Original
Article

Extraluminal Placement of a Bronchial Blocker Compared with Carbon Dioxide Artificial Pneumothorax in Infants Undergoing Video-Assisted Thoracoscopic Surgery

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Objective: To investigate the safety and effectiveness of extraluminal placement of a bronchial blocker compared with carbon dioxide (CO₂) artificial pneumothorax in infants undergoing video-assisted thoracoscopic surgery (VATS).

Methods: The study involved 33 infants (group A) who underwent one-lung ventilation (OLV) with extraluminal placement of a bronchial blocker and 35 other infants (group B) who underwent CO₂ artificial pneumothorax. Clinical characteristics, the degree of lung collapse, and complications were compared.

Results: The degree of lung collapse in group A was significantly higher than that in group B at T2 and T3. The mean arterial pressure (MAP) of group B was significantly lower than that of group A at 10 min and 30 min after OLV. The partial pressure of carbon dioxide (PaCO₂) of group B was significantly higher than that of group A at 30 min after OLV. The incidence of hypotension in group B was higher than that in group A.

Conclusion: Compared with CO₂ artificial pneumothorax, extraluminal placement of a bronchial blocker is associated with a better degree of lung collapse, fewer episodes of hypotension, and lower PaCO₂ accumulation during OLV in infants undergoing VATS.

Keywords: one-lung ventilation, bronchial blockers, artificial pneumothorax, infants, anesthesia

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Introduction

In recent years, video-assisted thoracoscopic surgery (VATS) has been increasingly used in pediatric thoracic surgery. In this process, lung collapse is induced to provide an adequate surgical field of vision. Carbon dioxide (CO₂) artificial pneumothorax is usually used to provide adequate operating conditions for VATS.^{1,2)} However, this method may lead to hemodynamic changes, such as hypotension.³⁾ The use of a bronchial blocker can induce collapse of the lung on the operative side of the chest, and the mode of ventilation can be changed at any time during the operation. The secretions of the lung on the operation side can be aspirated at any time, and this is widely applied with good effect.^{4,5)} A fiberoptic bronchoscope is used to locate the bronchial blocker after bronchial blocker placement. As the airway diameter of

infants is relatively small, it can be difficult to operate a fiberoptic bronchoscope. Bronchial blocker placement in the endotracheal tube may increase airway pressure and reduce sufficient ventilation. Therefore, during VATS in infants, a bronchial blocker can be placed outside the endotracheal tube for one-lung ventilation (OLV) to achieve lung collapse.^{6–9)} In this study, the data of infants undergoing VATS in our hospital from February 2018 to December 2020 were retrospectively analyzed to compare the safety and feasibility of extraluminal bronchial blocker placement and CO₂ artificial pneumothorax in those patients.

Materials and Methods

The Ethics Committee of Fujian Maternal and Child Health Hospital approved this retrospective study. The study was conducted following the Declaration of Helsinki's ethical standards and its later amendments.

From February 2018 to December 2020, data were collected for 68 children through the medical record system, including 33 children in the bronchial blocker group (group A) and 35 children in the CO₂ artificial pneumothorax group (group B). Relevant general clinical data were collected (**Table 1**). The inclusion criteria were as follows: the infants were aged 2–12 months and underwent video-assisted thoracoscopic pulmonary lobectomy or segmentectomy in our hospital; extraluminal placement of a bronchial blocker or artificial pneumothorax with CO₂ was used to induce lung collapse; the patient's diagnosis was congenital cystic adenomatoid malformation of lung sequestration; and the family members signed informed consent. The exclusion criteria were as follows: the patient had other severe medical conditions, such as pulmonary insufficiency or liver and kidney insufficiency; and the patient underwent alternative surgical procedures or was converted to conventional open-chest surgery intraoperatively. The family members of the patient were informed of the anesthetic and surgical options before the surgery, and different lung collapse options were selected according to the decision of the anesthesiologist and surgeon.

All infants fasted for 4 hours and did not consume water for 2 hours. They were sedated with 0.1 mg/kg midazolam by intravenous injection. After entering the operating room, physiologic saline was infused. Heart rate (HR), noninvasive blood pressure, electrocardiograph (ECG), and oxygen saturation (SpO₂) were monitored. Anesthesia was induced by intravenous injection

of propofol at 2–3 mg/kg, fentanyl at 3–5 µg/kg, and rocuronium at 0.6 mg/kg.

In group A, the glottis was exposed under a visual laryngoscope after mask inhalation of oxygen for 2 minutes. The smallest 5F bronchial blocker (Hangzhou Tappa Medical Technology Co., Ltd.) was inserted into the trachea. An uncuffed endotracheal tube was also inserted into the trachea to place the bronchial blocker outside the endotracheal tube. The endotracheal tube was adjusted approximately 1–2 cm away from the carina of the trachea using a fiberoptic bronchoscope, and the bronchial blocker was guided into the pulmonary region of the operative side. Next, 1–3 mL of air was injected into the cuff of the bronchial blocker. The anesthesiologist auscultated both sides of the chest, and disappearance of the sound of pulmonary respiration on the operative side indicated that the bronchial blocker was in the correct position. In this study, the bronchial blocker was placed in the right main bronchus, the right middle and right lower bronchus, the left main bronchus, and the left lower bronchus, depending on the surgical scope. After the cuff was deflated, the endotracheal tube was connected to the anesthesia machine for two-lung mechanical ventilation. The following pressure control mode was adopted: fraction of inspired oxygen (FiO₂) of 0.5, positive end-expiratory pressure (PEEP) of 3–5 mmHg, inspiratory/expiratory (I:E) ratio of 1:1.5, tidal volume (VT) of 8–10 mL/kg, respiratory frequency (R) of 30–35 times per minute, and oxygen flow rate of 2–3 L/min. Before skin incision, a fiberoptic bronchoscope was used again to determine the position of the cuff of the bronchial blocker. After entering the thoracic cavity on the operative side, the correct amount of air was injected into the cuff of the bronchial blocker, and one-lung mechanical ventilation was performed. The following pressure control mode was adopted: FiO₂ of 0.8–1.0, PEEP of 3–5 mmHg, I:E ratio of 1:1.5, VT of 6–8 mL/kg, R of 30–35 times per minute, and oxygen flow rate of 2–3 L/min. No additional CO₂ insufflation was needed in this group.

In group B, the endotracheal tube was inserted into the principal bronchus, and mechanical ventilation was performed after concordant auscultatory breath sounds were noted in both lungs. After the infant was in the lateral position, CO₂ insufflation (CO₂ pressure = 8–10 mmHg) was used to create an artificial pneumothorax inducing lung collapse on the operative field side.

After anesthesia induction, the radial artery and right internal jugular vein were punctured and catheterized.

Table 1 Clinical characteristics and outcomes of the two groups

	Group A	Group B	P value
Number	33	35	-
Male/female	18/15	19/16	-
Age (month)	6.9 ± 2.3	6.4 ± 2.7	0.79
Weight (kg)	7.0 ± 1.4	6.8 ± 1.5	0.59
OLV time (min)	51.5 ± 8.2	52.3 ± 8.4	0.68
Length of drainage (day)	2.5 ± 0.6	2.7 ± 0.6	0.43
Length of hospital stay (day)	5.4 ± 0.8	5.7 ± 0.7	0.49
Surgical procedure (case)			
Pulmonary lobectomy	14	14	0.90
Pulmonary segmentectomy	15	17	
Partial pulmonary resection	4	4	

OLV: one-lung ventilation

The invasive mean arterial pressure (MAP) was monitored, and blood samples were collected for blood gas analysis during the operation. Arterial oxygen saturation was maintained between 92% and 96%. Remifentanyl at 0.1–0.5 µg/kg/min and 1–3% sevoflurane were used to maintain anesthesia during the operation. The doses of remifentanyl and sevoflurane were adjusted according to the monitoring values of anesthesia depth and blood pressure changes, HR, and end-expiratory carbon dioxide partial pressure (PETCO₂). If necessary, rocuronium 0.1 mg/kg was added. Recruitment maneuvers were performed in all infants at the end of OLV before restarting two-lung ventilation (TLV). In this study, a lung recruitment maneuver was performed three times, with airway pressure sustained at 30 cm H₂O for 15–20 s.¹⁰⁾ Notably, during the operation, norepinephrine at 0.01–0.05 µg/kg/min was injected if the blood pressure was lower than 20% of the baseline. If the oxygen saturation was lower than 90%, OLV was suspended, double-lung ventilation was resumed, and the bronchial blocker position was checked. OLV was performed after oxygen saturation was restored to >96%.

Data were collected from the computer record system, which included (1) general data such as age, sex, weight, time of OLV, length of drainage, and length of hospital stay; (2) the MAP of infants before OLV (T1), 10 min after OLV (T2), 30 min after OLV (T3), and 10 min after the end of OLV (T4); (3) the degree of lung collapse evaluated by the operating surgeon using a verbal rating scale from 0 (no lung deflation) to 10 (complete collapse) at the time of T2 and T3;¹¹⁾ (4) incidence of intraoperative hypoxemia and hypotension during the operation (intraoperative hypoxemia was defined as SpO₂ <90%; hypotension was defined as blood pressure

lower than 20% of the baseline); and (5) incidence of postoperative complications including pneumothorax, atelectasis, and chylothorax. An arterial blood sample was taken for blood gas analysis at T1, T2, T3, and T4 by the anesthetist during the operation. The PaCO₂ value, oxygenation index (PaO₂/FiO₂), and alveolar-arterial oxygen partial pressure difference (PA-aO₂) data were also collected.

Statistical analysis

Statistical analysis was performed using SPSS software (25.0 Version, IBM Corp., Armonk, NY, USA). Data with a normal distribution were presented as the mean ± SD, and differences were evaluated using the t-test. Nonparametric data were presented as medians, and differences were evaluated using Wilcoxon rank sum. Categorical variables were shown as frequencies (percentages), and differences were evaluated using the χ^2 test. A comparison of pulmonary gas exchange parameters pre-OLV, during OLV, and post-OLV was performed using repeated-measures analysis of variance. A P value of less than 0.05 was defined as statistically significant.

Results

The clinical information and outcomes of the two groups are shown in **Table 1**. There were no significant differences between the two groups in terms of clinical parameters, including age, sex, weight, time of OLV, length of drainage, and length of hospital stay ($P > 0.05$). There were no significant differences between the two groups in length of drainage and hospital stay ($P > 0.05$). In group A, there were 14 cases of lobectomy, 15 cases

Table 2 The comparison of perioperative hemodynamics and the degree of lung collapse score of the two groups

	Group A	Group B	P value	Z value
T1				
MAP (mmHg)	58.7 ± 7.0	56.7 ± 8.1	0.27	-
HR (bpm)	122.1 ± 8.4	119.7 ± 11.3	0.30	-
T2				
MAP (mmHg)	57.12 ± 7.6	42.8 ± 4.9 ^a	<0.001	-
HR (bpm)	118.6 ± 10.2	120.8 ± 10	0.37	-
The degree of lung collapse score	9.0 (8.5,9.0)	8.0 (8.0,9.0) ^a	<0.001	-3.489
T3				
MAP (mmHg)	52.6 ± 7.6	41.9 ± 7.3 ^a	<0.001	-
HR (bpm)	124.5 ± 7.3	125.2 ± 10.7	0.75	-
The degree of lung collapse score	10.0 (9.0,10.0)	9.0 (9.0,9.0) ^a	<0.001	-4.195
T4				
MAP (mmHg)	57.9 ± 4.0	59.0 ± 3.4	0.17	-
HR (bpm)	119.3 ± 9.8	120.7 ± 9.5	0.54	-

HR: heart rate; MAP: the mean arterial pressure. ^aP <0.001 (indicate compared with group A).

of pulmonary segmentectomy, and 4 cases of partial pulmonary resection. In group B, there were 14 cases of lobectomy, 17 cases of pulmonary segmentectomy, and 4 cases of partial pulmonary resection. These results indicate that the two sets of data were comparable and homogeneous.

The comparison of the degree of lung collapse scores between the two groups is shown in **Table 2**. The degree of lung collapse score in group A was significantly higher than that in group B at T2 and T3 ($P < 0.001$, $z = -3.489$ and $P < 0.001$, $z = -4.195$). A comparison of hemodynamic parameters between the two groups is shown in **Table 2**. There was no significant difference in MAP between the two groups at T1 and T4, but the MAP in group B was significantly lower than that in group A at T2 and T3 ($P < 0.001$). There was no significant difference in HR between the two groups at any time point ($P > 0.05$). A comparison of the PaCO₂ between the two groups is shown in **Fig. 1**. There were no significant differences in PaCO₂ between the two groups at T1, T2, or T4. At T3, the PaCO₂ in group B was significantly higher than that in group A ($P < 0.001$).

The comparison of PaO₂/FiO₂ and PA-aO₂ between the two groups is shown in **Fig. 2**. There were no significant differences in PaO₂/FiO₂ and PA-aO₂ between the two groups at any time point. However, there were significant differences when comparing these parameters at different time points in each group. In group A, compared with T1, the PA-aO₂ gradient at T2 and T3 significantly

increased by 21.03 (95% CI: 18.61–23.45) and 22.67 (95% CI: 19.95–24.35), respectively ($P < 0.001$); the PaO₂/FiO₂ ratio at T2 and T3 significantly decreased by 175.33 (95% CI: 164.21–186.46) and 190.61 (95% CI: 181.02–200.19), respectively ($P < 0.001$). Compared with T4, the PA-aO₂ gradient at T2 and T3 significantly increased by 22.15 (95% CI: 19.95–25.38) and 23.79 (95% CI: 21.78–25.80), respectively ($P < 0.001$), whereas the PaO₂/FiO₂ ratio at T2 and T3 significantly decreased by 131.78 (95% CI: 125.90–137.67) and 147.06 (95% CI: 143.47–150.65), respectively ($P < 0.001$). In group B, compared with T1, the PA-aO₂ gradient at T2 and T3 significantly increased by 23.43 (95% CI: 21.28–25.57) and 26.91 (95% CI: 24.9–28.9), respectively ($P < 0.001$). The PaO₂/FiO₂ ratio at T2 and T3 significantly decreased by 161.46 (95% CI: 152.73–170.19) and 178.37 (95% CI: 168.99–187.75), respectively ($P < 0.001$). Compared with T4, the PA-aO₂ gradient at T2 and T3 significantly increased by 22.63 (95% CI: 20.50–24.77) and 26.11 (95% CI: 24.12–28.11), respectively ($P < 0.001$). The PaO₂/FiO₂ ratio at T2 and T3 significantly decreased by 159.40 (95% CI: 154.89–163.93) and 176.31 (95% CI: 172.69–179.94), respectively ($P < 0.001$).

The incidence of intraoperative and postoperative complications between the two groups is shown in **Table 3**. The hypotension incidence in group B was significantly higher than that in group A ($P < 0.001$). Hypoxemia occurred in both groups, but the incidence was not significantly different between the two groups ($P = 0.80$).

Fig. 1 The comparison of the PaCO₂ between the two groups. OLV: one-lung ventilation; PaCO₂: partial pressure of carbon dioxide

There was no significant difference in the incidence of postoperative complications between the two groups.

Discussion

Compared to conventional surgery, VATS is a minimally invasive method in thoracic surgery and has many advantages, such as a small incision, less postoperative pain, fewer postoperative complications, and a shorter hospital stay, promoting the postoperative recovery of patients. Therefore, VATS has been widely accepted by parents for their children.¹²⁾ During the operation, it is necessary to collapse the lung on the operating side to expose the surgical field of vision, which is needed and beneficial for the surgeon to perform the operation. Collapse of the unilateral lung during the operation can be implemented using a bronchial blocker, a double-lumen tube, and CO₂ artificial pneumothorax. However, the smallest model of the double-lumen tube is only suitable for children over 8 years old.¹³⁾ In VATS, intrathoracic positive pressure injection of CO₂ can provide adequate lung collapse and promote visual surgical field exposure. However, this approach may lead to the accumulation of CO₂.^{14,15)} The use of a bronchial blocker is suitable for infants and young children. In general, the bronchial blocker is placed in the endotracheal tube and located via fiberoptic bronchoscope guidance. When infants undergo endotracheal intubation, an endotracheal tube of ID 3.5# or 4.0# is generally selected. Due to the

limitation of the endotracheal tube's inner diameter, the fiberoptic bronchoscope and bronchial blocker cannot enter the endotracheal tube simultaneously. Therefore, we typically place the bronchial blocker outside the endotracheal tube to implement lung collapse for infants under 1 year old.

In this study, a single-lumen endobronchial tube was used for endotracheal intubation in both groups. Lung collapse was induced during surgery via the compression of CO₂ flow or the bronchial blocker to provide surgical vision and operating space. At T2 and T3, the degree of lung collapse of group A was greater than that of group B, indicating that the visual surgical field exposed by the bronchial blocker was better than that by CO₂ artificial pneumothorax. Zheng et al. and Yan et al. reported that the use of a bronchial blocker for lung collapse provides an adequate surgical field of vision.^{9,16)}

The MAP of group B was significantly lower than that of group A at T2 and T3. The incidence of intraoperative hypotension in group B was significantly higher than that in group A. Mohtar et al. also reported no significant change in blood pressure before, during, or after bronchial blocker use.¹⁷⁾ We considered that the pressure of CO₂ during CO₂ artificial pneumothorax could compress the heart and large blood vessels, resulting in reduced cardiac blood volume, transient or persistent hypotension, and even the need to use vasoactive drugs.¹⁸⁾

At T3, the PaCO₂ of group B was significantly higher than that of group A. We believe that this finding might have been due to the accumulation of CO₂ in the chest cavity after 30 minutes of artificial pneumothorax in the infants of group B, which resulted in acidosis and elevated PaCO₂ levels in the arterial blood. This observation was consistent with the research of Lin et al.¹⁴⁾ However, there was no significant difference in PaCO₂ between the two groups at T4. The reason might be due to the following operations performed at the end of the surgery. First, OLV was stopped immediately, and TLV was resumed. Second, the CO₂ was evacuated from the infants' thoracic cavity by the surgeon. Third, the infant was switched from the lateral to the supine position, and the lung recruitment maneuver was carried out. It has been reported that lung recruitment maneuvers improve oxygenation during OLV.¹⁹⁾ Therefore, the PaO₂ and PaCO₂ levels were significantly improved in blood gas analysis 10 minutes after OLV.

Fig. 2 The comparison of PaO₂/FiO₂ and PA-aO₂ between the two groups. A1 showed the comparison of PaO₂/FiO₂ at the time of T1, T2 and T4 between the two groups. A2 showed the comparison of PaO₂/FiO₂ at the time of T1, T3, and T4 between the two groups. B1 showed the comparison of PA-aO₂ at the time of T1, T2, and T4 between the two groups. B2 showed the comparison of PA-aO₂ at the time of T1, T3, and T4 between the two groups. FiO₂: fraction of inspired oxygen; OLV: one-lung ventilation; PA-aO₂: alveolar-arterial oxygen partial pressure difference; PaO₂: partial pressure of oxygen; PaO₂/FiO₂: Oxygenation Index; T1: before OLV; T2: 10 min after OLV; T3: 30 min after OLV; T4: 10 min after the end of OLV

Table 3 Comparison of intraoperative and postoperative complications of the two groups

Parameter	Group A	Group B	P value
Number of patient	33	35	-
Intraoperative hypotension: case (%)	5 (15.2)	28 (80) ^a	<0.001
Intraoperative hypoxemia: case (%)	15 (45.5)	17 (48.6)	0.88
Postoperative complications			
Pneumothorax: case (%)	2 (6.0)	3 (8.6)	0.71
Atelectasis: case (%)	1 (3.0)	2 (5.7)	0.61
Chylothorax: case (%)	1 (3.0)	0 (0)	0.31

^aP < 0.001 (indicate compared with group A).

During OLV in infants in the lateral decubitus position, the risk of hypoxemia is higher due to increased V/Q mismatch, compression of the healthy-side lung, and collapse of the surgical-side lung with higher oxygen consumption.^{20,21} In this retrospective study, we observed no significant differences in the PaO₂/FiO₂

ratio or PA-aO₂ between the two groups at any time point. However, at T2 and T3, the PaO₂/FiO₂ ratio decreased, and the PA-aO₂ increased in both groups, indicating a change in pulmonary function and a higher risk of hypoxemia in OLV. At T4, when OLV was stopped for 10 minutes, the PaO₂/FiO₂ ratio and PA-aO₂ returned

to baseline levels. Hale et al. reported the same situation in infants undergoing OLV.²²⁾ There was no significant difference in the incidence of hypoxemia between the two groups in this study. The state of hypoxemia can be quickly improved by suspending the operation and performing TLV.

This research had some limitations. The sample size included in this study was relatively small. In addition, our data collection might have been biased. Relatively, few indicators of pulmonary function were adopted in the study, a subjective rating scale was used to evaluate the degree of pulmonary collapse, and the data could have become biased in the recording process, which, in turn, could affect the accuracy of the results. In addition, this study was retrospective rather than a prospective case-control study, which limited its statistical power, but we still believe that the results have clinical significance. Future research should consider variations in factors and examine a larger sample to confirm our conclusions.

Conclusion

This study showed that OLV with extraluminal placement of a bronchial blocker is a safe choice for infants undergoing VATS. In our study, the gas exchange changed during OLV but returned to baseline following resumption of TLV. Compared with CO₂ artificial pneumothorax, OLV with extraluminal placement of a bronchial blocker appears to provide a higher degree of surgeon-rated lung collapse, fewer episodes of significant hypotension, and lower PaCO₂ accumulation for infants undergoing VATS.

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Disclosure Statement

All authors declared that they had no competing interests.

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