IDCases 22 (2020) e00977

Contents lists available at ScienceDirect

IDCases

journal homepage: www.elsevier.com/locate/idcr

are important to attain early therapy and disease control.

Miliary pattern pulmonary infiltrates in a diabetic patient: Coccidioidomycosis

Vanessa R. Wormser^{a,*}, Zi Ying Li^a, Robert Libke^b, Elham Rahmati^b

^a Department of Internal Medicine, University of California San Francisco-Fresno, Fresno, CA, United States

^b Department of Medicine, Division of Infectious Diseases, University of California San Francisco-Fresno, Fresno, CA, United States

ARTICLE INFO

ABSTRACT

Article history: Received 9 August 2020 Received in revised form 26 September 2020 Accepted 27 September 2020

Keywords: Coccidioidomycosis Miliary pattern pulmonary infiltrates Diabetes Central California

A 59-year-old Caucasian male with uncontrolled type II diabetes mellitus presented with two weeks of subjective fevers, dry cough, and worsening shortness of breath. Upon presentation, vitals were heart rate of 135 beats per min, afebrile, and tachypnea with O2 saturation 85 % on room air. His significant laboratory values were 17 % eosinophilia with white blood cells count of 10,200/µL, sodium 125mEq/L, glucose 327 mg/dL, and creatinine 1.7 mg/dL. HIV antibody/ antigen, blood cultures obtained prior to initiation of broadspectrum antimicrobials, tests for histoplasma, mycoplasma, and legionella urinary antigen were all negative. A chest radiograph showed diffuse reticulonodular pulmonary infiltrates. A subsequent computed tomography of the chest showed a right upper lobe cavitary lesion with a miliary pattern of diffuse pulmonary micronodularity (Fig. 1a, b). He required intubation and mechanical ventilation shortly after admission. Endotracheal aspirate cultures did not grow any bacterial, mycobacterial, or fungal pathogen. Coccidioidomycosis serology revealed positive IgM antibodies to Coccidioides by immunodiffusion suggestive of acute coccidioidomycosis infection. The serum was negative by complement fixation. On day 6 of hospitalization, the patient developed septic shock requiring vasopressors and acute kidney injury requiring hemodialysis. The patient's clinical course continued to deteriorate

* Corresponding author at: UCSF Fresno Medical Education Program, Internal Medicine, 155 N Fresno St., Fresno, CA, 93701, United States.

E-mail address: VWormser@fresno.ucsf.edu (V.R. Wormser).

despite intravenous liposomal amphotericin B and fluconazole and he eventually expired.

license (http://creativecommons.org/licenses/by-nc-nd/4.0/).

This is a case of miliary coccidioidomycosis. This case illustrates the importance of early suspicion for

coccidioidomycosis in patients from an endemic area. Early identification and recognition of the disease

© 2020 The Authors. Published by Elsevier Ltd. This is an open access article under the CC BY-NC-ND

Coccidioidomycosis is an endemic fungal infection in the Southwestern regions of the United States caused by dimorphic fungi *Coccidioides immitis or Coccidioides posadasii*, with *C. immitis* being more prevalent in California's Central Valley. Individuals living in endemic areas with diabetes, HIV, on chemotherapy or other immunosuppressive medications, or pregnant in the third trimester are all at risk of disseminated disease [1].

In correlation with clinical symptoms, serum serology, bodily fluid smear, and culture are used to establish the diagnosis of coccidioidomycosis. In our patient, the fungal cultures were negative due to a low sensitivity of coccidioidomycosis cultures in the setting of miliary coccidioidomycosis which is thought to occur through hematogenous or lymphatic spread and not due to a primary parenchymal lung disease. Sputum cultures are positive in less than 40 % of cases and lung biopsy is often required for diagnosis of miliary nodules [2]. Enzyme-linked immunoassays (EIA) and immunodiffusion are the most common tests used to make a diagnosis of coccidioidomycosis. In our patient, the serum was negative by complement fixation which may have been due to the timing of the test. The optimal detection of complement fixation peaks at approximately 1-2 months from the onset of symptomatic infection [3]. Polymerase chain reaction (PCR) assay has been used in our central California hospital laboratory to identify C. immitis in specimens including sputum, and bronchioalveolar lavage (BAL) with a more rapid result time of 4 h, and similar specificity and sensitivity when comparing other

2214-2509/© 2020 The Authors. Published by Elsevier Ltd. This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).



Case illustrated





https://doi.org/10.1016/j.idcr.2020.e00977



Fig. 1. (a) Coronal view of the CT chest showing right upper lobe pulmonary cavity accompanied by a miliary pattern of diffuse pulmonary micronodularity with associated mediastinal and hilar adenopathy. (b) Axial view of the miliary pattern of diffuse pulmonary micronodularity.

serological testing and fungal cultures [4]. PCR was not yet available during our patient's hospitalization.

In immunosuppressed individuals, disseminated disease such as miliary coccidioidomycosis may occur through hematogenous or lymphatic spread, and lead to respiratory failure [5].

A miliary pattern on chest imaging may be seen with metastatic cancer, tuberculosis, coccidioidomycosis, histoplasmosis, or candidiasis [6].

Amphotericin B and azole in combination or alone are used for the treatment of miliary coccidioidomycosis. Medical providers should optimize testing and management for immunocompromised patients in coccidioidomycosis endemic areas to reduce the risk of infection and the rate of dissemination.

Declaration of Competing Interest

No conflicts of interest. Nothing to Declare

Funding

Publication made possible in part by support from the UCSF Open Access Publishing Fund.

Consent

Informed consent was obtained for publication of this case report and accompanying images.

CRediT authorship contribution statement

Vanessa R. Wormser: Conceptualization, Writing - original draft. Zi Ying Li: Conceptualization, Writing - original draft. Robert Libke: Writing - review & editing. Elham Rahmati: Writing review & editing.

References

- [1] Sotello D, Marcella R, Audra F, et al. Coccidioidomycosis with diffuse miliary pneumonia. Baylor Univ Med Cent Proc 2016;29(1):39-41. [2] Bayer AS. Fungal Pneumonias: Pulmonary Coccidioidal syndromes (Part 2).
- CHEST 1981;79(6):686-91.
- [3] Blair JE, Mendoza N, Force S, et al. Clinical specificity of the enzyme immunoassay test for coccidioidomycosis varies according to the reason for its performance. Clin Vaccine Immunol 2013;20(1):95-8, doi:http://dx.doi.org/ 10.1128/CVI.00531-12.
- [4] Dominic D, Marilyn M, Bernadette D, et al. The utility of real-time polymerase chain reaction in detecting Coccidioides immitis among clinical specimens in the Central California san Joaquin Valley. Med Mycol 2019;57(August (6)):688-93.
- [5] Spinello IM, Munoz A, Johnson RH. Pulmonary coccidioidomycosis. Semin Respir Crit Care Med 2008;29(2):166-73.
- [6] PPTES Torres, Rabahi MF, Moreira MAC, et al. Tomographic assessment of thoracic fungal diseases: a pattern and signs approach. Radiol Bras 2018;51(Sep-Oct(5)):313-21.