Pneumothorax in a Patient Posted for Cervical Spine Surgery

Naina P Dalvi, Nilam D Virkar

ABSTRACT

A 54-year-old female posted for cervical laminectomy was started on antihypertensive drugs on admission. Magnetic resonance imaging showed cervical degeneration with posterior disk herniation at C3–C4 and disk bulge at L5–S1. After attaching the monitors, patient was premedicated and anesthetized. During mask ventilation, abdomen gradually distended. After intubation under vision, reduced air entry on right side and increased resistance was felt during manual ventilation. Salbutamol puff was given through endotracheal tube. Still air entry remained decreased on right side. X-ray and C-arm showed right-sided pneumothorax. Inter-costal drainage (ICD) was inserted in right 5th intercostal space in midaxillary line. Post-ICD X-ray showed significant expansion of right lung. Patient was ventilated and extubated after 4 hours. Highresolution computerized tomography confirmed the diagnosis. Surgery was rescheduled. On the 8th day, patient developed purulent drainage through ICD in the ward. She was diagnosed to have pulmonary Koch’s and was treated successfully.

Keywords: Laminectomy, Tension pneumothorax, Tuberculosis.

INTRODUCTION

Pneumothorax occurs when intrapleural pressures exceed the atmospheric pressure. It causes collapse of the lung and increase in the airway pressure. As the intrathoracic pressures increase more than cardiac filling pressures, it limits the stroke volume and cardiac output leading to life-threatening hypotension and hemodynamic instability. Intraoperative pneumothorax is a life-threatening emergency, which needs rapid intervention.

CASE REPORT

A 54-year-old female was admitted with pain in the neck radiating to both arms, tingling numbness in the fingers, and pain in the lower back radiating to lower limbs with difficulty in walking since 2 years.

On preoperative evaluation, blood investigations were normal; chest X-ray showed increased bronchovascular markings. She was diagnosed to have hypertension and was started on T. Amlodipine 5 mg twice daily and T. Losartan 12.5 mg once daily. Her two-dimensional echo showed mild pulmonary hypertension 40 mm Hg with ejection fraction 60%. Magnetic resonance imaging showed cervical degenerative spondylarthropathy with posterior cervical disk herniation at C3–C4 and canal stenosis of L2–L5 with disk bulge at L5–S1.

Patient was first operated for L5–S1 laminectomy. Postoperatively, she was electively ventilated due to prolonged duration of surgery and inadequate respiratory efforts and was extubated 12 hours later.

Patient was posted for cervical laminectomy after 30 days of the lumbar surgery. Preoperative pulse was 130/min, blood pressure (BP) 130/90 mm Hg, and arterial oxygen saturation 99%. Patient was taken to the operation table. Cardioscope, pulsoxymeter, and noninvasive BP monitor were attached. Patient was premedicated with Inj. Glycopyrrolate 0.2 mg, Inj. Midazolam 1 mg, and Inj. Pentazocine 18 mg. After giving Inj. Thiopental sodium 5 mg/kg, ventilation was checked and rocuronium 1 mg/kg was given. During mask ventilation with oxygen, nitrous oxide, and sevoflurane, resistance was felt and chest expansion was found to be inadequate. Head was repositioned and triple airway maneuver was performed. The resistance still persisted. Abdomen gradually distended, hence nitrous oxide was switched off. Under laryngoscopic vision, 7 no. cuffed polyvinyl chloride endotracheal tube was inserted. Air entry was found to be decreased on the right side. Increased resistance was felt during manual ventilation and airway pressure went up to 35 cm H2O. Endotracheal tube and circuit were checked thoroughly for any kink and displacement. Nasogastric tube was inserted to decompress the stomach. Abdominal distension though reduced persisted with resistance to ventilation. Salbutamol metered dose puff was given through endotracheal tube to rule out bronchospasm. Air entry remained persistently decreased on the right side.

1Additional Professor, 2Assistant Professor

1Department of Anaesthesia, HBTMC and Dr RN Cooper Hospital, Mumbai, Maharashtra, India

2Department of Anaesthesia, LTMMC and LTMG Hospital Mumbai, Maharashtra, India

Corresponding Author: Naina P Dalvi, Additional Professor, Department of Anaesthesia, HBTMC and Dr RN Cooper Hospital, Vile Parle, Mumbai, Maharashtra, India

Phone: +9199820711656, e-mail: dmaina@rediffmail.com
Arterial oxygen saturation was maintained throughout. Making use of C-arm and X-ray available in the operation theater, right pneumothorax was identified immediately (Fig. 1). Implantable cardioverter defibrillator was inserted in right 5th intercostal space in midaxillary line. Gush of air was heard. Implantable cardioverter defibrillator was kept in underwater seal. Post-ICD X-ray chest showed significant expansion of the right lung. Surgery was postponed and patient was extubated after 4 hours (Fig. 2).

High-resolution computerized tomography showed pneumothorax on right side with collapse. Few fibrobronchiectatic and fibrocalcific changes in left lung were seen. Subcutaneous emphysema was seen in right chest wall and right breast (Fig. 3).

On the 8th day, patient developed tachypnea and desaturation with purulent drainage through ICD and was started on higher antibiotics. The pleural effusion persisted even after that and hence was started on AKT. Patient responded well to AKT and was discharged after removal of ICD after 26 days.

DISCUSSION

Tension pneumothorax occurs when air enters the pleural space during inspiration, but owing to ball-valve action cannot escape during expiration. The most reliable signs of tension pneumothorax are tachycardia, decreased breath sounds, hyper-resonance, and hypotension associated with hypercarbia and hypoxia.

Pneumothorax occurring during general anesthesia is a rare event but potentially dangerous problem, accounting for less than 3% of anesthesia complications.1 The most common causes are regional blocks (40% of cases), airway instrumentation (19%), barotrauma (16%), and placement of central venous lines (7%).2,3 Patients with chronic obstructive pulmonary disease are at increased risk of spontaneous pneumothorax.4 It is seen commonly during laparoscopic procedures.5-7 Spontaneous rupture of preexisting bullae8 may lead to pneumothorax, especially in smokers and young males. Barotrauma due to mechanical ventilation associated with risk factors like high airway pressures, stiff lungs, volume cycling, unregulated manual inflation, obstruction of endotracheal tube, excessive positive end-expiratory pressure, and endobronchial intubation is a cause of pneumothorax in 0.5–38% of critically ill patients.9-11 Incidence of delayed pneumothorax has been reported to be 0.4% (asymptomatic in 22% and tension pneumothorax in 22%).12 Occasionally a tension pneumothorax may be bilateral.13,14

Under anesthesia, patient cannot complain of usual symptoms like respiratory distress, chest pain. Use of intermittent positive pressure ventilation (IPPV) with N₂O may increase the risk of tension pneumothorax.
The diagnosis of pneumothorax rests on high clinical suspicion if there is an unexpected cardiorespiratory decompensation. Pneumothorax needs to be differentiated from bronchospasm, pulmonary edema, pulmonary embolism, and pulmonary aspiration. Rapidly deteriorating general condition in spite of ventilation with 100% oxygen after general anesthesia in the absence of a predisposing cause should alert the anesthetist to the possibility of tension pneumothorax. The standard emergency treatment in these cases is a wide-bore needle placement in midclavicular line in 2nd intercostal space or if possible immediate ICD insertion.15,16

The diagnosis of pneumothorax in our patient was done immediately, before any hemodynamic instability could occur and was treated successfully. As our patient responded to AKT later, there could be a possibility of bulla secondary to tuberculosis that might have ruptured during IPPV leading to pneumothorax.

CONCLUSION

Intraoperative pneumothorax is a life-threatening emergency that requires rapid diagnosis and appropriate treatment.

REFERENCES