

# Assisted ventilation in COPD – association between previous hospitalizations and mortality

Anne Pernille Toft-Petersen<sup>1,2</sup>

Christian Torp-Pedersen<sup>1,3</sup>

Ulla Møller Weinreich<sup>1,4</sup>

Bodil Steen Rasmussen<sup>1,2</sup>

<sup>1</sup>Department of Clinical Medicine, Aalborg University, <sup>2</sup>Department of Anaesthesia and Intensive Care, Aalborg University Hospital, <sup>3</sup>Department of Health, Science and Technology, Aalborg University, Aalborg, <sup>4</sup>Department of Respiratory Medicine, Aalborg University Hospital, Denmark

**Background:** In general, previous studies have shown an association between prior exacerbations and mortality in COPD, but this association has not been demonstrated in the subpopulation of patients in need of assisted ventilation. We examined whether previous hospitalizations were independently associated with mortality among patients with COPD ventilated for the first time.

**Patients and methods:** In the Danish National Patient Registry, we established a cohort of patients with COPD ventilated for the first time from 2003 to 2011 and previously medicated for obstructive airway diseases. We assessed the number of hospitalizations for COPD in the preceding year, age, sex, comorbidity, mode of ventilation, survival to discharge, and days to death beyond discharge.

**Results:** The cohort consisted of 6,656 patients of whom 66% had not been hospitalized for COPD in the previous year, 18% once, 8% twice, and 9% thrice or more. In-hospital mortality was 45%, and of the patients alive at discharge, 11% died within a month and 39% within a year. In multivariate models, adjusted for age, sex, mode of ventilation, and comorbidity, odds ratios for in-hospital death were 1.26 (95% confidence interval [CI]: 1.11–1.44), 1.43 (95% CI: 1.19–1.72), and 1.56 (95% CI: 1.30–1.87) with one, two, and three or more hospitalizations, respectively. Hazard ratios for death after discharge from hospital were 1.32 (95% CI: 1.19–1.46), 1.76 (95% CI: 1.52–2.02), and 2.07 (95% CI: 1.80–2.38) with one, two, and three or more hospitalizations, respectively.

**Conclusion:** Preceding hospitalizations for COPD are associated with in-hospital mortality and after discharge in the subpopulation of patients with COPD with acute exacerbation treated with assisted ventilation for the first time.

**Keywords:** pulmonary disease, chronic obstructive, respiration, artificial, patient readmission, hospital mortality, critical care

## Introduction

COPD is a persistent, progressive airflow limitation associated with a chronic airway inflammation and is among the leading causes of mortality and morbidity worldwide, in fact predicted to be the third leading cause of death by 2020.<sup>1</sup>

An acute exacerbation of COPD (AECOPD), defined as worsening of symptoms beyond normal day-to-day variation, is associated with increased mortality.<sup>1–3</sup> In general, exacerbations are more frequent in severe COPD,<sup>4,5</sup> but recent studies show that the number of exacerbations varies extensively among patients with COPD with similar lung function in stable phases. Nevertheless, exacerbations are the best predictor of their own recurrence.<sup>3,4,6</sup> It has been suggested that an inherent “frequent-exacerbation phenotype” with a distinct pathophysiology might explain at least parts of the extensive clinical variability.<sup>4,7</sup> Previous accumulated exacerbations have been shown to predict

Correspondence: Anne Pernille Toft-Petersen  
Department of Clinical Medicine, Aalborg Universitetshospital, Sdr. Skovvej 15, 9000 Aalborg, Denmark  
Tel +45 97 66 18 64  
Fax +45 97 66 19 02  
Email aptp@rn.dk



not only new exacerbations but also mortality. This prediction is independent of stable phase lung function impairment, and mortality increases by each subsequent exacerbation.<sup>5,8–11</sup> It is not known whether the association between previous exacerbations and mortality is independent of the degree of acute illness. Patients subjected to assisted ventilation constitute a group whose acute illness is severe, and the association between previous exacerbations and mortality in this group has not been elucidated.

The overall short-term mortality among patients with acute exacerbations ranges from 2.1% to 20.4%,<sup>9,12,13</sup> but patients with severe acute COPD admitted to intensive care unit (ICU) are particularly at risk with a 6-month mortality ranging from 25% to 40%.<sup>12,14,15</sup> In-hospital mortality rates for patients being ventilated are 5%–25% but higher in relation to invasive mechanical ventilation (IMV) than to noninvasive mechanical ventilation (NIV).<sup>16–20</sup> Predictors of mortality in hospitalized patients with COPD<sup>12,21,22</sup> are age, comorbidities, and end-stage COPD with concomitant organ dysfunctions. Lung function impairment in stable phase is in itself a poor predictor of short-term mortality following exacerbations<sup>8,23</sup> but is associated with long-term mortality.<sup>24</sup> On the other hand, acidosis, which mirrors acute derangement, predicts short-term but not long-term mortality.<sup>12</sup> In patients admitted to ICU, acute derangement is of major importance regarding both short-term<sup>25,26</sup> and intermediate-term survival,<sup>15</sup> but long-term survival studies have been conflicting with regard to independent risk predictors.<sup>14,27–31</sup>

The aim of this study was to investigate whether the frequency of previous hospitalizations for AECOPD can predict in-hospital mortality and mortality after discharge in an inception cohort of patients with COPD treated with assisted ventilation for the first time. We hypothesized that frequent COPD hospitalizations were associated with excess mortality following assisted ventilation.

## Patients and methods

All hospitalizations from 2002 to 2011 were identified in the Danish National Patient Registry. Successive hospitalizations were merged if the discharge date of the first and the admission date of the second were identical. AECOPD hospitalizations were defined as hospitalizations with either a primary diagnosis of COPD (*International Classification of Diseases*, tenth edition [ICD-10]: DJ44) or a complex of either acute respiratory failure (ICD-10: DJ96) or pneumonia (ICD-10: DJ13–18) as the primary diagnosis and COPD as a secondary diagnosis as described by Thomsen et al.<sup>32</sup> In general, hospitalizations related to diseases of the respiratory system (respiratory

hospitalizations) were identified by a primary diagnosis within the entire ICD-10 DJ-spectrum. No attempts were made to distinguish between admissions for exacerbations with and without pneumonia. Patients <30 years of age were excluded to minimize inclusion of misclassified asthma. First time ventilation was defined as the first AECOPD hospitalization where assisted ventilation was performed, but first time ventilations in 2002 were discarded as no specific procedure code for NIV was accessible till 2003. In the National Register of Medicinal Product Statistics, all prescriptions in the year preceding first time hospitalizations were identified. By ATC code “R03”, drugs for obstructive airway diseases were identified. First time ventilation was classified as the index hospitalization if a prescription was redeemed within 1 year before. Any patient with an index hospitalization was included in the cohort (Figure 1). Mode of ventilation at the

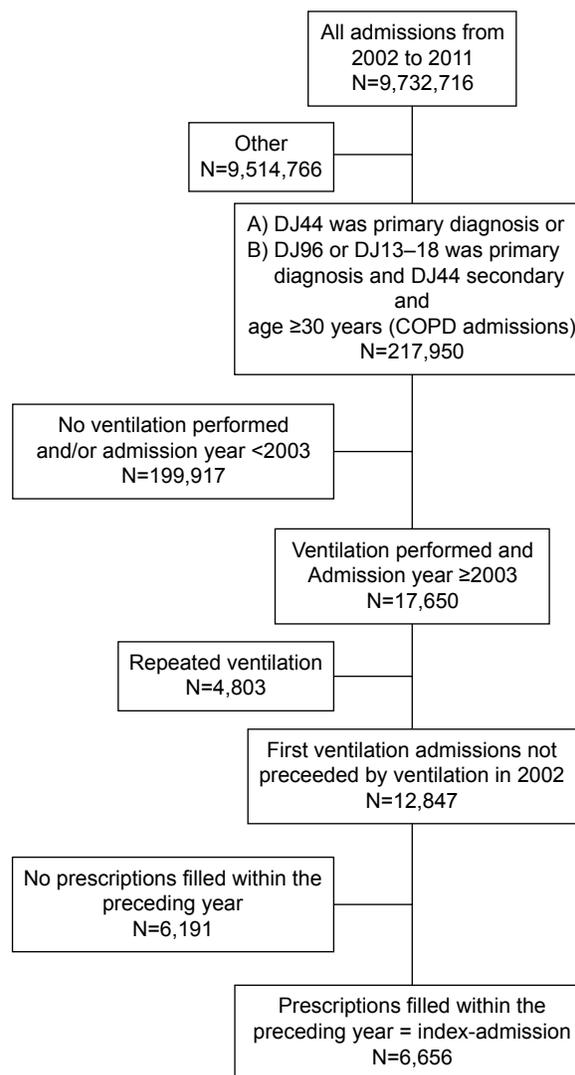


Figure 1 The inclusion of patients.

index hospitalization was characterized as either “IMV” (BGDA0) or “NIV” (BGDA1). The categories were mutually exclusive and hierarchically assigned so that IMV overruled NIV, if both were registered.

The number of AECOPD and respiratory system-related hospitalizations in the year preceding the index hospitalization was counted. Charlson comorbidity score based on diagnoses made during hospitalizations and outpatient visits 10 years prior to the discharge from the index hospitalization were assigned to individual diseases ad modum Radovanovic et al<sup>33</sup> and grouped in three with 1, 2, or 3+ points, respectively, as per definition all patients received one point for having COPD.

Information on day of death was obtained from the Danish Civil Registration System. Each patient was followed until the time of death or censored by December 31, 2013. Informed consent was not required according to Danish legislation.

General characteristics between groups were compared by chi-square test (character variables) and Kruskal–Wallis test (continuous variable). For analysis of in-hospital mortality, logistic regression was used, and for analysis of mortality beyond discharge, Cox proportional hazards models were used. In both models, continuous variables were tested for linearity, and in the Cox model, dependency of the estimates on time was evaluated by the Schoenfeld residuals. Interactions a priori were considered relevant, that is, interactions between previous hospitalizations, and age, sex, and mode

of ventilation, respectively, were tested for by comparisons of likelihood. *P*-values <0.05 were considered significant. Statistical analyses were performed in SAS 9.4© (SAS Institute Inc., Cary, NC, USA) and R© Version 3.1.1.

## Results

The population consisted of 6,656 patients (Table 1). Of these, 65% had not been hospitalized for AECOPD in the preceding year, 18% had been hospitalized once, 8% twice, and 9% thrice or more. The proportion excluded was 50% in the group of patients not hospitalized in the previous year and 44%, 41%, and 45% in the group hospitalized once, twice, and thrice, respectively. Mortality rates are shown in Tables 2 and 3. The annual number of first time ventilations for COPD exacerbation increased from 355 in 2003 to 1,236 in 2011.

Increasing age, high Charlson comorbidity score, and IMV were the risk factors for in-hospital mortality. The risk decreased during the study period. For all patients, in-hospital mortality was associated with the number of previous AECOPD hospitalizations. The odds ratios, compared to no previous hospitalizations, were 1.26 (95% confidence interval [CI]: 1.11–1.44) with one hospitalization, 1.43 (95% CI: 1.19–1.72) with two hospitalizations, and 1.56 (95% CI: 1.30–1.87) with three or more hospitalizations (Figure 2). The risk associated with three or more hospitalizations was significantly higher than the risk associated with one hospitalization (*P*=0.04).

**Table 1** Characteristics of included patients at baseline by number of COPD hospitalizations in the previous year

Characteristics	Number of admissions in the previous year				Total n=6,656
	0 n=4,333	1 n=1,213	2 n=538	3+ n=572	
Sex					
Male	2,481 (57.3%)	684 (56.4%)	305 (56.7%)	320 (55.9%)	3,790 (56.9%)
Age, year					
Median (IQR)	74.1 (66.4–79.8)	74.8 (67.5–80.6)	73.9 (67.1–80.0)	72.9 (65.1–77.9)	74.1 (66.5–79.8)
Charlson comorbidity score					
1	1,691 (39.0%)	429 (35.4%)	183 (34.0%)	216 (37.8%)	2,519 (37.8%)
2	1,051 (24.3%)	278 (22.9%)	131 (24.3%)	150 (26.2%)	1,610 (24.2%)
3+	1,591 (36.7%)	506 (41.7%)	224 (41.6%)	206 (36.0%)	2,527 (38.0%)
Time period					
2003–2005	856 (19.8%)	247 (20.4%)	120 (22.3%)	133 (23.3%)	1,356 (20.4%)
2006–2008	1,410 (32.5%)	412 (34.0%)	181 (33.6%)	191 (33.4%)	2,194 (33.0%)
2009–2011	2,067 (47.7%)	554 (45.7%)	237 (44.1%)	248 (43.4%)	3,106 (46.7%)
Mode of ventilation					
NIV	2,316 (53.5%)	726 (59.9%)	346 (64.3%)	389 (68.0%)	3,777 (56.7%)
Length of stay					
1–2 days	406 (9.4%)	126 (10.4%)	81 (15.1%)	64 (11.2%)	677 (10.2%)
3–7 days	1,023 (23.6%)	333 (27.5%)	157 (29.2%)	178 (31.1%)	1,691 (25.4%)
7+ days	2,904 (67.0%)	754 (62.2%)	300 (55.8%)	330 (57.7%)	4,288 (64.4%)

**Abbreviations:** IQR, interquartile range; NIV, noninvasive mechanical ventilation.

**Table 2** In-hospital mortality by number of COPD hospitalizations in the previous year

	Number of admissions in the previous year				Total n=6,656
	0 n=4,333	1 n=1,213	2 n=538	3+ n=572	
Mortality, n (%)					
Death before discharge	1,832 (42%)	585 (48%)	270 (50%)	286 (50%)	2,973 (45%)

Mortality after discharge from hospital was associated with female sex, increasing age, Charlson comorbidity score, and NIV. The risk decreased during the study period. For all patients, the number of previous AECOPD hospitalizations was associated with an increased mortality after discharge compared to no previous hospitalizations with a hazard ratio of 1.32 (95% CI: 1.19–1.46) with one hospitalization, 1.76 (95% CI: 1.52–2.02) with two hospitalizations, and 2.07 (95% CI: 1.80–2.38) with three or more hospitalizations (Figure 3). The risks associated with two and with three or more hospitalizations were significantly higher than the risk associated with one hospitalization ( $P<0.001$  and  $P<0.001$ , respectively).

The associations with both in-hospital and after-discharge mortality showed the same trends, though weaker, when, instead of AECOPD hospitalizations, respiratory hospitalizations were included in the model (Figure 4).

## Discussion

In this study of patients ventilated for COPD for the first time, an increasing number of hospitalizations in the preceding year were associated with increased overall mortality. We found significant odds ratios for in-hospital death ranging from 1.3 to 1.6 with increasing number of hospitalizations with COPD exacerbations within the previous year. To our knowledge, this association has not been studied in this subgroup of patients. A number of studies have examined the association between previous exacerbations and mortality in other COPD subpopulations, and in some of these, the association was far stronger than what we found.

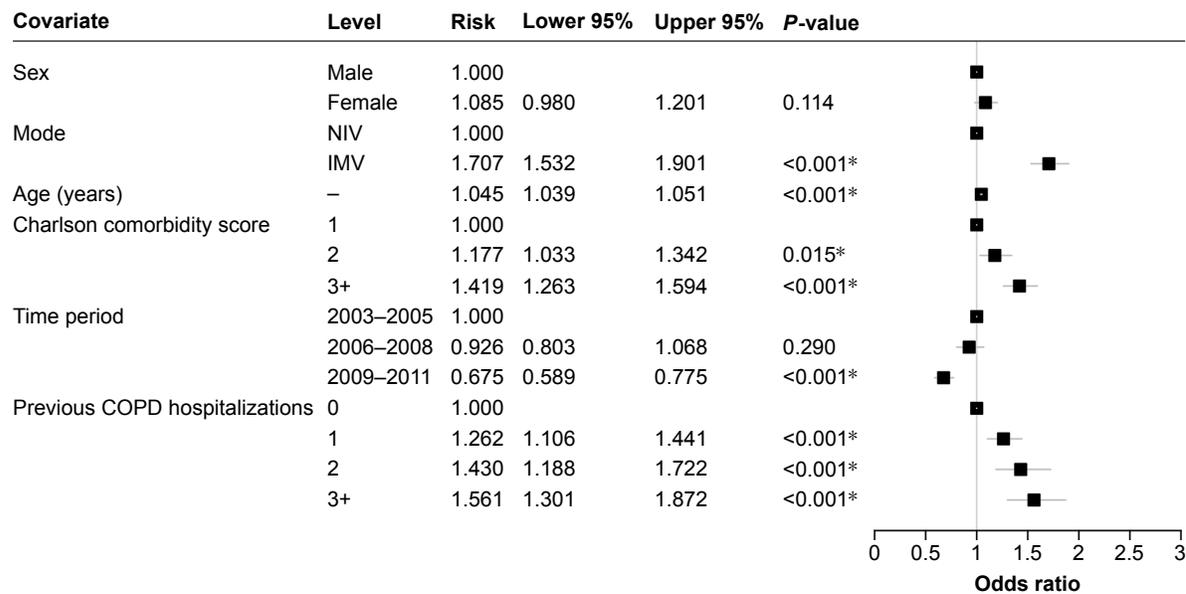
Hospital contact for exacerbation within the previous year was associated with adverse events within a month, odds ratio 3.9 (95% CI: 1.6–9.9).<sup>22</sup> Likewise, having had at least two

hospitalizations within the previous 2 years was associated with 30 days mortality, odds ratio 3.6 (95% CI: 2.5–5.2).<sup>34</sup> Among ICU admitted patients with COPD, two or more exacerbations within the previous year were associated with in-hospital mortality, odds ratio 9.1 (95% CI: 1.02–81.1).<sup>11</sup> Apart from different outcome measurements, the discrepancy in the magnitude of the correlations might be explained by different case mixes. Patients with end-stage COPD, which presumably have more frequent exacerbations, may to some extent have been excluded from our cohort due to advancement orders that ruled out ventilation. Whether the stage of stable phase COPD modifies the correlation between the number of hospitalizations and short-term mortality needs to be further evaluated.

Among patients with COPD in Denmark, Schmidt et al<sup>35</sup> found significant association between neither previous overall exacerbations nor previous hospitalized exacerbations and 30-day mortality after hospitalization for exacerbation upon adjustment for age, sex, comorbidities, and previous medication. Patients in this cohort, in contrast to those included in ours, were dissimilar with regard to the acute severity of the exacerbation due to a case mix of ventilated and nonventilated patients. Moreover, first time and repeated hospitalizations were evaluated together, and redeemed prescriptions for obstructive airway diseases medication were not a criterion for inclusion. Another study of 3-month mortality showed that previous hospitalizations were not a significant predictor of mortality upon inclusion of markers of acute derangement and performance status.<sup>36</sup> It is possible, though, that the decision of admitting a patient with any given degree of acute illness depends on performance status, thus masking an association between previous hospitalizations and mortality.

**Table 3** Mortality after discharge by number of COPD hospitalizations in the previous year

	Number of admissions in the previous year				Total n=3,683
	0 n=2,501	1 n=628	2 n=268	3+ n=286	
Mortality					
Death within 30 days	236 (9%)	99 (16%)	41 (15%)	44 (15%)	420 (11%)
Death within 1 year	827 (33%)	296 (47%)	158 (59%)	169 (59%)	1,450 (39%)



**Figure 2** Logistic regression model of in-hospital mortality.

**Note:** \*Significant at  $P < 0.05$ .

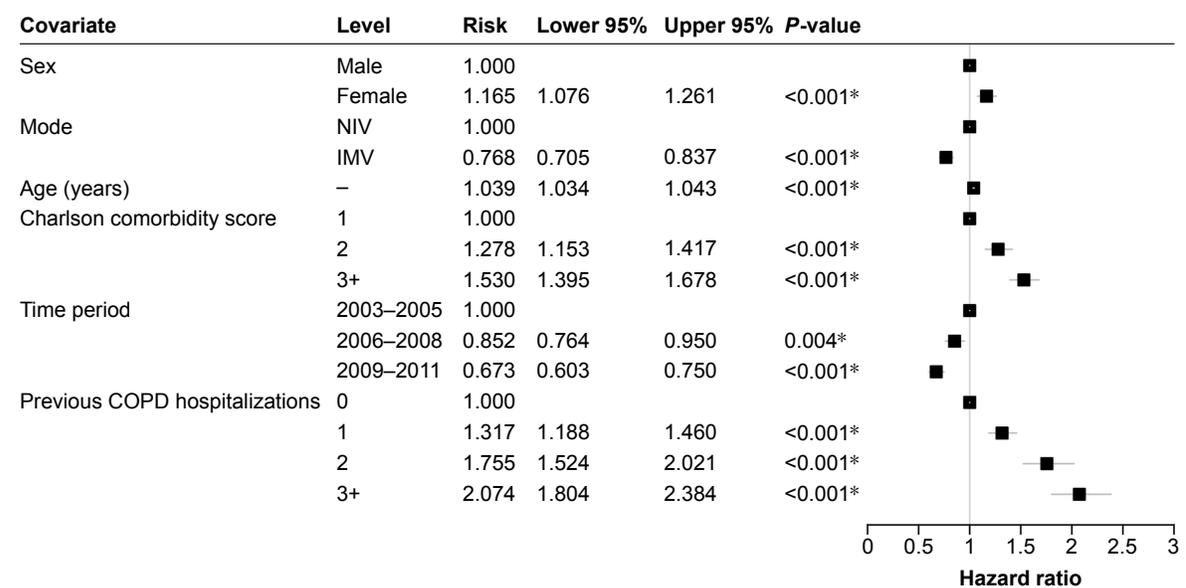
**Abbreviations:** NIV, noninvasive mechanical ventilation; IMV, invasive mechanical ventilation.

In our study population, mortality after discharge was associated with previous exacerbations with hazard ratios in the range of 1.3–2.0 for one and three or more hospitalizations, respectively. The aforementioned Danish study<sup>35</sup> also found a significant association between long-term mortality and the number of COPD exacerbations. The strength of the association is comparable to what we found and to what was found in a Spanish study.<sup>37</sup>

A large American study<sup>38</sup> found a lower hazard ratio after discharge of 1.14 by each previous hospitalization in a 1-year

period. However, the inclusion criteria as well as the case mix were profoundly different from ours, and in addition, immortal time bias might partly account for the estimate being lower than ours. No independent association was found in another study by Almagro et al,<sup>39</sup> but stable phase lung function impairment correlated strongly with prognosis and might have masked an association.

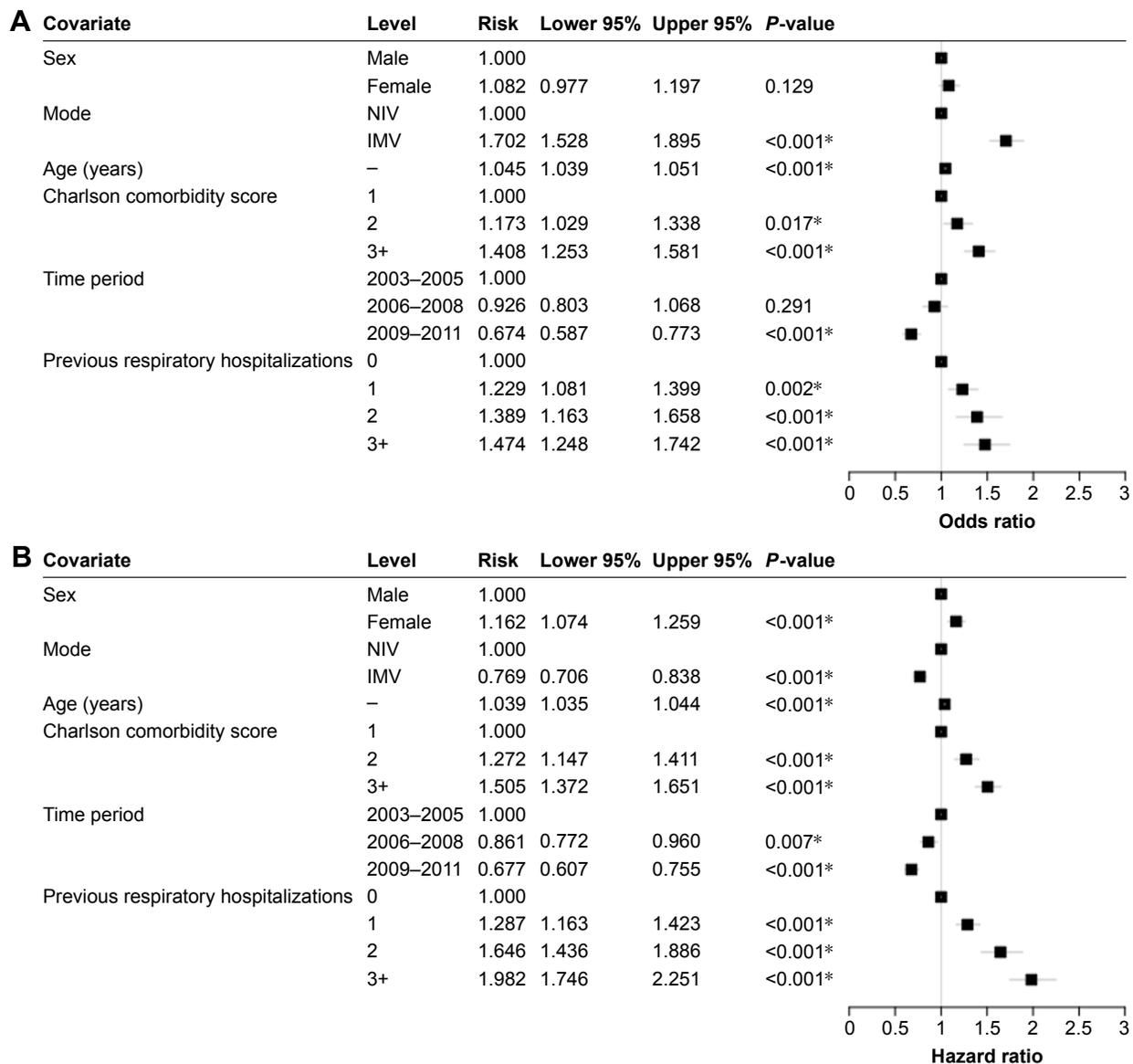
Our study confirmed the importance of age and comorbidity, while sex was only associated with mortality after discharge. Mortality risk decreased with time. IMV as



**Figure 3** Cox proportional hazards model of mortality after discharge.

**Note:** \*indicates significant at  $P < 0.05$ .

**Abbreviations:** NIV, noninvasive mechanical ventilation; IMV, invasive mechanical ventilation.



**Figure 4** Respiratory hospitalizations.

**Notes:** (A) Logistic regression model of in-hospital mortality, (B) Cox proportional hazards model of mortality after discharge. \*indicates significant at  $P < 0.05$ .

**Abbreviations:** NIV, noninvasive mechanical ventilation; IMV, invasive mechanical ventilation.

a mode of ventilation predicted death before discharge independently, whereas NIV predicted time to death after discharge. IMV is mainly initiated in patients with severe acute derangement, which probably accounts for the associated higher in-hospital mortality compared to NIV. That this relation is reversed after discharge can have several explanations, among those that patients discharged alive after IMV treatment had undergone harder selection.

In comparison with other studies, our estimate of in-hospital mortality of ventilated patients is high,<sup>16,31,40</sup> but this might partly be explained by inclusion in our study of patients with COPD with pneumonia, which confers a large additional risk.<sup>41</sup>

The association between treatment year and risk might be explained by increasingly aggressive strategies for stable phase medication and for ventilation during exacerbations. Assuming that both medication and ventilation have mainly gained a stronger foothold among less afflicted patients with COPD, gradual rarefication of mortality by the inherently favorable prognosis of this group can explain the improvement of prognosis.<sup>42</sup>

The strengths of our study are the large number of patients and the high reliability of the Danish medical registers, which is well established for a number of diagnoses,<sup>43</sup> among those the COPD diagnosis applied here (a positive predictive value of 92%),<sup>32</sup> procedures,<sup>44</sup> and vital status.<sup>45</sup> In addition,

the inherent ability to track unique patient courses virtually eliminates all but emigration as loss to follow-up.

Our study has noteworthy limitations. Misclassification is in our data set especially problematic with regard to the calculated comorbidity, where completeness of registration will be highest among frequently hospitalized patients. However, the use of Charlson comorbidity score has been shown to yield acceptable precision when calculated from registers,<sup>46</sup> although the sensitivity of secondary diagnoses in the National Patient Registry is known to be low.<sup>47</sup> We speculated whether the number of previous hospitalizations decided by our method could underestimate the number of hospitalizations as the propensity to register a COPD diagnosis in connection with a hospitalization might depend on the COPD stage. Such misclassification might even be differential as only previously admitted patients could be misclassified. To test this, we substituted respiratory hospitalizations for COPD hospitalizations, which did not materially change the results.

We chose to include only patients with COPD who had redeemed prescriptions for obstructive airway diseases medication within the year preceding ventilation. This excluded 49% of all patients whose hospitalizations would have qualified as index hospitalizations. This might reflect not only low compliance but also the inclusion of patients with breathing difficulties whose history was suggestive of COPD but who did not have symptoms on a daily basis. It is surprising and concerning that the proportion of patients who had not redeemed medication was almost equal in patients who had no previous admissions and in patients who had been repeatedly hospitalized. Exclusion in this case is a trade-off between generalizability and validity. It is generally assumed that previous medication acts as a proxy for previously recognized COPD, and thereby to some extent as a marker of stable phase disease severity. Previously, it has been shown that majority of the COPD population are unaware of their disease<sup>48</sup> and that a considerable proportion of even patients with advanced disease (25%) do not receive appropriate treatment.<sup>49</sup> Even so, in general, the unrecognized cases have less advanced disease, are hospitalized less frequently after the hospitalization where the diagnosis was established,<sup>50</sup> and have less severe symptoms prior to exacerbations.<sup>50</sup>

Other limitations to our study are the lack of information on lung function prior to the hospitalization, the degree of acute physiological derangement, and the extent of limitations in care. Differences in underlying acute and chronic lung function impairment might confound the association

between previous hospitalizations and mortality. However, we do consider this to be partly ameliorated by the inclusion of only such patients as had been offered ventilation upon clinical decision. Wildman et al<sup>51</sup> have shown that clinicians are overly pessimistic when predicting the survival of patients with COPD on hospitalization to specialized respiratory units or ICUs, so it must be assumed that the instigation of assisted ventilation, at least invasive ventilation, among patients with very severe lung function impairment is limited. As for variation in severity of present respiratory failure, any subject hospitalized to our selected study population had an overall acute derangement warranting assisted ventilation, and it must be assumed that the gas exchange was considerably impaired in most of our subjects. In pace with the dissemination of NIV as an accessible treatment option, NIV has to some extent become the interventional limit for a group of frail, elderly patients with often advanced COPD.<sup>19</sup> We did not have access to information on do-not-intubate-orders, but a previous Danish study found such orders in 10% of NIV patients.<sup>52</sup> Presumably, a subpopulation of patients with short expected life span diluted the survival in the NIV group.

## Conclusion

In conclusion, among the patients with COPD with acute exacerbation treated with assisted ventilation for the first time, the number of hospitalizations for COPD in the preceding year is associated with increased mortality in-hospital as well as after discharge. Our study did not aim at identifying patients in whom assisted ventilation is futile. Nonetheless, it emphasizes the importance of the clinical trajectory for the outcome following ventilation in a group of patients whose mortality is very high.

## Author contributions

APTP had full access to all of the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis. CTP, UMW, and BSR contributed substantially to the study design, data analysis and interpretation, and the writing of the manuscript. All authors contributed toward data analysis, drafting and critically revising the paper and agree to be accountable for all aspects of the work.

## Disclosure

The authors report no conflicts of interest in this work.

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