Chest CT findings of influenza virus-associated pneumonia in 12 adult patients

Jiro Fujita, a Shuji Bandoh, b Masahiro Yamaguchi, c Futoshi Higa, a Masao Tateyama a

a Department of Medicine and Therapeutics, Control and Prevention of Infectious Diseases (First Department of Internal Medicine), Faculty of Medicine, University of the Ryukyus, Okinawa, Japan. b Faculty of Medicine, Kagawa University, Kagawa, Japan. c Takamatsu Higashi Hospital, Kagawa, Japan.

Correspondence: Jiro Fujita, MD, PhD, Department of Medicine and Therapeutics, Control and Prevention of Infectious Diseases (First Department of Internal Medicine), Faculty of Medicine, University of the Ryukyus, 207 Uehara, Nishihara-cho, Okinawa 903-0215, Japan. Email: fujita@med.u-ryukyu.ac.jp

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Objective In this study, we describe the chest computed tomography findings of influenza virus-associated pneumonia in adult patients.

Methods Our retrospective study included 12 adult patients who had proven influenza virus-associated pneumonia.

Results Out of 12 patients, six were diagnosed as having pure influenza virus pneumonia, five as having bronchopneumonia caused by bacteria associated with influenza A infection, and one as having a cryptogenic organizing pneumonia associated with influenza A infection.

Conclusion Radiographic findings of influenza virus pneumonia in adult patients consist of ground-glass attenuation. Localized patchy consolidations were observed in cases of bronchopneumonia.

Key words Chest CT, influenza virus, pneumonia.

Introduction

The influenza virus is a common cause of lower respiratory tract infections in adults. It can occur during winter outbreaks and particularly during pandemics. Clinical pneumonia attributable to the influenza virus is uncommon, but when it does occur, secondary bacterial infection as well as the influenza A virus itself need to be considered as the possible cause.1 Influenza A virus pneumonia is usually mild, but it can be overwhelming in some patients and prove fatal within 24 hours after onset.2 Older patients and patients with underlying heart disease, chronic bronchitis or cystic fibrosis have an increased risk of developing pneumonia.3 Although isolated cases of influenza pneumonia in immunocompromised patients have been reported,4,5 surprisingly little is known about the radiologic manifestations and clinical pictures of influenza virus-associated pneumonia in adult patients6,7 or avian influenza.8,9

The aim of our study was to describe the radiographic and chest computed tomography (CT) findings of 12 adult patients with confirmed influenza virus-associated pneumonia.

Materials and methods

Patients

Our retrospective study included 12 adult patients who had proven influenza virus-associated pneumonia. The influenza virus infections were diagnosed using a rapid diagnostic kit between December 2003 and December 2006. All cases experienced an acute onset of high fever and symptoms compatible with an influenza virus infection. Influenza virus-associated pneumonia was diagnosed as having a positive influenza antigen result using a commercially available diagnostic kit, detection of infiltration by chest X-ray and chest CT, and the appropriate exclusion of other diagnoses.

The 12 patients in our study population were seven women and five men whose age ranged from 30 to 91 years (mean, 38 years; median, 32 years). Although some patients were very old, there was no pre-existing lung disease in those patients. However, case 7 and case 9 had severe bronchial asthma (Table 1).

Chest radiographs and chest CT scans of the chest were obtained for all patients. As all patients had a serious illness, chest CT scans were immediately performed for each...
Influenza virus-associated pneumonia was defined as the presence of infiltrates in chest radiographs and chest CT scans in patients with clinically proven influenza-virus infection confirmed by the rapid diagnosis kit.

Assessment of chest CT findings
Two observers reviewed the CT scans and reached a consensus decision about the pattern and distribution of the findings. When evaluating chest CTs, all clinical information was blinded before evaluating patterns of infiltration. Chest CT scans were assessed for the presence of ground-glass opacifications, consolidation, as well as large and small nodules. The presence of zonal (upper, middle, or lower and central or peripheral) and lateral (unilateral or bilateral) predominance was also assessed. Consolidation was defined as an area of opacification that obscured the underlying vessels; in contrast, ground-glass opacity was defined as a hazy increase in lung attenuation with no obscuration of underlying vessels.

With regard to evaluating patterns in the radiologic findings, it was very difficult to classify patterns in the chest X-rays, especially for patients with pleural effusion. Therefore, we used only chest CTs to assess patterns in the radiologic findings. Based on the two observers’ opinions, the patterns of chest CT scans were categorized as follows: interstitial pneumonia (mainly ground-glass opacity), bronchopneumonia (mainly patchy opacification) and a type compatible with cryptogenic organizing pneumonia (COP; only in this category did observers consider histological findings and steroid-responsiveness). If a lobular consolidation was observed, we considered that type of pneumonia to be bronchopneumonia. Lobular consolidation was considered present when the consolidation involved the entire secondary lobule but spared the adjacent lobules. If a ground-glass attenuation was mainly observed, we considered that type of pneumonia to be interstitial pneumonia.

Results
Seven of the twelve patients had other organisms identified in sputum or blood cultures (Table 1). In patients with

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Table 1. Patient characteristics

| Case | Age & sex | Background | Type | CRP/WBC | Sputum culture | Blood culture | Bacterial infection | Chest CT pattern | Therapy | Prognosis |
|------|-----------|------------|------|---------|----------------|--------------|-------------------|-----------------|---------|-----------|
| 1    | 70 M      | Steroid    | A    | ND      | –             | ND           | –                 | Interstitial pneumonia | Oseltamivir       | Alive     |
| 2    | 86 M      | Diabetes   | A    | 117/11 900 | –             | ND           | –                 | Interstitial pneumonia | Amantadine       | Died*     |
| 3    | 68 M      | Parkinson  | A    | 21/4100 | +             | Pseudomonas aeruginosa, MRSA| ND           | Interstitial pneumonia | Amantadine       | Alive     |
| 4    | 40 F      | Asthma     | A    | 147/12 700 | +             | ND           | –                 | Interstitial pneumonia | Oseltamivir + antibiotics + steroid | Alive     |
| 5    | 77 M      | CHD        | A    | 9.5/8200 | +             | –             | –                 | Interstitial pneumonia + effusion | Oseltamivir + antibiotics | Alive     |
| 6    | 79 M      | CHD        | A    | 154/6600 | +             | –             | –                 | Interstitial pneumonia + effusion | Oseltamivir + antibiotics | Died*     |
| 7    | 91 F      | Diabetes, hypertension | A | 4.5/8900 | –             | Streptococcus pneumoniae | – +            | Bronchopneumonia | Oseltamivir + antibiotics | Alive     |
| 8    | 41 F      | Diabetes   | A    | 118/26 200 | +             | MSSA          | MSSA +            | Bronchopneumonia | Oseltamivir + antibiotics | Alive     |
| 9    | 60 F      | Asthma     | A    | 184/16 300 | –             | MRSA + MSSA   | – +            | Bronchopneumonia | Oseltamivir + antibiotics | Alive     |
| 10   | 30 F      | –          | A    | 4.49/5000 | –             | Streptococcus pneumoniae | ND +            | Bronchopneumonia | Oseltamivir + antibiotics | Alive     |
| 11   | 82 F      | Diabetes   | A    | 2.46/5300 | –             | Gram stain (+), not identified | ND +            | Bronchopneumonia | Oseltamivir + antibiotics | Alive     |
| 12   | 38 F      | –          | A    | 8.85/14 300 | –             | ND           | –                 | Cryptogenic organizing pneumonia | Zanamivir | Alive     |

CHD, coronary heart disease; ND, not done; MRSA, methicillin-resistant Staphylococcus aureus; MSSA, methicillin-sensitive Staphylococcus aureus.

*Died of respiratory failure.
superimposed bacterial pneumonia, purulent sputum was observed.

Based on chest CT findings, six of the twelve patients were categorized as having pure influenza pneumonia (principally the interstitial pattern). Of these six patients, four had no pleural effusion (Figure 1), and two patients had pleural effusion (Figure 2). In these six patients, ground-glass opacities were the predominant finding; in five of the six, the opacities were bilateral and the opacities were unilateral in one patient. The ground-glass opacities showed upper lobe predominance in all six patients. *Pseudomonas aeruginosa* and methicillin-resistant *Staphylococcus aureus* (MRSA) were cultured in one patient with interstitial pneumonia. In this patient, the detection of *P. aeruginosa* and MRSA reflects colonization by these bacteria. As phagocytized bacteria were not detected by Gram stain of sputum from the patient we judged that these bacteria were not significantly important as causative pathogens. No bacteria were cultured in the remaining five patients who were categorized as having interstitial pneumonia.

Five of the twelve patients were categorized as having bronchopneumonia. Of the five patients categorized as having bronchopneumonia, bacteria were cultured from the sputum specimens of four patients: two showing *S. aureus* and two showing *Streptococcus pneumoniae*. In the remaining patient who was categorized as having bronchopneumonia, no bacteria were cultured even though Gram-positive rods as well as Gram-negative bacilli were observed in the sputum.

All patients were treated by oral oseltamivir. In addition, patients with superimposed bacterial infection and a patient with COP were treated by broad-spectrum antibiotics, which was then followed by deescalated antibiotic therapy based on data from the drug sensitivity test.

The chest CT findings for those categorized as having bronchopneumonia are shown in Figure 3. In all five patients, patchy consolidations are observable, and pneumothorax is complicated in one patient who had bronchopneumonia caused by *S. aureus* (Figure 3c).

One patient was categorized as having a pattern of COP, and had a biopsy that was histologically compatible with COP (Figure 4). In this case, our initial diagnosis was bacterial pneumonia. However, purulent sputum was not observed. In addition, although broad-spectrum antibiotics were administered, there was no improvement in the radiologic findings. Furthermore, as there have been a few case reports describing COP associated with influenza viral
infection, we decided to perform a lung biopsy. Steroid pulse therapy was very effective with this patient. None of the patients had mediastinal lymphadenopathy.

Discussion

Influenza is classified into three types (A, B or C) according to the nucleoprotein antigens and matrix protein components. Consistent with the present study, pure influenza pneumonia is usually reported in association with type A influenza viruses. Although fatal pneumonia can result from the primary viral infection, it has been reported that most deaths are related to bacterial superinfections.10

The radiographic findings of influenza virus pneumonia in immunocompetent patients have rarely been reported. Kim et al.6 evaluated the high-resolution CT findings of influenza virus pneumonia in two immunocompetent patients and reported that both lungs had areas of multifocal peribronchovascular or subpleural consolidation. For one patient, the chest CT scan shows multifocal peribronchiolar consolidation and ground-glass attenuation in both lungs. Some lesions have a lobular distribution. The other patient showed diffuse ground-glass opacities with irregular linear areas of increased attenuation as shown in our patients diagnosed with pure influenza pneumonia. In addition, Tanaka et al.7 reported the high-resolution CT findings of influenza virus pneumonia in a immunocompetent patient as consisting of bilateral areas of ground-glass attenuation with a lobular distribution.

Information about the radiographic findings of influenza pneumonia in immunocompromised patients is also limited. Leung et al.11 in their study of 59 cases of pulmonary infection in bone marrow transplant recipients included one case of influenza virus B pneumonia, a patient whose chest radiograph showed 20–30 nodules with diameters of 6–10 mm in the middle and lower zones of both lungs with no other associated findings. In a retrospective study of the radiographic findings of viral infections in 21 lung transplant recipients, the authors reported that one patient with influenza virus pneumonia showed bilateral homogeneous opacities that progressed to show patterns associated with acute respiratory distress syndrome, whereas the other patients had only minimal abnormalities.12 In addition, Oikonomou et al. reported that the radiographic findings consisted mainly of unilateral or bilateral patchy areas of consolidation with or without associated poorly defined nodular opacities.13

As influenza infection complicated by pneumonia is not very frequent, there may be some selection bias in our study. However, we checked every patient with an influenza virus infection which was diagnosed both clinically and by the detection of influenza antigen, and we selected those patients with chest X-ray abnormalities. In patients with
superimposed bacterial pneumonia, purulent sputum was observed. In addition, information obtained from Gram stain was also helpful in evaluating bacterial superinfection. In the present study, the chest CT patterns of 12 patients with influenza virus-associated pneumonia were demonstrated. Importantly, chest CT patterns, especially lobular consolidations, were very helpful in diagnosing bacterial-superimposed bacterial infections. In addition, diffuse ground-glass attenuation was frequently observed in pure influenza-virus pneumonia. As the treatment strategy for influenza virus-associated pneumonia is different in the presence and absence of a superimposed bacterial infection, evaluation of radiologic patterns by chest CT scans seemed to be clinically important. Furthermore, the clinician should be aware of the existence of COP associated with influenza virus infections in order to choose the appropriate treatment by corticosteroids.

Our study has two major limitations. First, the small number of patients does not allow us to make generalized statements about the range of potential abnormalities. Second, no correlation with pathological findings was possible because it was a retrospective study. However, we believe that physicians will find these chest CT findings very useful.

In summary, several patterns of chest CT findings of influenza A virus-associated pneumonia were demonstrated.

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