





High-flow nasal cannula therapy for hypoxemic respiratory failure in patients with COVID-19

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Abstract

Introduction: High-flow nasal cannula (HFNC) therapy in patients with hypoxemic respiratory failure due to COVID-19 is poorly understood and remains controversial.

Methods: We evaluated a large cohort of patients with COVID-19-related hypoxemic respiratory failure at the temporary COVID-19 hospital in Mexico City. The primary outcome was the success rate of HFNC to prevent the progression to invasive mechanical ventilation (IMV). We also evaluated the risk factors associated with HFNC success or failure.

Results: HFNC use effectively prevented IMV in 71.4% of patients [270 of 378 patients; 95% confidence interval (CI) 66.6–75.8%]. Factors that were significantly different at admission included age, the presence of hypertension, and the Charlson comorbidity index. Predictors of therapy failure (adjusted hazard ratio, 95% CI) included the comorbidity-age-lymphocyte count-lactate dehydrogenase (CALL) score at admission (1.27, 1.09–1.47; $p < 0.01$), Rox index at 1 hour (0.82, 0.7–0.96; $p = 0.02$), and no prior steroid treatment (0.34, 95% CI 0.19–0.62; $p < 0.0001$). Patients with HFNC success rarely required admission to the intensive care unit and had shorter lengths of hospital stay [19/270 (7.0%) and 15.0 (interquartile range, 11–20) days, respectively] than those who required IMV [104/108 (96.3%) and 26.5 (20–36) days, respectively].

Conclusion: Treating patients with HFNC at admission led to improvement in respiratory parameters in many patients with COVID-19.

Keywords: COVID-19; high-flow nasal cannula; hypoxia; invasive mechanical ventilation; Mexico; SARS-CoV-2 pneumonia

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Introduction

In the management of acute respiratory failure in patients with SARS-CoV-2 pneumonia, experts have agreed that following conventional oxygen therapy, the use of high-flow nasal cannula (HFNC) constitutes the next step in the therapeutic pyramid, before endotracheal intubation.¹ HFNC has emerged as an alternative, non-invasive respiratory support option that can reduce the

mortality rate as well as prevent or delay the need for intubation in patients with severe acute respiratory distress caused by SARS-CoV-2 pneumonia.² However, despite evidence of the beneficial effect of HFNC use in patients with acute respiratory distress,³ at present, there is limited evidence on the efficacy of HFNC in patients with severe SARS-CoV-2 pneumonia.⁴ While some concerns have been raised regarding the risk of dispersion of

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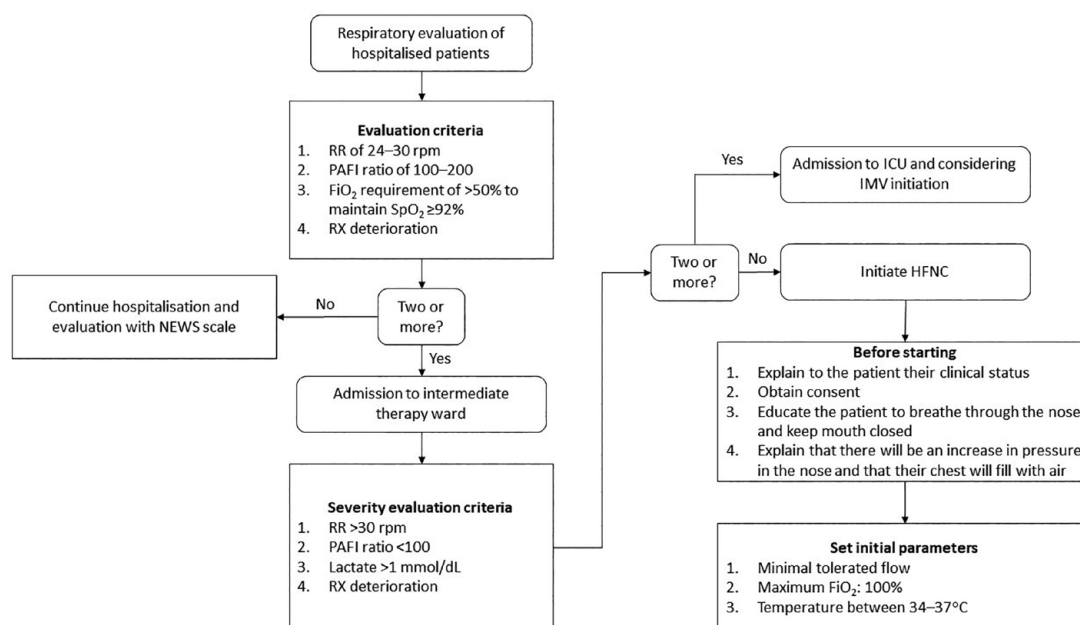


Figure 1. Clinical protocol for the management of hypoxemic respiratory failure in the temporary COVID-19 Hospital.

FiO₂, fractional inspired oxygen; HFNC, high-flow nasal cannula; ICU, intensive care unit; NEWS, National Early Warning Score; PAFI, ratio of PaO₂ over FiO₂; RR, respiratory rate; RX, radiography; SpO₂, oxygen saturation.

SARS-CoV-2 through bio-aerosols,⁵ the evidence to date challenges the presumption of aerosol generating procedures.^{6,7} Some concerns have also been raised regarding the risk of delayed intubation resulting in worse outcomes.⁸ Based on the known usefulness of HFNC in patients with acute respiratory failure,³ it is crucial to determine the efficacy of this non-invasive respiratory support method to avoid the need for invasive mechanical ventilation (IMV) in patients with SARS-CoV-2.

The present study, which was performed at the temporary COVID-19 hospital located in the Citibanamex Exhibition Center in Mexico City, aimed to evaluate the efficacy of HFNC in patients with hypoxemic respiratory failure due to severe SARS-CoV-2 pneumonia to reduce the risk of requiring IMV. We also aimed to identify the risk factors of disease progression among patients with severe SARS-CoV-2 pneumonia and treated with HFNC.

Methods

Study design

This prospective observational study was conducted between 16 May 2020 and 9 November

2020 in accordance with the Declaration of Helsinki. The study protocol was approved by an independent Ethical Review Board at the National Autonomous University of Mexico (FM/DI/099/2020). All patients provided written informed consent prior to participation.

Patients

Patients aged ≥18 years who were admitted to the temporary COVID-19 hospital with a confirmed diagnosis of COVID-19 [as verified by a positive polymerase chain reaction (PCR) test] and hypoxemic respiratory failure (PaO₂ ≤60 mmHg) due to severe SARS-CoV-2 pneumonia were included. For HFNC use, a respiratory rate of >24–30 breaths per minute, a PAFI ratio (PaO₂/FiO₂) of 100–200,⁹ and requiring a FiO₂ >50% to achieve an oxygen saturation (SpO₂) of ≥92% was necessary.

Procedures

After admission, patients were initially evaluated for treatment with HFNC (Figure 1). In addition, patients were also assessed for HFNC according to the Rox index (see Supplemental Figure 1 online). The Rox index is a measure of

hypoxemia severity that predicts the need for IMV.^{10,11} In this study, cut-off values of <2.85, <3.47, <3.85, and <4.88 after 2 hours, 6 hours, 12 hours, and 16 hours were considered failure of HFNC, as previously described.¹⁰ A Rox index of ≥ 4.88 after 16 hours was considered HFNC success.

HFNC (Precision Flow Plus, Vapotherm, New Hampshire, USA) was started with an oxygen flow rate of 40 l/min and FiO₂ of 100%, humidified and heated to 34–37°C. Patients were instructed to keep their mouths closed to avoid loss of gas flow. To conform with local standards of infection control, the use of N95 masks, ocular protection, surgical gowns, and disposable gloves were mandatory for all healthcare personnel.

After the initial respiratory evaluation, patients who were admitted to the intermediate therapy ward and fulfilled two or more of the following criteria were admitted to the intensive care unit (ICU): (1) respiratory rate >30 rpm, (2) PAFI ratio <100, (3) lactate >1 mmol/dl, and (4) radiographic deterioration (Figure 1). Patients who were admitted to the ICU were then considered for intubation according to ICU admission criteria. Among patients who initiated HFNC, admission to the ICU was based on the Rox index evaluation as follows: patients whose Rox index was <3.47 at 6 hours after initiating HFNC after a first HFNC failure, patients whose Rox index was <3.85 at 12 hours after initiating HFNC, and patients whose Rox index was <4.88 at 16 hours after initiating HFNC (see Supplemental Figure 1 online).

Variables

We evaluated demographic and clinical characteristics, duration of symptoms prior to admission, duration of HFNC or IMV, length of hospital stay, Rox index measurements, and respiratory and blood gas parameters at the start of HFNC and 16 hours after HFNC. The success of HFNC was defined as the non-progression from HFNC to IMV, with a CALL score cutoff of >6 points used to stratify the risk of progression in patients with COVID-19.¹²

Statistical methods

A convenience sampling approach was used to select patients for inclusion in the study.

Descriptive statistics were used for baseline demographic and clinical characteristics, with *n* (%) for categorical variables and median [interquartile range (IQR)] for continuous variables. The Wilcoxon rank-sum test was used to compare continuous data between patients who did or did not require IMV. A Chi-square or Fisher's exact test was used to compare categorical data between the groups. Statistical significance was set at a *p*-value <0.05.

Survival analysis (Cox regression) was fitted to calculate adjusted hazard ratios (HRs) and 95% confidence intervals (CIs) for IMV among patients treated with HFNC. A multivariate survival analysis was performed, adjusting for the following confounders: age, sex, CALL score, diabetes, hypertension, obesity, Rox index at 1 hour, treatment with steroids, absolute lymphocytes, D-dimer, PAFI, lactate, and PaCO₂. Only patients with complete clinical data were included in the survival analysis. Data were analyzed from the start of HFNC until failure (the need of IMV) or censure (discharge date). STATA v.15 (Stata Corp., College Station, TX, USA) and R version 3.6.3 were used for statistical analyses.

Results

Patients

The characteristics of patients included in this study are summarized in Table 1. A total of 378 patients were included in this study and data were collected between 16 May 2020 and 9 November 2020. The median (IQR) age was 54.5 (46–64) years, and 66.7% (*n*=252) were male. The proportions of patients with type 2 diabetes, hypertension, and obesity were 35.5% (*n*=134), 36.8% (*n*=139), and 47.6% (*n*=180), respectively.

Outcomes

A flow chart of the outcomes of 378 patients with hypoxemic respiratory failure who were treated with HFNC is shown in Figure 2. The HFNC success rate, defined as patients who did not require IMV, was 71.4% (*n*=270; 95% CI 66.6–75.8) compared with 28.6% (*n*=108; 95% CI 24.2–33.4) of patients who required IMV. Of the 270 patients who were successfully treated with HFNC, 262 patients (97.0%) were discharged, seven patients (2.6%) were referred, and one patient (0.4%) died. Among those 108 patients

Table 1. Patient characteristics.

	Total N=378	HFNC only n=270	HFNC + IMV n=108	p-value
Age, years	54.5 (46–64)	53 (45–61)	60 (51–70)	<0.0001
Sex				0.63
Female	126 (33.3)	92 (34.1)	34 (31.5)	–
Male	252 (66.7)	178 (65.9)	74 (68.5)	–
Diabetes	134 (35.5)	92 (34.1)	42 (38.9)	0.38
Uncontrolled diabetes ^a	71 (53.0)	44 (47.8)	27 (64.3)	0.08
Glucose, n=348	128 (106–172)	124 (105.5–166.5)	135 (107.5–187)	0.26
Hypertension	139 (36.8)	90 (33.3)	49 (45.4)	0.03
Uncontrolled hypertension ^b	30 (7.9)	17 (6.3)	13 (12)	0.30
BMI (kg/m ²)				0.80
Normal (18.5–24.9)	43 (11.4)	31 (11.5)	12 (11.1)	–
Overweight (25.0–29.9)	140 (37.0)	101 (37.4)	39 (36.1)	–
Obesity (≥30)	180 (47.6)	124 (45.9)	56 (51.9)	–
Unknown	15 (4.0)	14 (5.2)	1 (0.9)	–
Charlson comorbidity index				<0.0001
No comorbidities	162 (42.9)	130 (48.2)	32 (29.6)	–
Low-risk category (1–2)	178 (47.1)	121 (44.8)	57 (52.8)	–
High-risk category (≥3)	38 (10.1)	19 (7.0)	19 (17.6)	–
Duration of symptoms prior to admission, days	8 (5–11)	9 (6–12)	6 (4–8.5)	<0.0001
≤5 days	109 (28.8)	59 (21.8)	50 (46.3)	<0.0001
>5 days	268 (70.9)	210 (77.8)	58 (53.7)	<0.0001
Unknown	1 (0.3)	1 (0.4)	0 (0.0)	<0.0001
Time from admission to HFNC, days	1.99 (2.4)	1.8 (2.3)	2.5 (2.6)	0.02
Duration of HFNC, (days)	11 (4–16)	13 (10–18)	2 (1–3)	<0.0001
Steroid treatment at hospitalization	270 (71.4)	211 (78.2)	59 (54.6)	<0.0001
ICU admission	123 (32.5)	19 (7.0)	104 (96.3)	<0.0001
ICU duration, days	10 (5–19)	3 (2–3)	13 (7–20)	<0.0001
Hospital duration, days	18 (12–25)	15 (11–20)	26.5 (20–36)	<0.0001

Data are presented as n (%) or median (IQR)

^aUncontrolled diabetes defined as glucose >180 mg/dL.

^bUncontrolled hypertension defined as a blood pressure of >140/100 mmHg.

BMI, body mass index; HFNC, high-flow nasal cannula; ICU, intensive care unit; IMV, invasive mechanical ventilation; IQR, interquartile range.

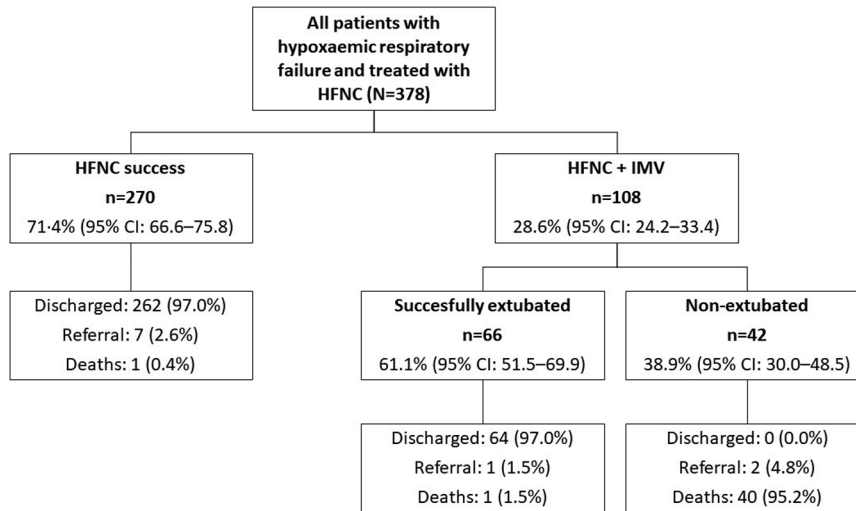


Figure 2. Flow chart of outcomes of patients treated with HFNC. HFNC success was defined as the non-progression from HFNC to IMV. HFNC, high-flow nasal cannula; IMV, invasive mechanical ventilation.

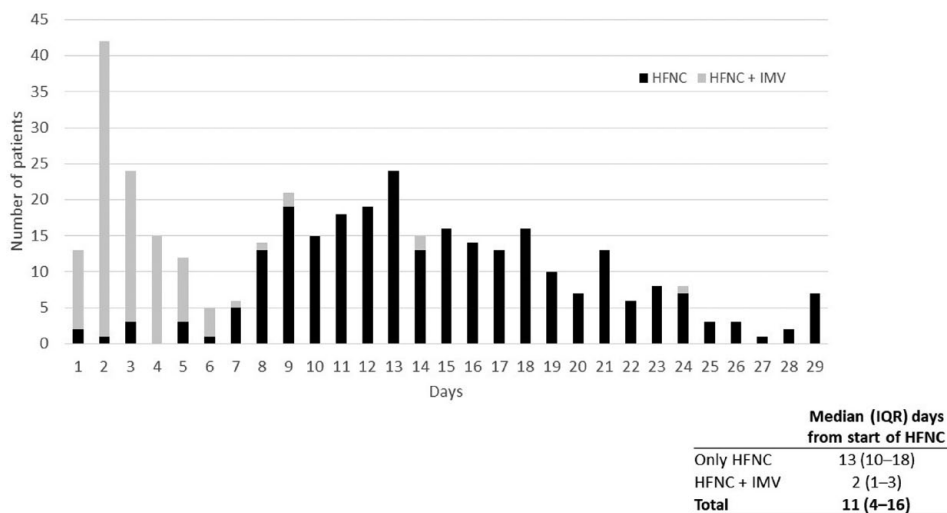


Figure 3. Proportion of patients with HFNC success and failure over time.

**p*-value < 0.0001.

HFNC, high-flow nasal cannula; IMV, invasive mechanical ventilation; IQR, interquartile range.

who required IMV, 61.1% ($n = 66$; 95% CI 51.5–69.9) were successfully extubated. Of those 66 patients who required IMV and who were successfully extubated, 64 patients (97.0%) were discharged, and one patient (1.5%) died. In addition, the proportion of patients with HFNC success increased over time from when HFNC treatment was started (Figure 3). The median (IQR) number of days of HFNC administration was 13 (10–18) days in patients who had success

compared with 2 (1–3) days in patients who then also required IMV.

When comparing patients who required IMV and those who did not require IMV, age and the presence of hypertension were significantly different ($p < 0.0001$ and $p = 0.03$, respectively). The Charlson comorbidity index categories also differed significantly between these two groups ($p < 0.0001$) (Table 1). Patients who only

Table 2. Laboratory results at admission and when HFNC was started.

	Total N = 378	Only HFNC n = 270	HFNC + IMV n = 108	p-value
Creatinine (mg/dl), n = 353				
At admission	0.9 [0.7–1.0]	0.8 [0.7–1.0]	0.9 [0.8–1.2]	0.002
At HFNC start	0.8 [0.7–1.0]	0.8 [0.7–1.0]	0.8 [0.6–1.2]	0.85
FiO ₂ (mmHg), n = 348				
At admission	35 [24–56]	35 [24–53]	32 [24–60]	0.82
At HFNC start	45 [32–75]	40 [30–60]	63 [34–94]	<0.0001
Lymphocyte (%) n = 362				
At admission	11.7 [6.9–18.6]	12.4 [7.7–18.6]	10.6 [6.3–17.4]	0.06
At HFNC start	11.5 [7.4–19.1]	12.5 [8.0–20.7]	8.8 [5.5–16.1]	<0.0001
D-dimer (ng/ml), n = 357				
At admission	550 [380–890]	555 [390–940]	550 [360–790]	0.32
At HFNC start	580 [370–970]	540 [350–860]	735 [490–1145]	<0.0001
Ferritin (µg/l), n = 279				
At admission	466.6 [227.3–836.8]	466.6 [229.2–770.6]	467.8 [219.8–961.7]	0.69
At HFNC start	457.4 [249.9–780.7]	438.4 [230.5–714.1]	556.4 [305.2–971.7]	0.03
Data are presented as median (interquartile range). FiO ₂ , fraction of inspired oxygen; HFNC, high-flow nasal cannula; IMV, invasive mechanical ventilation.				

received HFNC were treated for a significantly longer period of time *versus* those who eventually required IMV (13 *versus* 2 days, respectively; $p < 0.0001$). Patients with HFNC success rarely required admission to the ICU and had shorter lengths of hospital stay [19/270 (7.0%) and 15.0 days, respectively] than those who required IMV [104/108 (96.3%) and 26.5 days, respectively].

Laboratory results at admission and when HFNC was commenced are summarized in Table 2 and Supplemental Table 2 online. Patients who required IMV had less favorable laboratory parameters *versus* those who did not require IMV. It should be noted that there were significant differences at admission and when commencing HFNC in absolute lymphocyte counts ($p < 0.01$ and $p < 0.0001$, respectively). Although markers of inflammation (median D-dimer and ferritin levels) were not significantly different between

patients who only required HFNC and those who also required IMV at admission ($p = 0.32$ and $p = 0.69$, respectively), they were significantly different when commencing HFNC ($p < 0.01$ and $p = 0.03$, respectively).

Rox index values were significantly higher at each time point in patients who did not require IMV *versus* those who required IMV (Figure 4). In patients with HFNC success, Rox index values increased from 5.98 at baseline to 6.41, 6.83, 7.02, 7.37, 7.87, and 8.20 after 1, 2, 4, 6, 12, and 16 hours, respectively. In contrast, in patients with HFNC failure, Rox index values remained low from 5.40 at baseline to 5.70, 5.88, 5.76, 5.93, 5.74, and 5.62 after 1, 2, 4, 6, 12, and 16 hours, respectively. When comparing the Rox index values between 2 hours and 16 hours, the difference was significant in the HFNC only group, but not in the HFNC and IMV group ($p < 0.0001$ and $p = 0.91$, respectively).

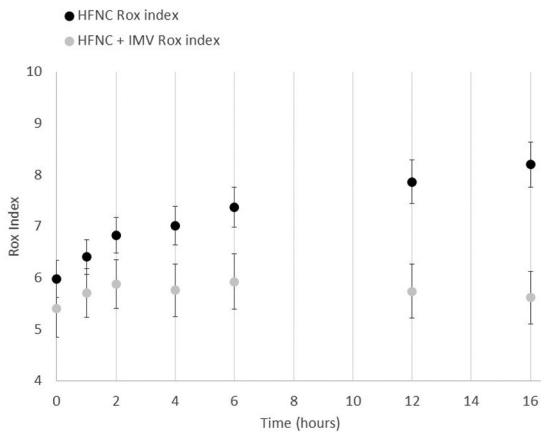


Figure 4. Rox index in patients with HFNC success and failure.

Rox index values at different time points in patients who did not require IMV [success of HFNC; black dots] *versus* those who required IMV (failure of HFNC; grey dots) among patients with hypoxemic respiratory failure who underwent treatment with HFNC. HFNC, high-flow nasal cannula; IMV, invasive mechanical ventilation.

The changes in arterial blood gas parameters from 2 hours after HFNC to 16 hours after HFNC are summarized in Supplemental Table 3 online. In patients with HFNC success, the median (IQR) ratio of SpO₂ over FiO₂ (SPFI) increased from 135.7 (115.3, 160.0) after 2 hours to 158.4 (127.2, 192.2) after 16 hours of HFNC. In patients with HFNC failure, the median (IQR) SPFI ratio decreased from 115.0 (98.0, 140.0) after 2 hours to 110.4 (96.5, 134.9) after 16 hours of HFNC. When comparing SPFI ratios between 2 hours and 16 hours, the difference was only significant in the HFNC only group and not in the HFNC and IMV group ($p < 0.0001$ and $p = 0.34$, respectively).

Finally, we show that the CALL score at admission (adjusted HR 1.27; 95% CI 1.09–1.47; $p < 0.01$), Rox index at 1 hour (adjusted HR 0.82; 95% CI 0.70–0.96; $p = 0.02$), and absence of treatment with steroids (adjusted HR 0.34; 95% CI 0.19–0.62; $p < 0.0001$) were all significant predictors of HFNC failure (Table 3).

Discussion

The present study is one of the largest observational prospective studies to evaluate the efficacy of HFNC use in patients with severe SARS-CoV-2 pneumonia. Our results showed that the HFNC success rate of 71.4% significantly

prevented escalation to IMV in patients with hypoxemic respiratory failure due to COVID-19. This finding is important, because if HFNC was not available at our temporary COVID-19 hospital, the majority of these patients would have required IMV, which is supported in the literature across several studies.^{13–17}

CALL score at admission predicted a linear increase in the risk of IMV, with an HR of 1.27 for every point increase. The Rox index at 1 hour after starting HFNC predicted an 18% decrease in the risk of IMV for every point increase; in contrast, steroid treatment predicted a 66% decrease in the risk of IMV compared with the absence of steroid treatment.

In a multicenter prospective observational study of 293 consecutive patients with severe COVID-19-related hypoxemic respiratory failure in South Africa, the HFNC success rate (defined as the proportion of patients successfully weaned from HFNC) was 47% (137/293 patients), which is substantially lower than the 71.4% we report in the present study.¹³ It was also reported that the median duration of HFNC was significantly higher in those with HFNC success *versus* those with HFNC failure ($p < 0.001$). In a study of 28 consecutive patients with hypoxemic acute respiratory failure due to SARS-CoV-2 infection in Italy, the HFNC success rate (defined as a reversal of hypoxemia or SpO₂ $\geq 92\%$), was 67.8% (19/28 patients).⁴ However, it was reported that only 17.8% of patients subsequently required IMV (*versus* 28.6% in the present study), although the small number of patients ($n = 28$) in that study is likely to be a confounding factor. In a larger multicenter, retrospective cohort study conducted in Wuhan, HFNC failure (defined as upgrading respiratory support to positive pressure ventilation or death) was reported in 46.5% of patients. Among these patients, 13 (30.2%) subsequently required IMV, with failure of HFNC associated with a higher mortality rate.¹⁸ Finally, in a small study of eight patients with severe and critical COVID-19 in China, it was reported that after 2 hours of HFNC use, the Rox index was ≥ 4.88 in 100% of patients and this remained above this cut-off for 12 hours.¹⁹ It was concluded that even in severe and critical patients who were experiencing hypoxemic respiratory failure, HFNC was successful. In a previous retrospective analysis of patients with moderate-to-severe hypoxemia due to highly suspected or proven

Table 3. Predictors of HFNC failure ($n=238$) (adjusted Cox regression model).

Variable	Adjusted HR (95% CI)	p-value
Age (per year increase)	1.003 (0.99–1.01)	0.53
Male <i>versus</i> female	0.73 (0.41–1.32)	0.28
CALL score (per 1-point increase)	1.27 (1.09–1.47)	<0.01
No diabetes (<i>versus</i> uncontrolled)	0.81 (0.43–1.51)	0.50
Diabetes (<i>versus</i> uncontrolled)	0.46 (0.16–1.29)	0.14
Hypertension (<i>versus</i> no hypertension)	1.08 (0.47–2.50)	0.85
Overweight (<i>versus</i> normal weight)	1.05 (0.38–2.93)	0.92
Obesity (<i>versus</i> normal weight)	1.49 (0.55–4.01)	0.43
Rox index 1 hour (per 1-point increase)	0.82 (0.70–0.96)	0.02
Treatment with steroids (<i>versus</i> no treatment)	0.34 (0.19–0.62)	<0.0001
Absolute lymphocytes (per unit increase)	0.99 (0.99–1.00)	0.24
D-dimer 550–1000 (<i>versus</i> normal levels)	1.23 (0.66–2.27)	0.51
D-dimer > 1000–1500 (<i>versus</i> normal levels)	1.52 (0.61–3.76)	0.37
D-dimer > 1500 (<i>versus</i> normal levels)	0.31 (0.09–1.09)	0.07
PAFI (per unit increase)	0.99 (0.99–1.00)	0.94
Lactate–mmol/l (per unit increase)	1.25 (0.80–1.97)	0.33
PaCO ₂ (per unit increase)	1.04 (0.98–1.11)	0.18

CALL, comorbidity–age–lymphocyte–and LDH; CI, confidence interval; HFNC, high-flow nasal cannula; HR, hazard ratio; PAFI, ratio of PaO₂ over FiO₂

COVID-19 infection who were treated with HFNC, 67 of 104 patients (64.4%) did not require IMV,²⁰ which is similar to the HFNC success rate (71.4%) in the present study. Overall, the present study findings are consistent with those reported in these previous studies in terms of HFNC success.^{4,13,18–20} However, direct comparisons cannot be made because of differences in the study design and definitions of HFNC success between the studies.

When investigating factors associated with HFNC failure, the present study found that the CALL score at admission was a significant predictor of HFNC failure. In contrast, the Rox index at 1 hour (per 1-point increase) and prior treatment with steroids were significant predictors of HFNC success. Our findings are consistent with those of a previous study, which reported that a Rox index

of 6 after HFNC commencement and the use of steroids were associated with HFNC success.¹³ The results of the RECOVERY trial on corticosteroid use in patients hospitalized with COVID-19 showed that early corticosteroid use (within ≤ 7 days of admission) reduced mortality and ICU admissions in these patients; however, no significant difference was shown in IMV rates between early and later corticosteroid use.²¹ Similar to the present study, the severity of hypoxemia and lower oxygen saturation at admission were reported to be factors associated with HFNC failure in previous studies.^{4,18} Other factors, including C reactive protein level⁴ and male sex,¹⁸ have also been previously associated with HFNC failure.

There are many potential advantages of HFNC use, including efficacy, less training needed for

health-care personnel, and lower cost compared with IMV. Furthermore, in hospitals with saturated critical care capacities, the early use of HFNC is likely to have a positive effect. In contrast, some of the potential barriers of HFNC use in low resource settings include the fact that the training of health-care personnel is needed, HFNC requires the cooperation of the patient for adequate use, and, as with IMV, there are limitations in the chain of supply.

The present study has some limitations, including those inherent to the observational, single-center study design and the retrospective analysis of patients according to the management/treatment algorithms, which may have introduced bias. In addition, we did not collect data on radiological implications.

Conclusions

In conclusion, the present study showed that 71.4% of patients with SARS-CoV-2 pneumonia and hypoxemic respiratory failure did not require IMV when treated with HFNC; this reinforces the benefits of the timely use of HFNC. HFNC led to improvement in respiratory parameters in many patients with COVID-19 and might reduce the length of stay in the hospital and the ICU. CALL score, the Rox index at 1 hour after starting HFNC, and the presence/absence of steroid treatment were identified as predictors of HFNC outcome. These results should be validated in prospectively registered, randomized controlled trials.

Authors' contributions

APC, ESL, MGN, RRVV, HHB, MAA, LERG, RVB, RVAW, BSO, MLRC, and LMC conceived and designed the study. HGR, JLG, LMJ, LAMJ, and RTC conceived and designed the study; drafted the manuscript; acquired, analyzed, and interpreted the data; and revised the manuscript for important intellectual content. All authors read and approved of the final version to be published and agree to be accountable for all aspects of the work.

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Conflict of interest statement

The Carlos Slim Foundation funded this study. APC, ESL, MGN, RRVV, HHB, LMJ, MAA, LERG, LMC, RVB, RVAW, BSO, and MLRC are full-time employees of the Temporary COVID-19 Hospital. HGR, JLG, LAMJ, and RTC are full-time employees of the Carlos Slim Foundation in Mexico. The authors declare no other conflicts of interest or outside funding from any other organizations.

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Availability of supporting data

Data will be made available upon reasonable request to the corresponding author.

Supplemental material

Supplemental material for this article is available online.

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