Scoliosis and bronchial obstruction

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Learning objectives:
- To understand the reasons for respiratory compromise in patients with severe scoliosis.
- To be familiar with the treatment of bronchial obstruction caused by severe scoliosis.

Pretest
- What is the most common cause of chronic dyspnea in patients with idiopathic scoliosis (IS)?
- What are the options for treating bronchial obstruction caused by scoliosis?

CASE PRESENTATION
A 52-year-old woman was referred to the authors' centre in 2003 for an unresolving pneumonia. She was known for severe IS treated surgically at 12 years of age. She was on albuterol/ipratropium combination (Flovent, GlaxoSmithKline, USA) inhalers for a clinical diagnosis of asthma, which, however, had not been substantiated by pulmonary function tests (PFTs). She also had a chronic, nonproductive cough but no chronic dyspnea. She had smoked one pack per day between 20 and 30 years of age. In the previous year, she had experienced at least five documented episodes of pneumonia; these had been diagnosed clinically and radiologically, although sputum cultures had been negative on several occasions.

Computed tomography of the chest revealed atelectasis and pneumonia of the right lower lobe and compression of the bronchus intermedius by a severe residual deformity of the thoracic spine (Figure 1A). Bronchoscopic revealed a significant stenosis of the bronchus intermedius (Figure 2A). There was no evidence of neoplasia. The pneumonia eventually resolved after a prolonged course of intravenous antibiotics, as documented clinically and radiologically. PFTs conducted electively were essentially normal despite the known obstruction in the bronchus intermedius, likely explained by the fact that the obstructed airway would not impact PFT results because it was not contributing whatsoever. Oxygen saturation on room air at rest was 98%. Because of technical details related to her previous surgery, the patient was not deemed to be a candidate for any further operative attempts at correcting the scoliosis. Therefore, the bronchial obstruction was managed with an indwelling metallic stent inserted in the bronchus intermedius. Although the stent was deployed successfully, the technique was difficult because of the proximity of the right upper lobe orifice to the stenotic segment (Figures 1B and 2B).

Stenting resulted in an almost complete re-expansion of the affected lung parenchyma, although there was no significant change in PFT results (Figure 3). Clinically, the chronic cough continued and the patient experienced occasional episodes of mild hemoptysis. There were no episodes of respiratory infection in the year following stent insertion and, thereafter, she has experienced one to two episodes of mild bronchitis per year, diagnosed on the basis of increased cough and a change in sputum, all of which responded readily to oral antibiotics and none of which required hospitalization. Two years after stent insertion, a small nodule of granulation tissue causing mild hemoptysis and partial obstruction of the middle and lower lobes had to be treated with cryotherapy without recurrence. The stent has always been in a good position in the bronchus intermedius and no further stent-related complications have occurred.

DISCUSSION
IS is a complex three-dimensional deformity involving both abnormal flexion and rotation of the thoracic vertebrae (1). Cobb’s angle is usually used to describe the geometric severity of IS. It approximately correlates with overall clinical severity (2) although, in reality, the exact relationship between scoliosis and lung function is highly complex.

Scoliosis significantly alters the geometry of the chest and, ultimately, decreases the three-dimensional range of motion of both the thoracic cage and spine during breathing (1). In addition, the deformity places respiratory muscles at a biomechanical disadvantage (2), ultimately leading to decreased compliance of the chest wall and a restrictive syndrome that may significantly impact respiratory function (1,2). The ultimate severity is a function of Cobb’s angle, the number...
of involved vertebrae, the location of the abnormal curvature and the loss of normal thoracic kyphosis (2). Long-standing pulmonary hypoinflation leads to retention of secretions and small airways disease, as well as atelectasis and, eventually, irreversible lung atrophy, further reducing volume (2). Although blood gas levels are often normal at rest, gas exchange may be markedly altered during exercise, with resulting hypoxemia and hypercapnia. Severe scoliosis (Cobb’s angle >70°), especially when secondary to neuromuscular disease, may eventually progress to chronic respiratory failure and pulmonary hypertension (2). However, even asymptomatic patients with mild scoliosis may experience significant ventilatory functional impairment during maximal exercise tolerance testing (3).

Although rare, proximal bronchial obstruction may also occur. This may either be due to direct compression by the deviated thoracic spine, or rotation and distortion of the airway because of severely altered thoracic anatomy (4). The patient will usually present with progressive dyspnea and recurrent episodes of localized pulmonary infection. Although evidence is scant, a review of published cases suggests that compression may occur more frequently on the right, especially at the level of the right mainstem bronchus. Cross-sectional imaging will document bronchial compression across the deformed spine, as well as reveal signs of atelectasis or infection in the affected bronchopulmonary segments (5). PFTs may show an obstructive pattern, although the overall picture is also highly dependent on the effects of altered chest wall configuration and mechanics on lung volumes. This likely accounts for the variable findings reported and, in our case, the lack of a clear impact of bronchial stenting on spirometry results despite restoration of bronchial patency (5).

The first step in the management of symptomatic bronchial obstruction is usually correction of the scoliosis (6). A conservative approach using bracing is worth considering in mild cases. Bracing has been used for decades, especially in adolescent scoliosis during the rapid growth phase (7). However, success in stabilizing or improving the scoliosis is modest, may not be permanent and requires the brace to be worn virtually continuously, often resulting in low compliance. Furthermore, the brace itself may worsen pulmonary restriction (2). Other nonsurgical interventions aimed at improving the deformity include exercise, biofeedback and electrostimulation; however, their effectiveness is limited at best (2).

Use of chest physiotherapy, postural drainage and airway clearance devices have been described in bronchial obstructive syndromes and patients with severe scoliosis, and may be used as ancillary measures (2).

If such a conservative approach fails, surgical correction of the scoliosis may be indicated to lift the bronchial obstruction. A comparable strategy has been used successfully in the treatment of postpneumonectomy syndrome, in which surgical repositioning of a widely displaced mediastinum relieves bronchial compression caused by an impingement of the bronchus between the descending aorta and spine (8). In scoliosis, the decision for surgical correction depends on several factors including the geometry of the scoliosis, age at presentation, severity of symptoms and general quality of life. Although an in-depth discussion of surgical techniques is beyond the scope of the present article, several points can be made. Most of these reliably restore chest wall architecture and correct spinal deformity (9-11). However, the results of surgery on overall pulmonary function are inconsistent and remain controversial (9-11). Most authors agree that spinal fusion through a posterior approach improves both lung volumes (vital capacity and total lung capacity) and function in both the short and long term (10,11). However, anterior approaches to spinal fusion that require thoracotomy as well as thoracoplasty techniques that result in disruption of the chest cavity both tend to cause worsening of lung function (10,11). It is also important to emphasize that there are no reliable predictors of the degree to which anatomical correction will restore bronchial patency (12). Thus, the net result of surgical treatment would depend both on the restoration of bronchial patency and changes in lung volumes and function.

In cases in which symptoms of bronchial obstruction persist, despite optimal correction of the scoliosis, endobronchial stenting has been used successfully (13,14). Stenting usually results in improvement of the obstructive pattern observed on PFTs, decreased evidence of volume loss on pulmonary imaging, improved gas exchange and improved dyspnea. However, self-expandable metal stents are not without risks and should be used cautiously. When treating benign disease, stents are usually reserved as a last resort in inoperable patients.

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Indwelling stents may lead to a variety of complications, which may require remediation by interventional bronchoscopy (15,16). Granulation tissue is the most frequent complication, occurring in approximately 40% to 50% of cases; it is usually managed with cautery, laser ablation, cryotherapy or balloon dilation (15,16). Intractable granulation may require stent removal in 12.9% of cases (15,16). Migration may occur in up to 5% of cases, and stent metal fatigue or fracture in a further 3% to 5% (15). Although the patient we describe is doing well, studies have shown that 25% of stents inserted for benign disease may eventually require removal for intractable complications (16). Although the stents are designed to be permanent, removal is still often possible, even after several years, but requires an experienced team and may be technically challenging (16).

Once the decision to use stenting has been made, these patients should be followed closely. Although we do not have a rigid post-stenting surveillance protocol, we perform follow-up bronchoscopies periodically and as needed, on a case-by-case basis.

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**REFERENCES**


**Post-test:**
- What is the most common cause of chronic dyspnea in patients with IS?
- Patients with symptomatic IS may have marked anatomical alteration causing a reduction in lung volumes and decreased efficiency of respiratory muscles. Therefore, a restrictive pattern is most commonly seen in these patients. Obstructive patterns are less frequent.
- What are the options for treating bronchial obstruction caused by scoliosis?

The first step in the management of symptomatic bronchial obstruction is usually correction of the scoliosis. A conservative approach with bracing and physiotherapy is reasonable in mild-to-moderate cases (Cobb's angle ≤70°) although the likelihood of success is modest. In cases of severe scoliosis (Cobb's angle >70°) or if a conservative approach fails, surgical correction of the scoliosis may be considered to lift the bronchial obstruction. Endobronchial stenting is a less invasive alternative that has shown efficacy in cases in which other options were not possible, although it should be considered with caution because of the potential for long-term complications.