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Patients of COVID-19 May Benefit from Sustained Lopinavir-Combined Regimen and the Increase of Eosinophil May Predict the Outcome of COVID-19 Progression

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Highlights

- Asymptomatic 2019-coronavirus disease patient with normal radiography throughout appeared in this study.
- Lopinavir has an positive effect on 2019-coronavirus disease patients.
- Eosinophil counts presented potentiality as predictor on the progression of 2019-coronavirus disease.

Abstract

Objectives: To explore the epidemiological information, clinical characteristics, therapeutic outcomes and temporal progression of laboratory findings in 2019-coronavirus disease (COVID-19) patients exposed to lopinavir.

Methods: We collected data from ten COVID-19 patients admitted between January 22, 2020 and February 11, 2020 at Xixi hospital in Hangzhou, China.

Results: Of ten patients, secondary, tertiary and quartus patients emerged, the incubation period was 3–7 days. Mainly initial symptoms were cough and low fever (37.3–38.0 °C). An asymptomatic case presented normal radiography, the others existed ground glass opacities. All cases (three transferred, seven discharged) exposed to lopinavir on initial hospitalization. Three patients stopped lopinavir using because of adverse effect, two of them deteriorated, one hospitalized longer than others who with sustained lopinavir using. Levels of potassium, albumin, lymphocyte were low, but increased persistently after treatment. Eosinophil values were low on initial hospitalization, then all returned to normal before discharge. Viral load of SARS-CoV-2, radiography and eosinophil improved continuously in 3–14, 6–8 and 7–9 days, respectively.

Conclusions: Increasing eosinophils may be an indicator of COVID-19 improvement. The COVID-19 patients may benefit from sustained lopinavir using. More researches on a larger scale are needed to verify these points.

Keywords: 2019-Coronavirus disease; Lopinavir; Asymptomatic infection; Eosinophil.

Introduction

At the end of 2019, coronavirus disease (COVID-19) was reported in Wuhan, China. COVID-19 was confirmed to be the pathogenic cause of severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2). The disease is highly contagious, and some cases infected with SARS-CoV-2 were rapidly fatal (The 2019-nCoV Outbreak Joint Field Epidemiology Investigation Team, 2020). More than 74000 cases from 31 provinces in China have been confirmed as of so far (National Health Commission, 2020). Some confirmed cases including American, France, Thailand, Japan and other twenty countries have also been reported. On January 27 of 2020, the World Health Organization (WHO) issued a global vigilance and worldwide surveillance of the outbreak from China, and provided the first edition of infection prevention and control (IPC) guidance, while this interim guidance needs more information to update (WHO, 2020). And for COVID-19, clinical, epidemiological and laboratory data, especially on temporal progression of abnormal laboratory findings remains limited. Herein, we reported epidemiological information, clinical characteristics, therapeutic outcomes and the temporal progression of laboratory findings in COVID-19 patients exposed to lopinavir (LPV) for use in case management.

Methods

Patients and data collection

Between January 22 and **February 11**, 2020, 10 patients whose disease confirmed by two consecutive positive results of SARS-CoV-2 RNA were screened for eligibility. These patients

were admitted through the outpatient clinic in Xixi hospital (a designated hospital for COVID-19, Hangzhou), or were transferred by ambulance from other hospitals in Hangzhou, and isolated in negative-pressure ward immediately. Protective equipment for aerosol-generating was taken by medical workers. This study was approved by the Institutional Review Board of Xixi Hospital.

Epidemiological information was obtained in detail from aspectant interview of these patients. Medical records and laboratory-test results were standardized as a uniform tabulation which were double checked independently by two researchers.

Radiologic assessment

The chest radiograph of each patient was evaluated without consulting clinical data. High-resolution computed tomography (HRCT) scanning of the chest was available and evaluated in all patients. The presence of opacification was classified as air-space shadowing (ground-glass opacity, focal consolidation or patchy consolidation), reticular or diffuse (all lung involvement). Pleural effusions were also marked.

Detection of SARS-CoV-2

Four respiratory samples (two nasal swabs and two throat swabs) were collected from each patient. A total of 300 μ L sample from nasal swab tubes (150 μ L per tube) was added into short-time cracking reaction fluid (MVR01, Liferiver, Shanghai, China), throat swabs were treated in the same way. Total RNA was extracted according to the manufacturer's instructions and 5 μ L RNA was used for real-time RT-PCR, which targeted E gene (first line screen assay), RdRP gene (confirmatory assay) and N gene (additional confirmatory assay) by using Multiple Fluorescence Quantitative PCR reagent. The reaction conditions of real-time RT-PCR started with 45 °C for 10 min and 95 °C for 3 min, followed by 45 cycles of amplification at 95°C for 15 s and 58°C for 30 s. The test results of SARS-CoV-2 were reported as positive (Cycle threshold; Ct \leq 35), weak

positive ($35 < Ct \leq 43$) and negative ($Ct > 43$).

Definitions

The definitions of this research were acute myocardial injury (AMI, troponin I $>0.2 \mu\text{g/L}$) (Dous et al., 2017), acute kidney injury (AKI, serum creatinine $>26.5 \mu\text{mol/L}$ in 48 hours or urine volume $<0.5\text{mL/kg/h}$ for 6 hours) (Kellum et al., 2012), respiratory failure (arterial partial oxygen pressure $<60 \text{ mm Hg}$) (Hamp et al., 2017), digestive adverse effect (the clinical symptom of diarrhea, nausea and vomiting), low potassium (serum potassium $<3.5 \text{ mmol/L}$) and hypoalbuminemia (serum albumin $<35 \text{ g/L}$) (Touma and Bisharat, 2019).

Statistical analysis

Continuous variables were represented with median (interquartile range, IQR). Linear regression model was used to estimate the variables change over time. All analyses were performed using the R software, version 3.3.1 (<http://www.R-project.org>).

Results

The median age of the 10 patients (6 females) was 42 (IQR, 34–50). Except as noted below, all patients had inconspicuous medical history. Patient 3 had hypertension and cardiovascular disease, was taking metoprolol and nifedipine. Patient 7 had chronic liver disease, was taking tenofovir. Nine patients had no smoking history. Patient 7 was a current smoker, 20 cigarettes per day for twenty years.

Contact history

The contact history of 10 patients was shown in Figure 1. Patient 1, a Hangzhou resident, was a clothing salesperson, who went to Wuhan garment market on January 11, 2020, and get back to Hangzhou on January 15. One day later, fever as the earliest symptom occurred. She was admitted to hospital A with pneumonia. Patient 6 was the husband of patient 1 and had close contact with

patient 1 for more than two days in home before the onset of symptom. The couple (patients 1 and 6) were transferred to hospital B (Xixi hospital) from hospital A on January 24 by ambulance after the laboratory test of SARS-CoV-2-RNA existed positive. Patients 2, 3, 4 and 5 were all Wuhan resident. They came to Hangzhou for a family reunion. Patients 3 and 5 have already had the symptom of fever when they were in Wuhan. Patient 7 was a Hangzhou resident, went to Wuhan for business investigation for two days. He had the symptom of chest congestion on 8th day after Wuhan exposure. Patient 8, a women living in Hangzhou, had a meeting with her colleagues from Wuhan on January 15. She had the symptom of fever on January 20. Patients 9 and 10 were a couple. Patient 10 went to Beijing for a conference for two days and had close contact with her colleagues who came from Wuhan, then returned to Hangzhou on January 13. Patient 9 lived with patient 10 in home for more than six days before the onset of symptom.

As shown in Figure 1, all patients had a direct or indirect history of Wuhan exposure. The maximum incubation period, which was estimated as the number of days from likely exposure to the onset of symptom, could be accurately assessed for patients 1, 6, 7, 8, 9 and 10 (five, three, seven, five, six and four days, respectively). The contact history of patients 2, 3, 4 and 5 was indistinct, that was not estimated precisely.

Clinical and other characteristics

Most patients presented cough and fever (mainly low fever, temperature ranges from 37.3 to 38.0) (Table 1). Four patients had the symptom of productive cough (white phlegm) and sore throat. Three patients reported headache and nausea. One patient only existed chest congestion (patient 7). None of the patients presented with rhinorrhea, rigor, diarrhea and dyspnea. Leukopenia (less than 3.5×10^9 leucocytes per liter) and lymphopenia (less than 1.1×10^9 lymphocytes per liter) were observed in two patients, respectively.

Among these patients, five patients (patients 1, 2, 3, 6 and 8) developed serious illness (resting oxygen saturation less than 93% or arterial partial oxygen pressure <60 mm Hg or respiratory rate more than 30 times per minutes), three (patients 1, 2 and 3) of them were transferred to the First Affiliated Hospital of Zhejiang University hospital for further treatment. **Seven patients (patients 4, 5, 6, 7, 8, 9 and 10) cured and discharged from hospital, on February 6, 8, 11, 2, 3, 2 and 7, respectively.** Two severe cases (patients 1 and 2) combined with multiple symptoms (cough, white phlegm, headache, nausea and fever) before admission. In discharged patients, patient 7 had just one symptom (chest congestion), while developed anxiety after admitted to hospital.

Five patients had not received any treatment before the onset of symptom. The results of cardiovascular, abdominal, and neurologic examination were normal in all patients, in consideration of their preexisting conditions.

Radiologic assessment

All patients had abnormal chest radiographs except patient 7. The main abnormalities in chest CT images were air-space shadowing: patchy ground-glass opacities (in patients 2, 3, 6 and 9) (Figure 2A) or patchy hyperdense areas (patients 1, 4, 5, 8 and 10) (Figure 2B) on the initial chest radiograph. The radiographic opacities were mostly in the lower lung zones (patients 1, 2, 3, 4, 5, 6, 8 and 9). Patient 6 presented with pleural effusions.

The radiographic lesions progressed throughout in size, extent and severity in three patients (patients 1, 2 and 3) after admission. Patchy consolidation was found in lower lung zone in patient 6. Some improvement on lesions occurred in patients 4, 5, 8, 9 and 10 within the first eight days after presentation. The CT-images of patient 8 on second, fourth, sixth and eighth day after admission were shown on Figure 2C, the progression of the lesions occurred on 4th day and the improvement appeared on 6th day, the opacities shrunk significantly on the 8th day after admission.

Laboratory findings

Figure 3 revealed the changes over time in fifteen laboratory indicators after the beginning of therapy. As can be seen from the figure, most values of temperature dropped below 37.5 degrees Celsius for four days treatment. During the therapeutic period, over half of the lymphocyte counts (normal range: $1.1\text{--}3.5 \times 10^9$ per liter), eosinophil counts (normal range: $0.02\text{--}0.52 \times 10^9$ per liter), artery potassium concentrations (normal range: 3.4–5.5 mmol/L) and serum albumin levels (normal range: 40–50 g/L) were under the lower threshold of normal range, but all presented a rising trend, particularly evident in eosinophils.

The eosinophil values of patients 1, 2 and 3 were all less than the lower bound of normal range before they were transferred to the First Affiliated Hospital of Zhejiang University (Figure 4). Other patients, except for patient 10, also had similar findings at the early stage of COVID-19. Eosinophil counts of patients 4, 5, 6, 7, 8 and 9 returned to normal in nine days after treatment (Figure 3), and were discharged within 7, 9, 12, 3, 2 and 4 days after the persistent improvement sign of eosinophil appeared (Figure 4). Two mild patients (patients 7 and 9) had a shorter improvement period of eosinophil than the other two severe cases (patients 6 and 8).

The results of SARS-CoV-2-RNA existed positive in patients 1, 2 and 3 before transferred. Patient 7 first showed the signs of improvement in virology on 3th day after treatment, and others presented such signs between 7 and 14 days after admission. The negative results of SARS-CoV-2-RNA of patients 7 and 9 occurred within 7 and 9 days after admission, earlier than that of patients 4, 5, 6, 8 and 10 (12, 14, 18, 11 and 11 days, respectively). During the period of lower eosinophil, SARS-CoV-2-RNA presented positive, and after eosinophil increased to normal, SARS-CoV-2-RNA turned to negative within 5 days, except for patient 6 (11 days).

Most values of C-reactive protein (CRP, normal range: 0–10 mg/L) were above the upper

bound of normal range, and over half of the fibrinogen levels (FIB, normal range: 1.8–3.5 g/L) were high during the observed period (Figure 3). Leucocyte count (normal range: $3.5\text{--}9.5 \times 10^9$ cells/L), platelet count (PLT, normal range: $125\text{--}350 \times 10^9$ cells/L), activated partial thromboplastin time (APTT, normal range: 23–32.6 s), total bilirubin level (normal range: 3.42–20.52 mmol/L), creatinine concentration (normal range: 41–81 $\mu\text{mol/L}$) and creatine kinase value (normal range: 24–170 U/L) were mainly normal in these cases (Figure 3). An increasing trend toward improved oxygen saturation, oxygen partial pressure (PO₂) and platelet count was found after initiating therapy.

Treatments and outcomes

The median duration between the onset of symptom and initiating therapy was five days (IQR, 3–6). All patients were treated with combined therapy (lopinavir, LPV, 400mg every twelve hours and interferon $\alpha 2b$ atomization inhalation, 5 million U twice daily) or single drug of LPV (in patient 7 only) (Table 2). Before the initiation of treatment, patients 1, 2, 4, 6 and 8 received antibiotics for a median of three days (IQR, 2–4) (Table 1). But not eased the symptoms of fever and cough.

As indicated in Table 2, the median observed period was 13 days (IQR, 4–17). During this period, all of the patients had no AMI and AKI, five patients reported digestive adverse effect (diarrhea or emesis). Seven patients had hypokalemia. All patients received oxygen inhalation through nasal tube except patient 7.

Patients 1 and 2 had severe digestive adverse effect (diarrhea and emesis) and hypokalemia within four days of treatment (LPV + interferon $\alpha 2b$). The combined therapy was changed empirically to interferon $\alpha 2b$ atomization inhalation (5 million U twice daily) and arbidol hydrochloride granules (AHG, 0.2g, three times daily) and human immunoglobulin intravenous

injection (IVIG, 20g every day) and methylprednisolone (40mg, every twelve hours). But, the clinical symptom and radiography still presented deterioration. Patient 3 existed complex underlying diseases and combined with mycoplasma pneumonia. Because becoming deterioration, patients 1, 2 and 3 were transferred to the First Affiliated Hospital of Zhejiang University within four days (Table 2).

Among discharged cases, patient 7 had unremarkable radiographic patterns throughout after admission. The opacities on CT images of patients 4, 5, 9 and 10 improved persistently after eight, seven, six and seven days of LPV-combined regimen exposure, respectively. Another discharged case (patient 8) presented severe diarrhea, hypokalemia, respiratory failure on the 4th day and hypoproteinemia on the 5th day of taking the medicine of LPV and interferon α 2b. In view of this, somac (one tablet daily), Human Albumin Solution (HAS, 10g per day), AHG (0.2g, three times daily), IVIG (20g every day) and methylprednisolone (40mg, every twelve hours) were added to the original treatment regimen, without stopping LPV. The adverse reaction, complications and radiographic opacities (Figure 2C) of patient 8 were significantly improved in two days. **Patient 6 stopped the usage of LPV due to digestive adverse reaction after four days of treatment. The period of improvement of radiographic lesions was 14 days, more longer than others (8 days). Because of the results of SARS-CoV-2 RNA presented weakly positive, he remained hospitalized when other six patients discharged. The result of SARS-CoV-2-RNA of patient 6 presented negative on 18th day after admission, later than other discharged cases. Fitness for discharge was based on improvement of chest radiographic evidence, with viral clearance in respiratory samples and abatement of fever for at least 3 days.**

Discussion

Our report of these 10 patients revealed that COVID-19 is a contagious and rapidly

progressive disease. Among these cases, secondary (patients 1, 2, 3, 4, 5 and 7), tertiary (patients 8 and 10) and quartus (patients 6 and 9) patient emerged. Four patients had a history of interaction with source patients in a room, which confirmed that contact (however trivial) is one of the routes of transmission. There was no further transmission among our healthcare workers in our hospital after we implemented appropriate isolation precautions involving use of isolation gowns, protective suits, N95 masks, gloves, and hands disinfecting before and after contacting with patients. Which protective approach has an effect on the decrease in transmission remains uncertain. These findings revealed that person-to-person transmission and interprovincial spread are possible, which consistent with the results of the lasted researches (Huang et al., 2020; Chan et al., 2020), control of travel and strict protection are essential pre-requisite.

The incubation period of COVID-19 was between 3 and 7 days in those patients could be accurately assessed. Such incubation period is similar to the recent report which estimated ranging from 3 to 6 days (Chan et al., 2020), but longer than that of SARS (Wang et al., 2004). This further complicates the epidemiological investigation and singles out at-risk exposure to source patients even without any symptoms as the key factor in the diagnosis of COVID-19.

In this cluster of 10 patients, most of them presented with fever, cough and bilateral lesions of lung. Two severe cases are combined with multiple early symptoms: cough, white phlegm, headache, nausea and fever. In seven discharged patients, we could not find a universal characteristic of clinical symptom before admission. But patient 7 had no symptom except for chest congestion and anxiety. It is also of note that the radiographic images of patient 7 was normal throughout until discharge. For the above points, it is suspected that chest congestion may be caused by anxiety. However, this findings is different from another asymptomatic infection that reported in a child aged 10 years without any clinical symptoms, but, existed ground-glass lung

opacities (Chan et al., 2020). From this point of view, asymptomatic patients like in our study will make the diagnosis of COVID-19 even more difficult, and diagnostic delays may further lead to the spread of the epidemic. However, whether asymptomatic infection actually exists needs more clinical evidences.

The radiological appearance of COVID-19 is mostly ground-glass opacity with patchy area, predominantly affecting the lower lobe. Focal absorption occurs in size, extent, and severity, following with features of the disease entering to fibrotic phase (Figure 2C). In most patients, radiological lesions changed evidently and rapidly in a few days after onset of symptom. Hence, timely radiographic observation and evaluation are necessary in realizing the progression and recovery of COVID-19.

Most patients had hypokalemia and digestive adverse effect in four days after the beginning of treatment, which could be caused by LPV exposure ((Dybul et al., 2002). In view of this, three severe cases (patients 1, 2 and 6) were taken off LPV empirically, instead, their therapies were replaced with glucocorticoid and interferon-backbone combination regimen. However, the conditions of patients 1 and 2 had no improvement, even aggravate after the withdrawal of LPV. **The result of SARS-CoV-2-RNA of patient 6 remained positive for four days after that of other six discharged cases who with continuous using of LPV turned to negative. Noticeably, one** severe case (patient 8) received support care and symptomatic treatment after the adverse reactions occurred, rather than stopping LPV, with improvement appeared in two days. The experience therefore suggests that receiving LPV is effective on disease recovery, the efficacy of glucocorticoid and interferon is not evident on cases infected with SARS-CoV-2 (Wang et al., 2020).

In this study, the findings of high CRP level (mostly between 10 and 100 mg/L) and increased

FIB concentration, consistent with that of recent study (Chan et al., 2020), the counts of white blood cell and PLT were normal in most cases, that was also shown in another research of COVID-19 (the normal percentage were 45% and 95%, respectively) (Huang et al., 2020). However, a large proportion of our cases presented low levels of potassium, albumin, lymphocyte and eosinophil in early stage of COVID-19, especially was that almost all eosinophil values were less than the lower threshold of normal range within the first week of hospitalization. This symptom could not be attributed to glucocorticoid exposure, since seven patients had not received such treatment at initial duration of admission. However, the reduction of eosinophils may be related to a mechanism of stress response in the condition of acute lung injury caused by SARS-CoV-2, which results in inhibiting the release of eosinophils in the marrow through glucocorticoid secretion. Interestingly, there was continuous improvements on eosinophils in the later stage of COVID-19, which was in sync with the symptom of radiographic and virology improvements in all discharged cases, and the time of eosinophil improvement of mild patients was earlier than that of severe cases, suggests that lower and increasing eosinophils may be the signs of COVID-19 progression and recovery. Combined with above abnormal indicators, whether it can be regarded as diagnostic characteristics of COVID-19 needs more research to determine. Although, such abnormalities existed in the early stage of COVID-19, there was an improving trend of these indicators after receiving LPV-combined regimen. This was further evidence that LPV has an positive effect on the patients infected with SARS-CoV-2.

Our study has limitations, the inherent shortcomings belong to retrospective observational single-center study, small sample size and short term follow-up make it difficult to address the relationships between COVID-19 and LPV, eosinophil, and reach a firm conclusion. All above limitations require further study to be overcome, nonetheless, our primary results provided

moderate yet important illumination for this topic.

Conclusions

Considering the information we described, taking appropriate isolated precautions with source patients suspected to have COVID-19 is vital important. Asymptomatic infection (as reported in patient 7) appears possible, strict infection control is crucial for avoiding the second pandemic. No effective therapy is known for COVID-19 at present. The use of LPV appears effective in this study, but digestive adverse effect and hypokalemia should be concerned by clinicians after the beginning of this drug. More clinical evidences are also needed imminently to assess the efficacy and safety of LPV in patients with SARS-CoV-2 infection. In addition, eosinophil counts presented potentiality as predictor on the development process of COVID-19 in this study. Further investigations and researches on a larger scale are worth to do to verify this point.

Authors' Contributors

LF and ZJ conceived, designed, and organized the study, interpreted the results, and drafted the manuscript. XAF helped supervise the study. The other authors contributed to collect the data on site.

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Ethics approval and consent to participate

This study was approved by the Institutional Review Board of Xixi Hospital. All data were anonymized to comply with the provisions of personal data protection legislation. Due to the retrospective nature of this study and due the fact that only historical medical data were collected, written informed consent was not required.

Conflict of interest statement

The authors declare that they have no competing interests.

Declarations of interest: none.

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Availability of data and materials

The data set used for this manuscript will be available from the corresponding author upon reasonable request.

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Figure 1. Contact history and timeline of the onset of symptoms in COVID-19 patients.

Figure 1

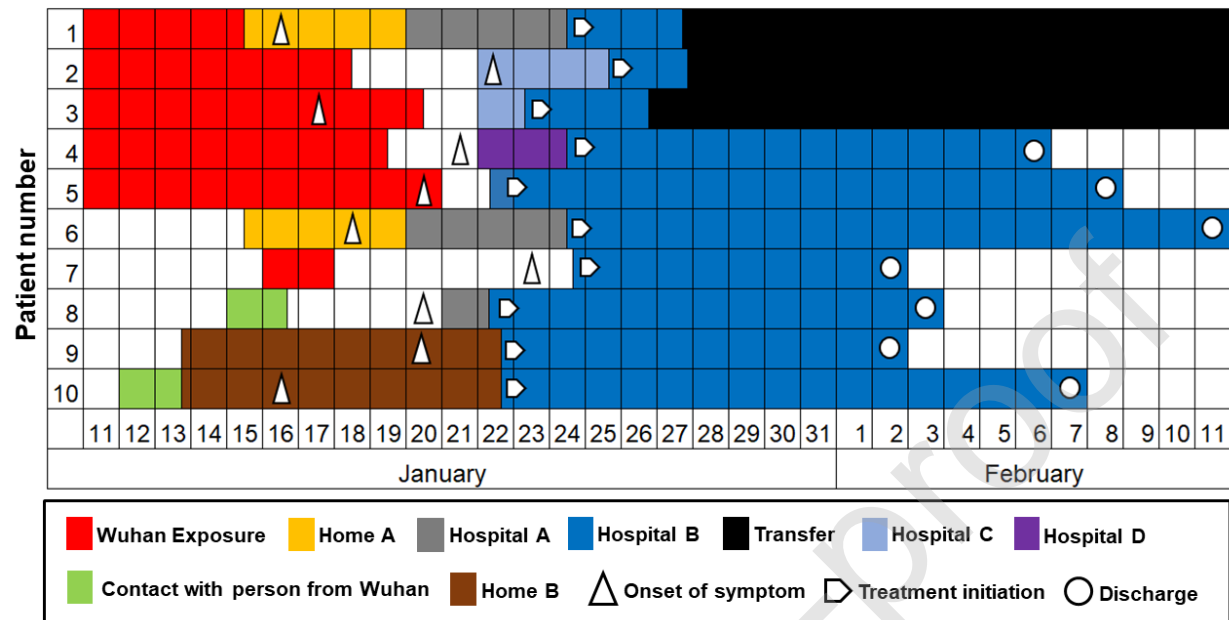


Figure 2. Chest radiographs showing patchy ground-glass opacities (panel A: patient 3), patchy hyperdense areas (panel B: patient 1) and multifocal opacity changes over time (panel C: patient 8).

Figure 2

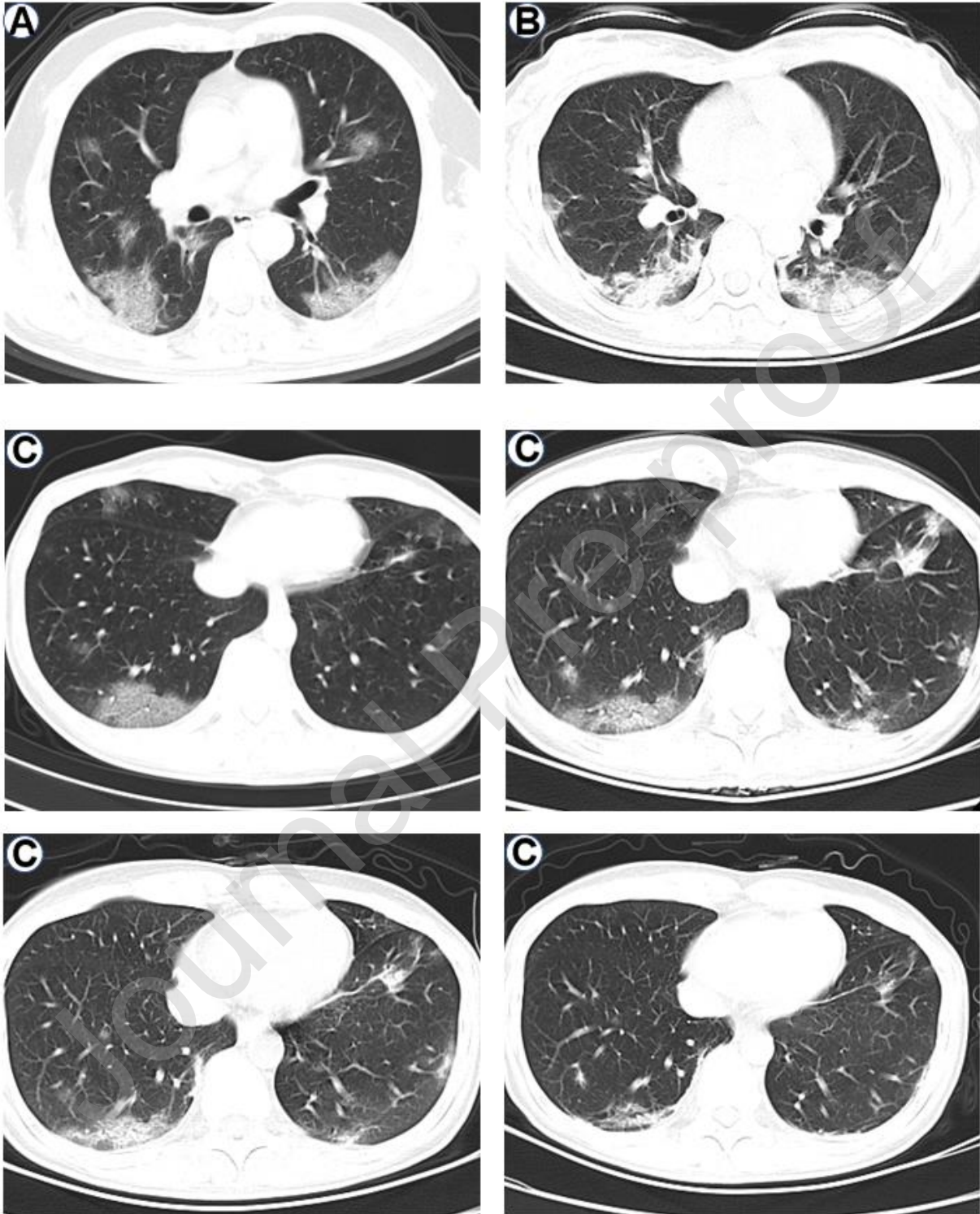


Figure 3. The temporal changes of laboratory indicators in COVID-19 patients after initiation of LPV-

combined therapy.

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Figure 3

○ Value — Fitted line 95% CI — Lower bound - - - Upper bound

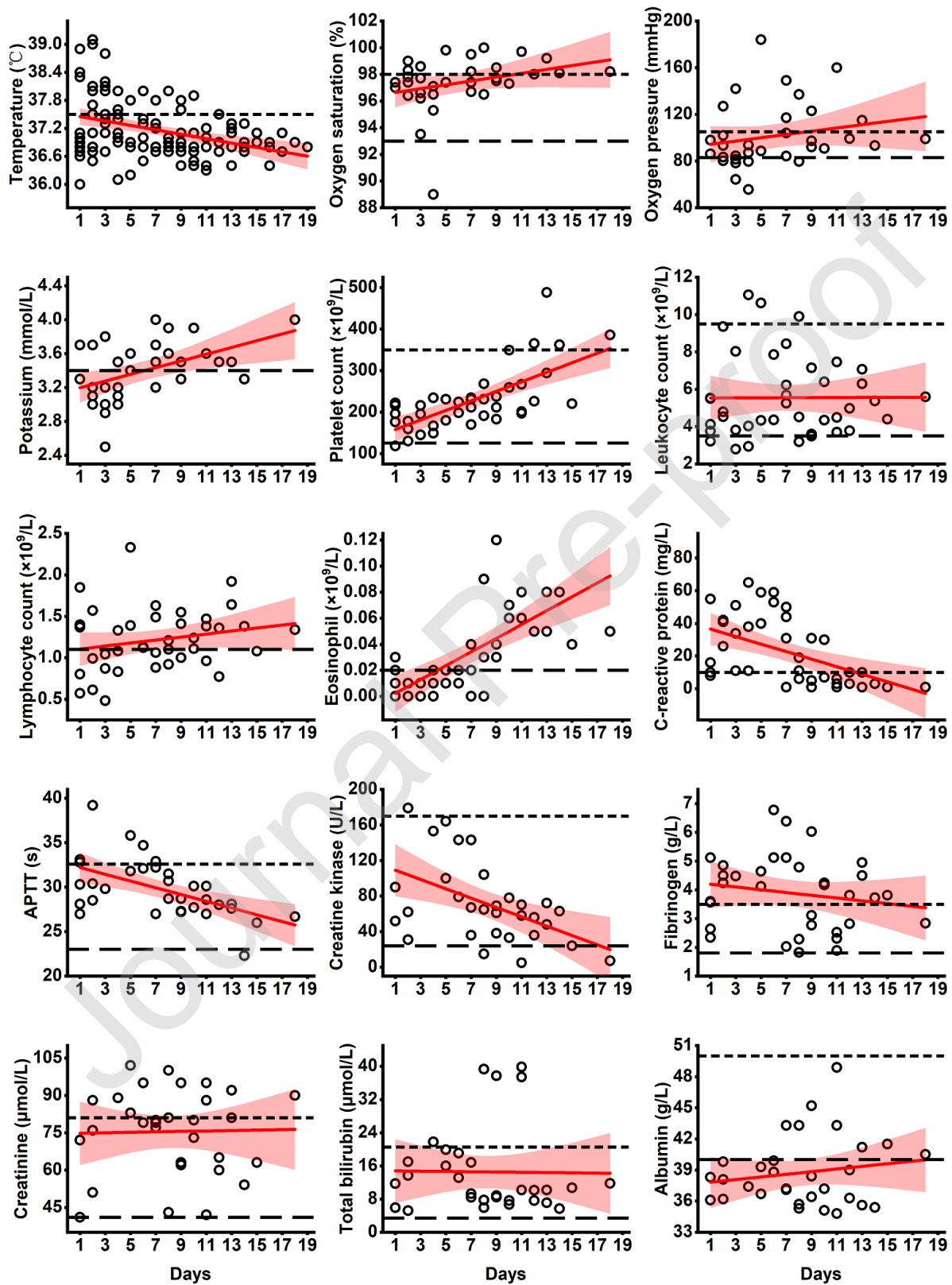
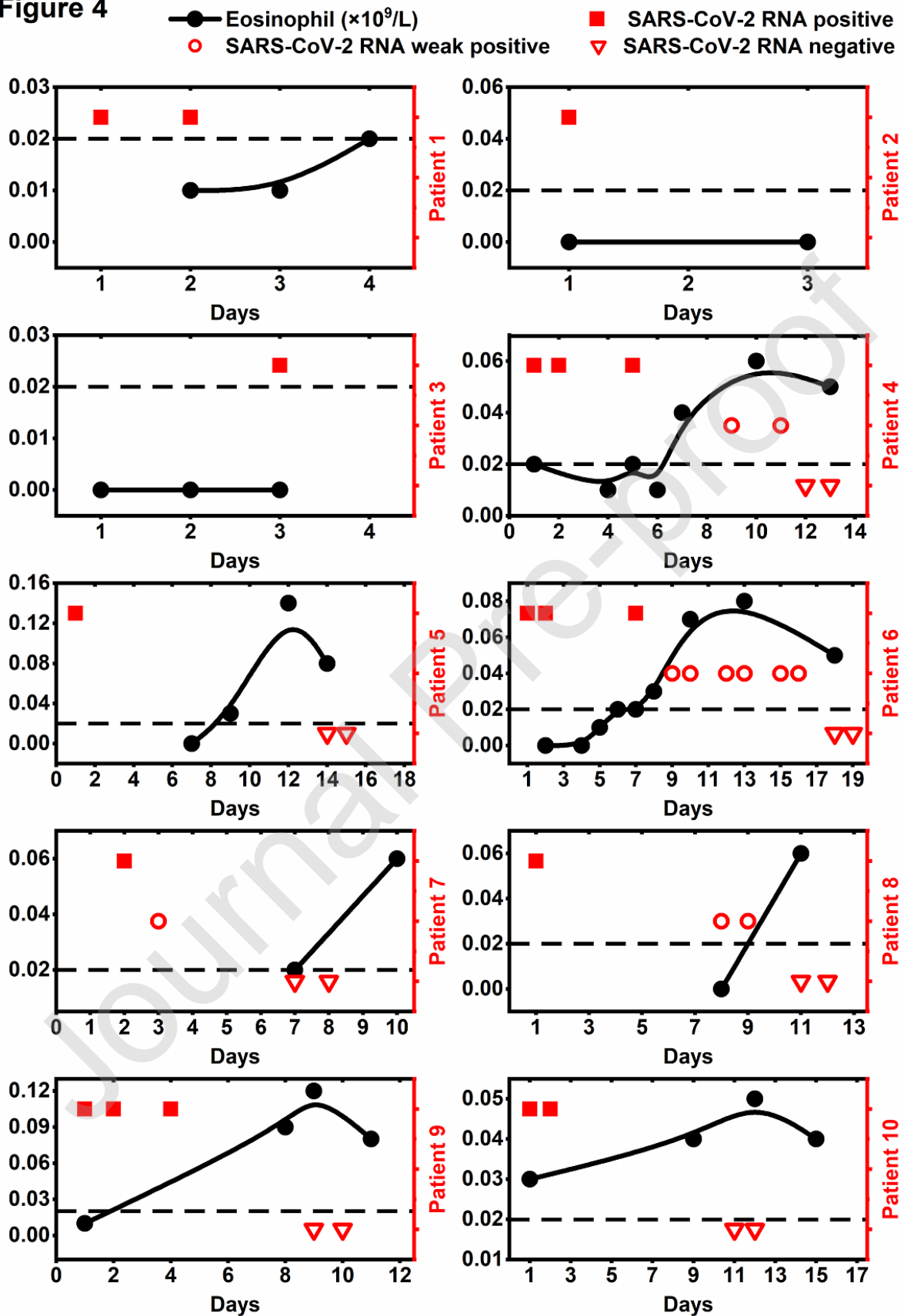


Figure 4. The changes over time on eosinophil level and SARS-CoV-2 viral load in each COVID-19 patient.

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Figure 4



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Table 1. Characteristics of COVID-19 Patients on Presentation

	Patient no.										Summary
	1	2	3	4	5	6	7	8	9	10	
Sex	F	F	M	F	F	M	M	F	M	F	—
Age (year)	45	30	62	53	51	47	40	33	34	35	42 (34–50)
Current smoking	-	-	-	-	-	-	+	-	-	-	1+
Symptoms											
Cough	+	+	+	+	+	+	-	+	+	-	8+
White phlegm	+	+	+	-	-	+	-	-	-	-	4+
Headache	+	+	-	+	-	-	-	-	-	-	3+
Nausea	+	+	-	-	-	-	-	+	-	-	3+
Sore throat	-	-	-	+	+	-	-	-	+	+	4+
Rhinorrhea	-	-	-	-	-	-	-	-	-	-	—
Rigor	-	-	-	-	-	-	-	-	-	-	—
Diarrhea	-	-	-	-	-	-	-	-	-	-	—
Chest congestion	-	-	-	-	-	-	+	-	-	-	1+
Fever	+	+	-	+	-	+	-	+	+	+	7+
Highest Temperature											
<37.3	-	-	+	-	+	-	+	-	-	-	3+
37.3-38.0	+	-	-	+	-	+	-	-	+	+	5+
38.0-39.0	-	+	-	-	-	-	-	+	-	-	2+
>39.0	-	-	-	-	-	-	-	-	-	-	—
Anxiety	-	-	-	-	-	-	+	-	-	-	1+
Dyspnea	-	-	-	-	-	-	-	-	-	-	—
Comorbidity											
Diabetes	-	-	-	-	-	-	-	-	-	-	—
Hypertension	-	-	+	-	-	-	-	-	-	-	1+
Cardiovascular disease	-	-	+	-	-	-	-	-	-	-	1+
Malignancy	-	-	-	-	-	-	-	-	-	-	—
Chronic liver disease	-	-	-	-	-	-	+	-	-	-	1+

Renal dysfunction	-	-	-	-	-	-	-	-	-	-	-	—
Physical signs												
WBC decreased ^a	-	-	-	+	-	-	-	-	+	-	-	2+
CRP increased ^b	-	+	+	+	-	-	-	-	-	-	-	3+
Lymphocyte decreased ^c	+	-	-	-	-	-	-	-	+	-	-	2+
Systolic pressure (mm Hg)	116	132	178	136	150	126	120	102	132	100		129 (117–135)
Respiratory rate (times/min)	19	20	18	17	17	19	19	18	18	18		18 (18–19)
Current medicine												
Antibiotics	+	+	-	+	-	+	-	+	-	-	-	5+
Antipyretics	+	-	-	-	-	+	-	+	-	-	-	3+
Exposure												
Wuhan exposure	+	+	+	+	+	-	+	-	-	-	-	6+
Contact with person from Wuhan	-	-	-	-	-	+	-	+	+	+	+	4+

A plus sign represents the data or symptom was present, and a minus sign indicates absent. WBC: white blood cell count.

a WBC decreased: below the normal range of WBC ($3.5\text{--}9.5 \times 10^9$ cells/L)

b CRP increased: above the normal range of C-reactive protein (0–10 mg/L)

c Lymphocyte decreased: below the normal range of lymphocyte ($1.1\text{--}3.2 \times 10^9$ cells/L)

Table 2. Treatments and outcomes of COVID-19 patients

	Patient No.										Summary
	1	2	3	4	5	6	7	8	9	10	
Complications											
Acute myocardial injury	-	-	-	-	-	-	-	-	-	-	—
Acute kidney injury	-	-	-	-	-	-	-	-	-	-	—
Respiratory failure	+	-	-	-	-	-	-	+	-	-	2+
Digestive adverse effect	+	+	-	-	-	+	-	+	-	+	5+
Low potassium	+	+	-	+	+	+	-	+	-	+	7+
Hypoalbuminemia	-	-	-	-	-	-	-	+	-	-	1+
Treatment											
LPV	+	+	+	+	+	+	+	+	+	+	10+
RH-interferon $\alpha 2b$	+	+	+	+	+	+	-	+	+	+	9+
Antibiotics	-	-	+	+	-	-	-	-	-	-	2+
Glucocorticoid	+	+	-	-	-	-	-	+	-	-	3+
Immunoglobulin	+	+	-	-	+	+	-	+	-	-	5+
Oxygen support											
Nasal cannula	+	+	+	+	+	+	-	+	+	+	9+
Non-invasive ventilation	-	-	-	-	-	-	-	-	-	-	—
Invasive mechanical ventilation	-	-	-	-	-	-	-	-	-	-	—
Prognosis											
Hospitalization days	4	3	4	14	18	19	10	13	12	17	13 (4–17)
Intensive transfers	+	+	+	-	-	-	-	-	-	-	3+
Chest radiographs											
Bilateral lesions of lung	+	+	+	+	+	+	-	+	-	+	8+

Pneumonia progression > 50%	+	-	-	-	-	-	-	-	-	-	1+
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LPV: lopinavir; RH-interferon α 2b: Recombinant Human Interferon α 2b Injection; Pneumonia progression > 50%: Lung pattern of the patient progressed more than 50% in 48 hours.

Journal Pre-proof