Original Article

Early enteral nutrition improves the outcome of critically ill patients with COVID-19: A retrospective study

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INTRODUCTION

Coronavirus disease 2019 (COVID-19) is a respiratory tract infection caused by a newly emergent coronavirus, SARS-CoV-2, that was first recognized in Wuhan, China, in December 2019. As of September 9, 2020, the total number of patients has risen to 27,688,740 around the world and the death toll has reached 899,315.¹ Whereas most people with COVID-19 develop mild or uncomplicated illness, approximately 14% develop severe disease requiring hospitalization and oxygen support, and 5% require admission to an intensive care unit (ICU).²

Nutrition support is well known to play an important role in the treatment of patients with severe respiratory diseases, but data on the nutritional status and nutrition support of critically ill patients with COVID-19 are scarce. Many studies and guidelines have suggested that early (within 24 or 48 hours of admission) enteral nutrition (EN) support can significantly decrease mortality in critically ill patients.³⁴ However, some critically ill patients with COVID-19 require initiation of nutrition support via a parenteral route because of gastrointestinal symptoms or for other medical reasons. In this study, we investigated the nutritional status of critically ill patients with confirmed COVID-19 and retrospectively compared the clinical outcomes of different routes of nutrition support.

METHODS

Patients

This single-center, retrospective study was performed at the ICU of Wuhan Union Hospital Tumor Center (Wuhan, Hubei), a designated COVID-19 hospital. This ICU was taken over by a volunteer medical team dispatched from Second Affiliated Hospital of Zhejiang University School of Medicine (Hangzhou, Zhejiang). In accordance with the deployment of the National Health Commission, a total of 29 critically ill patients with COVID-19 were admitted to this ICU from February 15, 2020 to February 29, 2020. All patients with laboratory confirmation of SARS-CoV-2 infection met the criteria for being critically ill. According to the Chinese Clinical Guidance for COVID-19 (7th Edition), criteria of critically ill is as fol-

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lows: 1. respiratory failure requiring mechanical ventilation. 2. shock. 3. multiple organ failure needed ICU monitoring. Two Patients who died within 48 hours after admission were excluded. A total of 27 patients were enrolled in the study. The patient enrollment process is shown in Figure 1.

**Study group definitions**

Patients were divided into two groups according to the route of nutrition support initiated in the first week of staying in the ICU: an EN group and a parenteral nutrition (PN) group.

**Data collection**

Demographic data, electronic medical records, radiological images and laboratory findings for all enrolled cases were reviewed in detail. We collected data on age, sex, initial symptoms, vital signs, mechanical ventilation strategies and chronic medical histories (such as hypertension, diabetes mellitus, renal dysfunction, hepatic dysfunction, malignancy, stroke, chronic pulmonary disease and surgery history).

**Disease severity and comorbidities**

The Acute Physiology and Chronic Health Evaluation II (APACHE II) were evaluated on the first day of admission to assess the severity of disease. Pneumonia Severity Index (PSI), which is a widely used and approved scale for assessing the severity of respiratory infectious diseases and predicting prognosis was assessed based on clinical status at admission and laboratory findings. Age-adjusted Charlson Comorbidity Index (ACCI) was used to assess the severity of comorbidities in these patients.

**Nutrition data**

Baseline body mass index (BMI) were collected as the patient's initial nutritional status. We collected nutrition data including daily energy and protein intake on the seventh day of admission, at which time the nutrition data could reflect the patient's true situation of early nutritional intervention.

**Nutrition screening tools**

Nutritional Risk Screening 2002 (NRS-2002) and Nutrition Risk in the Critically Ill (NUTRIC) were used to assess nutritional status. These measures effectively reflect the nutritional status of patients, including both current potential undernutrition and disease severity.7,8 An NRS-2002 score ≥3 indicates potential nutrition risk, whereas a score ≥5 indicates high nutrition risk. Because IL-6 is not routinely measured in our ICU, a NUTRIC score ≥5 suggested high nutritional risk. Laboratory results, such as serum albumin, total protein and hemoglobin were also recorded at baseline and seven days later as markers of nutritional status.

**Nutrition support procedures**

Nutrition support was delivered within 24 hours after admission. The physician on duty decided which nutritional support route to initiate. Some physicians followed the usual protocol for severe pneumonia and preferred EN if there were no contraindications. Others preferred PN for the following considerations: 1. There would be cough and vomiting in the process of gastric intubation, resulting in aerosol dispersion which would lead to unnecessary occupational exposure. 2. COVID-19 was often accompanied by gastrointestinal symptoms, such as diarrhea, nausea, vomiting and anorexia. Premature EN might aggravate diarrhea and gastroesophageal reflux leading to fluid and electrolyte imbalance and secondary bacterial infection.

In the EN group, patients received first-line EN through nasogastric or nasojejunal tubes with a nutrition pump. Two simple formulas were used to estimate daily...

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**Figure 1.** Flow diagram of patient recruitment.
energy and protein requirements: [25–30 kcal/kg/day]*body weight and 1.2 g/kg/day* body weight. In the first week, trophic (10–15 kcal/kg/day) EN was initiated and then patients were gradually transitioned to full-energy EN. Whole-protein preparations with relatively high caloric content were selected (SSPC, China or Nutricia, Netherlands). Patients with intestinal damage received predigested short peptide preparations (SSPC, China), and hyperglycemic patients received nutritional preparations beneficial for glycemic control (SSPC, China). While feeding, patients were seated in a semi-reclining position of 30°–45°. Prokinetic agents and probiotics were used to prevent regurgitation and dysbacteriosis. After 1 week, energy intake and nutrition status were reassessed. If patients were not receiving adequate energy or protein, the nutrition strategies were optimized, such as by adding supplemental PN.

In the PN group, 1 patient who could feed herself received partial parenteral nutrition (PPN) from the peripheral vein. The remaining 12 patients received total parenteral nutrition (TPN) via a central venous catheter for at least 7 days after admission. In the first week, a low-calorie strategy (<20 kcal/kg/day) was used. A commonly used formula was: Glucose: 50% to 60% of non-protein calories, which was adjusted according to blood glucose. Structural lipid emulsion (Fresenius-Kabi Germany): 250 mL/day. 11.4% Compound amino acids (SPSS China): 250–500 mL/day. Multi-trace elements (Fresenius-Kabi Germany), Fat-soluble and water-soluble vitamins (SPSS China) were also given at the same time. After 1 week, the physician reassessed the nutritional status of the patients, and decided whether to continue TPN, add EN, or switched to EN according to the patient’s situation.

Medication and oxygen therapy
All patients were treated in accordance with the principles recommended in the Chinese Clinical Guideline for COVID-19 Pneumonia Diagnosis and Treatment. Patients were given antiviral treatment within 24 hours of admission (ribavirin 500 mg 2 times a day + arbidol 200 mg 3 times a day). Antibiotics (quinolones, third generation cephalothins, β-lactamase inhibitor compounds et al, according to the bacterial culture results) were used if the patient had a complicated bacterial infection or if the patient was at high risk of secondary infection. Adequate fluid resuscitation and vasoactive drugs were administered immediately if patients showed signs of circulatory failure. Traditional Chinese medicine (Lianhua Qingwen capsules) were also used according to the patient’s condition.

Relief of hypoxemia was the core of our treatment. Patients with the following conditions: SpO₂ <93%; PaO2/FiO2 <300 mmHg (1 mmHg = 0.133 kPa); respiratory rate >25 times per min at bed; or remarkable progression on X-ray imaging were immediately given high-flow nasal cannula (HFNC) oxygen therapy or non-invasive ventilation. If there was no relief within 2 hours, tracheal intubation and mechanical ventilation would be performed in time.

Outcomes
The primary outcome was 28-day mortality rate after ICU admission. Secondary outcomes included hemoglobin, serum total protein and serum albumin levels at baseline and seven days after nutrition support.

Data analysis
Demographic information, disease and nutrition therapy related information, laboratory findings and clinical outcomes were recorded and analyzed according to variable type. Continuous variables are reported as mean (standard deviation), and categorical variables are reported as frequency (percentage). Shapiro-Wilk test was used for normality test. Normal distribution data were analyzed using an unpaired Student’s t-test between two groups. Non-normal distribution data were analyzed using the Mann-Whitney U test. Categorical variables were compared with chi-squared test or Fisher’s exact test. Kaplan–Meier survival curves were used to plot the survival curves of patients. The Cox proportional hazard regression model was used to evaluate potential associations between nutrition group (EN or PN) and 28-day mortality. To assess the impact of confounding factors, we first performed univariate regression analysis to analyze variables that might be associated with mortality rate, such as age, sex, disease duration, APACHE-II score, mechanical ventilation, comorbidities, NUTIRC score, and NRS2002 score. After that, route of nutrition support, age, and APACHE-II score were included as variables in the Cox regression analysis according to clinical experience and statistical results. A p value <0.05 was considered statistically significant. SPSS 25.0 software (IBM Inc., 2010, USA) was used for all analyses.

Ethics
The Ethics Commission of the Second Affiliated Hospital of Zhejiang University approved data collection for this study (Yan 2020–220). Due to the retrospective nature of the study, informed consent was waived.

RESULTS
Baseline characteristics
A total of 27 patients were recruited for this retrospective study, 14 in the EN group and 13 in the PN group. The baseline characteristics of the participants are shown in Table 1 and were similar between groups. The average age was 74.9±10.5 years; 18 (66.7%) were men. 23(85.2%) had chronic underlying diseases, primarily hypertension (14/27), heart disease (10/27), diabetes mellitus (7/27) and chronic obstructive pulmonary disease (7/27), average ACCI score was 4.1±2.0. The median duration from onset of symptoms to ICU admission was 8 days (4–23). The average APACHE II score was 19.2±4.8, average PSI score was 132±15.5. During the first week after admission, a total of 18 (66.7%) patients were treated with mechanical invasive ventilation, 7 (25.9%) patients were treated with non-invasive ventilation, and 2 (7.4%) patients were treated with high flow nasal cannulas for oxygen therapy.

Energy and protein intake
Data of energy and protein intake of patients in the two groups are shown in Table 1. There was no significant difference in BMI between the groups, and there was no
significant difference in energy or protein requirements according to the simple formulas. In the first week after admission, the daily energy intake of patients in both groups increased gradually. The EN group increased to 1400±229 kcal, reaching 87.3±6.7% of the target (25 kcal/kg/day), and the PN group reached 1224±264 kcal, reaching 87.3±6.7% of the target (25 kcal/kg/day). There was no significant difference between the two groups (p=0.078). However, in terms of daily protein intake, that of EN group (55.6±9.5 g) was much higher than PN group (37.2±12.7 g), with significant difference (p<0.001).

Nutritional markers and mortality
Using two nutritional screening tools, NRS2002 and NUTRIC, we found that the proportion of patients with an NRS2002 score ≥3 was 100% (27/27), and the proportion of patients with an NRS2002 score ≥5 was 96.3% (26/27). The ratio of patients with an NUTRIC score ≥5 was 63.0% (17/27). The 28-day mortality of the EN group was 50% (7/14), whereas that of the PN group was 76.9% (10/13). The survival curves of the two groups were represented by a Kaplan-Meier curve (Figure 2), and the Breslow test indicated a significant difference between groups (p=0.030).

Multivariate analysis
In addition to nutrition support modalities, we selected two other factors closely associated with prognosis—age and APACHE II score—on the basis of previous literature. These three factors were used as covariates. Cox proportional hazard regression analysis was performed to identify independent factors associated with prognosis. All three factors were found to be independent risk factors for 28-day mortality (Table 3).

DISCUSSION
Data on nutritional status and nutritional support in critically ill patients with COVID-19 are currently scarce. On the basis of information on other critically ill patients and
Table 2. Nutritional markers at baseline and 7 days later

<table>
<thead>
<tr>
<th></th>
<th>EN group (n=14)</th>
<th>PN group (n=13)</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>albumin (g/L)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Baseline</td>
<td>29.4±3.0</td>
<td>29.2±3.2</td>
<td>0.835</td>
</tr>
<tr>
<td>Day 7</td>
<td>31.8±4.8</td>
<td>28.1±5.1</td>
<td>0.030</td>
</tr>
<tr>
<td>Total protein (g/L)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Baseline</td>
<td>57.2±2.2</td>
<td>59.3±6.8</td>
<td>0.267</td>
</tr>
<tr>
<td>Day 7</td>
<td>63±6.5</td>
<td>60.0±9.7</td>
<td>0.450</td>
</tr>
<tr>
<td>hemoglobin (g/L)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Baseline</td>
<td>107±20.6</td>
<td>117±23.9</td>
<td>0.261</td>
</tr>
<tr>
<td>Day 7</td>
<td>106±22.1</td>
<td>110±21.7</td>
<td>0.671</td>
</tr>
</tbody>
</table>

EN: enteral nutrition; PN: parenteral nutrition

Table 3. Cox proportional hazards regression of 28-day mortality

<table>
<thead>
<tr>
<th>Independent variable</th>
<th>β</th>
<th>SE</th>
<th>Wald</th>
<th>HR (95% CI)</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nutrition support</td>
<td>1.71</td>
<td>0.549</td>
<td>9.64</td>
<td>0.182 (0.062-0.553)</td>
<td>0.002</td>
</tr>
<tr>
<td>Age</td>
<td>-0.086</td>
<td>0.041</td>
<td>4.39</td>
<td>0.918 (0.847-0.995)</td>
<td>0.036</td>
</tr>
<tr>
<td>APACHE II</td>
<td>0.248</td>
<td>0.080</td>
<td>9.60</td>
<td>1.28 (1.10-1.50)</td>
<td>0.002</td>
</tr>
</tbody>
</table>

SE: standard error; HR: hazard ratio; APACHE II: Acute Physiology and Chronic Health Evaluation II

other respiratory diseases, COVID-19 is expected to pose significant nutritional risks. A meta-analysis has reported that the prevalence of undernutrition risk in pulmonology department inpatients is 36.95%.10 An international, multicenter, prospective study has reported that the proportion of nutritional risk (NUTRIC ≥5) in critically ill patients admitted to the ICU is 57%.11 Our study found the incidence of nutritional risk in critically ill patients with COVID-19 to be 85.2%, a proportion much higher than that in previous studies, possibly due to 1. The average age of patients in this cohort was very high. 2. These patients had a very high incidence of comorbidities (80%). 3. According to the APACHE-II and PSI scores, these patients were in a very critical condition. These factors together may have significantly increased their nutritional risk.

The mortality rate in our cohort was higher than that previously reported,2 possibly because the patients in our cohort were significantly older (age 74.9 vs 59.7), were in more severe condition (APACHE II score 19 vs 17) and had higher ratios of mechanical ventilation (92.6% vs 71%). It is now generally believed that older patients with COVID-19 have significantly higher rates of critical illness and mortality. However, in this cohort, since most of the patients were elderly and the sample size was small, the mortality did not increase with the increase of age. Instead, several very elderly patients (maximum 97 years old) survived.

The importance of nutrition support for the treatment of critically ill patients is well known. Malnutrition may cause infections, muscle wasting, delayed recovery and increased mortality.12 However, which route of nutrition support should be initiated in critically ill patients remains controversial. A meta-analysis has indicated that the use of EN as opposed to PN may result in an important decrease in the incidence of infectious complications in critically ill patients and may be less costly.13 Casaer et al have shown that as a supplement to EN, early PN, compared with late PN, not only does not decrease mortality but also increases the risk of ICU infection, mechanical ventilation time and time for renal replacement therapy.14 However, a randomized, controlled, multicenter open-label, parallel-group study (NUTRIREA-2) has revealed that, compared with PN, isocaloric EN does not reduce mortality or the risk of secondary infections, but is associated with a greater risk of digestive complications.15 In addition, the CALORIES study has shown that patients with early PN and EN have a similar 30-day mortality.16 Another study has indicated that EN is associated with a higher overall mortality and a higher incidence of inadequate nutritional intake, complications associated with the delivery system and other feed-related morbidities than PN.17

In our study, initiation of EN appeared to be advantageous. According to the literature, we believe these findings may have occurred for the following reasons: 1. PN might impede blood glucose control. Some studies have suggested that hyperglycemia in critically ill patients may lead to an increased risk of infection and poorer outcomes.18,19 In the early stages of the outbreak, monitoring of blood glucose was not as frequent as usual, to avoid overexposure among medical staff. Therefore, intensive monitoring and tight control of blood glucose could not be achieved at that time. 2. Recently, some studies have identified the gastrointestinal tract as a potential pathway for SARS-CoV-2 infection.20,21 Some patients have gastrointestinal symptoms such as diarrhea, anorexia and nausea. In contrast, animal model studies and clinical studies have demonstrated that total PN may exacerbate inflammation of the intestinal epithelium and decrease intestinal barrier function, thus leading to bacterial translocation.22,23 The combination of SARS-CoV-2 infection and PN could lead to an increased risk of infection in patients. 3. Many of these critically ill patients had underlying diseases, particularly diseases affecting the circulatory system, such as cardiac dysfunction, which must be managed strictly through fluid transfusion. PN may aggravate these underlying diseases and cause internal environmental perturbation, thus affecting patient outcomes.

EN also is associated with certain risks.24 Therefore,
appropriate feeding procedures are crucial. There is little literature on COVID-19 nutrition support. We therefore drew on our experience with other severe respiratory diseases in clinical practice.

First, regarding the time of EN support initiation, it is necessary to initiate EN within 24 hours of admission for hemodynamically stable patients. The results of multiple meta-analyses have shown that early EN, compared with delayed EN, significantly decreases mortality and infection rates. Starting PN within 48 hours is also an appropriate option for patients with hemodynamic instability or the presence of EN contraindications.

Second, many studies have shown that severe underfeeding or overfeeding may increase mortality and prolong hospital stay and mechanical ventilation time, and should therefore be avoided as much as possible. However, some recent RCT studies have suggested that full-energy or trophic feeding has no significant effect on patient outcomes. We usually initiated trophic (10–15 kcal/h or no more than 500 kcal/day) nutrition support in the first week, because some studies have indicated that starting with trophic nutrition support may decrease the incidence of gastrointestinal intolerance. After a week, we reassessed the nutritional status and energy requirements and then gradually increased the nutrition support or added the other route of nutrition support. Monitoring and tight control of blood glucose should be considered necessary throughout the entire process of nutrition support.

Third, in the process of EN, regular evaluation of gastrointestinal function is required. The acute gastrointestinal injury grading system is usually used to assess the severity. Patient abdominal tension, bowel sounds and defecation, as well as vomiting, aspiration and other conditions, should be observed daily. If feeding intolerance is present, EN must be reduced or suspended.

Fourth, prone position ventilation is an important treatment for critically ill patients with COVID-19. A study has indicated that prone ventilation may result in more gastric retention and higher rates of vomiting and EN termination. However, in our experience, prone ventilation did not significantly increase the adverse effects of EN, but allowed for maintaining an appropriate body position and initiating feeding from a small volume. Some studies have obtained similar findings, in which patients in prone positions tolerated EN well.

Conclusions
The incidence of nutritional risk in critically ill patients with COVID-19 is very high and should be taken seriously. In addition, nutrition support should be provided promptly. Early EN (within 24 hours) may be beneficial to patient outcomes. Optimal feeding procedures may prevent the side effects of EN.

Limitations
COVID-19 is an emerging, rapidly evolving pandemic. Our healthcare system was under unprecedented pressure, therefore we were unable to conduct prospective research on nutrition support. Due to the retrospective nature of the study, randomization was not used to allocate patients to EN or PN groups, resulting in an inevitable bias. In this retrospective study, the sample size was small, the follow-up time was short and many clinical indicators could not be obtained. Larger prospective randomized controlled studies are needed to address this issue.

AUTHOR DISCLOSURES
The authors declare that they have no conflict of interest.

REFERENCES
13. Gramlich L, Kichian K, Pinilla J, Rodych NJ, Dhaliwal R, Heyland DK. Does enteral nutrition compared to parenteral nutrition result in better outcomes in critically ill adult