Hospitalized adult patients with 2009 influenza A (H1N1) in Beijing, China: risk factors for hospital mortality

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Abstract

Background: In April 2009, the pandemic influenza A(H1N1) virus emerged and spread globally. The objective of this study was to describe the independent risk factors for hospital mortality and the treatment effect of corticosteroids among patients with 2009 influenza A(H1N1) infection.

Methods: We retrospectively obtained clinical data of 155 adult patients with confirmed infection of 2009 influenza A(H1N1) in 23 hospitals in Beijing, China from October 1 to December 23, 2009. Risk factors for hospital mortality were identified with multivariate logistic regression analysis.

Results: Among the 155 patients, 90 (58.1%) were male, and mean age was 43.0 ± 18.6 years, and comorbidities were present in 81 (52.3%) patients. The most common organ dysfunctions included acute respiratory failure, altered mental status, septic shock, and acute renal failure. Oseltamivir was initiated in 125 patients (80.6%), only 16 patients received antiviral therapy within 48 hours after symptom onset. Fifty-two patients (33.5%) were treated with systemic corticosteroids, with a median daily dose of 80 mg. Twenty-seven patients (17.4%) died during hospital stay. Diabetes [odds ratio (OR) 8.830, 95% confidence interval [CI] 2.041 to 38.201, p = 0.004) and lactate dehydrogenase (LDH) level (OR 1.240, 95% CI 1.025 to 1.500, p = 0.027) were independent risk factors of hospital death, as were septic shock and altered mental status. Corticosteroids use was associated with a trend toward higher hospital mortality (OR 3.668, 95% CI 0.987 to 13.640, p = 0.052).

Conclusions: Hospitalized patients with 2009 H1N1 influenza had relative poor outcome. The risk factors at hospitalization may help clinicians to identify the high-risk patients. In addition, corticosteroids use should not be regarded as routine pharmacologic therapy.
corticosteroids treatment as well as its effect on hospital mortality were also evaluated.

**Methods**

The first case of 2009 influenza A(H1N1) in mainland China was identified on May 11, 2009 [8]. As a result, all hospitals were required to report every case to local health authorities and Center for Disease Control (CDC). In this retrospective study, cases were captured through the records in Beijing Health Bureau and Beijing CDC. All hospitals with H1N1 patients were identified and contacted for possible participation into the study. Twenty-three hospitals provided positive feedback, and constituted the study group.

Patients were eligible if they were (1) ≥ 18 years old; (2) admitted to any of the 23 participating hospitals from October 1 to December 23, 2009; (3) diagnosed as confirmed 2009 influenza A(H1N1) infection, according to case definitions developed by the World Health Organization [11]. Specifically, a confirmed case was defined by a positive result of a real-time reverse-transcriptase-polymerase-chain-reaction (RT-PCR) assay performed at a laboratory operated under the auspices of the Chinese CDC [8].

Case report form was initially developed by one investigator (BD), and then cycled among all investigators for feedback and pilot testing until consensus was reached. All reviewers were trained during a 1.5-hour course before the study, and then dispatched to all participating hospitals to review case records of all eligible cases.

For all patients, the following data were recorded: sex, age, major diagnosis, comorbidities, clinical presentation at symptom onset and on hospital admission, major laboratory results on hospital admission, complications, antiviral and antibiotic treatment, supportive treatment, and outcome. In particular, septic shock was defined according to the consensus definition of American College of Chest Physicians/Society of Critical Care Medicine [12], while acute respiratory failure or acute renal failure was defined if sequential organ failure assessment score for that particular organ was greater than two points [13].

Data were entered into a Microsoft Excel database (Microsoft, Seattle, Wash., USA) by a data manager (BD) under the supervision of the study steering committee (XM, YX, LJ, AL). Data were checked for inconsistencies and logical errors on entry, and queries were sent to the source hospital for resolution. The study protocol was approved by Institutional Review Board of Beijing Fuxing Hospital (IRB-2009-0135), and the need for informed consent was waived because of the retrospective study design.

Values are presented as the mean ± standard deviation or median (interquartile range) when appropriate (continuous variables), or as a percentage of the group from which they were derived (categorical variables). Continuous variables were compared with the use of the Student’s t-test or Mann-Whitney test. The chi-square test or Fisher’s exact test was used to compare categorical variables. For determination of independent predictors for hospital mortality, odds ratio was estimated on the basis of multivariate logistic regression analysis, including exploration for robustness and interactions using likelihood ratio tests. Two separate models using stepwise conditional forward entry were constructed - one for demography, comorbidities, clinical presentations, and laboratory tests on admission, and one for complications including organ failures and infections, if p < 0.1 in univariate analysis. Corticosteroids treatment was forced into the first model in order to examine the effect on patient outcome. Multivariate logistic regression analysis was also used to determine the independent predictors of corticosteroids treatment.

The Kaplan-Meier method was used to determine the probability of survival over the duration of follow-up and to generate survival curves, censoring at 30 days after symptom onset. All comparisons were unpaired and all tests of significance were two-tailed. A p value < 0.05 was considered as statistically significant.

**Results**

**Characteristics of study population**

From October 1 to December 23, 2009, 562 patients with 2009 influenza A(H1N1) infection who required hospitalization were reported to Beijing Health Bureau, among whom 64 patients died. Of all these patients, 287 patients were admitted into any one of the 23 participating hospitals, accounting for 51.1% of all reported cases. After excluding 132 paediatric patients, 155 adult patients were included in data analysis.

Among the 155 adult patients with confirmed 2009 influenza A(H1N1) infection, 90 (58.1%) were male, and mean age was 43.0 ± 18.6 years. Eighty-one patients (52.3%) had at least one comorbidity, including cardiovascular disease, chronic lung disease, diabetes, cerebral vascular disease, and malignancy. In particular, 12 patients were pregnant women, and 2 patients were postpartum women (Table 1).

The most common presenting symptom was fever (> 37.8°C) (93.5%), followed by cough (91.6%), dyspnea (47.1%), sore throat (29.7%), and myalgia (17.4%). The median time from symptom onset to hospital admission was 5 days. On hospital admission, 82 patients (52.9%) still complained dyspnea, and 129 patients (83.2%) were diagnosed as pneumonia according to chest X-ray. Forty of 141 patients (28.4%) who were tested had leukopenia (white cell count < 4.0 × 10⁹/L), while 19 patients
had leukocytosis (white cell count > 10.0 x 10^9/L), and 53 of 125 patients (42.4%) had lymphopenia (lymphocyte count < 0.8 x 10^9/L). Laboratory tests also suggested mild to moderately increased cardiac enzymes, liver enzymes, and total bilirubin (Table 1). Of 126 patients who were tested, 74 patients (58.7%) had elevated level of lactate dehydrogenase (LDH) (> 270 U/L). Alanine transaminase and aspartate transaminase were elevated in 32 of 111 (28.8%) and 69 of 115 (60.0%) patients, respectively.
Treatment and course of illness

Of all patients, 125 (80.6%) were treated with oseltamivir, although only 16 patients (10.3%) received antiviral therapy within 48 hours after symptom onset (Table 2). Broad-spectrum antibiotics were prescribed in 139 (89.7%) patients as empiric or definitive therapy.

The most common organ failures during hospital stay included acute respiratory failure (40.0%), altered mental status (12.9%), septic shock (11.6%), and acute renal failure (9.7%). Forty-three patients (27.7%) were treated with invasive mechanical ventilation, while noninvasive mechanical ventilation was used in 32 patients (20.6%). Vasopressors and inotropes were used in 28 (18.1%) and 9 (5.8%) patients, respectively.

Fifty-two patients (33.5%) were treated with systemic corticosteroids. Daily dose of corticosteroids ranged from methylprednisolone 12 mg to 320 mg (or equivalent dose), with a median dose of 80 mg (interquartile range, 80 mg to 160 mg).

Outcome and risk factors

Twenty-seven patients (17.4%) died during hospital stay, all within 28 days after admission (Figure 1). Causes of death included multiple organ failure (n = 14), refractory circulatory shock (n = 6), refractory respiratory failure (n = 5), and others (n = 2).

Compared with survivors, more nonsurvivors had hypertension (37.0% vs. 19.5%, p = 0.048). On hospital admission, nonsurvivors were more likely to be dyspneic (77.8% vs. 47.7%, p = 0.004), and lactate dehydrogenase (LDH) level was significantly higher (744 ± 759 vs. 356 ± 240 U/L, p = 0.041) (Table 1). During hospitalization, more nonsurvivors developed organ failure (acute respiratory failure, altered mental status, septic shock, and acute renal failure) and secondary infections (Table 2). Therefore, nonsurvivors were more likely to be treated with vasoactive agents, mechanical ventilation, and continuous renal replacement therapy (Table 2).

Risk factors for hospital mortality by multivariate logistic regression analysis included diabetes (odds ratio

Table 2 Hospitalized adult patients with 2009 Influenza A (H1N1) in Beijing: complications, definitive treatment, and supportive treatment

| Complications                        | Survivor (n = 128) | Nonsurvivor (n = 27) | Total (n = 155) | P value |
|--------------------------------------|--------------------|----------------------|-----------------|---------|
| Acute respiratory failure            | 39 (30.5%)         | 23 (85.2%)           | 62 (40.0%)      | 0.003   |
| Septic shock                         | 1 (0.8%)           | 17 (63.0%)           | 18 (11.6%)      | < 0.001 |
| Acute renal failure                  | 4 (3.1%)           | 11 (40.7%)           | 15 (9.7%)       | < 0.001 |
| Altered mental status                | 6 (4.7%)           | 14 (51.9%)           | 20 (12.9%)      | < 0.001 |
| Bacterial pneumonia                  | 15 (11.7%)         | 13 (48.1%)           | 28 (18.1%)      | < 0.001 |
| Bloodstream infection                | 0                  | 3 (11.1%)            | 3 (1.9%)        | 0.006   |
| Other infections                     | 0                  | 5 (18.5%)            | 5 (3.2%)        | < 0.001 |
| Antiviral agents                     | 107 (83.6%)        | 25 (92.6%)           | 132 (85.2%)     | 0.371   |
| Oseltamivir treatment                | 101 (78.9%)        | 24 (88.9%)           | 125 (80.6%)     | 0.259   |
| within 48 hrs of symptom onset      | 12 (9.4%)          | 4 (14.8%)            | 16 (10.3%)      | 0.507   |
| Broad-spectrum antibiotics           | 113 (88.3%)        | 26 (96.3%)           | 139 (89.7%)     | 0.309   |
| Corticosteroids                      | 35 (27.4%)         | 17 (63.0%)           | 52 (33.3%)      | < 0.001 |
| Vasopressors                         | 5 (3.9%)           | 23 (85.2%)           | 28 (18.1%)      | < 0.001 |
| Inotropes                            | 2 (1.6%)           | 7 (25.9%)            | 9 (5.8%)        | < 0.001 |
| Neuromuscular blockade               | 1 (0.8%)           | 7 (25.9%)            | 8 (5.2%)        | < 0.001 |
| Noninvasive ventilation              | 19 (14.9%)         | 13 (48.1%)           | 32 (20.6%)      | < 0.001 |
| Invasive ventilation                 | 19 (14.9%)         | 24 (88.9%)           | 43 (27.7%)      | < 0.001 |
| Continuous renal replacement therapy  | 4 (3.1%)           | 13 (48.1%)           | 17 (11.0%)      | < 0.001 |
Oseltamivir treatment did not result in significant improvement in patient outcome (Table 2). Among 125 patients treated with oseltamivir, initiation of antiviral therapy within 48 hours of symptom onset showed no influence on hospital mortality (4/16 vs. 20/109, p = 0.508).

Corticosteroids treatment
Corticosteroids treatment was associated with a trend toward higher hospital mortality (OR 3.668, 95% CI 0.987 to 13.640, p = 0.052) (Table 3). However, corticosteroids were used in more critically ill patients, as shown in Table 4. LDH level on hospital admission and acute respiratory failure were independent predictors of corticosteroids treatment (Table 4). Moreover, patients treated with a lower daily dose of corticosteroids (≤ 80 mg methylprednisolone or equivalent dose) exhibited a similar mortality rate compared with those treated with higher daily dose (9/30 vs. 8/22, p = 0.854).

| Table 3 Hospitalized adult patients with 2009 Influenza A (H1N1) in Beijing: risk factor for hospital mortality by logistic regression analysis |
|---------------------------------------------------------------|
| Model 1                                                                                   |
| Diabetes                                      | OR 8.830    |
| LDH on admission, per 100 U/L increase    | 2.041 - 38.201 |
| Corticosteroids treatment                  | 3.668  |
| Model 2                                                                                   |
| Septic shock                                      | OR 143.039 |
| Altered mental status                          | 10.355 - 1975.8 |
| Acute respiratory failure                      | 1.760 - 40.887 |
| CK-MB, creatine kinase-MB; CPK, creatine phosphokinase; CXR, chest-X ray; LDH, lactate dehydrogenase |

| Table 4 Comparison of patients treated with and without corticosteroids |
|---------------------------------------------------------------|
| No corticosteroids (n = 103) & Corticosteroids (n = 52) | Univariate analysis | Multivariate analysis |
| Comorbidity                                                                                   |
| Cardiovascular diseases                                      | 35 | 10 | 0.463 (0.208 - 1.031) |
| Chronic lung diseases                                       | 20 | 2  | 0.116 (0.037 - 0.740) |
| Symptoms at disease onset                                                                 |
| Dyspnea                                                     | 40 | 33 | 2.736 (1.373 - 5.452) |
| Hemoptysis                                                   | 9  | 10 | 2.487 (0.942 - 6.568) |
| On hospital admission                                      |
| Dyspnea                                                     | 42 | 40 | 4.841 (2.275 - 10.304) |
| Pneumonia on CXR                                            | 81 | 48 | 3.259 (1.060 - 10.025) |
| Laboratory tests on admission                              |
| CPK                                                          | 260 ± 447 | 577 ± 795 | 1.001 (1.000 - 1.002) |
| CK-MB                                                        | 20 ± 30 | 32 ± 33 | 1.014 (0.997 - 1.032) |
| LDH, per 100 U/L increase                                    |
| Acute respiratory failure                                   | 24 (23.3%) | 38 (73.1%) | 8.935 (4.160 - 19.187) |
| Septic shock                                                | 5 (4.9%) | 13 (25.0%) | 6.533 (2.183 - 19.551) |
| Acute renal failure                                         | 5 (4.9%) | 10 (19.2%) | 4.667 (1.503 - 14.486) |
| Altered mental status                                       | 9 (8.7%) | 11 (21.2%) | 2.842 (1.093 - 7.390) |
| Bloodstream infection                                       | 0  | 3  | 1.061 (0.992 - 1.135) |
| Other infections                                             | 1  | 4  | 8.500 (0.925 - 78.106) |
| Supportive treatment                                        |
| Vasopressors                                                | 9  | 19 | 6.013 (2.478 - 14.596) |
| Neuromuscular blockade                                       | 1  | 7  | 15.867 (1.896 - 132.776) |
| Noninvasive ventilation                                      | 11 | 21 | 5.666 (2.467 - 13.063) |
| Invasive ventilation                                         | 16 | 27 | 5.873 (2.742 - 12.578) |
analysis revealed that corticosteroids treatment was associated with higher hospital mortality rate in patients without acute renal failure (10/42 vs. 6/98, p = 0.0064) or altered mental status (8/41 vs. 5/94, p = 0.0242), and in patients with bacterial pneumonia (9/11 vs. 4/17, p = 0.0085). There was no systemic treatment effect on hospital mortality of either lower or higher daily dose of corticosteroids compared with no corticosteroids treatment (Additional file 1, Table S1).

No interaction between corticosteroids treatment and any complications (including acute respiratory failure, septic shock, acute renal failure, altered mental status, bacterial pneumonia, bloodstream infection, and other infections) or oseltamivir treatment was found.

### Discussion

In this retrospective study of hospitalized adult patients with confirmed infection of 2009 influenza A(H1N1), we have found that diabetes, serum level of LDH on hospital admission, septic shock, and altered mental status were independent predictors of hospital mortality. In addition, corticosteroids treatment is associated with a trend towards worse outcome, although it is more likely to be used in critical illness, as shown by an elevated LDH level and the presence of acute respiratory failure.

The mortality seen in our study is significantly higher than those of hospitalized patients in the United States [7], United Kingdom [14], and Australia [15]. The later studies often summarized the initial hospitalized patients in each country, when even mild cases were admitted due to uncertainty of disease progression and prognosis. In contrary, the patients in our study developed the illness about 5 months after the first case of 2009 influenza A(H1N1) in China [8], when only high-risk patients were hospitalized, while most mild cases were followed up in fever clinics. The above difference in admission policy may be reflected by the significantly longer median time from symptom onset to hospital admission (5 vs. 2 to 3 days), the higher proportion of patients with signs of pneumonia on Chest X-ray (83.2% vs. 29 to 40%), and the higher mortality rate in our study, which is similar to those observed in patients requiring intensive care support in Australia [16] and Canada [10].

Almost half of the 155 hospitalized patients in our analysis do not have any comorbidities. The absence of serious comorbidities emphasizes that young, relatively healthy adults were the primary population affected by 2009 influenza A(H1N1) infection. On the contrary, underlying medical conditions associated with complications from seasonal influenza have been consistently shown as risk factors for hospital admission [17], intensive care unit (ICU) admission [18,19], and death [17,19,20]. The commonly acknowledged comorbidities may be classified according to chronic medical conditions recognized by the Advisory Committee on Immunization Practices, including cardiovascular disease, pulmonary disease, liver disease, cancer, and diabetes [17]. Diabetes and stress-induced hyperglycemia are known to be associated with an increased risk of complications and death among critically ill patients [21]. In this study, when infected with 2009 influenza A(H1N1), patients with diabetes were at a higher risk of death, compared with those without diabetes. In a cohort of 1479 patients admitted to hospital with laboratory-confirmed pandemic (H1N1) influenza in Canada, Campbell reported that the risk of a severe outcome (including death and ICU admission) was greatest among patients with diabetes (relative risk 2.2, 95%CI 1.7 - 2.7) [18]. Likewise, when comparing 1266 hospitalized patients with the general population in France, Hanslik also found that diabetes was significantly associated with death (OR 3.5, 95%CI 2.5 - 5.1) [19].

Patients with increased serum level of LDH on hospital admission are at a higher risk of death during hospitalization. This is not a surprising finding, because, as a non-specific enzyme found ubiquitously in cells, the increased serum level of LDH probably indicates the degree of tissue necrosis, and hence the severity of the viral infection as well as pneumonia. The association of high level of LDH and severity of illness, or even death, although not found in patients with 2009 influenza A(H1N1), has been widely reported in patients with severe acute respiratory syndrome (SARS). For example, high LDH level on hospital admission was associated with development of acute respiratory distress syndrome [22], and death during hospital stay [23,24].

It is recommended that therapy with a neuraminidase inhibitor is especially important for patients with underlying risk factors and those with severe or progressive clinical illness [25]. However, we did not observe any treatment effect of oseltamivir in our analysis. Only 16 out of 155 patients (10.3%) in our study received oseltamivir treatment within 48 hours after symptom onset. In comparison, Jain and Louie found that early antiviral therapy was administered in 39% and 51% of patients, respectively [7,26]. Studies have shown that delayed antiviral treatment is associated with high risk of progression to severe disease such as ICU admission and respiratory failure, prolonged hospital stay, and even death [25,27]. It is also possible that the delay in antiviral treatment in our patients resulted in higher hospital mortality (17.4%) compared with the subgroup of adult hospitalized patients with 2009 influenza A(H1N1) in the United States [7,26].

Corticosteroids remain a controversy in the treatment of viral infection [28]. WHO strongly recommends that systemic corticosteroids should not be
Conclusions
We have found that hospitalized patients with 2009 influenza A(H1N1) have relatively poor outcome. Diabetes and elevated LDH level on hospital admission, as well as the presence of septic shock and altered mental status, represent independent risk factors for hospital death among these patients. Early recognition of these risk factors may help clinicians to identify the high-risk patients. In addition, corticosteroids should not be regarded as routine pharmacologic therapy, although this may need confirmation by further large scale prospective study.

Additional material

Additional file 1: Supplementary table S1. Table to show subgroup analysis of the effect of corticosteroids treatment on hospital mortality.

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Authors’ contributions
BD conceived of the study, participated in its design and coordination, had full access to all of the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis. XX conceived of the study, and participated in its design and coordination, and helped to draft the manuscript. YX participated in the design of the study, and helped to draft the manuscript. LJ participated in the design of the study, and helped to draft the manuscript. AL participated in the design of the study, and helped to draft the manuscript. JD participated in the design and coordination of the study, and helped to draft the manuscript. All authors reviewed and approved the final manuscript.

Competing interests
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