Endotracheal preocclusive lipoma: A rare reason of tracheal occlusion
Endotrakeal preoklusif lipom: Nadir bir trakeal tıkanıklık nedeni

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Introduction
Primary tracheal tumors are very rare, with an incidence of 0.2/100,000 of the population [1]. Benign tracheal tumors are a minority of these (10-20%). Endobronchial lipomas represent 0.1-0.5% of all pulmonary tumors [2]. There are few reports of tracheal lipomas in the literature, with most of the endobronchial lipomas located in the lower tracheobronchial tree [1,3].

Lipomas are composed exclusively of mature fat and comprise 0.1% of all benign lung tumors. Most intrathoracic lipomas are endobronchial [4]. Tracheobronchial lipomas arise from the submucosal fat of the tracheobronchial tree [4]. Lipomas can produce symptoms of airway obstruction such as a productive cough, wheezing, recurrent pneumonia, and bronchiectasis. Identification of fat on computed tomography (CT) within a lesion is suggestive of either a lipoma or hamartoma [5]. CT is highly specific and sensitive in the detection of fat and can be helpful in diagnosing tracheobronchial lipomas.

Endobronchial lipomas are more common than endotracheal lipomas. Although there are publications stating that lipomas are rarely found in the endobronchial system, we could not find any published papers giving statistical data about the location and incidence of endotracheal lipomas in the literature especially for the proximal trachea. Airway lipomas have a striking male dominance of 90% and usually present in patients at late middle age.

We report the case of a middle aged man with proximal preocclusive tracheal lipoma who had previously been treated for asthma. The diagnosis was only clarified after the onset of progressive dyspnea less responsive to the bronchodilators when the patient applied to the emergency room.
Case Report
A 62-year-old male patient applied to the emergency room with progressive shortness of breath. He had been smoking 1.5 packets of cigarettes per day for more than 20 years. His history showed a diagnosis of asthma 5 years ago but he had not been using the treatment regularly. Lately, he was suffering of dyspnea on exercise. He had no history of any recurrent pneumonia. Nine months earlier, he had the same symptoms with a cough and, on the basis of electrocardiography findings he was diagnosed as having a myocardial infarction with high troponin levels - initial troponin level at 20.7 progressively declining with his symptoms in ten days. On his latest admission, physical examination showed that he had stridor, slight peripheral cyanosis, dyspnea at rest, but auscultation showed no lung abnormality. His blood pressure was normal and his blood sugar level was slightly above normal. The other blood tests were within normal limits. The patient had no fever or infection on his arrival. The patient was monitored: the SPO2 level was 84%, oxygen treatment at 5 lt/min was applied but his response was slow.

On a chest CT scan, the superior part of the trachea showed an endotracheal soft tissue tumor obliterating more than 80% of the area of the lumen (Figures 1,2). The tumor was well-circumscribed, homogenous and hypodense, originating from the anterior endoluminal wall; the density was between –83 –110 HU. There was no parenchymal pathology in either lung field. Finally, an endotracheal lipoma was diagnosed.

The patient’s symptoms resolved after oxygen treatment, bronchodilators and steroid treatment, on follow-up. The patient was referred to surgery, but he rejected the operation.

Discussion
Benign tumors of the tracheobronchial system originate from the surface epithelium or the mesenchyme. Papillomas are the most frequent benign tumors in the group of benign tumors arising from the surface epithelium. Hamartomas, leiomyomas, lipomas, fibromas and neurogenic tumors are other benign tumors, and have a mesenchymal origin. Hamartomas are the most frequently seen tumors in this group and together with lipomas they are most commonly found in the left main stem bronchus of middle aged male smokers with clinical findings reported here [6] including a persistent cough, chest pain, dyspnea, recurrent fever and pneumonia, and sometimes with wheezing [7,8]. Hemoptysis is uncommon, owing to the avascular nature of lipomas, but can occur as a result of postobstructive infection [7].

In our patient the main symptom was dyspnea and previously he had been admitted to the coronary unit with high troponin and ECG findings supporting acute myocardial infarction and a history of asthma nine months earlier. In a long term smoker, clinical findings which mimic chronic obstructive pulmonary disease (COPD) or asthma, worsening of dyspnea in the supine position, short term improvement after bronchodilators and steroids can delay the diagnosis. In such cases, CT is the preferred screening modality to rule out endotracheobronchial lesions causing the symptoms.

Computed tomography is extremely valuable in the accurate diagnosis of endobronchial lipomas. They manifest as a pedunculated or a broad based homogeneous lesion with attenuation around -100 HU. Multidetector CT enables
the display of milimetric endobronchial tumors [9]. Bronchoscopic biopsies are frequently of no diagnostic value as the thick fibrous capsule could be misleading if atypical cells are found secondary to chronic inflammation.

In our review of the literature, we found only a few papers from Japanese and Brazilian [4,10] publications as case reports supporting the view that tracheal lipomas are extremely rare benign tumors. They could be mistaken and followed as obstructive lung disease.

References