The impact of chronic obstructive pulmonary disease on intensive care unit admission and 30-day mortality in patients undergoing colorectal cancer surgery: a Danish population-based cohort study

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ABSTRACT

Background and purpose: Chronic obstructive pulmonary disease (COPD) may increase the risk of postoperative complications and thus mortality after colorectal cancer (CRC) surgery, but the evidence is sparse.

Methods: We conducted this nationwide population-based cohort study in Denmark, including all patients undergoing CRC surgery in the period 2005–2011, identified through medical databases. We categorised the patients according to the history of COPD. We assessed the rate of complications within 30 days. We computed 30-day mortality among patients with/without COPD using the Kaplan-Meier method. We used Cox regression to compute HRs for death, controlling for age, gender, type of admission, cancer stage, hospital volume, alcohol-related diseases, obesity and Charlson comorbidity score.

Results: We identified 18 302 CRC surgery patients. Of these, 7.9% had a prior diagnosis of COPD. Among patients with COPD, 16.1% were admitted postoperatively to the intensive care unit, 1.9% were treated with mechanical ventilation, and 3.6% were treated with non-invasive ventilation. In patients without COPD, the corresponding proportions were 9.7%, 1.1% and 1.1%. The reoperation rate was 10.6% among patients with COPD and 8% among patients without COPD. 30-day mortality was 13% (95% CI 11.4% to 14.9%) among patients with COPD and 5.3% (95% CI 5.0% to 5.7%) among patients without COPD, corresponding to an adjusted HR of 1.7 (95% CI 1.5 to 2.0).

Conclusions: COPD is a strong predictor for intensive care unit admission and mortality after CRC surgery.

INTRODUCTION

Colorectal cancer (CRC) is a common disease, with a first-time diagnosis occurring at an average age of approximately 70 years.1 2 Surgical resection of the tumour has the potential to cure early-stage CRC, either alone or combined with chemotherapy and/or radiation.3 Overall, 30-day mortality after CRC resection has been estimated to be approximately 10%,4 with higher rates among elderly patients compared to younger patients.5

Many patients with CRC have concurrent chronic diseases, including chronic obstructive pulmonary disease (COPD). Evidence suggests that COPD is associated with substantially increased mortality after various types of surgery.4 6–10 Much of this evidence originates from studies of patients undergoing surgery for perforated peptic ulcers, cardiac procedures, hip fractures and abdominal aortic aneurysm repair.6–8 11 The studies have mainly included COPD as a covariate in analyses for other purposes like hospital variation in mortality4 have combined several types of cancer or lung diseases9 12 or focused only on men.10 However, evidence of an association between COPD and postoperative outcome after CRC surgery is limited.4 9 10 13

A primary reason for increased postoperative mortality among patients with COPD may be respiratory failure, leading to more frequent and longer intensive care unit (ICU) admissions.7 11 In addition, patients...
with a high comorbidity burden have in general been found to be at increased risk of surgical complications, leading in turn to a need for reoperation, but none of the former studies have examined the risk of post-operative admission to the ICU among patients undergoing CRC surgery.

We conducted a nationwide population-based cohort study to investigate the impact of COPD on ICU admissions, ICU treatments (mechanical ventilation, non-invasive ventilation (NIV), dialysis and treatment with inotropes/vasopressors), occurrence of reoperation and 30-day mortality following CRC surgery.

METHODS

Settings

This cohort study was conducted in Denmark, a country with 5.5 million inhabitants.

All Danish citizens have equal access to tax-funded medical care, and all CRC surgery is performed in public hospitals.

Since 1968, everyone living in Denmark has been assigned a personal civil registration number (CPR number) by the Danish Civil Registration System (CRS). The CRS is updated daily and contains information on gender, date of birth and vital status. We used the CPR number to link registries at the individual level.

Patients

We used the Danish Cancer Registry (DCR) to identify all patients with CRC diagnosed in the period from 1 January 2005 until 31 December 2011. This period was chosen because ICU admissions have been recorded since 2005. The DCR, established in 1943, covers all incident cancer cases in Denmark through notification from hospital departments, specialists and autopsy reports.

DCR data include CPR number, month and year of cancer diagnosis and tumour spread at the time of diagnosis. We classified CRC stages as localised (Dukes’ A and B), regional (Dukes’ C), metastasised (Dukes’ D) or unknown (see e-appendix 1).

We linked the patients with CRC to the Danish National Registry of Patients (DNRP), in order to identify date of CRC surgery. The DNRP, a nationwide registry established in 1977, contains records on 99.4% of all discharges from Danish hospitals. Emergency room and outpatient clinic visits have been included since 1995 and ICU admissions, including information on use of mechanical ventilation and other ICU treatments, since 2005.

The DNRP includes data on CPR number, hospital, department, discharge diagnoses and surgical and diagnostic procedures. Diagnoses were coded according to the International Classification of Diseases, 8th revision (ICD-8) until 1993 and 10th revision (ICD-10) thereafter. Our study cohort consisted of patients who underwent CRC surgery (see e-appendix 1 for surgical procedure codes). We categorised CRC surgery as open surgery, laparoscopic resection or other surgery; in addition, we investigated colon and rectal cancer surgery separately.

We also categorised type of admission as acute or elective because acute surgery has been shown to be associated with increased mortality. Furthermore, we divided hospitals according to volume: low volume (less than 500 CRC surgeries in the study period) and high volume (more than 500 CRC surgeries in the study period).

Patients with COPD

Within the cohort of CRC surgery patients, we searched the DNRP for any inpatient COPD diagnosis since 1977. In addition, we identified and included patients diagnosed with COPD in outpatient specialist clinics and emergency rooms since 1995 (ICD codes are provided in e-appendix 1).

Main outcome measures

Admission to the ICU, ICU treatments, reoperation and death

We obtained data from the DNRP on ICU admissions, use of mechanical ventilation, treatment with NIV, dialysis, inotropic treatment and reoperations within 30 days following surgery as a marker of surgical complications. To avoid including prophylactic ICU admissions as a marker of surgical complications, we only included ICU admission occurring at least 1 day after surgery. In addition, we divided the reason for reoperation according to the procedure code into five groups: infection, bleeding, wound rupture, anastomosis leakage and other reasons (procedure codes are provided in e-appendix 1). To investigate 30-day mortality, CRS data were used to follow patients from date of CRC surgery until death, emigration, 30 days postoperatively or the end of the study period, whichever came first.

Comorbidities

We used Charlson Comorbidity Index (CCI) scores to assess comorbidity levels in the study cohort. The CCI score is computed as the sum of points (between 1 and 6) assigned to each of the 19 diseases included in the index (see e-appendix 1). Information on comorbid diseases was obtained from the DNRP, including all inpatient and outpatient diagnoses within 5 years prior to date of CRC surgery. Patients were classified into three groups according to their CCI score: 0 points (‘low comorbidity level’); 1–2 points (‘moderate comorbidity level’); and 3 or more points (‘severe comorbidity level’). We excluded COPD and CRC from the CCI as they represent the exposure and the patient population under study. In addition to the CCI score, our analyses included data on alcohol-related diseases and obesity from the DNRP (see e-appendix 1).

Statistical analyses

We calculated frequencies of covariates (including categories of demographic characteristics, CCI scores, types of admission (elective or acute), types of surgery, hospital volume and cancer stage) according to the presence or absence of COPD. We computed the rates of patients...
with CRC who were admitted to the ICU 1–30 days after surgery, and received mechanical ventilation, NIV, dialysis, inotropes/vasopressors or underwent reoperation within 30 days after surgery. In a sensitivity analysis, we also computed the rate of ICU admission 0–30 days after surgery. We used Jeffrey’s method to compute 95% CIs for the rates.21 We estimated and plotted 30-day mortality for the rates.21 We estimated and plotted 30-day mortality

**RESULTS**

**Descriptive data**

We identified 18 302 CRC surgery patients, of whom 1449 (7.9%) had a COPD diagnosis. The proportion of patients with COPD was slightly higher among patients with colon cancer (8.3% of 12 053 patients with colon cancer) compared to patients with rectal cancer (7.1% of 6249 patients with rectal cancer).

Patients with CRC with COPD were older and had a higher comorbidity level than patients with CRC without COPD. Gender, admission type (elective/acute), surgery type and CRC stage at time of diagnosis were very similar in patients with and without COPD (table 1).

**ICU admission, ICU treatments and reoperation within 30 days after surgery**

Among patients with CRC with COPD, 16.1% (95% CI 14.3% to 18.1%) were admitted to the ICU 1–30 days after surgery, compared with 9.7% (95% CI 9.3% to 10.2%) of patients without COPD (table 2). In the sensitivity analysis including days 0–30 after surgery, we found an ICU admission rate of 29.0% (95% CI 26.7% to 31.4%) among patients with COPD and 17.6% (95% CI 17.0% to 18.2%) among patients without COPD (e-appendix 2).

Patients with CRC with COPD also received mechanical ventilation treatment more frequently (1.9%; 95% CI 1.3% to 2.8%) than patients without COPD (1.1%; 95% CI 1.0% to 1.3%) and more frequently received treatment with NIV (3.6%; 95% CI 2.7% to 4.7% vs 1.1%; 95% CI 0.9% to 1.2%; table 2).

More patients with CRC with COPD than those without COPD underwent reoperation (10.6% (95% CI 9.0% to 12.3%) vs 8.0% (95% CI 7.6% to 8.4%), particularly due to wound ruptures and bleeding (tables 2 and 3). Patients with COPD more frequently received inotropes/vasopressors than patients without COPD (table 2). In addition, ICU admissions, ICU treatments and reoperations were generally more frequent in the acute than in the elective setting, both among patients with CRC with and without COPD (e-appendix 2).

The results were similar when colon cancer and rectal cancer were considered separately (table 2).

In an analysis by year of CRC surgery, we found that the frequency of ICU admission declined for CRC surgery overall from 41.6% in 2005 to 25.8% in 2011 (Results in e-appendix 2). Among patients with CRC without COPD, 24% were admitted to the ICU in 2005 and 12.8% were admitted in 2011. Thus, while the frequency of ICU admission declined over time among patients with CRC overall, the frequency continued to be highest among patients with CRC with COPD.

In contrast, we found that the rate of patients treated with mechanical ventilation increased from 1.2% in 2005 to 2% in 2010 among patients with CRC with COPD. In patients with CRC without COPD, the frequency increased from 0.8% to 1.2%. The frequency of reoperation was stable throughout the study period, except for a higher frequency in 2005 (e-appendix 2).

**Mortality**

Thirty-day mortality after CRC surgery was 13.0% (95% CI 11.4% to 14.9%) among patients with CRC with COPD and 5.3% (95% CI 5% to 5.7%) among patients with CRC without COPD, corresponding to an adjusted HR of 1.7 (95% CI 1.5 to 2.0). Mortality in patients with CRC with COPD started to diverge from that in patients with CRC without COPD approximately 3 days after surgery (figure 1).

Thirty-day mortality was 14.8% (95% CI 12.8% to 17.2%) in patients with colon cancer with COPD and 6.2% (95% CI 5.8% to 6.7%) in patients with colon cancer without COPD, corresponding to an adjusted HR of 1.8 (95% CI 1.5 to 2.1).

Thirty-day mortality was 9.0% (95% CI 6.7% to 12.1%) in patients with rectal cancer with COPD and 3.7% (95% CI 3.3% to 4.2%) in patients with rectal cancer without COPD. The adjusted HR was 1.6 (95% CI 1.1% to 2.2%; table 4).

**Stratified analyses**

Stratified analyses showed that COPD had a particularly strong impact on 30-day mortality among patients less than 70 years of age (HR=2.0; 95% CI 1.3% to 3.0%),
among men (HR=1.9; 95% CI 1.6% to 2.4%), among patients undergoing elective surgery (HR=2.2; 95% CI 1.8% to 2.8%), among patients undergoing surgery in a low volume hospital (HR=2.1; 95% CI 1.4% to 3.0%) and among patients undergoing laparoscopic surgery (HR=2.9; 95% CI 1.8% to 4.7%; table 5).

**DISCUSSION**

**Key messages**

- Surgery in patients with CRC with COPD is more frequently associated with complications leading to ICU admission/treatments, and reoperations, compared to patients with CRC without COPD.

**Table 1** Characteristics of patients undergoing colon and rectal cancer surgery by presence/absence of a previous diagnosis of COPD, Denmark, 2005–2011

| Characteristic                  | Colon cancer | | Rectal cancer | |
|--------------------------------|--------------|---|---------------|---|
|                                | COPD         | Per cent | No COPD       | Per cent | COPD | Per cent | No COPD | Per cent |
| Total                          | 1006         | 100 | 11 047        | 100 | 443  | 100      | 5806    | 100 |
| Age group (years)              |              |     |               |     |      |          |         |     |
| 0–59                           | 40           | 4.0 | 1784         | 16.2| 32    | 7.2      | 1280    | 22.0 |
| 60–69                          | 178          | 17.7| 3090         | 28.0| 92    | 20.8     | 1934    | 33.3 |
| 70–79                          | 406          | 40.3| 3504         | 31.7| 202   | 45.6     | 1654    | 28.5 |
| 80+                            | 382          | 38.0| 2669         | 24.1| 117   | 26.4     | 938     | 16.2 |
| Gender                         |              |     |               |     |      |          |         |     |
| Female                         | 512          | 50.9| 5604         | 50.7| 173   | 39.1     | 2232    | 38.4 |
| Male                           | 494          | 49.1| 5443         | 49.3| 270   | 60.9     | 3574    | 61.6 |
| Charlson score                 |              |     |               |     |      |          |         |     |
| 0                              | 572          | 56.9| 8340         | 75.5| 272   | 61.4     | 4625    | 79.7 |
| 1–2                            | 335          | 33.3| 2141         | 19.4| 125   | 28.2     | 867     | 14.9 |
| >3                             | 99           | 9.8 | 566          | 5.1 | 46    | 10.4     | 314     | 5.4  |
| Charlson score                 |              |     |               |     |      |          |         |     |
| Myocardial infarction          |              |     |               |     |      |          |         |     |
| Congestive heart failure       |              |     |               |     |      |          |         |     |
| Peripheral vascular disease    |              |     |               |     |      |          |         |     |
| Cerebrovascular disease        |              |     |               |     |      |          |         |     |
| Dementia                       |              |     |               |     |      |          |         |     |
| Connective tissue disease      |              |     |               |     |      |          |         |     |
| Ulcer disease                  |              |     |               |     |      |          |         |     |
| Mild liver disease             |              |     |               |     |      |          |         |     |
| Diabetes I and II              |              |     |               |     |      |          |         |     |
| Hemiplegia                     |              |     |               |     |      |          |         |     |
| Moderate/severe renal disease  |              |     |               |     |      |          |         |     |
| Diabetes with complication     |              |     |               |     |      |          |         |     |
| Any tumour (except colon and rectal) |          |     |               |     |      |          |         |     |
| Leukaemia                      |              |     |               |     |      |          |         |     |
| Lymphoma                       |              |     |               |     |      |          |         |     |
| Moderate/severe liver disease  |              |     |               |     |      |          |         |     |
| Metastatic solid tumour (except colon and rectal) |          |     |               |     |      |          |         |     |
| AIDS                           |              |     |               |     |      |          |         |     |
| Type of admission              |              |     |               |     |      |          |         |     |
| Elective                       | 683          | 67.9| 7799         | 70.6| 379   | 13.5     | 5202    | 9.9  |
| Acute                          | 322          | 32.0| 3221         | 29.2| 60    | 85.6     | 572     | 89.6 |
| Unknown                        | 1            | 0.1 | 27           | 0.2 | 4     | 0.9      | 32      | 0.5  |
| Type of surgery                |              |     |               |     |      |          |         |     |
| Open                           | 636          | 63.2| 6685         | 60.5| 224   | 50.6     | 2838    | 48.9 |
| Laparoscopic                   | 257          | 25.6| 3236         | 29.3| 72    | 16.2     | 1160    | 20.0 |
| Other                          | 113          | 11.2| 1126         | 10.2| 147   | 33.2     | 1807    | 31.1 |
| Cancer stage                   |              |     |               |     |      |          |         |     |
| Localised                      | 450          | 44.7| 4291         | 38.8| 179   | 40.4     | 2048    | 35.3 |
| Regional                       | 202          | 20.1| 2928         | 26.5| 113   | 25.5     | 2033    | 35.0 |
| Metastasised                   | 214          | 21.3| 2418         | 21.9| 70    | 15.8     | 904     | 15.6 |
| Unknown                        | 140          | 13.9| 1410         | 12.8| 81    | 18.3     | 821     | 14.1 |

COPD, chronic obstructive pulmonary disease.
COPD is clearly associated with increased mortality, especially in patients with colon cancer.

Other studies
To the best of our knowledge, this is the first study to focus specifically on the impact of COPD on postoperative mortality and, in particular, intensive care treatment among patients with CRC. Nonetheless, our mortality results are in accord with those of a study by Osler et al who investigated hospital differences in mortality after CRC surgery.5 In that study, presence of COPD was associated with a 71% increase in 30-day mortality compared to CRC patients without COPD. Studies investigating the impact of COPD on mortality after other types of surgery have also demonstrated an association.6–8 22 For example, Ried et al7 found that 6.1% of patients with COPD died within 30 days of cardiac surgery, compared to 0.8% of patients without COPD. Moreover, a study by Christensen et al8 reported a 30-day mortality of 44% after surgery for peptic ulcer among patients with COPD and 25% among patients without COPD, corresponding to a 50% increased mortality in patients with COPD after adjustment for confounders.

Potential mechanisms
Several physiological mechanisms may explain our findings. Previous studies have shown that patients with COPD suffer more often from postoperative respiratory complications, such as pneumonia, than patients without COPD.13 22 This is supported by our findings of an association between COPD and an increased rate of ICU admission, mechanical ventilation and NIV. Nonetheless, the rates of mechanical ventilation and NIV among patients with COPD remained low, suggesting that a substantial proportion of ICU admissions were related to complications not requiring ventilation support. Furthermore, the higher rate of ICU admissions among patients with CRC with COPD, compared to patients without COPD, could possibly be explained by the fact that some hospitals previously routinely admitted patients with COPD to ICU after CRC surgery. Most hospitals have abandoned this practice, as indicated in our study, which found the most pronounced decline in the number of ICU admissions from 2005 to 2011 among patients with COPD.

Patients with severe COPD who experience exacerbations are usually treated with glucocorticoids, which slow wound healing.25 This could account for some postoperative complications experienced by patients with COPD.

Although patients with COPD are less likely to receive laparoscopic surgery, the impact of COPD on mortality was higher in those who had this type of surgery. This increased mortality impact could be due to perioperative hypoventilation when the pressure in the abdomen is increased during laparoscopic surgery, especially since patients with COPD already have decreased lung capacity.

We observed a relatively lower impact of COPD on mortality in acute patients than in elective patients, which may be explained by the higher absolute mortality risk in acute patients even without COPD.

| Table 2 | Postoperative treatment within 30 days after all colon and rectal cancer surgery by presence/absence of a previous diagnosis of COPD, Denmark, 2005–2011 |
|---------|------------------------------------------------|
|         | ICU admission (%) | NIV (%) | Mechanical Ventilation (%) | Reoperation (%) | Dialysis (%) | Treatment with inotropes/vasopressors (%) |
| Colorectal cancer | COPD | 16.1 (14.3 to 18.1) | 3.6 (2.7 to 4.7) | 1.9 (1.3 to 2.8) | 10.6 (9.0 to 12.3) | 10.9 (9.3 to 12.6) | 11.8 (10.2 to 13.6) |
|         | No COPD | 9.7 (9.3 to 10.2) | 1.1 (0.9 to 1.2) | 1.1 (1.0 to 1.3) | 8.0 (7.6 to 8.4) | 5.4 (5.1 to 5.8) | 6.2 (5.8 to 6.5) |
| Colon cancer | COPD | 17.0 (14.7 to 19.5) | 3.8 (2.7 to 5.1) | 2.2 (1.4 to 3.3) | 11.3 (9.4 to 13.5) | 12.5 (10.5 to 14.7) | 12.7 (10.7 to 14.9) |
|         | No COPD | 9.7 (9.2 to 10.3) | 1.2 (1.0 to 1.4) | 1.2 (1.0 to 1.4) | 7.8 (7.3 to 8.4) | 5.9 (5.5 to 6.4) | 6.4 (6.0 to 6.9) |
| Rectal cancer | COPD | 14.2 (11.1 to 17.8) | 3.2 (1.7 to 5.2) | 1.4 (0.5 to 2.3) | 8.8 (6.3 to 11.8) | 7.2 (5.0 to 10.0) | 9.7 (7.1 to 12.9) |
|         | No COPD | 9.6 (9.0 to 10.4) | 0.9 (0.6 to 1.1) | 1.0 (0.8 to 1.3) | 8.3 (7.6 to 9.0) | 4.5 (4.0 to 5.1) | 5.7 (5.1 to 6.3) |

Numbers in parentheses are 95% CI.
COPD, chronic obstructive pulmonary disease; ICU, intensive care unit; NIV, non-invasive ventilation.

| Table 3 | Postoperative reoperations within 30 days after all colorectal cancer surgery by presence/absence of a previous diagnosis of chronic obstructive pulmonary disease (COPD), Denmark, 2005–2011 |
|---------|-------------------------------------------------|
|         | Wound rupture (%) | Superficial/deep infection (%) | Superficial/deep bleeding (%) | Anastomosis leakage (%) | Other reason for reoperation (%) |
| COPD | Yes | 4.1 (3.2 to 5.3) | 0.9 (0.5 to 1.5) | 2.1 (1.4 to 2.9) | 3.5 (2.6 to 4.6) | 1.0 (0.5 to 1.6) |
|         | No | 2.0 (1.8 to 2.2) | 1.3 (1.1 to 1.5) | 0.8 (0.7 to 1.0) | 3.1 (2.9 to 3.4) | 0.7 (0.6 to 0.9) |

Numbers in parentheses are 95% CI.
A previous study found that patients with CRC with COPD are diagnosed and treated earlier than patients without COPD, so delay in diagnosis and treatment cannot therefore explain the poor prognosis.\textsuperscript{24}

\section*{Limitations}

In this nationwide population-based study conducted in a homogeneous healthcare system, we had complete DCR data and complete follow-up for death, allowing almost no risk of selection bias.\textsuperscript{25} Nonetheless, COPD may be under-reported in the DNRP, in which case we have included some true patients with COPD in the comparison cohort. Such misclassification would have caused us to conservatively bias the relative risk estimates. In the Danish study by Osler \textit{et al}, however, where COPD was defined using both hospitalisations and prescriptions, the COPD prevalence was 11.2\%, which is in accord with our findings.\textsuperscript{4} These findings may imply that under-reporting of COPD is not a major problem in our study. Furthermore, the coding of COPD and ICU admissions and treatments in the DNRP has been demonstrated to have a high-positive predictive value.\textsuperscript{26 27} However, the DNRP does not include detailed clinical data on severity of COPD, and we were therefore unable to evaluate any influence of COPD severity.

We controlled for several important sources of confounding, such as obesity, alcohol-related diseases and other comorbidities. The positive predictive value of the DNRP coding of all diseases included in the CCI is known to be high,\textsuperscript{28} but the index includes limited information on the severity of comorbidity. Despite this, the CCI performs well in colorectal cancer surgery patients.\textsuperscript{29} However, we cannot entirely rule out unmeasured confounding from smoking and incomplete measurement of comorbidity including obesity and alcohol-related diseases.

Although it is very likely that deaths occurring 30 days or less after CRC surgery are caused by complications, information on causes of death may have provided further insight into the mechanisms leading to increased mortality among patients with COPD. Unfortunately, we had no information on causes of death.

\begin{table}[h]
\centering
\begin{tabular}{|c|c|c|c|c|}
\hline
 & Colorectal cancer & Colon cancer & Rectal cancer & \\
\hline
Deaths & 189 & 899 & 684 & 215 \\
\hline
Mortality \% & 13.0 (11.4 to 14.9) & 17.0 (15.5 to 18.6) & 17.2 (15.8 to 18.6) & 9.0 (6.7 to 12.1) \\
\hline
Adjusted HR* (95\% CI) & 1.7 (1.5 to 2.0) & 1.8 (1.5 to 2.1) & 1.6 (1.1 to 2.2) & 3.7 (3.3 to 4.2) \\
\hline
\end{tabular}
\caption{Thirty-day mortality and HRs after colorectal cancer surgery by presence/absence of chronic obstructive pulmonary disease (COPD), Denmark, 2005–2011}
\end{table}
Finally and as mentioned above, rates of ICU admissions among patients with COPD may reflect clinical practice of routine ‘prophylactic’ admission rather than ICU admissions as a marker of postoperative complications; however, we addressed this by excluding ICU admission on the day of surgery. By excluding patients admitted to the ICU on the first postoperative day, we may also have excluded some patients with COPD with immediate surgical complications (e.g., patients not extubated in the operating room, but transferred to the ICU due to perioperative respiratory complications). Sensitivity analysis aimed to address this issue.

Our findings of a substantial impact of COPD on postoperative morbidity and mortality after CRC surgery underscore the need for close postoperative surveillance of patients with CRC with COPD.

In conclusion, we found an increased rate of ICU admission, mechanical ventilation, NIV and reoperation after CRC surgery in patients with COPD, compared with patients with CRC without COPD. In addition, COPD was associated with increased 30-day mortality after CRC surgery.

Table 5  Crude and adjusted HRs for death after colorectal cancer (CRC) surgery, comparing patients with chronic obstructive pulmonary disease (COPD) to those without

| Stratified by                          | COPD                              | Adjusted HR (95% CI) | Reference |
|---------------------------------------|-----------------------------------|----------------------|-----------|
| **Colorectal cancer**                 |                                   |                      |           |
|                                      | Crude HR (95% CI)                 | Adjusted HR (95% CI) |           |
| **Stratified by**                     |                                   |                      |           |
| Age group (years)                     |                                   |                      |           |
| 0–69                                  | 2.5 (1.5 to 4.1)                  | 2.0 (1.3 to 3.0)     | 1.0       |
| 70+                                   | 1.9 (1.6 to 2.3)                  | 1.6 (1.4 to 2.0)     | 1.0       |
| Gender                                |                                   |                      |           |
| Male                                  | 2.9 (2.4 to 3.6)                  | 1.9 (1.6 to 2.4)     | 1.0       |
| Female                                | 2.1 (1.7 to 2.7)                  | 1.5 (1.2 to 1.9)     | 1.0       |
| Charlson score                        |                                   |                      |           |
| 0                                     | 2.7 (2.2 to 3.4)                  | 2.1 (1.7 to 2.6)     | 1.0       |
| 1–2                                   | 1.7 (1.3 to 2.3)                  | 1.5 (1.1 to 1.9)     | 1.0       |
| 3+                                    | 1.7 (1.1 to 2.6)                  | 1.5 (1.0 to 2.3)     | 1.0       |
| Type of admission                     |                                   |                      |           |
| Elective                              | 3.3 (2.7 to 4.1)                  | 2.2 (1.8 to 2.8)     | 1.0       |
| Acute                                 | 1.8 (1.4 to 2.2)                  | 1.4 (1.1 to 1.7)     | 1.0       |
| Hospital volume                       |                                   |                      |           |
| Low (0–500)                           | 2.8 (2.0 to 4.0)                  | 2.1 (1.4 to 3.0)     | 1.0       |
| High (>500)                           | 2.5 (2.1 to 3.0)                  | 1.7 (1.4 to 2.0)     | 1.0       |
| Type of Surgery                       |                                   |                      |           |
| Open                                  | 2.8 (2.3 to 3.5)                  | 2.0 (1.6 to 2.4)     | 1.0       |
| Laparoscopic                          | 4.4 (2.8 to 7.0)                  | 2.9 (1.8 to 4.7)     | 1.0       |
| Other                                 | 1.4 (1.0 to 2.0)                  | 1.0 (0.7 to 1.5)     | 1.0       |
| CRC stage                             |                                   |                      |           |
| Localised                             | 2.9 (2.2 to 3.8)                  | 1.9 (1.4 to 2.5)     | 1.0       |
| Regional                              | 3.4 (2.3 to 5.0)                  | 2.1 (1.4 to 3.0)     | 1.0       |
| Metastasised                          | 2.4 (1.8 to 3.2)                  | 1.7 (1.2 to 2.2)     | 1.0       |
| Unknown                               | 1.7 (1.2 to 2.5)                  | 1.4 (0.9 to 2.0)     | 1.0       |

Contributors HTS, CFC, RE and AMP conceived the study idea and designed the study. LKA and CS collected the data. AMP analysed the data with help from CS, RE and CFC. All authors interpreted the findings. AMP reviewed the literature together with RE and CFC. AMP wrote the first draft, and all authors critically reviewed and edited the manuscript and approved the final version.

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Data sharing statement No additional data are available.

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