Association of Severe Obesity and Chronic Obstructive Pulmonary Disease With Pneumonia Following Non-Cardiac Surgery

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Abstract

Background: Pneumonia is the third most common surgical complication after urinary tract infection and wound infections. In addition to increased mortality, patients who develop postoperative pneumonia have a higher risk of prolonged hospital stay, intensive care unit (ICU) admissions, and higher healthcare costs. Obesity and chronic obstructive pulmonary disease (COPD) are both independent risk factors for the development and severity of postoperative pneumonia, although the combined effect of these comorbidities is unknown. Therefore, we evaluated whether the combination of severe obesity and COPD is associated with an increased risk of postoperative pneumonia.

Methods: We performed a multicenter retrospective cohort study of 365,273 patients aged 18 - 64 years who were either severely obese (body mass index (BMI) ≥ 40 kg/m²) or normal-weight (BMI between 18.6 and 24.9 kg/m²) and underwent general surgery, orthopedic surgery, neurosurgery, otolaryngology surgery, urology surgery, and vascular surgery in the American College of Surgeons (ACS) National Surgical Quality Improvement Program (NSQIP) participating hospitals from 2014 to 2018. We evaluated the combined effect of COPD and severe obesity on the risk for postoperative pneumonia, unplanned tracheal reintubation, and extended length of stay.

Results: The co-occurrence of severe obesity and COPD appeared to have a protective effect on the risk of postoperative pneumonia. In the presence of COPD, patients with severe obesity were 14% less likely to develop pneumonia compared to their normal-weight counterparts (2.9% vs. 4.4%; adjusted relative risk (RR): 0.76; 95% confidence interval (CI): 0.60, 0.95). In addition, in the presence of COPD, severe obesity conferred a lower risk for requiring an extended length of stay (37.6% vs. 47.9%; adjusted RR: 0.83; 95% CI: 0.78, 0.89).

Conclusions: Counterintuitively, the co-occurrence of severe obesity with COPD appeared to buffer the negative impact of COPD on postoperative pneumonia, unplanned tracheal reintubation, and prolonged hospital stay after noncardiac surgery. These findings are consistent with the obesity paradox and warrant further investigations.

Keywords: Severe obesity; COPD; Postoperative pneumonia; Unplanned tracheal reintubation

Introduction

Pneumonia is the third most common surgical complication after urinary tract infections and wound infections and is associated with significant perioperative morbidity [1, 2]. Patients who develop postoperative pneumonia have longer hospital stays, are more likely to be admitted to the intensive care unit (ICU) and are at a higher risk of early postoperative mortality [3-5]. The morbidity of pneumonia accounts for an additional 6 - 9 hospital days resulting in approximately $46,000 - $52,466 in hospital costs per patient after abdominal surgery [6]. In addition, under the Patient Protection and Affordable Care Act, hospitals with a high rate of 30-day readmissions for Medicare patients incur financial penalties [7]. Therefore, an improved understanding of complex risk factors associated with postoperative pneumonia may help identify these at-risk patients for targeted postoperative monitoring and rehabilitation.

Obesity and chronic obstructive pulmonary disease (COPD) are both independent risk factors for the development and severity of postoperative pneumonia [8-13]. Postoperative infections are highest among severely obese patients, potentially due to impaired immune function [14-17]. Severely obese patients may be at higher risk for aspiration because of their susceptibility to gastroesophageal reflux and hiatal hernia [18]. Furthermore, anatomical changes, including excess adipose tissue in pharyngeal walls, increased intra-abdominal pressure, and decreased chest wall compliance, may...
predispose patients with obesity to upper airway obstruction, reduced lung volume, airway collapse, atelectasis, and postoperative respiratory failure [18, 19]. In patients with COPD, increased susceptibility to pulmonary infection has been attributed to several factors, including impaired gas exchange and reduced mucociliary clearance, inflammation from airway instrumentation, preoperative airway bacterial colonization, and surgery-induced immunosuppression [20-22].

Therefore, it is reasonable to assume that the combined effect of obesity and COPD could potentiate the risk of postoperative pneumonia. Unfortunately, research examining both comorbidities in relation to postoperative pneumonia is currently unavailable. Therefore, this study evaluated whether the joint presence of obesity and COPD in patients increases their risk of postoperative pneumonia among a cohort of patients who underwent inpatient surgical procedures.

**Materials and Methods**

**Data source**

We analyzed a multicenter retrospective dataset from the American College of Surgeons National Surgical Quality Improvement Program (NSQIP) database from 2014 to 2018. The NSQIP is a multi-institutional surgical risk-adjusted, outcomes-based program that collects over 135 pre, intra, and postoperative clinical variables, including demographics, comorbidities, laboratory values, and tracks patient outcomes for up to 90 days postoperatively from > 600 participating hospitals [23]. In addition, trained surgical clinical reviewers identify data by current procedural terminology, collect, record data, and periodically perform random audits to ensure data accuracy and fidelity [23]. Our Institutional Review Board waived informed consent because our study was not considered human subjects research. This study was conducted in compliance with the ethical standards of the responsible institution on human subjects as well as with the Helsinki Declaration.

**Study design and population**

We performed a retrospective cohort study of patients aged between 18 and 64 years who were either severely obese (body mass index (BMI) ≥ 40 kg/m²) or normal-weight (BMI between 18.6 and 24.9 kg/m²) [24] and underwent general surgery, orthopedic surgery, neurosurgery, otolaryngology surgery, urology surgery, and vascular surgery. We excluded patients who had a history of dyspnea prior to surgery, required mechanical ventilation in the 48 h prior to surgery, were diagnosed with sepsis preoperatively, or who were admitted for emergency surgery.

**Outcome of interest**

The outcome of interest was postoperative pneumonia, diagnosed when predefined standardized criteria (clinical, radiological, or laboratory) are met. The radiological criteria consist of a chest radiological examination that demonstrates either an infiltrate, a consolidation, a cavitation, or a pleural effusion. The clinical (or laboratory) criteria include meeting one of two sections: 1) New onset of purulent sputum, or change in the character of sputum, or increased respiratory secretions, or increased suctioning requirements; new onset or worsening cough, dyspnea, or tachypnea; rales or rhonchi, worsening gas exchange; 2) Microscopic evidence of bacterial pneumonia. The clinical, radiologic, and microbiologic criteria used in the NSQIP were summarized here (Supplementary Material 1, www.jocmr.org). Our secondary outcome was unplanned tracheal reintubation within 30 days following surgery. We also explored the risk of extended length of postoperative hospital stay, defined as length of stay (LOS) longer than the 75th percentile of the study cohort.

**Statistical analysis**

We summarized categorical variables as frequencies and column percentages. Non-continuous variables were summarized as median (interquartile range (IQR)). We evaluated the risk of pneumonia in the presence or absence of COPD and according to whether patients were severely obese or normal-weight. To this end, we first divided the study population into four mutually exclusive groups: 1) Normal-weight patients without COPD (doubly unexposed); 2) Normal-weight patients with COPD; 3) Severely obese patients without COPD; and 4) Severely obese patients with COPD (doubly exposed). We then used log-binomial regression models to estimate the relative risk (RR) and its 95% confidence intervals (CIs) adjusting for covariates, selected a priori, based on our hypothesis of confounding the association between COPD and postoperative pneumonia: gender (female vs. male), current smoking status (within 1 year of surgery), American Society of Anesthesiologists (ASA) physical classification (≥ 3 vs. < 3), diabetes (noninsulin-dependent or insulin-dependent diabetes mellitus: yes vs. no), dyspnea (at rest or moderate exertion: yes vs. no), functional health status within 30 days of surgery (dependent vs. independent), history of chronic heart failure within 30 days of surgery (yes vs. no), hypertension (yes vs. no), corticosteroid use for a chronic condition (yes vs. no), and surgical specialty. We considered being statistically significant a P value of < 0.05. All analyses were performed with STATA 16 (StataCorp).

**Results**

**Demographic and clinical characteristics**

We analyzed a total of 365,273 patients who met inclusion criteria during the 5-year study period spanning between 2014 and 2018, of whom 9,329 (2.6%) had COPD (Table 1). Among patients without COPD, 50.4% (n = 179,451) were normal-weight, and 49.6% (n = 176,493) were severely obese. Among patients with COPD, 54.6% (n = 5,095) were normal-weight,
and 45.4% (n = 4,234) were severely obese. In the absence of COPD, severely obese patients were more likely to be female (70.1% vs. 54.8%), to have an ASA class ≥ 3 (72.1% vs. 39.9%), to have diabetes (23.6% vs. 7.2%), to have hypertension requiring medication (47.0% vs. 22.7%); but less likely to have smoked within 1 year of surgery (14.3% vs. 28.5%) and use steroid for a chronic condition (3.1% vs. 7.0%). Similar differences were observed between normal-weight and severely obese patients in the presence of COPD.

**Combined effects of obesity and COPD**

Out of the 365,273 total patients, 3,554 developed pneumonia (1.0%), 2,443 required unplanned tracheal reintubation following surgery (0.7%), and 100,785 required an extended length of hospital stay (27.6%). Overall, patients with COPD were more likely to develop pneumonia following surgery (3.7% vs. 0.9%; adjusted RR: 2.07; 95% CI: 1.84, 2.33). In addition, COPD conferred a higher risk of unplanned tracheal reintubation (2.4% vs. 0.6%; adjusted RR: 1.80; 95% CI: 1.56, 2.08) and extended LOS (43.2% vs. 27.2%; adjusted RR: 1.12; 95% CI: 1.08, 1.15) (Fig. 1).

The co-occurrence of severe obesity and COPD appeared to have a protective effect on the risk of postoperative pneumonia. Specifically, in the presence of COPD, patients with severe obesity were 14% less likely to develop pneumonia compared to their normal-weight counterparts (2.9% vs. 4.4%; adjusted RR: 0.76; 95% CI: 0.60, 0.95). The protective effect of severe obesity was also observed for the requirement of extended LOS (37.6% vs. 27.2%; adjusted RR: 0.83; 95% CI: 0.78, 0.89). In the presence of COPD, severe obesity conferred a lower risk of unplanned tracheal reintubation relative to normal weight, however, the statistical significance was lost after adjusting for baseline covariates (2.0% vs. 2.7%; adjusted RR: 0.79%; 95% CI: 0.60, 1.05) (Table 2).

**Discussion**

Although obesity and COPD are known factors for the development of postoperative pneumonia, we are unaware of a
previous study exploring their combined effects on the risk of postoperative pneumonia. Consistent with previous studies, we found that COPD conferred a higher risk of postoperative pneumonia with or without the presence of severe obesity [20, 25, 26]. However, counterintuitively, the presence of COPD carried a poorer prognosis in normal-weight patients relative to patients with severe obesity. Such a paradoxical relationship of severe obesity and COPD with postoperative pneumonia mirrors previous studies describing the perceived protection of obesity against some chronic debilitating diseases and post-surgical pulmonary complications [27-32]. Results from a retrospective cohort study on patients with COPD who used tobacco found that obesity was associated with a lower risk for mortality compared to normal weight [33]. Similarly, in another retrospective study of patients with pulmonary hypertension undergoing right-sided heart catheterization, patients with obesity had a lower risk of mortality compared to normal weight patients [34]. Our study findings together with data from these studies support and contribute to the growing body of evidence for the “obesity paradox”.

Several theories related to patient characteristics and provider behavior may explain why obesity appears to modify the association between COPD and postoperative pneumonia. One distinctive pathophysiological feature of COPD is lung hyperinflation due to loss of normal lung elastase and limited expiratory flow [35, 36]. These changes decrease vital capacity and tidal volume and limit the ability to respond to increased minute ventilation needs, as seen after surgery [37-40]. Obesity decreases chest wall compliance (due to increasing mass), decreasing the expiratory reserve volume and, therefore, functional residual capacity. This results in increased atelectasis which may predispose to hypoxemia following general anesthesia. Thus, each condition has unique pathophysiologic reasons to increase the risk of pneumonia postoperatively. However, almost 40 years ago, Ora et al demonstrated that obesity protected against both static and dynamic lung hyperinflation producing favorable pulmonary mechanics which were protective against dyspnea [40]. Thus, it is likely that pulmonary mechanical challenges of obesity and COPD offset one another to normalize the risk of postoperative pneumonia.

Due to increased muscle mass, obese patients may have a greater nutritional and metabolic reserve, particularly of amino acids [41]. This increased reserve may better resist the catabolic effects of COPD and the initial infectious process than healthy-weight counterparts [42, 43]. Furthermore, the increased nutritional reserves may help obese patients to withstand the alteration in metabolic and inflammatory stress that follows surgery [15, 44]. In support of this assertion, Windsor and colleagues, in their study of risk factors for postoperative pneumonia, reported that patients with preoperative protein depletion were significantly more likely to develop pneumonia following abdominal surgery [45].

The comorbidity of COPD and severe obesity may trigger heightened monitoring from clinicians, thus reducing the risk of preventable complications, including pneumonia. In addition to increased monitoring, there may be heightened awareness among clinicians for intervening at the earliest sign of a respiratory issue in these patients, thereby preventing the development and progression of postoperative pneumonia [35]. Alternatively, in an effort to prevent perioperative complications, surgical patient selection may have biased toward “healthy” obese patients in this non-bariatric surgical cohort.

Our findings should be interpreted with the limitations inherent in a retrospective cohort study design, including the possibility of errors as can be seen in any large dataset. First, we did not separate the diagnostic subtypes of postoperative pneumonia, expressly ventilator-associated (VAP) vs. hospital-acquired, which may have different risk profiles in the obese and COPD population. Second, we did not examine the type and extent of surgery in the study cohort. Evidence suggests that the likelihood of developing post-surgical nosocomial infections ranges between 1.5% and 57% depending on the type and extent of surgery [6]. It may be possible that the severely obese COPD patients in our dataset were more likely to have minor surgeries than the healthy-weight COPD cohort, thereby reducing their potential risk of pneumonia. Third, it is plausible that postoperative pneumonia diagnosis could be misclassified by overlapping diagnoses such as aspiration pneumonia, pulmonary edema, and pulmonary embolism. Even so, we have no reason to believe that such misclassification bias will skew towards a particular weight group, and it is unlikely to be widespread enough to explain away our findings. Fourth, we did not
explore the other possible causes of unintended tracheal reintubation such as anesthetic or surgical complications, nor did we explore other postoperative complications such as wound or urinary tract infections, and deep vein thrombosis (DVT), which may confound our findings on extended length of hospital stay. Finally, we could not determine from the NSQIP database, the severity of COPD among the study subjects based on the GOLD/ABCD classification. Given the information included in the database, we were unable to determine if chronic corticosteroid use was in place for the treatment of COPD.

Our retrospective study of a multi-institutional dataset revealed that the co-occurrence of severe obesity and COPD was associated with a lower rate of postoperative pneumonia. While various surgical complications are associated with COPD, its combination with severe obesity was counterintuitively associated with lower rates of extended hospital stay. However, we caution against interpreting this perceived protective effect of severe obesity in the presence of COPD to promote obesity or suggest excess weight gain in patients with COPD undergoing surgery. Although the underlying mechanism explaining the obesity paradox may differ across different health outcomes, our study provides additional evidence of the pulmonary mechanical consequences of the obesity paradox and warrant further research.

Supplementary Material

Suppl 1. NSQIP definition of pneumonia.

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The authors have no financial relationships relevant to this article to disclose.

Conflict of Interest

The authors have no conflict of interest relevant to this article to disclose.

Informed Consent

Our Institutional Review Board waived informed consent because our study was not considered human subjects research.

Author Contributions

Kwaku Owusu-Bediako helped with study design/conduction, data interpretation, manuscript preparation and editing. Kayla
Pfaff helped with manuscript preparation and editing. Nguyen K. Tran helped with data interpretation, manuscript preparation and editing. David L. Stahl helped with data interpretation, manuscript preparation and editing. Joseph D. Tobias helped with study design/conduction, statistical analysis, data interpretation, manuscript preparation and editing. Olubukola O. Nafiu and Christian Mpody helped with study design/conduct, statistical analysis, data interpretation, manuscript preparation and editing.

Data Availability

Any inquiries regarding supporting data availability of this study should be directed to the corresponding author.

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