Unexpected tracheal narrowing during general anesthesia in the prone position of Duchenne muscular dystrophy patient
-A report of two cases-

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Unexpected tracheal narrowing was observed in a patient with Duchenne muscular dystrophy during a corrective operation for thoracolumbar scoliosis. As the operating time progressed, peak airway pressure and end-tidal CO$_2$ increased gradually in the prone position. We found a floppy portion of the trachea using fiberoptic bronchoscopy (FB) in the prone position. We advanced a wire-reinforced tube toward the carina beyond the lesion. This allowed correction of the ventilatory abnormalities. We encountered another patient scheduled for the same operation. We performed FB in advance before the position change and observed a narrowed portion of trachea. We advanced the tracheal tube under FB beyond the pathologic portion and then moved the patient into the prone position. The operation was done successfully without any problems. (Korean J Anesthesiol 2013; 64: 456-459)

Key Words: Duchenne muscular dystrophy, Prone position, Tracheal narrowing.

Duchenne muscular dystrophy (DMD) is associated with a number of anesthetic problems including malignant hyperthermia, difficult intubation, upper airway obstruction and decreased lung function due to thoracolumbar scoliosis [1]. Upper airway obstruction can occur during surgery in DMD patients who experience difficult respiration in their daily lives [2]. But we found unexpected floppy tracheal wall during operations in DMD patients who did not report symptoms of respiration insufficiency in their daily life.

Thus we describe two cases of an unexpected tracheal narrowing during general anesthesia for the correction of scoliosis in DMD patients lying in the prone position, and how we successfully managed these potentially problematic situations with a simple measure.

Case Reports

Written statements granting permission of publication for the following cases were obtained from the parents or caregivers of the patients in nursing homes.
Case 1

A 12-year-old boy (height 125 cm, weight 18.8 kg) with Duchenne muscular dystrophy (DMD) was scheduled for the correction of thoracolumbar scoliosis. Cobb’s angle was 100°. Preoperative evaluation revealed that he was acceptable for general anesthesia, but he had decreased lung function. The FEV1 0.66 liter, FVC 0.73 liter, FEV1/FVC 90%). He did not report any symptoms of insufficient respiration in his daily life. He was supported on the operating table with multiple padding due to scoliosis and posterolaterally bent neck. Anesthesia was induced intravenously with propofol and rocuronium, and he was then intubated successfully with a wire-reinforced tracheal tube (internal dimension 5.5 mm, SPIRAL-FLEX® reinforced tubes, Hudson RCI, Temecula, USA). The tracheal tube was fixed at 17 cm. Both lung sounds were clear with manual ventilation. Peak inspiratory pressure (PIP) was 16 cmH2O with volume-controlled mechanical ventilation, tidal volume (TV) 160 ml, respiration rate (RR) 16 per minute and FiO2 0.5. Anesthesia was maintained with intravenous propofol and remifentanil starting at the targeted effect concentration of 3 μg/ml and 3.5 ng/ml respectively via target controlled infusion (TCI). After internal jugular vein and radial artery catheterization, the patient was moved into the prone position. In the prone position, PIP reached up to 30 cmH2O, and a low-pitched wheezing sound was detected in both lungs, although no secretion was observed from suctioning via tracheal tube. We switched the ventilator mode from the volume-controlled type to the pressure-controlled type. The expired TV ranged between 160-200 ml with set pressure at 25 cmH2O and RR 15 per minute. In this setting, arterial blood gas analysis (ABGA) was pH 7.43, PaO2 226 mmHg, and PaCO2 34 mmHg. A few minutes later, the expired TV gradually dropped to 120–140 ml at the same pressure-controlled ventilator setting. End tidal carbon dioxide concentration (EtCO2) increased gradually up to 50 mmHg. ABGA were at pH 7.25, PO2 224 mmHg and PCO2 60 mmHg. At this point, we decided to perform a fiberoptic bronchoscopy (LF-P, Intubation fiberscope, Olympus, Tokyo, Japan; FB). We found a narrowed, non pulsatile and nearly kissing tracheal wall between the carina and the tip of the tracheal tube using FB. Under the fiberscope guide, a wire-reinforced tracheal tube was advanced over the narrowed portion of the trachea, and we fixed the tube at 20.5 cm. The tip of the tracheal tube was placed just above the carina. Afterwards, with TV 160 ml, RR 16 per minute in the volume-controlled ventilation mode, PIP maintained below 20 cmH2O and EtCO2 was about 25 mmHg. The lung sounds of both lung fields were clear, so the remaining surgical procedure was done. At the end of the surgery, we changed the patient to the supine position. We set back the wire-reinforced tube to the previous position (17 cm) and observed the narrowed portion of trachea using FB. The narrowest part was acceptable for ventilation during positive ventilation and spontaneous respiration after the position change into supine. The emergence from general anesthesia was smooth without any complications.

Case 2

A 14 year-old male patient (height 158 cm, weight 50 kg) was scheduled to receive a corrective operation for thoracolumbar scoliosis. He also was diagnosed with DMD in his early childhood. When he was 9 years old, he started a wheelchair life due to muscle weakness. Preoperative evaluation revealed thoracolumbar scoliosis (Cobb’s angle 70°) and decreased lung function (FEV1 0.86 liter, FVC 1.37 liter, FEV1/FVC 63%). Except for these findings, all the other tests were in the normal range. He did not report any respiratory problems in his daily life. Anesthesia induction was performed with propofol, remifentanil, and a small dose of a rocuronium. The tracheal intubation was successful with a wire-reinforced tube. Airway

![Fig. 1. Fiberoptic bronchoscopic findings of the second case patient in the supine position. (A) A tracheal obstruction was observed in the mid-tracheal area. (B) Under bronchoscopic guidance, a tracheal tube was advanced to just above the carina. Thus tracheal patency was maintained in both the prone and supine position.](image-url)
pressures during controlled ventilation fell into normal range. But we immediately performed FB and observed a narrowed portion beyond the tip of the tracheal tube until the closing at the carina (Fig. 1). We advanced the tracheal tube along the fiberscope until its tip reached the carina. The sounds of both lung fields were clear. In the volume controlled ventilation mode (expired TV 400 ml, RR 13 per minute, FiO₂ 0.5), PIP was 18 cmH₂O, and EtCO₂ was 28 mmHg. After the patient was put into the prone position, we did FB again. It revealed the patency of the tracheal lumen in the prone position. During the operation, PIP maintained at a range of 22–26 cmH₂O without CO₂ retention. The surgery ended without any problems and at the end of anesthesia, FB confirmed the airway as narrow but acceptable for ventilation with retrieved endotracheal tube. During emergence and in the recovery room, the patient did not suffer from airway problems.

Discussion

DMD is the most common muscular dystrophy in children [1,3]. The disease process includes progressive degeneration of the skeletal, cardiac and smooth muscles. The anesthetic problems of DMD we may be confronted with are respiratory and cardiac complications, rhabdomyolysis, and malignant hyperthermia [1,2]. DMD patients usually present with concurrent pulmonary diseases preoperatively, and they are more susceptible to respiratory complications [4,5]. These complications range from minimal atelectasis to pulmonary edema and respiratory failure [4].

Tracheal obstruction, an unusual respiratory complication, has been reported before, and it was related to corrective scoliosis surgery in musculoskeletal diseases including DMD, Marfan syndrome, and congenital tracheomalacia [2,6,7]. Thoracolumbar scoliosis, a kind of abnormal thoracic configuration, may be related to tracheal obstruction, which results in an obstructive airway pattern in a pulmonary function test [8]. In thoracolumbar scoliosis, the deformity of the spine decreases the distance from the sternum to the thoracic vertebrae [9]. Moreover, corrective surgeries for scoliosis are usually performed in the prone position, and tracheal obstruction worsens with compressive pressure of the screw fixations. Additional deterioration of the spine curvature by corrective operation changes the structure of the mediastinum, and tracheal obstruction can be worsened by angulation, stretching, and the compression of vertebrae [10].

In the two cases listed above, although the patients had decreased lung function due to muscle weakness, they had no respiratory difficulty in their daily life. In the first patient, ABGA was tolerable at first, but it gradually worsened over time. We performed FB and found a narrowed, pulseless, nearly kissing tracheal wall. Then we advanced the wire-reinforced tracheal tube over the lesion in the prone position, and we solved the ventilatory problems. In the second case, we performed FB immediately after anesthesia induction based on the first experience. We also found partial tracheal obstruction and corrected it.

The problem we can expect with this maneuver includes accidental one-lung ventilation and mucosal edema of narrowed trachea. During corrective scoliosis surgery, the changes of thoracolumbar spine curvature could produce morphologic changes of tracheobronchial trees. Because we advanced the endotracheal tube until the tip of tube reached near the carina, unintentional one-lung ventilation could occur but fortunately did not. The second, the cuff or endotracheal...
tube itself could be a cause of mechanical irritation over the narrowed portion; mucosal edema could develop resulting in respiratory problems after extubation. To prevent this possible complication, we performed FB at the end of anesthesia with retrieved endotracheal tube depth and confirmed adequate patency of the narrowed tracheal portion and normal mucosal appearance. Maintaining adequate cuff pressure of the endotracheal tube and medication which reduces mucosal edema including steroids is recommended. Moreover, FB before extubation is highly recommended to prevent this complication.

We reviewed computed tomography of these patients to find predictive signs of tracheal obstruction which developed during general anesthesia, but not at other times. Usually the narrowest portion of mediastinum is around the manubrium level; the bony structures protect mediastinal contents from external forces. But these patients suffered compressed tracheal lumen around third or fourth thoracic vertebrae. At these levels, the bony support of the mediastinal structure was not usual because of the anatomical deformities including thoracic scoliosis (Fig. 2). In this situation, tracheal obstruction could develop in the prone position even though the patient did not have any prior respiratory problems.

In conclusion, when patients who have muscle weakness like DMD are scheduled for a surgical procedure in the prone position, FB is strongly recommended even though patients have no clinical symptoms and signs of respiratory insufficiency experienced in their daily life. Pre-operative imaging study and intraoperative FB help the anesthesiologist to recognize unexpected tracheal narrowing and that we would place the endotracheal tube at a proper depth to overcome tracheal narrowing.

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