An Alveolar–pleural fistula (APF) is communication between the alveoli and the pleural space that may result in intractable pneumothorax, severe infection, respiratory failure, physical weakness, and even death [1,2].

APF is usually not visible on bronchoscopy and diagnosed clinically [2,3]. In addition, APF rarely requires surgical intervention. However, when conservative treatment fails, interventions such as hydrogel application, blood patch, unidirectional valve and surgical repair are required [4–7].

We report a case of APF presenting with massive fluid regurgitated from an endotracheal tube during general anesthesia which was managed with bronchial suction, lung isolation and thoracostomy.

**CASE REPORT**

A 70-year-old male (170 cm, 59 kg) was admitted for a cystic mass (10.4 × 7.9 × 7.6 cm) of the liver and decided to undergo right hepatectomy. The patient had a medical history of hypertension and diabetes mellitus. Pre-operative transthoracic echocardiography revealed a left ventricular ejection fraction of 44% and severe hypokinesia of the left ventricle. Coronary angiography showed no significant stenosis. Pulmonary function tests demonstrated a restrictive ventilation defect. Positron emission tomography–computed tomography (PET–CT) revealed pleural effusion and consolidation in the right lower lobe (Fig. 1). Laboratory findings were as follows: hemoglobin, 12.9 g/dl, white blood cell, 7,100...
μl, platelet, 339,000 /μl, C-reactive protein, 11.1 mg/dl.

The patient’s vital signs were stable prior to general anesthesia. General anesthesia was induced with 50 mg of propofol and rocuronium 50 mg and maintained with 0.2 μg/kg/min of remifentanil and 5 vol% of desflurane. Tidal volume and respiration rate were set to 400 ml and 13 times/min, respectively. Positive end-expiratory pressure of 5 cmH₂O was applied, and peak inspiratory pressure (PIP) was 20 cmH₂O.

At four hours after the beginning of the operation, PIP abruptly increased to 32 cmH₂O and a large amount of serous fluid was regurgitated through the endotracheal tube (ETT) (Figs. 2 and 3). However, hypoxia did not occur at FiO₂ 0.5. PaO₂ and PaCO₂ were within normal range and oxygen saturation was 100% during the entire intra-operative period. The endotracheal tube was exchanged for a 35 Fr left double-lumen endotracheal tube (DLT) and tracheal suction was performed continuously. Approximately 1,000 ml of yellowish fluid was regurgitated through the right side lumen of the DLT within 30 min. Right-sided thoracostomy was performed following hepatectomy, and 300 ml of pleural effusion was drained through the chest tube. Air leakage was observed through the chest tube during positive pressure ventilation. We could not detect and localize abnormal lesion with bronchoscopy. Any gross defect of diaphragm was not observed at operation field. The oxygen saturation of the patient had been consistently maintained at 100%.

Total operation time was 7 h and estimated blood loss was about 9,000 ml. Nine liters of crystalloid fluid, 550 ml of colloid fluid, 5 units of packed red blood cells and 10 units of fresh frozen plasma were infused. Urine output during the operation was 790 ml. Post-operative hemoglobin level decreased to 9.1 g/dl after the operation.

The patient was transferred to the intensive care unit (ICU) with intubated state. Laboratory findings of the drained fluid from chest tube and the DLT revealed that it was parapneumonic exudate (pH 7.12, glucose 37 mg/dl, protein 3.31 g/dl, albumin 1.63 g/dl, lactate dehydrogenase 346 IU/L, cholesterol 18.2 mg/dl). The patient was diagnosed with pneumonia and treated with pleural drainage and antibiotics. The
volume of fluid drained through the ETT and the chest tube gradually decreased. The patient was extubated on postoperative day 5 and discharged from the ICU (Fig. 4).

DISCUSSION

An APF is a communication between the pulmonary parenchyma, distal to a segmental bronchus, and the pleural space. Common causes of APFs are invasive thoracic procedure such as lung biopsy, thoracentesis, and thoracic surgery. Other causes include necrotizing pneumonia and abscess, cancer, tuberculosis, infarction, bronchiectasis, trauma or barotrauma [2].

Without appropriate management, APF can lead to persistent air leaks, which are associated with prolonged hospital stays and high morbidity. Conservative managements of APF include ventilator strategies to reduce fistula flow, one lung ventilation, chest tube placement, or endobronchial valves. If these strategies are not successful, further interventions including chemical pleurodesis, autologous blood patch pleurodesis, or unidirectional valve device are needed [8,9].

There were several case reports of APF. The first reported case of APF detected during general anesthesia was described by Woo et al. [10] They suggested that APF developed before surgery due to thoracentesis, whereas our patient did not undergo any invasive procedure of lung. There are several case reports which is not associated with invasive thoracic procedures. El-Sameed et al. [7] reported cases of APF associated with tuberculosis. APF developed during positive pressure ventilation at ICU in a patient with pneumonia was reported by Wiaterek et al. [6]. Our case is different with these three cases because APF developed and detected with pleural effusion regurgitation during the surgery in a patient who had not undergone invasive thoracic procedure.

Any fistula between the pleural space and lung parenchyma, such as BPF or APF, pulmonary edema, and re-expansion pulmonary edema can be considered as a cause of regurgitated fluid through the endotracheal tube. Pulmonary edema was excluded from the differential diagnosis in this case because the regurgitated fluid was different from pink frothy sputum and oxygenation was maintained. In the present case, pleural effusion and consolidation in the right lung were observed on preoperative PET-CT. The patient had no respiratory symptoms preoperatively, and our patient did not undergo lung surgery or any thoracic procedures. Therefore, we considered that an APF occurred during the surgery and pleural effusion was regurgitated through a fistula in the right lung.

In addition, Manipulation of the liver during hepatectomy can facilitate the development of pleural effusion as a result of injury to the diaphragm, obstruction of the thoracic venous or lymphatic systems, or surgical manipulation of the hepatic coronary ligament [11–13]. The massive effusion observed during surgery in our patient could be attributed to hepatectomy.

We assumed that barotrauma due to mechanical ventila-
tion caused a fistula, with subsequent regurgitation of the pleural effusion. We immediately performed lung isolation with a DLT to protect the opposite side of the lung and reduce air leakage.

In conclusion, we presented a case of APF developed during general anesthesia presenting with massive exudate fluid regurgitated from an endotracheal tube that was managed with bronchial suction, lung isolation, and thoracostomy, and improved without surgical repair of the fistula.

CONFLICTS OF INTEREST

No potential conflict of interest relevant to this article was reported.

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