ORIGINAL RESEARCH



Effect of Single-Injection Thoracic Paravertebral Block via the Intrathoracic Approach for Analgesia After Single-Port Video-Assisted Thoracoscopic Lung Wedge Resection: A Randomized Controlled Trial

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ABSTRACT

Introduction: Pain is still severe after single-port video-assisted thoracoscopic (SPVAT) lung wedge resection. We observed the effect of single-injection thoracic paravertebral block (TPB) via the intrathoracic approach for analgesia after SPVAT lung wedge resection.

Methods: Sixty patients undergoing SPVAT lung wedge resection were randomly divided into a control group and an observation group. All patients underwent TPB via the intrathoracic approach at the T4 level with a scalp needle before closing the chest. The patients in the observation group received 20 ml 0.375% ropivacaine at the T4 level, and the patients in the control group received 20 ml of 0.9% saline. A patient-controlled intravenous analgesic (PCIA) pump with sufentanil was attached to all patients after surgery. The sufentanil consumption and number of PCIA presses in the first 24 h after surgery were recorded. The visual

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H. Tian · J. He Department of Thoracic Surgery, Ningbo Medical Center Lihuili Hospital, Ningbo, China analogue scale (VAS) scores (during rest and coughing) were recorded at 6 h, 12 h, 24 h, and 36 h after surgery. The incidence of adverse reactions after surgery were recorded.

Results: The sufentanil consumption in the observation group was significantly lower than that in the control group (34.2 \pm 1.9 μg vs. 52.3 \pm 2.3 μg; P < 0.001). The VAS score at 6, 12, and 24 h after surgery, the incidence of adverse reactions after surgery in the observation group were significantly lower than those in the control group (all P < 0.05). The number of PCIA presses in the observation group was significantly lower than that in the control group [0 (0–0) times vs. 3 (2–4) times, P < 0.001].

Conclusions: Single-injection TPB via the intrathoracic approach under thoracoscopic direct vision is easy to perform and can effectively alleviate postoperative pain after SPVAT lung wedge resection, with fewer adverse reactions.

Trial Registration: ChiCTR2000034726.

Keywords: Lung wedge resection; Postoperative analgesia; Single injection; Single-port video-assisted thoracoscopic surgery; Thoracic paravertebral block

Key Summary Points

Why carry out this study?

Pain is still severe after single-port thoracoscopic (SPVAT) lung wedge resection.

Can single injection local anesthetics into the paravertebral space via the intrathoracic approach under thoracoscopic direct vision alleviate postoperative pain?

What was learned from the study?

Single-injection TPB via the intrathoracic approach can alleviate postoperative pain and reduce adverse reactions.

Single-injection TPB via the intrathoracic approach under thoracoscopic direct vision is easy to perform and can effectively alleviate postoperative pain after SPVAT lung wedge resection with fewer adverse reactions.

DIGITAL FEATURES

This article is published with digital features, including a summary slide, to facilitate understanding of the article. To view digital features for this article go to https://doi.org/10.6084/m9.figshare.13402367.

INTRODUCTION

Traditional thoracotomy has great trauma and severe postoperative pain. At present, thoracoscopic minimally invasive surgery is the main method of thoracic surgery [1, 2]. In recent years, single-port video-assisted thoracoscopic (SPVAT) surgery has been widely used in thoracic surgery [3, 4], especially in lung wedge resection [5]. SPVAT lung wedge resection has the advantages of less incision, less pain and faster recovery. It was considered to have mild

postoperative pain, and ignored the need for postoperative analgesia, resulting in insufficient analgesic measures after surgery. In fact, the postoperative pain is still severe, which seriously affects the patient's postoperative recovery [6]. Currently, postoperative analgesia after thoracic surgery mainly includes thoracic epidural analgesia, patient-controlled intravenous analgesic (PCIA), thoracic paravertebral block (TPB) and erector spinae plane block [7]. These are mainly performed by well-trained anesthesiologists, and are mainly used in radical cancer surgery and other thoracic surgery. With the popularization of the concept of enhanced recovery after surgery, the surgical team began to pay attention to postoperative pain management. In recent years, percutaneous catheter TPB or intercostal nerve block under thoracoscopic direct vision has been widely favored by surgeons [8, 9]. The various analgesic methods mentioned above have their own advantages. but also have many complications and disadvantages [10-16]. Is there a simpler, quicker, effective and less complication analgesic method that can be used for postoperative analgesia after SPVAT lung wedge resection? Is it possible to inject local anesthetics(LAs) directly into the paravertebral space via the intrathoracic approach under thoracoscopic direct vision to achieve the purpose of TPB, thereby reducing postoperative pain? There is no report about it. Therefore, this study explored the analgesic effect of single-injection TPB via the intrathoracic approach under thoracoscopic direct vision for analgesia after SPVAT lung wedge resection.

METHODS

Patients

This study was approved by the Ethics Committee of Ningbo Medical Center, Lihuili Hospital, China (approval reference: KY2020PJ015). The study was performed in accordance with the Helsinki Declaration of 1964 and its later amendments. All participants provided informed consent to participate in the study. The trial was registered at www.chictr.

org.cn (registration number: ChiCTR2000034726). Sixty patients undergoing SPVAT lung wedge resection were enrolled between July 2020 and October 2020 in the Ningbo Medical Center Lihuili Hospital. The inclusion criteria for patients were as follows: age between 18 and 65 years, an American Society of Anesthesiologists (ASA) physical status of I ~ II, pneumothorax and bullae or lung nodules considered carcinoma in situ with a requirement for minimally invasive surgery. The exclusion criteria were patients with spinal deformity or spinal surgical history, allergies to LAs, and patients who did not consent to the procedure.

Patients were randomized divided into a control group and an observation group using a computer-generated random table, which was performed by an investigator who not involved in patient perioperative care and data collection. The investigator informed a nurse to prepare 20 ml of 0.375% ropivacaine or normal saline. Patients, surgeons, anesthesiologists, data acquisition personnel did not know the grouping.

Anesthesia

Standard monitoring was performed for electrocardiogram, blood pressure, heart rate, and oxygen saturation when the patients entered the operating theater. Rapid intravenous induction with 0.05 mg/kg midazolam, 0.2 µg/ kg sufentanil, 2 mg/kg propofol, and 0.6 mg/kg rocuronium was performed. After tracheal intubation, a bronchial occluder was optimally positioned using fiberoptic bronchoscope guidance. Mechanical Ventilation was given to ensure a tidal volume of 6 ml/kg, and a respiratory rate of 10-12 times/min. The intraoperative anesthesia maintenance was 8 mg·kg⁻¹ h⁻¹ propofol, 0.1 μg kg⁻¹ min⁻¹ remifentanil, and 0.2 mg (kg 0.5 h) rocuronium with a single intravenous injection. Adjusted the dose of the anesthetic to maintain the bispectral index between 45 and 55.

Surgery

A 4-cm incision was made between the four and five intercostals in the axillary midline of the two groups. SPVAT lung wedge resection was performed. The pulmonary nodules were confirmed as benign tumor or carcinoma in situ by intraoperative pathology.

Analgesia Methods

In the observation group, a scalp needle with an extended tube was inserted into the paravertebral space at the T4 level under thoracoscopic direct vision before closing the chest (Fig. 1). One centimeter adjacent to the vertebrae were inserted vertically 0.5 cm under the parietal pleura with the needle (Fig. 2), and 20 ml 0.375% ropivacaine was injected. The control group was injected with the same volume of 0.9% saline at T4 level. Observed for 5 min to make sure there was no hemorrhage or hematoma. The two groups were given 0.2 µg/kg sufentanil and 2 mg tropisetron 10 min before the end of surgery. At the end of the surgery, the PCIA pump was connected to patients to administer 1.5 μg·kg⁻¹ sufentanil and 5 mg tropisetron diluted to 100 ml with normal saline. The parameters were set as a continuous dose of 2 ml/h, a bolus dose of 1 ml, and a locking time of 20 min.

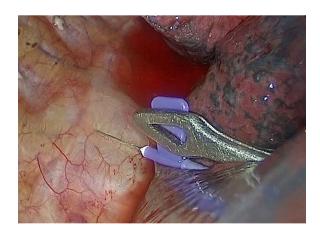


Fig. 1 TPB at T4 level via intrathoracic approach under thoracoscopic direct vision

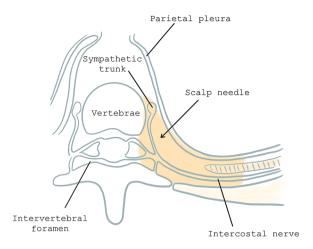


Fig. 2 Horizontal plane of paravertebral space. The *orange* area represents paravertebral space and site of scalp needle puncture

Data Collection

The primary outcome was the sufentanil consumption infused by the PCIA in the first 24 h after surgery. The secondary outcome was the visual analogue scale (VAS) during rest and while coughing at 6, 12, 24, and 36 h after surgery. The number of PCIA presses in the first 24 h after surgery were recorded. The incidence of adverse reactions such as nausea, vomiting, pruritus, somnolence, and atelectasis were recorded after surgery. The dermatomes of the sensory block were assessed 2 h after surgery using pinprick in the observation group.

Sample Size

The primary outcome was the sufentanil consumption infused by the PCIA in the first 24 h after surgery. Pilot testing using ten patients showed a mean [standard deviation] sufentanil consumption of 37.5 [4.66] μ g 24 h after surgery in the observation group and 43.5 [5.65] μ g in the control group. The requirement was 24 patients (12 in each group), which was calculated by the MedSci Sample Size Tools at a power of 0.8 with 0.05 alpha. Thus, to compensate for the possibility of missing data or dropouts, 30 patients were recruited for each group.

Statistical Analysis

Data were analyzed using SPSS version 23.0 (IBM Corp. Armonk, NY, USA). Continuous variables conforming to the features of normal distribution were represented as mean \pm standard deviation and analyzed using the t test. Repeated measurement data were analyzed using repeated ANOVA. Variables with a skewed distribution were presented as medians (quarand were compared using Kruskal-Wallis H test. Categorical data were presented as numbers (percentage) and analyzed using the Chi-square (χ^2) test. In all analvses, P < 0.05 was considered statistically significant.

RESULTS

A total of 60 patients were included and completed the study. All patients had successfully performed SPVAT lung wedge resection (Fig. 3). There were no significant differences of the patients' characteristics and surgical parameters between the two groups (Table 1).

Primary Outcome

The sufentanil consumption within the first 24 h after surgery in the observation group was significantly lower than that in the control group (34.2 \pm 1.9 μ g vs. 52.3 \pm 2.3 μ g; P < 0.001) (Fig. 4).

The Second Outcome

The VAS scores during rest and coughing at 6, 12 and 24 h were significantly lower than that at 36 h after surgery in the observation group. There was no significant difference in the VAS scores of rest and coughing at each time point in the control group. The VAS scores during rest and coughing in the observation group were significantly lower than those in the control group at 6, 12, and 24 h after surgery. There was no significant difference in VAS scores between the two groups at 36 h after surgery (Table 2).

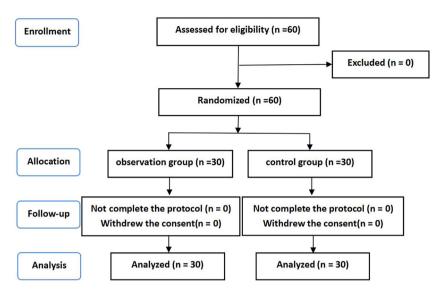


Fig. 3 Flow diagram of study

Table 1 Characteristics and surgical parameters

	Control group $(n = 30)$	Observation group $(n = 30)$	P value
Number of patients	30	30	
Type of disease			0.787
Pulmonary bulla or pneumothorax	10	11	
Pulmonary nodules	20	19	
Surgical site			0.796
Left	16 (53.3%)	15 (50.0%)	
Right	14 (46.7%)	15 (50.0%)	
Gender			0.795
Male	17 (56.7%)	16 (53.3%)	
Female	13 (43.3%)	14 (46.7%)	
Age (y)	48.9 ± 12.9	43.5 ± 11.9	0.100
Weight (kg)	65.4 ± 6.1	62.7 ± 7.1	0.115
Surgery time (min)	24.8 ± 4.5	25.2 ± 5.7	0.765

Data are presented as mean \pm standard deviation or number (%)

The observation group pressed the PCIA pump [0 (0-0) times] less than the control group [3 (2-4) times] (P < 0.001). The incidence of postoperative adverse reactions such as nausea and vomiting in the observation group were

significantly lower than those in the control group. No significant differences in the incidences of pruritus, somnolence, and atelectasis were observed between the two groups (Table 3). The dermatomes of the sensory block

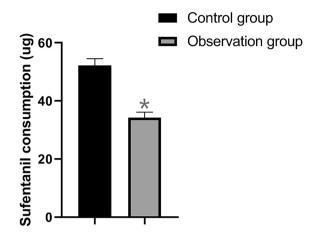


Fig. 4 Sufentanil consumptions within the first 24 h after surgery of the observation group and control group. $^*P < 0.001$

in the observation group were maintained at T1–T8 levels, and the main blocking area was at T2–T7 levels with 19 (63.3%) patients (Fig. 5).

DISCUSSION

Patients undergoing thoracic surgery have decreased postoperative lung function reserves. The severe pain after surgery prevents patients from breathing deeply with active coughing and sputum, resulting in the retention of respiratory secretions, causing hypoxemia, pneumonia, atelectasis, and even respiratory failure [17]. Adequate postoperative analgesia is conducive to early active cough and sputum disimprove lung function, postoperative complications, and promote postoperative recovery. SPVAT lung wedge resection has the advantages of less incision, shorter surgery time, less pain, and faster recovery. Therefore, the management of perioperative pain is often ignored, but in fact, the postoperative pain is still severe, which affects postoperative recovery [5, 18]. Currently, the commonly used postoperative analgesia is cumbersome and has many complications, and

Table 2 The VAS score after surgery

	0 7		
	Control group $(n = 30)$	Observation group $(n = 30)$	P value
During rest			
6 h after surgery	3.5 ± 0.7	2.5 ± 0.3	< 0.001*
12 h after surgery	3.4 ± 0.7	2.5 ± 0.2	< 0.001*
24 h after surgery	3.4 ± 0.7	2.4 ± 0.2	< 0.001*
36 h after surgery	3.2 ± 0.6	3.2 ± 0.5	0.805
F value	1.0	82.0	
P value	0.318	< 0.001#	
While coughing			
6 h after surgery	4.5 ± 0.4	3.1 ± 0.3	< 0.001*
12 h after surgery	4.8 ± 0.3	3.0 ± 0.3	< 0.001*
24 h after surgery	4.6 ± 0.3	3.0 ± 0.3	< 0.001*
36 h after surgery	4.4 ± 0.4	4.5 ± 0.4	0.637
F value	2.1	182.8	
P value	0.154	< 0.001#	

Data are presented as mean \pm standard deviation, *P < 0.001 Observation group vs Control group, *P < 0.001 36 h after surgery vs 6 h, 12 h, and 24 h after surgery

Table 3	The incid	lence of	adverse	reactions	after	surgery

	Control group $(n = 30)$ (%)	Observation group $(n = 30)$ (%)	P value
Nausea	12 (40.0%)	3 (10.0%)	0.007*
Vomit	9 (30.0%)	2 (6.7%)	0.020*
Pruritus	4 (13.3%)	0 (0.0%)	0.121
Somnolence	3 (10.0%)	0 (0.0%)	0.236
Atelectasis	3 (10.0%)	1 (3.3%)	0.605

Data are presented as number (%)*P < 0.05

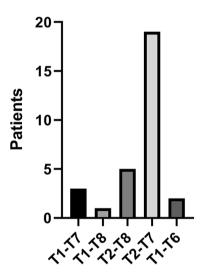


Fig. 5 Dermatomes of the sensory block in the patient of observation group. Frequency of sensory segments blockade to sharp touch at 2 h later after surgery

lacks a postoperative analgesia suitable for SPVAT wedge resection.

Currently, postoperative analgesia after thoracic surgery mainly includes thoracic epidural analgesia, PCIA, TPB, erector spinae plane block and intercostal nerve block [7, 9, 19]. Thoracic epidural analgesia is the gold standard for postoperative analgesia after thoracic surgery, but there are still complications such as epidural hematoma or infection, nerve damage or hypotension [11–13]. While PCIA is simple and convenient for postoperative care [14], but it requires large amounts of systemic opioids, with the adverse reactions such as nausea and vomiting [15]. TPB is a technique of injecting LAs into the paravertebral space, which produces a

similar effect to unilateral epidural block and effectively reduce postoperative pain [16, 20–22]. However, the requirements for ultrasound imaging and block technology are high, the operation space is limited and the difficulty is greater, the success rate is low. There is a possibility of puncturing the pleura and damaging the intercostal nerve or sympathetic chain [8, 21, 23, 24]. Compared with TPB, erector spinae plane block has a higher success rate, but requires higher ultrasound imaging and block technique [19]. Intercostal nerve block has a limited block range and can only suppress incision pain [9]. The above-mentioned analgesia methods require well-trained anesthesiologists, which are cumbersome, have more complications, and have a high failure rate. They are not suitable for SPVAT wedge resection.

Is there a simpler, faster operation, fewer complications, and an effective analgesic method that can apply postoperative analgesia after SPVAT wedge resection? Therefore, this study aimed to investigate the analgesic effect of single-injection TPB via the intrathoracic approach for analgesia after SPVAT lung wedge resection.

In this study, we used single-injection TPB at the T4 level. Before the end of the surgery, the scalp needle with the extended tube was inserted into the paravertebral space at the T4 level under thoracoscopic direct vision. One centimeter adjacent to the thoracic vertebrae were inserted vertically 0.5 cm under the parietal pleura and 20 ml 0.375% ropivacaine was injected. The dermatomes of the sensory block

in the observation group were maintained at T1-T8, and 19 (63.3%) were at T2-T7, indicating that the LAs was effectively spreading into the paravertebral space. The study by Gacio et al. reported ultrasound-guided single-injection paravertebral block allows proper control of acute pain with less intraoperative and postoperative consumption of opioids in major breast surgery [25]. The sufentanil consumption, VAS score, and number of PCIA presses in the observation group were significantly lower than those of the control group in the first 24 h after surgery. This fully showed that the postoperative pain of the observation group was significantly reduced, and once again proved that the single-injection TPB under thoracoscopic direct vision was indeed effective and feasible. LAs spread in the paravertebral spaces of 2-4 vertebral planes above and below the T4 segment, basically covering the area of surgery and incision, effectively reducing postoperative pain and reducing the amount of intravenous opioids after surgery. This is similar to the results of Gacio et al. [25]. The duration of the single injection TPB was related to the concentration of the LAs, generally within 12-24 h. The VAS scores during rest and coughing at 6, 12, and 24 h were significantly lower than that at 36 h after surgery in the observation group. This result also proved once again that the duration of a single block is within 24 h. There was no significant difference in VAS scores between the two groups at 36 h after surgery. This might be due to the following reasons: (1) SPVAT wedge resection is less traumatic, and the postoperative pain is mainly within 24 h after the surgery, and the pain is less after 24 h. (2) The endogenous analgesic system is activated when TPB block the transmission of noxious stimuli to the center nervous system at the level of the spinal cord, which can continue to play an analgesic effect. (3) The effect of TPB is gradually subsided, so that the patient's tolerance threshold for pain is increased, and hyperalgesia is avoided. This also suggests that after the effect of a single injection SITPB is over, it may still promote the improvement of postoperative pain through other ways.

Thoracoscopy imaging has magnification function and the thoracic vertebral body and

paravertebral tissue were fully exposed after lung atrophy. The vertical distance between the parietal pleura and the intervertebral foramen is about 1.0 cm. This ensures that the LAs is completely injected into the paravertebral space. We chose the transmural pleura to insert the needle vertically at a depth of 0.5 cm to avoid the risk of damaging the nerve root, or even entering the spinal canal. Single-injection TPB via the intrathoracic approach under thoracoscopic direct vision is easy to operate. It only requires checking the Computer tomography image structure of the block site before the block and it can be completed quickly by the surgeon. Compared with conventional multiports thoracoscopic surgery, SPVAT lung wedge resection has fewer incisions and faster recovery. In this study, a single-injection TPB can meet the need for postoperative analgesia. Compared with the single-injection TPB in this study, the operation of ultrasound-guided TPB or erector spinal plane block was relatively complicated, and the advantages and significance of analgesia after this kind of surgery might not be obvious.

Arunakul et al. found that PVB can reduce postoperative opioid requirement, pain, and severity of nausea and vomiting in modified radical mastectomy [26]. In this study, the incidence of nausea and vomiting in the observation group were significantly lower than those in the control group, which may be due to the poor analgesic effect of the control group, the consumption of sufentanil was more, the use of opioids was larger, so the incidence of adverse reactions was higher. There was no statistical significance of the incidence of pruritus and somnolence between the two groups of patients. This may be because the concentration of opioids is not enough to cause skin pruritus and somnolence. There was no atelectasis in the two groups, which may be due to less lung tissue resection, shorter surgery time and one lung ventilation time.

The present study had several limitations. First, this study did not observe local anesthetic poisoning, high epidural block and other block-related complications. Second, this study did not observe pain-related indicators such as sufentanil consumption beyond 24 h after

surgery. Lastly, this study did not compare with ultrasound-guided single thoracic paravertebral block or erector spinal plane block in terms of operation time, success rate, sensory level, and complications. These are also our next research directions.

CONCLUSIONS

In conclusion, a single-injection TPB via the intrathoracic approach under thoracoscopic direct vision is simple and quick to operate, which can effectively reduce postoperative pain after SPVAT wedge resection, with fewer adverse reactions.

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Authorship Contributions. LHH conceived and design the study, analyzed the data, and drafted the manuscript; XX, JXH helped collect data. HT, JXH performed the thoracic paravertebral block.

Disclosures. Lihong Hui, Xia Xu, Hui Tian and Jinxian He have nothing to disclose.

Compliance with Ethics Guidelines. This study was approved by Ethics Committee of

Ningbo Medical Center, Lihuili Hospital, China (KY2020PJ015). This study was performed in accordance with the Helsinki Declaration of 1964 and its later amendments. The trial was registered at www.chictr.org.cn (registration number: ChiCTR2000034726). All participants consented to participate in this study and signed informed consent forms.

Data Availability. The data are available from the corresponding author on reasonable request.

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