Introduction

Sarcoidosis is a multisystem granulomatous disorder with unknown etiology. The pathologic hallmark of sarcoidosis is the noncaseating granuloma in involved organs.\(^1\)

The prevalence of this disorder is not exactly known. In most cases it involves patients between the ages of 10 and 40. Almost always it involves the lungs.\(^2\) The common presenting symptoms are cough, dyspnea, and chest pain. In rare cases it may present with massive hemoptysis.\(^3\)

We report a 24 years old man with sarcoidosis who presented with massive hemoptysis in early course of the disease who did not have any lung parenchymal involvement.

Case Report

A 24 year old non-smoker male presented to our pulmonary clinic with a chief complaint of hemoptysis for 3 days. He did not have dyspnea. In physical examination the chest was clear bilaterally, without crackle or wheeze. Chest-X ray was normal. Chest CAT scan showed para tracheal and sub carina lymphadenopathy (Figure 1) without mass and cavitations. CAT scan did not show any evidence for pulmonary embolism.

Laboratory investigation showed: ESR = 50 mm, white blood cell count = 6500 mic/lit, Hg = 13 g/dl, Htc = 45 % and, Angiotensin-converting enzyme (ACE = 89 IU/lit (normal value < 40 IU/lit)). PPD skin test and acid fast bacilli was negative and combs wright was 1/40. Blood Culture for gram stain and fungal infection were negative. Body Box test was normal (TLC = 108 %, FEV1/FVC = 89 %, FEV1 = 90 %, RV/TLC = 112 %). DLCO did not show any specific finding and was in the normal range (DLCO = 90 %). Skin test and sputum culture for aspergillus were negative.

One day after admission, he developed a massive hemoptysis with more than 600 ml bleeding. Emergency rigid bronchoscopy was performed. Hemoptysis source was in the right lower bronchus. After bleeding was stopped, a fiberoptic broncoscopy was done which did not show any endobronchial lesion. Bronchoalveolar lavage for acid fast bacilli was negative. CT-angiography was negative for A-V malformation.

The patient underwent mediastinoscopy and all parathoracic and sub carina lymph nodes were removed. Pathology examination showed many granulomas composed of epithelium cell, polynuclear giant cells and lymphocytes. Asteroid bodies were seen in cytoplasm of some of the giant cells (Figure 2).

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Discussion

Sarcoidosis occurs worldwide with higher incidence in the United States and Sweden. Its incidence appears to be lower in Asia.\(^4\) The exact etiology and pathogenesis of Sarcoidosis remains unknown.\(^5\) Most frequently it involves lungs and presents with symptoms such as cough, dyspnea, and chest pain. It can also cause fatigue, malaise, fever, and weigh loss. Systemic symptoms are more common in older patients.\(^6\) Massive hemoptysis may occur only in less than 0.5 % of the patients.\(^7\) Hemoptysis can occur due to a necrotic sarcoidosis lesion, which could be treated with embolization or surgery.\(^8\) It can also occur due to micro aneurysm in bronchial arterioles which shunt to the pulmonary vein.\(^9\) In one report major hemoptysis was found to be due to involvement of the upper respiratory tract and ulceration of nasopharyngeal granuloma.\(^10\) Development of clinical bronchiectasis in advanced stages can also cause hemoptysis.\(^11\) In one report massive hemoptysis...
was due to granulomatosis artritis in small and medium size muscular arteries.12

Here we reported a case of sarcoidosis with massive hemoptysis with a favorable response to corticosteroid. Absence of lung parenchymal involvement and cavitation, differentiates our patient from other cases. We acknowledge the limitation in this report regarding the absence of lung biopsy and angiography to investigate granulomatosis arthritis and microaneurism.

Hemoptysis, especially massive hemoptysis in sarcoidosis is rare, by some patients may experience massive hemoptysis due to aspergilluma, bronchectasis, microaneurism and rarely granulomatosis artritis.

References


Figure 1. Para tracheal and sub carina lymphadenopathy

Figure 2. Granulomas composed of epithelium cell and polynuclear giant cells