

ARTICLE

The Potential of Low Molecular Weight Heparin to Mitigate Cytokine Storm in Severe COVID-19 Patients: A Retrospective Cohort Study

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On March 11, 2020, the World Health Organization declared its assessment of coronavirus disease 2019 (COVID-19) as a global pandemic. However, specific anti-severe acute respiratory syndrome-coronavirus 2 (SARS-CoV-2) drugs are still under development, and patients are managed by multiple complementary treatments. We performed a retrospective analysis to compare and evaluate the effect of low molecular weight heparin (LMWH) treatment on disease progression. For this purpose, the clinical records and laboratory indicators were extracted from electronic medical records of 42 patients with COVID-19 (21 of whom were treated with LMWH, and 21 without LMWH) hospitalized (Union Hospital of Huazhong University of Science and Technology) from February 1 to March 15, 2020. Changes in the percentage of lymphocytes before and after LMWH treatment were significantly different from those in the control group ($P = 0.011$). Likewise, changes in the levels of D-dimer and fibrinogen degradation products in the LMWH group before and after treatment were significantly different from those in the control group ($P = 0.035$). Remarkably, IL-6 levels were significantly reduced after LMWH treatment ($P = 0.006$), indicating that, besides other beneficial properties, LMWH may exert an anti-inflammatory effect and attenuate in part the “cytokine storm” induced by the virus. Our results support the use of LMWH as a potential therapeutic drug for the treatment of COVID-19, paving the way for a subsequent well-controlled clinical study.

Study Highlights

WHAT IS THE CURRENT KNOWLEDGE ON THE TOPIC?

☑ Our results strongly suggest low molecular weight heparin (LMWH) as an effective strategy in a therapeutic or combination therapy against coronavirus disease 2019 (COVID-19).

WHAT QUESTION DID THIS STUDY ADDRESS?

☑ LMWH exerts an anti-inflammatory effect by means of reducing IL-6 and increasing lymphocyte%. We, therefore, favor the use of LMWH as a potential therapeutic drug for the treatment of COVID-19.

WHAT DOES THIS STUDY ADD TO OUR KNOWLEDGE?

☑ A new therapeutic approach for COVID-19 was proposed based on the non-anticoagulant properties of LMWH.

HOW MIGHT THIS CHANGE CLINICAL PHARMACOLOGY OR TRANSLATIONAL SCIENCE?

☑ In view of the COVID-19 pandemic, our study will be of pronounced interest to a broad spectrum of clinicians and scientists of several disciplines focusing on translational and basic aspects related to COVID-19 and virology in general.

On March 11, 2020, the World Health Organization (WHO) declared its assessment of coronavirus disease 2019 (COVID-19) as a global pandemic. Severe acute respiratory syndrome-coronavirus 2 (SARS-CoV-2) is characterized by a long incubation period, high infectivity, and multiple routes of transmission.^{1,2} However, no effective medicines are currently available, so patients are treated symptomatically. A better understanding of the mechanisms of

pathological changes will help to screen potential drugs out of the currently available medications.

Several clinical studies revealed that cytokine storms are important mechanisms underlying disease exacerbation and death of patients with COVID-19.^{3–5} Particularly, IL-6 levels in severely ill patients were significantly higher than in mild cases.⁶ IL-6 is one of the core cytokines,⁷ contributing to many of the key symptoms of cytokine storm, such

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as vascular leakage, activation of the complement, and coagulation cascades, inducing disseminated intravascular coagulation.^{8,9} Reducing the levels of IL-6 and decreasing its activity may prevent or even reverse the cytokine storm syndrome,¹⁰ thereby improving the condition of patients with COVID-19.

Substantial studies have reported that low molecular weight heparin (LMWH) has various non-anticoagulant properties that play an anti-inflammatory role by reducing the release of IL-6.^{11–13} However, the anti-inflammatory effects of LMWH in COVID-19 are currently unknown. By analyzing the effect of LMWH in patients with COVID-19, our retrospective cohort study demonstrates, for the first time, the significant beneficial effect of LMWH in controlling cytokine storm and delaying disease progression (**Figure 1**).

METHODS

Research subjects

To investigate the therapeutic effect of LMWH on COVID-19, we conducted a retrospective cohort study. All cases in this study were located at Union Hospital, Tongji Medical College, Huazhong University of Science and Technology (Wuhan, Hubei Province, China), a designated treatment hospital for patients with COVID-19. This study was approved by the institutional review board of the hospital. In total, we retrospectively collected the electronic medical records of 42 patients with COVID-19, the admission data for these patients were from February 1, 2020, to March 15, 2020 (**Figure 2** shows the case inclusion flowchart) of which 21 patients underwent LMWH treatment (defined as the LMWH group, **Table 1** presents the LMWH medication), and 21 did not (defined as the control group), during hospitalization. As a designated hospital for the treatment of patients with COVID-19, our hospital received 850 patients during February 1, 2020, to March 15, 2020. After March 15, our hospital no longer undertook the treatment of patients with COVID-19. Case screening was performed after all patients were discharged from the hospital. Among these, 548 of nonsevere patients (diagnosed according to the New Coronavirus Pneumonia Diagnosis Program (7th edition) published by the National Health Commission of China) were excluded. Of the 302 severe patient group, 145 received LMWH of which 124 were excluded, as indicated in **Figure 2**. At this point, we went back to the 157 patients who were not treated with LMWH and decided to enroll the “first in list” 21 patients who were found suitable given the exclusion criteria that are presented in **Figure 2**. Notably, 32 patients who were not treated with LMWH were eligible for the study, but only the first 21 patients were actually included without any matching attempts or adherence to specific criteria.

Inclusion criteria were: (i) patients were diagnosed as having COVID-19 according to the New Coronavirus Pneumonia Diagnosis Program (7th edition) published by the National Health Commission of China, including any of the following: novel coronavirus nucleic acid was positive by real-time polymerase chain reaction fluorescence; the virus gene is highly homologous to a novel coronavirus; serum novel coronavirus specific IgM antibody and IgG antibody were positive, or serum novel coronavirus-specific IgG antibody

changed from negative to positive, or the recovery period was four times higher than the acute phase; (ii) the clinical classification was severe, including any of the following: shortness of breath, respiratory rate ≥ 30 bpm; blood oxygen saturation $\leq 93\%$ (at rest); PaO₂/FiO₂ ≤ 300 mm Hg; pulmonary inflammation that progresses significantly within 24–48 hours $> 50\%$; (iii) age ≥ 18 years old; (iv) no history of bronchiectasis, bronchial asthma, or other respiratory diseases; and (v) no immunosuppressant or glucocorticoid use during treatment.

Exclusion criteria: (i) patients with severe systemic diseases and other acute or chronic infectious diseases; (ii) patients with liver and kidney insufficiency or congenital heart disease; (iii) patients who had been treated with LMWH in the previous 3 months; (iv) patients with a history of mental illness; (v) pregnant or lactating women; (vi) patients clinically classified as critically ill or housed in the intensive care unit; and (vii) patients allergic to LMWH or contraindicated for LMWH.

Data collection

The basic information, complete blood count, coagulation profile, inflammatory cytokines, and serum biochemical indicators (including liver function, kidney function, lactate dehydrogenase, C-reactive protein (CRP) and electrolytes) of 42 patients with COVID-19 were retrospectively analyzed. Two researchers also independently reviewed the data collection forms to double-check the data collected.

Statistical analysis

Data analysis was performed using SPSS 22.0 statistical software. Data are expressed as mean \pm SD. GraphPad version 6.0 software was used for plotting. Differences between groups were evaluated using the unpaired two-sided Student's *t*-test for continuous measurement data, and the χ^2 test for count data. The Kruskal–Wallis nonparametric test was used for the comparisons between independent groups, and paired analysis was performed within groups (related samples). Differences of $P < 0.05$ were considered statistically significant.

RESULTS

General characteristics of patients with COVID-19

As shown in **Table 2**, the LMWH group consisted of 13 men and 8 women aged between 42 and 91 years (median age = 69.0 years), and the control group consisted of 14 men and 7 women aged between 40 and 84 years (median age = 69.0 years). There were no significant differences in comorbidities, onset symptoms, and antiviral treatment between the two groups. These results indicate that the general characteristics of the two groups of patients were consistent and comparable.

LMWH has no effect on the duration of conversion to negative and the length of patient hospitalization

As shown in **Table 2**, the number of days to convert virus to negative (time from admission to virus shedding) was 20.0 days (interquartile range (IQR) 11.0–31.0 days) in the LMWH group and 19.0 days (IQR 12.0–30.0 days) in the control group ($P = 0.46$); the difference between the two

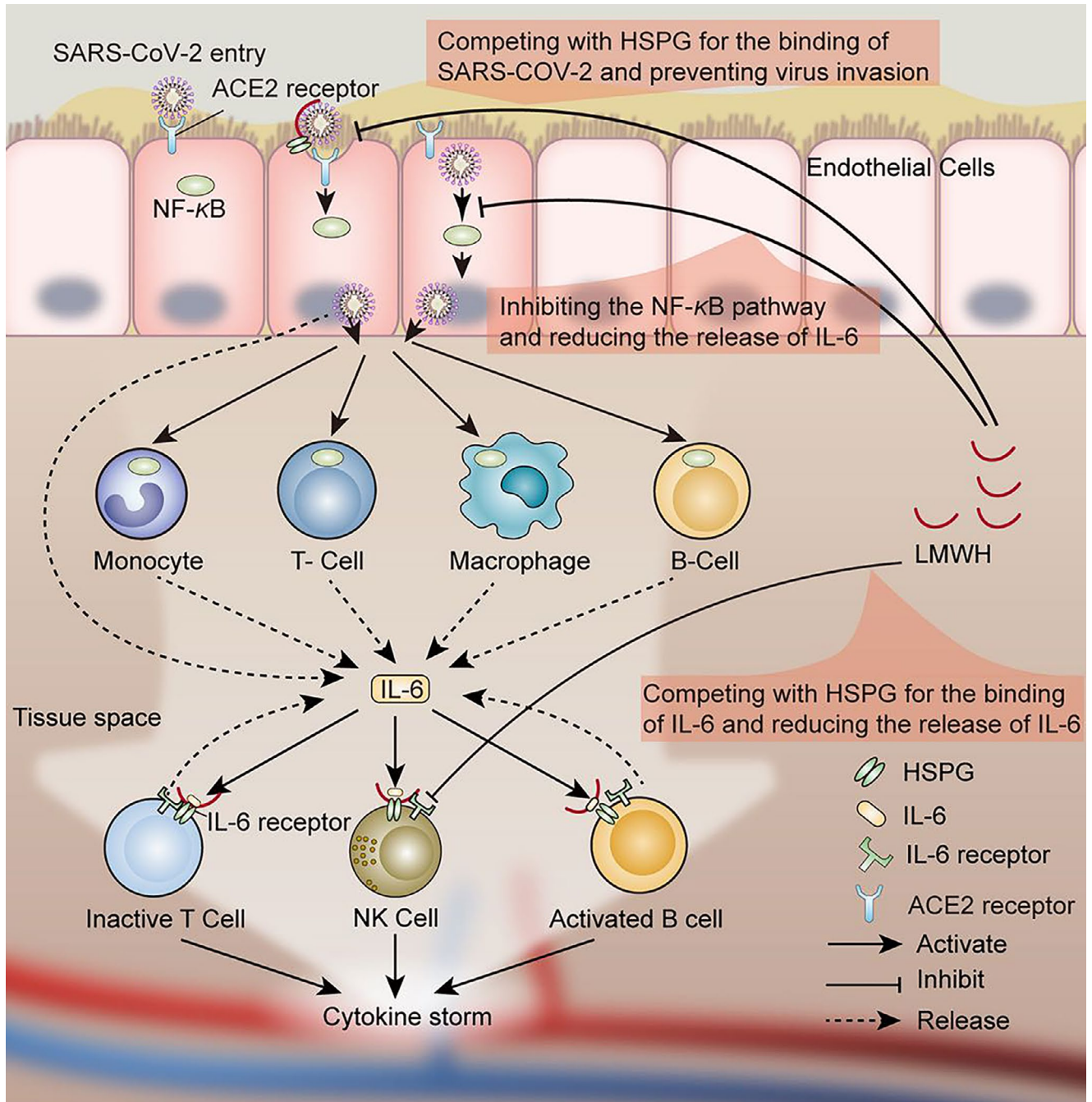


Figure 1 Possible mechanism of anti-inflammatory effects of low molecular weight heparin (LMWH) in patients with coronavirus disease 2019 (COVID-19). Under conventional antiviral treatment regimens, LMWH improves hypercoagulability, inhibits IL-6 release, and attenuates IL-6 biological activity. It has potential antiviral effects and helps delay or block inflammatory cytokine storms. LMWH can increase the lymphocyte% in the patients. The multiple effects of LMWH encourage its application for the treatment of patients with COVID-19. HSPG, heparin sulfate proteoglycan; SARS-CoV-2, severe acute respiratory syndrome-coronavirus 2.

groups was not significant. Similarly, the length of hospital stay was 29.0 days (IQR 17.0–42.0 days) in the LMWH group and 27.0 days (IQR 24.0–31.0) in the control group ($P = 0.41$); the difference between the two groups was not significant. Notably, all patients in the LMWH group and the control group showed overall improvement after treatment.

Effect of LMWH on cytokine levels in patients with COVID-19

As shown in **Figure 3a** there was no significant difference in IL-6 levels between the LMWH and control groups before treatment (47.47 ± 58.86 , 63.27 ± 96.27 , respectively, $P = 0.950$). In contrast, after LMWH treatment, the levels of IL-6 in the LMWH group were significantly lower compared

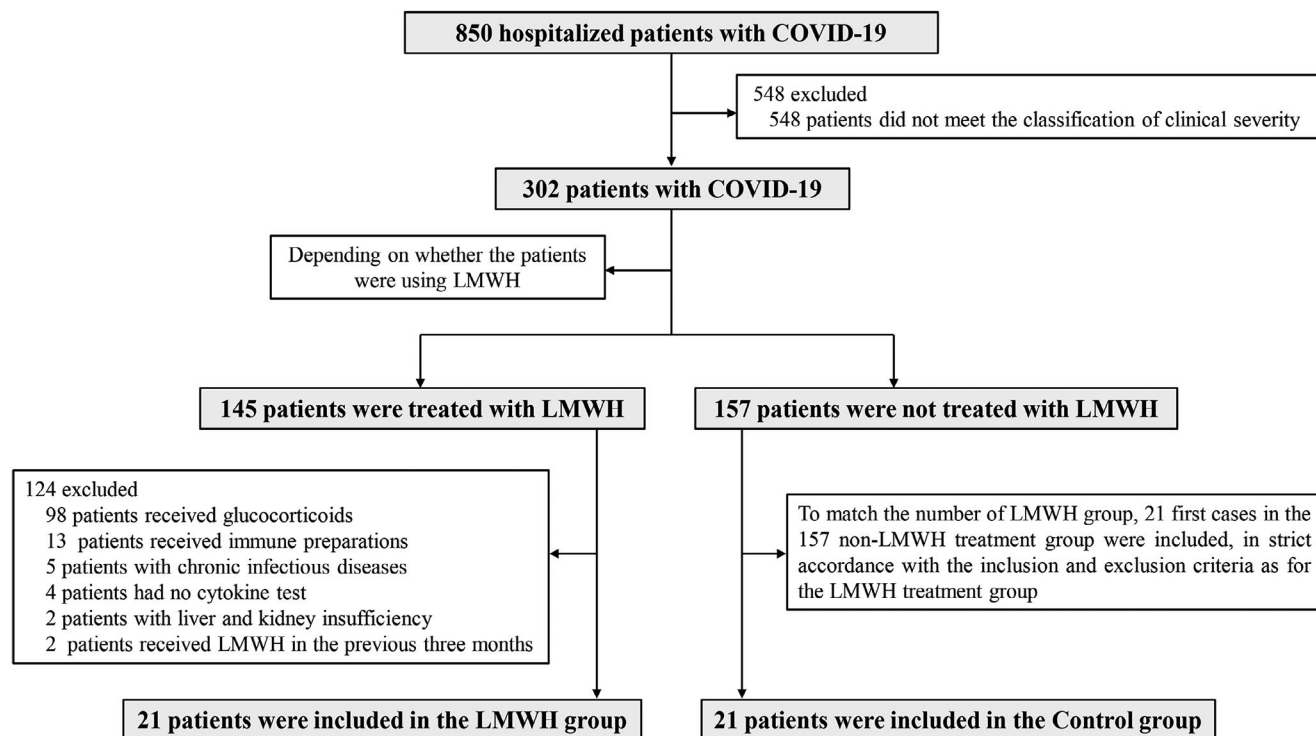


Figure 2 Flow chart for the inclusion and exclusion of patients with coronavirus disease 2019 (COVID-19). Based on strict inclusion and exclusion criteria, 42 patients with COVID-19 treated at the hospital between February 1, 2020, and March 15, 2020, were selected for the study, of which 21 underwent low molecular weight heparin (LMWH) treatment (LMWH group) and 21 did not (control group) during hospitalization.

with those in the control group (15.76 ± 25.71 , 78.24 ± 142.41 , respectively, $P = 0.00039$). Similarly, the changes in IL-6 levels in the LMWH group before and after LMWH treatment were significantly different from those in the control group (-31.71 ± 65.97 , 14.96 ± 151.09 , respectively, $P = 0.031$). However, there were no significant differences in the levels of IL-2, TNF- α , IL-4, IL-10, and IFN- γ between the LMWH treated and untreated groups (**Figure 3b-f**).

Effect of LMWH on blood routine characteristics

As shown in **Figure 4a-d**, there was no significant difference in red blood cells, white blood cells (WBCs), monocyte%, and neutrophil% levels between the two groups. **Figure 4e** reveals no significant difference in lymphocyte% (LYM%) between the LMWH and control groups before LMWH treatment (18.84 ± 8.24 , 22.42 ± 8.74 , respectively, $P = 0.144$). However, the changes in LYM% in patients of the LMWH group before and after LMWH treatment were significantly different from those in the control group (11.10 ± 9.50 , 3.08 ± 9.66 , respectively, $P = 0.011$).

Effect of LMWH on coagulation parameters

As shown in **Figure 4f-h**, there was no significant difference in thrombin time, activated partial thromboplastin time, and prothrombin time levels between the two groups before and after LMWH treatment. As shown in **Figure 4i,j**, the levels of D-dimer and fibrinogen degradation products (FDPs) in the LMWH group were significantly higher compared with those in the control group before treatment

(D-dimer: 3.75 ± 4.04 , 1.23 ± 1.15 , respectively, $P = 0.009$; FDP: 14.35 ± 14.6 , 4.05 ± 3.9 , respectively, $P = 0.002$). The changes in D-dimer and FDP levels in patients in the LMWH group before and after LMWH treatment were significantly different from those in the control group (D-dimer: -2.85 ± 3.90 , -0.05 ± 0.85 , respectively, $P = 0.002$; FDP: -9.05 ± 13.14 , -1.78 ± 3.15 , respectively, $P = 0.035$). However, there was no significant difference in fibrinogen (**Figure 4k**), antithrombin III (**Figure 4l**), and international normalized ratio (INR; **Figure 4m**) levels between the two groups.

Effect of LMWH on CRP levels

As shown in **Figure 4n**, LMWH treatment had no significant effect on CRP levels. There is no difference between the two groups of patients before LMWH treatment (31.15 ± 26.62 , 29.00 ± 23.79 , respectively, $P = 0.497$), nor after LMWH treatment (8.95 ± 10.44 , 8.76 ± 16.66 , respectively, $P = 0.620$). Consequently, there were no significant differences in the changes in CRP levels between the two groups of patients before and after LMWH treatment (-22.62 ± 23.79 , -20.23 ± 33.91 , respectively, $P = 0.660$).

DISCUSSION

Cytokine storms are associated with deterioration in several infectious diseases, including SARS and avian influenza,^{14,15} and IL-6 is one of the core cytokines that causes cytokine storms.⁷ In recent years, studies have revealed that heparin

Table 1 LMWH use in treating conditions of the 21 patients with COVID-19

LMWH group (n = 21)	Treatment with LMWH	Days of treatment
P1	Enoxaparin sodium injection 4000Axa IU q.d. i.h.	10
P2	Enoxaparin sodium injection 4000Axa IU q.d. i.h.	10
P3	Enoxaparin sodium injection 4000Axa IU q.d. i.h.	14
P4	Enoxaparin sodium injection 4000Axa IU q.d. i.h.	13
P5	Enoxaparin sodium injection 4000Axa IU q.d. i.h.	17
P6	Nadroparin calcium injection 4100Axa IU q.d. i.h.	9
P7	Enoxaparin sodium injection 4000Axa IU q.d. i.h.	2
P8	LMWH sodium injection 5000 IU once i.h.	1
P9	Enoxaparin sodium injection 4000Axa IU q.d. i.h.	16
P10	Enoxaparin sodium injection 4000Axa IU q.d. i.h.	19
P11	Enoxaparin sodium injection 4000Axa IU q.d. i.h.	14
P12	Enoxaparin sodium injection 2000Axa IU q.d. i.h.	19
P13	Enoxaparin sodium injection 2000Axa IU q.d. i.h.	22
P14	Nadroparin calcium injection 4100Axa IU q.d. i.h.	11
P15	Nadroparin calcium injection 4100Axa IU q.d. i.h.	13
P16	Enoxaparin sodium injection 4000Axa IU q.d. i.h.	8
P17	Nadroparin calcium injection 4100Axa IU q.d. i.h.	19
P18	LMWH sodium injection 5000 IU q.d. i.h.	8
P19	Enoxaparin sodium injection 4000Axa IU q.d. i.h.	8
P20	Enoxaparin sodium injection 4000Axa IU q.d. i.h.	7
P21	Enoxaparin sodium injection 4000Axa IU q.d. i.h.	10

Details of the dose, frequency, route of administration, and days of use of LMWH in the LMWH group. COVID-19, coronavirus disease 2019; LMWH, low molecular weight heparin.

has various non-anticoagulant properties, for example, LMWH can exert anti-inflammatory effects by reducing the release of IL-6.^{11-13,16}

IL-6 levels in severely ill patients with COVID-19 are significantly higher than in patients with mild disease.^{6,17} Transition from mild to severe conditions in patients with COVID-19 occur when cytokine levels reach and/or exceed a certain threshold, leading to a cytokine release syndrome.⁷ Hence, reducing IL-6 release is expected to attenuate the cytokine storm syndrome caused by the virus,⁸ thereby improving the condition of patients with COVID-19. LMWH was reported to reduce the release of IL-6 in the body by inhibiting the expression of NF- κ B.¹¹⁻¹³ Measuring the levels of proinflammatory cytokines in patients with COVID-19, we have found a marked decrease in the levels of IL-6 in the LMWH-treated patients compared with the patients without LMWH treatment ($P < 0.001$), consistent with the proposed protective effect of LMWH. Changes in other inflammatory factors were not statistically significant. In addition, IL-6 can bind to heparan sulfate (HS) on the cell surface, yielding a sufficiently high local concentration to activate signaling receptors,¹⁸ protect them against proteolysis, and promote paracrine action.^{16,19} An earlier study has reported that heparin binds IL-6, with affinity much higher than that of HS,¹⁶ thereby reducing its availability to its receptor complex. It, therefore, appears that LMWH reduces both the release of IL-6 and its biological activity.

In addition, there are other routes to explain a favorable effect of LMWH on patients with COVID-19. With the increasing interest in the use of heparin/LMWH for the treatment of COVID-19, our study signifies an additional activity of

heparin apart from anticoagulation.²⁰ Having demonstrated a marked reduction in IL-6 levels in the LMWH-treated patients, we further emphasize the well-documented anti-inflammatory/anti-sepsis effects of non-anticoagulant species of heparin.²¹

HS, a linear polyanionic polysaccharide, is a major constituent of all mammalian cells and tissues.²² It highly resembles heparin and LMWH in its structural properties and sugar composition.²² Importantly, HS has been known to serve as the first point of contact between target cells and a large number of human viruses (i.e., dengue virus, hepatitis C virus, HIV, human papillomavirus, and herpes viruses),^{23,24} including the SARS-CoV-2 virus.^{25,26} A very recent online paper has used surface plasmon resonance and circular dichroism and showed that the SARS-CoV-2 Spike S1 protein receptor binding domain interacts with heparin.²⁶ Heparin, LMWH, and heparin-like compounds have been shown to efficiently compete with HS and thereby attenuate viral attachment and infection,²⁷ providing a straightforward explanation for the antiviral effect of LMWH in clinical settings.

Importantly, as clarified below, recent studies expanded the established role of HS from a viral attachment molecule to an essential receptor required for entry. Heparin, LMWH, and non-anticoagulants species of heparin are known to inhibit the enzymatic activity of heparanase,²⁸ the sole HS-degrading endoglycosidase, shown recently to promote viral infection and spread.²⁹⁻³¹ It appears that heparanase behaves as a molecular switch in viral infection, which transforms the cell from a virus-permissive mode in which viral attachment and entry are favored, to a virus-detering mode, which allows for viral detachment and egress from

Table 2 General characteristics of all the included patients with COVID-19

Characteristics	LMWH group (n = 21)	Control (n = 21)	P value
Age, years	69.0 (42.0–91.0)	69.0 (40.0–84.0)	0.54
Sex	–	–	0.75
Female	8 (38%)	7 (33%)	–
Male	13 (62%)	14 (67%)	–
Comorbidity	13 (62%)	8 (38%)	0.12
Hypertension	8 (38%)	5 (24%)	0.32
Diabetes	6 (29%)	2 (10%)	0.12
Cardiovascular disease	5 (24%)	2 (10%)	0.21
Chronic obstructive lung disease	0	0	NA
Carcinoma	0	1 (5%)	0.31
Chronic kidney disease	0	0	NA
Other	4 (19%)	1 (5%)	0.15
Signs and symptoms			
Fever (temperature ≥ 37.3°C)	15 (71%)	13 (62%)	0.51
Cough	9 (43%)	7 (33%)	0.53
Sputum	6 (29%)	4 (19%)	0.47
Chest distress or asthma	11 (52%)	8 (38%)	0.35
Myalgia	2 (10%)	3 (14%)	0.63
Fatigue	8 (38%)	5 (24%)	0.32
Anorexia	6 (29%)	5 (24%)	0.73
Diarrhea	2 (10%)	1 (5%)	0.55
Nausea or vomiting	2 (10%)	1 (5%)	0.55
Respiratory rate ≥ 30 breaths per minute	0	0	NA
Pulse ≥ 125 beats per minute	0	0	NA
Systolic blood pressure < 90 mm Hg	0	0	NA
Antiviral therapy			
Arbidol	18 (86%)	20 (95%)	0.29
Recombinant human interferon α2B (aerosol inhalation)	6 (29%)	6 (29%)	1.00
Ribavirin	2 (10%)	0	0.15
Lopinavir/ritonavir	2 (10%)	0	0.15
Traditional Chinese medicine decoction	11 (52%)	9 (43%)	0.54
Disease progression			
Improved	21 (100%)	21 (100%)	NA
Invariable	0	0	NA
Deteriorative	0	0	NA
Time from hospitalization to virus shedding after the onset of the COVID-19, days	20.0 (11.0–31.0)	19.0 (12.0–30.0)	0.46
Hospital length of stay, days	29.0 (17.0–42.0)	27.0 (24.0–31.0)	0.41

Data are the median (IQR) or n (%). P values comparing the LMWH group and control group are from χ^2 test or unpaired two-sided Student's *t*-test. COVID-19, coronavirus disease 2019; LMWH, low molecular weight heparin; NA, not applicable.

There were no significant differences in age, sex, comorbidities, onset symptoms, time from hospitalization to virus shedding, length of hospital stay, antiviral treatment, and disease progression between the two groups.

cells.²⁹ Briefly, it was found that upregulation and activation of heparanase is a strategy common to a broad range of viral species (i.e., porcine reproductive and respiratory syndrome virus (PRRSV) and vaccinia virus) to increase egress, spread, and transmission.³¹ Interestingly, it appears that heparanase plays a role also in driving the undesirable cytokine storm discussed above. In individuals with SARS-CoV-2 infection, the level of inflammatory cytokines is markedly higher than normal and held responsible for the severity of the disease. Agelidis *et al.* documented that upon herpes simplex virus-1 infection, heparanase translocate to the nucleus of the infected cells and promotes inflammatory signaling,

mediated primarily via NF- κ B.³¹ In fact, transcription of IL-6 was significantly decreased after treatment with an inhibitor of heparanase enzymatic activity.³¹ LMWH, which inhibits heparanase activity,²⁸ may have a similar effect, possibly providing a mechanistic explanation for the decrease in IL-6 that we observed in the LMWH-treated patients. Collectively, the above considerations suggest that heparanase inhibitors (i.e., LMWH) may be an effective strategy in a therapeutic or combination therapy against viral infection, including COVID-19. Additional studies showed that inhibition of the glycolyx-degrading enzymes sialidase, cathepsin L and heparanase, using a combination therapy of zanamivir,

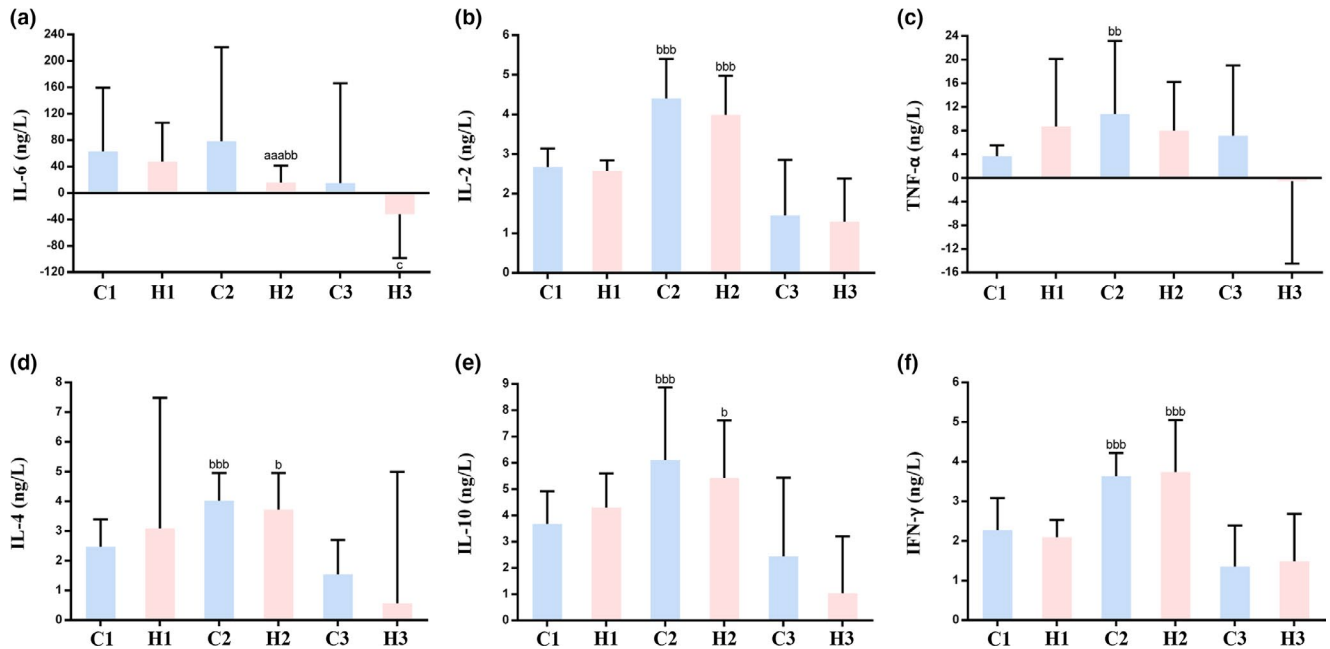


Figure 3 Effect of low molecular weight heparin (LMWH) on inflammatory cytokines in the included patients with coronavirus disease 2019 (COVID-19). (a–f) IL-6 (a), IL-2 (b), TNF- α (c), IL-4 (d), IL-10 (e), and IFN- γ (f) levels in the two groups of patients with COVID-19. Data are expressed as mean \pm SD ($n = 21$). C1 vs. H1 or C2 vs. H2, ^a $P < 0.05$, ^{aa} $P < 0.01$, ^{aaa} $P < 0.001$; C1 vs. C2 or H1 vs. H2, ^b $P < 0.05$, ^{bb} $P < 0.01$, ^{bbb} $P < 0.001$; C3 vs. H3, ^c $P < 0.05$, ^{cc} $P < 0.01$, ^{ccc} $P < 0.001$. (C1: control group, indices at admission; C2: control group, indices at discharge; C3: control group, changes in indices during hospitalization; H1: LMWH group, indices before LMWH treatment; H2: LMWH group, indices after LMWH treatment; H3: LMWH group, changes in indices before and after LMWH treatment).

cathepsin-L, and heparanase inhibitors, decreased vascular leakage after exposure to the influenza virus NS1 protein *in vitro* and *in vivo*.³² It will be interesting to see if an analogous therapeutic inhibition of glyocalyx breakdown can provide similar benefit clinically.

Several studies have recommended CRP and LYM% as indices for evaluating the effectiveness of clinical drugs or treatments.^{4,33,34} In the various analyses applied in this study, there was no statistically significant difference in CRP levels between the two groups, indicating that LMWH treatment has no effect on this parameter. Notably, the changes in LYM% in patients of the LMWH group before and after LMWH treatment were significantly different from those in the control group ($P < 0.05$), consistent with the results of Derhaschnig *et al.*³⁵ This result suggests that LMWH can increase LYM% in patients with COVID-19 and thereby improve their condition. Furthermore, it was reported that proinflammatory cytokines, such as TNF- α and IL-6, can induce lymphopenia.⁵ Hence, the decrease in IL-6 (Figure 3a) may contribute to the increase in LYM% observed in the LMWH-treated patients.

Tang *et al.* suggested a correlation among D-dimer, FDP, and COVID-19 severity.³⁶ However, there is currently no conclusive evidence supporting the use of D-dimer as an evaluation index.^{37–39} A broad sample analysis is required to determine whether D-dimer is associated with COVID-19 severity. Therefore, the present study does not consider this parameter as an evaluation index for disease progression. The average values of D-dimer and FDP before treatment was higher in the LMWH group than in the control group

(3.75, 1.23, respectively, $P < 0.01$; 14.35, 4.05, respectively, $P < 0.01$), therefore, LMWH was applied. Because this study is a retrospective analysis, we did not intervene in the type of treatment given to the patients, inferring that the purpose of medication in the LMWH group was to improve hypercoagulability. Because D-dimer and FDP are not considered as factors that designate patient's disease progression, their levels had no effect on subsequent analysis of the results. Notably, it appears (Table 2) that patients treated with LMWH had somewhat higher incidence of comorbidities and seemingly more frequent signs and symptoms of COVID-19. This may be due to the single-center retrospective study and the limited sample size, which may not fully reflect the overall characteristic status of patients in these aspects. Although there was no significant difference in baseline between the two groups, the lack of adjustment analysis was still a limitation of this study. These apparent effects are being evaluated in a prospective clinical study that evaluates the efficacy and safety of enoxaparin in the treatment of patients with COVID-19 (see below).

Importantly, LMWH exerts an anti-inflammatory effect by means of reducing IL-6 and increasing LYM%. We, therefore, favor the use of LMWH as a potential therapeutic drug for the treatment of COVID-19. We also suggest that non-anticoagulant species of LMWH that can be applied at high doses should be considered as a complement to conventional LMWH. To further support this conclusion, we are conducting a prospective clinical study to evaluate the efficacy and safety of one LMWH (enoxaparin) in the treatment of hospitalized adult patients with COVID-19 (Chinese

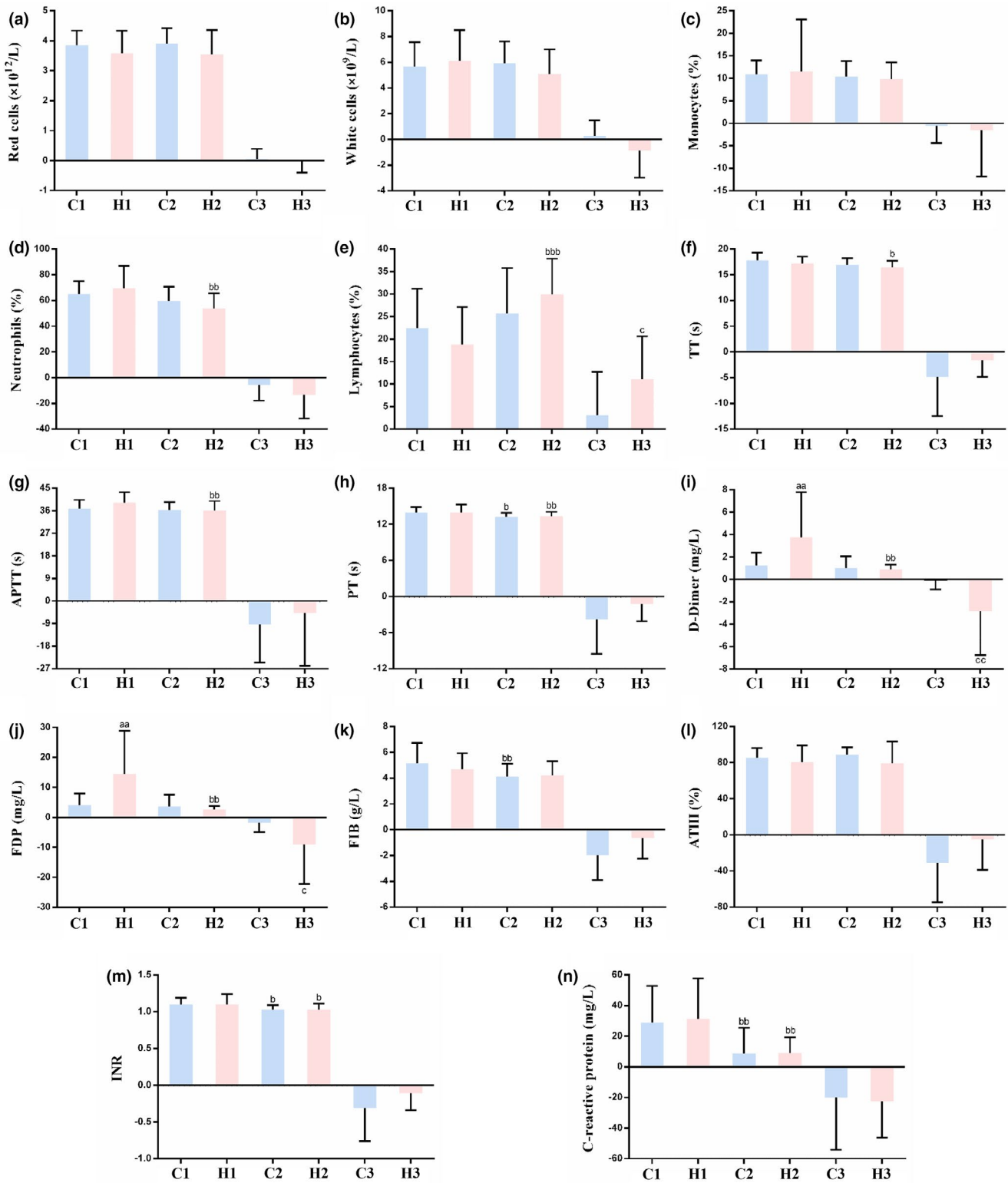


Figure 4 Effect of low molecular weight heparin (LMWH) on complete blood count, coagulation profile, and C-reactive protein (CRP) in the included patients with coronavirus disease 2019 (COVID-19). (a–n) Red blood cells (a), white blood cells (b), monocytes% (c), neutrophils% (d), lymphocytes% (e), thrombin time (TT, f), activated partial thromboplastin time (APTT, g), prothrombin time (PT, h), D-dimer (i), fibrinogen degradation products (FDP, j), fibrinogen (FIB, k), antithrombin III (AT III, l), international normalized ratio (INR, m) and CRP (n) levels in patients with COVID-19. Data are expressed as mean \pm SD ($n = 21$). C1 vs. H1 or C2 vs. H2, ^a $P < 0.05$, ^{aa} $P < 0.01$, ^{aaa} $P < 0.001$; C1 vs. C2 or H1 vs. H2, ^b $P < 0.05$, ^{bb} $P < 0.01$, ^{bbb} $P < 0.001$; C3 vs. H3, ^c $P < 0.05$, ^{cc} $P < 0.01$, ^{ccc} $P < 0.001$. (C1: control group, indices at admission; C2: control group, indices at discharge; C3: control group, changes in indices during hospitalization; H1: LMWH group, indices before LMWH treatment; H2: LMWH group, indices after LMWH treatment; H3: LMWH group, changes in indices before and after LMWH treatment.)

Clinical Trial Registry, number: chiCTR2000030700), with the objective of providing a more powerful reference for the treatment conditions.

This study still has several limitations. First, due to the retrospective design, we were unable to control the time intervals between examinations of the various indices in patients and the LMWH treatment schedule. Likewise, we could not estimate and manage the effective dose and timing of LMWH. Second, there were no critical cases in the two groups of patients; the treatment outcome of all cases was improvement and discharge, and there were no deaths. Finally, the findings are limited by the sample size and single-center design of our study.

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