CASE REPORT

Mycobacterium szulgai pulmonary infection in a vitamin Ddeficient patient: A case report

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Abstract

Closer attention should be paid to vitamin D status in patients with mycobacterial diseases.

KEYWORDS

case report, Mycobacterium szulgai, pulmonary infection, vitamin D deficiency

BACKGROUND 1

Mycobacterium szulgai (M szulgai) is an opportunistic and rare pathogen. Several risk factors play roles in the development of disease caused by this bacterium including immunodeficiency syndromes, previous Mycobacterium tuberculosis infection, and smoking. We reported a case of M szulgai respiratory infection in a 32-year-old male patient without any history of smoking, immunodeficiency syndromes or receiving immunosuppressive medications but the patient was severely deficient in vitamin D. Finally, the following treatment regimen was administered to the patient: isoniazid, rifampin, ethambutol, clarithromycin antimicrobial agents supplemented with vitamin B6 and vitamin D. The patient's clinical condition was improved after 10 months of treatment.

Vitamin D deficiency should be considered as a risk factor for *M* szulgai pulmonary infection in immunocompetent people who are daily exposed to very little sunlight due to weather, occupational, or religious reasons.

Nontuberculous mycobacteria (NTM) are widespread worldwide. These bacteria can be transmitted to humans from environmental sources such as soil and water with no reported cases of person-to-person transmission.¹

Mycobacterium szulgai is a slow-growing NTM and a rare human pathogen, representing < 0.2% of all NTM infections.² Mycobacterium szulgai pulmonary infections are infrequently reported in those with a history of chronic obstructive pulmonary disease (COPD), tuberculosis, or immunosuppression.³ Clinically, it is indistinguishable from *Mycobacterium* tuberculosis infection and neoplasm; therefore, it is critical

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to be diagnosed accurately.⁴ Moreover, there is no consensus recommendation for treatment of *M szulgai* infections, probably due to its low prevalence.⁵ Herein, we present a case of *M szulgai* pulmonary infection in an immunocompetent patient without any history of tuberculosis or consumption of immunosuppressive drugs but severely deficient in vitamin D that was successfully treated in Mashhad, Iran.

2 | CASE PRESENTATION

A 32-year-old man was referred to Ghaem Hospital affiliated to Mashhad University of Medical Sciences in February 2019 with symptoms of night sweats, cough, and chest pain with previous pulmonary infection *mycobacterium tuberculosis* diagnosis based on positive acid-fast bacilli smear test. Tuberculosis was diagnosed in August 2016. The patient was received a standard 6-month regimen including rifampin, isoniazid, ethambutol, and pyrazinamide. After the treatment, the result sputum smear and culture for acid-fast were negative. He had no history of alcohol use and smoking. The patient had a weight loss of 9 kg over 1 year. The patient worked in a clothing industry that was not exposed to sunlight. An initial test performed on the patient showed severe vitamin D deficiency. A serology test for HIV was negative. The chest X-ray revealed an increase in pleural thickness was seen in the left lung and less strongly in the right lung. Reticular lesions were observed in the field of lungs, specifically, at the top of the lungs (Figure 1A). High-resolution computed tomography (HRCT) revealed bronchiectasis, distractive changes, and multiple cysts with collapse in the left upper lobe. Bronchiectasis and cyst formation with tree-in-bud pattern were observed in the right upper lobe and less intensively in the right middle lobe and upper left lower lobe segment and with less intensity in the upper right lobe (Figure 1C). Laboratory findings (Table 1) combined with the clinical signs of the patient raised the possibility of Mycobacterium tuberculosis infection (TB). Therefore, direct smear microscopy for acid-fast bacilli (AFB) and mycobacterial culture were performed on the patient's sputum sample. All three samples obtained from the patient were smear-positive for AFB using Ziehl-Neelsen method. Also, mycobacterial culture of the patient's samples on Lowenstein-Jensen medium became positive for AFB 22 days after inoculation. To identify the grown bacterium, two independent PCR assays were performed on the DNA extracted from the bacterial colonies. One PCR assay had been designed to detect IS6110, a specific gene target for *M* tuberculosis complex, in the specimen and the other assay detected rpoB, a mycobacterium genus-specific target. The organism was identified as (NTM) due to obtaining negative and

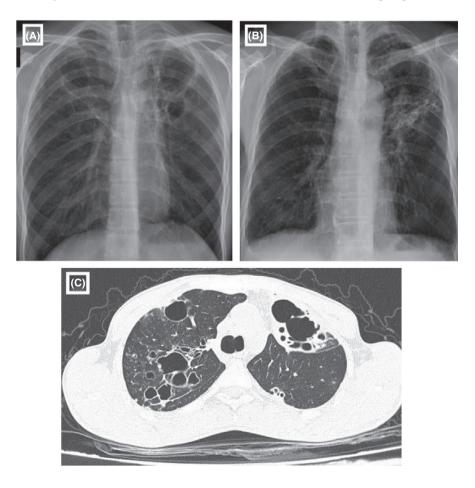


FIGURE 1 Thirty-two-year-old man with *M szulgai* respiratory infection. A, Fibrotic changes in the lungs with preference to the left lung. B, Follow-up chest radiograph 10 mo after treatment. C, Bronchiectasis and cyst formation with a tree-in-bud pattern in the right upper lobe and less strongly in the right middle lobe and upper left lower segments

TABLE 1 The results of the initial tests taken from the patient	t
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WBC (×10 ³ /UL)	12.4
RBC (×10 ³ /UL)	5.19
HB (g/dL)	11.8
BUN (mg/dL)	6
PLT (×10 ³ /UL)	523
Creatinine (mg/dL)	0.58
AST (units/L)	25
ALT (units/L)	55
Potassium (mEq/L)	4.6
Magnesium (mEq/L)	2.3
Sodium (mEq/L)	139
total bilirubin (mg/dL)	0.2
direct bilirubin (mg/dL)	0.1
vitamin D (ng/mL)	9
Neutrophils ($\times 10^3$ /UL)	10.16
Lymphocytes (×10 ³ /L)	2.11
Monocytes ($\times 10^3/L$)	0.13

Abbreviations: ALT, alanine aminotransferase; AST, aspartate aminotransferase; BUN, blood urea nitrogen; HB, hemoglobin; PLT, platelets; RBC, red blood cell count; WBC, white blood cell count.

positive results, respectively, for the former and the latter PCR assays. Results were also confirmed by sequencing of the *ITS* (16S-23S) (accession number MN124505), *rpoB* (accession number MN158122), and *hsp65* (accession number MN158123) genes of *M szulgai*. Based on the drug susceptibility test (DST) results, treatment with isonia-zid (225 mg), rifampicin (450 mg), ethambutol (825 mg), clarithromycin (1000 mg), vitamin B6 (10 mg), and vita-min D (1000 IU) oral daily was started. After 10 months of treatment, a sputum sample was taken from the patient, smear, and culture tests were negative. Chest pain and dyspnea were resolved and the patient gained 6 kg. Significant improvement was seen in the chest X-ray (Figure 1B). The patient's serum vitamin D also reached normal.

3 | **DISCUSSION**

According to the ATS (American thoracic society) and IDSA (infectious disease society of America) guidelines released in 2007, to identify NTM respiratory infection radiological, microbiological, and clinical findings are required. These include finding nodular or cavitary lesions on chest X-ray or bronchiectasis on HRCT and positive culture of at least two sputum specimens.⁶ Most cases of *M szulgai* respiratory infection have been previously reported to be fibrocavitary,⁵ whereas in our case was bronchiectasis with nodular lesions and a tree-in-bud view. In the bronchiectasis form, the disease is more scattered in

the middle lobe and lingual area, whereas in our case, most lesions were in the upper lobe. Lio et al⁷ showed that the bronchiectasis form of Mycobacterium avium pulmonary infection is more commonly seen in those with chronic obstructive pulmonary disease or gastrointestinal malignancies such as gastroesophageal reflux disease. Our patient had no history of any obstructive diseases or gastrointestinal malfunctions. In the majority of reported cases, M szulgai respiratory infections men over 50 years of age and have risk factors such as alcohol abuse, smoking, history of TB infection, low body mass index, or skeletal abnormalities of the chest.⁸ To our knowledge, the present patient is the youngest case of *M* szulgai pulmonary infection without any underlying disease with had severe vitamin D deficiency. Vitamins can affect different components of the innate immune system, and deficiencies lead to defects in the immune system and infections.⁹ The normal blood level of vitamin D is about 30 ng/mL, and vitamin D deficiency involves a blood level below 20 ng/mL. Douglas et al demonstrated that vitamin D plays an important role in immunological defense against mycobacterial infections. Vitamin D activates macrophages by blocking the intracellular growth of mycobacteria; on the other hand, vitamin D deficiency causes immune system defeat and growth Mycobacteria easily.¹¹ The main source of vitamin D for humans is sunlight. Because the patient spent a lot of time at the workplace and no history of eating vitamin D supplements and vitamin D rich diets, he had a severe vitamin D deficiency.

Unlike other NTM, *M szulgai* is sensitive to most anti-TB drugs and responds well to treatment regimens containing more than two anti-TB drugs.⁶ *Mycobacterium szulgai* is also sensitive to macrolides, aminoglycosides, and fluoroquinolones. In some reports, clarithromycin treatment with ethambutol and rifampicin or rifabutin has been found effective.⁷ Clarithromycin is one of the famous members of aminoglycosides which has been FDA approved in 1990. This therapeutic agent is recommending for various bacterial infection particularly mycobacterial diseases.¹⁰ According to review of the literatures, there are several evidence for successful treatment of clarithromycin in pulmonary infection caused by *M szulgai*.¹¹ On a 10-month follow-up, the patient had a negative smear and culture. Signs of improvement in the patient's chest X-ray visible (Figure 1B).

4 | CONCLUSION

Mycobacterium szulgai can infect immunocompetent individuals with risk factors such as vitamin D deficiency. Considering the positive effect of vitamin D on innate Immunity through the regulation of macrophage activity, supplementing regimens are highly recommended for the

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treatment of NTM infections. Closer attention should be paid to vitamin D status in patients with mycobacterial diseases.

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Consent statement: Published with written consent of the patient.

CONFLICT OF INTEREST

None declared.

AUTHOR CONTRIBUTION

HL: carry out the laboratory experiments and the acquisition of data. HL, MS, and EA: wrote the first draft of the manuscript. AKS: contributed to revise the article critically for important intellectual content. ZM, HL, and EA: collaboratively contributed to the concept of the original research resulted in this report. All authors reviewed and approved the final version of the manuscript.

ETHICAL APPROVAL

Ethical approval to report this case was obtained from Mashhad University of Medical Sciences Regional Ethics Committee, Mashhad University of Medical Sciences, Mashhad, Iran (Approval Number: IR.MUMS.fm.REC.1396.638).

DATA AVAILABILITY STATEMENT

The data that support the findings of this case are available from the corresponding author (Ehsan Aryan), upon reasonable request.

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