

Active smoking and COPD phenotype: distribution and impact on prognostic factors

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Purpose: Smoking can affect both the phenotypic expression of COPD and factors such as disease severity, quality of life, and comorbidities. Our objective was to evaluate if the impact of active smoking on these factors varies according to the disease phenotype.

Patients and methods: This was a Spanish, observational, cross-sectional, multicenter study of patients with a diagnosis of COPD. Smoking rates were described among four different phenotypes (non-exacerbators, asthma-COPD overlap syndrome [ACOS], exacerbators with emphysema, and exacerbators with chronic bronchitis), and correlated with disease severity (body mass index, obstruction, dyspnea and exacerbations [BODEx] index and dyspnea grade), quality of life according to the COPD assessment test (CAT), and presence of comorbidities, according to phenotypic expression.

Results: In total, 1,610 patients were recruited, of whom 46.70% were classified as non-exacerbators, 14.53% as ACOS, 16.37% as exacerbators with emphysema, and 22.40% as exacerbators with chronic bronchitis. Smokers were predominant in the latter 2 groups (58.91% and 57.67%, respectively, $P=0.03$). Active smoking was significantly associated with better quality of life and a higher dyspnea grade, although differences were observed depending on clinical phenotype.

Conclusion: Active smoking is more common among exacerbator phenotypes and appears to affect quality of life and dyspnea grade differently, depending on the clinical expression of the disease.

Keywords: COPD, phenotype, smoking, prognostic factors, quality of life

Introduction

COPD is characterized by chronic airflow limitation, which is not fully reversible, generally progressive, and associated with an inflammatory pulmonary response to toxic particles and gases. The systemic impact is severe.¹ This disease presents in a wide variety of forms, and the concept that distinct COPD phenotypes can be identified, each with its characteristic clinical, prognostic, and therapeutic impact, is well established.² In Spain, the GesEPOC guidelines (Spanish COPD guidelines published by the Spanish Society of Pulmonology and Thoracic Surgery, which aim to establish standard recommendations for the management of the disease in Spain) propose 4 phenotypes, each with a different prognosis and therapeutic approach: the non-exacerbator phenotype (with emphysema or chronic bronchitis); the asthma-COPD overlap syndrome (ACOS); the exacerbator phenotype with emphysema; and the exacerbator phenotype with chronic bronchitis.³

Tobacco use is widely accepted as the major risk factor for the development of COPD.⁴ Exposure to tobacco smoke may also act as a differential factor in the various phenotypic expressions of the disease, and evidence suggests that a phenotype derived

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from smoking may be different from one caused by other environmental factors.^{5–7} Active smoking has been associated with COPD severity, quality of life, and comorbidities, although some results are controversial. For example, some authors report a positive relationship between active smoking and a higher rate of admission due to acute exacerbation, but others have not confirmed this finding.^{8–11} The impact of smoking on patients' quality of life has also been a topic of discussion,^{12–17} with previous reports indicating a lower mental^{13,14} and physical¹⁶ functional status in current smokers and an improvement of health-related quality of life after smoking cessation,^{15,17} although other studies have shown the opposite results.^{18,19}

The primary objective of this study, conducted under the auspices of a large trial aimed primarily at describing the distribution and characteristics of the different COPD phenotypes in Spain, was to examine the cross-sectional prevalence of active smoking among the various COPD phenotypes; the secondary objective was to determine the relationships between the smoking habit, COPD phenotypes, and major demographic and clinical variables, such as quality of life, disease severity, and presence of comorbidities.

Patients and methods

Study design and sample

This was an observational, cross-sectional, multicenter study performed in Pulmonology departments and other specialist units (Internal Medicine, Family and Community Medicine, and Occupational Medicine) throughout Spain. The aim of this study was to determine the prevalence and characteristics of the 4 COPD phenotypes established by GesEPOC³ in Spain. Each investigator was requested to include the first 6–8 unselected (consecutive) patients that fulfilled the selection criteria during the 90-day recruitment period. The study was approved by the Clinical Research Ethics Committee of the Hospital San Pedro de Alc ntara in C ceres, and patients gave written informed consent before participating. In this study, we examined the rates of active smoking and its relationship with COPD phenotype and factors influencing disease prognosis (severity, comorbidities, and quality of life). Patients were included if they were over the age of 35 years and had a clinical and functional diagnosis of COPD according to Global Initiative for Chronic Obstructive Lung Disease (GOLD) or GesEPOC criteria at least 6 months before participation in the study. Patients with symptoms of COPD exacerbation or who had had an exacerbation within the previous 6 weeks which could interfere with their perception of quality of life and functional status,

those with any type of chronic respiratory disease other than COPD, or who had difficulty reading or understanding the study questionnaires were excluded from participation.

Sociodemographic and clinical variables

Each investigator was trained in the study procedures and questionnaires and used a case report form (CRF) to collect the demographic details of each patient and the clinical variables required for the study. The GesEPOC algorithm was used to determine COPD phenotype.³ This algorithm, classifies the phenotype according to the following rules: A) patients with <2 exacerbations in the previous year were classified as non-exacerbators; B) patients with a previous diagnosis of asthma were considered as ACOS; C) exacerbators with emphysema that have a clinical/radiological/functional diagnosis of emphysema; and D) exacerbators who experienced cough with expectoration for >3 months of the year over 2 consecutive years were classified as exacerbators with chronic bronchitis. The schematic algorithm was provided to all the investigators (Figure 1).

Patients were asked about their smoking habit and were classified as nonsmokers, ex-smokers, and active smokers. COPD severity was measured using the BODEx index.²⁰ Quality of life was assessed with the COPD Assessment Test (CAT).²¹ Finally, comorbidities present at the time of data collection were evaluated.

Statistical analysis

COPD phenotype distribution was calculated using a descriptive analysis, with percentages and binomial 95% confidence intervals (95% CIs). Qualitative variables were compared using Fisher's exact test, and quantitative groups and variables were compared with the Student's *t*-test (for 2 groups) or the one-way analysis of variance (ANOVA) (for 3 or more groups). A logistic regression method was used to model the probability of being a smoker in relation to comorbidities (arrhythmia, coronary artery disease, heart failure, hypertension, dyslipidemia, diabetes mellitus, osteoporosis, and dementia), CAT score, serious exacerbations, and BODEx index. After stepwise analysis, the adjusted odds ratios (ORs) were calculated using multivariate logistic regression. Only significant associations are shown in the tables. All calculations were performed using the SAS statistical package version 9.4 for Windows, establishing a level of significance of 0.05.

Results

A total of 268 respiratory medicine specialists, 177 primary care physicians, 34 internists, and 1 occupational

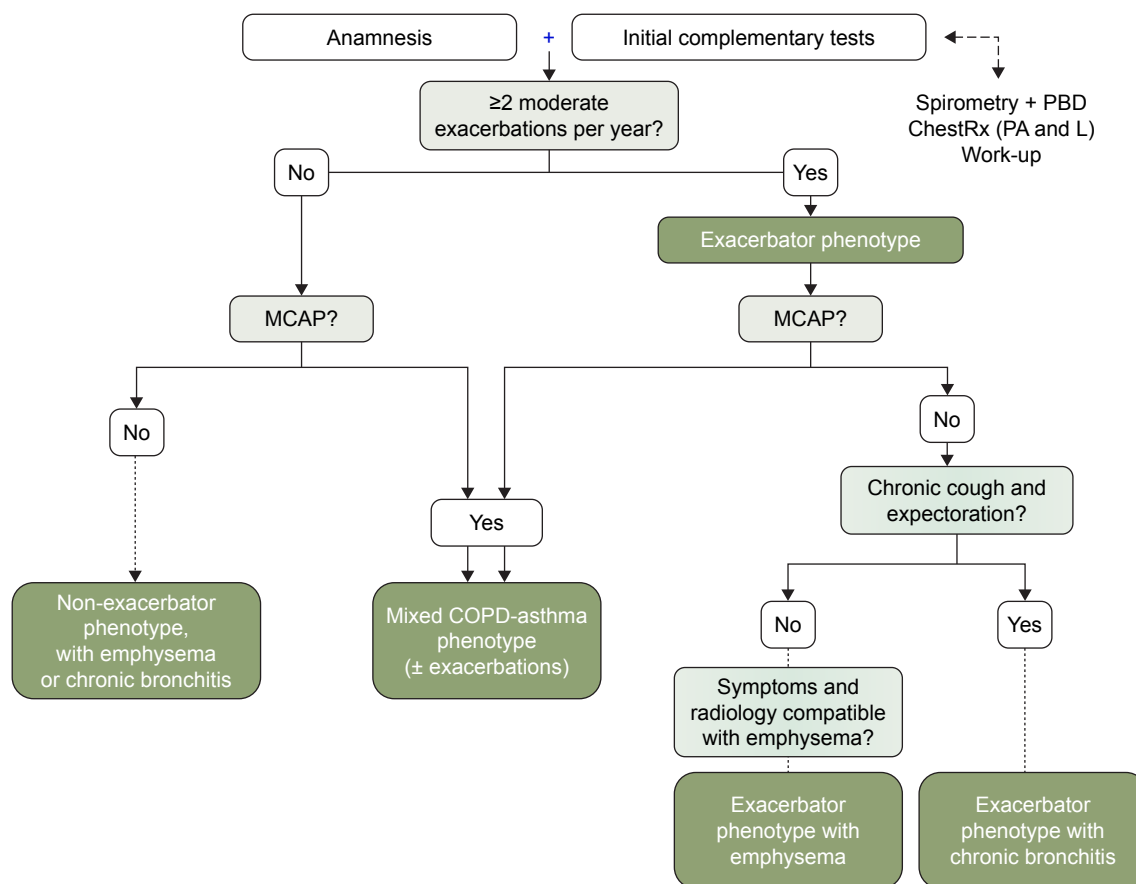


Figure 1 Diagnostic algorithm of the clinical phenotypes.

Note: Adapted from Miravittles M, Soler-Cataluña JJ, Calle M, et al. Spanish COPD Guidelines (GesEPOC): Pharmacological treatment of stable COPD. *Arch Bronconeumol*. 2012;48(7):247–257, with permission from the Sociedad Española de Neumología y Cirugía Torácica (SEPAR).³

Abbreviations: COPD, chronic obstructive pulmonary disease; MCAP, mixed COPD-asthma phenotype; ACOS, asthma-COPD overlap syndrome; PBD, bronchodilator test.

medicine specialist participated in this study. They recruited 1,610 patients (see Tables 1 and S1 for patients' characteristics), predominantly men (82.13%) with a mean age of 66.71 years (95% CI: 66.24–67.19). Of these, 55.41% (95% CI: 52.98–57.84) stated they were active smokers, with a smoking index of 65.02 pack-years (95% CI: 59.62–71.41), and a mean forced expiratory volume in 1 second (FEV₁) (%) of 54.70% (95% CI: 53.90–55.51). Seventy-one percent of patients had dyspnea grade ≤2, a mean BODEx index of 2.75 (95% CI: 2.65–2.85), and a mean CAT score of 22.04 (95% CI: 21.43–22.66). Finally, 83.79% had comorbidities, the most common being arterial hypertension (62.90%) and diabetes mellitus (31.7%).

With regard to phenotype distribution, 46.70% (95% CI: 44.26–49.14) were classified as non-exacerbators, 14.53% (95% CI: 12.81–16.25) as ACOS, 16.37% (95% CI: 14.56–18.18) as exacerbators with emphysema, and 22.40% (95% CI: 20.36–24.43) as exacerbators with chronic bronchitis (Figure 2). Significant differences were found in the distribution

of sexes by phenotype; the most pronounced being in the ACOS phenotype, in which there were almost twice as many women as men (21.7% vs 13.0%, respectively, $P=0.002$). Mean age among the phenotypes was significantly lower in the ACOS than that of exacerbators with chronic bronchitis (Table 2).

Active smoking

The frequency of active smoking by phenotype and its impact on quality of life, disease severity, and comorbidities were compared (Table 2). The group of nonsmokers accounted for only 2.9% of the total patient population; therefore, we decided to perform the remaining analyses on the basis of 2 groups only (active/nonsmokers + ex-smokers [nonactive smokers]). Even so, it is interesting to note that overall 9.1% of the female population had never smoked, whereas only 0.6% of all men had never smoked; this difference was statistically significant ($P<0.001$). Active smokers were predominant among the exacerbator phenotypes with emphysema and with chronic bronchitis (58.91% and 57.67%,

Table 1 Characteristics of the whole study population and stratification by smoking habit

Variables	Total population	Active smokers	Non smokers	P-value
Sex (n, %)				
Male	1,314 (82.1)	738 (56.2)	574 (43.8)	0.39
Female	286 (17.9)	152 (53.3)	133 (46.7)	
Age (years)	66.7 (66.2–67.2)	67.0 (66.4–67.7)	66.3 (65.6–67.0)	0.12
BMI (kg/m ²)	27.9 (27.6–28.1)	27.7 (27.4–28.0)	28.1 (27.7–28.4)	0.10
Dyspnea grade	1.8 (1.7–1.8)	1.8 (1.8–1.9)	1.7 (1.6–1.8)	0.003
Spirometry				
FVC (L)	2.5 (2.4–2.6)	2.5 (2.4–2.6)	2.5 (2.4–2.6)	0.89
FVC (%)	73.5 (72.7–74.2)	73.4 (72.3–64.4)	73.6 (72.7–74.2)	0.75
FEV ₁ (L)	1.3 (1.3–1.4)	1.4 (1.3–1.4)	1.3 (1.3–1.4)	0.56
FEV ₁ (%)	54.7 (53.9–55.5)	54.2 (53.1–55.4)	55.3 (54.1–56.5)	0.18
FEV ₁ /FVC	55.4 (54.7–56.0)	55.2 (54.3–56.1)	55.5 (53.9–55.5)	0.64
CAT score	22.0 (21.4–22.7)	20.5 (19.6–21.4)	23.3 (22.5–24.1)	<0.001
BODEx	2.7 (2.6–2.8)	2.8 (2.7–3.0)	2.7 (2.5–2.8)	0.15
Number of comorbidities	2.0 (1.9–2.1)	2.1 (2.0–2.2)	1.9 (1.8–2.0)	0.07

Note: Data presented as mean (95% confidence interval), unless otherwise stated.

Abbreviations: CI, confidence interval; BMI, body mass index; FVC, forced vital capacity; FEV₁, forced expiratory volume in 1 second; CAT, COPD assessment test; BODEx, body mass index, airflow obstruction, dyspnea and exacerbations.

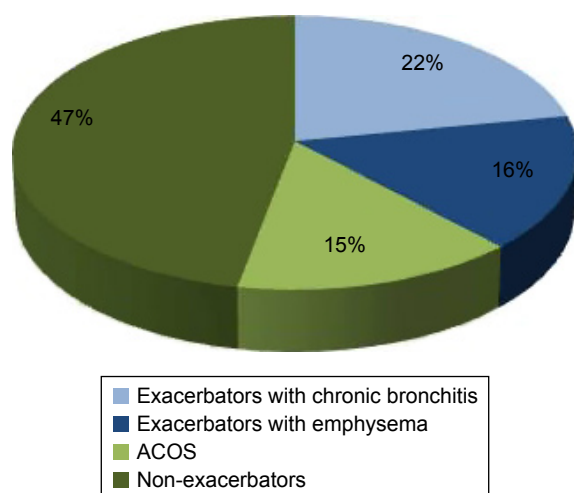
respectively, $P=0.03$). The impact of active smoking on patients' quality of life was found to be significantly positive, according to the CAT scores (20.51 [95% CI: 19.62–21.41] vs 23.33 [95% CI: 22.50–24.15; $P<0.001$]). This relationship remained significant for each of the items making up the CAT scale. When this correlation was analyzed for the different phenotypes, we found that while active smokers consistently had a better CAT score, it was statistically significant only in the more common phenotypes, that is, non-exacerbators and exacerbators with chronic bronchitis. In contrast, when the relationship between active smokers and dyspnea severity was analyzed, active smokers were seen to have a significantly greater dyspnea grade (1.83 [95% CI:

1.77–1.899] vs 1.69 [95% CI: 1.62–1.76]; $P=0.002$). When this relation was assessed according to the different phenotypes, it was only maintained in the group of exacerbators with chronic bronchitis (2.21 [95% CI: 2.09–2.33] vs 1.92 [95% CI: 1.80–2.00]; $P=0.001$). No differences were found in the BODEx score with respect to active smoking (2.82 [95% CI: 2.68–2.96] vs 2.67 [95% CI: 2.52–2.82]; $P=0.73$). Finally, no differences were observed in the presence of comorbidities according to smoking habit, although active smokers tended to have a higher comorbidity burden; this difference was significant among patients classified as exacerbators with chronic bronchitis (Table 2).

Multivariate analysis showed that higher CAT scores were negatively associated, whereas BODEx score was positively associated with active smoking (Table 3). Among comorbidities, osteoporosis was the only one associated with the probability of an individual being a smoker (OR =1.61, Wald 95% CI: 1.45–2.26).

Discussion

The results of this study are derived from a larger study designed to describe phenotype distribution among patients with COPD throughout Spain. In comparison with other studies of Spanish series,^{22,23} we found that while the proportion of patients with ACOS in our cohort was still around 15%, a greater proportion of patients were described as exacerbators (around 40% compared to 20% reported in other studies), and fewer were stable phenotypes (45% vs 65%). It is difficult to determine the reason for this discrepancy as the definition used to characterize the exacerbator

**Figure 2** Distribution by COPD phenotype.

Abbreviation: ACOS, asthma-COPD overlap syndrome.

Table 2 Patient characteristics and COPD severity according to phenotype and smoking habit

Variables	Non-exacerbators	ACOS	Exacerbators with emphysema	Exacerbators with chronic bronchitis
Sex (n, %)				
Male	618 (47.9)	167 (13.0)	209 (16.2)	296 (22.9)
Female	117 (42.2)	60 (21.7)	48 (17.3)	52 (18.8)
Age (years)	66.2 (65.5–67.0)	64.8 (63.5–66.2)	67.9 (66.7–69.0)	68.1 (67.1–69.1)
Spirometry				
FVC (L)	2.6 (2.5–2.7)	2.5 (2.3–2.7)	2.3 (2.2–2.4)	2.3 (2.2–2.4)
FVC (%)	74.4 (73.3–75.5)	75.1 (73.1–77.1)	71.3 (69.2–73.4)	72.1 (70.4–73.7)
FEV ₁ (L)	1.4 (1.4–1.5)	1.4 (1.3–1.5)	1.1 (1.1–1.2)	1.3 (1.2–1.3)
FEV ₁ (%)	56.6 (55.4–57.7)	60.8 (58.8–62.8)	46.6 (44.6–48.6)	52.6 (50.9–54.2)
FEV ₁ /FVC	56.0 (55.1–56.9)	60.0 (58.4–61.7)	49.3 (47.6–51.0)	55.3 (54.0–56.6)
Smoking habit (n, %)				
Active smokers	401 (54.6)	112 (48.9)	152 (58.9)	203 (57.7)
Non-active smokers	334 (45.4)	117 (51.1)	106 (41.1)	149 (42.3)
Dyspnea grade				
Active smokers	1.5 (1.4–1.6)	1.7 (1.5–1.9)	2.3 (2.2–2.5)	2.2 (2.1–2.3) ^a
Non-active smokers	1.4 (1.3–1.5)	1.6 (1.5–1.8)	2.3 (2.1–2.4)	1.9 (1.8–2.0)
CAT score				
Active smokers	16.6 (15.4–17.8) ^a	24.0 (22.1–26.5)	24.6 (22.7–26.5)	23.5 (21.4–25.7) ^a
Non-active smokers	20.0 (18.9–21.1)	25.3 (23.4–27.2)	26.8 (24.7–28.9)	26.8 (25.0–28.6)
BODEx				
Active smokers	2.1 (1.9–2.3)	2.2 (1.8–2.6)	4.2 (3.9–4.6)	3.5 (3.2–3.8)
Non-active smokers	2.0 (1.8–2.2)	2.3 (2.0–2.7)	4.3 (4.0–4.7)	3.2 (2.9–3.5)
Number of comorbidities				
Active smokers	1.7 (1.6–1.9)	2.1 (1.9–2.3)	2.1 (1.4–2.4)	2.7 (2.5–2.9) ^a
Non-active smokers	1.6 (1.4–1.7)	2.1 (1.8–2.3)	2.5 (1.9–2.4)	2.4 (2.2–2.6)

Notes: ^a $P < 0.05$ between active and non-active smokers in each phenotype category. Data presented as mean (95% confidence interval), unless otherwise stated.

Abbreviations: CI, Confidence Interval; ACOS, asthma-COPD overlap syndrome; CAT, COPD assessment test; BODEx, body mass index, airflow obstruction, dyspnea and exacerbations; FEV₁, forced expiratory volume in 1 second; FVC, forced vital capacity.

phenotype was the same in all studies (presence of at least 2 exacerbations in the last year or 1 hospitalization for acute exacerbation). The difficulty in precisely defining “exacerbation” may have contributed to this discrepancy. Evidence shows that exacerbations are often underreported because many patients with COPD may have difficulty in recognizing their symptoms as exacerbation, and accordingly, tend not to report them.^{24,25} Langsetmo et al, for example, evaluated the incidence of reported and unreported exacerbations in

a cohort of 421 patients with COPD and showed that less than one third of the exacerbations were reported, being that the number of symptoms at onset was the most important predictor of reporting.²⁴ However, Hurst et al, in one of the largest studies conducted to determine susceptibility to exacerbations in stable patients with COPD over a period of 3 years, established that the incidence of exacerbations was 22% in patients with moderate COPD, rising to 47% in severe cases.²⁵ According to the study authors, this incidence would have been much greater if untreated exacerbations had been computed along with treated events. In any case, the clinical implication of these results could be important. An action plan to help patients recognize changes in their symptoms, to implement self-care, and to self-initiate prescription in the event of an exacerbation has been suggested as a promising strategy.²⁶

Existing evidence shows not only that smoking is the primary etiological cause of COPD, but also that it can also lead to faster decline in lung function if the habit continues; even so, the proportion of smokers among patients with COPD remains higher than among the general population.²⁷

Table 3 Factors associated with active smoking in the multivariate analysis

Variables	OR	95% CI	P-value
CAT (11–20 vs 0–10)	1.141	0.769–1.693	0.51
CAT (21–30 vs 0–10)	0.700	0.472–1.038	0.08
CAT (31–40 vs 0–10)	0.301	0.187–0.487	<0.001
BODEx (3–4 vs 0–2)	1.202	0.914–1.581	0.19
BODEx (5–6 vs 0–2)	1.637	1.151–2.328	0.006
BODEx (≥ 7 vs 0–2)	2.051	1.031–4.080	0.040
Osteoporosis (yes vs no)	1.610	1.145–2.265	0.006

Abbreviations: OR, odds ratio; CI, confidence interval; CAT, COPD assessment test; BODEx, body mass index, airflow obstruction, dyspnea and exacerbations.

Recent studies performed in the USA found smoking rates between 35% and 45% among adult patients with COPD.^{28,29} In our series, the prevalence of active smokers is 55%, contrasting with reports recently published in Spain that set it at about 25%.^{22,23} Perhaps these differences reflect differences in data collection methods. Different ways of classifying a smoker as active may have a significant influence on the results of studies conducted in these patients, and hinder comparison among the different papers.³⁰ However, a good level of agreement has been shown between smoking statuses based on self-reported and serum cotinine measurements.³¹ In any case, the proportion of active smokers among patients with COPD is still unacceptably high, particularly when we consider that one of the key objectives established by GOLD guidelines in the management of these patients is smoking cessation.¹

Results on smoking and phenotype distribution among patients with COPD show that a significantly higher proportion of patients with exacerbator phenotype are active smokers. Previous data indicate that patients with greater susceptibility to viral infections are more prone to exacerbations, and this susceptibility appears to be associated with their smoking habit.³² The viral load during acute exacerbations of COPD has been detected as significantly higher than in stable state.³³ Moreover, a recent study has demonstrated an upregulated expression of ICAM-1 (major receptor for 60% of human rhinoviruses and *Haemophilus influenza*, two major pathogens in COPD) in the respiratory tract of smokers and especially in the airway epithelium in subjects with chronic airflow obstruction. This finding could reinforce the relation between active smoking, viral infection, and frequency of exacerbations.³⁴ However, conflicting data about this association still persists. For example, Hurst et al,²⁵ in a study that evaluated the impact of certain factors (ie, demographic and clinical characteristics, lung function, and laboratory and biomarkers values) on susceptibility to exacerbations found that while there was some relationship between active smoking and susceptibility to exacerbations, this association disappeared after multivariate analysis, showing that the link was not independent and may have been influenced by interactions with other variables, such as previous exacerbations, disease severity, health status, history of gastroesophageal reflux, and white cell count.²⁵ This discrepancy has also been observed in series recently published in Spain.^{22,23} Although Miravittles et al²² did report a higher rate of smoking among exacerbators, Cosio et al²³ failed to identify such an association. In our series, patients over the age of 50 years, who were significantly heavier

smokers, were predominant in these phenotypes, and this may have affected the results.

When the impact of smoking on patients' quality of life was analyzed, we were surprised to find that former smokers obtained a poorer CAT score. Moreover, the fact that the groups that smoked most, that is, older patients and exacerbators, reported a worse quality of life makes this finding even more difficult to understand. Although most studies have found an inverse relationship between smoking and quality of life, this has not always been the case. Some studies have found that while many smokers attempt to give up their habit, many others continue to smoke as they associate the habit with psychological well-being, relaxation, and relief of stress.^{18,19} These factors may lead to a mistaken perception of quality of life on the part of the patients themselves. Wijnhoven et al also observed that quality of life was positively affected by active smoking. These authors associated it with the fact that patients with a better perception of their disease (less symptomatic) may wait longer before quitting smoking than patients with more severe symptoms.³⁵ These differences might perhaps not have been revealed if the dynamics of this variable over time had been evaluated. Shields et al in a longitudinal study that analyzed the dynamics of smoking cessation between former smokers and nonsmokers concluded that a relatively long period of time was needed before quitters achieved level of comfortable life similar to those of individuals who have never smoked. Thus, the authors assert that it is difficult to extract conclusions on these measurements from cross-sectional studies.¹⁵ Ståhl et al also failed to find differences in quality of life associated with smoking habit among patients with COPD and suggest that the degree of disease severity is a metric with greater impact in this entity.³⁶ The discordance is deepened further by the fact that dyspnea severity is significantly associated with active smoking. Several studies have cast doubts on the equivalence between dyspnea grade and CAT score in the evaluation of disease involvement.^{37,38} Indeed, recent evidence shows that the isolated use of CAT as a tool to establish the impact and severity of COPD may be inappropriate, given the difficulty of establishing definite cutoff points for a sufficiently sensitive and specific classification of patients according to their severity.³⁹ Our results appear to support this premise. Finally, we should mention the limited representation among our series of nonsmokers, which prevented us from analyzing the influence of this population on the final results.

COPD is often associated with other diseases and a great deal has been written about the systemic effects of

the disease itself and how these clearly increase pressure on health care resources and the risk of death. COPD was associated with other diseases in up to 80% of the patients in our series, the most common comorbidity being arterial hypertension; these data coincide with the findings of most previously published studies.^{40,41} When comorbidities were analyzed according to the use of tobacco, it was found that patients with osteoporosis were more likely to be active smokers. Osteoporosis has been recognized as one of the major noncardiovascular comorbidities associated with COPD, occurring at rates of 4%–59%, depending on the diagnostic method employed.⁴² Many explanations have been given for this association, including age, sex, smoking habit, limited daily activity, chronic hypoxemia, or corticosteroid treatment. Smoking has consistently been associated with low bone density in healthy volunteers;⁴³ therefore, it is unsurprising that this relationship is stronger in patients with COPD, in whom other factors also combine to raise the risk of developing osteoporosis. It is also interesting to note the relationship that has been established between the emphysema phenotype and an increased risk of osteoporosis suggests a common mechanism between these two conditions.^{43,44} We ourselves did not find any specific relationship between phenotypes and osteoporosis, although active smokers were more predominant in phenotypes in which this disease may occur (nonexacerbators and exacerbators with emphysema).

Conclusion

Our study provides a general overview of how smoking, phenotype, and prognostic factors, such as quality of life or comorbidities, are interwoven in a large sample of Spanish patients with COPD. Some disparity between our data and those derived from similar recently published series may be due to the limitations to which all studies of this kind are subject (cross-sectional, participation of different specialists). Nevertheless, the results underline the importance of active smoking in the characterization of the disease.

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Disclosure

The authors report no conflicts of interest in this work.

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Supplementary materials

Table S1 Description of patient population included in the study

Demographics									
	N	NMiss	Median	SD	95% CI Low	95% CI High	Min	Median	Max
Age (years)	1,580	30	66.71	9.67	66.24	67.19	35.00	67.00	92.00
n									%
Age (years)									
35–60	42226.71								
61–65	28217.85								
66–75	56035.44								
+75	31620.00								
Total	1,580100.00								
Sex									
Male	1,31482.13								
Female	28617.88								
Total	1,600100.00								
	N	NMiss	Median	SD	95% CI Low	95% CI High	Min	Median	Max
Weight (kg)	1,587	23	77.77	14.53	77.05	78.48	40.00	78.00	183.00
Height (cm)	1,585	25	166.95	7.90	166.56	167.34	130.00	167.00	194.00
BMI (kg/m²)	1,581	29	27.86	4.72	27.63	28.10	13.52	27.68	68.36
Smoking status		Frequency		%		IC95% Inf %		IC95% Sup %	
Smoker		891		55.41		52.98		57.84	
Ex-smoker		677		42.10		39.69		44.51	
Nonsmoker		40		2.49		1.73		3.25	
		1,608		100.00					
	N	Median	SD	95% CI Low	95% CI High	Min	Median	Max	
Smoking index (pack/year)									
Smoker	849	65.02	94.95	58.62	71.41	1.00	40.00	900.00	
Ex-smoker	658	61.72	88.56	54.94	68.50	1.00	45.00	800.00	
Nonsmoker	0	–	–	–	–	–	–	–	–
Total	1,507	63.58	92.20	58.92	68.24	1.00	42.00	900.00	
Clinical characteristics									
	N	NMiss	Median	SD	95% CI Low	95% CI High	Min	Median	Max
Years from COPD diagnosis	1,505	105	8.95	7.93	8.55	9.35	0.00	7.00	89.00
	N	NMiss	Median	SD	95% CI Low	95% CI High	Min	Median	Max
FVC (mL)	1,234	376	2,499.94	869.44	2,451.38	2,548.50	300.00	2,530.00	4,310.00
FVC (%)	1,329	281	73.48	14.37	72.70	74.25	44.00	73.00	106.00
FVI (mL)	1,236	374	1,350.05	551.87	1,319.26	1,380.85	193.00	1,290.00	2,730.00
FVI (%)	1,345	265	54.70	15.08	53.90	55.51	26.00	55.00	87.00
FVI/FVC (%)	1,358	252	55.37	12.17	54.73	56.02	27.00	57.00	81.00
n									%
Dyspnea grade									
Grade 0	986.15								
Grade 1	57235.88								
Grade 2	56535.45								
Grade 3	31819.95								
Grade 4	412.57								
Total	1,594100.00								
	N	NMiss	Median	SD	95% CI Low	95% CI High	Min	Median	Max
Number of exacerbations in the previous year	1,567	43	0.95	1.25	0.88	1.01	0.00	1.00	10.00
BODEx	1,423	187	2.75	1.93	2.65	2.86	0.00	3.00	8.00

(Continued)

Table S1 (Continued)

					n	%				
BODEx										
BODEx 0–2					692					48.63
BODEx 3–4					449					31.55
BODEx 5–6					236					16.58
BODEx ≥7					46					3.23
Total					1,423					100.00
	N	NMiss	Median	SD	95% CI Low	95% CI High	Min	Median	Max	
CAT score	1,591	19	20.89	8.93	20.45	21.33	0.00	21.00	40.00	
					n	%				
COPD impact according to CAT										
None					55					3.46
Low					128					8.05
Intermediate					559					35.14
High					614					38.59
Very high					235					14.77
Total					1,591					100.00
Does the patient present comorbidities?										
Yes					1,349					83.79
No					261					16.21
Total					1,610					100.00
	N	NMiss	Median	SD	95% CI Low	95% CI High	Min	Median	Max	
No comorbidities	1,610	0	2.00	1.42	1.93	2.07	0.00	2.00	7.00	
					n	%				
No comorbidities										
None					261					16.21
1 comorbidity					354					21.99
2 comorbidities					460					28.57
3 comorbidities					306					19.01
+3 comorbidities					229					14.22
Total					1,610					100.00
Comorbidities										
Item	N			n	%		95% CI Low		95% CI High	
HBP	1,523			958	62.90		60.48		65.33	
Diabetes mellitus	1,479			461	31.17		28.81		33.53	
Chronic ischemic heart disease	1,448			205	14.16		12.36		15.95	
Arrhythmias	1,457			237	16.27		14.37		18.16	
Heart failure	1,458			229	15.71		13.84		17.57	
Osteoporosis	1,461			263	18.00		16.03		19.97	
Dyslipidemia	1,523			853	56.01		53.51		58.50	
Dementia	1,433			20	1.40		0.79		2.00	
COPD phenotype										
Phenotype A	Frequency			%		95% CI Low		95% CI High		
Phenotype B	736			46.70		44.26		49.14		
Phenotype C	229			14.53		12.81		16.25		
Phenotype D	258			16.37		14.56		18.18		
	353			22.40		20.36		24.43		
	1,576			100.00						

Abbreviations: BMI, body mass index; BODEx, body mass index, airflow obstruction, dyspnea and exacerbations; COPD, chronic obstructive pulmonary disease; CAT, COPD assessment test; FVC, forced vital capacity; FEV₁, forced expiratory volume in 1 second; HBP, high blood pressure; NMISS, number of missing entries.

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