Harmful effects of early hyperoxaemia in patients admitted to general wards: an observational cohort study in South Korea

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ABSTRACT

Objectives We evaluated the association between hyperoxaemia induced by a non-invasive oxygen supply for 3 days after emergency department (ED) arrival and the clinical outcomes at day 5 after ED arrival.

Design Observational cohort study.

Setting and patients Consecutive ED patients ≥16 years of age with available arterial blood gas analysis results who were admitted to our hospital were enrolled from January 2010 to December 2016.

Interventions The highest (PaO2MAX), average (PaO2AVG) and median (PaO2MED) PaO2 (arterial oxygen pressure) values within 72 hours and the area under the curve divided by the time elapsed between ED admittance and the last PaO2 result (AUC72) were used to assess hyperoxaemia. The AUC72 values were calculated using the trapezoid rule.

Outcomes The primary outcome was the 90-day in-hospital mortality rate. The secondary outcomes were intensive care unit (ICU) transfer and respiratory failure at day 5 after ED arrival, as well as new-onset cardiovascular, coagulation, hepatic and renal dysfunction at day 5 after ED arrival.

Results Among the 10 141 patients, the mortality rate was 5.8%. The adjusted ORs of in-hospital mortality for PaO2MAX, PaO2AVG, PaO2MED and AUC72 were 0.79 (95% CI 0.61 to 1.02; P=0.0715), 0.92 (95% CI 0.69 to 1.24; P=0.5863), 0.82 (95% CI 0.61 to 1.11; P=0.0205) and 1.53 (95% CI 1.25 to 1.88; P<0.0001). All of the hyperoxaemia variables showed significant positive correlations with ICU transfer at day 5 after ED arrival (P<0.05). AUC72 was positively correlated with respiratory failure, as well as cardiovascular, hepatic and renal dysfunction (P<0.05). PaO2MAX was positively correlated with cardiovascular dysfunction, PaO2MED and AUC72 were negatively correlated with coagulation dysfunction (P<0.05).

Conclusions Hyperoxaemia during the first 3 days in patients outside the ICU is associated with in-hospital mortality and ICU transfer at day 5 after arrival at the ED.

INTRODUCTION

Supplemental oxygen is frequently required by hypoxaemic patients and is frequently given in various clinical settings. Physicians tend to believe that oxygen is safe and beneficial for both non-hypoxaemic and hyperoxaemic patients.1 However, hyperoxaemia is associated with poor clinical outcomes.2 Patients in intensive care units (ICUs) exhibiting high arterial oxygen PaO2 and a high fraction of inspired oxygen (FiO2) experience more mortality than normoxaemic patients.3 Hyperoxaemia is associated with higher in-hospital mortality rates than normoxaemia in patients resuscitated from cardiac arrest.4–8

In addition, hyperoxaemia is associated with poor outcomes in patients with stroke, spontaneous subarachnoid haemorrhage and traumatic brain injury.9–11 A recent randomised controlled trial and a before-and-after study reported better outcomes in normoxaemic than hyperoxaemic ICU patients.12 13

Despite the evidence that hyperoxaemia is harmful, therapeutic strategies that prevent hyperoxaemia cannot be translated to all patients because the few relevant studies involved patients in ICUs or ventilator-assisted patients.3 9 12 14 A certain proportion of emergency department (ED) patients require supplemental oxygen. Of these, some require
mechanical ventilation and thus ICU admission. However, others receive supplemental oxygen non-invasively via a facial mask or nasal prong and are admitted to general wards. Only a few studies have explored the effects of hyperoxaemia on the clinical outcomes of ED patients. One study involved mechanically ventilated patients. Another study found that hyperoxaemia in ED patients induced by facial masks was harmful in those diagnosed with sepsis. However, that study involved a small number of patients, a limited disease spectrum and a single arterial blood gas (ABG) analysis result. No studies have evaluated the associations between hyperoxaemia and mortality in patients admitted to general wards. We hypothesised that hyperoxaemia induced by a non-invasive oxygen supply during the early treatment period would have adverse effects somewhat later in patients admitted to the general ward.

We evaluated the association between hyperoxaemia during the first 3 days after ED arrival and clinical outcomes day 5 after ED arrival.

METHOD
Study design and setting
We conducted a single-centre study at Gyeongsang National University Hospital, a tertiary teaching hospital located in the south central region of the Republic of Korea, from January 2010 to December 2016. This study was approved by our Institutional Review Board. All patients admitted to the ED are enrolled in the National Emergency Department Information System (NEDIS). The NEDIS was developed in 2003 to establish a national database of ED patients. The quality of the data was examined annually by the National Emergency Medical Centre, a government-funded, national ED control agency. In our ED, triage nurses and attending physicians entered patients’ data into the NEDIS, including basic demographic and time values, physiological values at ED arrival, symptoms, diagnosis, treatment details (including drugs and procedures), outcomes and other information. The data were organised using the standard NEDIS registry format in the hospital electronic medical records (EMRs). The validity of all data was checked by function modules within the system before the data are saved.

Participants
All consecutive patients ≥16 years of age with available ED ABG data who were admitted to the hospital with disease (not injury) during the study period were enrolled. We excluded patients with fewer than two PaO₂ results within 72 hours of ED arrival. We also excluded patients with a maximum value of PaO₂ within 72 hours (PaO₂max) of <60 mm Hg because we wished to compare hyperoxaemic and normoxaemic patients. Because we intended to assess delayed effects of hyperoxaemia, we also excluded patients who died prior to day 5 after ED arrival and who showed complications in the first 5 days (ICU transfer and respiratory failure, as well as new-onset cardiovascular, hepatic, renal and coagulation dysfunction).

The other exclusion criteria were transfer to other facilities after admission, discharge with no hope of recovery and left the hospital against medical advice. Patients with a hospital stay >90 days were excluded since their long stay may, potentially, be due to secondary problems. Moreover, we assumed that patients who were hospitalised >90 days would be free from the effects of early hyperoxaemia since there has been no study on how long the delayed effects of early hyperoxaemia last. Patients diagnosed with acute myocardial infarction were also excluded because supplemental oxygen is no longer recommended as a routine therapy in normoxaemic patients with acute myocardial infarction.

Data collection
The data were extracted from the hospital EMR system. The demographic values recorded were age and sex. The physiological values were systolic blood pressure, heart rate, breathing rate, body temperature, arterial oxyhaemoglobin saturation and mental status (assessed as alert, verbal, pain and unresponsive). We also extracted data from the Prehospital Record and List of Therapeutic Management sections of the EMR to determine whether a given patient had received supplemental oxygen therapy before ED arrival. All patients were given oxygen during their ED stays. The National Early Warning Score (NEWS) was calculated in each patient to access the severity of illness. The PaO₂ results determined by ABG analysis within 72 hours of ED arrival were collected. Time values between ED arrival and hospital discharge (dates of ED arrival, admission, ICU transfer, beginning of ventilator care, death and discharge) and final outcome (discharge, transfer, death or other) were also collected. Because we assessed complications of hyperoxaemia using the Sequential Organ Failure Assessment (SOFA) score components (cardiovascular, hepatic, renal and coagulation dysfunction), we evaluated the platelet count, and serum creatinine and bilirubin levels, at ED arrival and at day 5 after ED arrival, and determined whether a given patient received vasopressors. We also evaluated the use of mechanical ventilation therapy at day 5 after ED arrival to assess respiratory failure. Because the PaO₂:FIO₂ ratio could not be accurately calculated in the ED and general ward, respiratory failure was defined as the need for endotracheal intubation.

Hyperoxaemic variables
Among the hyperoxaemia metrics described by Helmerhorst et al, the highest (PaO₂max), average (PaO₂av) and median (PaO₂med) PaO₂ values within 72 hours and the area under the curve divided by the time elapsed between ED arrival and the last PaO₂ result (AUC_72) were used in this study. Using the PaO₂med as the starting value (t=0; ED arrival time) and the value at 72 hours (t=72), the AUC_72 was calculated using the trapezoid rule. Because no definition of hyperoxaemia has been established, we
used the following upper quintile values: 137 mm Hg for $\text{PaO}_{2\text{MAX}}$, 105 mm Hg for $\text{PaO}_{2\text{AVG}}$, 103 mm Hg for $\text{PaO}_{2\text{MED}}$ and 174 mm Hg for $\text{AUC}_{72}$.

**Study outcome**
The primary outcome was the 90-day in-hospital mortality rate. The secondary outcomes were ICU transfer and respiratory failure at day 5 after ED arrival, and new-onset cardiovascular, coagulation, hepatic and renal dysfunction (SOFA subscore ≥2) at day 5 after ED arrival.

**Data analysis**
Age was categorised as 16–39, 40–79 or ≥80 years. All continuous variables showed a skewed distribution, and are presented as medians with IQRs. The Mann-Whitney U test was used to compare continuous variables, and Pearson’s $\chi^2$ test for categorical variables.

Univariate logistic regression was performed using demographic and physiological data and the NEWS, laboratory values (platelet, creatinine and bilirubin levels and initial $\text{PaO}_2$) and hyperoxaemia variables. Variables that were significantly (p<0.01) associated with the outcome in univariate analyses were used in the multivariate logistic regression model. The adjusted ORs of these variables were calculated for each hyperoxaemia value to assess their association with in-hospital mortality.

The secondary outcomes (ICU transfer, respiratory failure and new-onset cardiovascular, hepatic, renal and coagulation dysfunction at day 5) were subjected to the same analyses as the primary outcome. The $\text{PaO}_{2\text{MAX}}$, $\text{PaO}_{2\text{AVG}}$, $\text{PaO}_{2\text{MED}}$ and $\text{AUC}_{72}$ values were subjected to multivariate analyses.

All p values were two-sided, and a p value of <0.05 was considered indicative of statistical significance. Analyses were performed using MedCalc V.17 (MedCalc Software BVBA, Ostend, Belgium) and Stata V.13 (StataCorp, LP, College Station, Texas, USA).

**Patient and public involvement**
No patients were involved in the research question and outcome.

![Figure 1](image-url) The study patients. AUC, area under the curve; ED, emergency department; ICU, intensive care unit.
Table 1 Baseline characteristics of the patients

| Characteristics                  | Total       | Missing data, n (%) |
|----------------------------------|-------------|---------------------|
| Number of patients               | 10,141      |                     |
| Age, year                        | 69.0 (57.0–78.0) | 0 (0)              |
| Age category, n (%)              |             |                     |
| 16–39                            | 786 (7.8)   | 0 (0)               |
| 40–79                            | 7,434 (73.3)| 0 (0)               |
| ≥80                              | 1,921 (18.9)| 0 (0)               |
| Male, n (%)                      | 6,040 (59.6)| 0 (0)               |

Physiological variables

| Systolic blood pressure, mm Hg | 130 (110–150) | 0 (0) |
| Heart rate, beats/min          | 90 (78–108)   | 0 (0) |
| Breath rate, breaths/min       | 20 (20–22)    | 4 (0.0) |
| Body temperature, °C           | 36.7 (36.4–37.2) | 10 (0.1) |
| SaO₂, %                        | 96 (93–98)    | 230 (2.3) |
| Consciousness (alert), n (%)   | 9,076 (89.5)  | 0 (0) |
| Supplemental oxygen, n (%)     | 4,012 (39.6)  | 0 (0) |
| NEWS                            | 4 (1–7)       | 236 (2.3) |

Initial laboratory results

| Platelet, ×10⁹/mm³           | 220 (162–287) | 56 (0.6) |
| Creatinine, mg/dL            | 0.87 (0.66–1.31)| 61 (0.6) |
| Bilirubin, mg/dL             | 0.68 (0.43–1.10)| 80 (0.8) |
| PO₂, mm Hg                   | 76 (59–96)    | 371 (3.7) |

Hyperoxaemia variables results

| PaO₂MAX                     | 99.0 (83.0–126.0) | 0 (0) |
| PaO₂AVG                     | 81.0 (68.9–99.0)  | 0 (0) |
| PaO₂MED                     | 80.0 (67.5–97.5)  | 0 (0) |
| AUC₇₂                       | 63.8 (23.2–153.2) | 0 (0) |
| Mortality, n (%)            | 584 (5.8)        | 0 (0) |

AUC₇₂, area under the curve divided by elapsed time between ED arrival and the last result of PO₂ within 72 hours; NEWS, National Early Warning Score; PaO₂MAX, average value of PO₂ within 72 hours; PaO₂AVG, highest value of PO₂ within 72 hours; PaO₂MED, median value of PO₂ within 72 hours; PO₂, PaO₂ of oxygen; SaO₂, oxyhaemoglobin saturation. Data are medians (IQR) unless otherwise stated.

Table 2 Univariate analysis of independent variables for 90-day in-hospital mortality

| Variable                        | OR       | P values |
|---------------------------------|----------|----------|
| Age 40–79 year                  | 2.63     | 0.0001   |
| Age ≥80 year                    | 3.20     | <0.0001  |
| Female                          | 0.54     | 0.05–0.65| <0.0001  |
| Systolic blood pressure         | 0.99     | 0.99–0.99| <0.0001  |
| Heart rate                      | 1.01     | 1.01–1.02| <0.0001  |
| Breathing rate                  | 1.07     | 1.05–1.09| <0.0001  |
| Body temperature                | 0.90     | 0.83–0.98| 0.0143   |
| SaO₂                            | 0.96     | 0.96–0.97| <0.0001  |
| Consciousness (non-alert)       | 1.96     | 1.57–2.45| <0.0001  |
| Supplemental oxygen             | 1.97     | 1.66–2.33| <0.0001  |
| NEWS                            | 1.14     | 1.11–1.16| <0.0001  |
| Platelet, ×10⁹/mm³              | 1.00     | 1.00–1.00| 0.0659   |
| Creatinine                      | 0.97     | 0.93–1.01| 0.1875   |
| Bilirubin                       | 1.15     | 1.12–1.17| <0.0001  |
| Initial PO₂                     | 0.99     | 0.99–1.00| 0.0005   |
| PaO₂MAX                         | 0.64     | 0.51–0.82| 0.0003   |
| PaO₂AVG                         | 0.57     | 0.45–0.74| <0.0001  |
| PaO₂MED                         | 0.54     | 0.42–0.69| <0.0001  |
| AUC₇₂                           | 1.59     | 1.32–1.92| <0.0001  |

AUC₇₂, area under the curve divided by elapsed time between ED arrival and the last result of PO₂ within 72 hours; NEWS, National Early Warning Score; PaO₂AVG, average value of PO₂ within 72 hours; PaO₂MAX, highest value of PO₂ within 72 hours; PaO₂MED, median value of PO₂ within 72 hours; PO₂, PaO₂ of oxygen; SaO₂, oxyhaemoglobin saturation.

No patients were involved in recruitment to this study. The study results will not be disseminated to study participants. This study is not a randomised controlled trial. No patients were involved in the study design or conduct of the study.

RESULTS

Baseline

Of the 228,326 patients who arrived at the ED during the study period, 32,821 met the inclusion criteria. After applying the exclusion criteria, 10,141 patients were eligible for analysis (Figure 1). Men accounted for 59.6% (60,40) of patients, and the median age of the study population was 69 (IQR: 57–78) years old. The total number of ABG samples was 37,908 and the mean number of ABG samples per patient was 3 (IQR 2–4) within 72 hours of ED arrival. The baseline characteristics of the patients are shown in Table 1.

Primary outcome

The results of univariate regression analyses are shown in Table 2. Patient age, sex and all physiological variables such as systolic blood pressure, heart rate, breathing rate, body temperature, arterial oxyhaemoglobin saturation and mental status were significantly associated with 90-day in-hospital mortality. Regarding the initial values of the laboratory values, bilirubin and PaO₂ were significantly associated with 90-day in-hospital mortality. The unadjusted ORs of PaO₂MAX, PaO₂AVG, PaO₂MED and AUC₇₂ were 0.64 (95% CI 0.51 to 0.82; p=0.0003), 0.57 (95% CI 0.45 to 0.74; p=0.0001), 0.54 (95% CI 0.42 to 0.69; p<0.0001) and 1.59 (95% CI 1.32 to 1.92; p<0.0001), respectively.

Because the values of the NEWS components (systolic blood pressure, heart rate, breathing rate, body temperature, SaO₂, supplementary oxygen and consciousness) had p values of <0.01 in univariate analyses, we subjected NEWS to multivariate regression analyses. The adjusted ORs of
PaO$_{2\text{MAX}}$, PaO$_{2\text{AVG}}$, PaO$_{2\text{MED}}$ and AUC$_{72}$ were 0.79 (95% CI 0.61 to 1.02; $p=0.0715$), 0.92 (95% CI 0.69 to 1.24; $p=0.5863$), 0.82 (95% CI 0.61 to 1.11; $p=0.2005$) and 1.53 (95% CI 1.25 to 1.88; $p<0.0001$), respectively (table 3).

**Secondary outcomes**

All of the hyperoxaemia variables were significantly positively correlated with ICU transfer at day 5 after ED arrival (table 4). Among the hyperoxaemia variables, AUC$_{72}$ had the highest OR for ICU transfer (4.03; 95% CI 3.25 to 5.01; $p<0.0001$). AUC$_{72}$ was positively correlated with respiratory failure as well as cardiovascular, hepatic and renal dysfunction. PaO$_{2\text{MAX}}$ was positively correlated with cardiovascular dysfunction. PaO$_{2\text{MAX}}$ and AUC$_{72}$ were negatively correlated with coagulation dysfunction (0.64; 95% CI 0.43 to 0.94; $p=0.022$ and 0.67; 95% CI 0.48 to 0.92; $p=0.015$).

**DISCUSSION**

We assessed the association between hyperoxaemia during the first 72 hours and the outcomes at day 5 after ED arrival. In univariate analyses, all of the hyperoxaemia variables showed significant correlations with the 90-day in-hospital mortality rate (table 2). After adjustment, only AUC$_{72}$ was significantly associated with the 90-day in-hospital mortality rate (table 3). AUC$_{72}$ was significantly positively correlated with ICU transfer, respiratory failure, cardiovascular dysfunction, hepatic dysfunction and renal dysfunction. PaO$_{2\text{MAX}}$ was significantly associated with ICU transfer and cardiovascular dysfunction. PaO$_{2\text{MAX}}$ and AUC$_{72}$ were negatively associated with coagulation dysfunction.

Only AUC$_{72}$ was significantly associated with 90-day in-hospital mortality in this study; the one-time hyperoxaemia values (PaO$_{2\text{MAX}}$, PaO$_{2\text{AVG}}$, and PaO$_{2\text{MED}}$) were not. Because AUC$_{72}$ is indicative of cumulative exposure to hyperoxaemia, it may reflect the degree of hyperoxaemia more accurately than the other variables.

The patients in this study were in a less-severe condition than those in previous studies. Our target population was patients who arrived at the ED but were not admitted to the ICU in the next 5 days. By contrast, previous studies involved only patients admitted to the ICU. The median NEWS was 4 (IQR: 2–7). We considered that a NEWS $\geq 5$ reflected critical illness.

We also used a non-invasive method of oxygen administration, unlike previous studies. Many patients in the ICU undergo mechanical ventilation. Mechanical ventilation may deliver a larger dose of oxygen in a more accurate manner. Patients who required higher oxygen levels, and those who were mechanically ventilated because of altered mentality or muscle weakness, were excluded from this study. We believe that this exclusion leads to a lower incidence of hyperoxaemia in this study than in previous studies. The single time exposure in non-critically ill patients under less-severe hyperoxaemic conditions may not be as harmful and may therefore fail to show statistical significance in terms of the mortality rate. Instead, ICU transfer was used as an indicator of an increase in clinical severity in this study. Therefore, AUC$_{72}$ is more suitable than the one-time hyperoxaemia values for assessing mortality and complications in these patients.

The significant association between AUC$_{72}$ and cardiovascular and hepatic dysfunction is consistent with Girardis et al., which found that strict oxygen use reduces the rates of mortality, shock and liver failure compared with conventional oxygen use. Because that previous study was conducted in the ICU, this is the first report of AUC$_{72}$ as an indicator of complications in non-ICU patients.

The PaO$_{2\text{MAX}}$ and AUC$_{72}$ values were associated with greater coagulation dysfunction. A previous study showed...
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that coagulation dysfunction, as determined by fibrin deposition, occurs in patients with hyperoxaemia-induced acute lung injury, but has not been investigated extensively. Thus, further studies should evaluate the association between coagulation dysfunction and hyperoxaemia.

Hyperoxaemia toxicity is caused by the production of reactive oxygen species (ROS), pulmonary toxicity, haemodynamic alterations and neurological damage. ROS lead to lipid peroxidation, protein oxidation, DNA damage and direct pulmonary damage mediated by damage to the alveolar capillary barrier. Another pulmonary complication includes pulmonary gas exchange impairment by adsorption atelectasis. Haemodynamic impairment by increased pulmonary vascular resistance is caused by the production of reactive oxygen species (ROS) and neuroendocrine changes, which enhance the production of reactive oxygen species (ROS), resulting in increased pulmonary vascular resistance. The lung is first affected because of higher oxygen tension. In a study on baboons, alveolar septal injury occurred following exposure to 60% O2 for 14 days. In rabbits, lung injury developed following exposure to moderate hyperoxaemia at a large tidal volume for 2 hours. In humans, symptoms can occur at 100% oxygen concentrations. After hyperoxia exposure, hyperoxic lung injury occurs, which affects the pulmonary vasculature. The lung is first affected because of higher oxygen tension. In a study on baboons, alveolar septal injury occurred following exposure to 60% O2 for 14 days. In rabbits, lung injury developed following exposure to moderate hyperoxaemia at a large tidal volume for 2 hours. In humans, symptoms can occur at 100% oxygen concentrations. After hyperoxia exposure, hyperoxic lung injury occurs, which affects the pulmonary vasculature.

Despite its limitations, this study was novel because we evaluated the association between hyperoxaemia and clinical outcomes. First, we could not rule out selection bias, as the study population was limited to patients who visited our ED and underwent blood gas analysis. Second, because the NEWS was calculated using the initial values in the ED rather than ICU-based severity scores, the assessment of severity may have been inaccurate; if so, this would introduce bias into the regression model. Third, this study was conducted in a single centre; thus, a further multicentre randomised controlled study is warranted. Fourth, we excluded the patients with poor outcomes prior to day 5 after ED arrival because there is no compelling evidence about the onset time of clinical outcome following hyperoxaemia. A further prospective randomised study is needed with regard to these patients. Fifth, this was an observational cohort study; although statistical associations were evident, causation cannot be inferred. Some relevant factors may not have been measured.

Table 4  Hyperoxaemia variables and adjusted ORs (95% CI) for secondary outcomes

| Variable | ICU transfer* | Respiratory failure† | Cardiovascular dysfunction‡ | Hepatic dysfunction§ | Renal dysfunction¶ | Coagulation dysfunction** |
|----------|--------------|---------------------|---------------------------|---------------------|------------------|-------------------------|
| PaO2MAX  | 2.81†† (2.26–3.50) | 1.45 (0.85–2.47) | 1.39‡‡ (1.09–1.78) | 0.99 (0.74–1.33) | 0.95 (0.72–1.26) | 0.64‡‡ (0.43–0.94) |
| PaO2AVG  | 2.04†† (1.61–2.57) | 1.13 (0.63–2.01) | 1.13 (0.87–1.47) | 0.96 (0.70–1.34) | 1.13 (0.84–1.53) | 1.08 (0.74–1.59) |
| PaO2MED  | 1.51†† (1.18–1.94) | 0.66 (0.34–1.30) | 1.02 (0.78–1.33) | 1.02 (0.74–1.40) | 0.97 (0.72–1.31) | 1.17 (0.81–1.69) |
| AUC72    | 4.03†† (3.25–5.01) | 2.40‡‡ (1.46–3.95) | 1.63‡‡ (1.29–2.07) | 1.53‡‡ (1.18–1.97) | 1.33‡‡ (1.05–1.68) | 0.67‡‡ (0.48–0.92) |

*Adjusted for systolic blood pressure, heart rate, body temperature, consciousness and supplemental oxygen.
†Adjusted for body temperature and consciousness.
‡Adjusted for age, sex, heart rate, SaO2, bilirubin and platelet count.
§Adjusted for age, sex, systolic blood pressure, SaO2, consciousness, supplemental oxygen, bilirubin, platelet and initial PO2.
¶Adjusted for NEWS, bilirubin, creatinine, platelet and initial PO2.
**Adjusted for sex, systolic blood pressure, heart rate, SaO2, consciousness, supplemental oxygen, bilirubin, platelet and initial PO2.
††P<0.0001.
‡‡P<0.05.
§§P<0.001.
AUC72, area under the curve divided by elapsed time between ED arrival and the last result of PO2 within 72 hours; ICU, intensive care unit; PaO2MED, median value of PO2 within 72 hours; PO2, PaO2 of oxygen. 
and outcomes in non-critically ill patients presenting to the ED.

CONCLUSION

Hyperoxaemia during the first 3 days in ED patients was associated with higher in-hospital mortality and more common ICU transfer at day 5 after ED arrival.

Contributors JHJ: conceptualised and designed the study, analysed the data, drafted the initial manuscript, critically reviewed the manuscript, approved the final manuscript as submitted and agreed to be accountable for all aspects of the work. DKK: conceptualised and designed the study, coordinated and supervised data collection, critically reviewed the manuscript, approved the final manuscript as submitted, and agreed to be accountable for all aspects of the work. TYK, CK, SHL, SCK, YJF: interpreted data, critically reviewed the manuscript, approved the final manuscript as submitted and agreed to be accountable for all aspects of the work.

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