Tracheobronchitis with stridor in a patient with ulcerative colitis

Zahava C. Farkas¹, Sevak Keshishyan², Raja Chandra Chakinala¹, Shalom Frager³, Faisal Saeed⁴, Yasmin Yusuf⁵, Chaitanya Shilagani⁶, Roxana Bodin¹, Kassem Harris², Wilbert S. Aronow²,⁴

¹Department of Medicine, ²Division of Pulmonary and Critical Care, ³Division of Gastroenterology and Hepatobiliary Diseases, ⁴Division of Cardiology, ⁵Department of Pathology, ⁶Department of Radiology, Westchester Medical Center and New York Medical College, Valhalla, NY, USA

Abstract: Bronchopulmonary involvement is a rare but well documented extraintestinal manifestation of inflammatory bowel disease (IBD). IBD-related pulmonary disease can range from subglottic stenosis to tracheobronchitis to interstitial lung disease and is often misdiagnosed on initial presentation. We present a case of tracheobronchitis with stridor in a 23-year-old-woman with well controlled ulcerative colitis (UC).

Keywords: Tracheobronchitis; ulcerative colitis (UC); inflammatory bowel disease (IBD)

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Introduction

Extraintestinal manifestations of inflammatory bowel disease (IBD) develop in 21% to 41% of patients with a higher incidence in those with Crohn’s disease (CD) as compared to ulcerative colitis (UC) (1,2). Bronchopulmonary involvement is rare, occurring in only 0.21% to 0.4% of patients with IBD. Any part of the respiratory system can be involved. Bronchiectasis is the most common reported finding, however, tracheobronchitis, pleural and parenchymal disease can also occur (3,4). The rare nature of this finding and lack of physician awareness is perhaps why it is often misdiagnosed as bronchial asthma on initial presentation (5). We present a case of UC-related tracheobronchitis in a patient who presented with stridor to emphasize the importance of maintaining a high clinical suspicion for pulmonary manifestations of IBD.

Case presentation

A 23-year-old woman presented to an outside hospital in acute respiratory distress with audible stridor. Three weeks prior, she had presented to that same hospital with shortness of breath, and was treated with levofloxacin for presumed community acquired pneumonia. She developed progressively worsening dyspnea, later associated with intermittent fever, night sweats, sore throat, voice hoarseness, productive cough and an unintentional nine-pound weight, prompting a return to the emergency department. This time she was noted to be tachypneic, in respiratory distress and with audible stridor.

Stridor in adults is commonly caused by an abscess, tumor, laryngomalacia, subglottic stenosis or upper airway swelling, such as anaphylaxis, tracheitis or respiratory papillomatosis. It can be a sign of a life-threatening emergency and treatment should not be delayed. In this case, the patient received intravenous dexamethasone, racemic epinephrine, helium-oxygen inhalation therapy and empiric broad-spectrum antibiotic coverage. Computed tomography of the neck and thorax showed diffuse circumferential nodular tracheal and central bronchial thickening (Figure 1), and nodules within the left upper lobe, thought to be secondary to infection versus inflammation in the setting of tracheobronchitis. Two days later she was transferred to the medical intensive care unit at our institution for ENT evaluation; by then, the previously noted stridor had resolved and she never required intubation.

Of note, this patient was a non-smoker with no history of pulmonary disease, sick contacts or known tuberculosis exposures. She emigrated from the Dominican Republic in 2009 but had no recent travel. Past medical history was
significant only for biopsy-proven UC, diagnosed 4 years prior, with one episode of inflammatory arthritis in her hands and knees 2 years after her initial UC diagnosis. She was previously taking mesalamine (4.8 g/day) but had discontinued it on her own 6 months prior to admission due to lack of symptoms. She did not recall ever receiving any treatments other than mesalamine.

Labs on arrival to our hospital were notable for a white blood cell count of 28,000 k/mm$^3$ (94% neutrophils, 5% lymphocytes); hemoglobin 7.5 g/dL; platelet count of 1,163 k/mm$^3$; procalcitonin 0.34 ng/mL; erythrocyte sedimentation rate 96 mm/h; C-reactive protein 17 mg/dL; antinuclear antibody titer 1:160; Anti-myeloperoxidase antibody was not reactive; Anti-proteinase-3 antibodies reactive. Her last UC flare was 3 months prior to this admission, however she still had intermittent diarrhea and was therefore tested for *Clostridium difficile* infection; she tested positive.

Flexible bronchoscopy was performed as part of the work-up for her initial stridor, and showed multiple papillomatous-like lesions throughout the trachea (Figure 2), lending to initial concern for tracheobronchial papillomatosis with parenchymal involvement. Tracheobronchial biopsies showed ulcerated mucosa with fibrinopurulent exudate and focal re-epithelialization without the detection of granuloma formation (Figure 3). Pathology was negative for papillomatosis, viral cytopathic changes, evidence of fungal organisms or malignancy. Workup for infectious etiologies including tuberculosis as well as other autoimmune pathology such as sarcoidosis or granulomatosis with polyangiitis, were negative.

Empiric treatment with high-dose corticosteroids led to marked improvement in her respiratory status. The dose of steroids was tapered and the patient remained symptom-free at discharge with plans to undergo repeat chest CT 4 weeks after initiation of steroids to monitor for resolution of lung nodules. Due to active *Clostridium difficile* infection, a colonoscopy was deferred. Biopsy and serologic findings, in the absence of an alternate etiology, support a diagnosis of UC-related lung disease. She was scheduled for

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**Figure 1** Contrast enhanced axial CT thorax in soft tissue (left) and lung (right) windows, demonstrates circumferential tracheal wall thickening above the level of the carina (A,B), circumferential tracheobronchial wall thickening at the level of the carina (C,D), as well as bronchial wall thickening below the level of the carina (E,F), as indicated by the red arrows.
outpatient pulmonary and GI appointments but failed to follow-up.

Discussion

IBD is well known to cause extraintestinal manifestations including uveitis, arthritis, erythema nodosum, pyoderma gangrenosum, sclerosing cholangitis, and hemolytic anemia (6,7). Aside from the occasional case report, airway-associated disease is rarely acknowledged as a complication of IBD. Lung involvement may be subclinical and incidentally found on PFTs or present as subglottic-glottic stenosis, tracheobronchitis, bronchiectasis, cryptogenic organizing pneumonia, or interstitial lung disease (3,8-10). A concrete link between lung disease and IBD is difficult to establish. Confounding factors include the fact that medications used for the treatment of IBD including sulfasalazine, mesalamine, methotrexate, and tumor necrosis factor-alpha inhibitors, are also known to cause airway disease. Thus, the association is typically made after other etiologies have been ruled out and based on histopathology findings from bronchoscopy. Also important to note is the fact that pulmonary manifestations may precede a diagnosis of IBD or occur during quiescent periods of the disease. Camus et al reported a retrospective review of 27 cases of UC-related pulmonary disease and found that 12% of those patients developed respiratory symptoms prior to the diagnosis of UC, and only 3% developed respiratory and gastro-intestinal symptoms concurrently (11).

Multiple studies have demonstrated that patients with IBD have abnormal pulmonary function tests (PFTs), as compared to healthy controls, even when they are clinically asymptomatic. Those with UC are more likely to have abnormal PFTs as compared to those with CD, and restrictive lung findings are more common than obstructive. The most common abnormality found was a decrease in diffusion capacity of the lung for carbon monoxide (DLCO). Changes in forced expiratory volume in one second (FEV1) and the inspiratory vital capacity (IVC) were also noted but varied depending on disease activity (12).

A wide variety of bronchoscopic findings have been reported including severe tracheal narrowing, ulcerated mucosa and an ulcerated cobblestone appearance (1). Typical histopathological findings include abundant inflammatory cells within the epithelium without evidence of granuloma formation (13).

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Systemic steroids are the mainstay of treatment for suspected IBD-related airway disease and led to rapid improvement in most cases with large airway involvement (3,11). Temporarily, a mixture of helium and oxygen (heliox) may be used in patients who present with airway obstruction symptoms due to tracheobronchitis (9). One case reported resolution of symptoms and improvement in tracheobronchial lesions on repeat bronchoscopy after treatment with inhaled corticosteroids alone (7). In more severe cases, the addition of cyclophosphamide to a steroid regimen was associated with improved outcomes (5). Treatment of the underlying IBD with mesalamine or other available disease controllers should be initiated simultaneously. In conclusion, tracheobronchitis is a rare extraintestinal manifestation of UC and may be life-threatening. Increased recognition amongst physicians is imperative as prompt diagnosis and aggressive management are the keys to a better outcome.

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Footnote

Conflicts of Interest: The authors have no conflicts of interest to declare.

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References
