Research Article

Clinical Characteristics of Diabetic Patients with COVID-19

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Background. Since December 2019, novel coronavirus (SARS-CoV-2-) infected pneumonia (COVID-19) has rapidly spread throughout China. This study is aimed at describing the characteristics of COVID-19 patients in Wuhan. Methods. 199 COVID-19 patients were admitted to Wuhan Red Cross Hospital in China from January 24th to March 15th. The cases were divided into diabetic and nondiabetic groups according to the history of taking antidiabetic drugs or by plasma fasting blood glucose level at admission, and the difference between groups were compared. Results. Among 199 COVID-19 patients, 76 were diabetic and 123 were nondiabetic. Compared with nondiabetics, patients with diabetes had an older age, high levels of fasting plasma glucose (FPG), D-dimer, white blood cell, blood urea nitrogen (BUN) and total bilirubin (TBIL), lower levels of lymphocyte, albumin and oxygen saturation (SaO2), and higher mortality ($P < 0.05$). The two groups showed no difference in clinical symptoms. Diabetes, higher level of D-dimer at admission, and lymphocyte count less than $0.6 \times 10^9 / L$ at admission were associated with increasing odds of death. Antidiabetic drugs were associated with decreasing odds of death. Treatment with low molecular weight heparin was not related to odds of death. Conclusion. The mortality rate of COVID-19 patients with diabetes was significantly higher than those without diabetes. Diabetes, higher level of D-dimer, and lymphocyte count less than $0.6 \times 10^9 / L$ at admission were the risk factors associated with in-hospital death.

1. Introduction

Since December 2019, novel coronavirus (SARS-CoV-2-) infected pneumonia (COVID-19) has rapidly spread throughout China and around the world [1–4]. The International Committee on Taxonomy of Viruses (ICTV) has named this virus SARS-CoV-2, with the disease termed COVID-19 [5]. The high infectivity of COVID-19 resulted in a rapid increase of new cases. Previous studies have described the epidemiological findings, clinical presentation, and clinical outcomes of patients with confirmed COVID-19 [6, 7]. However, specific information of patients with diabetes with COVID-19 remains unknown.

Diabetes mellitus (DM) is often identified as an independent risk factor for developing respiratory tract infections [8]. Studies have reported the relationship between blood glucose levels and the clinical course of severe acute respiratory syndrome (SARS) [9]. Up to now, information regarding the clinical characteristics of patients with diabetes with 2019 novel SARS-COV-2 pneumonia was scarce. In this study, the aim was to determine clinical symptoms, laboratory findings, and mortality of patients with diabetes and patients without diabetes in COVID-19, and to report on any difference.

2. Methods

2.1. Patients. The retrospective study was approved by the ethics committee of Wuhan Red Cross Hospital (No. 2020022). All patients with COVID-19 admitted to Wuhan Red Cross Hospital from January 24th to March 15th were enrolled. During this period, Wuhan Red Cross Hospital became a special designated hospital for the treatment of patients with COVID-19.
2.2. Definitions. COVID-19 was confirmed by detecting SARS-CoV-2 RNA in throat swab samples using a virus nucleic acid detection kit according to the manufacturer’s protocol (Shanghai BioGerm Medical Biotechnology Co., Ltd). All of the patients were admitted to the general fever ward excluding the intensive care unit. The cases were divided into diabetic and nondiabetic groups according to the history of taking anti-diabetic drugs or by plasma fasting blood glucose level at admission.

2.3. Data Collection. The case report form of COVID-19 was designed to document primary clinical data regarding previous medical history, clinical symptoms, laboratory findings, and clinical outcomes from electronic medical records. The following information was extracted for each patient: gender, age, medical history, clinical outcomes, and signs, symptoms, oxygen saturation, and laboratory findings at admission.

2.4. Data Analysis. Categorical data were described as percentages, and continuous data as median with interquartile range (IQR). χ² test for categorical data and Mann-Whitney U test for continuous data were used to compare variables between groups. All statistical analyses were performed using SPSS Statistics version 16.0 software. P < 0.05 was considered statistically significant.

3. Results

3.1. Baseline Characteristics. 199 patients with COVID-19 pneumonia were included in our study (Table 1). Among them, 76 were diabetic and 123 were nondiabetic. The median age of patients with COVID-19 pneumonia with diabetes was 67 years (IQR: 61–78), which was significantly higher than that of patients without diabetes (IQR: 47–68, P ≤ 0.001). The onset of initial symptom to hospital admission in patients with diabetes group was 10 days (IQR: 5–14), while patients without diabetes group was 12 days (IQR: 7–20, P = 0.036). Duration of hospital stays was similar for both groups (P = 0.968). Of 199 patients, fever (74.4%), cough (66.8%), dyspnea (26.8%), and fatigue (25.1%) were the most common symptoms, while diarrhea (3%), vomiting (2.5%), and dizziness (1%) were scarce. However, no statistically significant difference in all of clinical symptoms between diabetic and nondiabetic patients with COVID-19 pneumonia.

| Table 1: Baseline characteristics of patients with COVID-19 pneumonia admission to hospital. |
|---------------------------------|-----------------|-----------------|-----------------|-----------------|
|                                | Total (n=199)   | Diabetic (n=76) | Nondiabetic (n=123) | P Value        |
| Age, median (IQR), years       | 63 (50-75)      | 67 (61-78)      | 59 (47-68)       | ≤0.001         |
| Sex                             |                |                 |                  |                |
| Female                          | 110 (55.3)     | 37 (48.7)       | 73 (59.3)       | 0.142          |
| Male                            | 89 (44.7)      | 39 (51.3)       | 50 (40.7)       |                |
| Symptoms                        |                |                 |                  |                |
| Fever                           | 148 (74.4)     | 54 (71.1)       | 94 (76.4)       | 0.399          |
| Cough                           | 133 (66.8)     | 54 (71.1)       | 79 (64.2)       | 0.320          |
| Dyspnea                         | 53 (26.6)      | 22 (28.9)       | 31 (25.2)       | 0.562          |
| Fatigue                         | 50 (25.1)      | 16 (21.1)       | 34 (27.6)       | 0.289          |
| Anorexia                        | 32 (16.1)      | 9 (11.8)        | 23 (18.7)       | 0.201          |
| Nausea                          | 23 (11.6)      | 5 (6.6)         | 18 (14.6)       | 0.084          |
| Headache                        | 20 (10.1)      | 5 (6.6)         | 15 (12.2)       | 0.410          |
| Diarrhea                        | 6 (3.0)        | 1 (1.3)         | 5 (4.1)         | 0.651          |
| Vomiting                        | 5 (2.5)        | 1 (1.3)         | 3 (3.3)         | >0.999         |
| Dizziness                       | 2 (1.0)        | 1 (0.8)         | 1 (1.3)         |                |
| Onset of initial symptom to hospital admission (days), (IQR) | 10 (6-15) | 10 (5-14) | 12 (7-20) | 0.036 |
| Duration of hospital stay (days), (IQR) | 13 (8-20) | 13 (9-19) | 12 (7-22) | 0.968 |
| Clinical outcome                |                |                 |                  |                |
| Died                            | 18 (9.0)       | 11 (14.5)       | 7 (5.7)         | 0.036          |
| Alive                           | 181 (91.0)     | 65 (86.5)       | 116 (94.3)      |                |

COVID-19: coronavirus disease 2019; IQR: interquartile range; no: number.
level of oxygen saturation ($P \leq 0.001$), lymphocyte ($P = 0.036$), and albumin ($P = 0.002$).

### 3.3. Regression Analysis

We included 199 patients with complete data for all variables (181 survivors and 18 nonsurvivors) in the multivariable logistic regression model. We found that diabetes, higher level of D-dimer at admission, and lymphocyte count less than $0.6 \times 10^9/L$ at admission were associated with increasing odds of death. Antidiabetic drugs were associated with decreasing odds of death. Treatment with low molecular weight heparin was not related to odds of death (Table 3).

### 3.4. Clinical Outcome

There were 18 reported deaths in the COVID-19 patients (Table 1). Significant difference in mortality was found between patients with diabetes and without diabetes ($P = 0.036$). 11 of 76 patients with COVID-19 with diabetes died (14.5%), while 7 of 123 patients with COVID-19 without diabetes died (5.7%). Diabetes seems to increase the risk of death in patients with COVID-19 pneumonia (Log rank $P = 0.031$) (Figure 1).

### 4. Discussion

Coronavirus has received more attention compared to other causes of pneumonia, especially after the emergence of SARS and MERS. In certain risk factors, clinical manifestations, and clinical outcomes, COVID-19 was similar to SARS and MERS. It had been reported that a known history of diabetes was independent predictors for morbidity and death in patients with SARS [9]. In our study, 11 (14.5%) patients with COVID-19 pneumonia with diabetes died, while 7 (5.7%) patients with COVID-19 pneumonia without diabetes died ($P = 0.036$). Diabetes was associated with increasing odds of death. Antidiabetic drugs were associated with decreasing odds of death. Until now, large-scale analyses of clinical characteristics and outcome of patients with COVID-19 pneumonia with diabetes had been scarce. In this study, 199 COVID-19 patients were divided into diabetic and nondiabetic groups. We compared the clinical features, laboratory findings, and clinical outcome between the two groups.

The study found similar proportions of male and female patients in COVID-19 with and without diabetes.
The severity of lymphocytopenia may relate to COVID-19 at admission had only mild lymphocytopenia in patients with COVID-19. In our study, patients with diabetes had a lower level of lymphocyte. Lymphocytopenia is a prominent feature of critically ill patients with COVID-19 because targeted invasion by SARS-CoV viral particles damages the cytoplasmic component of the lymphocyte and causes its destruction [17]. Hence, we speculate that necrosis or apoptosis of lymphocytes also induces lymphocytopenia in critically ill patients with COVID-19. In our study, patients with diabetes with COVID-19 had signifi-
cantly higher than that of patients without diabetes. Diabetes, higher level of D-dimer, and lymphocyte count less than
7.6×10^9/L at admission were risk factors associated with death of adult patients with COVID-19 [7, 12, 20]. We
found that patients with COVID-19 with higher level of D-dimer at admission, especially those with diabetes,
are significantly associated with the risk of death. Treatment with low molecular weight heparin was not related
to odds of death. Magro et al. reported that severe COVID-19 infection was associated with microvascular
injury and thrombosis [21]. There is a need for further clinical trials using anticoagulants to determine whether
the application of anticoagulants is effective.

In conclusion, the mortality rate for diabetic patients with COVID-19, D-dimer levels increased dramatically. D-
dimer is an activation marker of fibrinolysis. Some studies have shown that D-dimer is a significant prognostic factor
in patients with pneumonia and sepsis [18, 19]. D-dimer is a marker of mortality in patients admitted to the emer-
gency department with suspected infection and sepsis [19]. In recent studies, the level of D-dimer was higher
in the death group than in the survival group, and elevated levels of D-dimer at admission were risk factors
for death of adult patients with COVID-19 [7, 12, 20].

Conflicts of Interest

The authors declare that they have no conflicts of interest.

Data Availability

The Excel data used to support the findings of this study are available from the corresponding author upon request.
Authors’ Contributions

GZL, QD, JLF, and FL collected the data. GZL, NX, and HQ prepared and revised the manuscript. GZL, NX, and QH were responsible for summarizing all data related to this study. Guozen Li and Qin Deng contributed equally to this work.

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