

Effect of Body Mass Index on Lung Function in Chinese Patients with Chronic Obstructive Pulmonary Disease: A Multicenter Cross-Sectional Study

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Objective: The aim of this study was to explain “obesity paradox” in chronic obstructive pulmonary disease (COPD) by evaluating the effect of body mass index (BMI) on lung function in Chinese patients with COPD.

Methods: A total of 1644 patients diagnosed with COPD were recruited from four Chinese tertiary hospitals and were divided into four groups including underweight, normal weight, overweight and obese according to BMI classification standard. The medical data of these patients were collected and used for the multiple linear regression analyses.

Results: After adjustment for age, sex, educational level, economic status, smoking status, alcohol consumption, duration of COPD history, events of acute exacerbation in previous year, hypertension, diabetes mellitus, cardiovascular disease, cerebrovascular disease and osteoporosis, BMI had a curvilinear correlation with the forced expiratory volume in the first second (FEV₁) in patients with Global Initiative for Obstructive Lung Disease (GOLD) 1–2 grade (first-order coefficient β , 0.09; 95% CI, 0.03–0.16; second-order coefficient β , –0.002; 95% CI, –0.003–0.001; $P < 0.01$). However, BMI had a positive correlation with FEV₁ in patients with GOLD 3–4 grade (β , 0.01; 95% CI, 0.008–0.017; $P < 0.01$) when BMI was used as a quantitative variable. When BMI was used as a qualitative variable, only FEV₁ in overweight group with GOLD 1–2 grade was significantly higher than that of normal weight group ($P < 0.01$). Interestingly, both overweight and obese groups had higher FEV₁ in GOLD 3–4 grade compared with normal weight group (β , 0.06; 95% CI, 0.02–0.11; β , 0.11; 95% CI, 0.04–0.18; $P < 0.01$). The effect of BMI on predicted percentage of FEV₁ (FEV₁%) was similar to that of FEV₁ in different GOLD grades.

Conclusion: Obesity only had a protective effect on lung function in COPD patients with GOLD 3–4 grade rather than GOLD 1–2 grade.

Trial Registry: ClinicalTrials.gov, No.: NCT 03182309, URL: www.clinicaltrials.gov.

Keywords: overweight, obesity

Introduction

Chronic obstructive pulmonary disease (COPD), characterized by restricted airflow, is a common disease in the older population. The prevalence of COPD was 13.7% for people over 40 years of age in China.¹ Obesity characterized by excessive accumulation of body fat can be evaluated by body mass index (BMI). Obesity is a common comorbidity of COPD, with estimates ranging from 6% to 54%.^{2–6} However, the prevalence of obesity in Chinese patients with COPD remains unclear.

The severity of COPD depends on degree of airflow limitation reflected by impaired lung function of patients. Body weight has an independent effect on lung function apart from tobacco exposure.⁷ Many studies indicated that obesity was associated with decreased lung function in the general population because of its effect on gas exchange, respiratory mechanics, muscular endurance and breath control.^{8,9} In addition, obesity could also contribute to decreased lung function through chronic low-grade inflammation.¹⁰ But less attention has been given to the association between obesity and lung function in patients with different diseases. Obesity is more likely associated with metabolic syndrome, type 2 diabetes mellitus, cardiovascular disease, obstructive sleep apnea and cancer, which all result in an increased risk of death.^{11,12} Allison et al's study from a multicenter prospective cohort suggested that obesity in patients with COPD might contribute to worse COPD-related health consequences such as acute exacerbation of COPD and dyspnea scores.¹³ Therefore, it is believed that obesity in patients with COPD should be treated with weight loss through exercise, which is conducive to improved lung function and reduced complications.¹⁴ However, there have been studies that suggested that "overweight" and obesity could prolong survival in patients with chronic heart or kidney diseases.^{15,16} This phenomenon is referred to as the "obesity paradox".¹⁵ The obesity paradox also existed in COPD patients because the mortality and exacerbation frequency of obese patients with COPD was lower than that of normal weight patients.¹⁷ It seems that obesity has a protective effect on the prognosis of COPD. Because of the obesity paradox, whether to treat obesity in COPD remains controversial for clinicians. The weight loss may improve cardiovascular outcomes but at the same time, the deteriorating respiratory outcomes will increase mortality of patients.¹⁸ Because the reference standards for obesity are different, it is necessary to explore the relationship between different BMIs and lung function of COPD patients in a Chinese population.

In this study, we tried to explain the obesity paradox by evaluating the effect of BMI on lung function of different Global Initiative for Obstructive Lung Disease (GOLD) grades in Chinese patients with COPD by multiple linear regression analyses.

Methods

Ethics Approval and Data Sharing Statement

This study was part of a national key research and development program about COPD, which was a national

clinical registration study initiated in January 2016 (Clinical Trials ID: NCT 03182309). The procedure of this study was reviewed and approved by the Ethics Committee of Renmin Hospital of Wuhan University (No.2017K-C014). This study was performed in accordance with the Declaration of Helsinki. Every patient included in this study was informed about the purpose of the study and written consent was signed. Regarding the data sharing, because this project will last for 5 years and multiple research centers participated, currently this project is not fully completed, our present study is only one part of this project, which focused on studying the effect of BMI on lung function in COPD patients based on current clinical data, so these data cannot be shared until the project is fully completed.

Subjects

All participants were enrolled in the primary cohort from four Chinese tertiary hospitals. All the participants were more than 40 years of age and were not pregnant, and had been diagnosed with COPD using a spirometer. The participants who had a history of bronchial asthma, bronchiectasis, pulmonary fibrosis, tuberculosis, cancer, renal dysfunction, neuro-muscular disease, or mental illness were excluded. The data of the patients who refused to complete lung function test, or if their results were considered as unqualified by the operators because the patients could not fully cooperate with lung function test, were considered as incomplete data and were excluded ($n=11$). Finally, 1644 participants met the inclusion criteria for the study and were classified into four groups including underweight ($\text{BMI}<18.5 \text{ kg/m}^2$), normal weight ($\text{BMI}:18.5\text{--}23.9 \text{ kg/m}^2$), overweight ($\text{BMI}:24\text{--}27.9 \text{ kg/m}^2$) and obese ($\text{BMI}\geq 28 \text{ kg/m}^2$) according to BMI classification standard for Chinese population.¹⁹

Data Collection

A post-bronchodilator FEV_1 /forced vital capacity (FVC) ratio of <0.7 was diagnosed as COPD according to the GOLD guidelines.²⁰ Lung function parameters, including the FVC, the FEV_1 , the predicted percentage of FEV_1 ($\text{FEV}_1\%$) and the $\text{FEV}_1/\text{FVC}\%$, were measured using a spirometer (Jaeger MasterScreen Body, Germany) after using bronchodilator.²¹ The other clinical data of participants including the general information, modified Medical Research Council (mMRC) dyspnea scale comorbidities such as hypertension, diabetes mellitus, cardiovascular disease and cerebrovascular disease, COPD history, events of COPD exacerbation in the previous year were collected

using the same standardized protocol from the multicenter study. Data could only be downloaded online by a third party, a special statistical expert who did not participate in the clinical study.

Statistical Analysis

Differences between multiple groups were compared using the Chi-squared test for categorical variables, the one-way ANOVA for normal quantitative variables and the Wilcoxon rank sum test for non-normal quantitative variables. The multiple linear regression models were used to determine the associations between different BMIs and lung function. Age, sex, educational level, economic status, smoking status, alcohol consumption, duration of COPD history, events of acute exacerbation in previous year, hypertension, diabetes mellitus, cardiovascular disease, cerebrovascular disease and osteoporosis were considered as covariates. Due to the curvilinear relation between the BMI and FEV₁ during the quantitative data exploration, a second-order term of BMI was added in the model when BMI was used as a quantitative variable in the formal analysis. If the coefficient of the second-order term was not significant, then only the regular first-order term of BMI was retained. Collinearity of independent variables was also assessed and was negative. All analyses were performed using SAS 9.4 (SAS Institute Inc, Cary, NC) and a $P < 0.05$ is considered statistically significant.

Results

Demographic Characteristics

The baseline characteristics of patients in this study are shown in Table 1. Among the 1644 COPD patients included in the study, a total of 198 patients were underweight (12.0%), 757 patients were normal weight (46.0%), 460 patients were overweight (28.0%) and 229 patients were obese (14.0%). There were no significant differences in gender or economic status in these four groups ($P > 0.05$).

Clinical Characteristics

There were significant differences in age, neck circumference, degree of education, smoking status, acute exacerbation of COPD in previous year, mMRC dyspnea scale, complication (hypertension, diabetes mellitus and cardiovascular disease), duration of COPD history, red blood cell count and hemoglobin in the four groups ($P < 0.05$) (Table 1). The FVC, FEV₁, FEV₁% and FEV₁/FVC% of

the patients in overweight and obese COPD groups were significantly higher than that of the patients in underweight and normal weight COPD groups ($P < 0.01$) (Figure 1A–D). However, there were no statistical differences in FVC, FEV₁, FEV₁% of the patients in overweight and obese COPD groups ($P > 0.05$).

The Effect of BMI on Lung Function of Patients with COPD

To investigate the effects of BMI on lung function in COPD patients, we analyzed the correlation between BMI and lung function parameters by multiple linear regression. As shown in Figure 2A, when BMI was used as a quantitative variable, BMI had a significantly curvilinear correlation (quadratic polynomial curve) with the FEV₁ (first-order coefficient β , 0.11; 95% CI, 0.06–0.16; second-order coefficient β , –0.002; 95% CI, –0.003–0.001; $P < 0.01$) after adjustment for age, sex, educational level, economic status, smoking status, alcohol consumption, duration of COPD history, events of acute exacerbation in previous year, hypertension, diabetes mellitus, cardiovascular disease, cerebrovascular disease and osteoporosis. When BMI was used as a qualitative variable and the normal weight group was designated as the reference level, both the overweight group (β , 0.22; 95% CI, 0.15–0.28; $P < 0.01$) and the obese group (β , 0.27; 95% CI, 0.18–0.36; $P < 0.01$) had higher FEV₁ levels than the normal weight group after adjusting for the previously mentioned covariates (Figure 2B). The effect of BMI on FEV₁% or FVC was similar to that of BMI on FEV₁ (Figure 2C and D).

The Effect of BMI on Lung Function in GOLD Grades

We further investigated whether lung function in GOLD 1–2 grade and GOLD 3–4 grade was affected by BMI using multiple linear regression analysis. As shown in Figure 3A, when BMI was used as a quantitative variable, BMI had a significantly curvilinear correlation with FEV₁ in GOLD 1–2 grade (first-order coefficient β , 0.09; 95% CI, 0.03–0.16; second-order coefficient β , –0.002; 95% CI, –0.003–0.001; $P < 0.01$) after adjustment for age, sex, educational level, economic status, smoking status, alcohol consumption, duration of COPD history, events of acute exacerbation in previous year, hypertension, diabetes mellitus, cardiovascular disease, cerebrovascular disease and osteoporosis. However, when BMI was used as

Table 1 Baseline Characteristics of COPD Patients with Different BMIs (n=1644)

Characteristics	Underweight (n=198)	Normal Weight (n=757)	Overweight (n=460)	Obese (n=229)	P value
Sex, male, n (%)	174 (87.9)	653 (86.3)	378 (82.2)	179 (78.2)	0.13
Age, yr	67.0±9.2	66.4±8.8	64.9±8.7	62.6±10.2	0.01
Neck Circumference, cm	35.1±2.7	37.4±2.6	39.5±3.0	42.3±4.3	<0.01
Degree of education, n (%)					0.03
Under college	182 (92.0)	662 (87.5)	385 (83.7)	198 (86.5)	
Above college	16 (8.0)	95 (12.5)	75 (16.3)	31 (13.5)	
Economic status, n (%)					0.51
Higher income	0 (0.0)	12 (1.6)	8 (1.7)	2 (0.9)	
Middle income	19 (9.6)	74 (9.8)	48 (10.4)	18 (7.9)	
Lower income	179 (90.4)	671 (88.6)	404 (87.9)	209 (91.2)	
Smoking status, n (%)					<0.01
Never	35 (17.7)	134 (17.7)	105 (22.8)	51 (22.3)	
Former	50 (25.3)	188 (24.8)	130 (28.3)	75 (32.7)	
Current	113 (57.0)	435 (57.5)	225 (48.9)	103 (45.0)	
Alcohol consumption, n (%)					0.05
Never	129 (65.2)	461 (60.9)	262 (57.0)	136 (59.4)	
Former	20 (10.1)	48 (6.3)	49 (10.6)	21 (9.2)	
Current	49 (24.7)	248 (32.8)	149 (32.4)	72 (31.4)	
FVC, L	2.35±0.81	2.60±0.88	2.88±0.97	2.80±1.02	<0.01
FEV ₁ , L	1.05±0.57	1.29±0.62	1.56±0.72	1.67±0.74	<0.01
FEV ₁ % Predicted	41.65±18.71	50.75±22.10	59.22±23.23	62.05±22.20	<0.01
FEV ₁ /FVC% Predicted	44.95±12.67	49.14±12.73	55.75±12.16	58.88±9.93	<0.01
Severity of COPD, n (%)					<0.01
GOLD 1	12 (6.1)	96 (12.7)	105 (22.8)	53 (23.1)	
GOLD 2	38 (19.2)	241 (31.8)	172 (37.5)	103 (45.0)	
GOLD 3	95 (48.0)	286 (37.8)	140 (30.4)	60 (26.2)	
GOLD 4	53 (26.7)	134 (17.7)	43 (9.3)	13 (5.7)	
mMRC dyspnea scale	2.02±0.96	1.79±0.96	1.58±0.97	1.55±1.00	<0.01
Hypertension, n (%)	43 (21.7)	257 (33.9)	218 (47.4)	147 (64.2)	<0.01
Diabetes mellitus, n (%)	7 (3.5)	60 (7.9)	69 (15.0)	51 (22.3)	<0.01
Cardiovascular disease, n (%)	34 (17.2)	163 (21.5)	131 (28.5)	1 (39.7)	<0.01
Cerebrovascular disease, n (%)	10 (5.1)	73 (9.6)	50 (10.9)	24 (10.5)	0.12
Osteoporosis, n (%)	3 (1.5)	24 (3.2)	14 (3.0)	4 (1.7)	0.45
COPD history, month, median (Q1,Q3)	60 (24, 120)	54 (14, 120)	36 (2, 84)	12 (0, 60)	<0.01
Events in previous year					
Acute exacerbation, n (%)	142 (71.7)	441 (58.3)	250 (54.3)	101 (44.1)	<0.01
NIV, n (%)	11 (5.6)	51 (6.7)	24 (5.2)	23 (10.0)	0.10
Blood cell analysis					
WBC, ×10 ⁹ /L	7.35±3.20	7.52±2.95	7.48±2.48	7.39±2.38	0.86
RBC, ×10 ¹² /L	4.30±0.56	4.50±0.61	4.56±0.54	4.73±0.59	<0.01
HB, g	129.2±16.16	136.7±17.5	140.2±16.17	143.0±17.4	<0.01
PLT, ×10 ⁹ /L	218.02±83.16	221.50±77.22	211.85±69.49	216.54±53.92	0.07

Notes: Continuous variables which are obviously skewed were presented as median (Q1, Q3), other continuous variables were described as means ±SD. Categorical variables were expressed as percentages. P value for the difference between the groups.

Abbreviations: BMI, body mass index; COPD, chronic obstructive pulmonary disease; FVC, forced vital capacity; FEV₁, forced expiratory volume in the first second; GOLD 1, FEV₁ ≥80% predicted; GOLD 2, FEV₁=50–79% predicted; GOLD 3, FEV₁=30–49% predicted; GOLD 4, FEV₁ <30% predicted; mMRC, modified Medical Research Council; NIV, non-invasive ventilation; WBC, white blood cells; RBC, red blood cells; HB, hemoglobin; PLT, platelets.

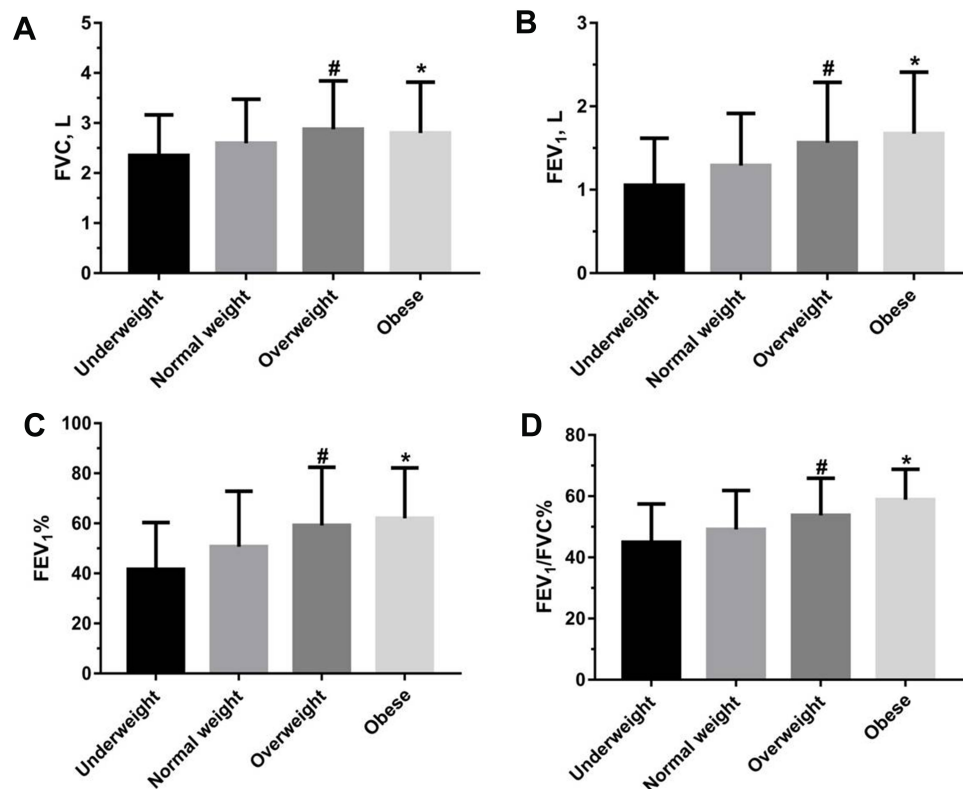


Figure 1 The differences in FVC, FEV₁, FEV₁% and FEV₁/FVC% of the four groups of patients with COPD (A–D). Statistical analysis was performed using one-way ANOVA test. The data were represented as mean values $\pm$ SD. [#] $p < 0.05$ indicating the statistical difference between overweight group with COPD and underweight or normal weight group. ^{*} $p < 0.05$ indicating the statistical difference between obese group with COPD and underweight or normal weight group.

Abbreviations: COPD, chronic obstructive pulmonary disease; FVC, forced vital capacity; FEV₁, forced expiratory volume in the first second; FEV₁%, predicted percentage of FEV₁; FEV₁/FVC%, percentage of FEV₁ and FVC.

a qualitative variable, FEV₁ in overweight COPD patients was significantly higher than that of normal weight patients after adjustment for previously mentioned covariates (β , 0.10; 95% CI, 0.03–0.18; $P < 0.01$) (Figure 3B). Interestingly, among the GOLD 3–4 grade patients, BMI did not have the quadratic polynomial curvilinear relationship but had straight linear positive correlation (β , 0.01; 95% CI, 0.01–0.02; $P < 0.01$) when BMI was used as a quantitative variable (Figure 3C). Moreover, when BMI was used as a qualitative variable, both overweight and obese had higher FEV₁ in GOLD 3–4 grade compared with normal weight after adjustment for previously mentioned concomitant variable (β , 0.06; 95% CI, 0.02–0.11; β , 0.11; 95% CI, 0.04–0.18; $P < 0.01$) (Figure 3D). BMI had similar effect on FEV₁% in GOLD grades of COPD patients (Figure 4A and B). When BMI was used as a quantitative variable, BMI had a significantly curvilinear correlation with FVC both in GOLD 1–2 grade (first-order coefficient β , 0.15; 95% CI, 0.06–0.23; second-order coefficient β , –0.003; 95% CI, –0.005–0.001; $P < 0.01$) and GOLD 3–4 grade (first-order coefficient β , 0.08; 95% CI,

0.01–0.15; second-order coefficient β , –0.001; 95% CI, –0.003–0.000; $P < 0.05$) after adjustment for previously mentioned variable (Figure 4C and D).

Discussion

Obesity was not only highly prevalent but also occurred more in COPD patients than in the general population.¹⁷ Our multicenter study showed that the prevalence of obesity in COPD was 14% in China. The prevalence of obesity in COPD reported by many research groups has variability.^{2–6} The difference mainly resulted from the difference in definition of obesity and the relatively lower prevalence of obesity in the general population of China. In addition, we found that obesity was most prevalent in COPD patients with GOLD 2 grade (45%) and least prevalent in COPD patients with GOLD 4 grade (5.7%), which were similar to the results reported by Steuten et al.⁵

In clinical practice, “underweight” is one of the most important concerns in COPD care. Being underweight might increase mortality risk of COPD patients as shown

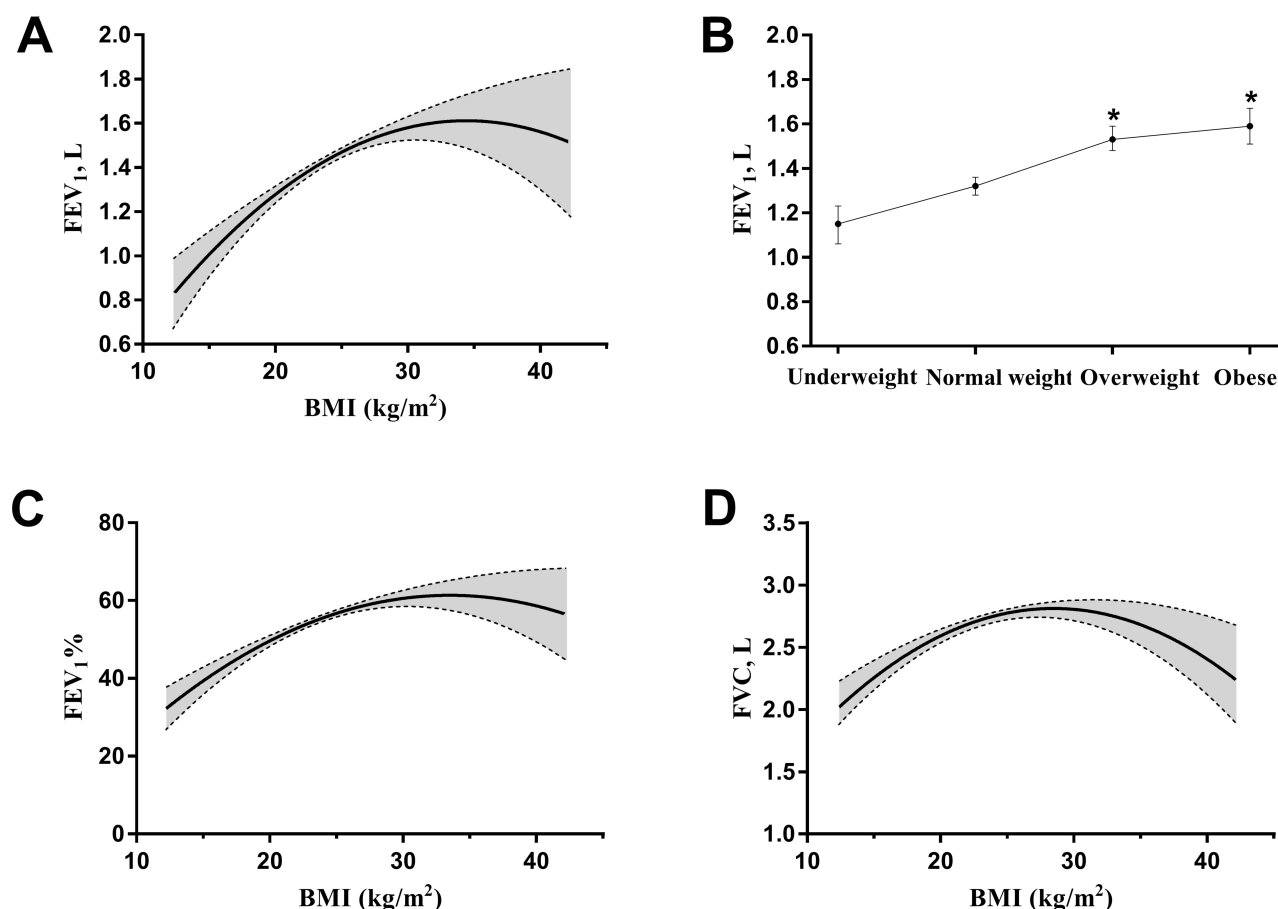


Figure 2 The effect of BMI on lung function in the patients with COPD. Statistical analysis was performed using multiple linear regression analysis. (A and B) BMI used as the quantitative and qualitative variable to assess the relation between BMI and FEV₁. (C and D) BMI used as the quantitative variable to assess the relation between BMI and FEV₁% or FVC. * $p < 0.05$ vs normal weight group.

Abbreviations: BMI, body mass index; COPD, chronic obstructive pulmonary disease; FEV₁, forced expiratory volume in the first second; FEV₁%, predicted percentage of FEV₁; FVC, forced vital capacity.

in previous studies.^{18,22} In our present study, we found that underweight patients with COPD had lower lung function, higher dyspnea scores and acute exacerbation frequency of COPD in previous year than obese patients with COPD. This result is consistent with that of another retrospective study of 774 COPD patients in China.²³ The phenomenon was explained by the fact that the underweight patients might have malnutrition, low muscularity and weak resistance against respiratory infection, all of which increased the risk of COPD.^{24–26} In addition, we found that underweight patients with COPD were older and their disease duration was longer, mainly because these patients had an increased frequency of exacerbation, leading to a faster decline in lung function and a poorer quality of life.²⁷

The most significant finding in our present study was that being overweight only had a positive correlation with lung function (FEV₁ and FEV₁%) in COPD patients with GOLD 1–2 grade, but both being overweight and

obese had a positive correlation with lung function in COPD patients with GOLD 3–4 grade. Our results indicated that obesity had a protective effect on lung function in GOLD 3–4 grade rather than GOLD 1–2 grade, which may partially explain the obesity paradox in COPD. A meta analysis including 62 studies showed that “overweight” or obesity may be detrimental to lung function in adults with or without asthma and the damage in asthma was less severe than in healthy people.²⁸ But the studies included in the meta analysis had mainly focused on subjects with or without asthma. Moreover, COPD is characterized by irreversible airflow obstruction, which is different from asthma. Our study indicated that obesity not only had no protective effect, but that further increased BMI, such as morbid obesity, may worsen lung function in GOLD 1–2 grade, which suggested that in the early stages of COPD, nutritional status was critical to maintaining the rate of decline in lung function, while excessive weight

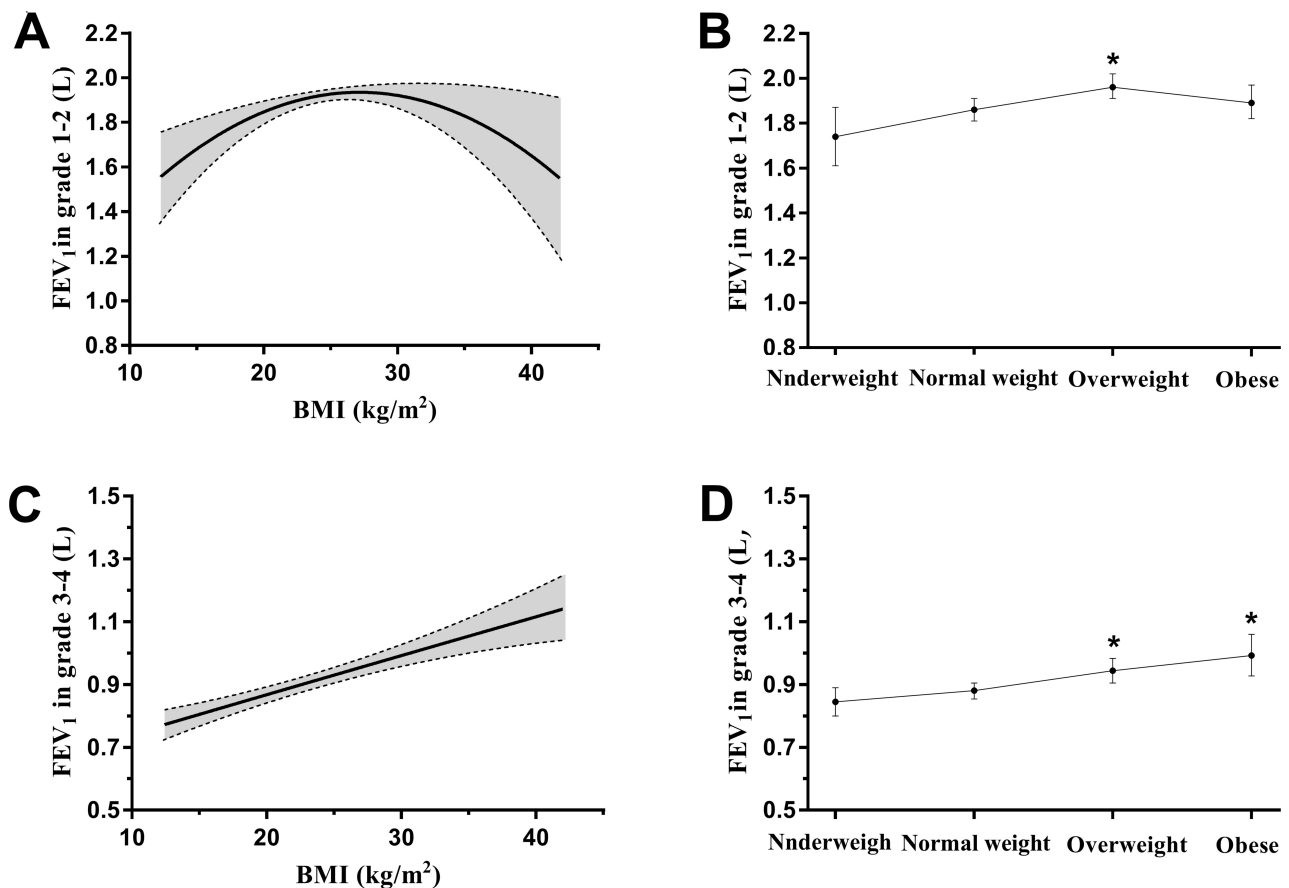


Figure 3 The effect of BMI on FEV₁ in GOLD grade of COPD patients. Statistical analysis was performed using multiple linear regression analysis. (A and B) BMI used as the quantitative and qualitative variable to assess the relation of BMI and FEV₁ in GOLD 1–2 grade. (C and D) BMI used as the quantitative and qualitative variable to assess the relation between BMI and FEV₁ in GOLD 3–4 grade. **p*<0.05 indicating the significant difference compared to normal weight group.

Abbreviations: BMI, body mass index; COPD, chronic obstructive pulmonary disease; FEV₁, forced expiratory volume in the first second; GOLD, global initiative for obstructive lung disease.

gain may be counter-productive. In addition to BMI, lung function was also influenced by gender, age, history and acute exacerbation frequency of COPD. Studies by others also showed that “overweight” was highly prevalent and had the lowest mortality risk in patients with of COPD,¹⁸ which was consistent with our results. The patients in GOLD 3–4 grade were liable to complicate with weight loss, mainly muscle loss, which could result in respiratory hypofunction and further worsen lung function and increase mortality. Obesity was not beneficial to lung function in the general population. Interestingly, our study found that increased BMI had a protective effect on lung function in GOLD 3–4 grade. Obese patients not only have more body fat, but also increased muscle mass. Nutritional status, which is of significant value for prognosis of COPD, played an important role in lung function in later stage COPD patients.¹⁸ Sami et al’s cross-sectional study that included 72 male patients with COPD also

showed that increased BMI had a positive relation with FEV₁ in GOLD 4 grade.²⁹ Landbo et al suggested that the relative risk for mortality seemed decreased in overweight and obese patients with COPD GOLD 3–4 grade, while it was increased in those with GOLD 1–2 grade.¹⁸ In addition, we found that the effect of BMI on FVC in patients with COPD had a curvilinear correlation in GOLD 1–2 grade and GOLD 3–4 grade. Our results suggested that increasing BMI to a certain threshold might impair FVC in COPD patients, which might be related to airflow limitation.

Obesity tends to cause many comorbidities such as cardiovascular and metabolic diseases.¹¹ The comorbidities are also relatively common in COPD.³⁰ In our present study we found that in COPD patients who were obese, rates of hypertension, diabetes mellitus and cardiovascular disease were significantly higher than in those with COPD with normal weight, which was in line with previous

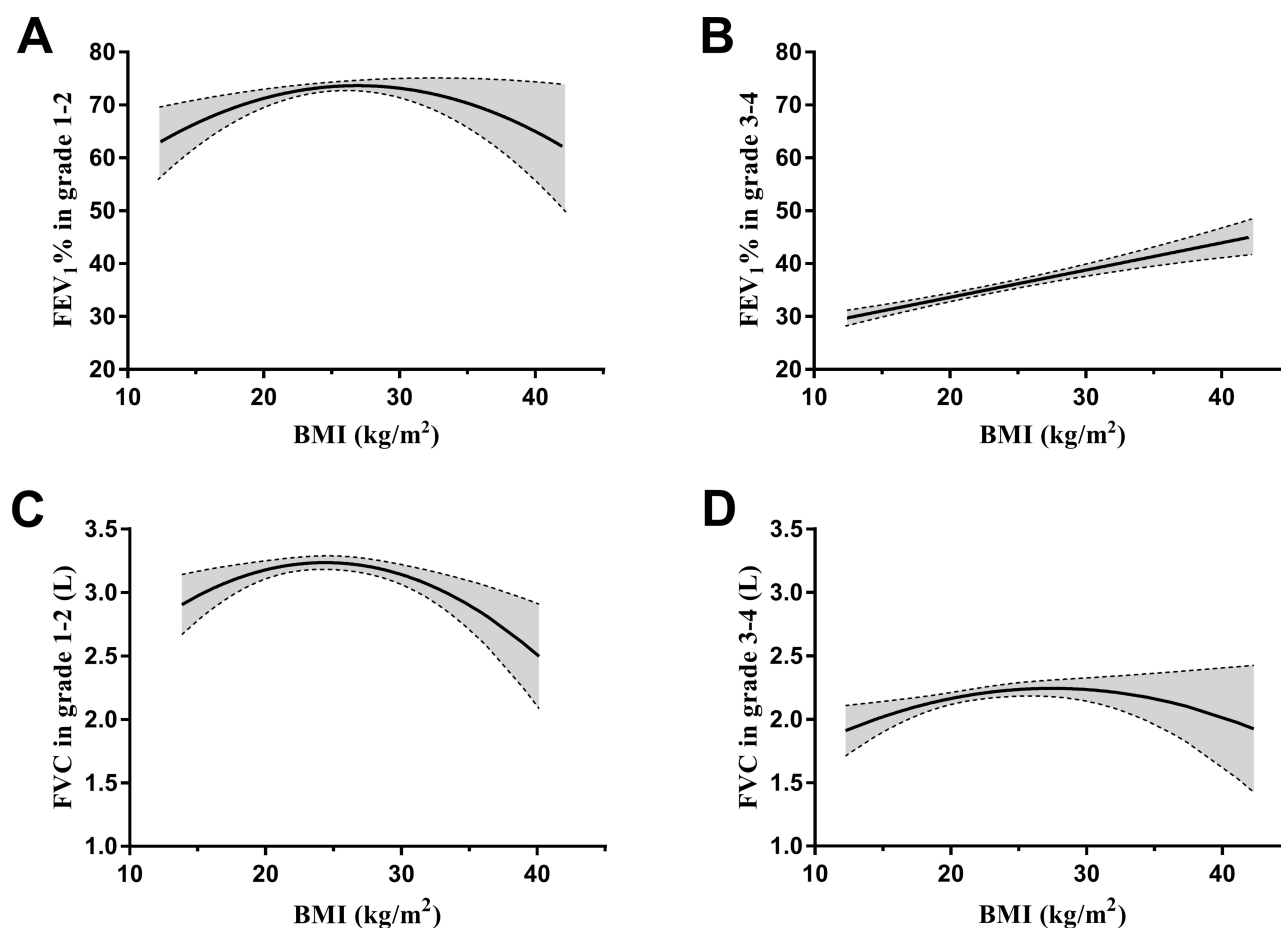


Figure 4 The effect of BMI on FEV₁% and FVC in GOLD grade of COPD patients. Statistical analysis was performed using multiple linear regression analysis. The curve and the 95% CI area were obtained from the regression equation by fixing each covariate at a certain level. (A and B) BMI used as the quantitative variable to assess the relation of BMI and FEV₁% in GOLD 1–2 grade and GOLD 3–4 grade. (C and D) BMI used as the quantitative variable to assess the relation between BMI and FVC in GOLD 1–2 grade and GOLD 3–4 grade.

Abbreviations: BMI, body mass index; COPD, chronic obstructive pulmonary disease; FEV₁%, predicted percentage of forced expiratory volume in the first second; FVC, forced vital capacity; GOLD, global initiative for obstructive lung disease.

studies.^{30,31} The mechanism may be related to increasing BMI, aging, smoking and inflammatory status in obesity. The comorbidities may explain the reason why obese patients with COPD have higher hospitalization rates.³² Therefore early detection and management of co-existing diseases may benefit quality of life and reduce mortality.

Although our study was not a prospective study, we found a low incidence of acute exacerbations in obese COPD patients in previous year, suggesting that obesity held some protective significance to COPD. In a recent study reported by Sun et al, they found that high BMI had a protective effect for rate of FEV₁ decline.³³ The possible mechanism was that obese patients with COPD might receive medical attention earlier than underweight patients, which was beneficial to maintain the progressive decline of lung function. In terms of physical

mechanisms, obese COPD patients may benefit from obesity by reducing lung hyperinflation which may have a beneficial effect on lung function.³⁴ However, in another meta analysis, the results indicated that the survival advantage seems to diminish, when the BMI is above 32 kg/m².¹⁷ Therefore, cut off point of BMI for advantages in different GOLD grades needs to be further elucidated, because of the intricate relation between BMI and lung function in patients with COPD.

We acknowledge that there are some limitations in this study. First, in this study, we used only a single measure method of BMI to assess obesity reflecting the nutritional status of patients, while the distribution of body composition (fat distribution or muscle content) was ignored. For example, fat free mass index which was associated with COPD prognosis might better capture body composition,

but the operation is more complicated than BMI.³⁵ Second, because our study was cross-sectional and had a relatively small sample size in subgroups, it was impossible to determine a causal relationship between BMI and lung function in patients with COPD. Finally, the data obtained from multiple centers may have heterogeneity, but it was minimized using standard instruments, trained researchers, and a centralized electronic data acquisition and management system.

Conclusion

Our results in this study indicated that obesity only had a protective effect on lung function in COPD patients with GOLD 3–4 grade rather than GOLD 1–2 grade, which may be used to guide refined management of BMI in COPD patients with different GOLD stages. In future, the pathophysiological mechanism of causal relationship between BMI and lung function of COPD patients needs to be further studied.

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Disclosure

The authors declare that they have no conflicts of interest for this work.

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