Research

Case mix and outcomes for admissions to UK adult, general critical care units with chronic obstructive pulmonary disease: a secondary analysis of the ICNARC Case Mix Programme Database

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Abstract

Introduction Chronic obstructive pulmonary disease (COPD) is a common cause of admission to intensive care units (ICUs) in the UK. This report describes the case mix and outcomes of these patients and explores associations of measures of case mix available in the first 24 hours with outcome.

Method We conducted a secondary analysis of a high quality clinical database, the ICNARC Case Mix Programme Database, of 129,647 admissions to 128 adult, general critical care units across England, Wales and Northern Ireland for the period from 1995 to 2001.

Results Nonsurgical admissions with COPD accounted for 3752 admissions (2.9% of all admissions). Patients were acidotic (median pH 7.26, interquartile range [IQR] 7.18–7.33), hypercapnic (median arterial CO₂ tension 8.7, IQR 6.9–10.7) and hypoxic (median arterial O₂ tension/fractional inspired oxygen gradient 22.9, IQR 17.2–29.6). Overall, 2775 (73.9%) were definitely intubated and 278 (7.4%) were probably intubated in the first 24 hours in the ICU. The median (IQR) ICU length of stay was 4.0 (1.6–9.4) days and the hospital length of stay was 16 (9–29) days. A total of 827 patients (23.1%) died in the admitting ICU and 1322 (38.3%) died during hospital admission. Age, presence of severe respiratory disease, length of stay in hospital before critical care admission, cardiopulmonary resuscitation within 24 hours before admission, intubation status in first 24 hours in critical care, pH, arterial oxygen tension/fractional inspired oxygen gradient, albumin, cardiovascular organ failure, neurological organ failure and renal organ failure all had independent associations with hospital mortality. Respiratory organ failure had a significant independent association with decreased hospital mortality.

Conclusion Nonsurgical patients with COPD represent an important group of patients admitted to UK ICUs. The presence of single organ respiratory failure in the first 24 hours in critical care identifies patients with a 70% chance of surviving to leave hospital.

Introduction

The prevalence of chronic obstructive pulmonary disease (COPD) in the UK has been estimated to be 1%, increasing to 5% in men aged 65–74 years and 10% in men older than 75 years [1]. COPD accounted for more than 100,000 hospital admissions in England in 2000/2001 and is the fifth most common cause of mortality in the UK, with 30,000 deaths each year [2]. Exacerbations of COPD may lead to respiratory failure requiring ventilatory support either noninvasively or invasively following endotracheal intubation. A recent UK study [3] suggested that clinicians involved in the emergency care of COPD patients had made a median of 10 intensive care unit (ICU) gatekeeping decisions for COPD patients in the preceding 12 months.

COPD is a progressive disorder that spans a continuum from early mild disease to severe and disabling terminal disease, and prognostic estimates can be important in informing critical...
care admission decisions [4]. Factors found to have an association with mortality have included those related to diminished functional reserve resulting from the COPD, such as impairments in activities of daily living (ADL) and factors related to the aetiology of respiratory failure [5]; severity of the acute illness, measured using either the acute physiology score of APACHE (Acute Physiology and Chronic Health Evaluation) II [6] or APACHE III [5,7]; and the presence of nonrespiratory organ failures [7,8].

The majority of studies reporting COPD ICU outcomes originate from the USA, and there have only been two recent studies from the UK; one reported on 42 intubated patients in a single centre [9] and another reported hospital mortality of 277 COPD ICU admissions from 24 ICUs in one region of the UK [10].

This report examines the outcomes of COPD patients admitted to ICUs across England, Wales and Northern Ireland, identified using a high quality clinical database. The case mix at ICU admission, outcome and activity associated with these admissions are described. The effect of factors, determined a priori, on hospital mortality is investigated.

Materials and methods

Case Mix Programme Database

Data were extracted for 129,647 admissions to 128 adult, general critical care units (ICUs including combined intensive care/high dependency units) from the Case Mix Programme Database (CMPD), covering the period from December 1995 to August 2001. Data were collected locally by trained data collectors according to precise rules and definitions, and the data underwent extensive validation before incorporation into the CMPD. The process of data validation and cleaning was described in detail previously [11].

Selection of data

A standard coding method that classifies information describing the reasons for admission was used to identify the patients, and has been described previously [12]. Surgical patients were identified and excluded when the primary or secondary reason for admission was a surgical code or if the patient was admitted from theatre having undergone all or part of a surgical procedure. Nonsurgical admissions were selected if they had a primary reason for admission of COPD, exacerbation of COPD or emphysema, or any of these three conditions as the secondary reason for admission when pneumonia, right ventricular failure, or left ventricular failure was the primary reason. Patients were assigned to one of three subgroups of COPD (pneumonia in a patient with COPD, right ventricular failure in a patient with COPD, or left ventricular failure in a patient with COPD) if the primary or secondary reason for admission was pneumonia, right ventricular failure, or left ventricular failure, respectively, in a patient in whom the other reason for admission was COPD, exacerbation of COPD, or emphysema. The primary and secondary reasons for admission were the diagnoses made by the clinicians and recorded in the patient notes. As such the diagnoses were not arrived at by applying preset diagnostic criteria but by using the working clinical diagnoses reached during the management of the patients, and will reflect the uncertainty inherent in clinical practice.

In addition, the intubation status of the COPD admissions was categorized on the basis of whether they were definitely intubated within the first 24 hours of admission to the Case Mix Programme (CMP) unit, probably intubated in the first 24 hours, or not intubated in the first 24 hours. The CMPD does not contain a response field solely for the identification of intubation status, but patients with an intubated fractional inspired oxygen (FiO₂) recorded and those defined as intubated in the field describing the intubation status associated with the lowest arterial oxygen tension (PaO₂) were identified as definitely intubated. Patients who were probably intubated had a response of ‘yes’ in the field for mechanical ventilation at admission, or a non-zero value for the lowest or highest ventilated respiratory rate. Patients not intubated in the first 24 hours in the CMP unit fulfilled none of the above criteria.

Data

Data were extracted on case mix, outcome and activity for nonsurgical COPD admissions to CMP units. Detailed description of the definitions used in the CMPD are described elsewhere [11], but the definitions of particular relevance to the COPD patients are outlined below.

Case mix

Age at admission and sex were extracted. A history of respiratory functional impairment and comorbidity was recorded using data collected as part of the chronic health evaluation section of the APACHE II score [13]. The respiratory functional impairment has two aspects: ‘severe respiratory disease’ is defined as patients with permanent shortness of breath with light activity due to pulmonary disease; and ‘home ventilation’ specifies whether a patient has used or uses home ventilation (excluding continuous positive airway pressure ventilation). Patients with liver disease included those with an episode of hepatic encephalopathy in the previous 6 months or who had evidence of portal hypertension. Patients with cardiovascular comorbidity included patients with fatigue, claudication or angina at rest, in whom any activity increased the symptoms and where the symptoms were ascribed to myocardial or peripheral vascular disease. Patients with chronic renal failure were those requiring chronic renal replacement therapy. Patients with haematological diseases included those with acute myelogenous leukaemia, acute lymphocytic leukaemia, multiple myeloma, chronic myelogenous leukaemia, chronic lymphocytic leukaemia, and lymphoma. Patients with immunosuppression included those with a congenital immunohumoral or cellular immune deficiency state and those receiving chemotherapy defined as the receipt of a drug that can lower
resistance to infection and those who have received radiotherapy. AIDS patients were those meeting the current World Health Organization definition. Patients with metastatic cancer had distant metastases documented by surgery, imaging, or biopsy. History of steroid treatment was defined as the receipt of 0.3 mg/kg, or greater, prednisolone (or equivalent) daily for the 6 months before critical care admission. Patients receiving cardiopulmonary resuscitation within 24 hours before unit admission were also identified.

The following physiological variables, selected a priori, were extracted from the data collected during the first 24 hours in the CMP unit: lowest pH; arterial carbon dioxide tension from the arterial blood gas with the lowest pH; \( \text{PaO}_2 / \text{FiO}_2 \) gradient from the arterial blood gas with the lowest \( \text{PaO}_2 \); and lowest serum albumin. Organ failures were defined according to Knaus [14] using data describing the patient's status during the first 24 hours in the CMP unit. Acute severity was also measured with the acute physiology scores from APACHE II and APACHE III, and the APACHE II and APACHE III scores.

**Outcome**

Survival data were collected at discharge from the CMP unit and at ultimate discharge from an acute hospital.

**Activity**

Length of stay in the CMP unit was calculated, as fraction of days, from the date/time of admission to the CMP unit and date/time of discharge from the CMP unit or death in the CMP unit. Total length of stay in hospital was calculated, in days, from the date of original admission to hospital and date of ultimate discharge from hospital or death in hospital. Transfers in from (out to) another critical care unit were identified as admissions whose source of admission to the CMP unit (destination following discharge from the unit) was any ICU or high dependency unit in the same or another hospital. Readmissions to the CMP unit in the same hospital stay were identified from the postcode, date of birth and sex, and confirmed by the participating units.

**Analyses**

The overall proportion of admissions to CMP units with nonsurgical COPD was calculated along with the proportion by subgroups: COPD with pneumonia; COPD with right ventricular failure; and COPD with left ventricular failure. Case mix, outcome and activity were described for the overall group and for the subgroups. Admissions of patients younger than 16 years, admissions of patients who stayed for less than 8 hours in critical care, and readmissions within the same hospital stay and transfers from other ICUs were excluded from the calculation of the APACHE II acute physiology score and APACHE III score.

The effects of case mix factors, specified a priori, on ultimate hospital mortality for nonsurgical admissions with COPD were investigated using logistic regression.

Readmissions within the same hospital stay and admissions for whom hospital outcome data were missing were excluded from all analyses relating case mix factors to ultimate hospital mortality. In addition to univariate analyses, the factors were entered into a multiple logistic regression model. Adjusted subgroup effects were calculated for the three specified subgroups from the multiple logistic regression model.

All analyses were performed using Stata 8.0 (StataCorp, College Station, TX, USA).

**Results**

**Case mix**

Of 129,647 admissions to 128 adult, general critical care units in the CMPD, 3755 (2.9%) of admissions were identified as nonsurgical COPD patients. Table 1 describes measures of case mix for all patients with COPD and the subgroups with pneumonia, right ventricular failure and left ventricular failure. Overall, 35% of patients had severe respiratory disease before hospital admission and 9% had received daily steroids in the 6 months before admission, although only 2% were receiving home ventilation. Comorbidity was relatively rare, with fewer than 1% of patients having any comorbidity other than cardiovascular disease, and with cardiovascular comorbidity present in 2.2% of the group as a whole, 5.9% of those with right ventricular failure and 7.6% of those with left ventricular failure. Patients spent a median of 1 day in hospital before admission to the CMP unit, and 8.3% of patients received CPR in the 24 hours before admission to the CMP unit. Of the group as a whole, 2775 (73.9%) were definitely intubated in the first 24 hours, with an additional 278 (7.4%) probably intubated in the first 24 hours. Patients were acidic, hypercapnic and hypoxic, and had low serum albumin (Table 1). For the group as a whole the median (interquartile range [IQR]) APACHE II and APACHE III scores were 18 (14–23) and 62 (49–78), respectively. Of all admissions, 12.2% had no organ failures, 49.4% had single organ failure (the majority, 45.2% of all admissions, being respiratory), 27.6% had two organ failures, and had 8.2% three or more organ failures.

**Outcome and activity**

Overall, 827 (23.1%) patients died in the admitting CMP unit and 1322 (38.3%) died during the hospital admission (Table 2). The median (IQR) length of stay in the CMP unit was 4.0 (1.6–9.4) days and in hospital it was 16 (9–29) days. Readmissions within the same hospital stay accounted for 4.5% of all admissions. A decision was made to withdraw all active treatment in 422 (11.3%) patients, with the withdrawal
## Table 1

### Case mix for admissions with chronic obstructive pulmonary disease by subgroup

|                          | All (n = 3752) | I. Pneumonia (n = 775) | II. Right VF (n = 147) | III. Left VF (n = 142) |
|--------------------------|----------------|------------------------|------------------------|------------------------|
| **Age (years; median [IQR])** | 67.8 (60.5–73.6) | 68.7 (61.4–74.0) | 68.8 (61.7–74.0) | 71.9 (67.2–75.1) |
| **Sex (n [%] male)**      | 1942 (51.8) | 401 (51.7) | 83 (56.5) | 81 (57.0) |

**Past medical history (n [%])**

- **Respiratory impairment**
  - Severe respiratory disease 1236 (35.0) 232 (31.6) 57 (42.2) 33 (25.2)
  - Home ventilation 70 (2.0) 6 (0.8) 6 (4.4) 2 (1.5)
  - Steroid treatment 305 (8.6) 61 (8.3) 7 (5.2) 8 (6.1)

- **Comorbidity:**
  - Liver disease 6 (0.2) 3 (0.4) 0 (0.0) 0 (0.0)
  - Cardiovascular disease 79 (2.2) 11 (1.5) 8 (5.9) 10 (7.6)
  - Chronic renal failure 11 (0.3) 1 (0.1) 0 (0.0) 0 (0.0)
  - Immunosuppression 27 (0.7) 3 (0.4) 1 (0.7) 1 (0.8)
  - Metastatic cancer 6 (0.5) 1 (0.4) 0 (0.0) 0 (0.0)

- **Hospital stay prior to unit admission (days; median [IQR])** 1 (0–3) 1 (0–3) 1 (0–2) 0 (0–2)

- **CPR prior to admission (n [%])** 308 (8.3) 45 (5.8) 20 (13.8) 12 (8.5)

- **Intubation status (n [%])**
  - Definitely intubated 2774 (73.9) 605 (78.1) 104 (13.8) 103 (72.5)
  - Probably intubated 278 (7.4) 56 (7.2) 13 (8.8) 9 (6.3)
  - Not intubated 700 (18.7) 114 (14.7) 30 (20.4) 30 (21.2)

- **Physiology**
  - Lowest pH (median [IQR]) 7.26 (7.18–7.33) 7.26 (7.18–7.33) 7.25 (7.19–7.33) 7.24 (7.15–7.32)
  - PaCO₂ (kPa; median [IQR]) 8.7 (6.9–10.7) 8.6 (6.9–10.6) 9.1 (7.2–11.7) 7.6 (6.4–10.4)
  - PaO₂/FiO₂ gradient (kPa; median [IQR]) 22.9 (17.2–29.6) 21.2 (15.3–27.8) 20.9 (17.2–26.2) 23.3 (15.9–29.3)
  - Lowest serum albumin (g/l; median [IQR]) 28 (23–33) 26 (21–31) 28 (25–32) 30 (25–34)

- **APACHE II**
  - APS (mean [SD]) 13.4 (5.6) 14.3 (5.5) 13.5 (5.1) 14.6 (6.2)
  - APS (median [IQR]) 12 (10–16) 13 (10–17) 13 (10–17) 14 (10–18)
  - Score (mean [SD]) 19.4 (6.4) 20.3 (6.3) 19.9 (5.8) 20.7 (6.8)
  - Score (median [IQR]) 19 (15–23) 19 (16–23) 19 (16–24) 20 (16–24)

- **APACHE III**
  - APS (mean [SD]) 52.9 (21.5) 55.9 (21.6) 53.4 (20.0) 56.8 (21.5)
  - APS (median [IQR]) 50 (39–64) 52 (42–68) 53 (41–68) 55 (42–68)
Case mix for admissions with chronic obstructive pulmonary disease by subgroup

| Score (mean [SD]) | 65.9 (23.0) | 69.3 (23.0) | 66.6 (21.8) | 71.6 (22.9) |
|-------------------|-------------|-------------|-------------|-------------|
| Score (median [IQR]) | 63 (50–78) | 66 (54–83) | 66 (52–80) | 69.5 (55–84) |

Organ system failures (n [%])

| 0 | 457 (12.2) | 85 (11.0) | 19 (12.9) | 15 (10.6) |
|---|------------|-----------|----------|-----------|
| 1 Respiratory | 1767 (45.2) | 362 (45.0) | 56 (37.4) | 55 (33.8) |
| 1 Other | 214 (4.4) | 60 (9.3) | 5 (4.1) | 9 (5.5) |
| 2 | 1036 (27.6) | 215 (27.7) | 50 (34.0) | 50 (35.2) |
| 3 or more | 309 (8.2) | 54 (7.0) | 17 (11.6) | 17 (12.0) |

| n | [15] reported a hospital mortality of 38.6% (95% CI 24.4–39.3%) and this study (40.6%, 95% CI 38.8–42.3%). Although almost all studies report hospital mortality, measures such as 30- or 60-day mortality are more robust for comparison purposes because the definition of hospital mortality may be influenced by local discharge practices. For example, Nevins and Epstein [6] reported a comparatively low hospital mortality of 27.7% (95% CI 21.1–35.2%) in 166 intubated COPD patients from one US centre, but 38% of patients were transferred to chronic care facilities. In contrast, Ely and coworkers [7] included patients with COPD exacerbations, patients who had right ventricular failure or pneumonia were excluded. In our study, patients with right ventricular failure had a significantly increased risk for hospital mortality after case mix adjustment.

Discussion

This study describes the outcomes of 3752 COPD patients admitted to UK ICUs over a 5-year period and includes analyses that describe the association between patient characteristics available within the first 24 hours of ICU admission and hospital outcome.

Overall, patients with COPD admitted to the 128 ICUs in this study had a 77% chance of surviving to leave ICU and a 61% chance of surviving to leave hospital. Comparisons of ICU outcomes between studies are difficult to interpret because ICU beds are used in different ways in different countries and the definition of hospital mortality may be influenced by local discharge practices. For example, Nevins and Epstein [6] reported a comparatively low hospital mortality of 27.7% (95% CI 21.1–35.2%) in 166 intubated COPD patients from one US centre, but 38% of patients were transferred to chronic care facilities. In contrast, Ely and coworkers [7] included patients with COPD exacerbations, patients who had right ventricular failure or pneumonia were excluded. In our study, patients with right ventricular failure had a significant, independent increased risk for death, and the exclusion of these patients may go some way to explaining the difference in hospital mortality in intubated patients between the Seneff study (31.8%, 95% CI 24.8–39.3%) and this study (40.6%, 95% CI 38.8–42.3%). Although almost all studies report hospital mortality, measures such as 30- or 60-day mortality are more robust for comparison purposes because the definition of hospital mortality may be influenced by local discharge practices. For example, Nevins and Epstein [6] reported a comparatively low hospital mortality of 27.7% (95% CI 21.1–35.2%) in 166 intubated COPD patients from one US centre, but 38% of patients were transferred to chronic care facilities. In contrast, Ely and coworkers [7] reported a hospital mortality of 38.6% (95% CI 24.4–
and found an overall hospital mortality of 28% (95% CI 24–32%) with a median (IQR) Simplified Acute Physiology Score II of 38 (31–49), as compared with 42 (35–52) for the intubated patients in the present study.

Differences in length of stay between studies are likely to be influenced by the case mix and supply side factors discussed above. The 522 mechanically ventilated COPD patients from the study by Esteban and coworkers [16] had a median (IQR) length of ICU stay of 8 (5–13) days, as compared with 4.0 (1.6–9.4) days observed in our study. However, the total hospital length of stay was similar in both studies, with the Esteban study having a median (IQR) hospital stay of 17 (10–27) days versus 16 (9–29) days in the present study. Longer ICU stay (median 7 days, IQR 4–14) and similar hospital stay (median 14 days, IQR 9–29) are also apparent in the study by Nevins and Epstein [6]. The greater proportion of the total hospital stay spent in European and North American ICUs compared with the UK may well reflect the lower proportion of all hospital beds designated for critical care in the UK.

This study demonstrates that age has an independent relationship with hospital death. This increased risk is similar to that observed in the SUPPORT study, which calculated the independent relative hazard of death to be 1.22 (95% CI 1.05–1.41) per 10-year increase [5]. A number of studies of COPD outcome [6,8,9,17] did not find a relationship between age and hospital mortality. There is potential for prognostic studies to produce conflicting results because of inaccuracy in the measurement of patient characteristics, inadequate measurement of and adjustment for confounding, and lack of power [18,19]. Most commentators suggest that a minimum of between 15 and 20 deaths will be required per risk factor analyzed [20,21]. In three of the studies that did not find age to predict mortality [6,8,9], age was analyzed as a univariate predictor with between 20 and 46 deaths. In the study by Breen and coworkers [17], in which age was analyzed in a multivariate logistic regression, there were only 15 hospital deaths.

Of chronic diseases existing before hospitalization, only severe respiratory disease had a significant independent relationship with hospital death. Patients with severe respiratory disease have shortness of breath performing most ADL, and the increased risk for mortality for patients with severe respiratory disease is consistent with the increased hazard of death associated with impairment in ADL identified in SUPPORT (relative hazard per additional ADL impairment: 1.14, 95% CI 1.03–1.26) [5]. Prior cardiovascular disease was only present in 78 patients (2% of the total) and did not have a significant independent association with outcome. Although a 2% prevalence of cardiovascular comorbidity may seem surprisingly low in a COPD population, it is should be remembered that, to qualify for this comorbidity, patients had to have impairment equivalent to New York Heart Association functional class IV. A total of 294 patients (7.8%) had received 0.3 mg/kg steroid daily for the 6 months prior to admission. Oral steroids have
been suggested to confer an increased risk for hospital mortality in a small study conducted in intubated COPD patients using univariate analysis [9]. It is possible that the lack of association of oral steroids with death in this study may represent a true absence of association. However, the use of oral steroids in a heterogeneous population, which might have included both COPD patients with a marked asthmatic component and a better than average prognosis, as well as very severe COPD patients with a worse than average prognosis, could lead to no net effect being detected, as of course could incomplete documentation of steroid use in the population as a whole.

In a study of 362 COPD admissions to US critical care units [22] the number of days in hospital before critical care admission was reported to be significantly associated with hospital mortality, although the increased risk per day was not reported. This study confirms an association between days in hospital before ICU and hospital mortality, but it shows the independent risk to be relatively small with patients having only a 2% (95% CI 1–3%) increased odds of hospital death for each day in hospital before ICU admission.

Brochard and coworkers [23] suggested that invasive ventilation was associated with increased mortality when compared with noninvasive ventilation, but there were differences in severity between groups. Seneff and coworkers [7] found that intubation on day 1 was not an independent predictor of hospital mortality ($P = 0.07$) in a multiple regression model, with 170 patients intubated on day 1 and 54 deaths among intubated patients. Afessa and coworkers [8] also failed to find an independent association between intubation and death in a study of 250 episodes of respiratory failure in 180 COPD patients, in which 153 patients episodes involved intubation and 31 patients died. In this study, 3052 patients were definitely or probably intubated on day 1 and intubation on day 1 was associated with an independent odds ratio (95% CI) of 1.76 (1.07–2.93) for hospital mortality. Given that the risk for ventilator-associated pneumonia has been estimated to be 3% per day [24], it would not be surprising to find increased hospital mortality associated with intubation, and it is possible that the Seneff and Afessa studies were underpowered to detect this effect.

Cardiovascular, neurological and renal organ failures were all associated with a significant, independent increased risk for hospital mortality. Treatment withdrawal was reported to occur in 11.3% of admissions, and if the development of organ failure were to precipitously trigger withdrawal decisions then interpretation of the prognosis associated with organ failure would be difficult. Treatment withdrawal was most common in patients with acute renal failure, but treatment withdrawal accounted for fewer than one-third of the deaths, and recalculating the risk for death after removing all patients with withdrawal decisions resulted in little change in the OR for death associated with acute renal failure. Respiratory organ failure had a significant independent association with lower hospital mortality, with patients with isolated respiratory failure having a hospital mortality of 29.8%, as compared with 33.8% for patients without any organ failures. It is possible that patients admitted to ICU with COPD but without respiratory organ failure had more severe illness in other organ systems. For example, although the regression model adjusts for the presence or absence of acute renal failure, the severity of renal failure may still constitute residual confounding. Afessa and coworkers [8] also used the Knaus organ failure classification [14] and found an independent increased odds of hospital death of 5.5 per organ failure (excluding respiratory organ failure). Seneff and coworkers [7] found that the development of nonrespiratory organ dysfunction was the major predictor of hospital mortality in 362 COPD admissions to US ICUs. This is also consistent with a study of patients with heterogeneous causes of respira-
Table 4

**Ultimate hospital mortality by organ system failures**

| Organ system failures | n   | Mortality (95% CI) |
|-----------------------|-----|--------------------|
| None                  | 441 | 33.8 (29.4–38.4)   |
| 1 Respiratory         | 1632| 29.8 (27.6–32.1)   |
| 1 Other               | 244 | 45.9 (39.5–52.4)   |
| 2                     | 991 | 43.8 (40.7–46.9)   |
| 3+                    | 303 | 70.0 (64.5–75.1)   |

CI, confidence interval.

Table 5

**Multiple logistic regression model of patient factors in relation to ultimate hospital mortality**

|                      | Deaths | n   | (%) | Unadjusted OR (95% CI) | Adjusted OR (95% CI) |
|----------------------|--------|-----|-----|------------------------|----------------------|
| **Age**              |        |     |     |                        |                      |
| <55                  | 103    | 469 | (22.0) | 1.59 (1.46 – 1.73) | 1.57 (1.44 – 1.73) |
| 55 – 60              | 108    | 395 | (27.3) |                        |                      |
| 60 – 64              | 187    | 556 | (33.6) |                        |                      |
| 65 – 69              | 281    | 745 | (37.7) |                        |                      |
| 70 – 74              | 325    | 747 | (43.5) |                        |                      |
| 75 – 79              | 293    | 537 | (54.6) |                        |                      |
| 80+                  | 96     | 162 | (59.3) |                        |                      |
| **Sex**              |        |     |     |                        |                      |
| Female               | 630    | 1742| (36.2) | Ref.                   | Ref.                 |
| Male                 | 763    | 1869| (40.8) | 1.19 (1.03 – 1.39) | 1.04 (0.88 – 1.22) |
| **Severe respiratory disease** | | | | | |
| No                   | 827    | 2204| (37.5) | Ref.                   | Ref.                 |
| Yes                  | 487    | 1200| (40.6) | 1.15 (0.98 – 1.34) | 1.23 (1.03 – 1.46) |
| **Home ventilation** |        |     |     |                        |                      |
| No                   | 1288   | 3337| (38.6) | Ref.                   | Ref.                 |
| Yes                  | 26     | 67  | (38.8) | 1.05 (0.62 – 1.78) | 1.16 (0.65 – 2.07) |
| **Steroid treatment in previous 6 months** | | | | | |
| No                   | 1204   | 3110| (38.7) | Ref.                   | Ref.                 |
| Yes                  | 110    | 294 | (37.4) | 0.96 (0.74 – 1.24) | 1.00 (0.75 – 1.33) |
| **Comorbidity: liver disease** | | | | | |
| No                   | 1309   | 3398| (38.5) | Ref.                   | Ref.                 |
| Yes                  | 5      | 6   | (83.3) | 8.22 (0.96 – 70.47) | 3.70 (0.39 – 35.37) |
| **Comorbidity: cardiovascular disease** | | | | | |
| No                   | 1282   | 3326| (38.5) | Ref.                   | Ref.                 |
| Yes                  | 32     | 78  | (41.0) | 1.03 (0.63 – 1.68) | 0.88 (0.51 – 1.50) |
| **Comorbidity: chronic renal failure** | | | | | |
| No                   | 1306   | 3393| (38.5) | Ref.                   | Ref.                 |
| Yes                  | 8      | 11  | (72.7) | 3.29 (0.82 – 13.17) | 2.21 (0.50 – 9.77) |
| **Comorbidity: immunosuppression** | | | | | |
### Table 5 (Continued)

#### Multiple logistic regression model of patient factors in relation to ultimate hospital mortality

|                  | No   | Yes   | Ref.  | Ref.  |               |               |
|------------------|------|-------|-------|-------|---------------|---------------|
|                  | 1299 | 15    | 1.92  | 1.75  | (0.89 – 4.17) | (0.71 – 4.28) |
| Comorbidity: metastatic cancer | 1310 | 4     | 3.28  | 2.50  | (0.60 – 17.96)| (0.43 – 14.66)|
|                  |      |       | per 1 day increase | per 1 day increase |
| Length of stay before unit admission (days) | 0    | 567   | 1.02  | 1.02  | (1.01 – 1.03) | (1.01 – 1.03) |
|                  | 1    | 256   | 1.75  | 1.75  | (0.71 – 4.28) | (0.71 – 4.28) |
|                  | 2    | 111   | 1.75  | 1.75  | (0.71 – 4.28) | (0.71 – 4.28) |
|                  | 3+   | 449   | 1.75  | 1.75  | (0.71 – 4.28) | (0.71 – 4.28) |
| CPR within 24 hours before admission | No   | 1207  | 2.38  | 2.38  | (1.83 – 3.08) | (1.83 – 3.08) |
|                  | Yes  | 176   | 2.38  | 2.38  | (1.83 – 3.08) | (1.83 – 3.08) |
| Intubation status | No   | 206   | 1.71  | 1.71  | (1.38 – 2.13) | (1.38 – 2.13) |
|                  | Yes  | 1187  | 1.71  | 1.71  | (1.38 – 2.13) | (1.38 – 2.13) |
| Lowest pH<sup>a</sup> | <7.18| 391   | 1.31  | 1.31  | (1.23 – 1.40) | (1.23 – 1.40) |
|                  | 7.18–7.25 | 294   | 1.31  | 1.31  | (1.23 – 1.40) | (1.23 – 1.40) |
|                  | 7.26–7.32 | 259   | 1.31  | 1.31  | (1.23 – 1.40) | (1.23 – 1.40) |
|                  | = 7.33 | 297   | 1.31  | 1.31  | (1.23 – 1.40) | (1.23 – 1.40) |
| PaCO<sub>2</sub> from ABG with lowest pH (kPa)<sup>a</sup> | <8.9 | 334   | 1.09  | 1.09  | (0.91 – 1.29) | (0.91 – 1.29) |
|                  | 6.9–8.6 | 316   | 1.09  | 1.09  | (0.91 – 1.29) | (0.91 – 1.29) |
|                  | 8.7–10.6 | 270  | 1.09  | 1.09  | (0.91 – 1.29) | (0.91 – 1.29) |
|                  | = 10.7 | 320   | 1.09  | 1.09  | (0.91 – 1.29) | (0.91 – 1.29) |
| PaO<sub>2</sub> /FiO<sub>2</sub> gradient (kPa)<sup>a</sup> | <17.2 | 425   | 1.17  | 1.17  | (1.09 – 1.26) | (1.09 – 1.26) |
|                  | 17.2–22.8 | 300   | 1.17  | 1.17  | (1.09 – 1.26) | (1.09 – 1.26) |
|                  | 22.9–29.5 | 293   | 1.17  | 1.17  | (1.09 – 1.26) | (1.09 – 1.26) |
|                  | = 29.6 | 294   | 1.17  | 1.17  | (1.09 – 1.26) | (1.09 – 1.26) |
| Lowest albumin (g/l)<sup>a</sup> | <20 | 178   | 2.73  | 2.73  | (2.11 – 3.54) | (2.11 – 3.54) |
|                  | 20–24 | 208   | 1.49  | 1.49  | (1.20 – 1.84) | (1.20 – 1.84) |
|                  | = 25 or not recorded | 1007 | 1.49  | 1.49  | (1.20 – 1.84) | (1.20 – 1.84) |
| Organ system failure: respiratory | No   | 335   | 1.09  | 1.09  | (0.87 – 1.36) | (0.87 – 1.36) |
|                  | Yes  | 1058  | 1.09  | 1.09  | (0.87 – 1.36) | (0.87 – 1.36) |

#### Organ system failure: cardiovascular
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Table 5 (Continued)

Multiple logistic regression model of patient factors in relation to ultimate hospital mortality

| No   | 881  | 2594 (34.0) | Ref. | Ref. |
|------|------|-------------|------|------|
| Yes  | 512  | 1017 (50.3) | 2.21 (1.87 – 2.62) | 1.48 (1.22 – 1.79) |

Organ system failure: neurological

| No   | 1066 | 2978 (35.8) | Ref. | Ref. |
|------|------|-------------|------|------|
| Yes  | 327  | 633 (51.7)  | 1.85 (1.53 – 2.24) | 1.36 (1.10 – 1.68) |

Organ system failure: renal

| No   | 1190 | 3346 (35.6) | Ref. | Ref. |
|------|------|-------------|------|------|
| Yes  | 203  | 265 (76.6)  | 5.17 (3.70 – 7.23) | 3.83 (2.67 – 5.47) |

Organ system failure: haematological

| No   | 1362 | 3550 (38.4) | Ref. | Ref. |
|------|------|-------------|------|------|
| Yes  | 31   | 61 (50.8)   | 1.65 (0.94 – 2.89) | 1.30 (0.69 – 2.45) |

Ultimate hospital mortality excludes admissions who were readmissions to the intensive care unit within the same hospital stay, admissions whose ultimate hospital discharge status was missing, and admissions for whom any of the entered risk factors were missing. *During first 24 hours following unit admission. ABG, arterial blood gas; CI, confidence interval; CPR, cardiopulmonary resuscitation; FiO2, fractional inspired oxygen; OR, odds ratio; PaCO2, arterial carbon dioxide tension; PaO2, arterial oxygen tension.

Non-surgical COPD patients account for 2.9% of all ICU admissions in the UK

Intubated patients have a median stay in intensive care of 6 days

COPD patients admitted to ICU have a mortality of 23% in the admitting ICU and 38% in hospital, which is similar to the mortality of patients with all diagnoses admitted to ICU

Key messages

- Non-surgical COPD patients account for 2.9% of all ICU admissions in the UK
- Intubated patients have a median stay in intensive care of 6 days
- COPD patients admitted to ICU have a mortality of 23% in the admitting ICU and 38% in hospital, which is similar to the mortality of patients with all diagnoses admitted to ICU

Competing interests

The author(s) declare that they have no competing interests.

Authors’ contributions

MW drafted the manuscript. DH performed the analyses and contributed to drafting the manuscript. All authors participated in the design and interpretation of the study, and critical revision of the manuscript, and read and approved the final manuscript.

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Contribution of COPD patients to the ultimate hospital mortality

| Organ system failure | No | Yes | OR (95% CI) |
|----------------------|----|-----|------------|
| Neurological         | 1066| 327 | 1.85 (1.53 – 2.24) |
| Renal                | 1190| 203 | 5.17 (3.70 – 7.23) |
| Haematological       | 1362| 31  | 1.65 (0.94 – 2.89) |

When patients with obstructive lung disease present with decompensated type II respiratory failure, it can be very difficult to distinguish those with ‘pure’ COPD (without any significant reversibility) from those with a mixture of COPD and asthma (patients with a major component of fixed obstruction that coexists with some reversibility). It is likely that patients with greater reversibility will tend to be younger, possibly respond better to steroids and have a better overall prognosis. Unfortunately, the CMP data do not allow this distinction to be explored, and there is no real agreement about how this distinction should be made even when a patient has had full lung function – a distinction that is even harder in the acute setting when information regarding lung function is rarely available. Additional prospective research may be required to further illuminate this problem.
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