD and compared by T-test. *Significant.

There are few limitations to the present study. Firstly, the efficiency of TTPB in patients undergoing high risk cardiac procedures was not determined in this study. Also, the sample size was modest, thus necessitating more clinical trials with a larger number of patients to identify the occurrence of rare side effects and we did not record any failures in the blocks because the blocks were delivered under ultrasound guidance by an expert.

Lastly, the blocks were performed after induction of anesthesia, so the sensory levels could not be measured.

Conclusion

For adult patients undergoing heart surgery, TTPB offers sufficient postoperative pain control, resulting in lower pain scores, lower postoperative analgesic requirements, faster extubation time, and shorter ICU stay length without any major consequences.

Funding

There is no funding to report.

Disclosure

The authors report no conflicts of interest in this work.

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