

# Mycobacterium Chimaera: A Rare Presentation

Jeffrey A. Miskoff <sup>1</sup>, Moiuz Chaudhri <sup>1</sup>

1. Medicine, Jersey Shore University Medical Center, Neptune City, USA

✉ **Corresponding author:** Jeffrey A. Miskoff, jamiskoff@yahoo.com

Disclosures can be found in Additional Information at the end of the article

---

---

## Abstract

Mycobacterium chimaera is an indolent nontuberculous mycobacterium which is abundant in soil, dust, and water. Although lacking recognition garnered by other mycobacteria (i.e., M. tuberculosis), it has been recognized as an emerging opportunistic threat to patients undergoing coronary bypass surgery and open heart procedures requiring extracorporeal devices. Here, we present a case of an individual initially seen in the inpatient setting without a history of such procedures or other risk factors commonly associated with mycobacterium-laden infections.

---

**Categories:** Internal Medicine, Infectious Disease, Pulmonology

**Keywords:** case report, coronary bypass surgery, mycobacterium, mycobacterium avium complex-mac, heater-cooler, open heart procedures, dna polymerase chain reaction, opportunistic infections, primary care

## Introduction

Mycobacterium chimaera is a slow-growing nontuberculous mycobacterium grouped with Mycobacterium intracellulare and Mycobacterium avium to make up the Mycobacterium avium complex. Evidence suggests that it is responsible for symptoms similar to those found in disseminated mycobacterial diseases, prosthetic valve endocarditis, ocular emboli, vertebral osteomyelitis, hepatitis, and renal dysfunction, along with other life-threatening conditions [1-3]. Additionally, it can cause pneumonia in patients with underlying respiratory complaints, such as cystic fibrosis and other immunodeficiencies [4].

## Case Presentation

A 67-year-old male presented to our care on December 26, 2017, for shortness of breath on exertion and persistent cough for three weeks. Prior to these symptoms, he was completely asymptomatic. Symptoms progressed gradually up to the extent that he was not able to leave his house. Furthermore, the patient experienced severe shortness of breath when he attempted to use the restroom. He described a productive cough with white to yellow sputum. Additionally, he felt feverish on three different occasions since the onset of symptoms. Although he had no chest pain, the patient experienced chest tightness and heaviness. The patient reported orthopnea and waking up in the middle of the night due to shortness of breath. The patient denied palpitations, chills, night sweats, dizziness, recent travel, contact with birds, exposure to tuberculosis, and contact with wild animals. On presentation, his blood pressure ranged from 95/57 to 125/63 mmHg, a heart rate of 98 beats per min, a respiration rate of 19 breaths per min, and temperature of 97.3 F. His oxygen saturation was 94% on room air. Pulmonary function testing revealed a Gold 3 category with a forced expiratory volume in 1 second (FEV-1) of 41%, forced vital capacity (FVC) of 69%, and an FEV-1/FVC ratio of 47%.

Received 05/03/2018

Review began 05/14/2018

Review ended 05/31/2018

Published 06/06/2018

© Copyright 2018

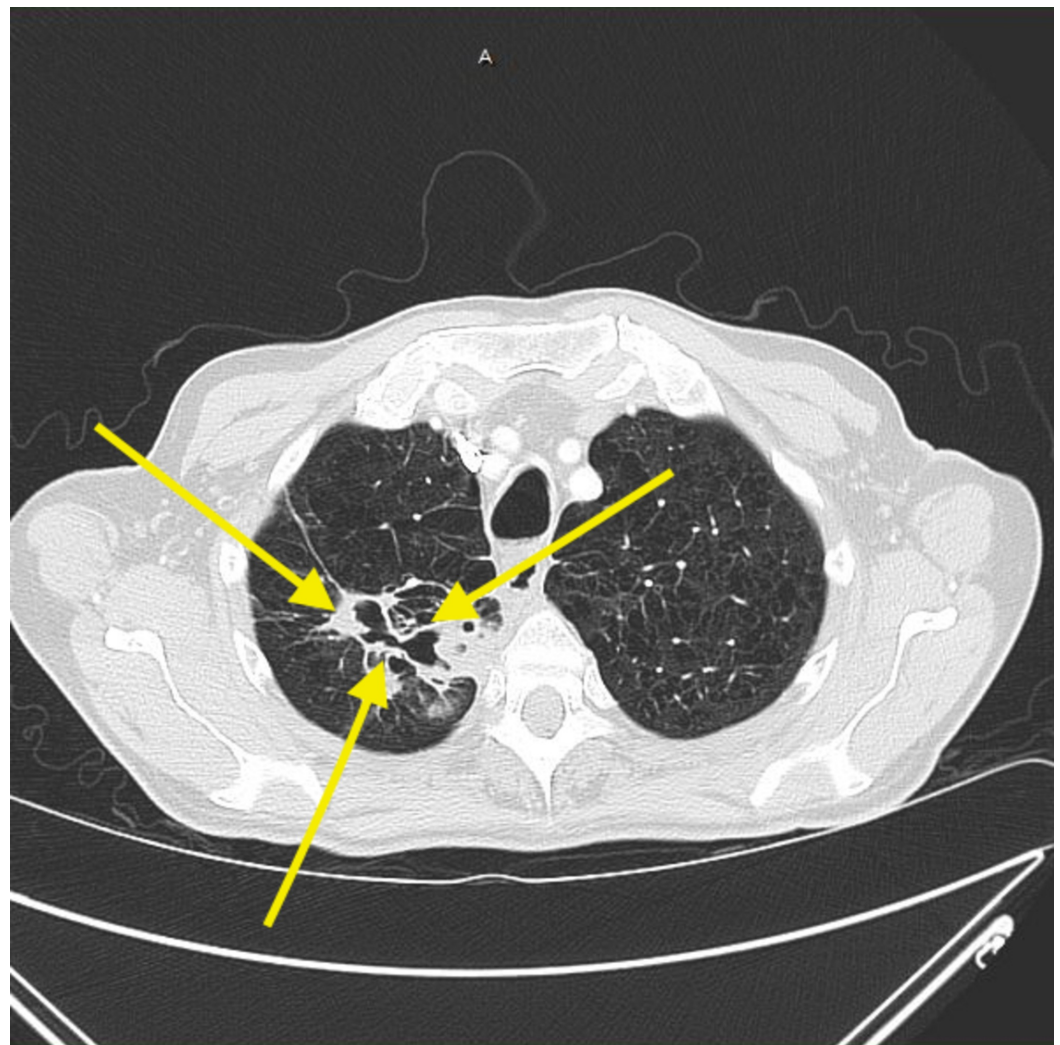
Miskoff et al. This is an open access article distributed under the terms of the Creative Commons Attribution License CC-BY 3.0., which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

### How to cite this article

Miskoff J A, Chaudhri M (June 06, 2018) Mycobacterium Chimaera: A Rare Presentation. Cureus 10(6): e2750. DOI 10.7759/cureus.2750

Lastly, the patient's bicarbonate, electrolytes, platelets, and transaminases were within normal limits. Physical examination findings of a barrel-shaped chest and poor air exchange, combined with the results of pulmonary function testing, supported a diagnosis of chronic obstructive pulmonary disease (COPD).

In addition, a chest x-ray exhibited right-sided apical opacity. The patient underwent computed tomography angiography (CTA), which was negative for pulmonary embolism and illustrated moderate to marked opacity in the right lung apex with cavitation extending up to the pleural surface. Additionally, patchy reticulonodular opacities consistent with infiltrates were present in the right upper, right middle, right lower, and left upper lobe (Figure 1). These findings led to a diagnosis of cavitory pneumonia, and blood cultures, along with mycoplasma IgM, were ordered. In addition, the patient tested negative for human immunodeficiency virus (HIV-1 and 2). The infectious disease team was consulted for their expertise and to ensure a thorough workup. Although the patient did not have true exposure to tuberculosis, the QuantiFERON®-TB Gold test (QFT-G) (Qiagen, Hilden, Germany) was ordered and the results for active and latent tuberculosis were negative. Furthermore, acid-fast bacilli (AFB) sputum cultures, fungal stains, serologies, and  $\beta$ -D-glucan were negative as well. Due to inconclusive results, a bronchoscopy with biopsies was performed which yielded a necrotic black specimen. A subsequent culture grew *Mycobacterium chimaera* and confirmed the diagnosis.



**FIGURE 1: Computed tomographic angiography (**

CTA of the patient's chest illustrating cavitation extending up to the pleural surface, along with

patchy reticulonodular infiltrates (yellow arrows).

Although a diagnosis was reached, this patient needed further workup to ensure complete resolution of this condition. The infectious disease team suggested optimizing the patient's respiratory regimen, following up with a chest CT in three months, and treating with a multidrug regimen for atypical mycobacterial infection if the patient clinically worsened.

## Discussion

*Mycobacterium avium* complex (gram-positive, nonmotile, and acid-fast) is a group of mycobacteria comprising of *intracellulare*, *avium*, and *chimaera*. *Mycobacterium chimaera* (*M. chimaera*) has been associated with low virulence, and until recently, it remained a mystery. Although *M. chimaera* was successfully isolated from water samples in 2012, it was not linked with infections in patients undergoing cardiac surgical procedures until 2013 [4]. Unfortunately, this organism did not come to the forefront until 2015, when this infectious agent was identified as a culprit responsible for two prolonged outbreaks involving prosthetic valves and associated systemic infection in the United States and Europe [1, 3]. Specifically, investigations revealed that the contaminated heater-cooler devices (Stockert 3T) came from one particular manufacturing facility, LivaNova PLC, in London [5]. The importance of heating-cooling units lies in their ability to regulate the body temperature of the patients during cardiac surgery; evidence suggests that the airborne transmission of aerosolized bacteria from the water tanks was responsible for this infection [6]. Isolates harvested from the bone marrow of a patient who underwent cardiac surgery were identical to two isolates from a single heating and cooling unit used in the same facility when the patient was operated on again [5-6]. These findings provided further evidence that strengthened the association between the units and *M. chimaera*. The research found that *M. chimaera* was predominantly found in water supply installations and could be diagnosed using 16S ribosomal RNA (rRNA) gene sequencing [3].

In addition to utilizing unique ribosomal sequencing, this group needs to be incubated for at least seven days at a specific temperature of 37° C for successful isolation [3]. Literature provides sufficient evidence associating *Mycobacterium chimaera* as the infectious organism in patients undergoing cardiac surgery (specifically, surgeries utilizing heater-cooler units) [7]. Our patient did not undergo coronary bypass surgery nor other procedures requiring these units, thus making this a unique presentation. Moreover, evidence suggests that the infection becomes clinically apparent after a lag time of months to years and this plays a significant role in *M. chimaera* remaining undiagnosed [8]. In a different study, investigators collected water and air samples from the units, wards, water circuits of heater-cooler units, and the operating room and isolated *M. chimaera* by utilizing random amplified polymorphic deoxyribonucleic acid (DNA) polymerase chain reaction (RAPD-PCR). Moreover, they determined that patients undergoing open heart surgery showed no signs of infection until 1.5 to 3.6 years after the procedure [3, 7]. Furthermore, the infection remains clinically dormant for years, making a diagnosis of *Mycobacterium chimaera* in a timely manner challenging [2, 6].

## Conclusions

*M. chimaera* is a rare species of atypical nontuberculous mycobacterium typically found in association with cardiac surgery patients. The above patient did not have the typical risk factors for *M. chimaera*, which makes this case unique. The decision to treat the patient or not presents several challenges, especially in light of severe emphysema and structural lung disease on imaging. Stabilizing the patient's COPD and concomitant consultation with an infectious disease specialist was the first step. Ultimately, the patient may require a prolonged multidrug regimen to eradicate the infection if there is any further clinical or radiographic deterioration.

Treatment of *M. chimaera* is the same as other *Mycobacterium avium* complex infections, including two to three antimicrobials for at least 12 months and often up to 18 months. Commonly used first-line drugs include macrolides, ethambutol, and rifamycins. Aminoglycosides, such as streptomycin and amikacin, are also used as additional or alternative agents. Treatment regimens may differ, especially if sensitivity testing reveals a resistant organism. The patient may also express concerns about potential adverse drug reactions and refuse or not tolerate a prolonged treatment regimen.

## Additional Information

### Disclosures

**Human subjects:** Consent was obtained by all participants in this study. **Conflicts of interest:** In compliance with the ICMJE uniform disclosure form, all authors declare the following:

**Payment/services info:** All authors have declared that no financial support was received from any organization for the submitted work. **Financial relationships:** All authors have declared that they have no financial relationships at present or within the previous three years with any organizations that might have an interest in the submitted work. **Other relationships:** All authors have declared that there are no other relationships or activities that could appear to have influenced the submitted work.

## References

1. Chand M, Lamagni T, Kranzer K, et al.: Insidious risk of severe mycobacterium chimaera infection in cardiac surgery patients. *Clin Infect Dis*. 2017, 64:335-42. [10.1093/cid/ciw754](https://doi.org/10.1093/cid/ciw754)
2. Kohler P, Kuster SP, Bloemberg G, et al.: Healthcare-associated prosthetic heart valve, aortic vascular graft, and disseminated *Mycobacterium chimaera* infections subsequent to open heart surgery. *Eur Heart J*. 2015, 36:2745-53. [10.1093/eurheartj/ehv342](https://doi.org/10.1093/eurheartj/ehv342)
3. Sax H, Bloemberg G, Hasse B, et al.: Prolonged outbreak of *Mycobacterium chimaera* infection after open-chest heart surgery. *Clin Infect Dis*. 2015, 61:67-75. [10.1093/cid/civ198](https://doi.org/10.1093/cid/civ198)
4. Jaworski R, Naumiuk Ł, Paczkowski K, et al.: *Mycobacterium chimaera* - a new threat for cardiac surgical patients?. *Kardiochir Torakochirurgia Pol*. 2017, 14:22-26. [10.5114/kitp.2017.66925](https://doi.org/10.5114/kitp.2017.66925)
5. Perkins KM, Laws A, Hasan NA, et al.: Notes from the field: *Mycobacterium chimaera* contamination of heater-cooler devices used in cardiac surgery - United States. *MMWR Morb Mortal Wkly Rep*. 2016, 65:1117-18. [10.15585/mmwr.mm6540a6](https://doi.org/10.15585/mmwr.mm6540a6)
6. Williamson D, Howden B, Stinear T: *Mycobacterium chimaera* spread from heating and cooling units in heart surgery. *N Engl J Med*. 2017, 376:600-602. [10.1056/NEJMc1612023](https://doi.org/10.1056/NEJMc1612023)
7. Sommerstein R, Rüegg C, Kohler P, et al.: Transmission of *Mycobacterium chimaera* from heater-cooler units during cardiac surgery despite an ultraclean air ventilation system. *Emerg Infect Dis*. 2016, 22:1008-13. [10.3201/eid2206.160045](https://doi.org/10.3201/eid2206.160045)
8. Walker J, Moore G, Collins S, et al.: Microbiological problems and biofilms associated with *Mycobacterium chimaera* in heater-cooler units used for cardiopulmonary bypass. *J Hosp Infect*. 2017, 96:209-20. [10.1016/j.jhin.2017.04.014](https://doi.org/10.1016/j.jhin.2017.04.014)