

## ORIGINAL ARTICLE

## Epidemiology and Genetics

# Clinical characteristics of 140 patients infected with SARS-CoV-2 in Wuhan, China

Jin-jin Zhang<sup>1</sup> | Xiang Dong<sup>1</sup> | Yi-yuan Cao<sup>2</sup> | Ya-dong Yuan<sup>3</sup> | Yi-bin Yang<sup>4</sup> | You-qin Yan<sup>5</sup> | Cezmi A. Akdis<sup>6</sup> | Ya-dong Gao<sup>1</sup>

<sup>1</sup>Department of Allergology, Zhongnan Hospital of Wuhan University, Wuhan, China

<sup>2</sup>Department of Radiology, Zhongnan Hospital of Wuhan University, Wuhan, China

<sup>3</sup>Department of Respiratory and Critical Care Medicine, Second Hospital of Hebei Medical University, Shijiazhuang, China

<sup>4</sup>Department of Respiratory and Critical Care Medicine, Zhongnan Hospital of Wuhan University, Wuhan, China

<sup>5</sup>Department of Infectious Disease, No. 7 Hospital of Wuhan, Wuhan, China

<sup>6</sup>Swiss Institute of Allergy and Asthma Research (SIAF), University of Zurich, Davos, Switzerland

## Correspondence

Ya-dong Gao, Department of Allergology, Zhongnan Hospital of Wuhan University, Wuhan 430071, Hubei, China.  
Email: gaoyadong@whu.edu.cn

## Abstract

**Background:** Coronavirus disease 2019 (COVID-19) caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) infection has been widely spread. We aim to investigate the clinical characteristic and allergy status of patients infected with SARS-CoV-2.

**Methods:** Electronic medical records including demographics, clinical manifestation, comorbidities, laboratory data, and radiological materials of 140 hospitalized COVID-19 patients, with confirmed result of SARS-CoV-2 viral infection, were extracted and analyzed.

**Results:** An approximately 1:1 ratio of male (50.7%) and female COVID-19 patients was found, with an overall median age of 57.0 years. All patients were community-acquired cases. Fever (91.7%), cough (75.0%), fatigue (75.0%), and gastrointestinal symptoms (39.6%) were the most common clinical manifestations, whereas hypertension (30.0%) and diabetes mellitus (12.1%) were the most common comorbidities. Drug hypersensitivity (11.4%) and urticaria (1.4%) were self-reported by several patients. Asthma or other allergic diseases were not reported by any of the patients. Chronic obstructive pulmonary disease (COPD, 1.4%) patients and current smokers (1.4%) were rare. Bilateral ground-glass or patchy opacity (89.6%) was the most common sign of radiological finding. Lymphopenia (75.4%) and eosinopenia (52.9%) were observed in most patients. Blood eosinophil counts correlate positively with lymphocyte counts in severe ( $r = .486, P < .001$ ) and nonsevere ( $r = .469, P < .001$ ) patients after hospital admission. Significantly higher levels of D-dimer, C-reactive protein, and procalcitonin were associated with severe patients compared to nonsevere patients (all  $P < .001$ ).

**Conclusion:** Detailed clinical investigation of 140 hospitalized COVID-19 cases suggests eosinopenia together with lymphopenia may be a potential indicator for diagnosis. Allergic diseases, asthma, and COPD are not risk factors for SARS-CoV-2 infection. Older age, high number of comorbidities, and more prominent laboratory abnormalities were associated with severe patients.

Jin-jin Zhang and Xiang Dong contributed equally to this work.

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## KEYWORDS

allergy, COVID-19, eosinophil, risk factor, SARS-CoV-2

## 1 | INTRODUCTION

A novel member of human coronavirus, newly identified in Wuhan, China, recently, now officially named as SARS-CoV-2 (severe acute respiratory syndrome coronavirus 2) by International Committee on Taxonomy of Viruses (ICTV) is a new strain of RNA viruses that has not been previously identified in humans. Studies have shown that the disease caused by SARS-CoV-2, recently named as COVID-19 (coronavirus disease 2019) by World Health Organization (WHO), could induce symptoms including fever, dry cough, dyspnea, fatigue, and lymphopenia in infected patients. In more severe cases, infections causing viral pneumonia may lead to severe acute respiratory syndrome (SARS) and even death.<sup>1-4</sup> Since the first report of COVID-19 in December 2019 in Wuhan, China, the outbreak of the disease is currently continuously evolving. Until February 16, 2020, the locations with confirmed SARS-CoV-2 cases include 25 countries. Globally, 73 332 cases were confirmed, including 72 528 cases in China and 804 cases outside of China.<sup>5</sup> A total of 1870 patients have died from this viral infection.<sup>5</sup>

It was established that the SARS-CoV-2 belongs to the beta-coronavirus 2b lineage in the phylogenetic tree. By examining the full-length genome of SARS-CoV-2, it was discovered that this novel virus shared 87.99% identity sequencing with the bat SARS-like coronavirus,<sup>6</sup> and it shared ~80% identity nucleotide with the original SARS epidemic virus.<sup>7</sup> Based on the preliminary information of this novel virus, it is considered that SARS-CoV-2 is the third zoonotic human coronavirus of the century.<sup>8</sup> In addition, clinical evidences have suggested that this virus is transmissible from person to person.<sup>9,10</sup> Currently, it is still unclear about the origins and possible intermediate animal vectors of SARS-CoV-2, as well as the mechanism of this virus, that is, spreading between persons.

Despite that many articles have established the clinical features of COVID-19 patients so far,<sup>1-4</sup> the allergy aspects and the information of the allergic disease-related laboratory findings of these patients have not been reported yet. Previous studies have discussed the possible relationship between viral infections, immune response, and allergy and asthma. Several types of viruses, such as rhinovirus (RVs) and respiratory syncytial virus (RSV), were the main focus of research,<sup>11-15</sup> whereas for coronavirus, and particularly COVID-19, there is no report in this context. It is generally established that allergy may increase the risk of virus-induced exacerbation of allergic diseases, such as asthma.<sup>16,17</sup> Genetic predisposition, deficient antiviral response, impairment of immune cell function, damage of epithelium, and cytokine and chemokine response may contribute to the synergistical promotion of allergic disease exacerbation by allergen and virus, especially for respiratory tract allergy.<sup>18-20</sup> However, according to the hygiene hypothesis in the allergy field, respiratory infections during early life may

play a protective role against the Th2-mediated allergic disease development.<sup>21</sup>

The prevalence of chronic obstructive pulmonary disease (COPD) in people  $\geq 40$  years old was 13.7% in China according to a recent study.<sup>22</sup> Respiratory tract virus infection is a common trigger for acute exacerbation of COPD.<sup>23</sup> The prevalence of SARS-CoV-2 infection in COPD patients is not clear. In addition, the influence of smoking behavior on the susceptibility to this virus has not been investigated.

This research aims to investigate the clinical and laboratory characteristic of hospitalized COVID-19 patients, including differences between severe and nonsevere patients, and to reveal the relationship between SARS-CoV-2 infection, immune response, allergy, and clinical manifestations, with a special focus on asthma, COPD, and smoking behavior.

## 2 | METHODS

### 2.1 | Patients' involvement and data collection

All hospitalized patients ( $n = 242$ ) (admission date from January 16 to February 3, 2020) in No. 7 Hospital of Wuhan, clinically diagnosed as "viral pneumonia" based on their clinical symptoms (fever or respiratory symptoms) with typical changes in chest radiology, were preliminarily involved in this study. No. 7 Hospital of Wuhan is one of the designated hospitals for the hospitalization of patients with COVID-19 and has been entrusted by Zhongnan Hospital of Wuhan University since January 2020. The pharyngeal swab specimens of these patients were collected and used for SARS-CoV-2 detection. Patients absent of or with negative SARS-CoV-2 test results were excluded from this study. All patients involved in this study were living in Wuhan during the outbreak period of COVID-19.

Demographic information, clinical characteristics (included medical history, exposure history, comorbidities, surgery history, signs, and symptoms), chest computed tomographic (CT) scan or X-ray results, and laboratory findings of each patient were obtained from the electronic medical record system of No. 7 Hospital of Wuhan and analyzed by three independent researchers. The access was granted by the director of the hospital. Patients with smoking history, COPD, and allergic diseases (including asthma, allergic rhinitis, food allergy, atopic dermatitis, and urticaria) were identified. For patients with smoking history, the amount of smoking, the years of smoking history, and the years of smoking cessation were individually collected. This study was approved by the institutional ethics board of Zhongnan Hospital of Wuhan University (No. 2020028).

The date of disease onset and hospital admission date, as well as the severity of COVID-19, were also recorded. The onset date

was defined as the day when any symptoms were noticed by the patients. Severity of COVID-19 was defined according to the diagnostic and treatment guideline for SARS-CoV-2 issued by Chinese National Health Committee (version 3-5). Severe COVID-19 was designated when the patients had one of the following criteria: (a) respiratory distress with respiratory frequency  $\geq 30$ /min; (b) pulse oximeter oxygen saturation  $\leq 93\%$  at rest; and (c) oxygenation index (artery partial pressure of oxygen/inspired oxygen fraction,  $\text{PaO}_2/\text{FiO}_2$ )  $\leq 300$  mm Hg.

## 2.2 | Laboratory testing

Patient pharyngeal swab specimens were collected for the SARS-CoV-2 viral nucleic acid detection using real-time reverse transcriptase-polymerase chain reaction (RT-PCR) assay. The viral nucleic acid testing for all patients was performed by the clinical laboratory from Zhongnan Hospital of Wuhan University in Wuhan, which is the designated laboratory for SARS-CoV-2 test for patients in No. 7 Hospital of Wuhan. Detailed protocol was described somewhere else.<sup>3</sup>

Medical laboratory finding results, including the numbers of leukocytes, lymphocytes, and eosinophils; percentages of lymphocyte and eosinophils; concentrations of D-dimer, C-reactive protein (CRP), procalcitonin (PCT), serum amyloid A (SAA), and serum creatine kinase; and detection of different pathogens (including *Mycoplasma pneumoniae*, *Chlamydia pneumoniae*, influenza A virus, influenza B virus, parainfluenza, Coxsackie virus group B, adenovirus, echovirus, respiratory syncytial virus, Epstein-Barr virus, and cytomegalovirus), were collected for each patient. All medical laboratory data were generated by the clinical laboratory of No. 7 Hospital of Wuhan. As disease progressed, the most updated secondary results for laboratory findings (included numbers and percentages of lymphocytes and eosinophils) during the hospital stay were also collected. The laboratory data for some patients were missing due to the absence of types of tests or delayed results.

## 2.3 | Statistical analysis

Categorical variables were summarized as frequencies and percentages, and continuous variables were described using median and interquartile ranges (IQR) values. To compare the continuous variables for data of different patient groups, two-tailed *t* test and *Mann-Whitney* test were used as appropriate. The frequencies of categorical variables were compared using the *chi-square* and *Fisher's* exact test as appropriate. *Spearman's* correlation test was used for calculation of correlation between different cell types. All statistical analyses and graphs were generated and plotted using GraphPad Prism version 7.00 software (GraphPad Software Inc). The tests with *P* value of  $<.05$  were considered statistically significant.

## 3 | RESULTS

### 3.1 | Demographics and clinical characteristics

A total of 140 patients diagnosed as COVID-19 were included in this study, with 82 patients categorized into nonsevere patients and 58 severe cases on admission. The median age for all patients was 57 years, ranging from 25 to 87 years old, and the majority (70%) of them were more than 50 years old. About half (50.7%) of patients were male. Since no patients had direct exposure history of Huanan wet markets or wildlife animals, we presumed all patients in this study were community-infected cases. The SARS-CoV-2 clustered within 15 (10.7%) patients in this study, whose family members or friends were also infected with SARS-CoV-2. Three hospital workers were infected. Ninety (64.3%) patients had at least one underlying comorbidity, the most common of which were chronic diseases, such as hypertension (30%) and diabetes (12.1%). Only two COPD patients were identified. Sixteen patients had self-reported medical history of drug hypersensitivity, and drug types include penicillin, cephalosporins, and specific Chinese traditional medicines. Two patients reported chronic urticaria. No self-reported other allergic diseases (including asthma, allergic rhinitis, food allergy, atopic dermatitis, and other type 2 allergic diseases) were declared. Thirty-eight (27.1%) patients had surgery experience(s), such as C-section, cholecystectomy, and appendectomy. Interestingly, there were only 2 current smokers and 7 past smokers (Table 1).

Symptoms of the patients on admission are shown in Table 2. The median interval from the onset of symptoms to hospital admission for all patients was 8 days (IQR, 6-11). The most commonly experienced symptoms were fever (91.7%), followed by cough (75%), fatigue (75%), and chest tightness or dyspnea (36.7%). 39.6% of them complained about gastrointestinal symptoms, including nausea, diarrhea, poor appetite, abdominal pain, belching, and emesis.

### 3.2 | Radiological and laboratory findings

Of the 135 patients with chest CT scan on admission, the majority (134, 99.3%) had abnormal results, showing typical images that were bilateral multiple ground-glass opacities or consolidation (Table 2, Figures 1-3). The blood cell test result of patients on the day of hospital admission showed normal leukocytes in most of the patients (68.1%), with 12.3% increased and 19.6% decreased numbers. Lymphopenia was common in 75.4% of the patients (Table 3). Interestingly, more than half of these patients (52.9%) had eosinopenia (eosinophil counts  $< 0.02 \times 10^9/\text{L}$ ). Other laboratory findings included higher concentration of C-reactive protein (91.9%), serum amyloid A (90.2%), and D-dimer (43.2%), while increased concentration of serum procalcitonin (34.7%) and creatine kinase (6.7%) was relatively less common in all patients. The results of secondary tests (median, 5 days after hospital

**TABLE 1** Demographics and baseline characteristics of patients with COVID-19

	All patients (n = 140)	Diseases severity		P value
		Nonsevere patients (n = 82)	Severe patients (n = 58)	
Age—median (range)	57 (25-87)	51.5 (26-78)	64 (25-87)	<.001
Age-groups—No. (%)				
<30 y	5 (3.6)	4 (4.9)	1 (1.7)	.002
30-49 y	37 (26.4)	28 (34.1)	9 (15.5)	-
50-69 y	69 (49.3)	41 (50.0)	28 (48.3)	-
≥70 y	29 (20.7)	9 (11.0)	20 (34.5)	-
Sex—No. (%)				
Female	69 (49.3)	44 (53.7)	25 (43.1)	.219
Male	71 (50.7)	38 (46.3)	33 (56.9)	-
Exposure history—No. (%)				
Familiar/cluster infections	15 (10.7)	11 (13.4)	4 (6.9)	.342
Hospital staff	3 (2.1)	3 (3.7)	0 (0)	.267
Comorbidity—No. (%)	90 (64.3)	44 (53.7)	46 (79.3)	.002
Hypertension	42 (30.0)	20 (24.4)	22 (37.9)	.085
Diabetes mellitus	17 (12.1)	9 (11.0)	8 (13.8)	.615
Fatty liver and abnormal liver function	8 (5.7)	4 (5.0)	4 (6.9)	.718
Chronic gastritis and gastric ulcer	7 (5.0)	5 (6.1)	2 (3.4)	.700
Coronary heart disease	7 (5.0)	3 (3.7)	4 (6.9)	.448
Hyperlipidemia	7 (5.0)	5 (6.1)	2 (3.4)	.700
Cholelithiasis	6 (4.3)	2 (2.4)	4 (6.9)	.232
Arrhythmia	5 (3.6)	1 (1.2)	4 (6.9)	.160
Thyroid diseases	5 (3.6)	1 (1.2)	4 (6.9)	.160
Electrolyte imbalance	4 (2.9)	0 (0)	4 (6.9)	.028
Urolithiasis	3 (2.1)	2 (2.4)	1 (1.7)	>.999
Stroke	3 (2.1)	1 (1.2)	2 (3.4)	.570
Chronic renal insufficiency	2 (1.4)	0 (0)	2 (3.4)	.170
Aorta sclerosis	2 (1.4)	1 (1.2)	1 (1.7)	>.999
Secondary pulmonary tuberculosis	2 (1.4)	0 (0)	2 (3.4)	.170
COPD	2 (1.4)	0 (0)	2 (3.4)	.170
Asthma and allergic diseases (self-reported)				
Asthma	0 (0)	0 (0)	0 (0)	-
Allergic rhinitis	0 (0)	0 (0)	0 (0)	-
Food allergy	0 (0)	0 (0)	0 (0)	-
Atopic dermatitis	0 (0)	0 (0)	0 (0)	-
Drug hypersensitivity	16 (11.4)	10 (12.2)	6 (10.3)	.735
Urticaria	2 (1.4)	1 (1.2)	1 (1.7)	>.999
Surgery history—No. (%)	38 (27.1)	19 (23.2)	19 (32.8)	.209
Cesarean section—No./total female No. (%)	10/69 (14.5)	9/44 (20.5)	1/25 (4.0)	.081
Cholecystectomy	9 (6.4)	3 (3.7)	6 (10.3)	.162
Appendectomy	7 (5.0)	3 (3.7)	4 (6.9)	.448
Tumor surgery	6 (4.3)	3 (3.7)	3 (5.2)	.692
Osteoarticular surgery	5 (3.6)	2 (2.4)	3 (5.2)	.649
Cranio-cerebral surgery	3 (2.1)	2 (2.4)	1 (1.7)	>.999

(Continues)

**TABLE 1** (Continued)

	All patients (n = 140)	Diseases severity		P value
		Nonsevere patients (n = 82)	Severe patients (n = 58)	
Cardiac intervention	3 (2.1)	0 (0)	3 (5.2)	.069
Ureterotomy	1 (0.7)	0 (0)	1 (1.7)	.414
Hysterectomy	1 (0.7)	0 (0)	1 (1.7)	.414
Hemorrhoidectomy	1 (0.7)	0 (0)	1 (1.7)	.414
Varicose vein surgery	1 (0.7)	0 (0)	1 (1.7)	.414
Smokers—No. (%)	9 (6.4)	3 (3.7)	6 (10.3)	.162
Past smokers	7 (5.0)	3 (3.7)	4 (6.9)	.448
Current smokers	2 (1.4)	0 (0)	2 (3.4)	.170
Smoking index				
<400	3	1	2	>.999
≥400	6	2	4	-

Note: P values denoted the comparison between nonsevere and severe subgroups. Smoking Index = cigarettes smoked per day × years of smoking. Abbreviations: COPD, chronic obstructive pulmonary disease; COVID-19, coronavirus disease 2019; IQR, interquartile range.

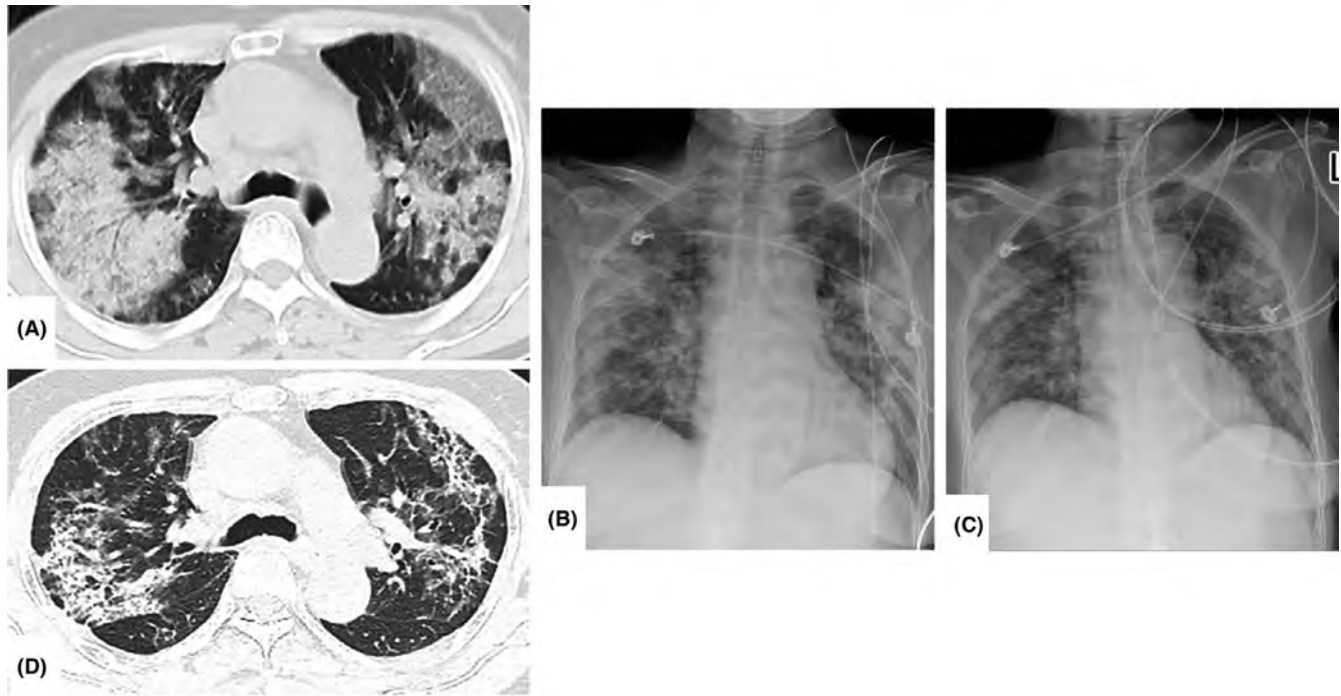
**TABLE 2** Symptomatic and radiological characteristics of patients with COVID-19

	All Patients	Diseases severity		P value
		Nonsevere patients	Severe patients	
Onset of symptom to hospital admission—median (IQR), days	8 (6-11)	8 (5-11)	7 (6-12)	.503
Signs and symptoms—No./total No. (%)				
Fever	110/120 (91.7)	59/67 (88.1)	51/53 (96.2)	.182
Cough	90/120 (75.0)	45/67 (67.2)	45/53 (84.9)	.023
Fatigue	90/120 (75.0)	51/67 (76.1)	39/53 (73.6)	.750
Chest tightness/dyspnea	44/120 (36.7)	20/67 (29.9)	24/53 (45.3)	.082
Gastrointestinal symptoms	55/139 (39.6)	31/82 (37.8)	24/57 (42.1)	.610
Nausea	24/139 (17.3)	19/82 (23.2)	5/57 (8.8)	.027
Diarrhea	18/139 (12.9)	9/82 (11.0)	9/57 (15.8)	.406
Anorexia	17/139 (12.2)	9/82 (11.0)	8/57 (14.0)	.588
Abdominal pain	8/139 (5.8)	2/82 (2.4)	6/57 (10.5)	.064
Belching	7/139 (5.0)	4/82 (4.9)	3/57 (5.3)	>.999
Emesis	7/139 (5.0)	5/82 (6.1)	2/57 (3.5)	.700
Chest CT images—No./total No. (%)				
Abnormal	134/135 (99.3)	77/78 (98.7)	57/57 (100.0)	>.999
Bilateral lung	121/135 (89.6)	68/78 (87.2)	53/57 (93.0)	-
Single lung—left	5/135 (3.7)	3/78 (3.8)	2/57 (3.5)	-
Single lung—right	8/135 (5.9)	6/78 (7.7)	2/57 (3.5)	-
Normal	1/135 (0.7)	1/78 (1.3)	0/57 (0)	-

Note: P values denoted the comparison between nonsevere and severe subgroups. Abbreviations: COVID-19, coronavirus disease 2019; IQR, interquartile range.

admission) showed decreased lymphocyte counts (median, 0.7 vs 1.1,  $P = .001$ ) were more common, and lymphocyte percentage (median, 10.3 vs 22.1,  $P < .001$ ) and the eosinophil percentage (median, 0.2 vs 0.8,  $P = .017$ ) were lower in severe patients (Table 4).

Besides SARS-CoV-2, other pathogens were also detected within some patients, including *Mycoplasma pneumoniae* (5, 8.6%), respiratory syncytial virus (1, 1.9%), and Epstein-Barr virus (1, 3.7%) (Table 5). However, no clinical and radiological signs of superinfection caused by these pathogens were identified.



**FIGURE 1** Chest X-ray and CT images of a 67-y-old woman with onset of cough and sputum on January 1, 2020, and progressively developed dyspnea. A, Transverse CT scan image on January 9 showing multiple lobular and segmental consolidation combined with ground-glass opacities diffusely distributed in bilateral lung field. B, Chest X-ray showing extended bilateral consolidation on January 12. C, The attenuation and the involvement of the consolidation decreased in chest X-ray of January 17 (D) CT scan on January 22 showing absorption of bilateral consolidation, scattered fibrous can be observed. The symptoms and dyspnea of the patient improved after treatment, and the patient was discharged on January 24

### 3.3 | Characteristics of severe cases

Severity was established based on respiratory functions on admission with one of the below criteria: respiratory frequency  $\geq 30$ /min, oxygen saturation  $\leq 93\%$  at rest, and oxygenation index  $\leq 300$  mm Hg (see methods). Median age was 64 years in severe cases, compared to 51.5 years in nonsevere cases ( $P < .001$ ). More comorbidities (79.3% vs 53.7%,  $P = .002$ ), higher median values of leukocyte count (5.3 vs 4.5,  $P = .014$ ), D-dimer (0.4 vs 0.2,  $P < .001$ ), CRP (47.6 vs 28.7,  $P < .001$ ), PCT (0.1 vs 0.05,  $P < .001$ ), and lower lymphocyte percentage (median, 0.7 vs 0.8,  $P = .048$ ) were found in severe cases, compared to nonsevere cases. No difference was identified for the occurrence rates of most signs and symptoms between nonsevere and severe patients, and only two symptoms (cough and nausea) were more commonly experienced in severe group ( $P = .023$  and 0.027, respectively). Increased leukocyte number ( $P = .003$ ), D-dimer ( $P = .004$ ), and PCT ( $P = .004$ ) were more commonly observed in severe patients (Tables 1-3).

### 3.4 | Correlation between blood eosinophil and lymphocyte counts

The absolute numbers of circulating eosinophils correlated positively with the numbers of lymphocytes for all 138 patients, both for the tests on the first day after hospital admission ( $r = .321$ ,  $P < .001$ )

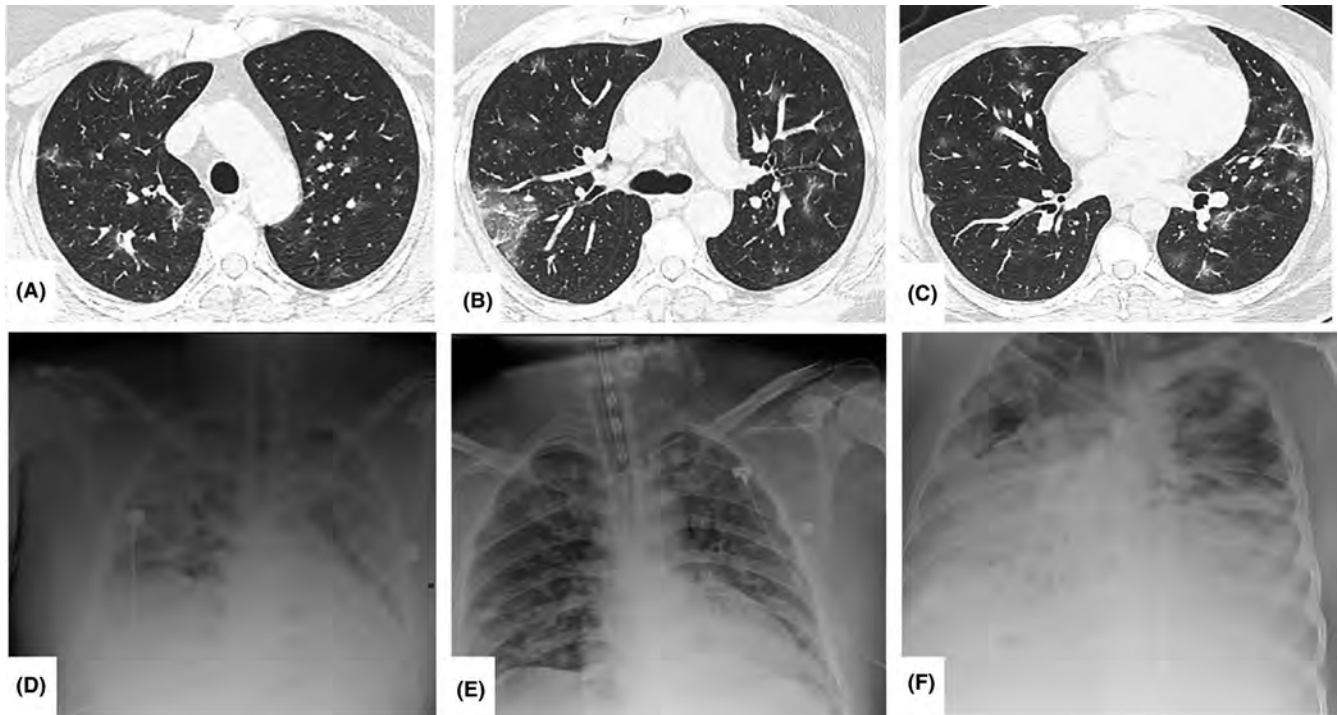
and for the secondary tests on 3 or more days afterward ( $r = .479$ ,  $P < .001$ ). Same correlation was also found between the two cell types, when patients were separated to severe ( $r = .250$ ,  $P = .066$  and  $r = .486$ ,  $P < .001$ , respectively) and nonsevere groups ( $r = .369$ ,  $P < .001$  and  $r = .469$ ,  $P < .001$ , respectively) (Figure 4). Although no significant correlations between the two cell types could be determined for the severe patients on the first day after hospital admission ( $r = .250$ ,  $P = .066$ ), this may be due to the extremely low values of eosinophils for the severe patients on that day.

## 4 | DISCUSSION

In this study including 140 community-infected COVID-19 patients, we found that most patients were middle- and old-aged, with almost 1:1 male-female ratio. Fever (91.7%), cough (75.0%), and fatigue (75.0%) were the most common symptoms in COVID-19 patients. More than 1/3 of the patients had chest tightness or dyspnea and gastrointestinal symptoms such as nausea, diarrhea, and anorexia. Radiologically, CT scan or X-ray showed bilateral ground-glass and patchy opacity in 89.6% patients. Infection with SARS-CoV-2 was confirmed by RT-PCR in all patients, and other pathogens were rarely identified in these patients.

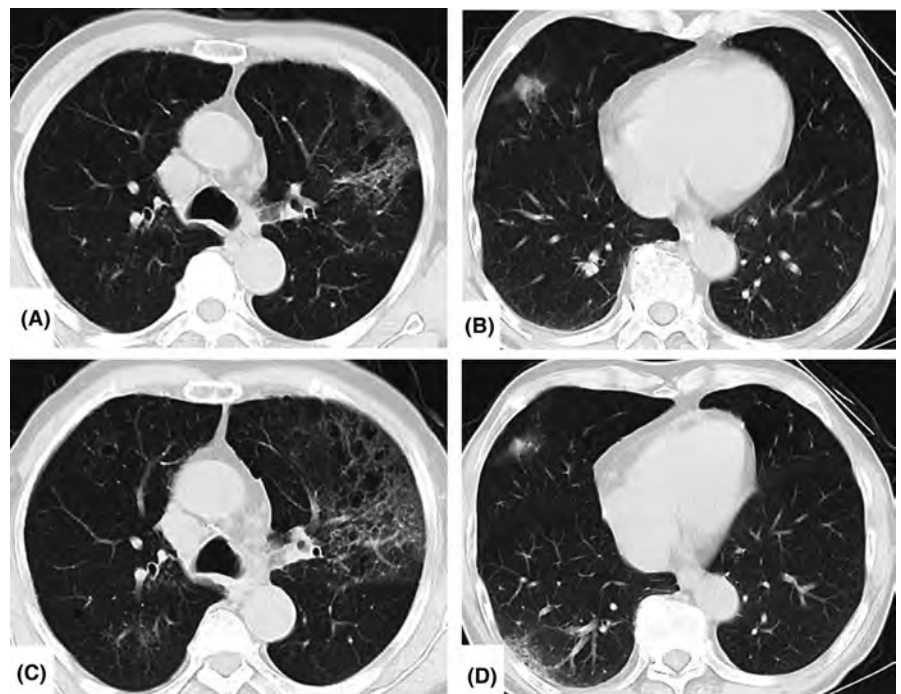
The median age of all patients was 57 years old, which is close to the data reported by Wang et al<sup>3</sup> (56.0 years) and Chen et al<sup>1</sup> (55.5 years), but older than that reported by Huang et al<sup>2</sup>





**FIGURE 2** Chest X-ray and CT scan images of a 36-y-old man without history of smoking. The patient had fever, cough, and diarrhea on January 6, 2020. A-C, Transverse CT images on January 9. A, Bilateral multiple ground-glass opacities, most of them are irregular small round lesions scattered in the lung field in upper lobe. B, Bilateral multiple irregular ground-glass opacities and a wedge-shaped opacity located in the right upper lobe under the pleura. C, Bilateral multiple irregular ground-glass opacities and a small nodular opacity located in the left lower upper lobe under the pleura. D, The symptoms of the patient deteriorated on January 12, chest X-ray showing bilateral diffuse patchy and consolidation, so-called “white lung.” E, Chest X-ray after intubation and mechanical ventilation on January 13, the attenuation lowered down, leaving scattered small irregular consolidation. F, On January 20, bilateral lung lesions deteriorated, and bilateral costophrenic angles were not clearly displayed, suggesting pleural effusion. The patient died on January 21

**FIGURE 3** CT images of a 76-y-old man with smoking history for 20 y, 1 pack/d, who has stopped smoking for 20 y. The patient developed fever and cough during hospitalization for stasis dermatitis. The SARS-CoV-2-induced pneumonia was confirmed with RT-PCR with samples from throat swab. A-B, CT images after 3 d of symptoms onset. A, Diffuse small round low-attenuation lesions associate with ground-glass opacity on left upper lobe. B, An irregular consolidation in right middle lobe. C, D, CT images after 6 d of symptoms onset. C, The opacity on left upper lobe extended. D, New ground-glass opacity under pleura developed in right lower lobe



(49.0 years). Severe patients were much older than nonsevere patients and associated with higher frequency of comorbidities. In

our report, 90 (64.3%) patients had comorbidities and 38 (27.1%) had surgery history. Hypertension (30%), diabetes mellitus (12.1%),

**TABLE 3** Laboratory results of patients with COVID-19

Laboratory parameters	All patients (n = 138)	Disease severity		P value
		Nonsevere patients (n = 82)	Severe patients (n = 56)	
Leukocytes ( $\times 10^9/L$ ; normal range 3.5-9.5)	4.7 (3.7-6.7)	4.5 (3.5-5.9)	5.3 (4.0-9.0)	.014
Increased—No./total No. (%)	17/138 (12.3)	4/82 (4.9)	13/56 (23.2)	.003
Decreased—No./total No. (%)	27/138 (19.6)	18/82 (22.0)	9/56 (16.1)	.513
Lymphocytes ( $\times 10^9/L$ ; normal range 1.1-3.2)	0.8 (0.6-1.1)	0.8 (0.6-1.2)	0.7 (0.5-1.0)	.048
Decreased—No./total No. (%)	104/138 (75.4)	58/82 (70.7)	46/56 (82.1)	.160
Lymphocyte percentage (%; normal range 20-50)	16.9 (9.2-26.0)	20.0 (12.5-28.4)	12.7 (7.7-22.0)	<.001
Eosinophils ( $\times 10^9/L$ ; normal range 0.02-0.52)	0.01 (0.0-0.05)	0.02 (0.008-0.05)	0.01 (0.0-0.06)	.451
Decreased—No./total No. (%)	73/138 (52.9)	39/82 (47.6)	34/56 (60.7)	.165
Eosinophils percentage (%; normal range 0.4-8)	0.3 (0.0-1.0)	0.5 (0.08-1.0)	0.2 (0.0-0.8)	.166
D-Dimer ( $\mu g/mL$ ; normal range 0-0.243)	0.2 (0.1-0.5)	0.2 (0.1-0.3)	0.4 (0.2-2.4)	<.001
Increased—No./total No. (%)	35/81 (43.2)	12/43 (27.9)	23/38 (60.5)	.004
C-reactive protein (CRP) (mg/L; normal range 0-3)	34.2 (12.5-67.4)	28.7 (9.5-52.1)	47.6 (20.6-87.1)	<.001
Increased—No./total No. (%)	125/136 (91.9)	72/81 (88.9)	53/55 (96.4)	.199
Procalcitonin (PCT) (ng/mL; normal range 0-0.1)	0.07 (0.04-0.1)	0.05 (0.03-0.1)	0.1 (0.06-0.3)	<.001
Increased—No./total No. (%)	41/118 (34.7)	16/68 (23.5)	25/50 (50.0)	.004
Serum amyloid A (SAA) (mg/L; normal range 0-10)	92.53 (44.6-161.3)	91.5 (24.9-163.2)	108.4 (54.1-161.6)	.600
Increased—No./total No. (%)	46/51 (90.2)	29/34 (85.3)	17/17 (100.0)	.156
Serum Creatine Kinase (U/L; normal range 40-200)	72.5 (52.2-115)	83.0 (56.0-112.0)	66.0 (38.5-144.0)	.192
Increased—No./total No. (%)	4/60 (6.7)	1/35 (2.8)	3/25 (12.0)	.298

Note: Data are shown as median (IQR); COVID-19, coronavirus disease 2019; IQR, interquartile range; P values denoted the comparison between nonsevere and severe subgroups.

**TABLE 4** Secondary lymphocyte and eosinophil counts in patients with COVID-19

Parameters	All patients (n = 138)	Disease severity		P value
		Nonsevere patients (n = 82)	Severe patients (n = 52)	
Days from the previous test—median (IQR), days	5 (3.0-7.3)	5 (4.0-10.0)	5 (3-7)	-
Lymphocytes ( $\times 10^9/L$ ; normal range 1.1-3.2)	0.9 (0.6-1.4)	1.1 (0.7-1.6)	0.7 (0.4-1.1)	<.001
Decreased—No./total No. (%)	71/118 (60.2)	31/66 (47.0)	40/52 (76.9)	.001
Lymphocyte percentage (%; normal range 20-50)	16.4 (8.1-26.4)	22.1 (13.5-29.5)	10.3 (4.3-16.6)	<.001
Eosinophils ( $\times 10^9/L$ ; normal range 0.02-0.52)	0.03 (0.0-0.08)	0.04 (0.01-0.08)	0.01 (0.0-0.09)	.157
Decreased—No./total No. (%)	49/118 (41.5)	22/66 (33.3)	27/52 (51.9)	.060
Eosinophils percentage (%; normal range 0.4-8)	0.5 (0.0-1.45)	0.8 (0.2-1.8)	0.2 (0.0-0.9)	.017

Note: Data are shown as median (IQR); COVID-19, coronavirus disease 2019; IQR, interquartile range; P values denoted the comparison between nonsevere and severe subgroups.

and cardiovascular diseases were the most common underlying diseases, consistent with other recent reports.<sup>1-3</sup> The prevalence of hypertension and diabetes in China was 23.2%<sup>24</sup> and 10.9%<sup>25</sup> in adults, which was slightly lower than the data in this study; this may be due to the large ratio of elder COVID-19 patients in the series. In general, aged people are more susceptible to COVID-19 and more likely to be severe than people younger than 50 years; this may be due to more health issues and comorbidities in this population.

In the present study, 50.7% of the patients were male; the percentage is lower than that reported by Huang et al<sup>2</sup> and Chen et al<sup>1</sup> with a male predominance (73.0%), but similar to that reported by Wang et al<sup>3</sup> (54.3%). This may be related to the occupational risk factors for men in wet market, considering that 66.0% patients in Huang's report and 49% patients in Chen's report had Huanan wet market exposure history. No patient in our report had a history of Huanan wet market exposure, indicating the cause of community infection of SARS-CoV-2; therefore, the female-male ratio tends



**TABLE 5** Pathogens identified in patients with COVID-19

Pathogens identified	Patients—No./ Total No. (%)
SARS-CoV-2 RT-PCR assay+	128/140 (91.4)
SARS-CoV-2 RT-PCR assay±	12/140 (8.6)
Mycoplasma pneumoniae IgM Ab+	5/58 (8.6)
Chlamydia pneumoniae IgM Ab+	0/58 (0.0)
Influenza A virus Ag+	0/23 (0.0)
Influenza B virus Ag+	0/23 (0.0)
Parainfluenza IgM Ab+	0/50 (0.0)
Coxsackie virus group B IgM Ab+	0/49 (0.0)
Adenovirus IgM Ab+	0/48 (0.0)
Echovirus IgM Ab+	0/49 (0.0)
RSV-IgM Ab+	1/52 (1.9)
EBV-IgM Ab+ (AU/mL; normal range 0-3)	1/27 (3.7)
CMV-IgM Ab+ (AU/mL; normal range 0-0.42)	0/25 (0.0)

Abbreviations: Ab, antibody; Ag, antigen; CMV, cytomegalovirus; COVID-19, coronavirus disease 2019; EBV, Epstein-Barr virus; IgM, immunoglobulin M; RSV, respiratory syncytial virus; RT-PCR, real-time polymerase chain reaction; SARS-CoV-2, severe acute respiratory syndrome coronavirus 2.

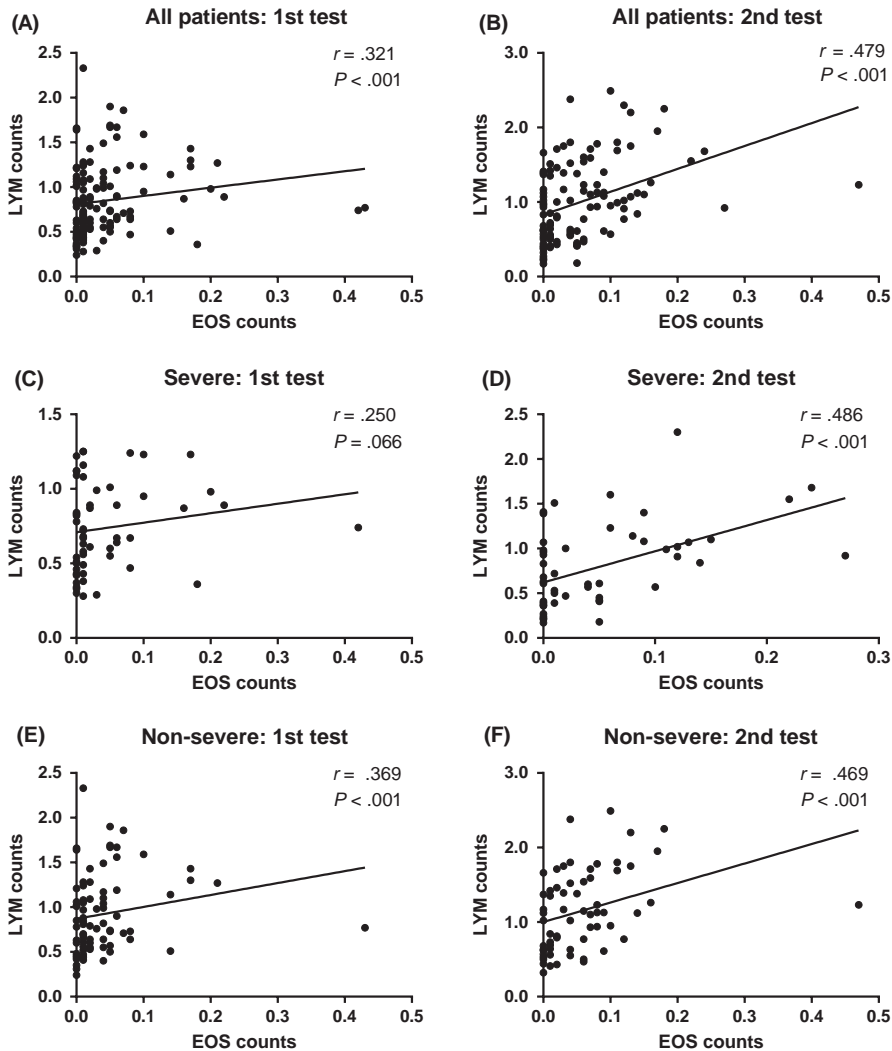
to be approximately 1:1. This is lower than that observed in SARS coronavirus-infected patients, which had a female predominance (61.0%).<sup>26</sup> In addition, there was no difference in the female-male ratio in severe patients.

No asthmatic patient was identified in this report, and only a few patients had self-reported drug hypersensitivity and urticaria. Other allergies such as allergic rhinitis, atopic dermatitis, and food allergy were not reported. Given the prevalence of asthma in China was 4.2%<sup>27</sup> and allergic rhinitis in Wuhan was 9.7%<sup>28</sup> in adults, asthma or allergy may not be the risk factor for the SARS-CoV-2 infection. Many previous studies have investigated the association between virus infection and asthma, showing that respiratory tract viral infections are associated with asthma exacerbations. About 62%-95% of children with acute wheezing episodes tested positive for respiratory viruses in both hospital and community settings; while the prevalence in adults was less, it was still in the range of 41%-78%.<sup>29</sup> Rhinoviruses (RVs) were the most frequently detected type of virus at all ages, whose infection could lead to more severe and longer-lasting lower respiratory tract symptoms. Virus-induced wheezing in infancy was often associated with asthma development in children.<sup>30</sup> In addition to RV, other respiratory tract viruses, such as respiratory syncytial virus (RSV), influenza viruses (IfVs), coronaviruses (CoVs), human metapneumoviruses (HMPVs), parainfluenza viruses (PIVs), adenoviruses (AdVs), and bocaviruses (BoVs), had all been detected in subjects with asthma exacerbations.<sup>23</sup> Nevertheless, findings about the association between asthma and coronaviruses were much less. CoV-OC43 and CoV-229E transmitted primarily during winter had been linked to asthma exacerbations in children and adults.<sup>30</sup> In a literature systemic review with 63 studies included, the pooled prevalence of CoV infection

during asthma exacerbations was 8.4% (95% CI, 5.1%-13.6%), ranking below RV (42.1%, the highest), RSV, herpes simplex virus (HSV), enterovirus (EnV), and IfV, and followed by cytomegalovirus (CV), BoV, PIV, metapneumovirus (MpV), and AdV. In Asia, the five major viruses associated with asthma were RV, EnV, IfV, PIV, and RSV.<sup>31</sup> Reported respiratory infection rates for CoV in children were usually from <1% to 9%, which varied between subtypes. In case-control studies, CoV was not more frequently presented in wheezing children in comparison with controls. Overall, CoV seemed to have a minor, if any, contribution to acute asthma exacerbations, and CoV infections were frequently co-infected with other viruses.<sup>29</sup> As for the other two types of CoVs, SARS-CoV and MERS-CoV which could cause severe respiratory disease in human, most established studies with animal models were found less relevant to asthma.<sup>32</sup> Intriguingly, a prospective clinical birth cohort study demonstrated that the number of early respiratory episodes was related to the risk of later development of asthma, irrespective of the virus type (eg, RV, RSV, and CoV), which indicated the specific viral trigger may not be a risk factor.<sup>33</sup>

On the other hand, it is also worth thinking whether allergic diseases could affect respiratory virus infections. A multicenter prospective study<sup>34</sup> examined the effects of the presence of atopy, which was defined as positive skin prick tests to one or more allergens, on the course of disease in children hospitalized with viral pneumonia. In that study, atopy was found in 21.4% patients, while it showed there was no effect of atopic sensitization on the severity of viral pneumonia in children. In addition, pollen, type 2 response, respiratory viruses, and the tissue milieu such as the type 2 or type 1 may all affect the development and severity of viral infections.<sup>16,17</sup> Also, history of allergic diseases such as ever wheezing, atopic dermatitis, and food/drug allergy was associated with severe pneumonia.<sup>34</sup> Taken together, virus infections have been associated with acute exacerbation of asthma, and allergy may not be a risk factor for virus infection. It is plausible that this concept may also apply to SARS-CoV-2; however, having no asthma patients and no respiratory allergies does not support this concept at least in this series of 140 patients.

Two (1.4%) patients had COPD in our report; the percentage is similar as that reported by Guan et al<sup>4</sup> (12/1099, 1.1%). In addition, only 9 (6.4%) patients had a history of smoking, and 7 of them were past smokers. It was reported that the prevalence of COPD in adults ≥40 years old was 13.7%,<sup>22</sup> and 27.3% of adults in China were current cigarette smokers (data in 2018).<sup>35</sup> The relationship between smoking and coronavirus infection is not clear, and the exact underlying causes of the lower incidence of COVID-19 in current smokers are still unknown. Previous study demonstrated that CoV was not frequently detected by RT-PCR in the exacerbation of COPD, as compared to other respiratory viruses such as RV, EnV, RSV, and IfV, indicating that coronavirus plays a minor role in the acute exacerbation of COPD.<sup>23</sup> Although our study found that COPD and smoking populations were less likely to be infected with SARS-CoV-2, but the outcome of SARS-CoV-2 infection in smokers may be more severe.



**FIGURE 4** Correlation between lymphocyte (LYM) and eosinophil (EOS) absolute numbers ( $\times 10^9/L$ ) in blood from COVID-19-infected patients. A, C, E: in admission; B, D, F: 2nd test after 3 or more days. Spearman's test was used to evaluate the correlation

Smokers and COPD patients are more susceptible to the infection of Middle East respiratory syndrome coronavirus (MERS-CoV). A recent study found that dipeptidyl peptidase IV (DPP4), the receptor for MERS-CoV, had a higher expression in smokers and COPD patients than in nonsmokers. In addition, the expression of DPP4 was inversely correlated with lung function and diffusing capacity parameters.<sup>36</sup>

Angiotensin-converting enzyme 2 (ACE2) is abundantly expressed in airway epithelial cells and was identified as a receptor for SARS-CoV, which plays a crucial role in SARS-CoV-induced lung injury.<sup>37</sup> ACE2 and other components of renin-angiotensin system are the core factors for the control of acute lung injury induced by SARS coronavirus. In vivo and in vitro studies demonstrated that ACE2 expression was downregulated once the disease process has been initiated.<sup>38</sup> Thus, ACE2 may have double effects on SARS-induced lung injury.<sup>39</sup> Initially, it acts as a receptor for the infection of SARS coronavirus, and then, its downregulation promotes lung injury. ACE2 was also identified as the receptor for the novel SARS-CoV-2.<sup>40</sup> The role of ACE2 in the pathogenesis of this new coronavirus-induced lung injury is still unknown. Therefore, further studies are required

to examine the expression of ACE2 in airway epithelia from COPD patients and current smokers.

No significant difference was identified between the severe (7 days) and nonsevere patients (8 days), regarding the median days from symptom onset to hospital admission. Almost all patients (91.7%) had fever at onset of symptoms, consistent with that reported by Huang<sup>2</sup> and Wang,<sup>3</sup> but much higher than that reported by Guan.<sup>4</sup> However, in Guan's study,<sup>4</sup> most patients (87.9%) developed fever during hospitalization. Therefore, fever was the most common symptom in patients with COVID-19. Cough was another common symptom in these patients. 75% patients in our report had cough; the number is similar as that reported by other studies.<sup>1-4</sup> The incidence of fatigue in this study was 75%, which is higher than that reported by Huang and Guan, but close to that reported by Wang. The incidence of gastrointestinal symptoms was 39.6% in our report, which is much higher than that reported by Huang and Guan, but similar as Wang's study. Radiologically, most common signs on CT scan were bilateral ground-glass or patchy opacity.<sup>1-4</sup> This is consistent with what we observed in our report (Figures 1-3). The distribution of the opacity in the lung was not different between severe and nonsevere patients.

Lymphopenia was common in patients of our study (75.4%); this is consistent with other reports.<sup>3,4</sup> Noticeably, the percentages, but not absolute counts of lymphocytes, were lower in severe patients when compared to nonsevere patients. This may be due to the increased total numbers of leukocytes in severe patients.

Interestingly, decreased eosinophil counts were also common in these patients (52.9%). No significant difference in the ratio of patients with decreased eosinophil counts between severe and nonsevere patients was identified. There was a positive correlation between eosinophil and lymphocytes numbers, especially for the second test during hospitalization. Therefore, decreased eosinophil count may be used as an indicator of SARS-CoV-2 infection in suspected patients. In those patients with typical symptoms and radiological changes with and without lymphopenia, decreased eosinophils may be an important diagnostic clue.

Other abnormal laboratory findings include increased level of serum CRP, SAA, PCT, D-dimer, and creatine kinase, indicating sustained inflammatory response and disturbed coagulation mechanism after infection with SARS-CoV-2. In addition, CRP, PCT, D-dimer concentration, and leukocyte counts were significantly higher in severe compared to nonsevere patients, which may represent more prominent inflammation in severe patients. Higher leukocyte count and PCT may also be due to secondary bacterial infection. More precautions should be taken in patients with high serum CK, which may be caused by direct effect of virus or indirect effect of hypoxia. In the present study, data in regard to the treatment and outcome of these patients were not finalized, since most of these patients are remaining hospitalized.

In summary, the study established the clinical and laboratory characteristics of 140 community-infected COVID-19 patients, showing an approximately 1:1 female-male ratio. Low prevalence of smokers and no allergic diseases despite of drug hypersensitivity and urticaria was self-reported by any patients, indicating that allergic diseases and smoking history may not be the susceptible factors for COVID-19. The positive correlation of blood eosinophil and lymphocyte counts suggests that eosinopenia along with lymphopenia may be a useful indicator for diagnosing COVID-19 in those patients with typical symptoms and radiological changes. Larger sample population is needed to further investigate the relationship between SARS-CoV-2 infection and allergic diseases.

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#### CONFLICT OF INTEREST

None of the authors have any conflict of interest to declare.

#### AUTHOR CONTRIBUTIONS

Jinjin Zhang, Xiang Dong, and Yadong Gao collected and analyzed the data, and prepared the manuscript. Yiyuan Cao contributed to the collecting and interpretation of radiological materials. Yadong Yuan, Yibin Yang, and Youqin Yan were involved in the patient management and organization work. Yadong Gao and Cezmi A. Akdis designed the study and reviewed the manuscript.

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