Impact of BMI on exacerbation and medical care expenses in subjects with mild to moderate airflow obstruction

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Background and objective: The rate of obesity is increasing in Asia, but the clinical impact of body mass index (BMI) on the outcome of chronic obstructive pulmonary disease (COPD) remains unknown. We aimed to assess this impact while focusing on the risk of exacerbation, health-care utilization, and medical costs.

Methods: We examined 43,864 subjects registered in the Korean National Health and Nutrition Examination Survey (KNHANES) database from 2007 to 2012, and linked the data of COPD patients who had mild to moderate airflow obstruction (n = 1,320) to National Health Insurance (NHI) data. COPD was confirmed by spirometry. BMI was used to stratify patients into four categories: underweight (BMI <18.5 kg/m²), normal range (18.5–22.9 kg/m²), overweight (23–24.9 kg/m²), and obese (≥25 kg/m²).

Results: Of the 1,320 patients with COPD with mild to moderate airflow obstruction, 27.8% had a BMI ≥25 kg/m². Compared with normal-weight patients, obese patients tended to experience fewer exacerbations (incidence rate ratio [IRR] 0.88; 95% CI 0.77–0.99; P = 0.04), although this association was not significant in a multivariable analysis. COPD-related health-care utilization and medical expenses were higher among underweight patients than the other groups. After adjustment, the risk of COPD-related hospitalization was highest among underweight and higher among overweight patients vs normal-weight patients (adjusted IRRs: 7.12, 1.00, 1.26, and 1.02 for underweight, normal, overweight, and obese groups, respectively; \( P = 0.01 \)).

Conclusion: Decreased weight tends to negatively influence prognosis of COPD with mild to moderate airflow obstruction, whereas higher BMI was not significantly related to worse outcomes.

Keywords: COPD, obesity, exacerbation, health-care utilization, medical expenses

Introduction

Chronic obstructive pulmonary disease (COPD) is a major cause of morbidity and mortality worldwide and has an increasing prevalence.1 COPD does not merely comprise progressive lung disease, but is complicated by associations with several comorbid conditions that contribute to a poor health status and increases in health-care utilization and expenses, hospitalization, and mortality. According to a previous epidemiologic report, more than 50% of patients with COPD have at least one comorbidity.2 Obesity, a known comorbidity of COPD, may contribute to a patient’s respiratory symptoms and reduced exercise capacity.3–5 Paradoxically, despite the propensity of obesity to increase the prevalence rates of various diseases and cause unfavorable outcomes in the general population, large cohort studies have demonstrated an association of obesity with reduced mortality among patients with COPD.6,7 In those studies, mortality was
highest among patients with a low body mass index (BMI), whereas overweight – but not extremely obese – patients exhibited improved survival. This finding suggests a survival benefit associated with increased BMI among overweight and mildly obese patients. However, previous studies of obesity among patients with COPD included symptomatic patients who received treatment and were followed regularly. The low prevalence of obesity in Asian populations relative to Western populations has led to few reports of the impact of BMI on COPD. However, obesity is increasing among Asians in response to increasingly Westernized diets. 

It is extremely difficult to enroll patients with mild, asymptomatic COPD who rarely visit hospital, in clinical trials or cohort studies, and this results in lack of data on the clinical features of those patients. However, we can evaluate these patients through our unique national insurance system. In the present study, we aimed to identify the clinical impact of BMIs on the outcome of patients with mild, asymptomatic COPD with mild to moderate airflow obstruction while focusing on the risk of exacerbation, health-care utilization, and medical expenses.

Methods
Source of data
In 1998, South Korea implemented a unique, compulsory, health insurance system for its population, and the corresponding National Health Insurance (NHI) database provides nationwide data with regard to illness and health-care utilization patterns. The Korean National Health and Nutrition Examination Survey (KNHANES) is conducted annually. In this study, we utilized KNHANES database records from 2007 to 2012 and merged with the NHI database for the same period. The KNHANES and NHI data of patients with COPD were linked by the Korean Centres for Disease Control and the Health Insurance Review and Assessment Service.

This study was exempted from the informed consent requirement because of its retrospective nature and was approved by the Institutional Review Board Committee of Kyung Hee University Hospital (IRB No. KHUH 2017-05-058).

Definitions of COPD and BMI category
COPD was defined as a spirometric airflow limitation (forced expiratory volume in 1 s [FEV₃]/forced vital capacity [FVC] <0.70) and only patients with COPD who met the following inclusion criteria were included: 1) age ≥40 years, and 2) FEV₃ ≥50%.

In Caucasians, obesity is defined as a BMI ≥30 kg/m²; however, the World Health Organization (WHO) has proposed a cutoff of 25.0 for Asian populations, and this criterion is recommended by the Korean Society for the Study of Obesity. We adopted this cutoff value in the present study and subdivided patients by BMI into four categories: underweight (BMI <18.5 kg/m²), normal range (18.5–22.9 kg/m²), overweight (23–24.9 kg/m²), and obese (≥25 kg/m²).

Clinical data measurements
We utilized and merged the Korean NHI (2008–2012) and KNHANES (2007–2012) data records of patients with COPD to investigate the impact of BMI on the risk of exacerbation and health-care utilization. The researchers were blinded to the subjects’ personally identifiable information.

The KNHANES provides researchers with a variety of clinical information parameters such as age, sex, spirometry result, and smoking status. Spirometry was conducted by an experienced technician and the result was checked by technicians who were working in tertiary affiliated hospitals. Comorbid conditions were identified by searching for the International Classification of Disease, Tenth Revision (ICD-10) codes in the Health Insurance Review and Assessment (HIRA) Service database. Moreover, we analyzed health-care utilization and expenses extracted from the NHI database. We set 2007 as the baseline and used data from that year to assess the previous exacerbation history, whereas the medical expenses analysis targeted data from 2008. A moderate exacerbation was defined as an outpatient clinic visit for worsening respiratory symptoms and the prescription of antibiotics and/or systemic steroids, whereas a severe exacerbation was defined as an exacerbation necessitating hospitalization or an emergency department visit. In addition to all-cause health-care utilization and medical expenses, we evaluated COPD-related utilization and expenses. Data with regard to inpatient and outpatient services and the ICD-10 code of COPD (J43.x–J44.x) with prescription of COPD-related medication were used.

All expenses are presented in US dollars (USD) with an exchange rate of 1 USD to 1,140 Korean won (May 2017 exchange rate).

Statistical analysis
Differences between the four BMI-stratified groups were assessed using the chi-square test for categorical variables and an analysis of variance for continuous variables. Because patients with COPD included in this study were detected by screening spirometry and usually had mild respiratory symptoms and relatively mild airflow limitation, exacerbation-related hospital visits were expected to be infrequent. Therefore, the incidence of exacerbation was analyzed using a negative binomial regression analysis model.
The multivariable analyses were adjusted for covariates such as age, sex, FEV\textsubscript{1}, and smoking pack-year. Then, we conducted subgroup analyses according to the percent predicted FEV\textsubscript{1}: FEV\textsubscript{1} ≥80%, and FEV\textsubscript{1} ≥50 but <80% predicted. The 5-year cumulative health-care utilization and medical expenses were expressed in terms of person-years and USD, respectively.

All analyses were two-sided and conducted at a significance level of 0.05 unless otherwise noted. All analyses were undertaken using SAS software, version 9.2 (SAS Institute Inc., Cary, NC, USA).

**Results**

**Characteristics of patients**

Among the 43,864 subjects who participated in KNHANES during the 6-year period, 1,320 were identified as COPD patients with mild to moderate airflow obstruction (Figure 1). Table 1 presents the demographic and clinical characteristics stratified by BMI category. Overall, 95.6% of patients were male, and 26.8% and 27.8% were overweight and obese, respectively. Younger age correlated significantly with an increased BMI, whereas the smoking rates were similar across BMI categories. Compared with a normal BMI, hypertension was more prevalent in the underweight and overweight/obesity groups, and osteoporosis was less prevalent in the obese group. The prevalence of other comorbidities was similar among the groups.

The groups did not differ with regard to a previous exacerbation history in the past year or initial treatment (including inhaler prescription). The EuroQol-5 Dimensions questionnaire (EQ-5D) scores were significantly lower in underweight patients, but did not differ among higher BMI groups. Furthermore, the underweight group had a low lung volume and more severe airflow limitation, whereas no significant trend was observed with increasing BMI (Table 2).

**Risk of exacerbation across the BMI categories**

Compared with the normal BMI group, the risk of moderate to severe exacerbation during 2008–2012 was significantly lower in the obese group (incidence rate ratio [IRR] 0.88; 95% CI 0.77–0.99; P-value, 0.04). However, after adjusting for the covariates of age, sex, percent predicted FEV\textsubscript{1}, and/or smoking pack-years, an increase in BMI did not correlate with a significant reduction in the exacerbation incidence (Table 3). In a subgroup analysis according to the percent predicted FEV\textsubscript{1} – FEV\textsubscript{1} ≥80%, and FEV\textsubscript{1} ≥50 but <80% predicted – an increasing BMI was similarly not found to correlate significantly with a reduced incidence of exacerbation in either the adjusted or unadjusted models (Table S1).

**Health-care utilization and cost**

COPD-related events (person-year) and medical expenses (USD) are presented in Table 4. The underweight group experienced more frequent hospitalization, outpatient visits, and pharmacy utilization as well as medical expenses, whereas these parameters did not differ markedly among the three higher-weight groups (Figure S1). Notably, the obese group exhibited trends toward the lowest health-care utilization rate and medical expenses, except for pharmacy costs.

Table 5 presents the IRRs of 5-year COPD-related health-care utilization. After adjusting for previously mentioned covariates, hospitalization was highest in the underweight group, although the overweight group exhibited an increased risk of hospitalization (adjusted IRRs: 7.12, 1.00, 1.26, and 1.02 for underweight, normal, overweight, and obese groups, respectively; P = 0.01). The IRRs for hospitalization did not differ significantly between the groups divided by the percent predicted FEV\textsubscript{1} (Table S2).

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**Figure 1** Flow diagram of the study participants.
Table 1 Comparison of demographic and clinical characteristics among patients with chronic obstructive pulmonary disease (COPD) stratified by body mass index (BMI)*

| BMI (kg/m²)      | P-value |
|------------------|---------|
| <18.5            | 18.5–22.9 | 23–24.9 | ≥25 |
| Subjects         | 37 (2.8) | 562 (42.6) | 354 (26.8) | 367 (27.8) |
| Age, years       | 70.5 ± 8.3 | 65.5 ± 9.2 | 64.5 ± 9.5 | 62.4 ± 9.6 | <0.01 |
| Sex, male (%)    | 33 (89.2) | 529 (95.9) | 344 (97.2) | 356 (97.0) | 0.08 |
| BMI (kg/m²)      | 17.5 ± 0.8 | 21.4 ± 1.1 | 23.9 ± 0.6 | 26.8 ± 1.5 | <0.01 |
| Smoking, pack-year | 33.5 ± 15.6 | 35.3 ± 21.0 | 35.1 ± 21.4 | 35.6 ± 20.9 | 0.94 |

Comorbid condition
- DM: 11 (29.7) | 205 (36.5) | 133 (37.6) | 148 (40.3) | 0.49
- HTN: 22 (59.5) | 274 (48.8) | 196 (55.4) | 234 (63.8) | <0.01
- Ischemic heart disease: 5 (13.5) | 49 (8.7) | 39 (11.0) | 39 (10.6) | 0.55
- Congestive heart failure: 3 (8.1) | 29 (5.2) | 13 (3.7) | 16 (4.4) | 0.54
- Arthritis: 5 (13.5) | 54 (9.6) | 34 (9.6) | 38 (10.4) | 0.87
- Osteoporosis: 11 (29.7) | 72 (12.8) | 46 (13.0) | 32 (8.7) | <0.01
- GERD: 19 (51.4) | 293 (52.1) | 201 (56.8) | 187 (51.0) | 0.41
- History of PTB: 0 (0) | 5 (0.9) | 6 (1.7) | 1 (0.3) | 0.22
- Depression: 9 (24.3) | 77 (13.7) | 49 (13.8) | 51 (13.9) | 0.35

History of exacerbation in the past year
- Moderate to severe: 28 (75.7) | 400 (71.2) | 250 (70.6) | 263 (71.7) | 0.93
- Frequency n/year: 4.1 ± 5.5 | 3.4 ± 6.4 | 3.0 ± 4.0 | 3.4 ± 4.4 | 0.40

Initial treatment, n (%)
- ICS and LABA: 1 (2.7) | 17 (3.0) | 7 (2.0) | 11 (3.0) | 0.79
- LAMA and/or LABA: 1 (2.7) | 4 (0.7) | 3 (0.9) | 0 (0) | 0.15
- SABA and/or SAM: 1 (2.7) | 23 (4.1) | 14 (4.0) | 17 (4.6) | 0.93

Note: *Data are presented as number (%) or mean ± SD.
Abbreviations: BMI, body mass index; DM, diabetes mellitus; GERD, gastroesophageal reflux disease; HTN, hypertension; ICS, inhaled corticosteroid; LABA, long-acting beta receptor agonist; LAMA, long-acting muscarinic receptor agonist; PTB, pulmonary tuberculosis; SABA, short-acting beta receptor agonist; SAM, short-acting muscarinic receptor agonist.

Discussion
To our knowledge, this is the first study to assess the relationship of BMI with the risk of exacerbation and health-care utilization among patients – and, particularly, Asian patients – with COPD with mild to moderate airflow obstruction. We found that underweight patients experienced more frequent clinic visits and hospitalization, whereas increase in BMI did not correlate significantly with increased COPD-related exacerbations.

Table 2 Comparison of the quality of life, laboratory findings, and pulmonary function among patients with chronic obstructive pulmonary disease (COPD)*

| BMI (kg/m²)      | P-value |
|------------------|---------|
| <18.5            | 18.5–22.9 | 23–24.9 | ≥25 |
| Subjects         | 37 (2.8) | 562 (42.6) | 354 (26.8) | 367 (27.8) |

Index of quality of life
- EQ-5D: 0.86 ± 0.18 | 0.92 ± 0.12 | 0.93 ± 0.11 | 0.93 ± 0.13 | <0.01

Laboratory test
- Total serum cholesterol, mg/dL: 167.3 ± 40.5 | 187.1 ± 36.1 | 187.6 ± 36.9 | 189.8 ± 35.1 | <0.01
- TG, mg/dL: 126.5 ± 116.3 | 147.4 ± 120.9 | 167.2 ± 102.9 | 191.2 ± 181.9 | <0.01
- LDL cholesterol, mg/dL: 109.5 ± 23.4 | 111.4 ± 36.5 | 109.8 ± 38.6 | 113.6 ± 32.0 | 0.93
- HDL cholesterol, mg/dL: 49.4 ± 12.2 | 46.8 ± 11.4 | 44.4 ± 11.0 | 42.6 ± 9.5 | <0.01

Lung function
- FVC, L: 3.4 ± 0.7 | 3.9 ± 0.7 | 4.0 ± 0.8 | 3.9 ± 0.7 | <0.01
- FVC, % predicted: 88.9 ± 15.0 | 92.5 ± 12.7 | 91.6 ± 13.0 | 87.4 ± 12.6 | <0.01
- FEV₁, L: 2.1 ± 0.5 | 2.4 ± 0.6 | 2.5 ± 0.6 | 2.5 ± 0.5 | <0.01
- FEV₁, % predicted: 76.0 ± 14.8 | 78.7 ± 12.9 | 79.6 ± 13.6 | 77.9 ± 12.0 | 0.18
- FEV₁/FVC: 0.62 ± 0.07 | 0.63 ± 0.06 | 0.63 ± 0.06 | 0.65 ± 0.05 | <0.01

Note: *Data are presented as number (%) or mean ± SD.
Abbreviations: EQ-5D, EuroQol-5 Dimensions questionnaire; FEV₁, forced expiratory volume in 1 s; FVC, forced vital capacity; HDL, high-density lipoprotein; LDL, low-density lipoprotein; TG, triglyceride; BMI, body mass index.
health-care utilization; notably, we observed similar results for medical expenses. A low BMI is known to be associated with an increased risk of hospitalization in patients with COPD, and our findings confirm this relationship. In this study, we used the obesity definition for Asians, but the results were not significantly different even when using the Western BMI category proposed by WHO.

Although our results did not show a significant association between obesity and a higher risk of exacerbation or increased health-care utilization, several reports have described the harmful effects of obesity on COPD outcomes, including respiratory-specific quality of life (QOL), exercise capacity, and dyspnea severity. Similarly, Lambert et al recently reported the negative effects of obesity on outcomes related to moderate to severe COPD, including QOL, dyspnea, exercise tolerance, and severe exacerbation. More recently, patients with COPD who had undergone bariatric surgery experienced fewer emergency department visits and hospitalizations for COPD exacerbation.

On the other hand, studies of asymptomatic COPD patients who had mild to moderate airflow obstruction are rare, as the affected patients generally do not experience sufficiently severe symptoms that would induce a voluntary hospital visit. Therefore, clinical practices find it extremely difficult to enroll those patients. However, we benefited from the use of the KNHANES to identify these patients. The patients diagnosed with COPD in the KNHANES database were asymptomatic. Their EQ-5D scores were almost normal, and almost none had received a previous COPD diagnosis or treatment. Most patients did not know their COPD status and were identified incidentally via spirometry screening. An extremely low frequency of inhaler prescriptions and low EQ-5D scores indicate that the population in this study truly comprised patients with asymptomatic COPD. Furthermore, it is difficult to follow up these patients. Despite the incidental diagnosis of COPD, patients who are largely asymptomatic have little motivation to participate in regular follow-ups. Similarly, routine hospital visits are very rare. Fortunately, however, we were able to connect patient information to the HIRA database. This allowed us to trace all health-care utilization and expense data without omission.

Few studies have investigated the characteristics and courses of patients with asymptomatic COPD who are expected to progress to symptomatic COPD over time.

Table 3 Incidence of moderate to severe exacerbation during a 5-year period

| BMI (kg/m²) | Crude IRR | 95% CI | P-value | Adjusted IRR | 95% CI | P-value | Adjusted IRR | 95% CI | P-value |
|-------------|-----------|--------|---------|-------------|--------|---------|-------------|--------|---------|
| <18.5       | 1.10      | 0.80–1.51 | 0.57     | 0.94        | 0.69–1.29 | 0.72 | 0.95        | 0.70–1.30 | 0.75    |
| 18.5–22.9   | Reference |        |         |             |         |        |             |         |         |
| 23–24.9     | 0.94      | 0.82–1.06 | 0.32     | 0.96        | 0.85–1.09 | 0.52 | 0.96        | 0.85–1.09 | 0.53    |
| ≥25         | 0.88      | 0.77–0.99 | 0.04     | 0.97        | 0.86–1.10 | 0.68 | 0.98        | 0.86–1.11 | 0.70    |

Notes: *Adjusted by age, sex, and FEV₁ (%). † Adjusted by age, sex, smoking pack-year and FEV₁ (%).
Abbreviations: IRR, incidence rate ratio; BMI, body mass index.

Table 4 Cumulative chronic obstructive pulmonary disease (COPD)-related health-care utilization and medical expenses during a 5-year period

| BMI (kg/m²) | P-value |
|-------------|---------|
|             |         |
| <18.5       |         |
| 18.5–22.9   |         |
| 23–24.9     |         |
| ≥25         |         |

Notes: *All costs are presented in US dollars (USD) with an exchange rate of 1 USD equal to 1,140 Korean won (exchange rate on May, 2017). † Data are presented as number (%) or mean ± SD.
Abbreviation: BMI, body mass index.
Table 5 Risk of chronic obstructive pulmonary disease (COPD)-related health-care utilization during a 5-year period

| BMI (kg/m²) group | Crude IRR (95% CI) | Adjusted IRR* (95% CI) | Adjusted IRR* (95% CI) | Adjusted IRR* (95% CI) |
|------------------|------------------|--------------------------|--------------------------|--------------------------|
|                  | Hospitalization  | Outpatient clinic visits | Pharmacy use             | Hospitalization  | Outpatient clinic visits | Pharmacy use             | Hospitalization  | Outpatient clinic visits | Pharmacy use             |
| <18.5            | 4.65             | (0.46–47.03)             | 1.22                     | (0.72–1.96)          | 4.93                     | (0.52–46.38)             | 0.83                     | (0.51–1.33)          | 7.12                     | (0.77–66.02)             | 0.82                     | (0.51–1.32)          | 0.90                     | (0.82–0.99)             |
| 18.5–22.9        | Reference        |                         |                          |                        |                          |                          |                          |                        |                          |                          |                          |                        |                          |                          |
| 23–24.9          | 1.17             | (0.46–2.96)             | 0.89                     | (0.71–1.06)          | 1.02                     | (0.42–2.50)             | 0.95                     | (0.76–1.12)          | 1.26                     | (0.50–3.16)             | 0.95                     | (0.79–1.15)          | 0.92                     | (0.88–0.96)             |
| ≥25              | 0.70             | (0.60–0.89)             | 0.73                     | (0.33–1.94)          | 0.79                     | (0.72–1.05)             | 0.87                     | (0.72–1.05)          | 1.02                     | (0.41–2.52)             | 0.87                     | (0.79–0.86)          | 0.82                     | (0.79–0.86)             |
| P for trend      | 0.25             |                          | 0.01                     |                          | 0.25                     |                          | 0.50                     |                          | 0.01                     |                          | 0.52                     |                          | <0.01                    |                          |

Notes: *Adjusted by age, sex, smoking pack-year, and FEV₁ (%).
Abbreviations: IRR, incidence rate ratio; BMI, body mass index.

In a general population, obesity can induce systemic comorbidities, such as diabetes mellitus and cardiovascular disease, and is known to correlate with increased mortality relative to that associated with normal weight.22 Paradoxically, increased weight and mild obesity have been reported to yield survival benefits among patients with COPD.6,7,23 Cao et al23 reported a meta-analysis of 22 studies with 21,150 individuals to evaluate the association between BMI and mortality in COPD. Relative to normal weight, decreased weight was associated with increased mortality (relative risk [RR] 1.34; 95% CI 1.01–1.78), whereas increased weight (RR 0.47; 95% CI 0.33–0.68) and obesity (RR 0.59; 95% CI 0.38–0.91) were associated with reduced mortality. Moreover, our results suggest a protective effect of increasing BMI against poor outcomes among COPD patients with mild to moderate airflow obstruction. The exact pathophysiology underlying this protective effect is poorly understood, despite increasing interest. However, the possible mechanisms underlying the positive effect of decreased weight on the unfavorable outcomes of COPD include sarcopenia and muscle weakness,24,25 inactivity,26 and systemic inflammation.26,27

In our study, obese patients tended to use medical care less frequently and had fewer medical expenses, compared with normal-weight patients. This relationship was more apparent among patients with relatively more severe airflow limitations. Our results are consistent with a previous study on the natural course of asymptomatic COPD patients with mild to moderate airflow obstruction, which suggested that patients with risk factors (eg, old age, lower BMI, reduced function, low EQ-5D scores) should be diagnosed and treated before disease progression.28

In practice, it is difficult to assess the prevalence of asymptomatic COPD – either alone or according to BMI – as these patients are not expected to visit hospitals frequently and clinicians might not conduct a spirometry screening for mild respiratory symptoms. Our study was made possible by access to large, well-designed national databases that encompass both demographic and functional test results. Furthermore, there was no missing data, and the NHI database allowed us to assess medical expenses. One interesting and valuable strength of this nationwide study is the potential for a long-term follow-up evaluation of the same group of patients. We could therefore assess the impact of BMI on progression to overt, symptomatic COPD from mild, asymptomatic COPD.

It remains unclear whether treating asymptomatic COPD could delay disease progression and change the outcome. However, the pathogenesis and continuum of disease progression suggest that early treatment might influence the course of disease. Despite evidence suggesting that obese patients with COPD have a better prognosis, we do not recommend being obese, as physical exercise and active rehabilitation are essential to improving exercise capacity, and proper nutritional support might play an important role. The finding that physically inactive and extremely obese patients with COPD had worse outcomes was very clear.

This may be the first study assessing the impact of BMI on the outcome of asymptomatic COPD patients with mild to moderate airflow obstruction, but several limitations must be considered. First, although we used various nationwide data, this was a retrospective study. Second, the KNHANES dataset does not include COPD-specific QOL assessment tools. Third, to clarify COPD diagnosis, COPD was defined by spirometry and smoking status. This might have resulted in male predominance of the sample population. Moreover, we did not consider biomass exposure-related airflow limitations because they are not easily quantified and might lead to a risk of over-diagnosis of COPD as well. Fourth, COPD was not defined by a fixed, irreversible airflow limitation because post-bronchodilator (BD) spirometric values were
not available. However, a previous large cohort study found that pre- and post-BD results were interchangeable.\textsuperscript{24,25} Moreover, the definition of exacerbation we used has not been validated. However, the same definition of exacerbation was applied in other studies.\textsuperscript{29,30} Lastly, all study participants were Korean; therefore, our results may not be generalizable to other populations. However, previous studies on BMI and COPD were mostly conducted in Western populations. Accordingly, Asia-specific data are scarce. Therefore, this study provides valuable information on the role of BMI in Asian patients with COPD.

**Conclusion**

In summary, we found a negative influence of decreased weight on the clinical outcomes of COPD patients with mild to moderate airflow obstruction, whereas an increasing BMI (relative to normal BMI) had no significant negative effect on prognosis.

**Disclosure**

The authors report no conflicts of interest in this work.

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## Supplementary materials

### Table S1 Subgroup analysis of the risk of exacerbation according to the severity of airflow limitation

| IRR (95% CI) | BMI (kg/m²) group | P-value |
|--------------|-------------------|---------|
|              | <18.5             | 18.5–22.9 | 23–24.9 | ≥25 |
| FEV₁ ≥80% predicted |                   |         |         |     |
| Crude IRR    | 1.01 (0.61–1.67)  | Reference | 0.90 (0.74–1.08) | 0.88 (0.72–1.07) | 0.51 |
| Adjusted IRRα | 0.88 (0.54–1.45)  | 0.89 (0.75–1.07) | 0.98 (0.81–1.19) | 0.65 |
| Adjusted IRRβ | 0.89 (0.54–1.46)  | 0.90 (0.75–1.07) | 0.98 (0.81–1.20) | 0.66 |
| 50% ≤ FEV₁ < 80% predicted |                   |         |         |     |
| Crude IRR    | 1.16 (0.77–1.74)  | Reference | 0.98 (0.82–1.17) | 0.88 (0.74–1.04) | 0.33 |
| Adjusted IRRα | 0.99 (0.67–1.47)  | 1.02 (0.86–1.22) | 0.99 (0.84–1.16) | 0.98 |
| Adjusted IRRβ | 0.99 (0.67–1.47)  | 1.02 (0.86–1.21) | 0.98 (0.84–1.16) | 0.98 |

**Notes:** αAdjusted by age, sex, and FEV₁ (%). βAdjusted by age, sex, smoking pack-year, and FEV₁ (%).

**Abbreviations:** IRR, incidence rate ratio; FEV₁, forced expiratory volume in 1 s; BMI, body mass index.

### Figure S1 Cumulative chronic obstructive pulmonary disease (COPD)-related medical costs per person-year stratified by body mass index (BMI).

**Note:** Medical costs are presented in US dollars (USD) with an exchange rate of 1 USD equal to 1,140 Korean won (exchange rate on May, 2017).
Table S2 Subgroup analysis of the risk of chronic obstructive pulmonary disease (COPD)-related health-care utilization during a 5-year period according to the forced expiratory volume in 1 s (FEV₁)

| IRR (95% CI) | BMI (kg/m²) |  |  |  |  |
|--------------|-------------|--------|--------|--------|--------|
|              | <18.5       | 18.5–22.9 | 23–24.9 | ≥25    |  |
| **FEV₁ ≥80% predicted** |                   |               |        |        |  |
|                | Reference    | Reference    | Reference | Reference |  |
| Crude IRR       |             |             |          |          |  |
| Hospitalization | 7.20 (0.20–265.07) | 1.24 (0.33–4.66) | 0.57 (0.14–2.32) | 0.33    |
| Outpatient clinic visits | 0.52 (0.28–0.96) | 1.04 (0.85–1.26) | 0.89 (0.72–1.10) | 0.11    |
| Pharmacy use     | 0.33 (0.15–0.73) | 1.04 (0.79–1.37) | 0.90 (0.67–1.19) | 0.08    |
| Adjusted IRR     |             |             |          |          |  |
| Hospitalization | 10.04 (0.29–348.56) | 1.08 (0.30–3.92) | 0.58 (0.14–2.38) | 0.20    |
| Outpatient clinic visits | 0.45 (0.24–0.82) | 1.05 (0.87–1.27) | 0.97 (0.79–1.19) | 0.05    |
| Pharmacy use     | 0.27 (0.12–0.59) | 1.05 (0.80–1.36) | 1.00 (0.76–1.33) | 0.03    |
| **50% ≤ FEV₁ < 80% predicted** |                   |               |        |        |  |
|                | Reference    | Reference    | Reference | Reference |  |
| Crude IRR       |             |             |          |          |  |
| Hospitalization | 2.57 (0.15–42.62) | 1.70 (0.50–5.77) | 0.84 (0.27–2.63) | 0.63    |
| Outpatient clinic visits | 1.58 (0.98–2.55) | 0.86 (0.70–1.07) | 0.73 (0.59–0.89) | <0.01   |
| Pharmacy use     | 1.53 (0.79–2.96) | 0.78 (0.59–1.04) | 0.64 (0.49–0.84) | <0.01   |
| Adjusted IRR     |             |             |          |          |  |
| Hospitalization | 2.68 (0.18–40.95) | 1.36 (0.41–4.48) | 1.21 (0.39–3.73) | 0.86    |
| Outpatient clinic visits | 1.22 (0.79–1.90) | 0.89 (0.72–1.09) | 0.86 (0.71–1.05) | 0.25    |
| Pharmacy use     | 1.14 (0.61–2.14) | 0.83 (0.63–1.09) | 0.78 (0.60–1.01) | 0.22    |
| Adjusted IRR     |             |             |          |          |  |
| Hospitalization | 3.14 (0.21–46.92) | 1.64 (0.49–5.54) | 1.37 (0.44–4.21) | 0.73    |
| Outpatient clinic visits | 1.21 (0.78–1.87) | 0.86 (0.70–1.05) | 0.85 (0.70–1.03) | 0.16    |
| Pharmacy use     | 1.11 (1.00–1.23) | 0.79 (0.75–0.84) | 0.73 (0.69–0.78) | <0.01   |

Notes: *Adjusted by age, sex, and FEV₁ (%). †Adjusted by age, sex, smoking pack-year, and FEV₁ (%).

Abbreviations: IRR, incidence rate ratio; BMI, body mass index; FEV₁, forced expiratory volume in 1 s.