Tracheobronchial Calcification with Sinus Destruction

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Abstract

A 34-year-old lady presented with history suggestive of bronchial asthma & allergic rhinosinusitis. Computerised tomogram (CT) of thorax showed nodular calcification of trachea with sparing of the posterior wall. CT paranasal sinus revealed sinusitis with bone destruction. Bronchoscopy and bronchial biopsy was consistent with the diagnosis of tracheobronchopathia osteochondroplastica (TPO). Diagnosis of atrophic rhino sinusitis was established based on the diagnostic criteria. Sinobronchial symptoms have been described in TPO, but destruction of sinus due to atrophic rhinosinusitis has not been reported.

Keywords: Tracheobronchopathia osteochondroplastica (TPO); Atrophic rhinosinusitis; Tracheobronchial calcification

Introduction

Many diseases can lead to tracheobronchial calcification. Clinico-radiological correlation can identify the etiology of tracheobronchial calcification in most cases.

Tracheobronchopathia Osteochondroplastica (TPO) is one of the rare causes of tracheobronchial calcification. Sinobronchial involvement due to TPO has been documented in past, however CT sinus findings in cases of TPO have not been described earlier.

Case Report

A 34 year old non-smoker, homemaker was referred for the management of bronchial asthma. She complained of moderate-severe rhinitis, sinusitis, progressively increasing dyspnea on exertion and cough for 7 years. She denied history of paroxysmal breathlessness, emergency hospital visits or use of inhaled bronchodilators.

There was no history of recurrent lung infections in childhood. She had a history of atopy in self. Her paternal uncle had a history of asthma. She also had recurrent purulent nasal discharge and anosmia for the last two years. She did not have any systemic complaints and there was no past history of sinus surgery.

At presentation her vitals were normal. General physical examination was unremarkable. Examine of para-nasal sinuses (bilateral maxillary and frontal) showed tenderness. There was nasal crusting and patulous nasal passage on nasal examination.

Examination of ear was normal. Respiratory system examination showed bilaterally equal breath sound with polyphonic rhonchi auscultated all over the chest. Other system examination was normal.

Routine microscopic examination of her urine sample was normal. She had a chest radiograph done 2 weeks ago from elsewhere, showed bilateral mid-zone non-homogeneous opacities (Figure 1a). These opacities had reduced at presentation and had subsided after 1 month (Figure 1b).

Her routine investigations showed hemoglobin 11.8 gm/dl, total leucocyte count of 10400/mm3 with differential counts of polymorphs 67%, lymphocytes 29% and eosinophil 4%. Her biochemical test results were normal. Total immunoglobulin (Ig) E level was 600 IU/ml.

Spirometry showed obstructive defect i.e. ratio of forced expiratory volume in the first second to forced vital capacity of 61% and forced expiratory volume in the first second of 1.68 l (59% predicted) with significant bronchodilator reversibility of 390 ml and 17%. Serology for aspergillus showed anti-aspergillus IgG level of 7.8 IU/ml (normal<8), anti-aspergillus IgE level of 0.22 IU/ml (normal<0.35) and anti-aspergillus IgM level of 4.20 IU/ml (normal<8.00).

Computerised tomogram (CT) of thorax revealed diffuse thickening of walls of trachea, main and segmental bronchi along with submucosal nodular calcifications causing irregular luminal narrowing. Posterior wall of trachea was spared (Figure 2).
Peripheral subsegmental areas of atelectasis and consolidation were also seen in bilateral upper lobes. Tomogram of para-nasal sinuses (PNS) showed pansinusitis with erosive destruction of walls of sinuses and turbinates, which appeared atrophic (Figure 3). Cytoplasmic and peripheral antineutrophil antibodies were 2.19 (normal<7 IU/ml) and 3.22 (normal<7 IU/ml) respectively.

She was further advised positron emission tomography (PET) scan which showed metabolically inactive irregular mural thickening of anterior and lateral wall of trachea, bilateral main and segmental bronchi with multiple calcified nodules predominantly in left upper, lower lobe and right middle bronchi (Figure 4). There was no activity in paranasal sinuses.

Bronchoscopy revealed multiple white to yellowish hard nodules with cobblestone appearance projecting into the lumen of tracheobronchial tree. The nodules were observed all along the tracheal lumen, main bronchi, segmental bronchi and subsegmental bronchi with sparing of subglottic region and posterior wall of trachea (Figure 5). The trachea and bronchi were narrowed. The subsegmental bronchi could not be negotiated due to narrowing.

Histopathological examination of biopsy showed spicules of lamellar bone, fat cells and bits of metaplastic squamous epithelium (Figure 6). She was treated with long acting beta agonists, inhaled steroids and steroidal nasal spray because of atopic symptoms and good bronchodilator reversibility. Atrophic rhinosinusitis was managed conservatively with nasal irrigation with normal saline. The patient is taking prescribed medication regularly and has reported significant improvement in symptoms. She is on regular follow up for possible requirement of intervention for tracheobronchial nodule and atrophic rhinosinusitis.
younger individuals and females. Asymptomatic until the tracheal diameter falls below 1 cm. Others have frequently in males over 50 years, although cases have been reported in females with sparing of the subglottic trachea and the posterior wall. An abnormal chest radiograph led to further investigation. Pulmonary function test may be normal or show an obstructive pattern as seen in our case [2].

She was diagnosed to have atrophic rhinosinusitis as per diagnostic criteria suggested by Ly TH et al. [3]. These criteria are:

- Patient-reported recurrent epistaxis or episodic anosmia;
- Physician-documented nasal purulence, nasal crusting, chronic inflammatory disease of the upper airway
- Two or more sinus surgeries

If a patient with chronic rhinosinusitis demonstrates two or more clinical features for 6 months and longer it indicates atrophic rhinosinusitis [3]. She satisfied two of the criteria i.e. episodic anosmia and nasal crusting. Large roomy nasal cavity on nasal examination corroborated with the diagnosis of atrophic rhinosinusitis.

Chest radiograph in TPO is usually normal, but there can be areas of atelectasis and ill-defined opacities [4] as in our case. The CT thorax in our patient showed nodular calcification of tracheobronchial tree with sparing of the subglottic trachea and the posterior wall. Differential diagnosis of tracheobronchial calcification are tracheomalacia, saber sheath trachea, Mounier-Kuhn syndrome, Wegener's granulomatosis, endobronchial sarcoidosis, inflammatory bowel disease, papillomatosis, amyloidosis, relapsing polychondritis, and TPO [5]. Tracheomalacia has lunate-shaped trachea, saber sheath trachea has narrowed coronal diameter whereas, Mounier-Kuhn has increased tracheobronchial diameter. The tracheal anatomy apart from narrowing was normal in our case. Wegener's granulomatosis almost always affect upper airways and subglottic trachea [6]. Kidneys are involved in 80% cases. Elevation of cytoplasmic antineutrophil antibodies is common. Moreover, there are metabolically active areas of inflammation on PET scan in Wegener's granulomatosis. In our case the lesions were metabolically inactive. Sarcoidosis and inflammatory bowel disease have systemic symptoms, which were absent in our case. Papillomatosis and amyloidosis tends to involve the airway concentrically, as opposed to TPO, which spares the posterior wall. The only other disease apart from TPO, which spare the posterior wall, is relapsing polychondritis. In relapsing polychondritis however besides tracheal calcification there is recurrent chondritis of ear, nose and respiratory tract along with inflammatory non-erosive polyarthritis and ocular inflammation. Also, in relapsing polychondritis there is high attenuation thickening of tracheal and bronchial walls along with destruction of tracheal cartilaginous rings, whereas presence of focal, coarse calcification/ossification favors TPO [7]. The clinical-radiological findings, thus suggested the diagnosis of TPO.

Though, CT chest and clinical correlation was highly suggestive of TPO, sinus destruction on CT PNS could not be explained by TPO. The sinus destruction corroborated with the clinical diagnosis of atrophic rhinosinusitis [8]. A few case reports of TPO with sinobronchial symptoms have been described [9]. Occurrence of atrophic rhinitis has been observed in a few case series as well [1,10]. It has also been suggested that the presence of cough, dyspnea, hemoptysis, or recurrent sinobronchial infections in a patient with atrophic rhinitis should raise the suspicion of TPO [11]. But TPO with atrophic rhinosinusitis has not been described so far. It is possible that if previous cases of atrophic rhinitis and sinobronchial symptoms had been studied in detail, diagnosis of atrophic rhinosinusitis could have been established. There is a possibility that atrophic rhinosinusitis is secondary to TPO, as chronic inflammatory conditions are known to be associated with atrophic rhinosinusitis. More cases of TPO are required to be studied in detail with CT PNS to establish that rhinosinusitis in TPO is secondary.

The diagnosis of TPO is made on bronchoscopy. The classical findings are sessile cartilaginous and/or bony nodules with normal overlying mucosa sparing the posterior wall, producing a beaded appearance. This appearance is also called rock garden or cobblestone appearance of tracheobronchial tree 8 as seen in our case. Biopsy of these nodules is essential to confirm the diagnosis. The biopsy confirmed the diagnosis of TPO in our case. Treatment of TPO is necessary for symptomatic cases only, as no definitive management is available and the prognosis is generally good. Conservative management with antibiotics is required for those with recurrent respiratory tract infections, while others who develop obstructive symptoms may require interventions like removal of nodules by forceps, laser photoevaporation or cryotherapy or external beam irradiation or endobronchial stent placement or linear tracheoplasty [1,12].

Discussion

Tracheobronchopathia Osteochondroplastica (TPO) is an idiopathic, benign disorder of large airways. Till now approximately 400 cases have been reported in literature [1]. It occurs more frequently in males over 50 years, although cases have been reported in younger individuals and females. The severity of symptoms in TPO depends on the extent of airway narrowing arising due to submucosal nodules projecting in the airway lumen. TPO is a slowly progressive disorder, remains stable for many years. Majority of TPO patients are asymptomatic until the tracheal diameter falls below 1 cm. Others have cough, breathlessness, wheezing, hemoptysis and recurrent respiratory infections. Often they are diagnosed as asthma or chronic bronchitis. Our patient too was being referred for the management of asthma. But, atypical presentation i.e. absence of paroxysmal dyspnea and presence of abnormal chest radiograph led to further investigation. Pulmonary function test may be normal or show an obstructive pattern as seen in our case [2].

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