Characteristics of Postoperative Pain After VATS and Pain-Related Factors: The Experience in National Cancer Center of China

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Purpose: No final conclusion has yet been reached on characteristics of postoperative pain and pain-related factors after video-assisted thoracoscopic surgery (VATS). This study was designed to explore features of acute severe pain and chronic post-surgical pain (CPSP), and the pain-related factors of VATS.

Patients and Methods: Data of patients who underwent VATS for lung cancer in Cancer Hospital, Chinese Academy of Medical Sciences between March 2017 and January 2019 were reviewed in this retrospective study. A numerical rating scale (NRS) was used for evaluating the intensity of postoperative pain including no pain (NRS=0), mild pain (NRS=1–3), moderate pain (NRS=4–6), and severe pain (NRS=7–10). Pain intensity was assessed daily within a week after operation, and also evaluated at 3 months postoperatively.

Results: One hundred and five (3.4%) of the 3072 patients enrolled experienced severe pain (NRS=7–10) on the 1st day after operation, and 17 (0.6%) on the 2nd day. Smoking history, three-port VATS, prolonged operation time, and without patient-controlled analgesia (PCA) were correlated to increased incidence of severe pain. Among all patients, 237 (7.7%) cases generated CPSP, and VATS type, operation time, duration of drainage, and severe pain on the 1st day were four independent risk factors related to CPSP.

Conclusion: Patients seemed to experience a lower incidence of acute severe pain and CPSP after VATS than traditional open surgery. Acute severe pain was correlated with smoking history, VATS type, operation time, and PCA; VATS type, operation time, duration of drainage, and severe pain on the 1st day postoperatively were four independent risk factors of CPSP.

Keywords: VATS, acute severe pain, chronic post-surgical pain, NRS, risk factor

Introduction

As one of the most common malignant diseases, lung cancer has a high incidence rate and threatens human health and life seriously.¹ A study reported that the age-standardized incidence rates of lung cancer were from 33.3 to 66.8 per 100,000 among males and 10.5 to 37.4 per 100,000 among females in most of the Americas, Europe, and Oceania populations during the period 2003–2007.² A report from the National Cancer Center of China showed that the crude incidence rate of lung cancer was 54.66/100,000, the age-standardized incidence rates by China population and by World population were 35.13/100,000 and 34.86/100,000, respectively.³ Surgical resection is still the most basic and important clinical treatment method at present.⁴ With the development of science and technology, and also with the

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improvement of medical services level, the traditional open thoracotomy has been gradually replaced by video-assisted thoracoscopic surgery (VATS) for resectable lung cancer.⁵ Postoperative acute chest pain which can lead to various adverse outcomes is a head-scratching problem. A large number of studies have focused on postoperative acute pain in order to analyze the characteristics of pain and find treatment strategy. VATS, as the minimally invasive surgery (MIS), may provide patients with the relieved acute pain compared to traditional open surgery in theory. No final conclusion has yet been reached on the characteristics of acute severe pain and pain-related factors after VATS however.

Chronic pain has been recognized as pain that persists past normal healing time, while chronic post-surgical pain (CPSP) is defined as pain that develops after a surgical procedure and persists at least 3 months after surgery, with the exclusion of all other causes of pain (infection, recurring malignancy) as well as pain from a preexisting pain problem.⁶ Patients are likely to complain of a variety of pain symptoms, such as aching, numbness, stabbing, cramping, and burning. Although MIS, the reported CPSP incidence after VATS still ranges from 20% to 47%.^{7,9} The risk factors related to CPSP have not been clearly established, especially for VATS.

This study was designed for the purpose of exploring the features of pain, including both acute severe pain and CPSP after VATS, and analyzing the related factors of postoperative pain with the aim of receiving better methods to relieve pain for lung cancer patients undergoing VATS.

Patients and Methods

Patients

This retrospective study was approved by the Institutional Review Board of National Cancer Center of China, Cancer Hospital, Chinese Academy of Medical Sciences. All subjects or their representatives gave their written informed consent after receiving a full explanation of the nature of the study. This study was conducted in accordance with the Declaration of Helsinki.

Clinical data of patients who underwent VATS for lung cancer in Cancer Hospital, Chinese Academy of Medical Sciences between March 2017 and January 2019 were reviewed and analyzed. All data were collected anonymously and confidentially and all procedures were in compliance with the Declaration of Helsinki. This

retrospective study was approved by the Institutional Review Board of National Cancer Center of China, Cancer Hospital, Chinese Academy of Medical Sciences. Exclusion criteria included being younger than 18-year old, suffering long-term chronic pain before operation, open surgery, reoperation after VATS, and tumor recurrence during the follow-up period.

Operation

VATS procedure was performed for all of the persons recruited. Operation types of lobectomy, segmentectomy, and wedge resection were included in this study, and total pneumonectomy was excluded. The procedures of single-port or three-port VATS were performed by 12 experienced surgeons, and intravenous patient-controlled analgesia (PCA) with opioids was administered to patients whose consent had been obtained before surgery.

Evaluation of Pain

A numerical rating scale (NRS=0–10) was used for evaluating the intensity of postoperative pain. The pain intensity scale was categorized as no pain (NRS=0), mild pain (NRS=1–3), moderate pain (NRS=4–6), and severe pain (NRS=7–10). The patients were requested to report the pain intensity twice on the 1st, 2nd, and 3rd day after operation (at 9 am and 3 pm), the most serious pain (when coughing, moving, rest or others) was collected and the mean score was used as the pain intensity indicator for analyzing for each day, then the pain intensity was recorded as no, mild, moderate, or severe according to the average NRS score. On the 4th to 7th days, the pain intensity was evaluated once (9 am) per day; telephone follow-up was used if the patients were discharged within 1 week. The pain intensity was assessed at 3 months after operation during the follow-up period.

Definition of CPSP

We defined CPSP similarly to Macrae:¹⁰ 1) Pain that developed after surgery. 2) Other causes for the pain should had been excluded (eg, continuing malignancy or chronic infection). 3) Exacerbation of a pre-existing problem as a cause for the pain had been ruled out. 4) The duration of pain was more than 3 months, and the $NRS \geq 3$. The highest intensity (when coughing, moving, or others) which was evaluated as the chronic pain intensity was collected for analyzing.

Treatment Strategy of Postoperative Pain

For patients having PCA, intravenous analgesics were selected for those with severe pain (NRS=7–10); for patients without PCA, intravenous analgesics including Parecoxib Sodium or Flurbiprofen Axetil was routinely provided within 2 days after operation. Oxycodone and Acetaminophen tablets for oral administration were available for patients during late hospitalization.

Statistical Analysis

SPSS 16.0 (IBM, Chicago, IL, USA) was used for data analysis. Quantitative variables were analyzed with the Student *t*-test according to data distribution (age, BMI, operation time in Table 1; age, BMI, operation time and duration of drainage in Table 2). Categorical variables were analyzed by Chi-square test or Fisher exact test, as appropriate. A *P*-value less than 0.05 was considered significant. Univariate analyses were performed to identify risk factors predicting severe pain and CPSP. Multivariable logistic regression was used to identify the independent factors of CPSP. After univariate analyses, variables including patient's general conditions and operation-related outcomes with a *P*-value less than 0.05 were included in a multivariable logistic regression model.

Table 1 Comparisons Between Patients with and without Severe Pain on 1st Day After Operation

Parameters	Severe Pain (NRS≥7)	No-Severe Pain (NRS<7)	P-value
Gender (male/female), cases	33/72	1134/1833	0.159
Age, years	57.2±9.8	56.8±11.7	0.746
BMI, Kg/m ²	25.5±2.7	25.9±3.2	0.280
Smoking before operation (yes/no), cases	69/36	1435/1532	<0.001
VATS type (single-port/three-port), cases	15/90	2421/546	<0.001
Operation type (lobectomy/segmentectomy/wedge resection), cases	63/19/23	1959/560/448	0.247
Operation time, min	139.2±20.5	133.4±24.2	0.015
Surgical site (left/right), cases	36/69	1071/1896	0.704
Number of chest tube (1/2), cases	103/2	2919/48	0.819
PCA (yes/no), cases	60/45	2069/898	0.006

Abbreviations: BMI, body mass index; NRS, numerical rating scale; PCA, patient-controlled analgesia.

Table 2 Comparisons Between Patients with and without CPSP

Parameters	CPSP	No-CPSP	P-value
Gender (male/female), cases	73/164	1094/1741	0.018
Age, years	55.7±10.8	56.9±11.7	0.112
BMI, Kg/m ²	26.0±2.7	25.9±3.3	0.675
Smoking before operation (yes/no), cases	143/94	1361/1474	<0.001
VATS type (single-port/three-port), cases	37/200	2399/436	<0.001
Operation type (lobectomy/segmentectomy/wedge resection), cases	147/54/36	1875/525/435	0.262
Operation time, min	138.6±21.3	133.2±24.2	<0.001
Surgical site (left/right), cases	85/152	1022/1813	0.955
Number of chest tube (1/2), cases	232/5	2790/45	0.541
Duration of drainage, days	3.7±0.7	3.4±0.8	<0.001

Abbreviations: BMI, body mass index; CPSP, chronic post-surgical pain; VATS, video-assisted thoracoscopic surgery.

Results

A total of 3072 patients were collected for analyzing in this study. Of these patients, 1167 (38%) were male and 1905 (62%) were female; age ranged from 28 to 75 years, with an average of 56.8±11.7 years old; 1504 (49.0%) patients had a smoking history before operation. Lobectomy, segmentectomy, and wedge resection were performed for 2022 (65.8%), 579 (18.8%), and 471 (15.3%) patients, respectively. Non-small cell lung cancer was confirmed for all patients by postoperative pathology. Single-port VATS was conducted in 2436 (79.3%) patients and three-port VATS in 636 (20.7%) patients. A total of 3022 (98.4%) patients had one chest tube and 50 (1.6%) had two tubes after operation, and the retention time of the chest tube was from 2 to 5 days.

The mean NRS score decreased with the passage of time, the change was shown in Figure 1. One hundred and five (3.4%) and 17 (0.6%) patients experienced severe pain on the 1st and 2nd day after operation, respectively (Figure 2). As shown in Table 1, some factors were related to the acute severe pain on the 1st day postoperatively, smoking history was related to a higher incidence of severe pain, while three-port VATS, prolonged operation time, and being without PCA could also lead to increased incidence of severe pain.

The incidence of CPSP was 7.7% (237 cases) in this study. As shown in Table 2, the risk factors related to CPSP included female patients, having smoking history,

Figure 1 The change of mean NRS (numerical rating scale) scores from the 1st to the 7th day postoperatively.

three-port VATS surgery, longer operation time, and prolonged duration of drainage. We also found that severe pain on the 1st and 2nd day after operation was closely related to CPSP (Table 3), 81.9% and 88.2% of patients with acute severe pain on the 1st and 2nd day developed CPSP, respectively.

Table 4 illustrated that VATS type, operation time, duration of drainage, and severe pain on the 1st day were four independent risk factors related to CPSP.

Discussion

VATS has become a selected operation type during recent years. Without compromising short-term and long-term outcomes, VATS can provide several benefits to patients compared with traditional open thoracotomy.¹¹ Post-surgical pain is one of the most common surgical complications, despite the advances in understanding pain and the development of new modalities for pain management.¹² Several studies demonstrate that 41–69% of patients have experienced moderate-to-severe pain after operation.^{13,14}

Whether VATS can reduce the pain intensity or not is still uncertain.

Data of 312 patients who received open thoracotomy or VATS were analyzed in a recent study designed by Wang et al,⁷ their results conveyed that the pain intensity was 3.91 ± 2.12 at rest and 5.73 ± 2.26 when moving in the VATS group on the 1st day after operation, and both of them were higher than in the open group. Whereas a study designed by Wildgaard et al¹⁵ gave the opposite results; their study confirmed that patients who underwent VATS had decreased pain intensity during the first 72 hours. In our study, the mean NRS scores were 2.60 ± 1.31 , 2.06 ± 1.11 , and 1.34 ± 0.88 on the 1st, 2nd, and 3rd day after operation, and the incidence of severe pain was 3.4% and 0.6% during the first 2 days. No patients experienced severe pain on the 3rd day after operation.

We also employed further analysis about the high-risk factors related to severe pain, and the results showed that patients with a smoking history had a high incidence of severe pain. Maybe smoking before the operation lead to a bad cough and the cough induced serious pain. Single-port

Figure 2 The different percentages for each pain intensity within the first week after operation.

VATS contributed to a lower incidence of severe pain compared with three-port VATS in our study, which was similar to previous results by Sun et al.¹⁶ Their results demonstrated that increased BMI, smoking history, and multi-port VATS had high incidence of moderate–severe pain during the first 24 hours after operation. There was another study which illustrated that besides pain intensity, the functional areas such as physical function, role function, emotional function, social function, and the overall health status of the patients with single-port were significantly better than those with three-port.⁶ PCA had ideal analgesic effect, patients with PCA had lower incidence of severe pain after operation.

CPSP was defined initially by Macrae and Davies⁸ as a persistent pain which has existed for a minimum of 2 months following a surgical procedure. There are many suggestions to revise this definition by the International Association of Pain Study (IAPS). Currently, 3 months is considered the minimal accepted duration to diagnose a patient with CPSP. CPSP places considerable physical and psychological burdens on individuals. An early study on CPSP reported that 22.5% of 5130 chronic pain patients seen in 10 UK clinics indicated major surgery as its source.¹⁷ Since then, studies have specified that the incidence of CPSP ranged from 10% to 70% depending on the surgery types.^{18,19} Wang et al²⁰ reported that 64.5%

Table 3 Univariate Analyses for the Relationship Between Severe Pain and CPSP

Pain Intensity After Operation	CPSP (n=237)	No-CPSP (n=2835)	P-value
Severe pain on the 1st day, cases			<0.001
Yes	86	19	
No	151	2816	
Severe pain on the 2nd day, cases			<0.001
Yes	15	2	
No	222	2833	

Abbreviation: CPSP, chronic post-surgical pain.

Table 4 Logistic Regression Analysis for the Independent Factors of CPSP

Related Factors	Incidence of CPSP			
	RR	95% CI		P-value
Gender	1.352	0.940	1.945	0.103
Smoking before operation	1.282	0.911	1.806	0.154
VATS type	4.536	3.736	5.508	<0.001
Operation time	1.008	1.001	1.016	0.024
Duration of drainage	1.372	1.108	1.697	0.004
Severe pain on 1st day	25.825	13.448	49.593	<0.001
Severe pain on 2nd day	3.080	0.464	20.449	0.244

Abbreviations: CPSP, chronic post-surgical pain; VATS, video-assisted thoracoscopic surgery.

of patients who underwent thoracotomy experienced CPSP, and Peng et al⁶ reported that 24.9% of patients who underwent open and VATS thoracotomy generated CPSP. Reported incidence of CPSP after VATS ranged from 13.3% to 47%,²¹ and the incidence of CPSP was 7.7% in our study. We thought that removing patients with chronic pain history from this study might be a critical reason for the lower incidence of CPSP.

Several operation-related factors have an influence on CPSP. Peng et al⁶ reported that age <60 years old, female gender, and prolonged duration of postoperative chest tube drainage (≥ 4 days) were related to a high incidence of CPSP. Bayman²¹ reported that severity of acute pain was associated with chronic pain after thoracic surgery. Buchheit²² reported that prolonged duration of chest tube drainage had an increased incidence of CPSP. Our results demonstrated that female patients, smoking history, three-port VATS, longer operation time, and prolonged duration of drainage were correlated to the high incidence of CPSP, meanwhile, we found that patients who experienced severe pain during the first 2 days postoperatively had a higher chance to experience CPSP. VATS type, operation time, duration of drainage, and severe pain on the 1st day were four independent risk factors for high incidence of CPSP.

The shortcomings of the study were that no open surgery data was collected and analyzed, so we could not compare the differences including characteristics of acute pain and CPSP, and influence factors between open surgery and VATS.

Conclusion

VATS seems to have lower incidence of acute severe pain and CPSP, with smoking history, VATS type, operation time, and PCA being related to acute severe pain; VATS type, operation time, duration of drainage, and severe pain on the 1st day postoperatively are independent risk factors of CPSP.

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Disclosure

The authors report no conflicts of interest in this work.

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