Research article

Chest imaging of H7N9 subtype of human avian influenza

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Abstract

Background: Human infection with avian influenza A H7N9 virus is an acute respiratory infectious disease, which usually causes severe pneumonia with a high mortality. Chest radiographs and Computed Tomography (CT) are principal radiological modalities to assess the lung abnormalities.

Objectives: The goal of this study was to investigate the chest images characteristic of H7N9 subtype of human avian influenza.

Materials and methods: The clinical and imaging data of 11 cases diagnosed as H7N9 subtype of human avian influenza were collected from 4 cities in the southern region of the Yangtze River, China. The chest imaging manifestations were analyzed by the assigned expert group. The analyzed cases include 7 males and 4 females aged from 20 to 84 years, with a mean of 55.6 years. The clinical symptoms were mainly fever (100%, 11/11) and cough (72.7%, 8/11).

Results: Segmental or lobar ground-glass opacity (GGO) or consolidation was shown in 8 cases (72.7% or 8/11). Air bronchogram was found in 7 cases (63.6% or 7/11). The lesions developed into multiple or diffuse in both lungs rapidly at the progressive stage. The reticulation shadows were shown after some lesions absorbed at the stable stage.

Conclusions: The characteristic imaging demonstrations of H7N9 subtype of human avian influenza are segmental or lobar exudative lesions at lungs at the initial stage, which rapidly progress into bilateral distribution at lungs at the progressive stage.

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Keywords: Human avian influenza; H7N9; Radiography; Tomography; Computed X-ray

1. Introduction

Human infection with avian influenza A H7N9 virus is an acute respiratory infectious disease. It firstly emerged in Anhui Province and Shanghai City, P.R. China in February, 2013 [1], and more cases were then reported in the Eastern China [2]. The disease is characterized by acute onset and high mortality, which poses a great threat to human health. In this paper, we
retrospectively reviewed chest radiographs and Computed Tomography (CT) findings in 11 patients that had confirmatively been diagnosed with H7N9 subtype of human avian influenza.

2. Materials and methods

2.1. Subjects

This study retrospectively reviewed the clinical and imaging data of 11 patients with H7N9 subtype of human avian influenza (Table 1) from 4 cities in the southern region of the Yangtze River, China. The diagnostic criteria of human infection with avian influenza A H7N9 virus was established by the National Health and Family Planning Commission of China (formerly Ministry of Health). The results of real-time reverse transcriptase polymerase chain reaction (RT-PCR) of nasal swabs and aspirates of all the 11 patients were positive for H7N9 avian influenza virus. All patients were approved by the expert board of Provincial Department of Health in Jiangsu or Zhejiang, China.

The study subjects included 7 males and 4 females with their ages ranging from 20 to 84 years and a mean of 55.6 years. All 11 patients had initially presented with influenza-like symptoms, including fever (100%, 11/11), cough (72.7%, 8/11), expectoration (45.5%, 5/11), chest tightness (18.2%, 2/11), muscular pain (18.2%, 2/11), shortness of breath (9.1%, 1/11), fatigue (9.1%, 1/11), sore throat (9.1%, 1/11) and vomiting (9.1%, 1/11). Seven of the 11 subjects had the following medical histories: thoracic surgery with excision of thymoma (n = 1), pregnancy (n = 1), hypertension (n = 3) and chronic bronchitis (n = 2). The other 4 patients had no significant medical history. The initial WBC count was normal in 7 patients, low in 3 patients and high in 1 patient.

After onset of the illness, all 11 patients were hospitalized for 1–10 days and received antiviral and respiratory support therapies.

2.2. Imaging technologies

Posteroanterior radiographs were obtained using Axiom Aristos (Siemens Healthcare) with a constant technique of 125Kv and automatic mA adjustment. MDCT was performed on 64-MDCT scanner (Somatom Sensation, Siemens healthcare), 128-MDCT scanner (Somatom Definition, Siemens healthcare) and 4-MDCT scanner (Brightspeed excel, GE healthcare). The scan protocol was as the following: 120 kV, automatic mA adjustment (about 115mAs), pitch of 0.9, matrix of 512 × 512, thickness of 5.0–6.0 mm, interval of 1.4 mm, and 0.75 mm reformation.

All 11 patients had serial bedside anteroposterior-projection follow-up radiographs at an interval of 2 or 3 days, while only 7 patients required follow-up CT scans. Bedside anteroposterior-projection radiographs were obtained with a mobile unit (Philips Healthcare), using a standard exposure factor of 80 kV and 5 mAs.

2.3. Image analysis

The chest radiographs and MDCT images were independently reviewed by two experienced radiologists and final interpretations were achieved by consensus. The anatomic distribution was characterized as either unilateral or bilateral. The extent of abnormality was graded as focal, multifocal and diffuse. The abnormalities were characterized as consolidation (opacity with obscuration of the underlying vessels), ground-glass opacity (GGO) which increased attenuation without obscuration of the underlying vessels, nodules opacities and reticulation. The presence of enlarged lymph nodes and pleural effusions were also recorded.

3. Results

3.1. Initial image findings

The initial chest radiographs were abnormal in all 11 patients (Table 2) (Fig. 1A). The lung abnormalities were unilateral in 45.5% or 5/11 patients (4/5 in left lung and 1/5 in right lung) and bilateral in the rest of 6 patients. The abnormalities were found in one lobe in 4 patients, two lobes in 2 patients, three lobes in 2 patients and four lobes in 3 patients, respectively.

<table>
<thead>
<tr>
<th>Case</th>
<th>Gender</th>
<th>Age (year)</th>
<th>Onset symptoms</th>
<th>Temperature (°C)</th>
<th>WBC (× 10⁹/L)</th>
<th>Interval between initial imaging and onset(d)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>F</td>
<td>32</td>
<td>Fever, cough, sputum</td>
<td>39.6</td>
<td>2.0</td>
<td>4</td>
</tr>
<tr>
<td>2</td>
<td>M</td>
<td>72</td>
<td>Fever, vomiting</td>
<td>39</td>
<td>6.2</td>
<td>3</td>
</tr>
<tr>
<td>3</td>
<td>F</td>
<td>20</td>
<td>Fever, cough, sputum</td>
<td>39</td>
<td>3.8</td>
<td>1</td>
</tr>
<tr>
<td>4</td>
<td>M</td>
<td>84</td>
<td>Fever, cough, sputum</td>
<td>39</td>
<td>5.27</td>
<td>6</td>
</tr>
<tr>
<td>5</td>
<td>M</td>
<td>72</td>
<td>Fever, cough, chest tightness, fatigue</td>
<td>39.4</td>
<td>7.42</td>
<td>7</td>
</tr>
<tr>
<td>6</td>
<td>M</td>
<td>60</td>
<td>Fever, chest tightness, shortness of breath, pharyngalgia</td>
<td>39.6</td>
<td>4.1</td>
<td>10</td>
</tr>
<tr>
<td>7</td>
<td>M</td>
<td>56</td>
<td>Fever, muscle ache</td>
<td>39.5</td>
<td>11.19</td>
<td>6</td>
</tr>
<tr>
<td>8</td>
<td>M</td>
<td>70</td>
<td>Fever, cough</td>
<td>38.6</td>
<td>4.7</td>
<td>9</td>
</tr>
<tr>
<td>9</td>
<td>F</td>
<td>25</td>
<td>Fever, cough, sputum, muscle ache</td>
<td>39.9</td>
<td>7.9</td>
<td>1</td>
</tr>
<tr>
<td>10</td>
<td>M</td>
<td>65</td>
<td>Fever, cough</td>
<td>38.6</td>
<td>7.0</td>
<td>4</td>
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<tr>
<td>11</td>
<td>F</td>
<td>56</td>
<td>Fever, cough, sputum</td>
<td>39.4</td>
<td>2.6</td>
<td>4</td>
</tr>
</tbody>
</table>

Table 1
Summary of the clinical data and initial examination of the 11 patients with H7N9 subtype of human avian influenza.
The predominant chest imaging finding was segmental or lobar consolidation (72.7% or 8/11), among which three (3/8) showed multi-nodular opacities (Fig. 2A), and two (2/8) presented with combined segmental ground-glass opacity. Segmental ground-glass opacity was demonstrated in 2 patients (18.2% or 2/11). Multi-patchy shadows presented in 1 patient (9.1% or 1/11).

Air bronchogram was a commonly seen lung abnormality (63.6% or 7/11). A sharp boundary due to the interlobar fissure was also manifested (36.4% or 4/11).

Thickened adjacent pleura could be seen in four patients and pleural effusions were found in two patients. Lymph node enlargement was not significant in any of the eleven cases.

3.2. Follow-up imaging findings

During 3—15 days after onset, eight patients showed obvious progress of the conditions compared to initial abnormalities (Fig. 1B). Firstly, lesions increased in size, progress from unilateral to bilateral lungs or diffuse distribution (Fig. 2B). Secondly, density of abnormalities developed from GGO to consolidation. The patients at this stage were characterized as acute respiratory distress syndrome (ARDS), with occurrence of death in 2 cases and no obvious change in other 3 cases.

After 15 days, three of the rest 9 patients (3/9) were improved compared with the former lesions (Fig. 2C—2D). The abnormalities demonstrated as reticulation with a decreased extent. One of these three patients had segmental consolidation in the left upper lobe and segmental consolidation in bilateral lower lobes at the initial scanning, but was improved afterwards. One patient (1/9) who had previously undergone thoracic surgery for excision of thymoma showed segmental consolidation in the left upper lobe by the initial radiograph and CT scan seven days after onset (Fig. 3A), and diffuse GGO and consolidation at both lungs by the bedside radiograph ten days after onset (Fig. 3B). The extent and density of the abnormalities remained unchanged during the following twenty days (Fig. 3C—3D). The extent of abnormalities of the other five patients also remained unchanged but with invariable or slightly deteriorated clinical conditions. Death occurred in another 4 cases at this stage.

4. Discussion

On March 2013, a subtype of avian influenza virus causing human infections, H7N9, was identified in Eastern China. H7 subtypes (H7N2, H7N3 and H7N7) of avian influenza viruses are commonly found and are often limited to mild illness, among which only one case of death was reported in...
To our knowledge, this is the first time that H7N9 subtype has infected human and caused death. Preliminary studies have shown that the H7N9 subtype of avian influenza virus is novel reassortants and more virulent in humans than other H7 viruses [4,5].

Infection of H7N9 subtype of avian influenza virus typically manifests itself as flulike symptoms such as fever, cough, sore throat, headache and body aches [1,6]. The clinical findings of the 11 patients from our group that had been confirmatively diagnosed as infection of H7N9 subtype of avian influenza virus reveal that rapid progress is characteristic of this disease. A high incidence of severe pneumonia is a common abnormality, with all eleven patients suffering from persistent fever (body temperature of 39 °C or above), and accompanying dyspnea or ARDS. WBC count may be normal or slightly decrease [2].

In our study, the pulmonary lesions of H7N9 infection of human avian influenza were radiologically demonstrated to be segmental or lobar GGO or consolidation combined with air bronchogram at the initial stage without a predominant distribution, which are inconsistent with findings from previous studies [7]. We believe that the pathological mechanism of infection of H7N9 subtype of avian influenza virus may resemble to other viral pneumonia and is based on alveolar exudation. The lesions can develop into either multiple or diffuse at both lungs with rapid progress at the progressive stage, or reticulation when the abnormalities improve later into the later stage (after 20 days) [5,6]. According to the evolution process by radiology, infection of H7N9 subtype of avian influenza virus can be divided into three stages. The initial stage is the first 3 days after onset. The progressive stage is the period of d 3-15 after onset. Period since d 15 after onset is defined as the stable stage. The stable stage is protracted for some cases due to the remaining viral infection in addition with bacterial infections. Pleural effusions are rarely seen and lymph node enlargement is not significant.

The imaging demonstrations of H7N9 subtype infection of avian influenza virus may resemble to other pneumonia including novel swine influenza A (H1N1) virus (S-OIV) infection, severe acute respiratory syndromes (SARS), bacterial pneumonia and highly pathogenic H5N1 subtype human avian influenza virus infection. The main imaging findings of S-OIV infection and SARS are unilateral or bilateral GGO or focal areas of consolidations with a predominant peribronchovascular and subpleural distribution [8–10], while H7N9 subtype avian influenza infection presents pulmonary segment or lobar exudative lesions without obvious distinctive distribution. Bacterial pneumonia usually shows lobar consolidation and increased WBC clinically, which can recover quickly by antibiotic therapy. The imaging findings of H7N9 subtype infection of avian influenza virus resemble to those of H5N1 subtype infection of avian influenza virus, both of which have pulmonary segment or lobar exudative lesions as the predominant imaging findings except that multifocal consolidations have a predilection at lower lobes in cases of H5N1 subtype infection of avian influenza virus [11].
However, differential diagnosis relies on virus isolation and detection [6].

This study has several limitations. Firstly, it is a retrospective study based on a series of 11 cases. Secondly, none of the patients underwent lung biopsy or autopsy that allowed radiographic—histopathologic correlation.

In summary, the imaging findings of H7N9 subtype infection of avian influenza virus in this series of patients tends to be pulmonary segment or lobar consolidation and mostly progress rapidly into diffuse exudation.

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