Clinical characteristics and survival analysis in critical and non-critical patients with COVID-19 in Wuhan, China: a single-center retrospective case control study

Ran Tian\(^1,10\), Wei Wu\(^1\), Chunyao Wang\(^2,10\), Haiyu Pang\(^3,10\), Zhiyu Zhang\(^4,10\), Haopeng Xu\(^4,10\), Qingfeng Luo\(^5,10\), Peng Gao\(^1\), Jihua Shi\(^5\), Wenbin Li\(^1\), Hao Qian\(^1\), Fan Guo\(^1\), Taisheng Li\(^6\), Zhengyin Liu\(^6\), Jinglan Wang\(^7\), Xiang Zhou\(^8\), Yan Qin\(^9\), Xiaowei Yan\(^1\) & Shuyang Zhang\(^1\)*

Since the outbreak of COVID-19 in China at the end of 2019, the world has experienced a large-scale epidemic caused by the SARS-CoV-2. The epidemiological and clinical course of COVID-19 patients has been reported, but there have been few analyses about the characteristics, predictive risk factors, and outcomes of critical patients. In this single-center retrospective case–control study, 90 adult inpatients hospitalized at Tongji Hospital (Wuhan, China) were included. Demographic, clinical, laboratory tests, and treatment data were obtained and compared between critical and non-critical patients. We found that compared with non-critical patients, the critical patients had higher SOFA score and qSOFA scores. Critical patients had lower lymphocyte and platelet count, elevated D-dimer, decreased fibrinogen, and elevated high-sensitivity C-reactive protein (hsCRP), and interleukin-6 (IL-6). More critical patients received treatment including antibiotics, anticoagulation, corticosteroid, and oxygen therapy than non-critical ones. Multivariable regression showed higher qSOFA score and elevation of IL-6 were related to critical patients. Antibiotic usage and anticoagulation were associated with decreased in-hospital mortality. And critical grouping contributed greatly to in-hospital death. Critical COVID-19 patients have a more severe clinical course. qSOFA score and elevation of IL-6 are risk factors for critical condition. Non-critical grouping, positive antibiotic application, and anticoagulation may be beneficial for patient survival.
confirmed deaths\textsuperscript{2}. The World Health Organization (WHO) has claimed COVID-19 as a global pandemic\textsuperscript{3}. And health systems in all countries are currently faced with serious challenges.

Although we have acquired a deeper understanding of the disease through autopsy and virological study, no specific treatment seems to be definitive and effective for preventing the disease progression and death of critical patients. According to the diagnosis and treatment guidelines released by China Health and Medical Commission\textsuperscript{4}, patients can be divided into four types clinically: mild, ordinary, severe and critical. The clinical manifestation and required treatment vary greatly between different types. Mild and ordinary patients need only supportive treatment, and self-healing cases have been reported, but most of the critically ill patients need mechanical ventilation or even extra-corporeal membrane oxygenation (ECMO) and continuous renal replacement therapy (CRRT). The mortality rate reported in different literatures is between 1–4\%\textsuperscript{5–7}, but the severity of the illness and the mortality rate can be underestimated due to a large number of asymptomatic infections and mild patients. Most current researches are descriptive studies on patients admitted to hospitals, yet there have been few analyses about the outcomes and risk factors of critical patients until now.

Here, we present the clinical course and outcomes of a group of critical patients with COVID-19, and attempt to identify risk factors for disease progression and in-hospital mortality in these patients. We aim to find some predictive risk factors for early warning, to provide opportunities for timely medical intervention by simple and effective assessment.

**Results**

We collected 90 inpatients diagnosed with COVID-19 at the Tongji Hospital (Wuhan, China) from Jan 28th to Feb 28th. Of all the patients, 45 are critical patients and 45 are non-critical patients. All patients were discharged or died before the date of data collection. 32 of the 90 patients died and 48 were discharged. (Fig. 1) The median age of all the patients was 64 years (56–70), ranging from 26 to 92. 48 patients were males and 42 were females. No significant differences were observed between the two groups in terms of age, gender, and comorbidities. (Table 1).

Vital signs at the time of admission were analyzed. The critical patients had faster heart rate (102 ± 19, per min) and respiratory rate (26, 22–35, per min), higher SOFA score (5, 4–7) and qSOFA score(2, 1–2). More patients in the critical group had symptoms of fever, expectoration, and dyspnea, but most of the symptoms like cough, fatigue, chill, etc. were non-specific and showed no significance between groups (not all shown in the table).

The mortality rate in the critical group was 67\%, which was significantly higher than the non-critical group. The median time from onset to admission in the critical group was 16 days (10–25), longer than that in the non-critical group. There was no significant difference in the time of hospitalization and the entire course of disease between the two groups.

Results of blood test at admission for all patients were obtained and analyzed (Table 2). It was found that the critical group had higher white blood cell (10.25, 7.96–15.14, ×10\textsuperscript{9}/L) and neutrophil count (9.21,
Elevated alanine aminotransferase (ALT, 29.0, 20.0–45.5, U/L), total bilirubin (12.2, 7.8–18.9, μmol/L), hypoalbuminemia (29.6, 5.4, g/L), and hyperuricemia (7.6, 5.6–12.7) were also more commonly observed in the critical group. Meanwhile, critical patients are more likely to show prolonged partial thromboplastin time (PT, 16.7, 15.1–18.2, s), increased D-dimer, and decreased fibrinogen. In terms of inflammatory factors, the proportion of patients with elevated ferritin and elevated hsCRP and IL-6 in the critical group was also larger.

More than two-thirds of the 90 patients (n = 62) received varying level of oxygen therapy support, of which 36 patients had mechanical ventilation. The critical group has higher requirements for oxygen therapy support. More patients need high-flow oxygen inhalation, prone ventilation, BiPAP and mechanical ventilation. The usage of antiviral drugs is similar between the two groups. More patients were treated with antibiotics, intravenous immunoglobulin (IVIG), and glucocorticoid in the critical group; more critical patients received treatment for complications, including renal replacement therapy and anticoagulation. 6 of the patients received anti-IL-6 treatment (5 died) and 4 received ECMO (3 died).

To find the possible indicators for patients’ group, potential influential factors including differences between the two groups and factors related to clinical diagnosis and treatment were screened. Age, gender, qSOFA scores, low lymphocyte (less than 0.8), high D-dimers (greater than 1), and high IL-6 (greater than twice the upper limit) were analyzed in conditional logistic regression using COX survival analysis. We found that qSOFA scores and increased IL-6 were significantly associated with critical group (Table 3).

6.77–13.10, × 10⁹/L) and lower lymphocyte count. Platelet counts are also lower.

| Table 1. Demographic and clinical characteristics of patients on admission. Data are median (IQR), average (SD) or n (%). P values comparing patients are from Student’s t test, chi-square test, or Fisher’s exact test. HR heart rate, RR respiratory rate, SBP systolic blood pressure, DBP diastolic blood pressure, SOFA Sequential Organ Failure Assessment score, qSOFA quick Sequential Organ Failure Assessment Score. *The data is normally distributed, thus average (SD) are used. †Limited sample number: non-critical 33 samples, critical: 44 samples. *Limited sample number: critical 43 samples.

| Total (n = 90) | Non-critical (n = 45) | Critical (n = 45) | p-value |
|----------------|-----------------------|-------------------|---------|
| Age, years     | 64 (56–70)            | 63 (59–70)        | 64 (56–71) | 0.984 |
| Sex            |                       |                   |          | 0.398 |
| Male           | 48 (53%)              | 22 (49%)          | 26 (58%) |       |
| Female         | 42 (47%)              | 23 (51%)          | 19 (42%) |       |
| Comorbidity    |                       |                   |          |       |
| Cardiovascular disease | 11 (12%) | 4 (9%) | 7 (15.6%) | 0.334 |
| Hypertension   | 38 (42%)              | 19 (42%)          | 19 (42%) | 1.000 |
| Diabetes mellitus | 17 (19%) | 9 (20%) | 8 (18%)  | 0.788 |
| Chronic obstructive pulmonary disease | 4 (4%) | 2 (4%) | 2 (4%)  | 1.000 |
| Chronic kidney disease | 1 (1%) | 0 (0%) | 1 (1%)  | 1.000 |
| Cerebrovascular disease | 6 (7%) | 2 (4%) | 4 (9%)  | 0.677 |
| Malignance     | 10 (11%)              | 8 (18%)           | 2 (4%)   | 0.044 |
| Disease progression and outcome |                   |                   |          |       |
| Time between illness onset and hospital admission | 14 (7–22) | 12 (5–15) | 16 (10–25) | 0.004 |
| Time between hospital admission and outcome | 16 (9–26) | 18 (13–24) | 12 (5–29) | 0.455 |
| Time between illness onset to outcome* | 31 (21–42) | 30 (22–34) | 32 (19–49) | 0.422 |
| Death          | 32 (36%)              | 2 (4%)            | 30 (67%) | <0.001 |
| Vital signs    |                       |                   |          |       |
| HR*, min⁻¹     | 102 (19)              | 93 (15)           | 110 (19) | <0.001 |
| RR, min⁻¹      | 20 (20–28)            | 20 (18–20)        | 26 (22–35) | <0.001 |
| SBP*, mmHg     | 130 (20)              | 132 (19)          | 128 (21) | 0.288 |
| DBP*, mmHg     | 80 (13)               | 82 (13)           | 78 (13)  | 0.172 |
| SOFA           | 3 (1–5)               | 1 (1–2)           | 5 (4–7)  | <0.001 |
| qSOFA          | 1 (0–2)               | 0 (0–0)           | 2 (1–2)  | <0.001 |
| Symptoms†      |                       |                   |          |       |
| Fever          | 34 (38%)              | 11 (24%)          | 23 (40%) | 0.009 |
| Cough          | 50 (57%)              | 25 (57%)          | 25 (58%) | 0.807 |
| Dyspnea        | 27 (31%)              | 4 (9%)            | 23 (54%) | <0.001 |
| Expectoration  | 32 (36%)              | 11 (24%)          | 21 (49%) | 0.017 |
| Fatigue        | 44 (50%)              | 24 (53%)          | 20 (47%) | 0.522 |
glucocorticoid usage, BiPAP, mechanical ventilation, etc. (data not shown). Corticosteroid was the first reported drug that can lower 28-day mortality of patients hospitalized with COVID-19 among those who were receiving either invasive mechanical ventilation or oxygen alone at randomization but not among those receiving no respiratory support\(^8\), which supported our findings that corticosteroid (\(p = 0.016\)), non-invasive ventilation (\(p = 0.014\)) and invasive ventilation (\(p < 0.001\)) were considered to be linked with in-hospital mortality. Then, five factors, especially treatment factors, were selected in multivariate regression analysis, among which age, disease grouping, antibiotics and anticoagulation were related to death. (Fig. 2. and Table 4.)

### Table 2. Laboratory test results and treatment. Data are median (IQR), average (SD) or n (%). \(^*\)The data is normally distributed, thus average (SD) are used.

|                     | Total (n = 90) | Non-Critical (n = 45) | Critical (n = 45) | p     |
|---------------------|---------------|-----------------------|------------------|-------|
| White blood cell, \(\times 10^9/\text{L}\) | 7.56 (5.08–10.55) | 5.90 (3.97–7.25) | 10.25 (7.96–15.14) | <0.001 |
| Neutrophil, \(\times 10^9/\text{L}\) | 6.04 (3.36–9.46) | 3.72 (2.40–5.49) | 9.21 (6.77–13.10) | <0.001 |
| Lymphocyte, \(\times 10^9/\text{L}\) | 0.84 (0.48–1.22) | 1.04 (0.80–1.43) | 0.54 (0.33–0.89) | <0.001 |
| Hemoglobin, g/L      | 120.5 (105.8–132.3) | 119 (107–132) | 124 (105–136) | 0.470  |
| Platelet, \(\times 10^{12}/\text{L}\) | 192.50 (136.25–285.25) | 230.00 (164.00–310.50) | 159.00 (102.00–235.50) | 0.005  |
| ALT, U/L             | 24.5 (14.0–45.0) | 19.0 (11.0–38.5) | 29.0 (20.0–45.5) | 0.043  |
| Total bilirubin, \(\mu\text{mol}/\text{L}\) | 10.5 (7.3–15.8) | 9.9 (6.8–12.8) | 12.2 (7.8–18.9) | 0.047  |
| Serum creatinine, \(\mu\text{mol}/\text{L}\) | 70.0 (56.5–91.3) | 64.0 (57.5–80.5) | 79.0 (50.5–106) | 0.211  |
| BUN, mmol/L          | 5.1 (3.9–8.1) | 4.1 (3.4–4.9) | 7.6 (5.6–12.7) | <0.001 |
| Albumin*, g/L        | 31.7 (5.4) | 33.8 (4.6) | 29.6 (5.4) | <0.001 |
| INR                  | 1.1 (1.0–1.4) | 1.0 (1.0–1.1) | 1.3 (1.2–1.5) | <0.001 |
| PT, s                | 14.5 (13.7–17.1) | 13.7 (13.4–14.3) | 16.7 (15.1–18.2) | <0.001 |
| APTT, s              | 39.4 (35.7–44.4) | 38.7 (35.7–45.1) | 40.8 (35.5–43.7) | 0.704  |
| D-dimer >1 mg/L      | 66 (73%) | 25 (56%) | 41 (91%) | <0.001 |
| Fibrinogen, g/L      | 4.7 (3.3–6.0) | 5.25 (4.1–6.2) | 3.9 (2.6–5.4) | 0.005  |
| Fcr >400 ug/L        | 77 (86%) | 33 (73.3%) | 44 (97.8%) | 0.003  |
| hsCRP >3 mg/L        | 77 (86%) | 34 (76%) | 43 (96%) | 0.007  |
| IL-6 >14 pg/mL       | 50 (56.2%) | 15 (29%) | 37 (84%) | <0.001 |

### Table 3. Risk factors associated with critical patients. \(^*\)The data is normally distributed, thus average (SD) are used.

| Treatment       | OR     | 95% CI          | p      |
|-----------------|--------|-----------------|--------|
| Antibiotics     | 47 (52%) | 16 (36%) | 31 (69%) | 0.002  |
| Antiviral       | 49 (54%) | 21 (47%) | 28 (62%) | 0.138  |
| Anticoagulation | 25 (28%) | 0 (0%) | 25 (56%) | <0.001 |
| Corticosteroid  | 43 (48%) | 9 (20%) | 34 (76%) | <0.001 |
| IVIG            | 38 (42%) | 3 (7%) | 35 (78%) | <0.001 |
| Anti-IL6 therapy| 6 (7%) | 1 (2%) | 5 (11%) | 0.203  |
| CRRT            | 7 (8%) | 0 (0%) | 7 (16%) | 0.012  |
| High flow oxygen| 13 (17%) | 1 (2%) | 14 (31%) | 0.001  |
| Prone ventilation| 14 (16%) | 1 (2%) | 13 (16%) | 0.001  |
| BiPAP           | 11 (12%) | 2 (4%) | 9 (20%) | 0.024  |
| Invasive ventilation | 36 (40%) | 1 (2%) | 35 (78%) | <0.001 |
| ECMO            | 4 (4%) | 0 (0%) | 4 (9%) | 0.117  |
COVID-19 has now become a global pandemic, and medical systems in many countries are facing serious challenges. Survival of critical and non-critical patients are significantly different, so the early identification of critical patients is very important. This retrospective research shows that critical patients have more severe clinical situation and worse prognosis, which requires medical support of high grade, including higher level of oxygen therapy, supportive therapy and organ replacement therapy. As seen in other literatures, the risk of complications is also higher in critical patients, including respiratory failure, acute respiratory distress syndrome (ARDS), secondary infections, myocardial injury, liver and kidney dysfunction, etc. 9,10. This is consistent with our observations during clinical process.

There are several clinical manifestations worth mentioning. For example, dyspnea is not the most significant symptom of COVID-19, but it is relatively prominent in critical patients, so it may be a suggestive clue. Critical patients have significantly lower lymphocytes compared with non-critical ones, which is consistent with autopsy results 11, indicating a more severe bone marrow suppression and lymphocyte failure 12. In addition, worse coagulation function is more likely to occur in critical patients, and the disseminated intravascular coagulation (DIC) indicators like platelet count, PT, and D-dimer have changed significantly. Acrotic gangrene was observed in several patients (not shown in the data), indicating a potential DIC. By comparing the pathogenesis of SARS, MERS, and other viral pneumonias, we find that some critical patients may need low-molecular-weight heparin anticoagulant therapy 13. During our clinical practice, the gangrene in critical patients did significantly improve after low-molecular-weight heparin treatment. Also, data analysis also provided supportive evidence that anticoagulation may be beneficial for patient outcomes.

**Table 4.** Risk factors associated with in-hospital mortality. Note: for univariant, P values are from Log rank test, for multivariant, P values are from Cox regression. HR hazard ratio, CI confidence interval, IVIG intravenous immunoglobin, CRRT continuous renal replacement therapy, ECMO extracorporeal membrane oxygenation.

| Risk Factors               | Univariate survival analysis | Multivariate survival analysis |
|----------------------------|------------------------------|--------------------------------|
|                           | p-value                      | Hazard Ratio (95% CI)         | p-value |
| Age                       | –                            | 1.05 (1.02–1.09)             | 0.002   |
| Sex                       |                              |                               |         |
| Male                      | 0.039                        | 2.01 (0.935–4.334)            | 0.074   |
| Female                    | Referee                      |                               |         |
| Critical/non-critical      | <0.001                       | 25.70 (5.51–119.92)           | <0.001  |
| Antibiotics               | 0.572                        | 0.405 (0.18–0.91)             | 0.029   |
| Antiviral                 | 0.341                        |                               |         |
| Anticoagulation            | 0.226                        | 0.465 (0.22–0.99)             | 0.048   |
| Corticosteroid            | 0.016                        |                               |         |
| IVIG                      | 0.208                        |                               |         |
| Anti-IL6 therapy          | 0.148                        |                               |         |
| CRRT                      | 0.285                        |                               |         |
| High flow oxygen          | 0.313                        |                               |         |
| Non-invasive ventilation  | 0.014                        |                               |         |
| Invasive ventilation      | <0.001                       |                               |         |
| ECMO                      | 0.326                        |                               |         |
| Prone ventilation         | 0.490                        |                               |         |

**Discussion**

COVID-19 has now become a global pandemic, and medical systems in many countries are facing serious challenges. Survival of critical and non-critical patients are significantly different, so the early identification of critical patients is very important. This retrospective research shows that critical patients have more severe clinical situation and worse prognosis, which requires medical support of high grade, including higher level of oxygen therapy, supportive therapy and organ replacement therapy. As seen in other literatures, the risk of complications is also higher in critical patients, including respiratory failure, acute respiratory distress syndrome (ARDS), secondary infections, myocardial injury, liver and kidney dysfunction, etc. 9,10. This is consistent with our observations during clinical process.

There are several clinical manifestations worth mentioning. For example, dyspnea is not the most significant symptom of COVID-19, but it is relatively prominent in critical patients, so it may be a suggestive clue. Critical patients have significantly lower lymphocytes compared with non-critical ones, which is consistent with autopsy results 11, indicating a more severe bone marrow suppression and lymphocyte failure 12. In addition, worse coagulation function is more likely to occur in critical patients, and the disseminated intravascular coagulation (DIC) indicators like platelet count, PT, and D-dimer have changed significantly. Acrotic gangrene was observed in several patients (not shown in the data), indicating a potential DIC. By comparing the pathogenesis of SARS, MERS, and other viral pneumonias, we find that some critical patients may need low-molecular-weight heparin anticoagulant therapy 13. During our clinical practice, the gangrene in critical patients did significantly improve after low-molecular-weight heparin treatment. Also, data analysis also provided supportive evidence that anticoagulation may be beneficial for patient outcomes.
In addition to the risk factors like elder age that have been reported, we have found that high qSOFA scores and IL-6 elevation may help to identify critical patients. The qSOFA score can make quick evaluation of patient condition based on vital signs and consciousness. Although previous research revealed that qSOFA score may have limited utility for predicting mortality in an ICU setting compared with SOFA score or SIRS criteria, we believe that its convenience may have great value for practical application in current situation especially for primary medical institutions and emergency with insufficient medical resources. The elevation in IL-6 suggests the possible role of cytokine storm in the progression of COVID-19 and potential therapeutic targets.

Treatment is another field we paid attention to, although no specific treatment has been proven effective. We mainly provided supportive treatment according to clinical symptoms. The use of more antibiotics in critical patients, combined with relatively high neutrophil count, indicates more possibility of secondary opportunistic or drug-resistant infections during the long period of disease and mechanical ventilation. Pathological examination also confirmed that bacterial infection is an important pathological process which may aggravate alveolar injury and ventilation dysfunction among dead patients. Multivariate regression analysis showed that positive antibiotic usage may be beneficial for patient outcomes. Glucocorticoid is another controversy. Although it can be used as an anti-shock and anti-inflammatory agent, the use of glucocorticoid may contribute to infection. The experience of SARS has deepened our understanding of the role of glucocorticoid in severe viral pneumonia, but the balance between the suppressive effects of glucocorticoid on immunity and the positive effects of its anti-inflammatory role requires further research.

This study will provide possible supportive evidence for potential treatments by comparing effect made by different treatments on patients’ survival curves. Antiviral drugs, anti-IL-6 therapy and proper glucocorticoid usage may all have potential therapeutic effects, as many clinical trials are still ongoing. We are urgently expecting some promising results, which is of vital importance in clinical course.

Our study has some limitations. First, only 90 patients with confirmed COVID-19 were included; the suspicious and undiagnosed cases were excluded in the analyses. It would be better to absorb more patients to gain a comprehensive understanding of 2019-nCoV. Second, the lack of availability to some medical records limited our analyses of certain data. Some blood tests have not been performed in all patients for realistic reasons. In addition, since some of the critical patients were transferred to our hospital in urgent need for medical support due to the outbreak of COVID-19, it is difficult to evaluate the effect of previous treatment, and may lead to some unknown bias to sample selection. Moreover, this study is a retrospective case–control design of a single center and a rather small sample size may limit our selection of potential risk factors in the multiple regression analysis to some degree. The possibility of selection bias may exist and the results need careful interpretation.

In conclusion, this study compared the clinical characteristics between critical and non-critical COVID-19 inpatients, and qSOFA score and elevation of IL-6 are risk factors for critical condition. In multivariate survival analysis, the treatment of antibiotics and anticoagulation were significant factors for in-hospital mortality.

Methods
Study design and participants. This study is a retrospective case–control study, including adult (≥18 years old) inpatients hospitalized at Tongji Hospital (Wuhan, China) from Jan 9th to Feb 28th. All adult patients who were diagnosed with COVID-19 according to WHO interim guidance were screened. All patients were discharged or died before the date of data collection. These patients were divided into critical and non-critical group which include mild, ordinary and severe patients. The diagnosis was made according to the diagnosis and treatment guidelines (7th ed) released by China Health and Medical Commission. Mild patients are defined as patients with minor clinical symptoms and no imaging manifestations. Ordinary patients are defined as patients with typical clinical and imaging manifestations. Severe patients are defined as patients meeting at least one of the following criteria: respiratory rate >30/min, SpO2 less than 93% at rest, PaO2/FiO2 less than 300 mmHg, rapidly progressive lung imaging lesions. Critical patients are defined as patients who need mechanical ventilation or shock or have other organ failure.

The criteria for discharge were absence of fever for at least 3 days, substantial improvement in both lungs in chest CT, clinical remission of respiratory symptoms, and two throat-swab samples negative for SARS-CoV-2 RNA obtained at least 24 h apart.

Personal identifiable information was removed from all cases during the study to protect privacy. The study was approved by the Research Ethics Commission of Tongji Hospital (Wuhan, China) and the requirement for informed consent was waived by the Ethics Commission. All methods were performed in accordance with the relevant guidelines and regulations.

Data collection. Epidemiological, demographic, clinical, laboratory, treatment, and outcome data were extracted from electronic medical records in the hospital form by two physicians independently using a standardized collection. The data consistency and accuracy were checked by a third researcher.

Laboratory procedures. Methods for laboratory confirmation of COVID-19 have been described elsewhere. According to the latest guideline, COVID-19 can be diagnosed by either serum antibody or nucleic acid detection. The SARS-CoV-2 RNA was detected by Centers for Disease Control and Prevention, and the detection of serum antibody was done by local health institutions.

Routine blood examinations were complete blood count, coagulation profile, serum biochemical tests (including renal and liver function, creatine kinase, lactate dehydrogenase, and electrolytes), myocardial enzymes, inflammatory factors (including high-sensitivity C-reactive protein (hsCRP), interleukin-1 (IL-1), interleukin-2 (IL-2), interleukin-6 (IL-6), interleukin-8 (IL-8), interleukin-10 (IL-10) tumor necrosis factor α (TNF α) and...
serum ferritin), and procalcitonin. Baseline examination data were obtained for all patients. Frequency of examinations was determined by the treating physician according to disease progression.

**Statistical analysis.** The continuous data were expressed by mean (SD) or median (IQR) depending on whether they are normally distributed. They were tested by t-test or Mann-Whitney U test depending on normal distribution and homogeneity in variance. The Shapiro–Wilk test was used to test whether the continuous data were normally distributed. Categorical data were expressed by number (percentage), tested by Chi-square or Fisher’s exact test. Multivariate analysis used logistic regression. For survival analysis, univariate analysis was Kaplan–Meier analysis, and multivariant analysis was COX regression. The significance was defined as p value below 0.05. All the data analysis was conducted in SPSS 21.0 (IBM Corporation, Armonk, NY, USA).

**Data availability**

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

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**Author contributions**

R.T., W.W., C.W., Q.L., P.G., S.Z. collected all the clinical data and contributed to the structure and writing of the discussion part. H.P., Z.Z. and H.X. analyzed and interpreted the patient data and were also major contributor in writing the manuscript. J.S., W.L., H.Q. and F.G. contributed in patient screening, data collection and manuscript proofreading. T.L., Z.L., J.W., X.Z., Y.Q. and X.Y. provided professional opinions regarding infectious disease, respiratory disease and critical care. They also contributed to the background and interpretation of the data. All authors read and approved the final manuscript.

**Competing interests**

The authors declare no competing interests.

**Additional information**

**Correspondence** and requests for materials should be addressed to S.Z.

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