

SCIENTIFIC REPORTS

OPEN

A scoring system to detect fixed airflow limitation in smokers from simple easy-to-use parameters

Esther Román-Conejos¹, Antonio Palazón-Bru², David Manuel Folgado-de la Rosa², Manuel Sánchez-Molla³, María Mercedes Rizo-Baeza⁴, Vicente Francisco Gil-Guillén² & Ernesto Cortés-Castell⁵

Received: 20 March 2018

Accepted: 8 August 2018

Published online: 06 September 2018

No validated screening method currently exists for Chronic Obstructive Pulmonary Disease (COPD) in smokers. Therefore, we constructed a predictive model with simple parameters that can be applied for COPD screening to detect fixed airflow limitation. This observational cross-sectional study included a random sample of 222 smokers with no previous diagnosis of COPD undertaken in a Spanish region in 2014–2016. The main variable was fixed airflow limitation by spirometry. The secondary variables (COPD factors) were: age, gender, smoking (pack-years and Fagerström test), body mass index, educational level, respiratory symptoms and exacerbations. A points system was developed to predict fixed airflow limitation based on secondary variables. The model was validated internally through bootstrapping, determining discrimination and calibration. The system was then integrated into a mobile application for Android. Fifty-seven patients (25.7%) presented fixed airflow limitation. The points system included as predictors: age, pack-years, Fagerström test and presence of respiratory symptoms. Internal validation of the system was very satisfactory, both in discrimination and calibration. In conclusion, a points system has been constructed to predict fixed airflow limitation in smokers with no previous COPD. This system can be integrated as a screening tool, though it should be externally validated in other geographical regions.

Chronic obstructive pulmonary disease (COPD) is a major public health problem, making it the fourth leading cause of death worldwide¹. Despite its associated morbidity and mortality and the considerable expenditure of resources generated by COPD, it is underdiagnosed worldwide^{2–4}.

The main risk factor for COPD is smoking⁵. Other factors include age, male gender and respiratory symptoms⁶. COPD has also been associated with certain chronic comorbidities such as hypertension, cardiovascular problems, and diabetes^{7,8}.

The most commonly accepted method to confirm COPD is forced spirometry, which is considered positive with a post-bronchodilator forced expiratory volume in the first second (FEV₁)/forced vital capacity (FVC) ratio < 0.7. This is a simple, reproducible technique with low individual cost^{9,10}. Although it is indicated in patients with respiratory symptoms, in normal clinical practice it is underutilized¹¹, thereby explaining part of the underdiagnosis. Thus, forced spirometry does not appear to be a good screening technique that can be applied in all smokers regardless of their symptoms because, in most cases, these symptoms become evident when the disease is already established or when they already affect quality of life, which is associated with a greater severity of the disease^{12,13}.

As there is no established early screening program, alternatives for the early detection of COPD have been proposed. These may prove useful prior to confirmation by spirometry to reduce the high prevalence of underdiagnosis. There are only three models to predict COPD in patients with a high risk of developing the disorder but with no prior history of lung disease (Table 1). Although these models are easy to use in general clinical

¹Raval Health Centre, Generalitat Valenciana, Elche, Alicante, Spain. ²Department of Clinical Medicine, Miguel Hernández University, San Juan de Alicante, Alicante, Spain. ³Department of Medical Management, University General Hospital of Elche, Elche, Alicante, Spain. ⁴Department of Nursing, University of Alicante, San Vicente del Raspeig, Alicante, Spain. ⁵Department of Pharmacology, Pediatrics and Organic Chemistry, Miguel Hernández University, San Juan de Alicante, Alicante, Spain. Esther Román-Conejos and Antonio Palazón-Bru contributed equally. María Mercedes Rizo-Baeza, Vicente Francisco Gil-Guillén and Ernesto Cortés-Castell jointly supervised this work. Correspondence and requests for materials should be addressed to A.P.-B. (email: antonio.pb23@gmail.com)

Reference	Population	Variables	Clinical use	EPV ≥ 10	Continuous variables	Missing data	Model building strategy	Discrimination
López-Varela <i>et al.</i> ³⁸	COPD risk	Gender, age, pack-years, dyspnoea, chronic phlegm and cough, and previous spirometry	Scoring system	Yes	Categorization	Not mentioned	Used all their co-variables	AUC, sensitivity and specificity, no bootstrapping
Llordés <i>et al.</i> ³⁹	Smokers	Age, gender, BMI, pack-years, profession of risk, expectoration, dyspnoea, cold complications, dyspnoea treatment and cardiovascular disease	Scoring system	No	Categorization	Not mentioned	Selection based on bivariate analysis	AUC, sensitivity and specificity, no bootstrapping
Price <i>et al.</i> ^{40,41}	COPD risk	Age, BMI, pack-years, cough due to the weather, phlegm, breathlessness and allergies	Scoring system	No	Categorization	Not mentioned	Stepwise selection based on bivariate analyses, factor analysis and AIC	AUC, sensitivity and specificity, no bootstrapping

Table 1. Analysis of the published studies that developed a predictive model to diagnose chronic obstructive pulmonary disease in risk populations (screening). Abbreviations: A.I.C., Akaike's Information Criterion; A.U.C., area under the receiver operating characteristic curve; B.M.I., body mass index; C.O.P.D., chronic obstructive pulmonary disease; E.P.V., events-per-variable; C.O.P.D. risk is defined as being current or former smoker. Note that it is entirely possible that the mentioned limitations could be overcome by a subsequent successful external validation.

practice as they employ variables that are easy to obtain and the risk of COPD is calculated with a scoring system, they nevertheless suffer from important limitations concerning their development (estimation) and their internal validation. Concerning the former, two of the models present overfitting (events-per-variable < 10), none of the three analysed the functional form of the continuous predictors and the treatment of missing data is not given; in addition to which only one model selected the predictors based on the overall goodness of fit of the model. Concerning the validation, this was not done by bootstrapping in any of the models and the calibration was not done with smooth curves to estimate the observed probability of COPD^{14–16}. Accordingly, no real model to predict COPD is available for use as a screening tool developed with the best and recommended statistical technique.

As a response to the need to implement a screening method that can differentiate smokers with a higher risk of fixed airflow limitation prior to onset of established symptoms, the objective of this study was the elaboration and internal validation (bootstrapping) of a prediction model for fixed airflow limitation, using the recommended statistical methodology^{14–16}, with simple parameters that can be implemented in routine clinical practice in a systematic way, without incurring an increase in care time. In addition, to facilitate the implementation of this model, it will be integrated into a mobile phone application in the Android operating system (*COPD predictor* in Google Play), which could be used even by patients. With all this, we will have a screening tool that, provided it is externally validated in other populations, could be integrated into the protocols and clinical guidelines for COPD, prior to the definitive diagnosis by forced spirometry and respiratory symptoms. Whilst COPD screening is not currently recommended in smokers, this is because the techniques to do so are time consuming and expensive. Therefore, in this study we present an alternative that overcomes these limitations and leaves spirometry for confirmation in high-risk cases.

Method

Study Population. Smokers aged 40–75 years with no previous diagnosis of COPD whose primary health care is delivered at the Raval Health Centre in Elche (Alicante). This centre covers a population of 20,284 adults and is a primary care centre of the Spanish Public Health System which provides universal and free coverage.

Study design and participants. This cross-sectional observational study was carried out during 2014–2016 with the objective of determining the underdiagnosis of COPD resulting from the underuse of forced spirometry and with this to construct a predictive model. We consulted the outpatient database of this centre and excluded those patients who did not meet the following inclusion criteria: smokers, age 40–75 years, and with no active diagnosis of COPD in their medical record. From the list obtained in this consultation, a group of patients was selected by random sampling (random number table) and contacted by telephone, asking them over the phone for their informed consent to participate in the study. The patients who agreed to participate were asked to perform forced spirometry, fill out questionnaires on COPD risk factors, and sign the written informed consent. Just under 10% of those selected refused to participate (generally due to incompatible work schedules, $n = 24$, 9.8%). These patients were passed over and the next person in the random table was selected.

Variables and measurements. The primary variable was fixed airflow limitation performed with a validated Dataspir 110 A spirometer, according to the regulations of the main Spanish medical societies and the criteria established in the GOLD guide (post-bronchodilator FEV₁/FVC ratio < 0.7)^{12,17}.

Secondary variables were collected according to the risk factors associated with the diagnosis of COPD: older age, male gender, number of cigarettes smoked and for how long, low body mass index (BMI), low educational level, presence of respiratory symptoms and exacerbations during the last year^{4,8,18–20}. In the interview details were obtained about: gender; age; educational level (0 = primary, 1 = secondary-intermediate degree, 2 = higher studies-University); smoking history in pack-years, calculated as the number of cigarettes smoked per day

multiplied by the number of years smoking divided by 20; Fagerström test, considering three levels of dependence: mild (0–3 points), moderate (4–6 points) and severe (7–10 points)²¹; presence of symptoms or respiratory problems (aphonia, cough, catarrh, bronchitis, etc.) requiring medical consultation during the last year and level of exacerbation, described as a sustained deterioration in the patient's baseline clinical condition, beyond the usual daily variability, which appears acutely, and is accompanied by increased dyspnoea and expectoration and a change in the appearance of sputum, or any combination of these three symptoms, requiring a therapeutic change and classified as mild (no episodes of bronchitis in the last year), moderate (episodes treated by primary care physician) and severe (treated in the emergency area and/or hospitalisation)²².

Sample size calculation. Since the objective of this study was to construct a predictive model of a binary event through a binary logistic regression model, the sample size had to verify that the ratio between the number of events and the number of predictors of the model was greater than or equal to 10²³.

Statistical analysis. Qualitative variables were described using absolute and relative frequencies, whilst quantitative variables were expressed through means and standard deviations. Our variables had no missing data. Associations between the primary variable and the secondary variables were assessed using the χ^2 test and the *t*-test. A logistic regression model was constructed with a maximum number of five predictors (57 patients with fixed airflow limitation). Taking into account that we had a total of 8 predictors [considering educational level and the Fagerström test as linear predictors, as they did not show a quadratic trend (Wald test)], we checked all possible combinations of 1 to 5 predictors, selecting the one with the highest discriminating capacity, that is, the one that gave a maximum area under the receiver operating characteristic curve (AUC). Thus, the AUC was calculated in a total of 218 combinations. The optimum combination was adapted to a points system using the Framingham study methodology²⁴ which, through a weighting of the model coefficients and a categorization of the predictors, associates a score to each variable and the sum of these scores gives an event probability. Once the points system was developed, it was internally validated through bootstrapping (1000 random samples), since this is the most recommended technique²⁵. Discrimination was determined in each of the 1000 samples (the points system can differentiate fixed airflow limitation) and calibration (verifying that the prediction of the model corresponds to reality). Discrimination is addressed by calculating the AUC, whilst calibration is evaluated through the construction of smooth curves (linear splines) with the Hosmer-Lemeshow test, which is appropriate for the recommended level of calibration (moderate)¹⁵. All analyses were performed with a type I error of 5%, and for each relevant parameter its associated confidence interval (CI) was calculated. Statistical packages used were SPSS Statistics 24 and R 2.13.2.

Ethical approval. This project was approved by the Clinical Trials Ethics Committee of the Department of Health of the Generalitat Valenciana (General University Hospital of Elche, Alicante) at its meeting on 20 March 2014, in accordance with the International Guidelines for Ethical Review of Epidemiological Studies and the recommendations of the Spanish Society of Epidemiology on the review of the ethical aspects of epidemiological research studies, respecting the rules set forth in Law 15/1999 on Personal Data Protection in research studies. The committee approved the age range of the patients in the study (40–75 years), as the highest prevalence of COPD occurs at these ages.

How This Fits In. The diagnosis of COPD is customarily made using forced spirometry and respiratory symptoms, but this does not appear to be a good screening technique that can be extended to all smokers. We elaborated and internally validated a prediction model for fixed airflow limitation with simple parameters that can be implemented in routine clinical practice in a systematic way. Care time would not be increased, since the model will be implemented in a mobile app for Android. With all this, we will have a screening tool that could be integrated, after external validation, in the protocols and clinical guidelines for COPD.

Results

Spirometry was performed in a total of 222 patients, of which 119 (53.6%) were men; 57 (25.7%) patients had fixed airflow limitation. Table 2 shows the values of the variables analysed; also shown are the results of the bivariate analysis which revealed male gender, age, educational level, nicotine dependence (Fagerström), cigarette consumption, presence of respiratory symptoms and exacerbations to be significant risk factors ($p < 0.05$). The same table shows the coefficients of the multivariate model with the optimum combination to predict fixed airflow limitation. We highlight that this combination included: age, educational level, Fagerström test, pack-years smoked and presence of respiratory symptoms. Figure 1 illustrates the adaptation of the model to a points system (educational level was not present when we adapted the model to a scoring system). Figure 2 depicts the logistic tendency for the probability of fixed airflow limitation as the overall score increases.

The internal validation of the points system using the bootstrap technique was very satisfactory. First, in Fig. 3 we can see that the discrimination is adequate, as the central value of the AUC distribution is 0.80. Subsequently, Fig. 4 shows that the observed versus expected probabilities. Note that most of the scores had small errors, except when the patient had 6 points (15.36%), but these patients had a proportion of 10.8% in the sample and overall this was appropriate (Hosmer-Lemeshow test: $p = 0.492$). Consequently, the points system has been internally validated with satisfactory calibration and discrimination.

Discussion

Summary. This study developed a simple prediction tool to assess fixed airflow limitation in smokers with no previous diagnosis of COPD. The model is simple to use because the variables are easy to measure, risk is assessed by a points system and its implementation in a mobile application makes calculations even easier. The model has been internally validated in our setting through the most recommended statistical methods. Underdiagnosis of

Variable	Total n = 222 n(%) / $\bar{x} \pm s$	Fixed airflow limitation n = 57 (25.7%) n(%) / $\bar{x} \pm s$	p-value	Adj. OR (95% CI)	p-value
Male gender	119(53.6)	37(31.1)	0.047	N/M	N/M
Age (years)	56.8 $\pm$ 10.4	63.8 $\pm$ 9.2	<0.001	1.08(1.03–1.14)	0.001
Educational level*:					
Primary Secondary University	78(35.1) 87(39.2) 57(25.7)	29(37.2) 19(21.8) 9(15.8)	0.011	1.03(0.61–1.75)	0.902
Fagerström test (dependence)*:					
Low Moderate High	90(40.5) 113(50.9) 19(8.6)	12(13.3) 33(29.2) 12(63.2)	<0.001	2.06(1.07–3.99)	0.032
Smoking pack-years	28.8 $\pm$ 13.4	38.6 $\pm$ 11.8	<0.001	1.04(1.01–1.07)	0.021
Respiratory symptoms	88(39.6)	41(46.6)	<0.001	4.36(2.01–9.46)	<0.001
Exacerbation*:					
Mild Moderate Severe	147(66.2) 67(30.2) 8(3.6)	22(15.0) 28(41.8) 7(87.5)	<0.001	N/M	N/M
BMI (kg/m ²)	26.6 $\pm$ 4.8	27.1 $\pm$ 5.8	0.387	N/M	N/M

Table 2. Predictive model for fixed airflow limitation in smokers with no previous diagnosis of chronic obstructive pulmonary disease in primary health care. Abbreviations: Adj. O.R., adjusted odds ratio; B.M.I., body mass index; C.I., confidence interval; n(%), absolute frequency (relative frequency) N/M, not in the model; $\bar{x} \pm s$, mean $\pm$ standard deviation. *Analysed in the multivariate model as a quantitative variable. Goodness-of-fit of the multivariate model (likelihood ratio test): $\chi^2 = 11.5$, $p < 0.001$.

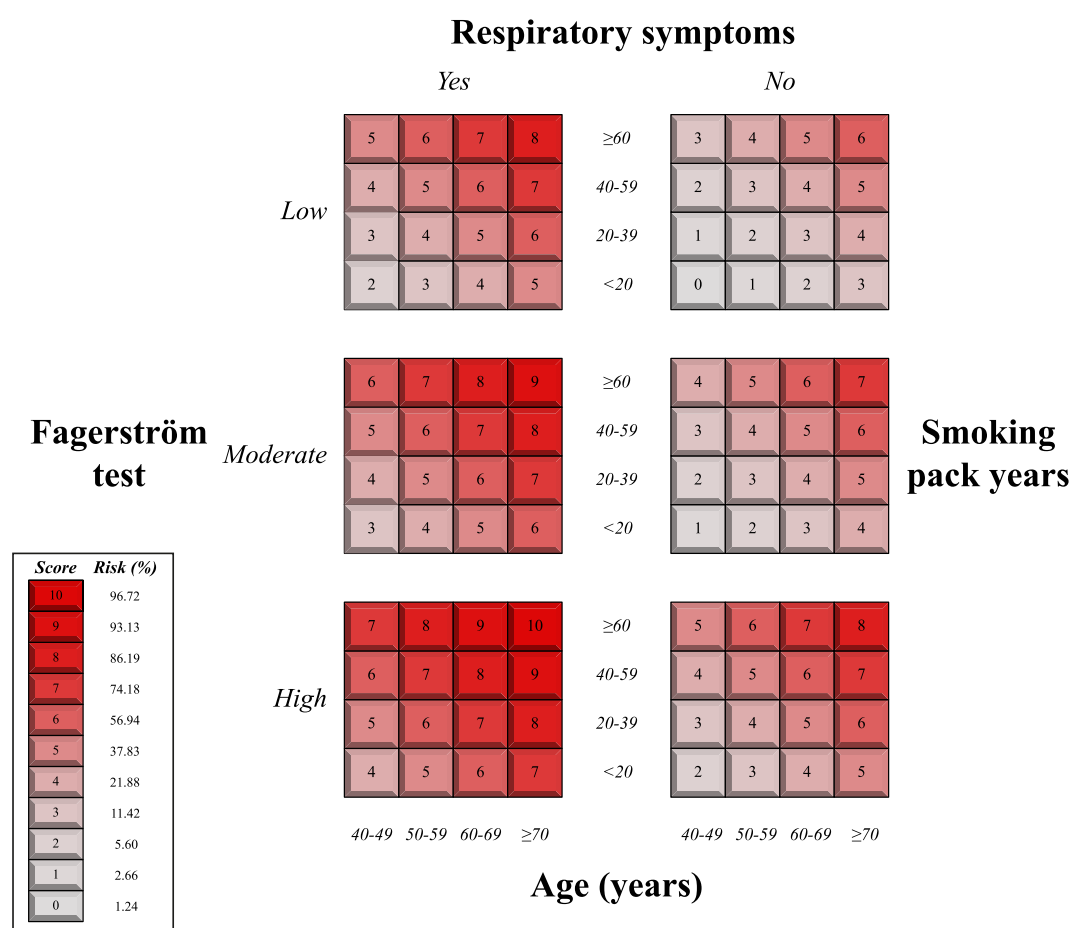


Figure 1. Scoring system to predict the diagnosis of fixed airflow limitation in smokers. The box on the left shows the overall scores with their associated risk of fixed airflow limitation.

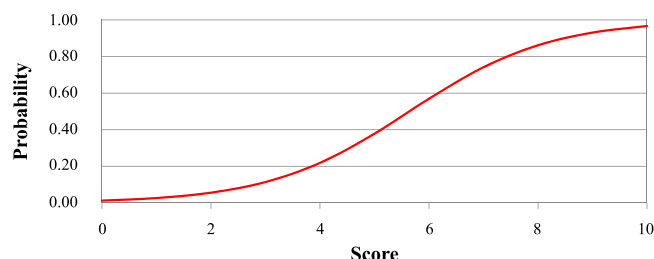


Figure 2. Predicted probability of fixed airflow limitation using our scoring system.

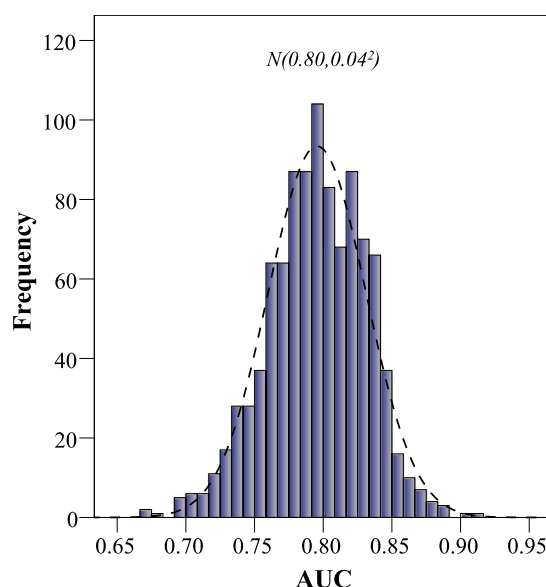


Figure 3. Area under the Receiver Operating Characteristic curve distribution for the validation of our scoring system using the bootstrap method. Abbreviations: AUC, area under the Receiver Operating Characteristic curve; CI, confidence interval.

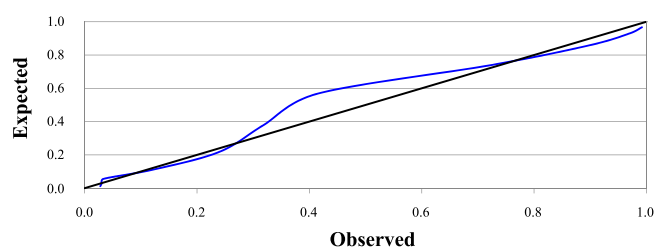


Figure 4. Smooth calibration plots for the validation of our scoring system using the bootstrap method. The black line represents perfect calibration and the grey line indicates the results of our calibration. The error (observed-expected in %) in each score was (ordered from 0 to 10 points; in parentheses we indicate the proportion of patients with each score): 1.53 (7.7%), 0.34 (5.9%), −2.30 (18.9%), 1.08 (17.6%), 2.11 (14.0%), −6.05 (12.6%), −15.36 (10.8%), −1.53 (5.0%), 4.04 (5.4%), 3.95 (1.8%) and 2.48% (0.5%).

COPD in our population was 25.7% (fixed airflow limitation), with the following associated risk factors: male gender, older age, lower educational level, higher level of nicotine dependence (Fagerström), higher cigarette consumption, presence of respiratory symptoms and high level of exacerbations.

Strengths and limitations. The main strength of our study is the development and internal validation of a prediction model for COPD screening in patients prior to diagnostic confirmation by spirometry and respiratory symptoms. Although others have developed a prediction model, their models have several limitations for use in clinical practice (Table 1). We also highlight the statistical methodology used since we chose the combination of explanatory variables of the multivariate model with the highest discriminating capacity. Additionally, we used the most recommended techniques for the validation of a predictive model of a binary event^{14–16,25}.

Selection bias was minimized by randomly choosing the sample from among all smokers in the corresponding health area, with very low exclusion for not wanting to participate (about 10%). To minimize information bias, the tests were conducted by the principal investigator using validated questionnaires and instruments. To avoid possible confounding bias, we applied well-calibrated multivariate models with high discriminating capacity. Our main limitations were sample size (limiting the number of predictors) and lack of external validation (future line of research).

We could have used the lower limit criterion instead of setting the FEV₁/FVC ratio threshold at 0.7. However, it has been seen in populations similar to ours that establishing the diagnosis of COPD with the lower limit could exclude a high number of patients with significant clinical impact and high consumption of healthcare resources²⁶. We have also used a total of 8 explanatory variables to predict fixed airflow limitation, with a greater number of factors contributing to this problem. Nevertheless, the mathematical model had very good discrimination and good calibration.

Another point to take into account is that we did not have a sufficient sample size to externally validate the points system constructed. For this, a completely different sample should be available with at least 100 patients with fixed airflow limitation¹⁶. Our team is collecting a new sample for this purpose, and this type of study should also be carried out in other geographical areas to determine whether the predictive model is satisfactory for detecting fixed airflow limitation.

We would like to note that our points system is only applicable to patients without COPD who are current smokers and between the ages of 40 and 75 years. Patients with COPD who have no history of smoking have also been excluded, as they already have the disease and there is no point in screening. The exclusion of ex-smokers and elderly people could lead to an increase in the prevalence of fixed airflow limitation, and it should be verified in another study whether the model is applicable to these people, since the variables age, Fagerström test, and pack-years can be evaluated in these excluded patients. If similar results are obtained, the target population of our predictive model could be expanded.

Comparison with existing literature. The technique used to confirm the presence of COPD is still spirometry and respiratory symptoms¹², but its use is limited and the effectiveness of extending it to the entire population is unknown¹¹. Consequently, several authors have proposed the possibility of first using questionnaires aimed at detecting and classifying patients at high risk of COPD, in conjunction with the intermediate use of a pocket spirometer, and followed by confirmation with conventional spirometry in cases with low FEV₁/FEV₆ figures^{11,27}.

The older predictive models had major statistical limitations (Table 1). We used the recommended guidelines for developing a predictive model^{14–16} in order to develop a simple algorithm based on the data obtained in our study indicating the population at risk of COPD and that could go undetected. The tool can be used quickly in daily clinical practice by the primary care physician, or even with the development of a mobile application (app) that facilitates self-diagnosis by the patient (fixed airflow limitation), leading directly to the need for spirometry and assessment of respiratory symptoms to confirm the presence or absence of COPD and thus preventing the high levels of underdiagnosis. Consequently, this tool can be very useful, since the estimated worldwide prevalence of COPD is 1% in the general population and 8–10% in those aged over 40 years²⁸, and between 2.1% to 26.1% in Europe depending on country, method and population²⁹.

Regarding risk factors associated with underdiagnosis, age and smoking are already recognized as the main risk factors for underdiagnosis^{3,12,20,30–34}. Other studies refer to a higher risk of underdiagnosed COPD in cases of low educational level³, low socioeconomic status²⁰, previous respiratory symptoms or high BMI (≥ 30 kg/m²)², whilst others associate it with low BMI¹¹. Underdiagnosis has also been associated with increased comorbidity, especially cardiovascular^{5,19,32,35,36}. Therefore, we can confirm correspondence with the data found in our study.

Implications for research and/or practice. Our study provides a scoring system that is very easy to use in daily clinical practice that can facilitate the request for spirometry based on a calculated risk, making it a good model for COPD screening. For its use, once the result has been obtained in smokers who have attended the health care centre (opportunistic screening), the physician must evaluate the need for confirmation by spirometry, according to patient characteristics and cost-effectiveness. In addition, understanding their COPD risk may well encourage smokers who are already considering quitting to take the definitive step³⁷. Because of its ease of use, this scoring system can be extended to the entire population through mobile applications (apps), enabling patients to easily understand their risk and thus serving as a coercive measure for quitting. In other words, the points obtained for age cannot be modified, but it is visible on the scale that quitting smoking can improve scores, both in nicotine dependence and in respiratory symptoms. It is also a tool that can save time in primary care consultations and in the use of spirometry by applying the technique only in cases with a high probability of fixed airflow limitation. Finally, this model (and those in Table 1) should be externally validated in other populations and its cost-effectiveness verified in order to be able to extend its use in primary care in other centres and clinical settings. It also raises issues that open lines of investigation in the action against smoking and target population.

Data Availability

The data that support the findings of this study are available from Esther Román-Conejos but restrictions apply to the availability of these data, which were used under license for the current study, and so are not publicly available. Data are however available from the authors upon reasonable request and with permission of the Clinical Trials Ethics Committee of the Department of Health of the Generalitat Valenciana (General University Hospital of Elche, Alicante).

References

- Lozano, R. *et al.* Global and regional mortality from 235 causes of death for 20 age groups in 1990 and 2010: a systematic analysis for the Global Burden of Disease Study 2010. *Lancet* **380**, 2095–2128 (2012).
- Casas Herrera, A. *et al.* PUMA Team. COPD Underdiagnosis and Misdiagnosis in a High-Risk Primary Care Population in Four Latin American Countries. A Key to Enhance Disease Diagnosis: The PUMA Study. *PLoS One* **11**, e0152266 (2016).
- Bernd, L. *et al.* BOLD Collaborative Research Group, the EPI-SCAN Team, the PLATINO Team, and the PREPOCOL Study Group. Determinants of underdiagnosis of COPD in national and international surveys. *Chest* **148**, 971–985 (2015).
- Quach, A. *et al.* Prevalence and underdiagnosis of airway obstruction among middle-aged adults in northern France: The ELISABET study 2011–2013. *Respir. Med.* **109**, 1553–1561 (2015).
- Soriano, J. B. *et al.* Recent trends in COPD prevalence in Spain: a repeated cross-sectional survey 1997–2007. *Eur. Respir. J.* **36**, 758–765 (2010).
- Sørheim, I. C. *et al.* Gender differences in COPD: are women more susceptible to smoking effects than men? *Thorax* **65**, 480–485 (2010).
- Clotet, J., Gómez-Arbonés, X., Ciria, C. & Albalad, J. M. Spirometry is a good method for detecting and monitoring chronic obstructive pulmonary disease in high-risk smokers in primary health care. *Arch. Bronconeumol.* **40**, 155–159 (2004).
- Soriano, J. B. *et al.* High prevalence of undiagnosed airflow limitation in patients with cardiovascular disease. *Chest* **137**, 333–340 (2010).
- García-Río, F. E. Normativa SEPAR. *Arch. Bronconeumol.* **49**, 388–401 (2013).
- Clotet, J. *et al.* Espirometría como método de cribado y de intervención antitabaco en fumadores de alto riesgo en atención primaria. *Aten. Prim.* **44**, 328–334 (2012).
- van Schayck, C. P. *et al.* Comparison of existing symptom-based questionnaires for identifying COPD in the general practice setting. *Respirology* **10**, 323–333 (2005).
- Vogelmeier, C. F. *et al.* Global Strategy for the Diagnosis, Management, and Prevention of Chronic Obstructive Lung Disease 2017 Report. *Respirology* **22**, 575–601 (2017).
- Llordés, M. *et al.* Prevalence, Risk Factors and Diagnostic Accuracy of COPD Among Smokers in Primary Care. *COPD* **12**, 404–412 (2015).
- Collins, G. S., Mallett, S., Omar, O. & Yu, L. M. Developing risk prediction models for type 2 diabetes: a systematic review of methodology and reporting. *BMC Med.* **9**, 103 (2011).
- Van Calster, B. *et al.* A calibration hierarchy for risk models was defined: from utopia to empirical data. *J. Clin. Epidemiol.* **74**, 167–176 (2016).
- Palazón-Bru, A., Folgado-de la Rosa, D. M., Cortés-Castell, E., López-Cascales, M. T. & Gil-Guillén, V. F. Sample size calculation to externally validate scoring systems based on logistic regression models. *PLoS One* **12**, e0176726 (2017).
- Sociedad Española de Medicina de Familia y Comunitaria & Sociedad Española de Neumología y Cirugía Torácica. *Atención Integral al paciente con Enfermedad Pulmonar Obstructiva Crónica (EPOC): Desde la Atención Primaria a la Especializada*. Barcelona, (SEMFYC ediciones, 2010).
- Hill, K. *et al.* Prevalence and underdiagnosis of chronic obstructive pulmonary disease among patients at risk in primary care. *CMAJ* **182**, 673–678 (2010).
- Almagro, P. *et al.* Underdiagnosis and prognosis of chronic obstructive pulmonary disease after percutaneous coronary intervention: a prospective study. *Int. J. Chron. Obstruct. Pulmon. Dis.* **10**, 1353–1361 (2015).
- Kainu, A. *et al.* COPD in Helsinki, Finland: socioeconomic status based on occupation has an important impact on prevalence. *Scand. J. Public Health* **41**, 570–578 (2013).
- Fagerström, K. & Schneider, N. Measuring nicotine dependence a review of the Fagerström tolerance questionnaire. *J. Behav. Med.* **12**, 159–182 (1989).
- Calle, M., Morales, B. & Rodríguez, J. L. Exacerbación de la EPOC. *Arch. Bronconeumol.* **46**, 21–25 (2010).
- Pavlou, M. *et al.* How to develop a more accurate risk prediction model when there are few events. *BMJ* **351**, h3868 (2015).
- Sullivan, L. M., Massaro, J. M. & D'Agostino, R. B. Sr. Presentation of multivariate data for clinical use: The Framingham Study risk score functions. *Stat. Med.* **23**, 1631–1660 (2004).
- Steyerberg, E. W. *et al.* Internal validation of predictive models: efficiency of some procedures for logistic regression analysis. *J. Clin. Epidemiol.* **54**, 774–781 (2001).
- Izquierdo Alonso, J. L., De Lucas Ramos, P. & Rodríguez Glez-Moro, J. M. grupo de estudio CONSISTE. The use of the lower limit of normal as a criterion for COPD excludes patients with increased morbidity and high consumption of health-care resources. *Arch. Bronconeumol.* **48**, 223–228 (2012).
- Pleasant, R. A. *et al.* Targeting Persons With or At High Risk for Chronic Obstructive Pulmonary Disease by State-based Surveillance. *COPD* **12**, 680–689 (2015).
- Halbert, R. J. *et al.* Global burden of COPD: systematic review and meta-analysis. *Eur. Respir. J.* **28**, 523–532 (2006).
- Atsou, K., Chouaid, C. & Hejblum, G. Variability of the chronic obstructive pulmonary disease key epidemiological data in Europe: systematic review. *BMC Med.* **9**, 7 (2011).
- Markun, S., Rosemann, T., Dalla-Lana, K. & Steurer-Stey, C. The Impact of Case Finding on the Recruitment Yield for COPD Research in Primary Care: An Observational Study. *Respiration* **92**, 308–315 (2016).
- Menezes, A. M. *et al.* Chronic obstructive pulmonary disease in five Latin American cities (the PLATINO study): a prevalence study. *Lancet* **366**, 1875–1881 (2005).
- Utsugi, H. *et al.* Associations of lifelong cigarette consumption and hypertension with airflow limitation in primary care clinic outpatients in Japan. *Respir. Investig.* **54**, 35–43 (2016).
- Lindberg, A., Bjerg, A., Rönmark, E., Larsson, L. G. & Lündback, B. Prevalence and underdiagnosis of COPD by disease severity and the attributable fraction of smoking Report from the Obstructive Lung Disease in Northern Sweden Studies. *Respir. Med.* **100**, 264–272 (2006).
- Tinkelman, D. G., Price, D., Nordyke, R. J. & Halbert, R. J. COPD screening efforts in primary care: what is the yield? *Prim. Care Respir. J.* **16**, 41–48 (2007).
- Yamasaki, A. *et al.* COPD is frequent in conditions of comorbidity in patients treated with various diseases in a university hospital. *Int. J. Chron. Obstruct. Pulmon. Dis.* **5**, 351–355 (2010).
- Názara-Otero, C. A. & Balloira-Villar, A. The continuum of COPD and cardiovascular risk: A global scenario of disease. *Clin. Investig. Arterioscler.* **27**, 144–147 (2015).
- Parkes, G., Greenhalgh, T., Griffin, M. & Dent, R. Effect on smoking quit rate of telling patients their lung age: the Step2quit randomised controlled trial. *BMJ* **336**, 598–600 (2008).
- López Varela, M. V. *et al.* PUMA Team. Development of a simple screening tool for opportunistic COPD case finding in primary care in Latin America: The PUMA study. *Respirology* **21**, 1227–1234 (2016).
- Llordés, M. *et al.* Which is the Best Screening Strategy for COPD among Smokers in Primary Care? *COPD* **14**, 43–51 (2017).
- Price, D. B. *et al.* Symptom-based questionnaire for identifying COPD in smokers. *Respiration* **73**, 285–295 (2006).
- Price, D. B., Tinkelman, D. G., Nordyke, R. J., Isonaka, S. & Halbert, R. J., COPD Questionnaire Study Group. Scoring system and clinical application of COPD diagnostic questionnaires. *Chest* **129**, 1531–1539 (2006).

Acknowledgements

The authors thank Maria Repice and Ian Johnstone for help with the English language version of the text.

Author Contributions

E.R. designed the study, collected the data and drafted the manuscript. A.P. participated in the study design, performed the statistical analysis and helped draft the manuscript. D.M.F. participated in the study design, developed the mobile application and reviewed critically the manuscript. M.S, M.M.R. V.F.G. and E.C. participated in the study design and reviewed critically the manuscript. All the authors approved the final version of the manuscript to be considered for publication.

Additional Information

Competing Interests: The authors declare no competing interests.

Publisher's note: Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.



Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The images or other third party material in this article are included in the article's Creative Commons license, unless indicated otherwise in a credit line to the material. If material is not included in the article's Creative Commons license and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this license, visit <http://creativecommons.org/licenses/by/4.0/>.

© The Author(s) 2018