The efficacy and tolerance of prone positioning in non-intubation patients with acute hypoxemic respiratory failure and ARDS: a meta-analysis

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Abstract
Background and aims: The application of prone positioning with acute hypoxemic respiratory failure (AHRF) or acute respiratory distress syndrome (ARDS) in non-intubation patients is increasing gradually, applying prone positioning for more high-flow nasal oxygen therapy (HFNC) and non-invasive ventilation (NIV) patients. This meta-analysis evaluates the efficacy and tolerance of prone positioning combined with non-invasive respiratory support in patients with AHRF or ARDS.

Methods: We searched randomized controlled trials (RCTs) (prospective or retrospective cohort studies, RCTs and case series) published in PubMed, EMBASE and the Cochrane Central Register of Controlled Trials from 1 January 2000 to 1 July 2020. We included studies that compared prone and supine positioning with non-invasive respiratory support in awake patients with AHRF or ARDS. The meta-analyses used random effects models. The methodological quality of the RCTs was evaluated using the Newcastle–Ottawa quality assessment scale.

Results: A total of 16 studies fulfilled selection criteria and included 243 patients. The aggregated intubation rate and mortality rate were 33% [95% confidence interval (CI): 0.26–0.42, \(I^2 = 25\%\)], 4% [95% CI: 0.01–0.07, \(I^2 = 0\%)\], respectively, and the intolerance rate was 7% [95% CI: 0.01–0.12, \(I^2 = 5\%\)]. Prone positioning increased PaO2/FiO2 [mean difference (MD) = 47.89, 95% CI: 28.12–67.66; \(p < 0.00001\), \(I^2 = 67\%\)] and SpO2 [MD = 4.58, 95% CI: 1.35–7.80, \(p = 0.005\), \(I^2 = 97\%\)], whereas it reduced respiratory rate (MD = −5.01, 95% CI: −8.49 to −1.52, \(p = 0.005\), \(I^2 = 85\%\)). Subgroup analyses demonstrated that the intubation rate of shorter duration prone (\(<=5\) h/day) and longer duration prone (\(>5\) h/day) were 34% and 21%, respectively; and the mortality rate of shorter duration prone (\(<=5\) h/day) and longer duration prone (\(>5\) h/day) were 6% and 0%, respectively. PaO2/FiO2 and SpO2 were significantly improved in COVID-19 patients and non-COVID-19 patients.

Conclusion: Prone positioning could improve the oxygenation and reduce respiratory rate in both COVID-19 patients and non-COVID-19 patients with non-intubated AHRF or ARDS.

The reviews of this paper are available via the supplemental material section.

Keywords: acute hypoxemic respiratory failure, acute respiratory distress syndrome, meta-analysis, prone positioning

Introduction
Acute hypoxemic respiratory failure (AHRF) develops from numerous pulmonary and extrapulmonary diseases or disease processes and it is a common reason for admission to the intensive care unit (ICU). A large proportion of
patients with AHRF meet the criteria for acute respiratory distress syndrome (ARDS).\(^1\)-\(^3\) Non-invasive respiratory support delivered via face mask, high-flow nasal oxygen therapy (HFNC) and non-invasive ventilation (NIV) is usually the treatment to correct hypoxia in AHRF and ARDS.\(^3\) However, certain patients exhibit further progress and require invasive mechanical ventilation. Previous studies have indicated that the intubation rate of patients with AHRF is 42–58%.\(^4\),\(^5\) Invasive mechanical ventilation may lead to increased risk of ventilator associated pneumonia and utilize medical resources for a long time period, especially during the outbreak of COVID-19. The aggregated mortality rate of patients with AHRF and ARDS may reach 27–45%.\(^6\)-\(^11\) The need for invasive mechanical ventilation may be associated with high mortality.\(^12\) Many factors are associated with high mortality, such as pneumothorax, as it is difficult to maintain the plateau pressure <30 cmH\(_2\)O and the driving pressure <15 cmH\(_2\)O. Therefore, it is necessary to actively seek other intervention measures to reduce the intubation rate and improve the prognosis of patients.

The prone positioning leads to alterations in the distribution of alveolar ventilation, improved matching of local ventilation and perfusion and reduction in regions of low ventilation/perfusion ratios through gravitational effects and reduction of ventilator-induced lung injury.\(^13\)-\(^16\) Several studies have shown that the prone positioning can improve the gas exchange and progression in patients with AHRF or ARDS.\(^17\)-\(^20\) All these studies included patients with AHRF or ARDS undergoing intubation for mechanical ventilation and most of them were under sedation. Valter et al.\(^21\) reported that four awake patients with hypoxemia were in the prone positioning without sedation and without intubation. All the patients were well tolerated with rapid improvement of PaO\(_2\). Previous studies demonstrated that the number of patients in the prone positioning combined with non-invasive respiratory support has gradually increased in recent years.\(^22\)-\(^36\) Notably during the outbreak of COVID-19,\(^22\)-\(^33\) it was reported that the prone positioning could improve oxygenation and reduce intubation rate in patients with AHRF or ARDS as well as delay or reduce the need to admit to the ICU.\(^23\) These studies suggested that the prone positioning may be used to avoid invasive mechanical ventilation and may be beneficial in several awake patients with AHRF or ARDS.

At present, the studies that have examined the prone positioning combined with non-invasive respiratory support in awake patients with AHRF or ARDS are mainly observational studies, cohort studies and case reports. Large sample-size randomized controlled trial (RCT) studies are deficient and the results are not completely consistent. The purpose of this meta-analysis was to analyze the efficacy and tolerance of prone positioning combined with non-invasive respiratory support in patients with AHRF or ARDS.

**Methods**

**Search strategy**

We searched prospective or retrospective cohort studies, RCTs and case series published in PubMed, EMBASE and the Cochrane Central Register of Controlled Trials from 1 January 2000 to 1 July 2020. The search strategy used was as follows: (respiratory distress syndrome OR ARDS) AND (Respiratory Insufficiency OR AHRF) AND (prone positioning OR prone positioning) AND (Non-invasive Ventilation OR NIV) AND (high-flow nasal cannula OR HFNC) AND Oxygen therapy. The language of publication was restricted to English. In addition, the reference lists of all primary studies and review articles were evaluated for additional relevant studies.

**Study selection**

The inclusion criteria were as follows: (i) cohort studies or case series; (ii) adult (\(\geqslant 18\) years old) patients with AHRF or ARDS and in waking state; (iii) prone positioning combined with non-invasive respiratory support (non-invasive mechanical ventilation, high flow nasal canula, venturi mask, conventional oxygen therapy); (iv) outcomes including at least one of the following measures: aggregated mortality rate, intubation rate, tolerability, prior to and following difference of arterial oxygen tension/fraction of inspired oxygen (PaO\(_2\)/FiO\(_2\)) ratios, peripheral oxygen saturation (SpO\(_2\)) and respiratory rate.

The exclusion criteria were as follows: (i) patients who did not meet the screening criteria; (ii) studies that were not in English or commentaries, reviews, duplicate publications from the same study; (iii) data that could not be extracted by the statistical methods or non-targeted outcomes.
**Data extraction and study quality**
The Newcastle–Ottawa quality assessment scale (NOS) checklist (Supplementary material Figure S1 online) was used to assess the quality of eligible studies. By using this scale, each study was assessed on seven items and categorized into three groups as follows: selection, study design and outcomes. Stars were awarded for each quality item and the highest quality studies were awarded seven stars. A study was considered to be of good, normal and poor quality when it achieved 6–7, 3–5 and 0–2 stars, respectively.

**Statistical analysis**
The meta-analysis was performed using available data from the primary studies with the R software (R version 4.0.2; Comprehensive R Archive Network, CRAN 2020) and the Review Manager software (RevMan version 5.3; Nordic Cochrane Centre, Cochrane Collaboration, 2014). The mean difference (MD) was used to describe continuous data, while the risk ratios used were for dichotomous data. These data were assessed in the median-interquartile range and were transformed into standard mean difference format for further comparison. The inverse-variance method was used to pool the mean differences to yield an overall effect size with 95% confidence intervals (CIs).

The results were analyzed by the random-effects model and were presented in a forest plot. The $I^2$ statistical index (ranges from 0% to 100%) was used to measure the heterogeneity among the studies in each analysis, with values of 25%, 50% and 75% corresponding to degrees of low, moderate and high heterogeneity, respectively. Publication bias was assessed with the Egger’s test and funnel plot. In addition, subgroup analysis was also performed to investigate different effects of prone positioning duration and etiology on treatment outcomes. A $p$-value of less than 0.05 ($p<0.05$) was considered for significant differences.

Two investigators (Xu and Tan) independently extracted data and assessed the validity of the included studies. The definitive inclusion of the trials was made following review of the full text of the articles, including publication date, study type and design, oxygen therapy method, prone positioning duration and outcomes. Discrepancies were resolved by a consensus.

**Results**

**Search results**
A total of 275 relevant publications were obtained from the databases and a total of 16 articles which were published from 2003 to 2020 were included following further screening, with four of those being cohort studies and the other 12 case series. The main characteristics of all included articles are presented in Table 1. A total of 243 patients were included in the present meta-analysis and all of them were awake adult patients (57 ± 11 years old) with AHRF or ARDS, of which 195 patients (80%) were infected by COVID-19. The patients were admitted to the hospital and received different methods of non-invasive respiratory support, of which 32 patients (13.1%) were on NIV and 39 (16%) on HFNC (HFNC/venturi mask), 86 patients (35.4%) used nasal cannulas/facial masks, and the oxygen therapy methods for the remaining 86 patients (35.4%) were not clear. The process of searching and screening is described in the flow chart (Figure 1).

**Literature quality and bias assessment**
The quality of the included literature was assessed by a modified NOS checklist and the results are shown in Table 1. All articles were of medium quality ($\geq 3$ stars) or above and three articles were considered as high-quality studies ($\geq 6$ stars). The Egger’s test results ($p<0.001$) and asymmetric funnel plot (Egger’s test funnel plots) suggested the presence of publication bias (Figure 2).

**Intubation rate**
A total of 71 out of the 243 patients (29.2%) from 16 studies were ultimately intubated and received invasive mechanical ventilation with an aggregated intubation rate of 0.33 [95% confidence interval (CI): 0.26–0.42, $I^2=25\%$]. Among these patients, 195 patients from 11 studies had COVID-19-related AHRF or ARDS, with an aggregated intubation rate of 0.32 (95% CI: 0.23–0.43, $I^2=36\%$) compared with 0.33 (95% CI: 0.20–0.54, $I^2=9\%$) noted in 48 non-COVID-19 patients from five studies. In 10 studies, a total of 214 patients performed prone positioning $\leq 5$ h/day (shorter duration prone), and their intubation rate was 0.34 (95% CI: 0.25–0.45, $I^2=44\%$), whereas 29 patients from six other studies who performed prone
| Reference          | Study Type | Cause          | Patients Age (mean ± SD) | No. of patients | Therapy Duration (mean) | Oxygen therapy method | Outcomes | Mortality | Modified NOS quality score |
|--------------------|------------|----------------|--------------------------|-----------------|-------------------------|-----------------------|----------|-----------|---------------------------|
| Bellone and Basile | Cohort     | Non-COVID-19   | 59 ± 5                   | 3               | 6 h/day                 | HFNC                  | 0        | 0         | 3/7                       |
| Caputo et al.      | Case series| COVID-19       | 57 ± 7                   | 42 of 56        | 3 h/day                 | NIV, HFNC or face masks| 18       | N/A       | 5/7                       |
| Coppo et al.       | Cohort     | COVID-19       | 56 ± 12                  | 10              | 16 h/day                | HFNC or nasal cannulas| 2        | 0         | 5/7                       |
| Damarta et al.     | Case series| COVID-19       | 50 ± 10                  | 20              | 4 h/day                 | HFNC or NIV           | 9        | 1         | 6/7                       |
| Ding et al.        | Cohort     | Non-COVID-19   | 59 ± 6                   | 1 of 10         | >5 h/day                | Nasal cannulas or HFNC| 1        | 1         | 5/7                       |
| Elharrar et al.    | Cohort     | COVID-19       | 66 ± 10                  | 24              | >3 h/day                | HFNC or nasal cannulas| 5        | 0         | 5/7                       |
| Huang et al.       | Cohort     | COVID-19       | 59 ± 3                   | 3               | 8 h/day                 | HFNC and NIV          | 1        | N/A       | 3/7                       |
| Ng et al.          | Case series| COVID-19       | 66                       | 1 of 10         | >5 h/day                | HFNC                  | 1        | 1         | 5/7                       |
| Pérez-Nieto et al. | Case series| Non-COVID-19   | 36 ± 14                  | 6               | 5 h/day                 | HFNC or NIV           | 2        | 1         | 5/7                       |
| Sartini et al.     | Case series| COVID-19       | 59 ± 6                   | 15              | 3 h/day                 | NIV                   | 1        | 1         | 4/7                       |
| Sztajnbok et al.   | Case series| COVID-19       | 40 ± 4                   | 2               | 8 h/day                 | Face masks            | 0        | 0         | 3/7                       |
| Thompson et al.    | Cohort     | COVID-19       | N/A                      | 29              | 5 h/day                 | Nasal cannulas or face masks| 12       | 3         | 5/7                       |
| Tu et al.          | Case series| COVID-19       | 51 ± 11                  | 9               | 4 h                     | HFNC                  | 2        | 0         | 6/7                       |
| Valter et al.      | Case series| Non-COVID-19   | 55 ± 27                  | 4               | 3 h                     | Three NIV and one unknown| 0        | 1         | 4/7                       |
| Scaravilli et al.  | Case series| Non-COVID-19   | 58 ± 23                  | 15              | 2 h                     | Nasal cannulas, HFNC or NIV| 2        | 3         | 5/7                       |
| Xu et al.          | Case series| COVID-19       | 50 ± 9                   | 10              | >16 h/day               | HFNC                  | 0        | 0         | 5/7                       |

Mortality was defined as early all-cause mortality used for all studies where available. HFNC includes high flow nasal cannula and venturi mask. Non-COVID-19: acute hypoxemic respiratory failure or acute respiratory distress syndrome caused by drowning, sepsis, trauma and other diseases. COVID-19: coronavirus disease 2019; HFNC, high flow nasal cannula; N/A, not applicable; NIV, non-invasive ventilation; NOS, Newcastle–Ottawa quality assessment scale; PP, prone positioning; SD, standard deviation.
Figure 1. Selection of studies for the meta-analysis [Preferred Reporting Items for Systematic Reviews and Meta-Analyses].

Figure 2. Funnel plots of the proportion versus the standard error of intubation proportion (a) and mortality proportion (b). Studies included in the meta-analysis are represented by the circles.
Table 2. Comparison of different subgroups.

| Subgroup         | No. of studies | No. of patients | Mean ± SD or (95% CI) | PP ≤5 h/day           | PP >5 h/day           |
|------------------|----------------|----------------|-----------------------|-----------------------|-----------------------|
| **COVID-19**     |                |                |                       |                       |                       |
| Basic PaO2/FiO2  | 5              | 195            | 0.33 (0.20–0.54)      | 10                    | 0.30 (0.18–0.43)      |
| Intubation rate  | 5              | 108            | 0.34 (0.25–0.50)      | 9                     | 0.34 (0.25–0.50)      |
| Mortality rate   | 5              | 138            | 0.00 (0.00–0.00)      | 6                     | 0.01 (0.00–0.02)      |
| Improvement in the PaO2/FiO2 | 4 | 61             | 52.02 (15.36–87.61)  | 4                     | 52.02 (15.36–87.61)  |
| Improvement in SpO2 | 5 | 112            | 5.23 (1.25–9.22)     | 4                     | 4.85 (1.22–9.28)     |
| Change in respiratory rate | 4 | 48             | −5.00 (−5.28–−4.73)  | 2                     | −5.00 (−5.28–−4.73)  |
| Intolerance rate | 5              | 81             | 0.06 (0.00–0.13)      | 2                     | 0.06 (0.00–0.13)      |

| **Non-COVID-19** |                |                |                       |                       |                       |
| Basic PaO2/FiO2  | 4              | 45             | 0.38 (0.19–0.57)      | 4                     | 0.38 (0.19–0.57)      |
| Intubation rate  | 4              | 40             | 0.23 (0.10–0.46)      | 10                    | 0.23 (0.10–0.46)      |
| Mortality rate   | 4              | 138            | 0.00 (0.00–0.00)      | 9                     | 0.00 (0.00–0.00)      |
| Improvement in the PaO2/FiO2 | 4 | 61             | 52.02 (15.36–87.61)  | 4                     | 52.02 (15.36–87.61)  |
| Improvement in SpO2 | 5 | 112            | 5.23 (1.25–9.22)     | 4                     | 4.85 (1.22–9.28)     |
| Change in respiratory rate | 4 | 48             | −5.00 (−5.28–−4.73)  | 2                     | −5.00 (−5.28–−4.73)  |
| Intolerance rate | 5              | 81             | 0.06 (0.00–0.13)      | 2                     | 0.06 (0.00–0.13)      |

Basic PaO2/FiO2, PaO2/FiO2 measured before prone positioning. Non-COVID-19, acute hypoxemic respiratory failure or acute respiratory distress syndrome caused by drowning, sepsis, trauma and other diseases. Significant difference was found between COVID-19 patients and non-COVID-19 patients (p<0.05).

Positioning >5h/day (longer duration prone) had an intubation rate of 0.21 (95% CI: 0.10–0.45, P=0%) (Table 2).

### Mortality rate

A total of 15 out of the 186 patients (8.1%) from 14 studies were ultimately deceased with an aggregated mortality rate of 0.04 (95% CI: 0.01–0.07, P=0%). Among these patients, 138 patients from nine studies had COVID-19-related AHRF or ARDS with an aggregated mortality rate of 0.03 (95% CI: 0.00–0.07, P=0%) compared with 0.08 (95% CI: 0.01–0.16, P=0%) in 48 non-COVID-19 patients from five studies. A total of 160 patients in nine studies performed prone positioning ≤5h/day and their aggregated mortality rate was 0.06 (95% CI: 0.02–0.11, P=23%), whereas 26 patients from five other studies who performed prone positioning >5h/day exhibited an aggregated mortality rate of 0.00 (95% CI: 0.00–0.08, P=0%) (Table 2).

### Improvement in the PaO2/FiO2 ratios

The PaO2/FiO2 ratios of a total of 101 patients were recorded prior to and following prone positioning in eight studies. Meta-analysis indicated that the PaO2/FiO2 ratios of the patients following prone positioning were significantly higher than prior to prone positioning (MD = 47.89, 95% CI: 28.12–67.66, p < 0.00001, I² = 67%). Further subgroup analysis according to the causes of disease formation indicated significant improvement in PaO2/FiO2 ratios both of COVID-19 (MD = 52.06, 95% CI: 5.36–98.76, p = 0.03, F = 81%) and non-COVID-19 patients (MD = 47.11, 95% CI: 21.16–73.06, p = 0.0004, F = 56%) (Figure 3).

### Improvement in SpO2

A total of six studies recorded SpO2 changes of 155 patients prior to and following prone positioning and a significant improvement was noted (MD = 4.58, 95% CI: 1.35–7.80, p = 0.005, F = 97%). Further subgroup analysis according to the causes of disease formation indicated significant improvement in SpO2 both for COVID-19 patients (MD = 5.23, 95% CI: 1.25–9.22, p = 0.01, F = 98%) and non-COVID-19 patients (MD = 2.00, 95% CI: 0.92–3.08, p = 0.0003, F incalculable) (Figure 4).
The changes in the respiratory rate of 87 patients prior to and following prone positioning were recorded in six studies.\cite{21,25,28-30,34} Meta-analysis indicated that prone positioning significantly reduced patient respiratory rate (MD $= -5.01$, 95% CI $= -8.49$ to $-1.52$, $p = 0.005$, $I^2 = 85\%$) (Figure 5).

Intolerance rate

A total of 12 out of the 116 patients (10.3%) from eight studies\cite{22-24,27-30,34} could not tolerate the prone positioning treatment with an aggregated intolerance rate of 0.07 (95% CI: 0.01–0.12, $P = 5\%$) (Figure 6). Among these patients, 81 patients from six studies exhibited COVID-19 related AHRF or ARDS with an aggregated intolerance rate of 0.06 (95% CI: 0.00–0.13, $P = 20\%$) compared with 0.11 (95% CI: 0.01–0.22, $P = 0\%$) in 35 non-COVID-19 patients from two studies. A total of 103 patients in five studies performed prone positioning $\leqslant 5\text{ h/day}$ and their aggregated intolerance rate was 0.09 (95% CI: 0.02–0.16, $P = 0\%$), whereas 13 patients from three other studies who performed prone positioning $> 5\text{ h/day}$ exhibited an aggregated intolerance rate of 0.00 (95% CI: 0.00–0.12, $P = 0\%$) (Table 2).

Discussion

A total of 243 awake patients with non-intubated AHRF or ARDS for prone positioning, whose data were derived from 16 studies, were included in our meta-analysis. The aggregated intubation rate was 33\%, the aggregated mortality rate was 4\% and the aggregated incidence of intolerance was 7\%. The prone positioning could significantly improve the oxygenation in awake patients with AHRF or ARDS and reduce their respiratory rate. The subgroup analysis indicated that the PaO$_2$/FiO$_2$ ratios and SpO$_2$ were significantly improved both in COVID-19 patients and non-COVID-19 patients. The intubation rate of shorter duration prone ($\leqslant 5\text{ h/day}$) and longer duration prone ($> 5\text{ h/day}$) was 34\% and 21\%, respectively; the aggregated mortality rate was 6\% and 0\%, respectively.

The PROSEVA trial in 2013 demonstrated that the prone positioning could significantly reduce the mortality of patients with ARDS\cite{38} and two
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subsequent meta-analyses indicated that the prone positioning combined with invasive mechanical ventilation could reduce the mortality of patients with severe ARDS compared with that of patients in the supine positioning (40% versus 45%, 41% versus 47%, respectively). However, it is not clear whether the prone positioning combined with non-invasive respiratory support improves the prognosis of awake patients with AHRF or ARDS. A previous study suggested that prone positioning combined with non-invasive respiratory support could delay intubation and reduce the intubation rate, while another study demonstrated that it did not improve patient outcome. The studies selected for our meta-analysis were mainly observational studies, cohort studies and case reports, including 243 awake and non-intubated patients with AHRF or ARDS. The aggregated intubation rate was 33% and the aggregated mortality rate was 4%. A previous multicenter and open-label study by Frat et al. included 330 patients with AHRF who received non-invasive respiratory support and demonstrated that the intubation rate was 42% and the mortality rate 17%. However, due to lack of RCTs, it is unclear whether the prone positioning reduces the intubation rate and mortality in awake patients with AHRF or ARDS.

Previous studies have shown that prolonged duration of prone positioning correlated with improved prognosis of patients with ARDS undergoing invasive mechanical ventilation. Two recent meta-analyses indicated that the duration of prone positioning higher than 12 h or 16 h could reduce the mortality of patients. The current guidelines recommend that the duration of prone positioning should last at least 12 h or 16 h. All patients included in the studies were awake and probably could not tolerate prolonged prone positioning. The duration of prone positioning was different in each study and varied from less than 1 h to more than 16 h, and the number of patients with the duration of prone positioning more than 12 h was very small. We tried to find the time point that balances effectiveness and tolerance like invasive mechanical ventilation. We divided the patients into a shorter duration prone group (⩽5 h/day) and longer duration prone group (>5 h/day) according to the general distribution of the duration of prone positioning in the studies and our experience in patient tolerance time. The aggregated intubation rate of the shorter duration prone group and longer duration prone group was 34% and 21%, respectively; the aggregated mortality rate was 6% and 0%, respectively. This may be related to the mild

Figure 4. Effect of prone positioning on SpO2 among patients with non-intubated AHRF or ARDS caused by COVID-19 and non-COVID-19 reasons [AHRF or ARDS caused by drowning, sepsis, trauma, and other diseases].

AHRF, acute hypoxemic respiratory failure; ARDS, acute respiratory distress syndrome; CI, confidence interval; COVID-19, coronavirus disease 2019; IV, inverse variance; PP, prone positioning; SD, standard deviation.
condition of patients who can tolerate >5 h/day of prone positioning. However, we could not make a statistical comparison between the two groups because of different interventions in the studies. Different implementation schemes, such as duration and interval, and different clinical experience of each center may lead to different results. The current data is limited and RCT studies are needed to assess these outcomes in future.

The results indicated that prone positioning combined with non-invasive respiratory support could improve oxygenation in patients with AHRF or ARDS, which was similar to the results obtained by prone positioning in patients with invasive mechanical ventilation. Recent studies demonstrated that prone positioning may provide benefits to patients with early ARDS severe ARDS and with localized infiltrates among

Figure 5. Effect of prone positioning on respiratory rate among patients with non-intubated AHRF or ARDS.
AHRF, acute hypoxemic respiratory failure; ARDS, acute respiratory distress syndrome; CI, confidence interval; PP, prone positioning; SD, standard deviation.

Figure 6. Intolerance rate of prone positioning among patients with non-intubated AHRF or ARDS caused by COVID-19 and non-COVID-19 reasons (AHRF or ARDS caused by drowning, sepsis, trauma, and other diseases). Events: patients ceased prone positioning in advance for the unbearable discomfort brought by prone positioning.
AHRF, acute hypoxemic respiratory failure; ARDS, acute respiratory distress syndrome; CI, confidence interval; COVID-19, coronavirus disease 2019.
patients with invasive mechanical ventilation. The response of different causes of ARDS to the prone positioning was also different.\textsuperscript{50-55} The number of patients included in the present study was small and most of them were COVID-19 patients. A limited number of patients with other causes of pneumonia, trauma and drowning exhibited significantly improved $\text{PaO}_2/\text{FiO}_2$ ratios and $\text{SpO}_2$ for the two groups. However, the awake patients with AHRF or ARDS selected in these studies demonstrated incomplete data regarding the parameters starting time of the prone positioning (early versus late), severity of hypoxemia, different causes (pulmonary versus extrapulmonary), the type of infiltrates (focal versus diffuse) and other relevant data. In addition, a lack of RCTs was also noted and patients who benefited more from prone positioning required further studies in the future.

Several observational studies demonstrated that the oxygenation could be continuously improved when patients returned from the prone to the supine positioning with limited duration of oxygenation.\textsuperscript{45,56,57} The three studies included in our meta-analysis demonstrated that the $\text{PaO}_2/\text{FiO}_2$ ratios of patients were decreased following returning from the prone to the supine positioning (1 h, 6–8 h and 6–12 h, respectively).\textsuperscript{25,27,34} Because the patient’s condition was mild and the duration of prone position was relatively short, the duration of oxygenation improvement after returning to the supine positioning was limited. Previous studies have found that the prolonged protocol of prone positioning was often adopted in patients with invasive mechanical ventilation and most of the patients were under sedation, which maintains the patient in the prone position for a prolonged time period while minimizing the number of rotations per day, thus minimizing the riskiest protocol.\textsuperscript{58} However, the abbreviated protocol may be another option for awake patients with AHRF or ARDS which allows the flexibility of keeping the patient prone for several hours and supine for a certain time, and then repeats the above steps. The abbreviated protocol of prone positioning may facilitate the maintenance of oxygenation because each duration of supine positioning is short, minimizes pressure-related skin ulceration and intolerance, and does not increase operational risk as the patients are awake, but may increase oxygen consumption for multiple daily repositioning maneuvers. Further clinical studies are required to determine whether the abbreviated protocol of prone positioning is more suitable for awake patients with AHRF or ARDS.

The tolerance of 116 patients to the prone positioning was reported in eight studies included in our meta-analysis. The results indicated that intolerance occurred in 12 patients with prone positioning, with an aggregated intolerance rate of 7\%. The following intolerances were mainly noted: discomfort feeling, non-cooperation and aggravated cough; there was an absence of serious adverse complications. Previous studies demonstrated that the prone positioning combined with invasive mechanical ventilation could increase the risk of pressure sores,\textsuperscript{59} endotracheal tube displacement,\textsuperscript{18,20} obstruction of the endotracheal tube, dislodgement of the thoracostomy tube\textsuperscript{17} and venous access loss.\textsuperscript{19} Patients included in the present study were awake and the main complication was subjective intolerance. Overall, the patients exhibited optimal tolerance and the method used was feasible and safe. No risk of endotracheal tube displacement and low incidence of pressure scores was noted in patients with the prone positioning combined with non-invasive respiratory support compared with those with invasive mechanical ventilation, which may be associated with the patient’s waking state, the absence of sedation and the short duration of the prone positioning. Further studies are required in order to assess the potential of reducing patient intolerance and to improve patient comfort through psychological intervention, anti-anxiety or sedative drugs and palliative therapy for awake patients in the prone positioning.

The present study exhibits several limitations. First, the patients included were mainly patients with COVID-19, which may affect the results obtained. Second, our results were based on relatively small trials and the quality of the evidence in the studies was low. A lack of RCTs was noted, which may reduce accuracy and heterogeneity. Finally, the data in most of the studies were incomplete and our preliminary analysis lacked the starting time of the prone positioning (early versus late), severity of hypoxemia and the different causes responsible for the type of infiltrates (focal versus diffuse). Oxygen device differs from one study to another as well as $\text{FiO}_2$. The time of prone positioning was very different in different studies and the number of patients with longer duration prone was also small. Whether the prone positioning could reduce the intubation rate and mortality of patients remained unclear.
Conclusions
In summary, our meta-analysis demonstrated that the aggregated intubation rate and the mortality rate of patients with non-intubation AHRF or ARDS for prone positioning was 33% and 4%, respectively, whereas the intolerance rate was 7%. The prone positioning could improve the oxygenation both in COVID-19 patients and non-COVID-19 patients and reduce the respiratory frequency of awake patients with non-intubation AHRF or ARDS.

Author contributions
Wei Tan, Dong-yang Xu, Meng-jiao Xu, Zan-feng Wang, Li-li Li: Formal analysis; Resources; Writing-original draft.

Bing Dai, Li-li Li, Hong-wen Zhao, Wei Wang, Jian Kang: Methodology; Supervision; Writing review & editing.

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Supplemental material
The reviews of this paper are available via the supplemental material section.

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