



Case Report

Pulmonary melioidosis misdiagnosed as pulmonary tuberculosis

Chandan Kumar Sheet^{1,*}, Saibal Ghosh¹, Subhankar Chatterjee¹, Biplab Chandra¹¹Dept. of Pulmonary Medicine and Critical Care, Calcutta Heart Clinic & Hospital, Bidhannagar, Kolkata, West Bengal, India

ARTICLE INFO

Article history:

Received 11-07-2023

Accepted 07-10-2023

Available online xx xx xxxx

Keywords:

Melioidosis

Burkholderia pseudomallei

Tuberculosis

ABSTRACT

Background: *Burkholderia pseudomallei* is a facultative Gram-negative saprophytic bacterium commonly found in soil or contaminated water causing melioidosis. Melioidosis can mimic various other disease due to its heterogenous clinical manifestations and different organ involvement. Because of its versatility it is called as “the great imitator” and remains challenging to diagnose. We report a case of melioidosis misdiagnosed and treated as pulmonary tuberculosis.

Case Presentation: A 54-year-old male non-smoker with history of diabetes admitted with persistent cough, breathlessness and hemoptysis for 5 months. Initially there was pleuritic chest pain and high-grade fever. He was treated with multiple intravenous broad-spectrum antibiotics and anti-tubercular therapy multiple times in the local hospitals based on clinical symptoms and radiological manifestation though Sputum examination for AFB and Gene xpert for Mycobacterium tuberculosis was negative. Chest radiology showed multiple thick-walled cavities with pericavitary consolidation along with patchy infiltrative opacities. BALF culture identified *Burkholderia pseudomallei*. The respiratory morbidity was resolved using antibiotics based on antibiotic susceptibility tests.

Conclusion: This case study described a case of melioidosis in adult male with diabetes and engaged in farming presented with diverse and indistinct clinical manifestations that mimics many other diseases. Definitive diagnosis was made by isolation *Burkholderia pseudomallei*, in culture collected through bronchoscopic examination.

This is an Open Access (OA) journal, and articles are distributed under the terms of the [Creative Commons Attribution-NonCommercial-ShareAlike 4.0 License](https://creativecommons.org/licenses/by-nc-sa/4.0/), which allows others to remix, tweak, and build upon the work non-commercially, as long as appropriate credit is given and the new creations are licensed under the identical terms.

For reprints contact: reprint@ipinnovative.com

1. Introduction

Gram-negative bacillus *Burkholderia (Pseudomonas) pseudomallei* (Whitmore bacillus) is the causative organism for a rare infectious disease called Melioidosis which could affect whole body but the most commonly affected organ is the lung followed by spleen, skin and soft tissue. It could manifest in acute, subacute, or chronic forms.¹ It is an emerging, potentially life-threatening infection in India as well as south East Asia.

2. Case Presentation

A 54 years male from Assam, India, and farmer by occupation admitted with persistent cough, breathlessness and hemoptysis for 5 months. Initially there was pleuritic type chest pain and high-grade fever. He was treated with multiple intravenous broad-spectrum antibiotics in the local hospitals. Sputum examination for acid-fast bacilli (AFB) and Gene xpert for Mycobacterium tuberculosis was done which came out to be negative. He was started with anti-tubercular therapy (HREZ regime (consisting of isoniazid (H), rifampicin (R), ethambutol (E) and pyrazinamide (Z)) initially then stopped after 2 months and again started with HREZ) based on clinical symptoms and radiological manifestation. Despite anti tubercular therapy there was

* Corresponding author.

E-mail address: chandansheet1986@gmail.com (C. K. Sheet).

24 recurrent hemoptysis and he presented to us after 5 months
 25 with worsening of symptoms. He was diabetic with good
 26 glycemic control and non-smoker.

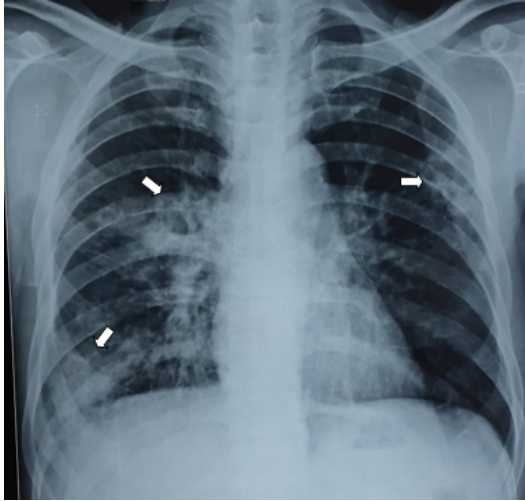


Fig. 1: CXR on initiation of symptoms (done outside our hospital) showing multiple small cavitary lesion with pericavitary consolidation in bilateral upper lobe, and right lower lobe and right parahilar region (white arrow).

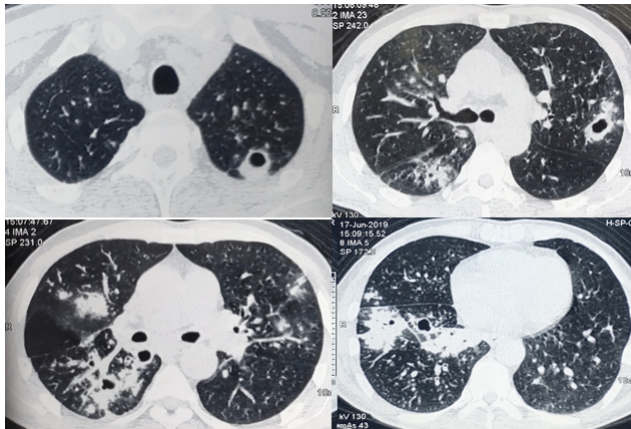


Fig. 2: HRCT chest at presentation: showing multiple thick-walled cavities with pericavitary consolidation in RUL, RLL, LUL and LLL with parenchymal infiltrations and tree in bud appearance

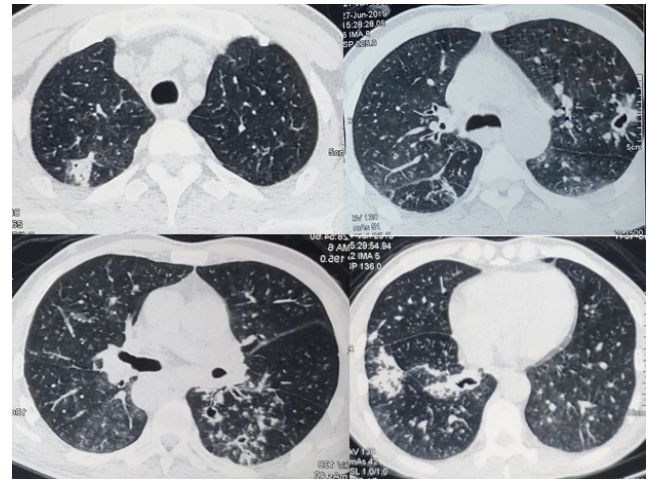


Fig. 3: HRCT chest after 10 days of proper antibiotic therapy showing resolving consolidation, cavitary lesion and decreased infiltrations

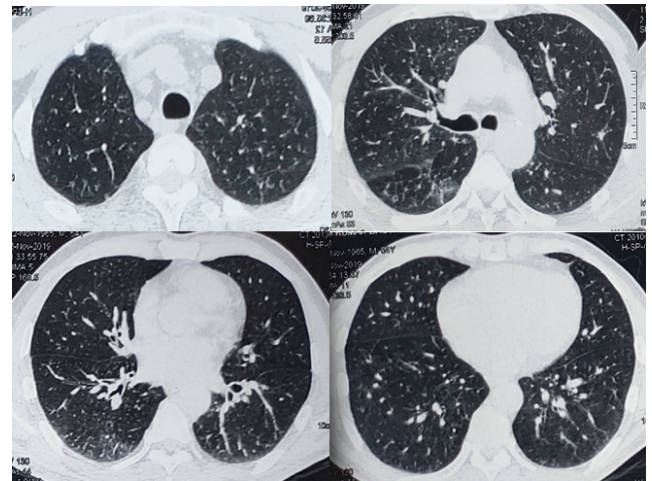


Fig. 4: HRCT after 9 months showing complete resolution of lung opacities

27 On general examination, he was febrile and ill looking.
 28 There was pallor but no clubbing, palpable lymph node
 29 or skin lesion present. Examination of respiratory system
 30 revealed bilateral coarse inspiratory crepitation in both
 31 lungs with an isolated area of bronchial breath sound
 32 without any evidence of pleural effusions. No abnormalities
 33 were detected on cardiac and abdominal examinations.

34 On admission there was microcytic hypochromic anemia
 35 (Hb-8.4%), Neutrophilic leukocytosis (WBC - 14900,
 36 Neutrophil-92%), raised erythrocyte sedimentation rate
 37 (ESR) of 132 mm/hr. and C- reactive protein of 236 mg/l.

Chest X-ray (Figure 1) showed multiple cavitary lesion
 with pericavitary consolidation in bilateral upper lobe
 and right lower lobe. and multiple patchy infiltrates in both
 lung fields. HRCT chest (Figure 2) showed multiple thick-
 walled cavities with pericavitary consolidation in left upper
 lobe (LUL), Right upper lobe (RUL), right lower lobe
 (RLL) along with patchy infiltrative opacities. Mantoux test
 with 5TU and Sputum examination for AFB were negative.
 Sputum bacterial culture and sensitivity showed no growth.
 He was started with empirical intravenous Piperacillin and
 tazobactam and doxycycline. However, symptoms including
 fever persisted. Subsequently, his clinical condition was
 worsened with elevated inflammatory markers. Possibility
 of infective endocarditis was excluded by a transthoracic
 echocardiography. Tests for antinuclear antibody (ANA),

53 antinuclear cytoplasmic antibodies (ANCA) and retroviral
54 screening were also negative. Subsequently, a bronchoscopy
55 was done and bronchoalveolar lavage (BAL) taken.
56 BAL culture grew *Burkholderia pseudomallei* sensitive
57 to ceftazidime, and cotrimoxazole. BAL for GeneXpert
58 for mycobacterium tuberculosis, galactomannan and fungal
59 culture were negative. BAL cytology for malignant cell was
60 negative.

61 According to the antibiotic sensitivity pattern, he was
62 started with IV ceftazidime for 10 days and significant
63 clinical and radiological improvement (Figure 3) was found.
64 He was discharged with oral cotrimoxazole twice daily for
65 next 12 weeks.² Repeat HRCT after 9 months revealed
66 total radiological resolution and no recurrence of symptoms
67 (Figure 4).

68 3. Discussion

69 *B. pseudomallei* (causative organism of Melioidosis) is
70 a facultative Gram-negative saprophytic bacterium and
71 is commonly found in soil or contaminated water.³
72 Risk factors for melioidosis include chronic alcohol use,
73 diseases (such as diabetes mellitus, thalassemia and renal
74 disease), immunosuppressive therapy including steroids and
75 occupational exposure to contaminated soil or water.^{4,5} As
76 our patient was known diabetic and farmer by occupation
77 with a possibility of contact with the contaminated water
78 and soil. This may have predisposed him to melioidosis
79 infection.

80 It could present with diverse clinical manifestations and
81 organ involvement depending upon the duration of infection
82 and could mimic many diseases (earning a name “the great
83 imitator”).⁶ The Darwin study found pneumonia to be the
84 most common presentation of melioidosis (50%) followed
85 by genitourinary infection (14%), skin infection (13%),
86 non-specific bacteremia (11%), and less commonly septic
87 arthritis or osteomyelitis (4%) and neurological melioidosis
88 (3%).⁷ Because of this, its clinical diagnosis remains a
89 challenge. Acute melioidosis, usually rapidly progressive
90 and predominantly affects upper lobes with early cavitation.
91 On the other hand, in subacute and chronic forms, it
92 could mimic tuberculosis in radiological examination, with
93 involvement of upper lobe and/or patchy alveolar infiltrate
94 with cavities or fibroreticular lesions.⁸

95 4. Conclusions

96 This case study described a case of melioidosis in adult
97 male with diabetes and engaged in farming presented with
98 diverse and indistinct clinical manifestations that mimics
99 many other diseases. Definitive diagnosis was made by
100 isolation *Burkholderia pseudomallei*, in culture collected
101 through bronchoscopic examination.

102 5. Declaration of Patient Consent

103 Identity and confidentiality of the patient maintained
104 properly; proper informed consent has been obtained as
105 well.

106 6. Source of Funding

107 None.

108 7. Conflict of Funding

109 None.

110 8. Acknowledgments

111 Contributors: CKS, SG: clinical care of the patient. SC,
112 CKS: compiling data and interpretation, figures, manuscript
113 writing. CKS, SC & BC: revision and final approval of
114 manuscript. Final approval of the version published has been
115 agreed by SG, SC, BC & CKS.

116 References

- 117 1. Orphanet. Melioidosis; 2023. [Last accessed 30/09/2023]. Available
118 from: [https://www.orpha.net/consor/cgi-bin/OC_Exp.php?lng=en&
119 Expert=31202](https://www.orpha.net/consor/cgi-bin/OC_Exp.php?lng=en&Expert=31202).
- 120 2. Centers for disease control and prevention. Melioidosis; 2023.
121 [Last accessed 30/09/2023]. Available from: [https://www.cdc.gov/
122 melioidosis/treatment/index.html](https://www.cdc.gov/melioidosis/treatment/index.html).
- 123 3. Wiersinga WJ, Virk HS, Torres AG, Currie BJ, Peacock SJ, Dance
124 DAB, et al. Melioidosis. *Nat Rev Dis Prim*. 2018;4:17107.
125 doi:10.1038/nrdp.2017.107.
- 126 4. Currie BJ, Fisher DA, Howard DM, Burrow JNC, Lo D, Selvanayagam
127 S, et al. Endemic melioidosis in tropical northern Australia: a 10-
128 year prospective study and review of the literature. *Clin Infect Dis*.
129 2000;31(4):981-6.
- 130 5. Suputtamongkol Y, Chaowagul W, Chetchotisakd P, Lertpatanasuwun
131 N, Intaranongpai S, Ruchtrakool T, et al. Risk factors for melioidosis
132 and bacteremic melioidosis. *Clin Infect Dis*. 1999;29(2):408-13.
- 133 6. Chang CY. A case report of pulmonary melioidosis with the air crescent
134 sign. *Radiol Infect Dis*. 2000;7(1):31-4.
- 135 7. Abramson S. The air crescent sign. *Radiology*. 2001;218(1):230-2.
- 136 8. Reecheaipichitkul W. Clinical manifestation of pulmonary melioidosis
137 in adults. *Southeast Asian J Trop Med Public Health*. 2004;35(3):664-
138 9.

139 Author biography

140 **Chandan Kumar Sheet**, Consultant  [https://orcid.org/0000-0002-4173-
141 2377](https://orcid.org/0000-0002-4173-2377)

142 **Saibal Ghosh**, Consultant

143 **Subhankar Chatterjee**, Consultant

144 **Bioplal Chandra**, Consultant

Cite this article: Sheet CK, Ghosh S, Chatterjee S, Chandra B.
145 Pulmonary melioidosis misdiagnosed as pulmonary tuberculosis. *IP
Indian J Immunol Respir Med* 2023;8(3):1-3.