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Haoxiang Li

Affiliated Hospital of Jiangsu University

Jianguo Zhang

Affiliated Hospital of Jiangsu University

Jinhui Zhang

Affiliated Hospital of Jiangsu University

Ling Yang

Affiliated Hospital of Jiangsu University

Dong Wang

Affiliated Hospital of Jiangsu University

Li Zhao

Affiliated Hospital of Jiangsu University

Xia Deng

Affiliated Hospital of Jiangsu University

Guoyue Yuan (✉ yuanguoyue08@163.com)

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**Clinical Characteristics and Prognosis of patients with COVID-19 combined
with or without diabetes, hypertension or coronary**

Haixiang Li¹, Jianguo Zhang², Jinhui Zhang², Ling Yang¹, Dong Wang¹, Li Zhao¹,
Xia Deng¹, Guoyue Yuan¹

(1. Department of Endocrinology, 2. Department of Intensive Care Unit, Affiliated
Hospital of Jiangsu University, 438, Jiefang Road, Zhenjiang, Jiangsu 212001,
China)

corresponding author:

Guoyue Yuan, M.D, PhD,

Department of Endocrinology, Affiliated Hospital of Jiangsu University, 438, Jiefang
Road, Zhenjiang, Jiangsu 212001, China

Mobile: +8613505289352, Fax number: 86-511-85019237

Email: yuanguoyue@hotmail.com, yuanguoyue08@163.com

Abstract

Background: This study was to investigate the clinical characteristics and prognosis of COVID-19 patients combined with or without major chronic diseases like diabetes, hypertension or coronary.

Methods: We retrospectively analyzed 183 patients with COVID-19 diagnosed at First People's Hospital of Jiangxia District (FPHJD) in Wuhan, China attended by Affiliated Hospital of Jiangsu University supporting medical team from February 1, 2020 to March 15, 2020. Patients were divided into simple COVID-19 group(n=134), COVID-19 combined with diabetes, hypertension or coronary group(n=49). Besides, COVID-19 patients with diabetes, hypertension or coronary were further classified into severe pneumonia group(n=23) and common pneumonia group(n=26), death group(n=17) and survival group(n=32). The prognosis of COVID-19 patients was evaluated by analyzing the clinical data and the results of laboratory tests.

Results: 183 patients were included in this study, of whom 166 were discharged and 16 died in hospital. 49 (26.92%) patients had a comorbidity, with hypertension being the most common [37 (20.33%) patients], followed by diabetes [25 (13.74%) patients] and coronary heart disease [4 (2.2%) patients]. Compared with simple COVID-19 group, the proportion of history of chronic respiratory system disease, age, D-dimer, procalcitonin, C-reactive protein, myoglobin, cardiac troponin I, creatine kinase MB, lactate dehydrogenase, white blood cell count, neutrophil count, neutrophil percentage, blood urea nitrogen, creatinine and mortality rate were significantly higher in COVID-19 combined with chronic diseases group, whereas lymphocyte count, lymphocyte percentage and alanine transferase were significantly lower in COVID-19 combined with chronic diseases group. Among COVID-19 patients with chronic diseases, D-dimer, procalcitonin, C-reactive protein, myoglobin, cardiac troponin I, lactate dehydrogenase, white blood cell count, neutrophil count, neutrophil percentage, blood urea nitrogen, death rate was significantly higher in severe pneumonia group than common pneumonia group. While lymphocyte count and lymphocyte percentage were significantly lower in severe pneumonia group than common pneumonia group. Besides, we found that the proportion of history of chronic respiratory system disease,

D-dimer, procalcitonin, myoglobin, cardiac troponin I, creatine kinase MB, lactate dehydrogenase, neutrophil count, neutrophil percentage, blood urea nitrogen were significantly higher in death group compared with survival group, whereas lymphocyte count and lymphocyte percentage were significantly lower in survival group. In COVID-19 combined with chronic diseases group, univariate logistic regression showed that the risk for severe pneumonia were D-dimer, C-reactive protein, lactate dehydrogenase, white blood cell count, neutrophil count and neutrophil percentage. Univariate logistic regression also showed that the risk for death were D-dimer, lactate dehydrogenase, white blood cell count, neutrophil count, neutrophil percentage and blood urea nitrogen. Multivariate regression logistic showed that lactate dehydrogenase were independent risk factors for death among COVID-19 patients combined with chronic diseases. Cox regression analysis showed that compared with simple COVID-19 group, the RR(95% CI) in COVID-19 patients combined with diabetes, hypertension, and coronary were 2.187 (1.141~4.191) for death ($P<0.05$).

Conclusion: Among COVID-19 patients combined with diabetes, hypertension or coronary, the risk factors for severe pneumonia were D-dimer, C-reactive protein, lactate dehydrogenase, white blood cell count, neutrophil count and neutrophil percentage, whereas the risk factors for death were D-dimer, lactate dehydrogenase, white blood cell count, neutrophil count, neutrophil percentage and blood urea nitrogen. Moreover, lactate dehydrogenase were independent risk factors for death. The mortality rate of COVID-19 patients combined with diabetes, hypertension or coronary was higher than that of simple COVID-19 patients.

Key words: COVID-19; diabetes; hypertension; coronary

Background: Coronavirus Disease 2019 (COVID-19) has quickly progressed to a global health emergency. COVID-19 is a new type of coronavirus which belongs to the Betacoronavirus genera with the capacity for rapid mutation and recombination.[1]. The main clinical symptoms of COVID-19 are fever, fatigue, and a dry cough, and in severe cases, multiple organ failure[2-3]. COVID-19 has numerous transmission channels and humans are highly susceptible to infection. Cases of COVID-19 have been found across the whole of China and overseas[4]. Recent studies have found that COVID-19 patients with chronic diseases like diabetes, hypertension and coronary have higher critical illness rate and mortality rate[5-6]. However, the risk factors for mortality and a detailed clinical course of illness have not been well described. This paper reviews and summarizes the epidemiological and clinical characteristics of COVID-19 patients combined with or without diabetes, hypertension or coronary in order to provide a reliable basis for early diagnosis and treatment.

Methods

Study population

This retrospective cohort study included patients with COVID-19 diagnosed at the First People's Hospital of Jiangxia District (FPHJD) in Wuhan, China attended by Affiliated Hospital of Jiangsu University supporting medical team from February 1, 2020 to March 15, 2020. In this study, patients who were hospitalised for COVID-19 and had a definite outcome (dead or discharged) at the early stage of the outbreak. The study was approved by the Research Ethics Commission of FPHJD (Approval No. 2020021) and written consent was waived by Ethics Commission of FPHJD.

Data collection

After taking medical history, necessary investigations like blood examinations which included complete blood count, serum biochemical tests (including renal and liver function, creatine kinase, creatine kinase MB and lactate dehydrogenase), myocardial enzymes(including myoglobin and cardiac troponin I), D-dimer, C-reactive protein

and procalcitonin were performed upon hospital admission. All data were checked by two physicians (JZ and JZ).

Definitions

All patients were up to 18 years old and received throat swab samples which were gathered for SARS-CoV-2 RNA detection by gene sequencing or real-time RT-PCR and the results were positive at least once. Diabetes mellitus was defined as a medical history of diabetes or the use of oral hypoglycemic medication or insulin or patients with a fasting glucose ≥ 7.0 mmol/L or a two-hour postprandial serum glucose ≥ 11.1 mmol/L[7]. Hypertension was defined as a medical history of hypertension or the use of antihypertensive, a systolic blood pressure of ≥ 140 mmHg and/or a diastolic blood pressure of ≥ 90 mmHg[8]. Coronary diagnosis was defined as ischemic symptoms or coronary computed tomographic angiography (CTA) or percutaneous coronary intervention (PCI) and defined as angiographic evidence of more than 50% luminal narrowing in at least one segment of a main epicardial coronary artery[9]. The illness severity of pneumonia was defined according to the Chinese management guideline for COVID-19 (version 6.0)[10]. All the subjects were excluded from malignancy, pregnancy, blood disease, autoimmune disease and patients who failed to complete relevant blood examinations.

Statistical Analysis

All statistical analyses were performed using SPSS version 16.0 (SPSS Inc., Chicago, IL). Data were summarized as means \pm standard deviations for normally distributed variables, medians plus percentiles (25th; 75th) for nonnormally distributed variables, and frequencies for categorical variables. For comparisons between two groups, independent Student t test was used. Categorical variables were examined by χ^2 test. To explore the risk factors associated with severe pneumonia and death, univariable and multivariable logistic regression models were used. To identify the chronic diseases affecting the mortality rate, we used a multivariate Cox regression analysis

model. All calculated *p* values were two-sided, and *p* values < 0.05 were considered statistically significant.

Results

Clinical and biochemical characteristics between simple COVID-19 group and COVID-19 combined with chronic diseases group

Compared with simple COVID-19 group, the proportion of history of chronic respiratory system disease, age, D-dimer, procalcitonin, C-reactive protein, myoglobin, cardiac troponin I, creatine kinase MB, lactate dehydrogenase, white blood cell count, neutrophil count, neutrophil percentage, blood urea nitrogen, creatinine and mortality rate were significantly higher in COVID-19 combined with chronic diseases group, whereas lymphocyte count, lymphocyte percentage and alanine transferase were significantly lower in COVID-19 combined with chronic diseases group (Table 1).

Table 1: Clinical and biochemical findings of patients on admission

	simple COVID-19 group (n=134)	COVID-19 combined with chronic diseases group (n=49)	<i>P</i> value
Sex(F/M)	65/69	30/19	0.127
history of chronic respiratory system disease(yes/no)	1/134	3/46	0.022*
Age(year)	50.37±16.45	66.49±12.24	<0.001*
D-dimer, mg/L	0.37(0.19,0.84)	0.67(1.26,1.89)	0.009*
Procalcitonin, ng/ml	0.06(0.04,0.13)	0.10(0.05,0.25)	0.005*
C-reactive protein, mg/L	10.39(1.16,46.96)	40.49(12.93,85.15)	0.001*
myoglobin, U/L	25.82(21.00,55.43)	49.37(25.60,217.25)	<0.001*
cardiac troponin I, µg/L	0.01(0.01,0.01)	0.02(0.01,0.03)	<0.001*
creatinine kinase MB	13.55(10.38,17.23)	15.9(11.10,26.25)	0.064
creatinine kinase, U/L	72.60(44.50,122.00)	77(50.50, 146.00)	0.668
lactate dehydrogenase, U/L	215.30(179.50,333.75)	266(209.30,394.70)	0.009*
white blood cell count, ×10 ⁹ /L	6.20(4.40,8.40)	7.60(5.30,11.95)	0.007*
neutrophil count, ×10 ⁹ /L	4.44(2.75,6.32)	5.89(4.07,10.54)	0.001*
neutrophil percentage, %	70.59±13.99	80.27±9.52	<0.001*
lymphocyte count, ×10 ⁹ /L	1.06(0.71,1.58)	0.73(0.55,1.19)	0.003*
lymphocyte percentage, %	17.75(10.48,30.15)	12.20(6.87,16.55)	<0.001*
platelet count, ×10 ⁹ /L	210.00(150.75,260.50)	179.00(122.00,286.50)	0.11

aspartate aminotransferase, U/L	26.60(19.43,38.88)	26.50(20.40,38.85)	0.857
alanine aminotransferase, U/L	22.80(15.20,42.18)	18.50(12.60,31.80)	0.032*
blood urea nitrogen, mmol/L	4.45(3.40,5.50)	5.80(4.50,11.05)	<0.001*
Creatinine, μ mol/L	63.35(51.88,81.05)	73.60(60.60,104.50)	0.001*
mortality rate	21/134	16/49	<0.001*

F: female; M: male.

Data are means \pm SD, n, and median (25th and 75th percentiles).* P <0.05.

Clinical and biochemical characteristics among COVID-19 patients with chronic diseases

Among COVID-19 patients with chronic diseases, D-dimer, procalcitonin, C-reactive protein, myoglobin, cardiac troponin I, lactate dehydrogenase, white blood cell count, neutrophil count, neutrophil percentage, blood urea nitrogen and mortality rate were significantly higher in severe pneumonia group than common pneumonia group. While lymphocyte count and lymphocyte percentage were significantly lower in severe pneumonia group than common pneumonia group (Table 2). Besides, we found that the proportion of history of chronic respiratory system disease, D-dimer, procalcitonin, myoglobin, cardiac troponin I, creatine kinase MB, lactate dehydrogenase, neutrophil count, neutrophil percentage, blood urea nitrogen were significantly higher in death group compared with survival group, whereas lymphocyte count and lymphocyte percentage were significantly lower in survival group (Table 3).

Table 2: Clinical and biochemical findings between common pneumonia group and severe pneumonia group

	common pneumonia group (n=26)	severe pneumonia group (n=23)	<i>P</i> value
Sex(F/M)	14/12	16/7	0.26
history of chronic respiratory system disease(yes/no)	0/26	3/20	0.057
Age(year)	65.19 \pm 11.48	67.96 \pm 13.15	0.436
D-dimer, mg/L	0.38(0.20,0.68)	1.87(0.78,7.33)	<0.001*
Procalcitonin, ng/ml	0.06(0.03,0.14)	0.16(0.10,0.46)	<0.001*

C-reactive protein, mg/L	18.30(2.26,65.91)	60.53(37.07,106.56)	0.002*
myoglobin, U/L	39.33(22.23,84.75)	126.50(26.89,276.70)	0.044*
cardiac troponin I, µg/L	0.01(0.01,0.02)	0.02(0.01,0.18)	0.007*
creatine kinase MB	14.95(11.15,21.95)	17.10(10.60,34.60)	0.218
creatine kinase, U/L	69.80(46.00,118.50)	92.00(53.00,157.00)	0.352
lactate dehydrogenase, U/L	240.50(198.75,272.58)	446.40(226.00,611.00)	<0.001*
white blood cell count, ×10 ⁹ /L	6.55(5.35,7.98)	9.10(4.80,14.90)	0.044*
neutrophil count, ×10 ⁹ /L	4.87(4.03,6.01)	8.36(4.09,13.75)	0.013*
neutrophil percentage, %	75.00±8.38	86.23±6.95	<0.001*
lymphocyte count, ×10 ⁹ /L	1.08(0.69,1.48)	0.57(0.49,0.73)	<0.001*
lymphocyte percentage, %	16.73±6.61	7.80±4.32	<0.001*
platelet count, ×10 ⁹ /L	191.00(141.00,310.00)	161.00 (110.00,233.00)	0.087
aspartate aminotransferase, U/L	25.25(19.25,36.43)	28.60 (20.30,44.60)	0.435
alanine aminotransferase, U/L	20.65(13.75,31.50)	16.60 (11.60,33.90)	0.561
blood urea nitrogen, mmol/L	5.20(4.15,8.13)	7.60 (5.50,13.10)	0.014*
Creatinine, µmol/L	68.60(55.83,97.93)	92.60(63.70,111.10)	0.103
mortality rate	0/26	16/23	<0.001*

F: female; M: male.

Data are means±SD, n, and median (25th and 75th percentiles).* *P*<0.05.

Table 3: Clinical and biochemical findings between survival group and death

	group		<i>P</i> value
	survival group (n=32)	death group (n=17)	
Sex(F/M)	19/13	11/6	0.715
history of chronic respiratory system disease(yes/no)	0/32	3/14	0.014*
Age(year)	66.22±12.25	67.00±12.58	0.834
D-dimer, mg/L	0.50(0.20,0.74)	1.93(0.82,7.70)	0.001*
Procalcitonin, ng/ml	0.07(0.04,0.18)	0.15(0.09,0.47)	0.015*
C-reactive protein, mg/L	30.16(4.70,89.75)	58.99(28.59,81.11)	0.118
myoglobin, U/L	40.43(23.70,117.20)	149.00(31.97,373.30)	0.049*
cardiac troponin I, µg/L	0.01(0.01,0.02)	0.02(0.02,0.55)	0.001*
creatine kinase MB	13.30(11.05,21.83)	23.20(11.85,35.65)	0.048*
creatine kinase, U/L	74.50(49.75,137.00)	86.00(47.00,167.00)	0.698
lactate dehydrogenase, U/L	248.00(208.95,280.03)	493.00(244.00,627.50)	0.002*
white blood cell count, ×10 ⁹ /L	6.65(5.25,9.08)	8.80(6.10,15.40)	0.095
neutrophil count, ×10 ⁹ /L	5.15(4.00,7.89)	7.36(5.02,14.11)	0.034*
neutrophil percentage, %	77.05±9.11	86.32±7.19	0.001*
lymphocyte count, ×10 ⁹ /L	0.85(0.58,1.41)	0.06(0.49,0.75)	0.004*
lymphocyte percentage, %	15.18±7.02	7.56±4.37	<0.001*

platelet count, $\times 10^9/L$	184.50(139.00,301.75)	146.00(102.00,212.00)	0.106
aspartate aminotransferase, U/L	24.90(19.55,35.70)	34.30(23.90,55.20)	0.106
alanine aminotransferase, U/L	19.80(13.45,30.60)	16.60(12.10,16.20)	0.975
blood urea nitrogen, mmol/L	5.30(4.23,9.88)	7.00(5.55,16.20)	0.02*
Creatinine, $\mu\text{mol/L}$	69.20(56.40,95.80)	96.10(62.55,138.10)	0.057

F: female; M: male.

Data are means \pm SD, n (%), and median (25th and 75th percentiles). * $P < 0.05$.

Univariate and multivariate analysis of the risk factors for severe pneumonia and death in COVID-19 patients combined with chronic diseases

The distribution of COVID-19 patients combined with diabetes, hypertension or coronary was showed in Table 4. We calculated the individual comorbidity percentage in common pneumonia group and severe pneumonia group, survival group and death group, and plotted them versus each other as shown in Figure 1 and Figure 2. In COVID-19 combined with chronic diseases group, univariate logistic regression showed that the risk for severe pneumonia were D-dimer, C-reactive protein, lactate dehydrogenase, white blood cell count, neutrophil count and neutrophil percentage(Table 5). Univariate logistic regression also showed that the risk for death were D-dimer, lactate dehydrogenase, white blood cell count, neutrophil count, neutrophil percentage and blood urea nitrogen(Table 6). Multivariate logistic regression showed that lactate dehydrogenase were independent risk factors for death among COVID-19 patients combined with chronic diseases(Table 7).

Table 4: Distribution of COVID-19 patients combined with diabetes, hypertension or coronary

Items	Death group(n=16)	Suivival group(33)
Diabetes	3	9
Hypertension	4	18
Coronary	0	0
Diabetes & hypertension	7	4
Diabetes & coronary	0	0

Hypertension & coronary	2	0
Diabetes & Hypertension & coronary	0	2

Table 5: Univariate logistic regression analysis of the risk factors for severe pneumonia in COVID-19 patients combined with chronic diseases

variable	β	SE	Wald	<i>P</i> value	OR	95% CI
D-dimer	0.513	0.223	5.318	0.021*	1.671	1.080~2.584
C-reactive protein	0.019	0.007	6.553	0.01*	1.019	1.004~1.034
lactate dehydrogenase	0.013	0.005	7.962	0.005*	1.013	1.004~1.023
white blood cell count	0.167	0.074	5.108	0.024*	1.181	1.022~1.365
neutrophil count	0.220	0.086	6.577	0.01*	1.246	1.053~1.473
neutrophil percentage	0.197	0.056	12.188	<0.01*	1.217	1.090~1.359
lymphocyte count	-2.844	1.046	7.391	0.007*	0.058	0.007~0.452
lymphocyte percentage	-0.343	0.096	12.703	<0.01*	0.710	0.588~0.857

* $P < 0.05$.

Table 6: Univariate logistic regression analysis of the risk factors for death in COVID-19 patients combined with chronic diseases

variable	β	SE	Wald	<i>P</i> value	OR	95% CI
D-dimer	0.344	0.127	7.296	0.007*	1.41	1.099~1.809
lactate dehydrogenase	0.009	0.003	9.581	0.002*	1.009	1.003~1.015
white blood cell count	0.128	0.064	3.974	0.046*	1.137	1.002~1.290
neutrophil count	0.159	0.071	5.039	0.025*	1.173	1.020~1.384
neutrophil percentage	0.143	0.048	8.863	0.003*	1.154	1.050~1.267
lymphocyte count	-3.334	1.358	6.031	0.014*	0.036	0.002~0.510
lymphocyte percentage	-0.253	0.079	10.285	0.001*	0.777	0.665~0.907
blood urea nitrogen	0.126	0.062	4.128	0.042*	1.135	1.004~1.281

* $P < 0.05$.

Table 7: multivariate analysis of the risk factors for death in COVID-19 patients combined with chronic diseases

variable	β	SE	Wald	<i>P</i> value	OR	95% CI
lactate dehydrogenase	0.008	0.004	4.352	0.037	1.008	1.000~1.015

Figure 1. Comorbidity percentage in common pneumonia and severe pneumonia patients was plotted versus each other. Diagonal (gray, dotted) indicated a hypothetically equal percentage between the two groups.

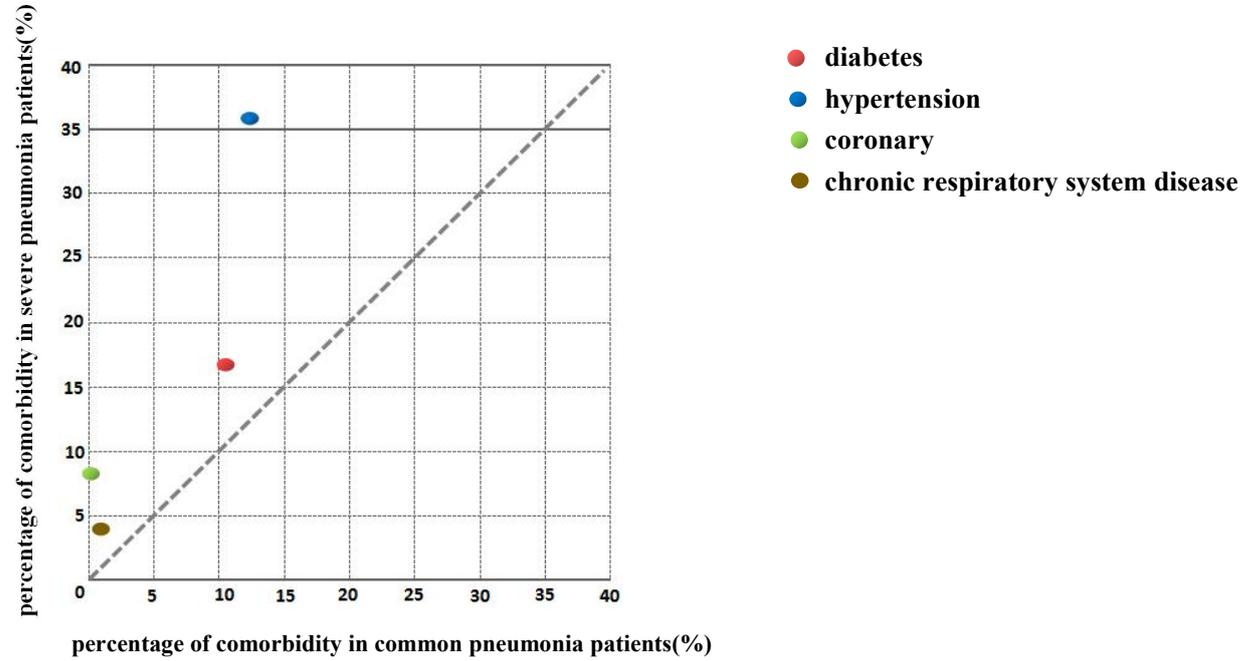
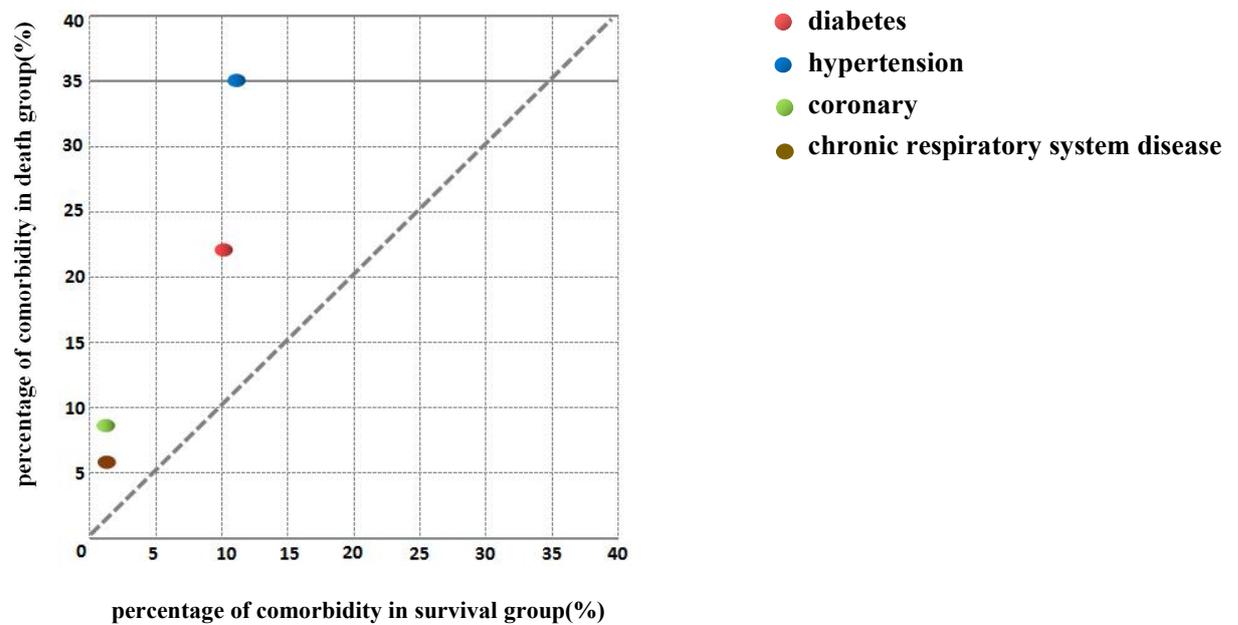


Figure 2. Comorbidity percentage in death and survival group was plotted versus each other. Diagonal (gray, dotted) indicated a hypothetically equal percentage between the two groups.

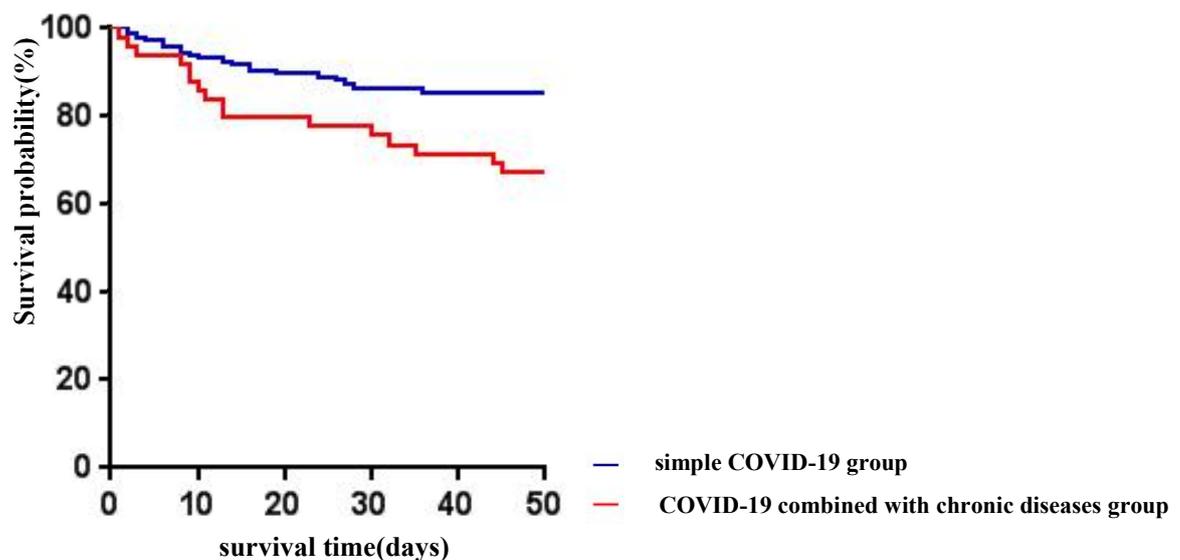


Cox regression analysis and survival curve

Cox regression analysis showed that compared with simple COVID-19 group, the RR

(95% CI) in COVID-19 combined with chronic diseases group were 2.187 (1.141~4.191) for death ($P<0.05$). The results of survival analysis revealed that the survival time of patients in the COVID-19 combined with chronic diseases group was significantly shorter than that in the simple COVID-19 group (Fig 4).

Figure 3. The survival curves of the simple COVID-19 group and COVID-19 combined with chronic diseases group



Discussion

Patients with hypertension, diabetes and coronary heart disease have worse clinical outcomes when infected with coronavirus [11-14]. From previous studies of the fatal cases of severe acute respiratory syndrome coronavirus (SARS-CoV) pneumonia, the comorbidities of hypertension, diabetes and coronary heart disease were found to be dangerous factors that resulted in death [15-16]. Also, a systematic analysis of 637 Middle East respiratory syndrome coronavirus (MERS-CoV) cases suggests that diabetes and hypertension are equally prevalent in approximately 50% of the patients while coronary heart disease are present in 30% [17]. However, the mechanisms for high morbidity and mortality of patients with comorbidities are unknown. The SARS-CoV-2, a positive strand RNA virus, has been seen to infect humans through the angiotensin converting enzyme -2 (ACE-2) receptor [18-19]. The ACE-2 receptor is a part of the dual system renin-angiotensin-system (RAS) consisting of

ACE-Ang-II-AT1R axis and ACE-2-Ang-(1-7)-Mas axis. In metabolic disorders and with increased age, it is known that there is an upregulation of ACE-Ang-II-AT1R axis with a downregulation of ACE-2-Ang-(1-7)-Mas axis. ACE-2-Ang-(1-7)-Mas axis has anti-inflammatory and antifibrotic effects on the respiratory system and anti-inflammatory, antioxidative stress, and protective effects on vascular function, protects against myocardial fibrosis, nephropathy, pancreatitis, and insulin resistance. The already strained ACE-2-Ang-(1-7)-Mas in metabolic disorders is further stressed due to the use of the ACE-2 by the virus for entry, which can affect the prognosis in terms of respiratory compromise[20]. Many scholars believe that this may be one of the possible mechanisms for the high mortality of COVID-19 in patients with underlying metabolic conditions and they believe that ACE-2-based therapy has been proposed as a potential therapeutic approach in COVID-19 pneumonia[21].

There is growing literature exploring that myocardial injury is one of the important pathogenic features of COVID-19[22-25]. In a cohort of 52 critically ill adult patients with SARS-CoV-2 pneumonia who were admitted to the intensive care unit (ICU) of Wuhan Jin Yin-tan hospital, 12 patients (23%) had cardiac injury which was defined as an elevated serum level of high-sensitivity cardiac troponin I (hs-TnI)[26]. In concert with previous studies, we found that increased cardiac biomarkers mainly myoglobin and cardiac troponins T in the COVID-19 patients combined with chronic diseases especially those with severe pneumonia or death. The proposed mechanisms of myocardial injury are direct damage to the cardiomyocytes, systemic inflammation, myocardial interstitial fibrosis, interferon mediated immune response, exaggerated cytokine response by Type 1 and 2 helper T cells, in addition to coronary plaque destabilization, and hypoxia[27-29]. We suggest that myocardial injury is a problem that cannot be ignored in patients with COVID-19. It has also been proposed that in patients with COVID-19 had increased coagulation activity, marked by increased D-dimer concentrations[30]. Study has proved that D-dimer might help in early recognition of these high-risk patients and also predict outcome[31]. We also observed that D-dimer is one of the risk factors for severe pneumonia or death among COVID-19 patients combined with chronic diseases. Contributory mechanisms

include systemic pro-inflammatory cytokine responses that are mediators of atherosclerosis directly contributing to plaque rupture through local inflammation, induction of procoagulant factors, and haemodynamic changes, which predispose to ischaemia and thrombosis[32]. However, due to the limitations of existing evidence, future research is needed to elaborate on the potential mechanisms.

In this study, lactate dehydrogenase (LDH) were independent risk factors for death among COVID-19 patients combined with chronic diseases. LDH is a cytoplasmic glycolytic enzyme found in all most every tissue. Its elevation generally indicates tissue damage. Raised LDH was a common findings in patients infected with MERS-CoV [33–35]. High LDH levels has previously been shown to be an independent prognostic indicator of SARS infection which can help clinicians to predict adverse clinical outcome [36-38]. It was also reported to be one of the factors tightly associated with mortality of acute respiratory distress syndrome (ARDS) [39]. Our finding of increased LDH in severe pneumonia group and death group indicated the possibility of subclinical tissue damage. As the disease progresses, not only the damage of lung but also involvement in multiple tissues and organs can be observed in severe patients, which indicates systemic organ damage caused by the excessive reaction of the immune response. Thus, we suggest that LDH levels could be used as a surrogate marker help to locate damaged tissues or organs.

Our study has some limitations. First, we no doubt missed patients who were asymptomatic or had mild cases and who were treated at home, so our study cohort may represent the more severe end of Covid-19. Second, some specific information was missing in this study, such as symptoms, chest CT scans, supportive treatment and living status. Third, data generation was clinically driven and not systematic. Last but not least, interpretation of our findings might be limited by the sample size, which may have some impact on the statistical results. Based on the limitations above, a multicenter study will be needed to expand the sample size and to conduct more rigorous randomized controlled trials.

Conclusions:

In summary, our results indicate that the risk factors for severe pneumonia were D-dimer, C-reactive protein, lactate dehydrogenase, white blood cell count, neutrophil count and neutrophil percentage, whereas the risk factors for death were D-dimer, lactate dehydrogenase, white blood cell count, neutrophil count, neutrophil percentage and blood urea nitrogen among COVID-19 patients combined with diabetes, hypertension or coronary. Lactate dehydrogenase were independent risk factors for death. The mortality rate of COVID-19 patients combined with diabetes, hypertension or coronary was higher than that of simple COVID-19 patients.

Declarations

Authors' Contributions

H. L. contributed to literature search and drafting the manuscript. J.L. and J.Z. contributed to data collection and literature search. L.Y., D.W., L.Z. and X.D. contributed to make grammatical revisions and polish the language. G.Y. contributed to the study conception and final approval of the manuscript. G.Y. is the guarantors of this work and, as such, had full access to all data in the study and take responsibility for the integrity of data and the accuracy of the data analysis. Here, we show our sincere respect to all medical staff around the world gathered in an emergency and ran to the front line of the epidemic who put their personal safety aside and fought hard with the sick devil.

Acknowledgements

All the authors of this manuscript have made substantial contributions to this work.

Competing Interests

None of the authors have any potential conflict of interests associated with this research.

Availability of data and materials

The datasets used and/or analysed during the current study are available from the

authors on request.

Ethics approval and consent to participate

The study was approved by the Research Ethics Commission of FPHJD (Approval No. 2020021) and written consent was waived by Ethics Commission of FPHJD, and all participants provided written informed consents.

Consent for publication

All participants provided written informed consents.

Disclosure

Each of the authors acknowledges that he or she participated sufficiently in the work to take public responsibility for its content.

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Figures

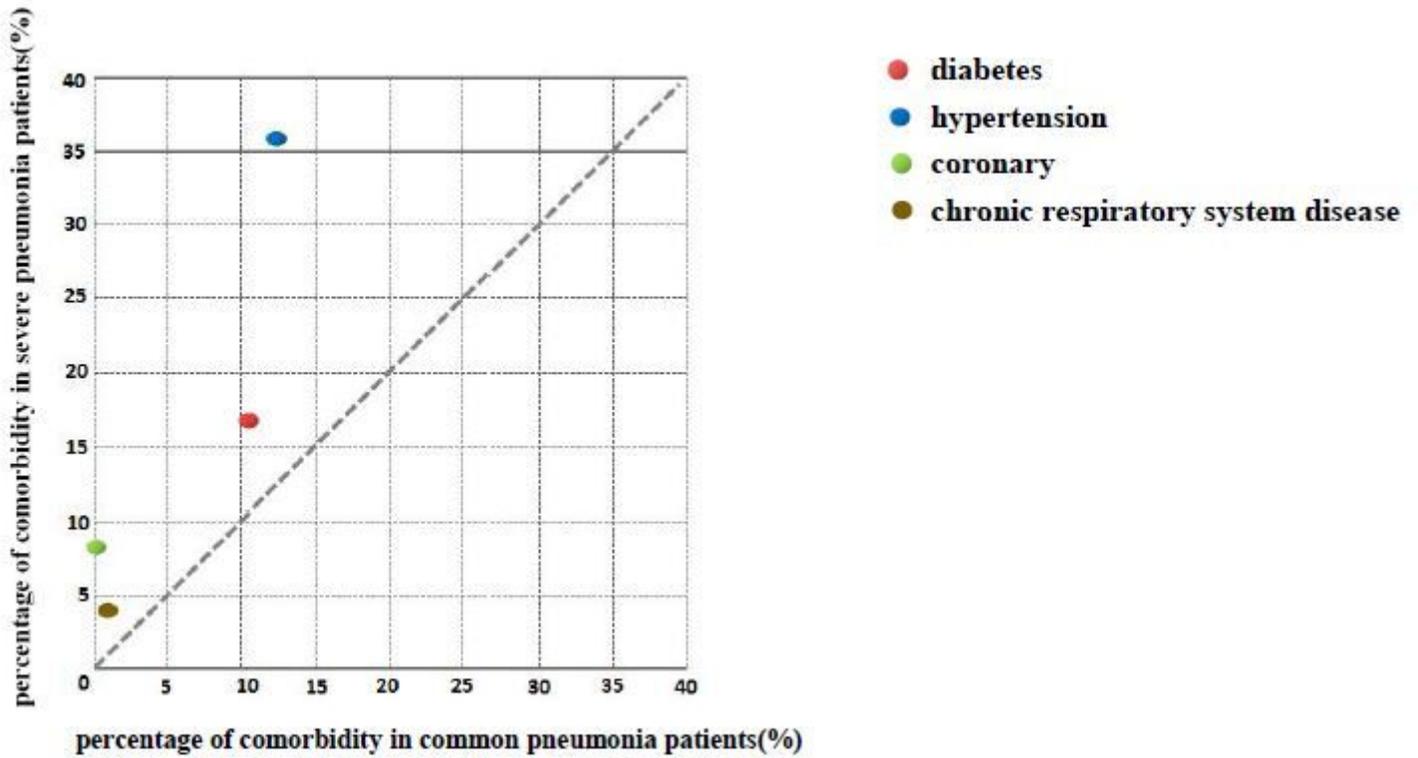


Figure 1

Comorbidity percentage in common pneumonia and severe pneumonia patients was plotted versus each other. Diagonal (gray, dotted) indicated a hypothetically equal percentage between the two groups.

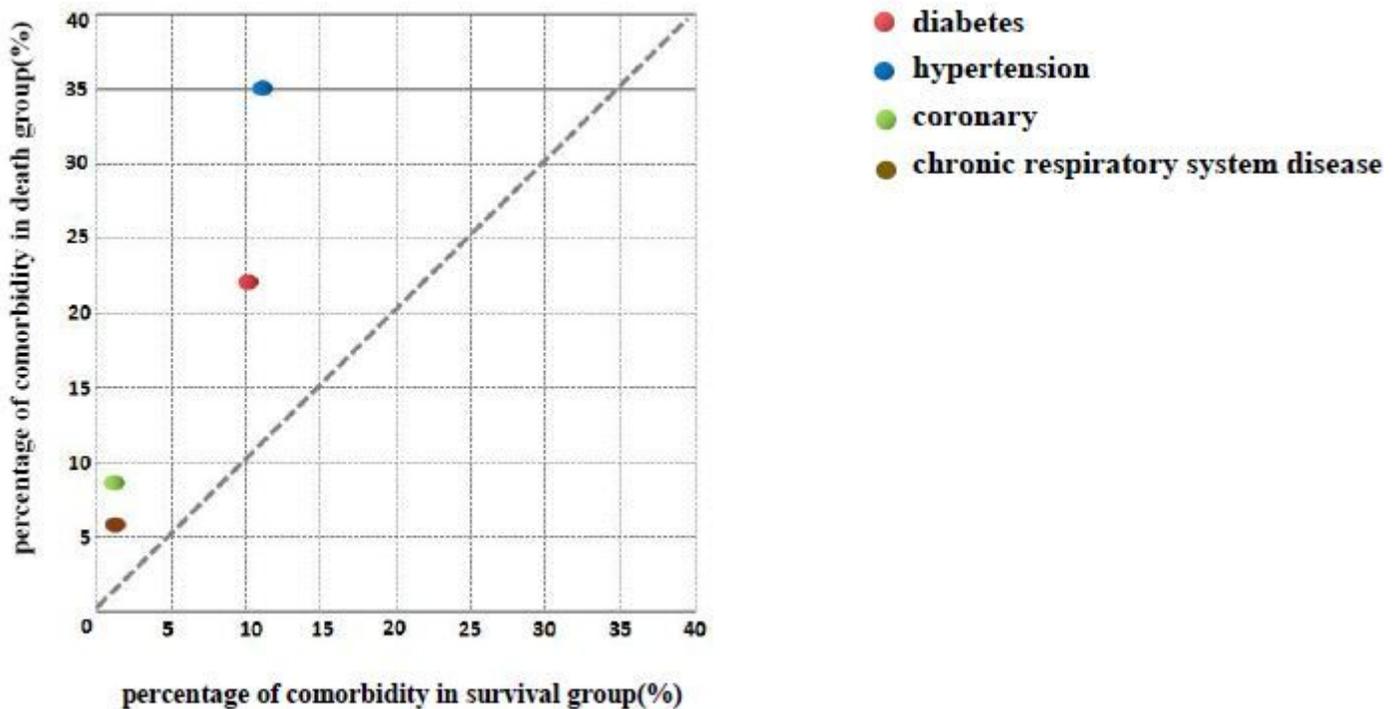


Figure 2

Comorbidity percentage in death and survival group was plotted versus each other. Diagonal (gray, dotted) indicated a hypothetically equal percentage between the two groups.

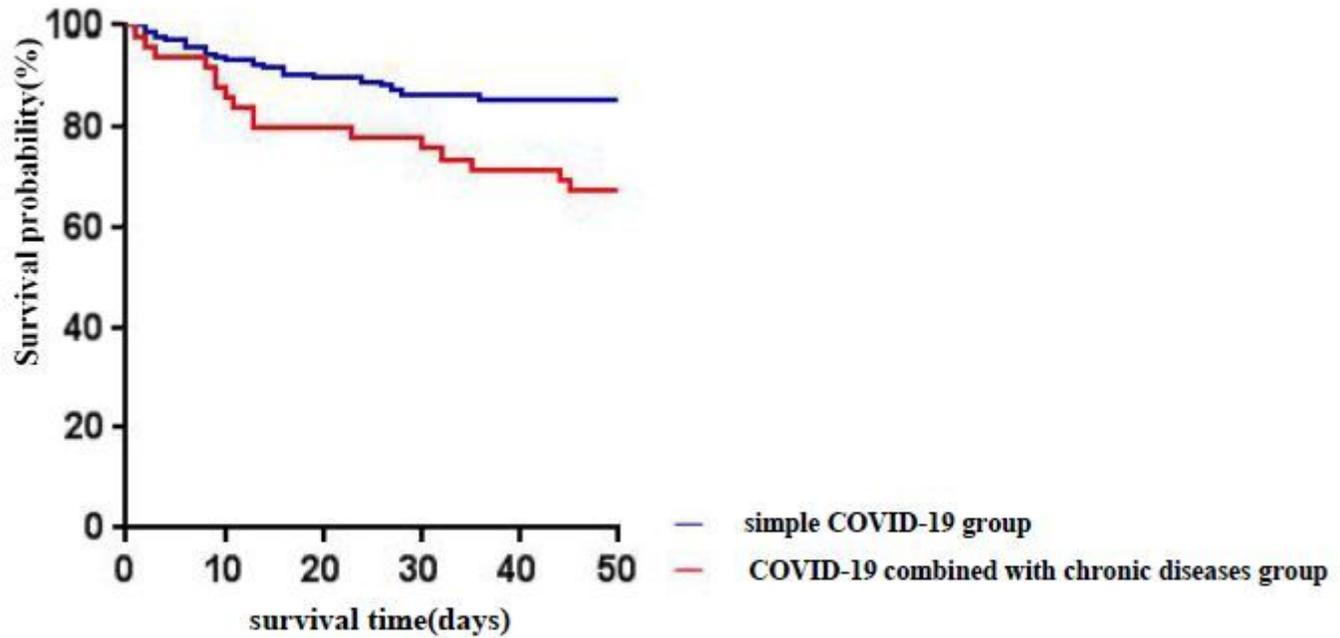


Figure 3

The survival curves of the simple COVID-19 group and COVID-19 combined with chronic diseases group