Since January 2020 Elsevier has created a COVID-19 resource centre with free information in English and Mandarin on the novel coronavirus COVID-19. The COVID-19 resource centre is hosted on Elsevier Connect, the company's public news and information website.

Elsevier hereby grants permission to make all its COVID-19-related research that is available on the COVID-19 resource centre - including this research content - immediately available in PubMed Central and other publicly funded repositories, such as the WHO COVID database with rights for unrestricted research re-use and analyses in any form or by any means with acknowledgement of the original source. These permissions are granted for free by Elsevier for as long as the COVID-19 resource centre remains active.
Pulmonary tuberculosis and COVID-19 coinfection: Hickam's Dictum revisited

Larry Ellee Nyanti, Zhun Han Wong, Benjamin Sachdev Manjit Singh, Andrew Kean Wei Chang, Ahmad Tirmizi Jobli, Hock Hin Chua

PII: S2213-0071(22)00075-2
DOI: https://doi.org/10.1016/j.rmcr.2022.101653
Reference: RMCR 101653

To appear in: Respiratory Medicine Case Reports

Received Date: 5 February 2022
Revised Date: 12 March 2022
Accepted Date: 13 April 2022

Please cite this article as: Nyanti LE, Wong ZH, Sachdev Manjit Singh B, Chang AKW, Jobli AT, Chua HH, Pulmonary tuberculosis and COVID-19 coinfection: Hickam's Dictum revisited, Respiratory Medicine Case Reports (2022), doi: https://doi.org/10.1016/j.rmcr.2022.101653.

This is a PDF file of an article that has undergone enhancements after acceptance, such as the addition of a cover page and metadata, and formatting for readability, but it is not yet the definitive version of record. This version will undergo additional copyediting, typesetting and review before it is published in its final form, but we are providing this version to give early visibility of the article. Please note that, during the production process, errors may be discovered which could affect the content, and all legal disclaimers that apply to the journal pertain.

© 2022 Published by Elsevier Ltd.
Pulmonary tuberculosis and COVID-19 coinfection: Hickam’s Dictum revisited

Author list (family name in all-caps)

Larry Ellee NYANTI1,2 MRCPI larryen90@yahoo.com ORCID: 0000-0002-5790-3919
Zhun Han WONG1 MRCPI zhunhan@hotmail.com
Benjamin SACHDEV MANJIT SINGH1 MRCP benjaminsachdev@gmail.com ORCID: 0000-0002-0978-8602
Andrew Kean Wei CHANG1 MRCP andrchang@gmail.com ORCID: 0000-0003-2557-170X
Ahmad Tirmizi JOBLI3 MMed jatirmizi@unimas.my ORCID: 0000-0002-8499-3315
Hock Hin CHUA1 FRCP hhchua2009@gmail.com ORCID: 0000-0001-6004-4273

Affiliation

1Infectious Disease Unit, Medical Department, Sarawak General Hospital, Jalan Hospital, 93586 Kuching, Sarawak, Malaysia.
2Faculty of Medicine and Health Sciences, Universiti Malaysia Sabah, Kota Kinabalu, Sabah, Malaysia.
3Faculty of Medicine and Health Sciences, University Malaysia Sarawak, Kota Samarahan, Sarawak, Malaysia.

Corresponding author:
Larry Ellee Nyanti
Email: larrynyanti@ums.edu.my
Address: Faculty of Medicine and Health Sciences, Universiti Malaysia Sabah, Kota Kinabalu, Sabah, Malaysia.
List of abbreviations

| Abbreviation | Full Form |
|--------------|-----------|
| tuberculosis | TB        |
| pulmonary tuberculosis | PTB     |
| computed tomography | CT       |
| World Health Organization | WHO      |
| nasopharyngeal and oropharyngeal | NPOP   |
| computed tomography pulmonary angiogram | CTPA |
| high resolution computed tomography | HRCT   |
| ground glass opacities | GGO      |
| anti-tuberculous therapy | ATT    |
| interferon-gamma release assay | IGRA |

Abstract

COVID-19 and pulmonary tuberculosis (PTB) coinfection is associated with increased mortality and presents a unique diagnostic challenge to the clinician. We describe three cases of newly diagnosed PTB in COVID-19 patients treated at our centre and their clinical and radiological features. The challenges associated with diagnosis and management are also explored. Patient 1 was a case of smear positive, endobronchial tuberculosis incidentally diagnosed due to CT changes, and eventually made good recovery. Patient 2 was a case of COVID-19 who succumbed but was diagnosed posthumously due to a positive sputum culture for tuberculosis. Patient 3 showed radiographic features of PTB and was treated empirically for TB. In conclusion, COVID-19 and PTB coinfection should be suspected in the presence of constitutional symptoms, prior immunocompromised states, prolonged respiratory symptoms or fever, or unresolved radiological abnormalities, more so in regions where TB is endemic.

Keywords

COVID-19, Pulmonary Tuberculosis, Coinfection, Endobronchial Tuberculosis, Case Series
1.1 Introduction

COVID-19 was declared a pandemic in 2020, whereas tuberculosis (TB) was declared a global health emergency by the World Health Organization (WHO) in 1993 [1]. Both diseases manifest as respiratory illnesses with varying severities and display similar symptoms [2], reflecting Hickam’s dictum which states that multiple symptoms and signs may be due to more than one disease [3]. TB is endemic in the state of Sarawak, Malaysia, with a 2016 incidence of 104.23 per 100,000 population, far above the national average of 81.3 per 100,000 population [4]. Pulmonary TB (PTB) and COVID-19 coinfection has been well-reported [2,5,6,7,8,9] with meta-analysis showing increased risk of mortality [10]. We describe three cases of newly diagnosed PTB in COVID-19 patients treated at our tertiary centre, discuss their varied clinical and radiological presentations, and elucidate the challenges associated with diagnosis and management of PTB and COVID-19 coinfection. All three patients were infected with COVID-19 before the introduction of COVID vaccines.

2.1 Case presentation

2.1.1 Patient 1

A 61-year-old woman with a history of total abdominal hysterectomy and salpingooopherectomy for cervical carcinoma initially visited the surgical clinic with complaints of abdominal bloating and loss of weight for a month, associated with a recent dry cough for one week. She denied shortness of breath, fever, haemoptysis, sore throat, anosmia or dysgeusia. Computed tomography (CT) of the abdomen and pelvis showed no intrabdominal abnormalities, however, ground glass opacities were observed at bilateral lower zones of the lung. Rapid nasopharyngeal and oropharyngeal (NPOP) antigen assay for COVID-19 was positive, and she was admitted to the isolation ward.

On admission, she was apyrexial and normotensive, but was noted to be tachycardic (heart rate of 128 beats per minute), tachypnoeic (respiratory rate of 22 breaths per minute) and had an oxygen saturation of 89% under room air. Arterial blood gas taken under room air revealed type 1 respiratory failure (pH 7.46, PO$_2$ 67mmHg, PCO$_2$ 30mmHg, HCO$_3$ 28 mmol/L). Electrocardiogram showed atrial fibrillation with fast ventricular response, but no ischemic or myocarditis changes. Chest radiograph revealed bilateral nodular and airspace opacities (Figure 1A).

Dexamethasone and lopinavir/ritonavir were administered as per local COVID-19 treatment guidelines at the time. She required invasive mechanical ventilation for three days. Endotracheal aspirates sampled for bacterial culture and respiratory virus PCR panel (QIAstat-Dx, QIAGEN, Maryland, United States) were negative. Post-extubation, she...
remained oxygen dependent and computed tomography pulmonary angiogram (CTPA) was performed on day 7 of admission. CTPA showed pulmonary embolism in the segmental branch of right descending pulmonary artery, generalized, diffuse peri-bronchovascular ground glass changes bilaterally, interlobular and intralobular septal thickening at bilateral upper lobes, focal consolidation at the lateral basal segment of the left lower lobe, and tree-in-bud nodules bilaterally (Figure 2A). Sputum smear examination was positive for acid fast bacilli. Molecular testing for *M. TB* nucleic acids (Cepheid Xpert MTB/RIF, https://www.cepheid.com) revealed no rifampicin resistance. She was commenced on anticoagulant and anti-tuberculous therapy (ATT) consisting of rifampicin, isoniazid, pyrazinamide, and ethambutol, and finally weaned off to room air on day 32 of admission. Follow-on high resolution computed tomography (HRCT) of the thorax at day 40 of admission showed resolution of ground glass changes and absence of septal thickening. However, there were increased diffuse centriflobular lung nodules with traction bronchiectasis at bilateral lower lobes, consistent with endobronchial TB (Figure 2B). She was discharged well and completed a six-month course of ATT. Chest radiograph taken 3 months post-discharge showed resolving opacities (Figure 1B).

### 2.1.2 Patient 2

A 63-year-old woman with underlying type 2 diabetes mellitus presented to a district clinic with loss of weight and loss of appetite for the last two months, two weeks of lethargy and poor oral intake, one week of fever and productive cough with whitish sputum, and rapidly worsening shortness of breath for the past two days. She was normotensive but tachycardic (pulse rate 115 beats per minute) and tachypnoeic (respiratory rate 28 breaths per minute) and had an oxygen saturation of 96% under room air. She was hyperglycaemic (13 mmol/L) and pyretic (temperature of 38.5 degree Celsius). Auscultation of lungs revealed reduced air entry over the right lower zone, and chest radiograph revealed bilateral lower zone airspace opacities with blunting of the right costophrenic angle (Figure 3A). She was then referred to our centre for further care. On arrival to the emergency department, arterial blood gas revealed type 1 respiratory failure despite on 40% supplementation via Venturi mask (pH 7.5 PO$_2$ 69 mmHg PCO$_2$ 29 mmHg HCO$_3$ 22 mmol/L) and she required bi-level positive airway pressure ventilation.

NPOP sample sent for COVID-19 polymerase chain reaction testing was positive. Dexamethasone, lopinavir/ritonavir and favipiravir were administered as per local COVID-19 guidelines at the time. In view of constitutional symptoms, three successive sputum samples were submitted for smear examination, but were negative for acid fast bacilli. The sputum samples were then cultured in Lowenstein-Jensen media. Her pyrexia subsided and she was weaned off to
room air by day 10 of admission. However, on day 12 of admission, she complained of chills and breathlessness, and her saturation dropped to 80% under room air. Repeated chest radiograph showed progression of airspace opacities involving mid zones bilaterally (Figure 3B). Intravenous meropenem was commenced. She succumbed on day 17 of admission. Posthumously, mycobacterium TB complex was isolated in her sputum sample after seven weeks. First-line drug sensitivity testing confirmed susceptibility to isoniazid, rifampicin, ethambutol, and streptomycin.

2.1.3 Patient 3
A 78-year-old man with underlying hypertension presented to the emergency department with dry cough, loss of weight and intermittent fever for two months with recent progressive shortness of breath for the past two weeks. He was normotensive and afebrile, and not tachycardic. He had a normal oxygen saturation of 98% under room air but was mildly tachypnoeic (respiratory rate of 21 breaths per minute). Clinically, he appeared cachetic. Chest radiograph showed bilateral nodular and airspace opacities with left mid zone consolidation (Figure 4A). NPOP sample sent for COVID-19 polymerase chain reaction testing was positive, and he was admitted for isolation and treated with oral favipiravir. ATT was immediately commenced in view of suggestive chest radiograph and constitutional symptoms. Subsequently, sputum smear examination was positive for acid fast bacilli. Chest radiograph on day 14 of admission showed partial resolution of the left mid zone consolidation with residual nodular opacities (Figure 4B). He was discharged well to continue his TB treatment as an outpatient. He did not require oxygen supplementation throughout his admission.

3.1 Discussion
Individuals with active PTB are at higher risk of contracting COVID-19 due to alterations in lung immunity driven by attenuated interferon-gamma host response to SARS-CoV-2 virus [11]. Meanwhile, reactivation of latent TB in the setting of COVID-19 infection is plausible, given that the two diseases augment each other with a transient decrease in cellular immunity [12]. Meta-analysis showed that the summary proportion of active PTB among COVID-19 patients is higher than WHO estimates for annual incidence of tuberculosis in high tuberculosis burden countries such as China, India, Nigeria, Philippines, and South Africa [10]. This is not surprising, given that both COVID-19 and PTB spread through airborne transmission of close contacts [10]. Furthermore, stay-at-home recommendations and national lockdown policies may have exacerbated crowded living conditions among underprivileged communities [13].
A large meta-analysis of 43 studies by Aggarwal et al. showed that patients with active PTB and COVID-19 carried a two-fold higher relative risk of death, which was comparable to relative risk of mortality in COVID-19 patients with diabetes, hypertension, and cardiovascular diseases [10]. In resource-limited settings, poverty and malnutrition might play an important role in increasing morbidity and mortality [8]. The proportion of PTB in our cohort of COVID-19 patients is 0.6%, or 3 out of 498 cases, presenting over a span of five months. This proportion is not generalizable as this was a single-centre study conducted in a limited time frame and did not include the entirety of COVID-19 patients admitted to non-tertiary quarantine centres or other tertiary hospitals in the state.

This case series represents varying scenarios in which TB might present in the setting of COVID-19. One of the uncommon forms of pulmonary tuberculosis is endobronchial TB, as described in the case of patient 1. While PTB commonly presents with cough (88%), loss of weight (58.1%), loss of appetite (57.3%), fever (56.4%), night sweats (30%), and haemoptysis (30%), endobronchial TB tends to present with dry cough, chest discomfort, and less commonly, wheezing or stridor in the presence of bronchial stenosis [14,15]. Further obscuring the diagnosis of TB are the shared x-ray abnormalities of endobronchial TB and COVID-19. Endobronchial TB may present as patchy parenchymal infiltrates, consolidations, segmental and lobar collapse, bronchiectasis, and consolidation [16] while COVID-19 present as ground glass opacities, coarse linear opacities, and consolidations [17].

CT may prove essential in differentiating endobronchial TB and COVID-19. Endobronchial TB tends to present with centrilobular nodules, tree-in-bud appearance, consolidations, thickening of the bronchial wall and interlobular septa [14], while COVID-19 tends to show ground glass opacities (GGO), intralobular septal thickening, consolidations, and parenchymal bands [18,19]. Interestingly, in one case series, nearly half the HRCT in coinfection cases had no TB characteristic changes [8]. In the absence of CT, lung ultrasonography may prove useful - one small study demonstrated that COVID-19 subpleural lesions differed significantly from similar ones observed in bacterial pneumonia, pulmonary abscess, tuberculosis, atelectasis, and cardiogenic pulmonary edema [20]. However, whether tuberculosis changes can be differentiated from COVID-19 on LUS has yet to be demonstrated.

Resolution of GGO tends to occur after day fourteen of diagnosis of COVID-19, whereas consolidation and intralobular septal thickening tend to persist [19]. By 2 months post-COVID diagnosis, a third of COVID-19 patients showed complete resolution, while nearly half had residual parenchymal bands [18]. Similarly in our patient, the repeated CT on
day 40 showed resolution of her initial consolidation and septal thickening. There are no known studies on the short-term effect of TB treatment on CT abnormalities in endobronchial TB. One study demonstrated complete resolution of changes in 60% (33 out of 55) of patients after treatment course completion [21]. Nodules (inter and intra-lobular) are not a common CT feature of COVID-19 infection [18,19], hence should raise suspicion of an alternate diagnosis, such as TB. The differences between the two CTs in patient 1 may demonstrate not only the resolution of COVID disease, but also an unmasking of underlying endobronchial TB.

Patient 2 illustrates a case of severe COVID-19 complicated by hospital acquired pneumonia with posthumous diagnosis of smear-negative, culture-positive PTB. The absence of acid-fast bacilli on respiratory samples may have reflected poor sampling quality or low tuberculous load in the lungs. The use of high dose steroids has been associated with TB reactivation; our patient was given dexamethasone for the treatment of COVID-19 infection which may have contributed to a worsened outcome [22]. Given her constitutional symptoms and initial chest radiograph showing unilateral pleural effusion, follow-on rapid diagnostic methods like Xpert MTB/RIF may have contributed to a more favourable outcome in this case by ensuring early commencement of ATT. Xpert MTB/RIF has been quoted to have a detection rate of 95% [8]. Another method is interferon-gamma release assay (IGRA) (QIAGEN, https://www.qiagen.com), which has been reported to pick up TB in Xpert MTB/RIF negative cases [23]. While several countries have proposed bidirectional screening of COVID-19 and PTB, implementation is difficult in resource-limited settings [10]. In resource-limited settings, we propose sputum smear for acid-fast bacilli as preliminary screening in COVID-19 patients with high risk for PTB.

Finally, patient 3 illustrates a clinical scenario that allows for a high index of suspicion for TB, and thus, early commencement of ATT. A history of chronic loss of weight and fever is not a feature of acute COVID-19 infection and warrants further investigation for TB. The resolution of changes on chest radiograph after two weeks may reflect either resolution of COVID-19 infection or response to ATT, nevertheless, a repeat chest radiograph and sputum examination after treatment course is warranted.

4.1 Conclusion

In TB-endemic regions, COVID-19 and PTB coinfection should be suspected in the presence of constitutional symptoms, prior immunocompromised states, prolonged respiratory symptoms or fever, unresolved radiological abnormalities, or
a prolonged dependence on oxygen supplementation. Clinicians are reminded that Hickam’s Dictum remains relevant to this day.

**Declarations**

**Ethics approval and consent to participate**

This case series was registered via National Medical Research Register Malaysia (NMRR ID: NMRR ID-21-02248-OBB). Written consent was obtained from the patients.

**Consent for publication**

Written consent was obtained from the patients for publication of this case series and accompanying images.

**Availability of data and materials**

Data sharing is not applicable to this article as no datasets were generated or analyzed during the current study.

**Declaration of interests**

None

**Funding**

The authors declare that no funding was used for the publication of this manuscript.

**Authors’ contributions**

LEN and ZHW initiated the idea for case reporting. LEN, BSMS, AKWC, AJT and HHC prepared the final copy of the manuscript. LEN, ZHW, BSMS, AKWC, AJT and HHC were involved in the overall management of the patients. All authors have read and approved the final manuscript.

**Acknowledgements**

The authors would like to thank the respiratory team of Sarawak General Hospital for their input on patient care, the Faculty of Medicine and Health Sciences, University Malaysia Sabah for sponsoring this paper’s publication fees, and the Director General of Health Malaysia for permission to publish this paper.
References

1. Dara M, Sotgiu G, Reichler MR. et al. New diseases and old threats: lessons from TB for the COVID-19 response. Int J Tuberc Lung Dis. 2020 May 1;24(5):544-545. doi: 10.5588/ijtld.20.0151.

2. Musso M, Di Gennaro F, Gualano G, et al. Concurrent cavitary pulmonary TB and COVID-19 pneumonia with in vitro immune cell anergy. Infection. 2021 Jan 17;4–4. doi: 10.1007/s15010-021-0.

3. Mani N, Slevin N, Hudson A. What Three Wise Men have to say about diagnosis. BMJ. 2011 Dec 20;343:d7769. doi: 10.1136/bmj.d7769. PMID: 22187188.

4. Malaysian Healthcare Performance Unit, Malaysian Health at a Glance: 2018, Ministry of Health Malaysia: Putrajaya.1576-
y.

5. Sy KTL, Haw NJL, Uy J. Previous and active tuberculosis increases risk of death and prolongs recovery in patients with COVID-19. Infect Dis (Lond). 2020 Nov-Dec;52(12):902-907. doi: 10.1080/23744235.2020.1806353.

6. Motta I, Centis R, D'Ambrosio L, et al. TB, COVID-19 and migrants: preliminary analysis of deaths occurring in 69 patients from two cohorts. Pulmonology. 2020. https://doi.org/10.1016/j.pulmoe.2020.05.002

7. He G, Wu J, Shi J, et al. COVID-19 in TB patients: A report of three cases. 2020. J Med Virol, 92: 1802-1806. https://doi.org/10.1002/jmv.25943Tadolini M, Codecasa LR, García-García Jé-Mía, et al. Active TB, sequelae and COVID-19 co-infection: first cohort of 49 cases. Eur Respir J 2020; in press (https://doi.org/10.1183/13993003.01398-2020).

8. Tadolini M, Codecasa LR, García-García Jé-Mía, et al. Active TB, sequelae and COVID-19 co-infection: first cohort of 49 cases. Eur Respir J 2020; in press (https://doi.org/10.1183/13993003.01398-2020).

9. Stochino C, Villa S, Zucchi P, et al. Clinical characteristics of COVID-19 and active TB co-infection in an Italian reference hospital. Eur Respir J. 2020 Jul 30;56(1):2001708. doi: 10.1183/13993003.01708-2020.

10. Aggarwal AN, Agarwal R, Dhooria S, Prasad KT, Sehgal IS, Muthu V (2021) Active pulmonary tuberculosis and coronavirus disease 2019: A systematic review and meta-analysis. PLoS ONE 16(10): e0259006. https://doi.org/10.1371/journal.pone.0259006

11. Petrone L, Petruccioli E, Vanini V, Cuzzi G, Gualano G, Vittozzi P, et al. Coinfection of tuberculosis and COVID-19 limits the ability to in vitro respond to SARS-CoV-2. Int J Infect Dis. 2021:In press. Epub 2021/03/14. https://doi.org/10.1016/j.ijid.2021.02.090

12. Singh A, Gupta A, Das K. Severe acute respiratory syndrome Coronavirus-2 and pulmonary TB coinfection: double trouble. Indian J. Med. Spec. 2020. DOI: 10.4103/INJMS.INJMS_72_20
13. Zumla A, Marais BJ, McHugh TD, et al. COVID-19 and TB-threats and opportunities. Int J Tuberc Lung Dis. 2020 Aug 1;24(8):757-760. doi: 10.5588/ijtld.20.0387.

14. Ariffin F, Ahmad Zubaidi AZ, Md Yasin M, et al. Management of pulmonary TB in health clinics in the Gombak district: How are we doing so far? Malays Fam Physician. 2015 Apr 30;10(1):26-33. PMID: 26425292; PMCID: PMC4567890.

15. Kashyap S, Solanki A. Challenges in endobronchial TB: from diagnosis to management. Pulm Med. 2014;2014:594806. doi: 10.1155/2014/594806.

16. Shahzad T, Irfan M. Endobronchial TB-a review. J Thorac Dis. 2016 Dec;8(12):3797-3802. doi: 10.21037/jtd.2016.12.73.

17. Jacobi A, Chung M, Bernheim A, et al. Portable chest X-ray in coronavirus disease-19 (COVID-19): A pictorial review. Clin Imaging2020;64:35-42. doi:10.1016/j.clinimag.2020.04.001 pmid:32302927

18. Tianhe Ye, Yanqing Fan, Jiacheng Liu et al. Follow-up Chest CT findings from discharged patients with severe COVID-19: an 83-day observational study, 12 May 2020, PREPRINT (Version 1) available at Research Square [https://doi.org/10.21203/rs.3.rs-27359/v1]

19. Pan F, Ye T, Sun P, et al. Time Course of Lung Changes at Chest CT during Recovery from Coronavirus Disease 2019 (COVID-19). Radiology. 2020 Jun;295(3):715-721. doi: 10.1148/radiol.2020200370.

20. Huang Y, Wang S, Liu Y, et al. A preliminary study on the ultrasonic manifestations of peripulmonary lesions of non-critical novel coronavirus pneumonia (COVID-19) (February 26, 2020). Available at: https://doi.org/10.2139/ssrn.3544750

21. Yanardag H, Tetikkurt C, Tetikkurt S, et al. Computed tomography and bronchoscopy in endobronchial TB. Can Respir J. 2003 Nov-Dec;10(8):445-8. doi: 10.1155/2003/496296.

22. Patil S, Jadhav A. Short Course of High-dose Steroids for Anaphylaxis Caused Flare Up of Tuberculosis: A Case Report. J Transl Int Med. 2019 Mar 29;7(1):39-42. doi: 10.2478/jtim-2019-0008.

23. Tham S, Lim W, Lee C, et al. Four Patients with COVID-19 and TB, Singapore, April–May 2020. Emerging Infectious Diseases. 2020;26(11):2763-2765. doi:10.3201/eid2611.2002752.

Figure Legends
Figure 1. Chest radiograph of patient 1 on admission demonstrates nodular and air space opacities bilaterally (Panel A). These opacities significantly resolve on day 12 of admission (Panel B). No cavity, enlarged hilar node, or pleural effusion is seen in both chest radiographs.
Figure 2. Representative slices (axial view) of CT thorax for patient 1 demonstrates ground-glass changes in both upper lobes with associated nodular changes bilaterally on day 7 of admission (Panel A), followed by resolution of the ground-glass changes, leaving residual nodular changes in both lung fields on day 40 of admission (Panel B).
Figure 3. Chest radiograph of patient 2 on admission demonstrates right pleural effusion with minimal air space opacities at bilateral lower zones (Panel A) which progresses to involve the mid zone bilaterally on day 12 of admission (Panel B). There is no significant enlarged hilar node bilaterally.

Figure 4. Chest radiograph of patient 3 on admission demonstrates predominantly nodular with some air space opacities bilaterally (Panel A). Some nodules coalesce to form a larger patch of consolidation at the left midzone. No cavity, air-fluid level, effusion or enlarged hilar node is seen. Repeated chest radiograph at day 12 of admission shows partial resolution of these changes (Panel B).