

Probiotics use is associated with improved clinical outcomes among hospitalized patients with COVID-19

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Abstract

Background and aims: Currently, there are no definitive therapies for coronavirus disease 2019 (COVID-19). Gut microbial dysbiosis has been proved to be associated with COVID-19 severity and probiotics is an adjunctive therapy for COVID-19. However, the potential benefit of probiotics in COVID-19 has not been studied. We aimed to assess the relationship of probiotics use with clinical outcomes in patients with COVID-19.

Methods: We conducted a propensity-score matched retrospective cohort study of adult patients with COVID-19. Eligible patients received either probiotics plus standard care (probiotics group) or standard care alone (non-probiotics group). The primary outcome was the clinical improvement rate, which was compared among propensity-score matched groups and in the unmatched cohort. Secondary outcomes included the duration of viral shedding, fever, and hospital stay.

Results: Among the propensity-score matched groups, probiotics use was related to clinical improvement rates (log-rank $p=0.028$). This relationship was driven primarily by a shorter (days) time to clinical improvement [difference, -3 (-4 to -1), $p=0.022$], reduction in duration of fever [-1.0 (-2.0 to 0.0), $p=0.025$], viral shedding [-3 (-6 to -1), $p<0.001$], and hospital stay [-3 (-5 to -1), $p=0.009$]. Using the Cox model with time-varying exposure, use of probiotics remained independently related to better clinical improvement rate in the unmatched cohort.

Conclusion: Our study suggested that probiotics use was related to improved clinical outcomes in patients with COVID-19. Further studies are required to validate the effect of probiotics in combating the COVID-19 pandemic.

Keywords: clinical improvement, COVID-19, gut microbiota, probiotics

Received: 11 April 2021; revised manuscript accepted: 29 June 2021.

Introduction

The coronavirus disease 2019 (COVID-19) has been considered a global pandemic.¹ Fever, fatigue, and cough are the most common clinical features of COVID-19, but gastrointestinal symptoms (e.g., diarrhea, nausea, vomiting) are increasingly recognized among COVID-19 patients.^{2,3} Previous studies have shown that positive severe acute respiratory syndrome coronavirus (SARS-CoV-2) in the stool of COVID-19 patients was

associated with longer illness duration.⁴ Angiotensin-converting enzyme 2 has been found to be abundant in the epithelia of the lungs and intestine, suggesting that there is a subtle link between them.⁴ The gut microbiota have been proved to play a critical role in health through the “gut-lung axis”. Thus, SARS-CoV-2 might also affect the gut microbiota in COVID-19. Indeed, microbial dysbiosis was found in some patients with COVID-19.⁵ Imbalance of intestinal flora

Ther Adv Gastroenterol

2021, Vol. 14: 1–9

DOI: 10.1177/
17562848211035670

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can lead to dysbiosis, which might influence the outcome of the clinical features. A novel and more targeted method may be needed to modulate gut microbiota as one of the treatments for COVID-19 and its comorbidities.

Currently, no definitive drugs are recommended for the treatment of COVID-19. Developing effective new therapies for curing COVID-19 is time-consuming. Therefore, using existing approved therapies or treatment modalities is seen as a more cost-effective strategy for reducing the severity of COVID-19. In early February 2020, China's National Health Commission and National Administration (version 5) recommended the use of probiotics, gut microecological modulators, in COVID-19 patients to maintain the balance of intestinal microecology.⁶ Probiotics have shown good results in regulating innate immunity and improving inflammatory conditions.⁷

We hypothesized that probiotics modulate the gut microbiota to protect the respiratory system and would be related to clinical improvement in COVID-19. Thus, we conducted a propensity-score matched retrospective cohort study to assess the association of in-hospital use of probiotics with clinical outcomes.

Methods

Population

We performed a propensity-score matched retrospective cohort study at the Shenzhen Third People's Hospital (Guangdong, China) from 11 January 2020 to 1 April 2020. Adults (age 18 years or older) were eligible for the study if they tested positive for SARS-CoV-2 according to World Health Organization (WHO) interim guidance. Exclusion criteria were pregnancy or lactation; conditions with gastrointestinal or systemic diseases known to affect gut microbiota composition such as gastrointestinal cancer, inflammatory bowel disease; incomplete information. The definition of disease severity of COVID-19 was based on the Chinese guideline for the management of COVID-19.⁶ The study protocols were approved by the Ethics of Research Committee of the Shenzhen Third People's Hospital (No. 2020-193). A STARD checklist is shown in the Supplemental material online.

Standard care comprised, as necessary, supplemental oxygen, high-flow nasal cannula (HFNC), antiviral agents, antibiotics, corticosteroids, mechanical ventilation, according to the Chinese guideline for the management of COVID-19.⁸ Inclusion criteria for the adjuvant probiotics treatment were: mild to severe patients with COVID-19; there was no history of probiotics or antibiotics use for at least 1 month; did not have allergy to probiotics. Probiotic treatment was administered as 630 mg (three pills) twice daily, using live combined *Bifidobacterium*, *Lactobacillus*, and *Enterococcus* capsules (Bifico) (Shanghai, China). Each capsule is of 210 mg, which contains *Bifidobacterium*, *Lactobacillus* and *Enterococcus* (1.0×10^7 CFU for each ingredient). There was no connection with the probiotic treatment with the gastrointestinal symptoms of COVID-19 patients. The primary exposure was use of probiotics, classified as present if probiotics were received within 48 h of hospital admission and otherwise as absent. The duration of oral probiotics administration was defined as the time from probiotics treatment initiation to viral shedding or death. In the control group, the initial time for the patients without probiotics administration was defined as the time of viral shedding or death.

Clinical data were collected from the Shenzhen Third People's Hospital (Guangdong, China). Data reliability was verified by two researchers (TWL and LNZ) and difference in interpretation was adjudicated by a third researcher (JYW).

Outcome measures

The primary outcome was the time (days) to clinical improvement. The definition of time to clinical improvement was the time from any treatment initiation to live discharge or an improvement of two points on a seven-point scale, whichever came first according to the previous study.⁸ Second outcomes included the duration of hospital stay, fever, viral shedding, and gastrointestinal symptoms.

Laboratory measures

Throat-swab samples were collected for SARS-CoV-2 detection according to WHO interim guidance. Complete blood count, D-dimer, serum biochemical tests, interleukin-6 (IL-6), procalcitonin, CD4⁺ and CD8⁺ T cell count were

examined. The reference values were as follows: white blood cell (WBC) count: $(4-10) \times 10^9$; platelet count: $(100-300) \times 10^9$ /l; lymphocyte count: $(0.8-4.0) \times 10^9$ /l; IL-6: (0-7) pg/ml; albumin: 40-55 g/l; D-dimer: 0-0.5 μ g/ml; aspartate aminotransferase (AST): 0-45 g/l; alanine transaminase (ALT): 0-45 g/l; procalcitonin: <0.1 ng/ml; cluster of differentiation (CD) 4⁺ T cell count: >400/mm³; CD8⁺ T cell count: >150/mm³.

Statistical analysis

Patients in the cohort were classified according to whether probiotics were used or not (probiotics *versus* non-probiotics). Continuous and categorical variables were shown as medians [interquartile ranges (IQRs) or standard deviations (SDs)] and *n* (%), respectively. No imputation was performed for missing data, because the patients with insufficient information had been removed during the patient selection. Patients who had a probiotics approach were matched to those who did not receive probiotics therapy 1:1 by greedy matching on the logit of propensity scores with a caliper of 0.2 SD.⁹ Variables thought to be potential confounders were enrolled into the propensity score and included socio-demographics [age, sex, body mass index (BMI)], clinical characteristics (disease severity, hypertension, diabetes, coronary disease, radiological findings), laboratory tests (WBC counts, lymphocyte count, IL-6, D-dimer, ALT, AST, CD4⁺ and CD8⁺ T cell count), clinical treatments [antibiotics, corticosteroid, intravenous immunoglobulin, oxygen therapy, HFNC, non-invasive mechanical ventilation (NIMV), invasive mechanical ventilation (IMV)], and ICU admission. Standardized differences for all covariates were calculated before and after matching, with 10% or more considered indicative of imbalance.¹⁰ The primary outcome was described by Kaplan–Meier plot and compared with a log-rank test after matching. For all outcomes, time to different clinical events was treated as continuous variable, and the difference between the probiotics and non-probiotics groups was analyzed by Hodges–Lehmann estimation in the matched cohort. To further validate the effects of probiotics on clinical improvement, Cox proportional hazard model with hazard ratio (HR) and 95% confidence interval (CI) was used in the unmatched cohort after adjustment for confounding factors. Analysis were performed by R software (version 3.6.1, Vienna, Austria). All tests were two-sided with a significance level of 0.05.

Results

Baseline characteristics of patients before matching

A total of 411 COVID-19 patients were enrolled. Eighteen patients were excluded as they were less than 18 years old, 15 patients had both gastrointestinal disease and tumor, and 10 patients had incomplete information. The final cohort included 375 patients, among which 179 patients received standard care plus probiotics (the probiotics group) and the other 196 cases were treated with standard care alone (the non-probiotics group). The clinical characteristics are shown in Table 1.

Before probiotics treatment, the number of patients with diarrhea was 20 (10.2%) in the non-probiotics group and 13 (7.3%) in the probiotics group ($p=0.10$); the number of patients with abdominal pain was three (1.5%) in the non-probiotics group and three (1.7%) in the probiotics group ($p=0.12$); the number of patients with nausea and vomiting was eight (4.1%) in the non-probiotics group and four (2.2%) in the probiotics group ($p=0.12$). After probiotics treatment, gastrointestinal symptoms of these patients improved and symptoms disappeared. We did not observe any side effects of probiotic usage. Compared with patients who received probiotics therapy, those who had not received probiotics had lower rates of mild type (5.6% *versus* 8.4%; standardized difference, 0.11) and normal radiological findings (5.1% *versus* 8.4%; 0.13), as well as a lower lymphocyte count [median, 1.2×10^9 /l (1.0–1.7) *versus* 1.3×10^9 /l (1.0–1.7); 0.23]. With respect to clinical therapy, patients in the non-probiotics group had higher rates of antibiotics (38.8% *versus* 32.4%; 0.13), corticosteroids (34.7% *versus* 17.9%; 0.39), intravenous immunoglobulin (34.7% *versus* 20.7%; 0.32), oxygen therapy (75.0% *versus* 57.0%; 0.39), HFNC (4.1% *versus* 2.2%; 0.11), NIMV (19.9% *versus* 12.3%; 0.21), and ICU admission (9.2% *versus* 5.6%; 0.14) as compared with those in the probiotics group. There was no significant difference in antibiotics use between the two groups. The type, number, and combinations of administered drugs are shown in Supplemental Table 1.

Primary outcome after matching

A total of 150 patients (83.8%) who had undergone a probiotics approach were each matched to

Table 1. Demographic and clinical characteristics of the COVID-19 patients at baseline before and after matching.^a

Characteristic	Before matching			After matching		
	Non-probiotics group n = 196	Probiotics group n = 179	SDM ^b	Non-probiotics group n = 150	Probiotics group n = 150	SDM ^b
Male sex, n (%)	105 (53.6)	88 (49.2)	0.09	76 (50.7)	77 (51.3)	0.01
Age, median (IQR), years	50 (36–62)	48 (36–59)	0.01	50 (37–62)	49 (35–60)	0.00
BMI, median (IQR)	23.6 (21.7–26.1)	22.9 (20.8–25.1)	0.08	23.3 (21.4–25.6)	23.2 (21.3–25.3)	0.01
Disease severity, n (%)						
Mild	11 (5.6)	15 (8.4)	0.11	10 (6.7)	11 (7.4)	0.03
Moderate	174 (88.8)	157 (87.7)	0.03	135 (90.0)	134 (89.3)	0.02
Severe	11 (5.6)	7 (3.9)	0.08	5 (3.3)	5 (3.3)	0.0
Comorbidity, n (%)						
Hypertension	36 (18.4)	31 (17.3)	0.03	27 (18.0)	31 (20.7)	0.07
Diabetes	22 (11.2)	15 (8.4)	0.09	16 (10.7)	14 (9.3)	0.05
Coronary disease	10 (5.1)	6 (3.4)	0.08	7 (4.7)	6 (4.0)	0.03
Laboratory test, median (IQR)						
WBC count, ×10 ⁹ /l	4.6 (3.8–5.7)	4.8 (3.8–6.0)	0.09	4.7 (3.9–5.8)	4.7 (3.6–5.0)	0.00
Lymphocyte count, ×10 ⁹ /l	1.2 (1.0–1.7)	1.3 (1.0–1.7)	0.23	1.3 (1.0–1.7)	1.3 (1.0–1.7)	0.02
IL-6, pg/ml	10.2 (3.9–20.7)	6.3 (3.0–17.1)	0.01	8.8 (3.4–17.3)	8.5 (3.0–18.9)	0.00
D-dimer, µg/ml	0.4 (0.3–0.6)	0.3 (0.2–0.5)	0.04	0.4 (0.2–0.6)	0.4 (0.3–0.6)	0.07
ALT, g/l	21.0 (15.0–30.8)	18.0 (12.0–28.0)	0.00	20.0 (13.0–29.0)	19.0 (13.0–29.5)	0.00
AST, g/l	25.0 (19.3–31.0)	22.0 (19.0–28.0)	0.01	24.0 (19.0–30.0)	23.0 (19.0–29.3)	0.01
T cell count, /ml	967 (592–1261)	1076 (711–1389)	0.00	988 (628–1373)	1043 (678–1359)	0.00
CD4 ⁺ T cell count, /ml	519 (377–735)	586 (413–780)	0.00	545 (390–759)	567 (409–780)	0.00
CD8 ⁺ T cell count, /ml	343 (204–470)	386 (240–539)	0.00	348 (200–504)	377 (231–514)	0.00
Radiological findings, n (%)						
Normal	10 (5.1)	15 (8.4)	0.13	9 (6.0)	11 (7.3)	0.05
Ground-glass opacity	174 (88.8)	157 (87.7)	0.03	129 (86.0)	133 (88.7)	0.08
Bilateral pulmonary infiltration	147 (75.0)	131 (73.3)	0.04	108 (72.0)	114 (76.0)	0.09
Duration of illness onset to any treatments initiation, median (IQR), days	3 (2–6)	3.5 (2–7)	0.02	3 (2–7)	3 (1–5)	0.05
Treatments, n (%)						
Antibiotics	76 (38.8)	58 (32.4)	0.13	54 (36.0)	49 (32.7)	0.07
Antiviral treatment	196 (100.0)	179 (100.0)	0.00	150 (100.0)	150 (100.0)	0.00
Corticosteroids	68 (34.7)	32 (17.9)	0.39	32 (21.3)	32 (21.3)	0.00

(continued)

Table 1. (continued)

Characteristic	Before matching			After matching		
	Non-probiotics group <i>n</i> = 196	Probiotics group <i>n</i> = 179	SDM ^b	Non-probiotics group <i>n</i> = 150	Probiotics group <i>n</i> = 150	SDM ^b
Intravenous immunoglobulin	68 (34.7)	37 (20.7)	0.32	36 (24.0)	36 (24.0)	0.00
Oxygen therapy	147 (75.0)	102 (57.0)	0.39	101 (67.3)	102 (68.0)	0.02
HFNC	8 (4.1)	4 (2.2)	0.11	5 (3.3)	4 (2.7)	0.04
NIMV	39 (19.9)	22 (12.3)	0.21	21 (14.0)	22 (14.7)	0.02
IMV	9 (4.6)	7 (3.9)	0.03	6 (4.0)	7 (4.7)	0.03
ICU admission	18 (9.2)	10 (5.6)	0.14	9 (6.0)	10 (6.7)	0.03

^aGreedy matching occurred 1:1 on the logit of a propensity score with a caliper of 0.2 SD.
^bSDMs of 0.1 or more represent meaningful difference in covariates between groups before and after matching.
 ALT, alanine transaminase; AST, aspartate aminotransferase; BMI, body mass index; CD, cluster of differentiation; HFNC, high-flow nasal cannula; ICU, intensive care unit; IL-6, interleukin-6; IMV, invasive mechanical ventilation; IQR, interquartile range; NIMV, non-invasive mechanical ventilation; SDM, standard difference of the mean; WBC, white blood cell.

a patient who had not received probiotics therapy with standardized difference of less than 10% for all covariates, suggesting an adequate match (Table 1). Compared with the non-probiotics group, the cumulative improvement rate was higher in the probiotics group with a log-rank *p* value of 0.028 (Figure 1). Patients in the probiotics group also have an accelerated time to clinical improvement after matching [median, 18 days (IQR: 14–28) versus 21 days (IQR: 17–29)] with a median difference of –3 days (95% CI: –4 to –1; *p* = 0.022) [Figure 2(a) and Table 2].

Secondary outcomes after matching

The secondary outcomes of the probiotics and non-probiotics groups after matching are shown in Table 2. Among patients who experienced fever symptoms, the duration of fevers in the probiotics group was also shorter than that in the non-probiotics group (median, 6 days *versus* 7 days) with a significant difference in median –1 day (IQR: –2 to 0 days; *p* = 0.025) [Figure 1(b)]. The probiotics group was superior to the non-probiotics group in shortening the duration of hospital stay [median, 17 days *versus* 20 days; difference, –3 days (95% CI: –5 to –1), *p* = 0.009] [Figure 1(c)] and the duration of viral shedding [median, 15 days *versus* 18 days; difference, –3 days (95% CI: –6 to –1), *p* < 0.001] [Figure 1(d)]. However, there was no significance in the ICU stay length, duration of NIMV, duration of IMV, and duration of oxygen support between

the propensity-score matched groups (*p* > 0.05) (Table 2).

Effects of probiotics on clinical improvement in the unmatched cohort

The effects of probiotics on clinical improvement analyzed by Cox regression analyses in the unmatched cohort are shown in Table 3. A crude model of univariate analysis showed that the application of probiotics in COVID-19 patients was dramatically related to a better clinical improvement (HR, 1.35; *p* = 0.006). Using further multivariate analyses we found that patients who underwent probiotics therapy were likely to experience a favorable clinical improvement compared with those not treated with probiotics, even after adjusting for age, sex, BMI [Model 1; adjusted HR (aHR), 1.33; *p* = 0.007]; further adjusted for disease severity, comorbidity, radiological findings based on Model 1 (Model 2; aHR, 1.32; *p* = 0.012); based on Model 2, further adjusted for laboratory test results (Model 3; aHR, 1.28; *p* = 0.026). We further additionally adjusted for clinical treatments and ICU admission based on Model 3 (Model 4; aHR, 1.30; *p* = 0.017).

Discussion

In the present study, we found that probiotics added to standard care was superior to standard care alone in time to clinical improvement. The significant reduction in duration of viral shedding

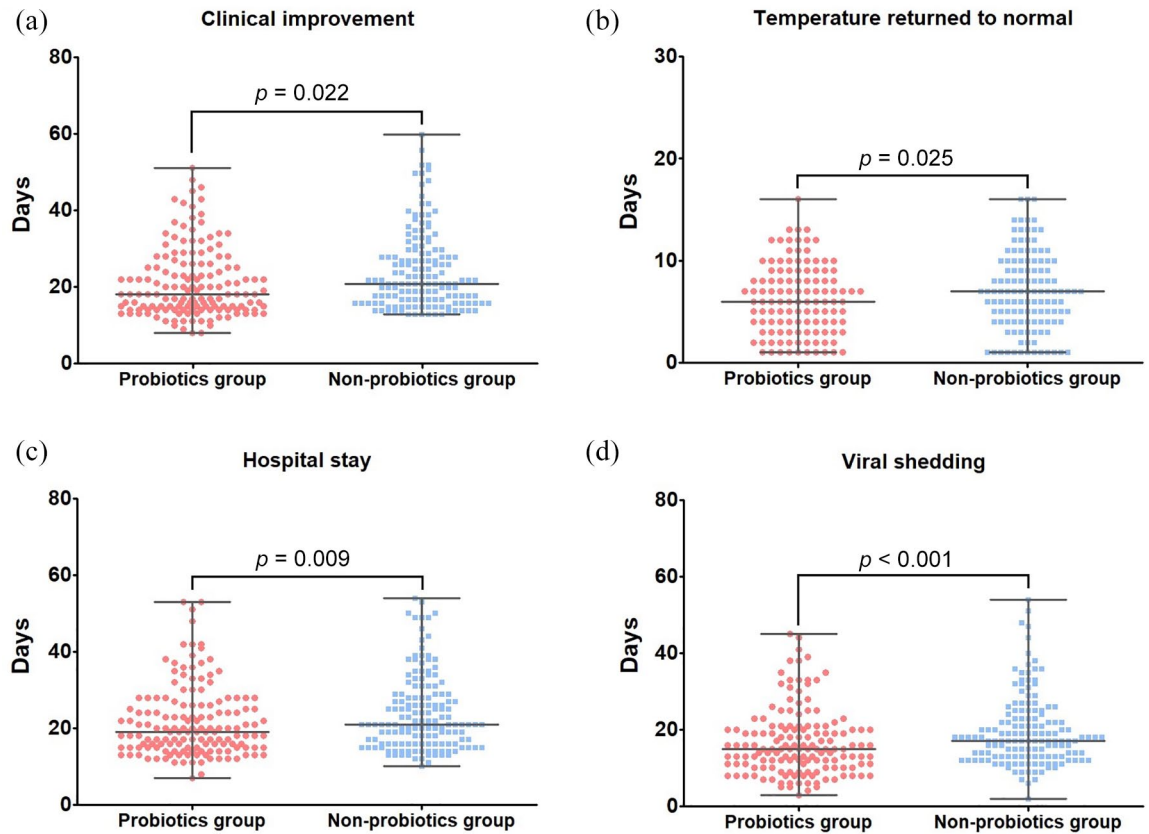


Figure 1. Secondary outcomes after matching. (a) Time to clinical improvement, (b) time to temperature return to normal, (c) duration of hospital stay, (d) duration of viral shedding.

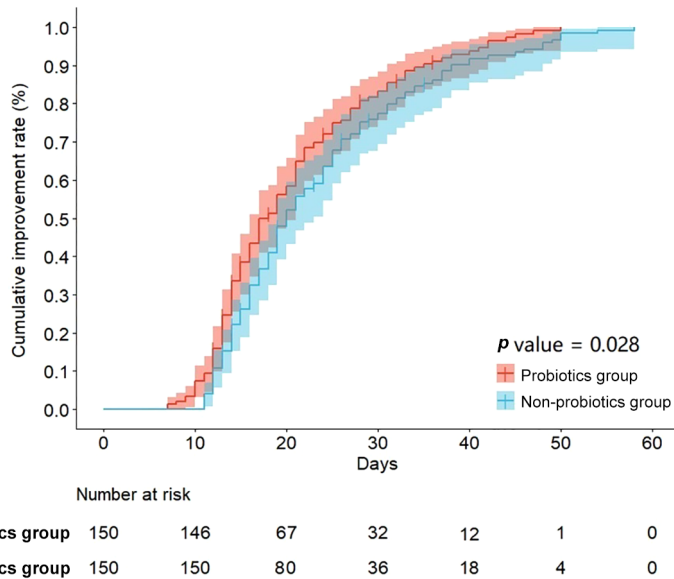


Figure 2. Kaplan-Meier estimates of cumulative clinical improvement rate.

was related to clinical improvement as shown by the significant reductions in duration of fever and hospital stay.

Gastrointestinal symptoms with a 70% relative increased risk for positive detection of COVID-19. The presence of diarrhea was associated with the severity of respiratory symptoms.² The gut microbiota are considered to have an important role on the innate and adaptive immune system.¹⁶ Gut-lung crosstalk may affect the gut microbiota in patients with COVID-19.⁴ It has been shown that modulating intestinal flora can avoid early replication of influenza virus in lung epithelia and reduce ventilator-associated pneumonia.^{17,18} Currently, no specific treatment has been recommended for COVID-19. Probiotics have been recommended for maintain the balance of gut microbiota in COVID-19 patients and prevent secondary bacterial infection.⁶ Probiotics, including bacteria and

Table 2. Clinical outcomes after the propensity-score matching.

Characteristic	Total N=300	Non-probiotics group n=150	Probiotics group n=150	Difference	p value
Primary outcome					
Time to clinical improvement, days	20.0 (15.0–28.0)	21.0 (17.0–29.0)	18.0 (14.0–28.0)	-3 (-4 to -1)	0.022
Secondary outcomes					
Hospital stay, days	21.0 (15.0–28.0)	22.0 (16.0–31.0)	19.0 (15.0–25.0)	-3 (-5 to -1)	00.009
Duration of viral shedding, days	16.0 (12.0–22.0)	18.0 (13.0–24.0)	15.0 (10.0–20.0)	-3 (-6 to -1)	<0.001
Duration of fever, days	7.0 (4.0–9.0)	7.0 (4.0–10.0)	6.0 (3.0–9.0)	-1.0 (-2.0 to 0.0)	00.025
ICU stay, days	15.0 (10.0–20.0)	16.0 (13.0–30.0)	13.0 (7.0–17.0)	-4 (-9 to 2)	0.397
Duration of NIMV, days	6.0 (4.75–8.0)	6.0 (4.0–8.0)	7.0 (5.0–9.5)	1 (-1 to 3)	0.545
Duration of IMV, days	19.0 (10.5–28.5)	24.0 (19.0–33.0)	12.0 (7.0–25.5)	-11 (-20 to 1)	0.615
Oxygen support, days	14.0 (8.0–21.0)	14.5 (7.25–20.0)	14.0 (8.0–21.0)	0 (-3 to 2)	0.081
Data are median (interquartile range). ICU, intensive care unit; IMV, invasive mechanical ventilation; NIMV, non-invasive mechanical ventilation.					

Table 3. Time-varying analysis for the effect of probiotics on clinical improvement in the unmatched cohort.

Models	Hazard ratio (95% CI)	p value
Crude model	1.35 (1.09–1.67)	0.006
Model 1 ^a	1.33 (1.09–1.67)	0.007
Model 2 ^b	1.32 (1.06–1.64)	0.012
Model 3 ^c	1.28 (1.04–1.59)	0.026
Model 4 ^d	1.30 (1.04–1.61)	0.017

^aModel 1 was adjusted for age, sex, and body mass index.

^bModel 2 was additionally adjusted for disease severity, comorbidity (including hypertension, diabetes, and coronary disease), and radiological findings (including normal, ground-glass opacity, and bilateral pulmonary infiltration).

^cModel 3 was additionally adjusted for laboratory test results (including white blood cell count, lymphocyte count, interleukin-6, D-dimer, alanine transaminase, aspartate aminotransferase, T cell count, CD4⁺ T cell count, and CD8⁺ T cell count).

^dModel 4 was additionally adjusted for clinical treatments (including antibiotics, corticosteroid, intravenous immunoglobulin, oxygen therapy, high-flow nasal cannula, non-invasive mechanical ventilation, and invasive mechanical ventilation), and intensive care unit admission. CI, confidence interval.

yeast, are living microorganisms that have been shown to be beneficial to human health.¹⁷ *In vivo* experiments and clinical trials expand our current understanding of the critical role of probiotics in human intestinal flora-related diseases. Many clinical trials have shown that probiotics can shape the intestinal flora, making it possible to control a variety of intestinal diseases and promote overall health.¹⁹ The presence of gastrointestinal symptoms was related to a 70% increased risk for positive detection of COVID-19.

COVID-19 infection affects lung tissue and intestines, thereby activating inflammation. Probiotics supplementation may improve the ability of the gastrointestinal microbiota to regulate immune activity, thereby preventing viral infections, including COVID-19. Although many experimental and clinical studies support the possible role of probiotic microorganisms in protecting the host against viral infections (e.g., colds and flu),^{20–23} no research has reported the use of probiotics to treat or prevent COVID-19. In our study, higher percentages of antibiotics, antiviral treatment corticosteroids, intravenous immunoglobulin, oxygen therapy, HFNC, NIMV, IMV, and ICU admission in the non-probiotics group before matching

imply that disease was much more severe in the non-probiotics group compared with the probiotic group. Similarly, higher percentages of antibiotics were found in the non-probiotics group after matching. We found that patients in the probiotics plus standard care group had shorter time to clinical improvement. The significant reduction in duration of viral shedding was related to clinical improvement as shown by the significant reductions in duration of fever and hospital stay. These results may lead to a better understanding of probiotics use in COVID-19 patients. According to our study, the indication of probiotics is mild to severe patients with COVID-19 and the contraindication is patients who are allergic to probiotics. The strains selection may contain *Bifidobacterium*, *Lactobacillus*, and *Enterococcus*, based on our current study. The dosage of probiotics is given according to the specifications. However, these findings are observational, and we cannot exclude the possibility of unmeasured confounders or hidden bias that account for the association between probiotics use and improved outcomes. Further randomized controlled trials (RCTs) are needed to prospectively explore the benefit of probiotics in combating the COVID-19 pandemic. Our study may also provide a theoretical basis for the next prospective randomized controlled studies. Currently, three registered trials to evaluate probiotics administration to COVID-19 patients are ongoing and these studies could help to address the scientific questions.²⁴⁻²⁶

Our study has several limitations. First, the cohort study design limits interpretation of results and confirmation of efficacy will require controlled trials; second, this study was limited to one single medical institution in Shenzhen and therefore may have limited external generalizability to other medical settings; third, critically ill patients are underrepresented, which did not allow the generalization of our findings to critical cases.

In summary, our study has demonstrated that the use of probiotics was associated with a shorter time to clinical improvement including fever, hospital stay, and viral shedding in hospitalized COVID-19 subjects. This information may help clinicians to consider the use of probiotics in patients with mild to severe COVID-19. Further RCTs are needed to prospectively explore the benefit of probiotics in combating the COVID-19 pandemic.

Author contributions

Contributors: LNZ, HQH, XL, and CZC served as co-first authors. TWL, JYW, and ZQW served as co-corresponding authors. TWL and ZQW designed the study, interpreted the results, the accuracy of the data analysis, and drafted the manuscript. LNZ, JYW, XL, and HQH contributed to data collection, data analysis, and data interpretation. XBX, CLW, SCY, QH, FW, FH, and GMS contributed to literature search and data collection. All authors approved the final version of the manuscript.

Conflict of interest statement

The authors declare that there is no conflict of interest.

Data availability statement

The data supporting the findings of this study are available from the corresponding author upon reasonable request.

Funding

The authors disclosed receipt of the following financial support for the research, authorship, and/or publication of this article: This work is supported by the Guangdong Basic and Applied Basic Research Foundation (No. 2020B1515020004, No. 2021A1515011056, No. 2018A0303130269), the National Natural Science Foundation of China (81873404); Guangdong Medical Science and Technology Research Fund (No. B2021206, No. A2018162); Project of Young Innovative Talents in Colleges and Universities in Guangdong Province (No. 2018KQNCX095); Shenzhen Key Medical Discipline Construction Fund (No. SZXK076) and Sanming Project of Medicine in Shenzhen (No. SZSM201612014).

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Supplemental material

Supplemental material for this article is available online.

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