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**Unilateral pulmonary hemorrhage caused by negative pressure pulmonary edema: A case report**

Park HJ *et al*. Unilateral pulmonary hemorrhage

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**Abstract**

BACKGROUND

Unilateral pulmonary hemorrhage is typically reported in young and healthy men with upper respiratory tract obstruction during anesthesia in special situations. Negative pressure in the lungs is created, resulting in negative pressure pulmonary edema (NPPE).

CASE SUMMARY

A 78-year-old male patient diagnosed with spinal stenosis was admitted to receive a unilateral laminectomy with bilateral decompression. The patient had been diagnosed with hypertension four years earlier and asthma more than 70 years earlier. We experienced a unilateral alveolar hemorrhage associated with NPPE that occurred in a longstanding asthma patient who bit the intubated endotracheal tube for a short period during posture change at the end of surgery. Because diffuse alveolar hemorrhage accompanied by NPPE was caused in this case by airway obstruction in an older patient with asthma without known risk factors, anesthesiologists should be careful not to induce airway irritation during anesthesia awakening in asthma patients.

CONCLUSION

Because diffuse alveolar hemorrhage accompanied by NPPE can occur, anesthesiologists should take care not to induce airway irritation.

**Key Words:** Airway; Anesthesia; Asthma; Hemorrhage; Pulmonary edema; Case report; Surgery

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**Core Tip:** In this case, because diffuse alveolar bleeding occurred after negative pressure pulmonary edema without risk factors, anesthesiologists should be careful not to induce irritation during anesthesia awakening in asthma patients.

**INTRODUCTION**

Diffuse alveolar hemorrhage (DAH) is characterized by damage to the alveolar microvessels and accumulation of red blood cells in the alveoli. DAH is a syndrome in which respiratory failure occurs due to hemoptysis, anemia, and hypoxemia[1,2]. Pulmonary imaging may be important in the diagnosis. The most frequent causes of DAH are forms of systemic vasculitis such as anti-glomerular basement antibodies, granulomatosis with polyangiitis, microscopic polyangiitis, capillaritis, vasculitis, or complications that can occur in patients with diseases such as Wegener’s granulomatosis, Goodpasture’s syndrome, and systemic lupus erythematosus[3].

DAH may also be associated with surgery and anesthesia. It has been reported that, especially in young, healthy, and muscular men with upper respiratory tract obstruction during anesthesia, negative pressure in the lungs is created, resulting in negative pressure pulmonary edema (NPPE) and, in rare cases, bilateral pulmonary hemorrhage[3]. It has also been reported that one-sided alveolar or pulmonary hemorrhage may occur in cases of the development of negative intrathoracic pressure owing to upper respiratory tract obstruction or administration of a large amount of fluid to a patient in the lateral position or in the presence of a disease such as right atrial failure[3,4]. DAH, although often spontaneously resolving, can sometimes lead to fatal complications.

In the case we report here, we attempted to remove the patient’s endotracheal tube under deep anesthesia at the end of surgery in order to avoid irritation of the airway in a patient who had lived with asthma for many years. However, we report this case with a literature review and informed consent to publication was obtained from the guardian because we experienced a unilateral DAH associated with NPPE that occurred after a short period of airway obstruction after the patient bit the intubated endotracheal tube during posture change.

**CASE PRESENTATION**

***Chief complaints***

A 78-year-old male patient who complained of low back pain was diagnosed with spinal stenosis.

***History of present illness***

He was admitted to receive a unilateral laminectomy with bilateral decompression.

***History of past illness***

No other past history was reported.

***Personal and family history***

The patient had been diagnosed with hypertension four years earlier and asthma more than 70 years earlier and was prescribed fluticasone furoate and vilanterol inhalers for the treatment of asthma; however, these medications were not used consistently.

***Physical examination***

Physical examination of the patient was not significant.

***Laboratory examinations***

The results of pulmonary function tests were forced vital capacity 2.53 L (61%), forced expiratory volume at 1 s 1.37 L (52%), and forced expiratory volume at 1 s/forced vital capacity 54.

***Imaging examinations***

Chest radiographs showed subsegmental atelectasis in the left lower lobe and mild cardiomegaly (Figure 1). Transthoracic echocardiography performed prior to surgery showed concentric left ventricular hypertrophy, trivial mitral regurgitation, and a mild relaxation abnormality with other findings within normal limits. As premedication, hydrocortisone 100 mg was administered on the morning of surgery. Lung sounds confirmed after the patient entered the operating room were normal, and anesthesia was initiated after prophylactic administration of salbutamol inhalants.

***Anesthesia and follow-up***

General anesthesia was induced with thiopental 250 mg and fentanyl 100 mcg, and the patient was intubated without special difficulty after administration of rocuronium 50 mg. Anesthesia was maintained at 50% N2O and 5%-7% desflurane. The operation was performed with the patient in the prone position, and there were no special events during the operation. The arterial blood gas analysis performed during the operation showed FiO2 0.5 to pH 7.420, pO2 242.5, pCO2 36.7, and base excess-0.6, and hemoglobin level was 12.2 g/dL. At the end of the operation, sugammadex 200 mg was administered for reversal of muscle relaxation, the patient’s position was changed from prone to supine under deep anesthesia, and the bispectral index score at that time was in the 40s. Subsequently, several aspiration attempts were made to remove the secretions from the trachea and oral cavity in order to manage the tracheal tube, but the patient was clenching his teeth and did not open his mouth. At this time, airway obstruction occurred for several minutes, making ventilation difficult. Oxygen saturation decreased by 40% on pulse oximetry, even though 50% of oxygen was provided and the concentration of inhalation anesthetics were not reduced. An additional 200 mg sugammadex was administered owing to severe airway obstruction caused by the patient clenching his teeth. After a few minutes, the patient stopped clenching his teeth, but a large amount of bloody discharge from the endotracheal tube was observed.

**FINAL DIAGNOSIS**

The final diagnosis of the presented case is unilateral pulmonary edema and diffuse alveolar hemorrhage.

**TREATMENT**

Following the presentation of unilateral pulmonary edema with diffuse alveolar hemorrhage in the operating room, furosemide 40 mg was administered immediately based on suspected NPPE due to airway obstruction. The patient was ventilated with 100% oxygen, but the oxygen saturation was maintained at 90% or less. Chest imaging (Figure 2) performed after transfer to the intensive care unit revealed haziness of the entire right lung, and bronchoscopy showed bloody secretions with red bubbles in the right lung(Figure 3). However, no active bleeding or endobronchial lesions were found. Arterial blood gas analysis performed after transfer to the intensive care unit showed pH 7.30, pO2 72.9, pCO2 40.6, and HCO3 20.1, (FiO2 1.0) with hemoglobin of 11.4 g/dL. On the day of surgery, norepinephrine and vasopressin were used because the patient’s blood pressure decreased in the intensive care unit. Tranexamic acid and methylprednisolone were administered for pulmonary hemorrhage and pulmonary edema. Extubation was performed on the eighth postoperative day.

**OUTCOME AND FOLLOW-UP**

The patient was transferred to the general ward on the tenth postoperative day. On the subsequent chest radiography, most of the radiologic haziness of the right lung had disappeared (Figure 4). The patient was discharged without any pathologic changes and clinical complications after a week after moving to the general ward.

**DISCUSSION**

The patient in this case had been diagnosed with asthma many years previously. To remove the endotracheal tube gently without irritating the airway following the surgery, we changed the patient’s posture under deep anesthesia. During this process, the patient severely bit the tube, and we were unable to provide ventilation for several minutes. NPPE occurred owing to severe negative intrathoracic pressure due to airway obstruction caused by the severe biting of the endotracheal tube, and a large quantity of bloody secretion was released into the intubation tube. At first, NPPE was suspected, and treatment was actively performed, including fluid restriction, diuretic administration, exhalation positive breathing, and 100% oxygen administration. Chest imaging performed after transfer to the intensive care unit showed diffuse haziness in the right lung field and persistent hemoptysis. Bronchoscopy findings showed no abnormal airway damage with the exception of bloody secretion in the bronchus or airways.

According to Mondoni *et al*[5], early bronchoscopy helps detect bleeding sources. They suggested that if there is rebleeding, the bronchoscopy must be performed again to examine the damaged airway area. Hemoptysis caused by damage to the airways due to temporary negative pressure is reported to naturally cease over time. Bhavani-Shankar *et al*[6] reported airway bleeding and hemoptysis caused by temporary intrathoracic negative pressure resulting in airway damage from airway obstruction due to various causes. This may be misdiagnosed as pulmonary edema. The authors suggested that hemoptysis may be caused by aspiration pneumonitis, airway injury, polyps, arteriovenous malformation, and tumors.

de Prost *et al*[7] reported that NPPE was a rare (5%, 6/112) cause of DAH. Contou *et al*[1] surveyed 149 patients over 36 years (1980-2015) in whom DAH was suggested and found that 18 cases (12%) of DAH occurred in 15 patients through NPPE. According to the study, almost 17 cases (94%) showed acute respiratory failure with no other organ failure or circulatory failure. Hemoptysis was found in 17 out of 18 cases (94%) and was the most common symptom, but in only one case was more than 200 mL per day observed. Anemia was observed in ten cases (56%), and the average hemoglobin reduction was 1.5 g/dL. It was reported that nine cases (50%) satisfied the DAH triad (hemoptysis, anemia, and pulmonary infiltrates). In this case, hemoptysis was the main symptom and anemia was not severe. The authors reported that 16 cases (89%) showed alveolar or interstitial opacity of both lungs. Only two cases (11%) demonstrated alveolar or interstitial opacity in the right lung. In the present case, diffuse opacity was observed in the right lung.

Schwartz *et al*[8] reported pulmonary hemorrhage after NPPE in surgical patients for the first time, and Goldenberg *et al*[9] reported two patients with tricuspid and pulmonary insufficiency among six post-laryngospasm NPPE patients. Dolinski *et al*[3] reported pulmonary hemorrhage in patients with mild tricuspid regurgitation due to increased pulmonary artery pressure after NPPE. In this case, the patient had evidence of trivial mitral regurgitation in the transthoracic echocardiography performed before surgery, but the clinical symptom was not specific. It is thought that a more detailed examination is needed to determine whether this mild mitral regurgitation was the cause of unilateral DAH in this case, but such an examination was not performed in this patient. In 2016, Han *et al*[10] reported negative pressure pulmonary hemorrhage that occurred after laryngospasm in the postoperative period. In 2016, Choe *et al*[11] reported DAH after transfemoral cerebral angiography in patients taking clopidogrel, and there is a potential for DAH associated with neuro-intervention in patients with hemoptysis and respiratory failure. It was argued that patients who are taking anticoagulants should carefully taper off the medicine prior to surgery and then begin it again during or after surgery. In addition, the author stated that it is better to protect less damaged lungs through active lung management such as oxygen therapy, bronchodilators, endotracheal intubation, and mechanical ventilation management in the treatment of patients with DAH.

NPPE by acute upper respiratory tract obstruction has been reported in about 11% of cases[12]. Risk factors for postoperative obstruction include obesity, short neck, obstructive sleep apnea, upper respiratory surgery, hypertrophy, and difficulty in tracheal intubation. While any factor that can cause gastrointestinal obstruction can cause NPPE, the most common cause of NPPE during anesthesia is postoperative laryngospasm. It is reported that NPPE occurs within a few minutes of airway obstruction and improves quickly after a day or two with active treatment. Steroids, diuretics, and bronchodilators are known to help, but their actions in these cases are not clear[12].

Papaioannou *et al*[13] reported that hemoptysis caused by NPPE, which occurs during obstruction of the upper airways, such as cases of laryngospasm, may be caused by negative-pressure tracheobronchial injury or alveolar-capillary disruption. It is also said that tension applied to the pulmonary-capillary wall may be caused by alveolar-capillary membrane rupture. The authors stated said that administration of succinylcholine 80 mg and administration of an unconsciousness inducing agent can prevent airway obstruction caused by laryngospasm[13]. Both laryngospasm and biting were possible causes of the patient’s airway obstruction in the present case. The authors chose sugammadex for muscle relaxation reversal instead of succinylcholine in this case, focusing more on the visible biting rather than laryngospasm as the cause of airway obstruction, and the patient quickly stopped biting and was able to undergo ventilation.

Young male athletes are known to be at risk. The syndrome occurs rarely in patients with endotracheal intubation but may also occur if the patient bites the endotracheal tube during waking[14]. Additional causes include excessive endotracheal or laryngeal aspiration as well as abnormal positioning of the mask. However, there are few reports of pulmonary hemorrhage accompanied by NPPE due to upper respiratory tract obstruction[1,8]. Schwartz *et al*[8] reported that at the end of surgery, negative intrathoracic pressure caused by upper airway obstruction after extubation causes hypoxic respiratory failure. The authors noted that postanesthetic laryngospasm, strangulation, hanging, foreign body airway obstruction, and obstructive sleep apnea may be factors associated with NPPE.

Contou *et al*[1] reported that the mechanisms of development of NPPE were negative pressure formation in the thoracic cavity, increased hydrostatic pressure in systemic and pulmonary capillaries, and mechanical stress in the alveolar capillaries. According to Contou *et al*[1], first -2 to -5 cmH2O intrathoracic negative pressure (-100 cmH2O in severe cases) increases venous return from peripheral blood vessels to the right atrium, simultaneously increasing pulmonary blood flow, increasing transmitral capillary pressure, and increasing pulmonary capillary and alveolar cavity pressure and stress. The hydrostatic pressure rises. These changes increase the transcapillary pressure gradient, and the fluid exudes into the interstitial space. The authors also reported that elevated central venous pressure interferes with lymphatic blood circulation.

Second, according to Contou *et al*[1], the systemic pressure increases owing to the secretion of norepinephrine due to hypoxemia, hyper carbon dioxide, and anxiety, which increases the amount of pulmonary blood flow. These effects reduce right ventricular dilatation, ventricular wall movement, and cardiac output. As the vascular resistance of the whole body increases, the resistance of the left ventricular wall increases, and the left ventricle prevents the blood flow. Lastly, the authors reported that effusion from capillaries due to physical alveolar-capillary membrane damage caused alveolar edema and bleeding.

Unilateral pulmonary edema is a very rare complication. This complication is mainly reported in cases of heart disease such as severe mitral insufficiency or congestive heart failure, and there are few reports in patients without heart disease. The syndrome can be caused by prolonged lying on the lateral side or by excessive administration of fluids in this position and can occur after treatment in cases such as pleural effusion or pneumothorax, or after re-expansion. MacLeod syndrome, unilateral pulmonary agenesis, talc pleurodesis, trauma, epilepsy, upper airway obstruction, pulmonary venous obstruction from mediastinal fibrosis, post-lobectomy, unilateral mainstem intubation, nitrogen mustard, heroin-related pulmonary edema, and pregnancy may all be causes of the syndrome[15]. The patient in the present case was not in the lateral position and had no specific trigger except for asthma from which he had been suffering for a long time.

As such, negative unilateral pulmonary edema and unilateral pulmonary hemorrhage may result from many causes. In patients with acute respiratory failure with hemoptysis and unilateral alveolar opacity, if clinical findings are not consistent with pneumonia, then it is important to determine the presence of heart disease. Rapid initiation of active treatment is important.

In summary, DAH is mainly caused by airway obstruction after surgery in young and healthy male athletes. The patient in the present case demonstrated acute respiratory failure accompanied by hemoptysis and alveolar opacity on the right side with no bronchial damage. We suggest that biting was the possible cause of the patient’s airway obstruction in the present case, and negative unilateral pulmonary edema with unilateral alveolar hemorrhage may result from airway obstruction. It is known that in older individuals, there is a high possibility that there will not be enough negative pressure to destroy the pulmonary capillaries, but the patient in this case was an elderly male patient who had been diagnosed with asthma 70 years earlier, considered to be at increased risk, and with damaged sensitivity to the endothelial tissue of the bronchioles.

**CONCLUSION**

In a 78-year-old man treated with asthma for a long time, change to the supine position at the end of general anesthesia resulted in severe hypoxic respiratory failure and increased airway pressure after acute upper respiratory obstruction due to severe endotracheal tube biting. The patient experienced a large amount of bloody secretions. Ultimately, the patient was discharged without complications after appropriate treatment for negative pressure pulmonary hemorrhage that occurred after endotracheal tube biting. Anesthesiologists should take care not to induce airway irritation during the emergence from anesthesia because DAH accompanied by NPPE was caused in this case by airway irritation in an older patient with asthma without known risk factors.

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**Footnotes**

**Informed consent statement:** Written informed consent was obtained from the guardian for publication of this report and any accompanying images.

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**CARE Checklist (2016) statement:** The authors have read the CARE Checklist (2016), and the manuscript was prepared and revised according to the CARE Checklist (2016).

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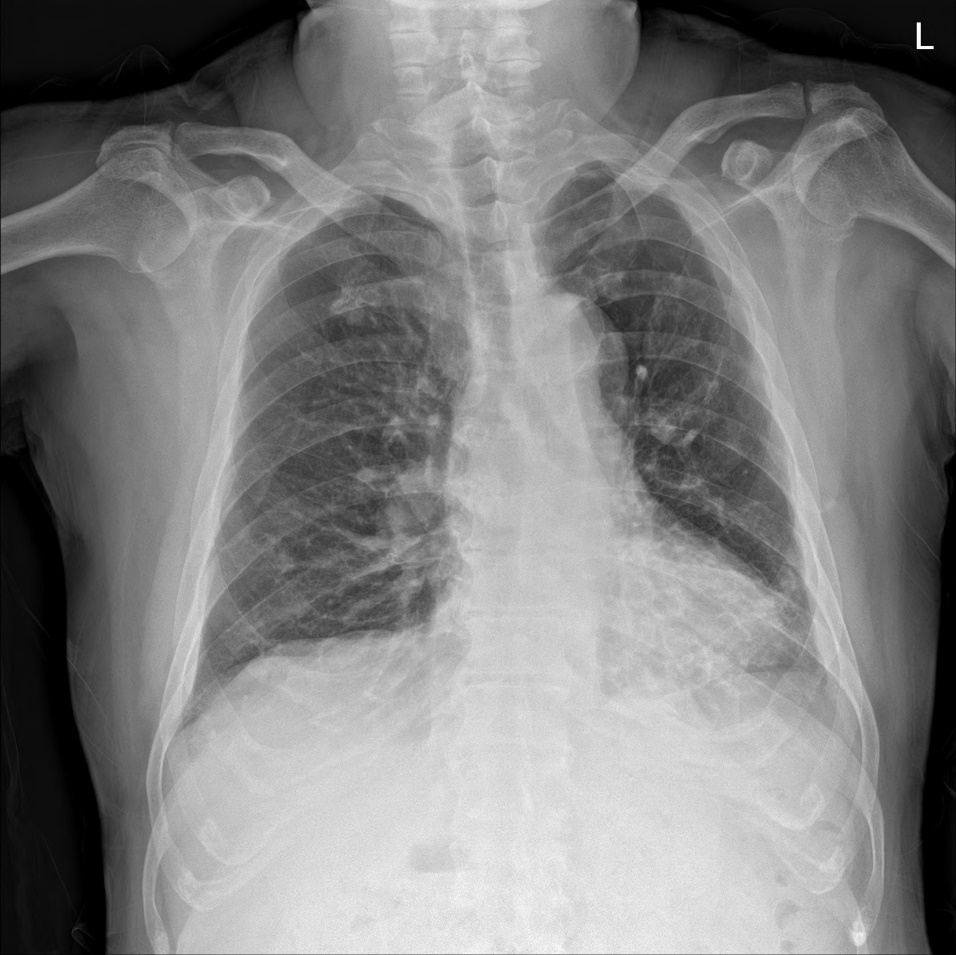
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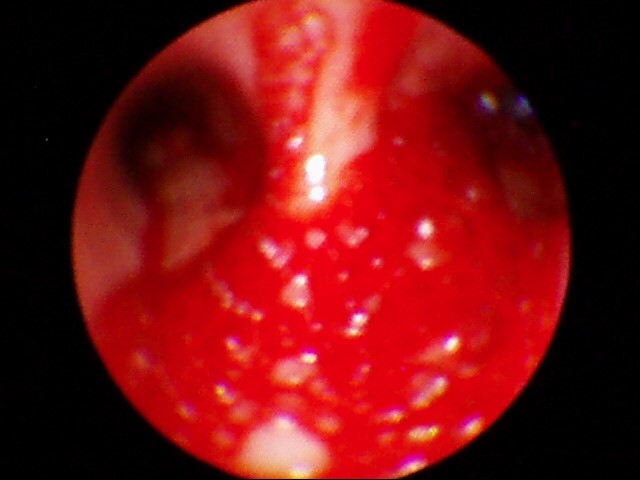
**Figure Legends**



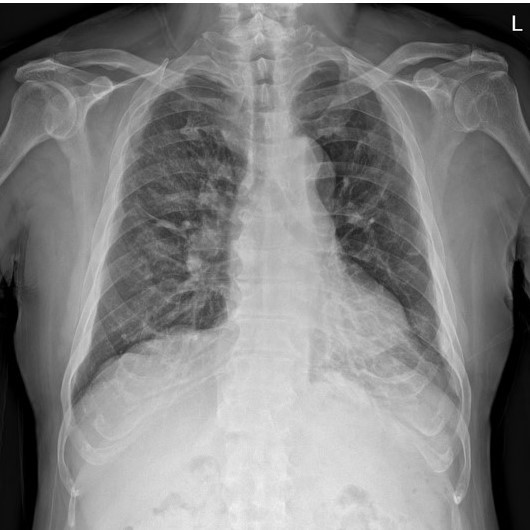
**Figure 1 Preoperative chest radiograph.** Chest radiographs showed subsegmental atelectasis in the left lower lobe and mild cardiomegaly.



**Figure 2 Postoperative chest radiograph at the intensive care unit.** Immediate postoperative chest X-ray revealed diffuse haziness of the entire right lung field.



**Figure 3 Postoperative bronchoscopic examination.** Postoperative bronchoscopy showed bloody secretions with red bubbles in the right lung.



**Figure 4 Postoperative chest radiographic examination on the eighthpostoperative day.** After recovery, most of the radiologic haziness of the right lung had disappeared in the chest radiograph.