

Effect Of Oxygen Administration Protocols On Carbon Dioxide Accumulation In Emergency Pneumonia Patients

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Abstract

Oxygen therapy is essential in the management of emergency pneumonia, but differing administration protocols may influence carbon dioxide CO₂ retention and overall respiratory safety. This research aims to evaluate the effects of various oxygen delivery strategies on CO₂ levels in 120 adult patients presenting with emergency pneumonia. Participants were randomly assigned to one of three groups: Low-Flow Nasal Cannula (LFNC), Medium-Flow Face Mask (MFM), or Titrated Oxygen (TOX) targeting optimal oxygen saturation (SpO₂). Transcutaneous carbon dioxide (TcCO₂) levels were measured at baseline, 20, 40, and 60 minutes, alongside continuous monitoring of SpO₂, Respiratory Rate (RR), and arterial blood gases. The primary outcome was the temporal change in TcCO₂, while secondary outcomes assessed the proportion of patients exceeding clinically significant CO₂ thresholds. Statistical analysis was performed using repeated measures ANOVA with post-hoc comparisons (SPSS), with significance set at P < 0.05. Results demonstrated protocol-dependent differences in CO₂ accumulation. At 60 minutes, mean TcCO₂ rose by 42.0±3.2 mmHg in the Medium-Flow mask group (MFM), the Titrated Oxygen group (TOX), and the Low-Flow Nasal Cannula group (LFNC). Clinically significant elevations in TcCO₂ (≥4 mmHg) occurred in medium-flow patients, compared to the titrated group and the low-flow group. These findings suggest that titrated and low-flow oxygen protocols minimize CO₂ retention while maintaining adequate oxygenation, highlighting the importance of individualized oxygen therapy in emergency pneumonia to prevent hypercapnia and optimize respiratory safety.

Keywords: Oxygen Therapy, Carbon Dioxide (CO₂) Accumulation, Emergency Care, Pneumonia, Oxygen Administration Protocols.

INTRODUCTION

Pneumonia is an acute lung parenchymal respiratory infection causing inflammation, impaired gas exchange, and life-threatening consequences. Unfortunately, pneumonia remains one of the most common causes of morbidity and mortality worldwide and is particularly dangerous for adults who present to the emergency room [1]. Oxygen therapy is a necessary management strategy for acute pneumonia, with the goal for the clinician to correct hypoxemia and achieve adequate tissue oxygenation. In the clinical environment, there are various delivery systems for oxygen therapy, including low-flow nasal cannulas, faces masks, and patient-modulated titration systems [2].

Despite the above-mentioned benefits of oxygen therapy, it can lead to retention of carbon dioxide (CO₂) in patients with pneumonia, ventilation, lung compliance, and increased dead space. The retention of carbon dioxide will exacerbate respiratory distress and change acid-base balance and subsequently lead to hypercapnic respiratory failure [3]. Therefore, understanding different oxygen modalities and how they affect CO₂ dynamics is critical so that it optimizes patient safety and outcomes in such a challenging emergency environment.

Low-flow oxygen is thought to be safe and well tolerated but is inadequate to treat severe hypoxemia [4]. In contrast, medium- or high-flow systems can result in excessive oxygenation and oxygen-induced hypercapnia in patients with compromised respiratory mechanics. Titrated oxygen delivery has focused on adequate oxygen saturation (SpO₂) and minimizing risk, and therefore is attractive for its accuracy and lower risk of hypercapnia [5].

However, its use depends on continuous monitoring as well as availability of equipment and training, and this availability continues to widely vary among emergency departments. Persistent hurdles continue to exist, including inadequate standardized treatment protocols and little application of real-time monitoring to early recognize accumulation of CO₂ [6]. Because of the presented hurdles, the need for systematic comparisons to assess the safety and effectiveness of respective oxygen-delivery approaches for emergency pneumonia is critical. Protocols that can maintain oxygenation and avoid accumulation of CO₂ represent advancement in personalized treatments with better respiratory safety for this at-risk population.

This research assesses the impact of various oxygen administration protocols on CO₂ build-up and respiratory safety in emergency pneumonia patients.

RELATED WORK

Research analyzed relationships between ambient CO and respiratory diseases and outpatient visits per day in Lanzhou [7]. A time-series design used 2014–2019 visits (1,096,115) in an equipoising regression and distributed lag models. For each 1 mg/m³ increase in CO, there was an increase of 1.23% for total respiratory visits and 2.53% for asthma visits. Cold season and female sex were found to be associated with stronger effects. Limitations included a single-city study, residual confounding, and fixed-site monitoring uncertainty.

The research examined the role of Superoxide Dismutase type 1 (SOD1) in lipoid pneumonia, using event-control learning reconsidering intervention and 134 cases and 134 matched controls, adjusted with logistic regression and Receiver Operating Characteristic (ROC) curves. Higher SOD1 was associated with lower odds of lipoid pneumonia, ROC 0.739 with sensitivity 59.7% and specificity 82.4% [8]. Results indicated diagnostic utility in addition to routine markers. Limitations included a single-center design, retrospective data, and unmeasured confounding.

A turn-on Near-Infrared (NIR) fluorescent probe for CO tracking in infectious pneumonia and Cd²⁺ stress in plants was brought into research. The method included synthesis, spectroscopic characterization, and imaging in biological systems, published in *Journal of Hazardous Materials* 495:139135. Outcomes explained NIR emission with CO-induced signal enhancement and successful in vivo/in situ imaging demonstrations [9]. The reporting emphasized the dual applicability of the findings across both infection method and plant stress models. Constraints were present in the abstract-level revelation of performance measures and limited public access to complete quantitative parameters. The investigation determined if preoperative Fluorodeoxyglucose Positron Emission Tomography (FDG-PET) visual uptake in interstitial pneumonia fields predicted acute exacerbation following lung resection. 193 cases of lung cancer received PET-Computed Tomography (CT) and qualitative Interstitial Pneumonia (IP)-field uptake scores. Acute Exacerbation (AE) incidence was 10.0% with FDG accumulation versus 0%; 90-day mortality was 6.9% versus 0% [10]. Multivariable analysis identified FDG accumulation as an independent risk factor for AE. Retrospective single-institutional data and subjective visual scores were limitations.

Research discussed lung cancer operation following COVID-19 infection in the context of simple interstitial pneumonia and restraining ventilatory damage. Treatment included left basal segmentectomy 7 weeks from the time of infection, with serial pulmonary function monitoring subsequently at 61.7 at 12 weeks [11]. The procedure was completed without AE-IP in early follow-up. Limitations were single-patient data and limited generalizability. Research compared baricitinib with methylprednisolone in severe pneumonia. The hospital cohort included 182 patients compared to control vs standard care [12]. The combination enhanced peripheral SpO₂ and resulted in a lower oxygenation index compared to the control group; adverse reactions were not significantly different. Drawbacks were non-blinded design specifics and missing granular effect sizes in open summaries.

MATERIALS AND METHODS

The research was conducted as a prospective, randomized controlled trial to ascertain how various oxygen delivery strategies affect CO₂ accumulation in patients who present to the emergency department with pneumonia. Adult patients requiring oxygen therapy at the time of admission were considered eligible. The participants were arbitrarily allocated to one of three interference arms and observed for transcutaneous CO₂ and associated respiratory parameter changes. The research was conducted in adherence to ethical guidelines, and informed consent was provided by all patients before they participated.

Participation

A total of 120 adult patients (≥18 years) with pneumonia based on clinical presentation, radiographic evidence, and laboratory results were included.

Inclusion Criteria

Adult patients who were 18 years and older were eligible if they had confirmed cases of acute pneumonia as identified by clinical assessment, radiography, and laboratory work. The participants needed to require emergency oxygen therapy, which was given as an SpO₂ of less than 92% on room air, and must be hemodynamically stable during presentation.

Exclusion Criteria

Patients with a history of long-standing respiratory illnesses like Chronic Obstructive Pulmonary Disease (COPD), Obstructive Sleep Apnea (OSA), or Interstitial Lung Disease (ILD), or who were intubated or

needed mechanical ventilation upon admission, were excluded. Exclusion was also made for patients with neuromuscular disorders impacting the respiratory system, patients with established severe hemodynamic instability or shock, and those who refused to give informed consent to participate in the research.

Group Allocation

The eligible patients were arbitrarily allocated in a 1:1:1 ratio to one of the three intervention groups through a computer-generated random sequence with sealed opaque envelopes. The Low-Flow Nasal Cannula Group (LFNC) was administered oxygen at 2–4 L/min through nasal cannula, the Medium-Flow Face Mask Group (MFM) was administered oxygen at 6–8 L/min through a simple face mask, and the Titrated Oxygen Group (TOX) was administered dynamically titrated oxygen through a nasal cannula or Venturi mask to keep (SpO_2) between 92% and 96%. Patient and clinician blinding was not possible because the intervention consisted of overtly different methods of oxygen delivery; outcome assessors and data analysis were blinded to group assignments to minimize assessment bias.

Data Collection

A total of 120 patients were recruited and divided similarly into the three oxygen supply protocols (n = 40 per group). All patients were monitored for 60 minutes uninterrupted once oxygen therapy was initiated. Transcutaneous CO_2 was assessed by a calibrated transcutaneous monitor at baseline, 20, 40, and 60 minutes, whereas SpO_2 was measured with continuous pulse oximetry. Respiratory Rate (RR) was counted manually at each time point, and arterial blood gas was obtained at baseline and at 60 minutes to validate CO_2 trends. All the measurements were taken by trained nursing personnel according to standardized clinical protocols.

Measures

The main outcome of the research was temporal alteration in transcutaneous CO_2 levels among the three groups during the 60-minute period of monitoring. Secondary outcomes were the proportion of patients who had developed clinically significant CO_2 retention, that is, a rise of ≥ 4 mmHg above baseline, as well as assessment of oxygenation status by maintenance of (SpO_2) within the optimal range, and assessment of alteration of RR and arterial blood gas parameters.

Sample Size Calculation

The sample size was determined using a difference of at least 2 mmHg in $TcCO_2$ increase between groups, given a standard deviation of 2.5 mmHg, and $\alpha = 0.05$, and power $(1-\beta) = 0.8$. The total number of patients in each group, therefore, was at least 35. For the dropout rate, this was increased to 40 patients per group, resulting in a total of 120 patients recruited.

Statistical Analyses

Each of the statistical analyses was carried out using IBM SPSS version 25.0. Continuous variables were reported as mean \pm standard deviation with a check for normal distribution before testing. Using repeated measures ANOVA, transcutaneous CO_2 changes over time were compared across the three intervention groups. For significant group changes, post-hoc pair-wise comparisons with Bonferroni adjustment were used to find specific intergroup differences, which were evaluated, and a two-tailed p value < 0.05 was considered statistically significant.

RESULT AND DISCUSSION

This research is intended to assess the effect of different strategies of oxygen administration on CO_2 load in emergency pneumonia patients in connection with maximising respiratory safety and clinical effectiveness in adult patients.

Temporal Change in $TcCO_2$

$TcCO_2$ levels increased over time in each group, with differing rates of change in the group and significant grouping differences. Table 1 and Figure 1 display the changes in $TcCO_2$ over time for the oxygen groups.

Table 1: Time-dependent Change in $TcCO_2$ across the LFNC, MFM, and TOX groups.

Time (Minutes)	LFNC	MFM	TOX
0	41.2 \pm 3.4	41.5 \pm 3.2	41.1 \pm 3.6
20	41.5 \pm 3.3	42.9 \pm 3.4	41.9 \pm 3.4
40	41.8 \pm 3.2	44.2 \pm 3.5	42.3 \pm 3.3
60	42.0 \pm 3.2	45.3 \pm 3.6	42.6 \pm 3.4

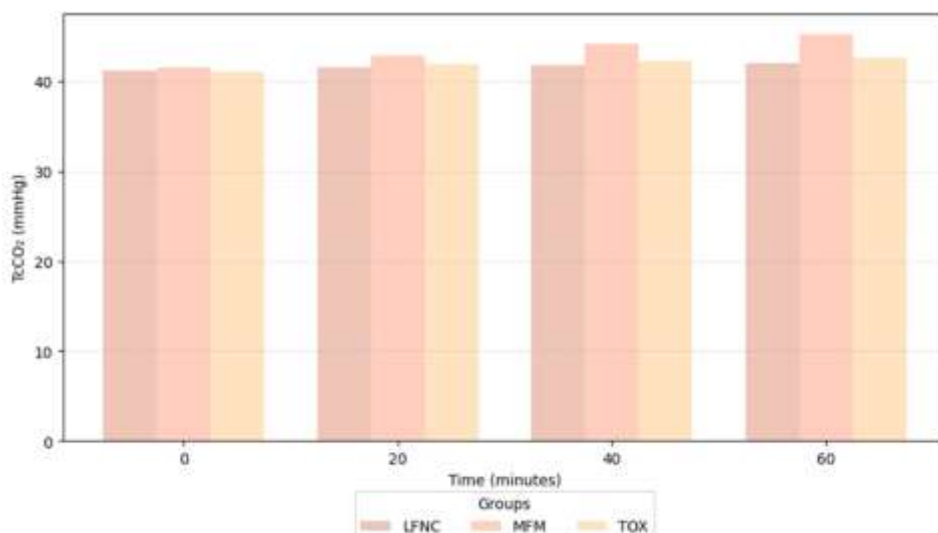


Figure 1: Protocol-dependent differences in TcCO₂ accumulation in patients with pneumonia. All groups had slow and steady increases across each status over 60 minutes; however, the MFM group exhibited the largest increase, suggesting a higher degree of CO₂ retention. LFNC and TOX reported moderate increases, suggesting that these methods were more effective at reducing CO₂ build-up while providing stabilization for these patients.

Clinically Important CO₂ Accumulation

The percentage of patients with TcCO₂ increase ≥ 4 mmHg differed profoundly among groups. Table 2 displays the patients exceeding the TcCO₂ threshold by oxygen groups.

Table 2: Number and Proportion of Patients Exceedingly Clinically Significant TcCO₂ Threshold Across Oxygen Administration Groups

Oxygen Groups	Patients Exceeding Threshold, n (%)
LFNC	3 (8%)
MFM	17 (42%)
TOX	6 (15%)

A higher proportion of patients in the MFM group was observed to have crossed the threshold as compared to those in the LFNC or TOX groups. This specifies that MFM usage is linked with an increased risk of CO₂ retention, while LFNC and TOX groups reflected a safer profile with fewer patients reaching the critical mark.

Oxygen Saturation and Respiratory Rate (SpO₂ and RR)

All groups preserved SpO₂ $\geq 92\%$. There were no significant differences in SpO₂ among groups over time. RR increased slightly in the medium-flow mask group but this was not significant. Table 3 shows the SpO₂ and RR over time across oxygen administration groups.

Table 3: Comparison of RR and SpO₂ among three oxygen administration regimens: TOX, MFM, and LFNC.

Time (min)	SpO ₂ (%) LFNC	SpO ₂ (%) MFM	SpO ₂ (%) TOX	RR (LFNC)	RR (MFM)	RR (TOX)
0	90.5±2.1	90.8±2.3	90.7±2.2	22.1±3.6	22.5±3.8	22.3±3.5
20	93.0±1.8	94.2±2.0	93.8±1.9	21.8±3.4	23.0±3.7	22.0±3.3
40	93.5±1.7	94.5±1.9	94.0±1.8	21.5±3.2	23.2±3.6	22.0±3.2
60	93.8±1.6	94.7±1.8	94.3±1.7	21.5±3.1	23.5±3.5	22.1±3.2

SpO₂ levels at the baseline (0 min) were comparable among groups (LFNC: 90.5 ± 2.1%, MFM: 90.8 ± 2.3%, TOX: 90.7 ± 2.2%), with corresponding RR values being approximately 22 breaths/min. With the passage of time, SpO₂ continued to improve in each group, and this was 93.8 ± 1.6% LFNC, 94.7 ± 1.8% MFM, and 94.3 ± 1.7% TOX at 60 min, with MFM experiencing the maximum rise. RR in the LFNC and TOX groups was mostly constant (21.5–22.1 breaths/min), while it increased slightly with MFM, reaching 23.5 ± 3.5 at 60 min. In general, MFM showed more improvement in oxygenation but was also related to a smaller increase in respiratory rate than LFNC and TOX.

This research assessed the impact of various oxygen administration protocols on CO₂ build-up and respiratory safety in emergency pneumonia patients. In spite of the large body of research on pneumonia, there are some gaps. The effect of ambient carbon monoxide on pneumonia outcomes and oxygen therapy effectiveness is not yet fully investigated (7). Predictive imaging strategies, such as PET-based visual assessment, are not yet combined with oxygen therapy protocols (10). Pharmacological treatments combined with oxygen therapy, such as baricitinib with corticosteroids, are not yet fully evaluated in clinical trials (12). This research evaluated the effect of different oxygen treatment protocols on CO₂ accumulation and respiratory safety in emergency pneumonia patients.

CONCLUSION

The research presented that oxygen treatment protocols affected CO₂ retention in emergency pneumonia patients. Medium-flow face masks led to the greatest CO₂ retention, whereas low-flow nasal cannula and titrated oxygen reduced hypercapnia. All groups ensured proper oxygenation, but individualized oxygen delivery was more effective in providing respiratory safety. The results highlighted the need to choose proper oxygen strategies to avoid CO₂ retention and maximize patient outcomes in emergency departments. At 60 minutes, the mean TcCO₂ was 42.0±3.2 mmHg in the MFM, 45.3±3.6 mmHg in the TOX, and 42.6±3.4 mmHg in the LFNC. Clinically significant elevations in TcCO₂ (≥4 mmHg) occurred in 42% of medium-flow patients, compared to 15% in the titrated group and 8% in the low-flow group

Limitations and Future Work

The research is limited by its brief observation time and single-center design, which can influence the external validity of the results. Future work should investigate long-term consequences of oxygen protocols, involve larger multicenter populations, and evaluate effects on clinical outcomes and safety.

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