



## Influenza A-associated severe pneumonia in hospitalized patients: Risk factors and NAI treatments



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### ABSTRACT

**Objective:** The risk factors and the impact of NAI treatments in patients with severe influenza A-associated pneumonia remain unclear.

**Methods:** A multicenter, retrospective, observational study was conducted in Zhejiang, China during a severe influenza epidemic in August 2017–May 2018. Clinical records of patients (>14 y) hospitalized with laboratory-confirmed influenza A virus infection and who developed severe pneumonia were compared to those with mild-to-moderate pneumonia. Risk factors related to pneumonia severity and effects of NAI treatments (monotherapy and combination of peramivir and oseltamivir) were analyzed. **Results:** 202 patients with influenza A-associated severe pneumonia were enrolled, of whom 84 (41.6%) had died. Male gender (OR = 1.782; 95% CI: 1.089–2.917;  $P = 0.022$ ), chronic pulmonary disease (OR = 2.581; 95% CI: 1.447–4.603;  $P = 0.001$ ) and diabetes mellitus (OR = 2.042; 95% CI: 1.135–3.673;  $P = 0.017$ ) were risk factors related to influenza A pneumonia severity. In cox proportional hazards model, severe pneumonia patients treated with double dose oseltamivir (300mg/d) had a better survival rate compared to those receiving a single dose (150 mg/d) (HR = 0.475; 95%CI: 0.254–0.887;  $P = 0.019$ ). However, different doses of peramivir (300 mg/d vs. 600 mg/d) and combination therapy (oseltamivir-peramivir vs. monotherapy) showed no differences in 60-day mortality ( $P = 0.392$  and  $P = 0.658$ , respectively).

**Conclusions:** Patients with male gender, chronic pulmonary disease, or diabetes mellitus were at high risk of developing severe pneumonia after influenza A infection. Double dose oseltamivir might be considered in treating influenza A-associated severe pneumonia.

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## Introduction

Intense seasonal influenza epidemics were observed during the 2017/2018 seasons in countries around the world, leading to increased influenza-associated hospitalizations (Hammond et al., 2018). Similarly, the 2017–2018 winter epidemic in China was severe with higher levels of influenza like illness (ILI) activity, and influenza-related hospitalization exceeded the peak levels of the previous three influenza seasons (Chinese National Influenza Center, 2019). Pandemic and seasonal influenza infections can progress to severe pneumonia and cause substantial mortality (Paules and Subbarao, 2017). Patients who are elderly, obese, pregnant or with comorbidity are at high risks of hospitalization, ICU admission and mortality (Mertz et al., 2013; Zhang et al., 2019).

In severe cases, influenza virus can induce alveolar epithelium cell death, further leading to respiratory failure, acute respiratory distress syndrome (ARDS) and multiorgan dysfunction (Short et al., 2014). Antiviral treatment is the primary approach against severe influenza, with neuraminidase inhibitors (NAIs) (oseltamivir and peramivir) most frequently used. The World Health Organization (WHO) recommends a standard dose of NAIs for non-complicated influenza (World Health Organization, 2010). There is substantial variation in clinical treatment courses, and it is not known whether a standard dose of NAIs is adequate for severe influenza (Hurt and Kelly, 2016). Higher dose oseltamivir treatment was observed with rapid reduction of virus load in influenza B patients (Lee et al., 2013). However, studies of double dose oseltamivir and oseltamivir-peramivir combination therapy in hospitalized and ICU patients found no improved clinical outcomes (South East Asia Infectious Disease Clinical Research Network, 2013; Noel et al., 2017; Wang et al., 2018). Nevertheless, previous observational studies were affected by small sample sizes with limited applicability to severe influenza pneumonia (Dixit et al., 2015; Lee et al., 2013).

The objective of our study was to systematically assess clinical characteristics, treatments and outcomes of hospitalized patients with influenza A-associated severe pneumonia as compared to those with mild-to-moderate pneumonia, and to evaluate the risk factors related to pneumonia severity and identify the impact of different NAI regimes.

## Materials and methods

### Study design

A multicenter, retrospective, observational study involving 13 teaching hospitals in Zhejiang, China between August 2017 and May 2018 was conducted. Patients (>14 years) hospitalized with influenza A infection and who developed severe pneumonia were included. Influenza A virus was confirmed by real-time polymerase chain reaction (RT-PCR) or immunofluorescence assay. We collected 202 patients from the same season with mild-to-moderate pneumonia due to influenza A, who had complete clinical records, for comparison. A team of trained physicians reviewed all medical records and collected the following data: demographics, comorbidities, clinical manifestations, laboratory and chest radiographic features on admission, treatments and clinical outcomes. Ethical approval was obtained from the research ethics committee of the First Affiliated Hospital, Zhejiang University School of Medicine, and informed consents were waived due to the non-interventional nature of our study.

## Definitions

Pneumonia was defined according to the American Thoracic Society-Infectious Diseases Society of America (ATS/IDSA) guideline (Mandell et al., 2007). Classification as severe pneumonia needed to meet one of the major criteria or at least three minor criteria. Major criteria included (1) invasive mechanical ventilation, (2) septic shock with the need for vasopressors, and minor criteria were (1) respiratory rate  $\geq 30$  breaths per min; (2)  $\text{PaO}_2/\text{FiO}_2 \leq 250$ ; (3) multilobar infiltrates; (4) confusion or disorientation; (5) blood urea nitrogen  $\geq 20$  mg/dL; (6) leucocyte count  $< 4 \times 10^9$  cells/L; (7) platelet count  $< 100 \times 10^9$  cells/L; (8) core temperature  $< 36$  °C; (9) hypotension requiring aggressive fluid resuscitation (Mandell et al., 2007). Patients with pneumonia that did not meet the criteria of severe pneumonia were classified as mild-to-moderate pneumonia. Septic shock was considered in patients who required a vasopressor to maintain mean arterial pressure  $\geq 65$  mmHg and serum lactate  $> 2$  mmol/L despite adequate fluid resuscitation (Singer et al., 2016). Secondary infection was defined as the recurrence of symptoms and signs of infection along with positive cultures of bacterial/fungal from lower respiratory tract specimens and/or blood after 48 h admission (Rothberg and Haessler, 2010).

### Statistical analysis

Categorical variables were presented as percentage and compared by Chi-square test or Fisher's exact test. Continuous variables were expressed as median (interquartile ranges) and compared with *t* test or Mann-Whitney *U* test. Significant or clinically plausible variables were included in multivariate logistic regression with enter method to assess risk factors associated with influenza A pneumonia severity. Results were presented as odds ratio (OR) and 95% confidence interval (CI). A stepwise Cox proportional hazards model was used to estimate the cumulative survival rate among different doses of oseltamivir treatments in severe pneumonia group. Covariates included significant or potential clinically relevant baseline variables. Statistical analyses were carried out by SPSS (version 22.0) and a *P* value  $< 0.05$  (2-tailed) was considered significant.

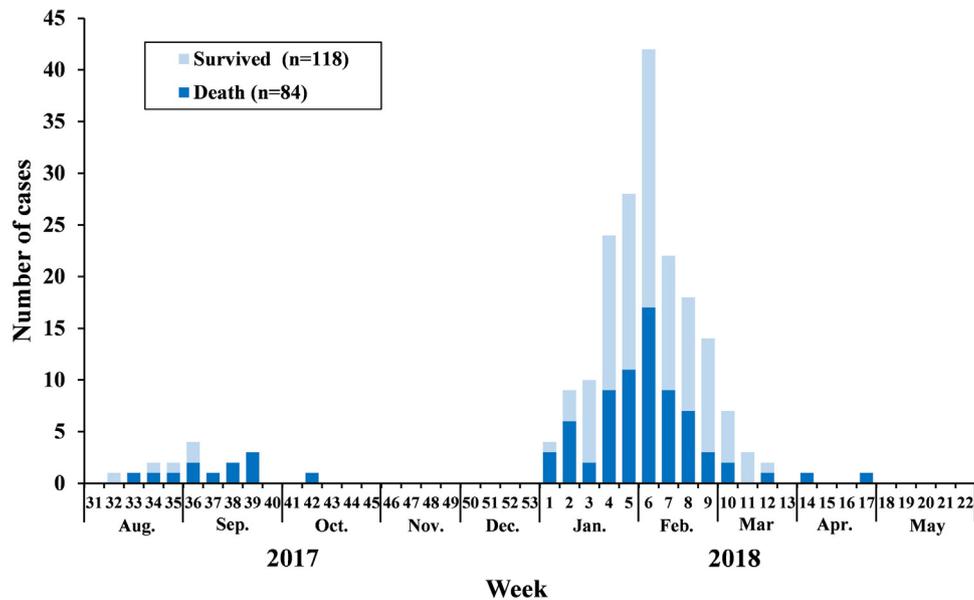
## Results

### Prevalence of influenza A-associated severe pneumonia

During the study period between August 2017 and May 2018, 4468 hospitalized patients from 13 hospitals in Zhejiang were infected with influenza A virus and 202 (4.5%) developed severe pneumonia. Of 202 severe cases, 84 (41.6%) had died. Our study included a small number of the severe pneumonia cases (*n* = 16) from August to September 2017 along with a majority collected during the peak (*n* = 183) between January and February 2018 (Figure 1).

### Clinical characteristics and outcomes

The proportion of patients aged  $\geq 65$  years old was significantly higher in the severe pneumonia group (*P* = 0.022) (Table 1). Comorbidities including hypertension, chronic pulmonary disease, diabetes mellitus, immunosuppression and symptoms including fever, sputum, chest distress and hemoptysis were more



**Figure 1.** Influenza A-associated severe pneumonia patients in Zhejiang, China, through August 2017 to May 2018. Weekly number of patients that were hospitalized.

**Table 1**

Clinical characteristics of 404 patients with influenza A-associated pneumonia according to disease severity.

Variables	Mild-to-moderate pneumonia (n = 202)	Severe pneumonia (n = 202)	P values
<b>Demographics</b>			
Age (median [IQR]) (yr)	61 (48–72.3)	66 (50.8–76)	<b>.041</b>
Age ≥65y, n (%)	84 (41.6)	107 (53.0)	<b>.022</b>
Male gender, n (%)	118 (58.4)	141 (69.8)	<b>.017</b>
Smoking history, n (%)	63 (31.2)	70 (34.7)	.459
<b>Comorbidities, n (%)</b>			
Any comorbidity	120 (59.4)	139 (68.8)	<b>.049</b>
Hypertension	67 (33.2)	87 (43.1)	<b>.040</b>
Chronic pulmonary disease <sup>a</sup>	24 (11.9)	50 (24.8)	<b>.001</b>
Diabetes mellitus	22 (10.9)	42 (20.8)	<b>.006</b>
Malignancy	10 (5.0)	8 (4.0)	.630
Coronary heart disease	11 (5.4)	12 (5.9)	.830
Chronic liver disease	6 (3.0)	13 (6.4)	.100
Chronic renal disease	6 (3.0)	12 (5.9)	.148
Immunosuppression <sup>b</sup>	1 (0.5)	14 (6.9)	<b>.001</b>
<b>Clinical manifestations, n (%)</b>			
Fever	178 (88.1)	190 (94.1)	<b>.036</b>
Cough	184 (91.1)	190 (94.1)	.255
Sputum	160 (79.2)	178 (88.1)	<b>.015</b>
Chest distress	55 (27.2)	150 (74.3)	<b>&lt;.0001</b>
Gastrointestinal	24 (11.9)	26 (12.9)	.763
Hemoptysis	5 (2.5)	14 (6.9)	<b>.034</b>
<b>Laboratory results on admission</b>			
Leukocyte count (median [IQR]) (mm <sup>3</sup> )	6 (4–8.9)	7.2 (4.5–10.1)	<b>.015</b>
Lymphocyte count (median [IQR]) (mm <sup>3</sup> )	0.9 (0.6–1.3)	0.5 (0.3–0.7)	<b>&lt;.0001</b>
Hemoglobin (median [IQR]) (g/L)	125 (112–138)	123 (102–137.8)	.307
Platelet count (median [IQR]) (mm <sup>3</sup> )	165 (125–231)	143 (105–211)	<b>.013</b>
Creatine kinase (median [IQR]) (UI/L)	83 (43–192.3)	143 (51.8–358.8)	<b>.001</b>
Aspartate transaminase (median [IQR]) (UI/L)	29 (21.3–43)	51 (30–93)	<b>&lt;.0001</b>
Creatinine (median [IQR]) (μmol/L)	73 (61–88)	77 (61–102)	.066
Lactate dehydrogenase (median [IQR]) (UI/L)	247 (191–324)	429 (277.8–710.8)	<b>&lt;.0001</b>
D-dimer (median [IQR]) (mg/L)	0.52 (0.31–0.91)	1.52 (0.77–3.08)	<b>&lt;.0001</b>
PaO <sub>2</sub> (median [IQR]) (mmHg)	79 (64–103)	62.8 (50.1–81.9)	<b>&lt;.0001</b>
K <sup>+</sup> (median [IQR]) (mmol/L)	3.9 (3.6–4.3)	3.9 (3.5–4.3)	.673
C-reactive protein (median [IQR]) (mg/L)	48.2 (21–99.1)	90 (42.9–148)	<b>&lt;.0001</b>
Procalcitonin (median [IQR]) (ng/mL)	0.12 (0.05–0.4)	0.7 (0.2–3)	<b>&lt;.0001</b>
<b>Chest radiographic features on admission</b>			
Infiltrations involving bilateral lungs, n (%)	78 (38.6)	188 (93.1)	<b>&lt;.0001</b>

IQR, interquartile range.

<sup>a</sup> Chronic pulmonary diseases include chronic obstructive pulmonary disease, asthma and bronchiectasis.

<sup>b</sup> Immunosuppression includes receiving radiotherapy, chemotherapy or corticosteroid therapy within one month before symptom onset.

**Table 2**

Treatment and clinical outcomes of 404 patients with influenza A-associated pneumonia according to disease severity.

Variables	Mild-to-moderate pneumonia (n = 202)	Severe pneumonia (n = 202)	P values
<b>Treatments</b>			
Invasive mechanical ventilation, n (%)	0 (0)	140 (69.3)	<0.001
Extracorporeal membrane oxygenation, n (%)	0 (0)	13 (6.4)	<0.001
Continuous renal replacement therapy, n (%)	0 (0)	11 (5.4)	.001
Antibiotic treatment, n (%)	199 (98.5)	200 (99.0)	.653
Corticosteroid treatment, n (%)	67 (33.2)	159 (78.7)	<0.001
NAIs treatment, n (%)	192 (95.0)	191 (94.6)	.823
NAIs administered ≤48 h from illness onset, n (%)	23 (11.4)	17 (8.4)	.318
Duration of NAIs treatment (median [IQR]) (d)	6 (4–9.8)	7 (5–10)	<0.001
<b>Clinical outcomes</b>			
Septic Shock, n (%)	0 (0)	39 (19.3)	<0.001
Secondary bacterial infection, n (%)	1 (0.5)	57 (28.2)	<0.001
ICU admission, n (%)	0 (0)	135 (66.8)	<0.001
60-day mortality, n (%)	0 (0)	84 (41.6)	<0.001
Length of stay in hospital (median [IQR]) (d)	9 (6–13.3)	16 (10–24)	<0.001

IQR, interquartile range; NAI, neuraminidase inhibitor; ICU, intensive care unit.

commonly seen in the severe pneumonia group. Among laboratory results on admission, higher levels of leukocyte count, creatine kinase, aspartate transaminase, lactate dehydrogenase, D-dimer, C-reactive protein, procalcitonin and lower levels of lymphocyte count, platelet count and PaO<sub>2</sub> were observed in severe pneumonia group. The chest image of 188 (93.1%) patients in the severe pneumonia group showed infiltrations involving bilateral lungs, which was significantly higher than those in mild-to-moderate pneumonia group ( $P < 0.001$ ) (Table 1).

Supportive therapies provided for the severe pneumonia group include invasive mechanical ventilation (69.3%), extracorporeal membrane oxygenation (6.4%) and continuous renal replacement therapy (5.4%). A higher percentage of corticosteroid treatment was observed in the severe pneumonia group (78.7% vs. 33.2%,  $P < 0.001$ ), but no difference in treatments using antibiotics and NAIs between the two groups. The proportion of NAIs administered ≤48 h from illness onset in mild-to-moderate pneumonia group was higher than the severe pneumonia group, although no statistical difference was found (11.4% vs. 8.4%,  $P = 0.318$ ). Patients with severe pneumonia had significantly higher rates of septic shock ( $P < 0.001$ ) and secondary infection ( $P < 0.001$ ) comparing to mild-to-moderate pneumonia. 135 (66.8%) patients in the severe pneumonia group were admitted into the ICU and 84 (41.6%) had died (Table 2).

#### Risk factors for influenza A-associated pneumonia severity

In the multivariate regression analysis, risk factors associated with influenza A-associated pneumonia severity were as follows: male gender (OR = 1.782; 95% CI: 1.089–2.917;  $P = 0.022$ ), chronic pulmonary disease (OR = 2.581; 95% CI: 1.447–4.603;  $P = 0.001$ ) and diabetes mellitus (OR = 2.042; 95% CI: 1.135–3.673;  $P = 0.017$ ), whereas age, smoking history and hypertension were not the independent risk factors (Table 3).

#### NAIs treatment on influenza A-associated severe pneumonia

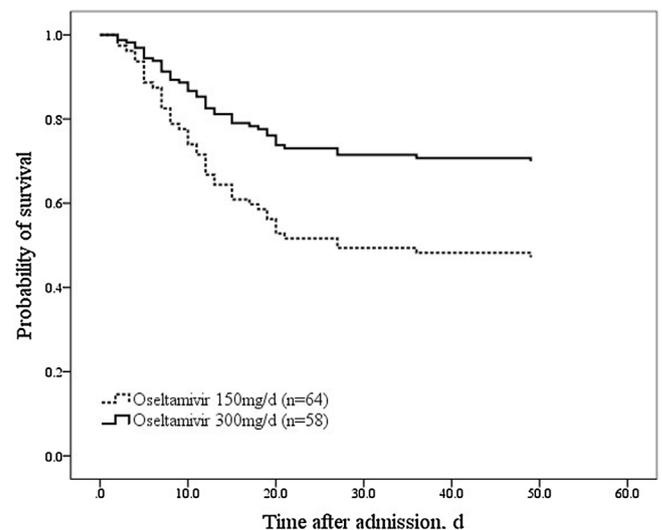
Of 191 NAI-administered patients in the severe pneumonia group, 122 (63.9%) received oseltamivir monotherapy, 40 (20.9%) received peramivir monotherapy and 29 (15.2%) received oseltamivir-peramivir combination therapy. Demographic information showed that patients treated with double dose oseltamivir (300 mg/d) were younger ( $P < 0.001$ ) and had less comorbidities ( $P = 0.002$ ) compared to those with single dose (150mg/d) treatment (Table S1). In adjusted Cox regression survival analysis, double dose oseltamivir (300mg/d) recipients

**Table 3**

Multivariate analysis for risk factors associated with influenza A pneumonia severity.

Factors	OR	95% CI	P values
Age	1.000	0.986–1.013	.967
Male gender	1.782	1.089–2.917	.022
Smoking history	0.780	0.470–1.293	.335
Chronic pulmonary disease	2.581	1.447–4.603	.001
Diabetes mellitus	2.042	1.135–3.673	.017
Hypertension	1.283	0.811–2.030	.287

OR, odds ratio; CI, confidence interval.



**Figure 2.** Survival plot classified by oseltamivir treatments in patients with influenza A-associated severe pneumonia. Cox regression for adjusted survival in patients treated with 150 mg/d and 300 mg/d oseltamivir. Covariates included oseltamivir treatment (300 mg/d vs. 150 mg/d; HR = 0.475, 95%CI: 0.254–0.887,  $P = 0.019$ ), age, sex, any comorbidity, oseltamivir initial time and SOFA score. SOFA: sequential organ failure assessment.

had a higher cumulative survival rate compared to single dose (150mg/d) recipients (HR = 0.475, 95%CI: 0.254–0.887,  $P = 0.019$ ; Figure 2). However, there were no significant differences in 60-day mortality between patients who received double dose (300 mg/d) and single dose (150 mg/d) peramivir treatment ( $P = 0.392$ ) (Table S2), and among oseltamivir-peramivir combination therapy, oseltamivir monotherapy and peramivir monotherapy ( $P = 0.658$ ) (Table S3).

## Discussion

Our study found that 4.5% (202/4468) of inpatients had influenza A-associated severe pneumonia and 1.9% (84/4468) of them died during the 2017–2018 influenza season. The overall in-hospital mortality during 2017–2018 was marginally higher than the 1.2% in-hospital mortality of seasonal influenza during 2011–2017 in China (Fu et al., 2019), although consistent with a global report of 1.5% in-hospital mortality of influenza respiratory tract infection in 2017 (GBD 2017 Influenza Collaborators, 2019). Fatality due to influenza A-associated severe pneumonia was substantially higher (84/202, 41.6%) than fatality reported among ICU admitted patients during 2009–2016 (26.4–20.6%) (Shah et al., 2015; Alvarez-Lerma et al., 2016; Chao et al., 2017). The high mortality of influenza A-associated pneumonia observed in our study suggests that more effective treatments toward this certain population are urgently warranted.

Improved survival requires the identification of factors associated with severe pneumonia following influenza infection and timely treatments, with host factors and treatment regimens being key drivers of severity. Previous studies reported underlying chronic medical conditions of the host, such as cardiac disease, chronic pulmonary disease, immunosuppression, diabetes mellitus, obesity and pregnancy, were major factors associated with influenza severity (Chow et al., 2019). Similarly, our study found patients with comorbidities including chronic pulmonary disease and diabetes mellitus had a higher predisposition to critical influenza (Mertz et al., 2013; Zhang et al., 2019). Decreased lung compliance is related to persistent viral RNA detection, which contribute to the higher risk of progression to severe disease in these patients (Chow et al., 2019). In our study, male gender was an independent risk factor related to pneumonia severity, which might be attributed to the higher frequency of comorbidities and intense immune states observed in males (Bonmarin et al., 2015; Shah et al., 2015). Virus factors driving enhancing disease severity in patients may include virus type/subtype and key genetic mutations (Kash and Taubenberger, 2015). Regrettably, few hospitals in our study subtyped the influenza A virus, therefore we were unable to assess the basis for routine influenza subtyping for patients with severe pneumonia. According to national surveillance reports, influenza A(H3N2) virus was predominant during July to October 2017 in Southern China, and influenza A(H1N1)pdm09 and B (Yamagata lineage) viruses gradually showed up and became dominant between January and February 2018 (Chinese National Influenza Center, 2019).

NAIs are the predominant countermeasure available for influenza infection (Chow et al., 2019). In this study, we mainly focused on different NAI treatments for patients with influenza A-associated severe pneumonia. NAIs are most beneficial when commenced within 48 h from illness onset when viral replication is active (Lee et al., 2009). However, only a few patients (9.9%) had received NAIs within 48 h of illness onset in our study. A higher rate of NAI treatment within 48 h was observed in the mild-to-moderate pneumonia group, indicating the importance of prompt diagnosis and NAIs treatment following infection.

The WHO guideline recommends a higher dose of oseltamivir (e.g. 300 mg/d in adults) to treat critically ill patients. However, whether a higher dose could provide additional protection against severe influenza remains controversial due to the implications in stockpile (Hurt and Kelly, 2016). Studies conducted in avian influenza A(H7N9) infected patients (Wang et al., 2018), ICU-admitted patients with untyped influenza infection (Noel et al., 2017) and pediatric patients with severe influenza (South East Asia Infectious Disease Clinical Research Network, 2013) showed that double dose oseltamivir (300 mg per day) did not present additive effects on clinical outcomes (duration of mechanical ventilation

day, length of hospitalization, length of ICU stay or mortality) compared with the standard dose. In our study, after adjusting for covariates, double dose oseltamivir was correlated with improved survival in patients with influenza A-associated severe pneumonia, whereas different doses of peramivir and combination therapy of oseltamivir-peramivir showed no differences in survival.

Higher viral loads and prolonged viral shedding are observed in patients with severe influenza (Li et al., 2010), and an increased dose of oseltamivir in animal models has shown correlation with decreased mortality (Govorkova et al., 2007). Lee et al observed a faster RNA decline rate in influenza B virus infected inpatients (n = 10) who were treated with 300 mg/d oseltamivir (Lee et al., 2013), while Kumar et al. found that a triple dose oseltamivir in ICU patients (n = 18) was associated with better virus clearance, but observed no differences in clinical outcomes (Kumar, 2013). These observational studies were small with a few ICU patient populations, while our study focused on patients with community acquired influenza A-associated severe pneumonia, a well-defined population and a substantially larger sample size than previous studies. In addition, our study was conducted in the same influenza season during the predominant circulation of A(H1N1)pdm09 subtype, therefore virus variations would have limited influence on the potency of NAIs. Our study urges further research on virus clearance and pharmacokinetics of different dose oseltamivir treatments in influenza A-associated severe pneumonia.

Peramivir is an intravenous NAI favored for seriously ill patients or those unable to tolerate oral antivirals (Wester and Shetty, 2016). Reduction of mortality was insignificant between treatment of 300 mg or 600 mg peramivir once daily, which was congruent with other studies in severe influenza (de Jong et al., 2014; Ison et al., 2014). This might be potentially explained by the high-dose treating preference towards more severe patients, however the group size in our study was small (n = 43). In addition, our study found no significant difference in 60-day mortality among monotherapy and oseltamivir-peramivir combination therapy. Similarly, studies of A(H7N9) hospitalized patients comparing oseltamivir-peramivir combination to monotherapy found no additional benefits in virologic or clinical outcomes (Wang et al., 2018; Zhang et al., 2016). Research on seasonal influenza and A(H1N1)pdm09 infection also showed no synergistic effect between oseltamivir-zanamivir bithery and monotherapy (Duval et al., 2010; Escuret et al., 2012). Due to the similarity in chemical structure, NAIs combination treatment might not exert a greater antiviral effect (Duval et al., 2010). Novel antiviral drugs such as Baloxavir Marboxil (Hayden et al., 2018) with different mechanisms would be an option for potential combination therapy.

There are several limitations in this study. First, as an observational and retrospective study, certain biases, such as the lost information of patients with incomplete records, unstandardized disease course on patients' admission, and physician's treatment preference, are difficult to completely rule out. Secondly, our study had a small sample size, and it is underpowered to evaluate the effects of different NAIs treatments in subgroups. Our result regarding the benefit of double dose oseltamivir treatment in influenza A-associated severe pneumonia must be interpreted carefully and within the limitation of the study design. In addition, the virologic response is a direct indicator in evaluating the effects of different NAI regimens. However, we were unable to assess the daily changes of viral RNA concentrations due to the retrospective nature of our study.

## Conclusions

A high rate of mortality due to influenza A-associated severe pneumonia was observed among hospitalized patients in China during a peak infection in the 2017–2018 season. Patients who developed influenza A-associated pneumonia were more likely to

be of male gender, with chronic pulmonary disease and diabetes mellitus. Our study urges closer attention to this high-risk group during influenza season, offering a chance for prompt NAs treatment, preventing progressing into severe pneumonia. We also provided supportive evidence for double dose oseltamivir in treating patients with influenza A-associated severe pneumonia.

### Declaration of interest

None.

### Authors contributions

Yu Chen and Shufa Zheng designed the study and reviewed the manuscript. Qianda Zou conducted data analysis and drafted the manuscript. Xiaochen Wang and Sijia Liu involved data collection, data analysis and manuscript revising. Jiaqi Bao and Fei Yu conceived the study and interpreted data. Wei Wu, Xianjun Wang, Bo Shen, Tieli Zhou, Zhigang Zhao, Yiping Wang, Ruchang Chen, Wei Wang, Jianbo Ma, Yongcheng Li, Xiaoyan Wu, Weifeng Shen and Fuyi Xie contributed in data collection from each hospital. Dhanasekaran Vijaykrishna reviewed the manuscript. All authors read and approved the final manuscript.

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### Appendix A. Supplementary data

Supplementary material related to this article can be found, in the online version, at doi:<https://doi.org/10.1016/j.ijid.2020.01.017>.

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