

PULMONARY PERSPECTIVE

Early Diagnosis and Treatment of Chronic Obstructive Pulmonary Disease

The Costs and Benefits of Case Finding

Shawn D. Aaron^{1*}, Maria Montes de Oca^{2,3*}, Bartolome Celli⁴, Surya P. Bhatt⁶, Jean Bourbeau⁷, Gerard J. Criner⁸, Dawn L. DeMeo⁵, David M. G. Halpin⁹, MeiLan K. Han¹⁰, John R. Hurst¹¹, Jamuna K. Krishnan¹², David Mannino¹³, Job F. M. van Boven¹⁴, Claus F. Vogelmeier¹⁵, Jadwiga A. Wedzicha¹⁶, Barbara P. Yawn¹⁷, and Fernando J. Martinez¹⁸

¹The Ottawa Hospital Research Institute, Department of Medicine, University of Ottawa, Ottawa, Ontario, Canada; ²Universidad Central de Venezuela, Caracas, Venezuela; ³Hospital Centro Médico de Caracas, Caracas, Venezuela; ⁴Pulmonary and Critical Care Medicine and ⁵Channing Division of Network Medicine, Brigham and Women's Hospital, Harvard Medical School, Boston, Massachusetts; ⁶Pulmonary, Allergy and Critical Care Medicine, University of Alabama at Birmingham, Birmingham, Alabama; ⁷Department of Medicine, McGill University, Montreal, Quebec, Canada; ⁸Lewis Katz School of Medicine at Temple University, Philadelphia, Pennsylvania; ⁹University of Exeter Medical School, University of Exeter, Exeter, United Kingdom; ¹⁰Division of Pulmonary & Critical Care, University of Michigan, Ann Arbor, Michigan; ¹¹UCL Respiratory, University College London, London, United Kingdom; ¹²Division of Pulmonary and Critical Care, Weill Cornell Medicine, New York, New York; ¹³College of Medicine, University of Kentucky, Lexington, Kentucky; ¹⁴Department of Clinical Pharmacy & Pharmacology, University Medical Center Groningen, Groningen Research Institute for Asthma and COPD, University of Groningen, Groningen, The Netherlands; ¹⁵Philipps-Universität Marburg, German Center for Lung Research, Marburg, Germany; ¹⁶National Heart and Lung Institute, Imperial College London, London, United Kingdom; ¹⁷Department of Family Medicine and Community Health, University of Minnesota, Minneapolis, Minnesota; and ¹⁸Cornell Medical College, New York, New York

ORCID IDs: 0000-0002-7266-8371 (B.C.); 0000-0003-2009-4406 (D.M.G.H.); 0000-0002-7246-6040 (J.R.H.); 0000-0002-1460-5276 (J.K.K.); 0000-0003-3646-7828 (D.M.); 0000-0003-2368-2262 (J.F.M.v.B.).

Undiagnosed Chronic Obstructive Pulmonary Disease: A Global Problem

In 2020, the prevalence of clinician-diagnosed chronic obstructive pulmonary disease (COPD) among U.S. adults aged ≥ 18 years was 5.6% (95% confidence interval [CI], 5.2–6.2) (1). This proportion is likely to be just the tip of an iceberg because studies from across the world suggest that as many as 70–90% of individuals with COPD remain undiagnosed (2). The 2007–2012 National Health and Nutritional Examination Survey found obstructive lung disease in 13.4% of randomly selected American adults; however, 71% were undiagnosed (3).

A collaborative study assessed the prevalence of undiagnosed COPD in 27 countries. Among 30,874 participants, the prevalence of COPD (FEV₁/FVC below the lower limit of normal) was 9.7%, and 81.4% of COPD cases were undiagnosed (4). The

problem of COPD underdiagnosis was worse in low- and middle-income countries (LMICs). Centers in Turkey, India, Nigeria, the Philippines, Colombia, Nepal, Peru, and Uganda reported that more than 90–95% of COPD cases were undiagnosed (4, 5).

The Clinical and Societal Burden of Undiagnosed COPD

Individuals with undiagnosed COPD are afflicted by poor quality of life, experience exacerbations like those seen in patients with diagnosed COPD, and are at increased risk of death (6, 7). In addition, their respiratory symptoms lead to work absenteeism, productivity loss, and consumption of healthcare resources (8). This perspective reviews the burden of disease in people with COPD who have not been diagnosed, considers why a diagnosis is not made or is

delayed, and summarizes how patients could be identified and diagnosed earlier.

Symptoms and Health-related Quality of Life

A Canadian study of individuals with symptomatic undiagnosed COPD identified via case finding compared with age-matched control subjects showed that adjusted mean between-group differences in COPD Assessment Test (CAT), St. George's Respiratory Questionnaire (SGRQ), and 36-item Short Form Health Survey total scores were 13, 33, and –20 points, respectively (6). These values exceed the established minimal clinically important differences for these questionnaires. The study concluded that undiagnosed COPD was associated with greater symptom burden, poorer disease-specific health-related quality of life (HRQOL), and poorer overall general health status than in age-matched control subjects (6). Similar findings have been

(Received in original form November 19, 2023; accepted in final form February 14, 2024)

*These authors are co-primary authors.

Author Contributions: Conception and design: S.D.A., M.M.d.O., F.J.M., and M.K.H. Drafting the manuscript for important intellectual content: S.D.A., M.M.d.O., F.J.M., M.K.H., B.C., S.P.B., J.B., G.J.C., D.L.D., D.M.G.H., J.R.H., J.K.K., D.M., J.F.M.v.B., C.F.V., J.A.W., and B.P.Y.

Correspondence and request for reprints should be addressed to Maria Montes de Oca, M.D., Centro Médico de Caracas, Anexo B, San Bernardino, Caracas 1011, Venezuela. E-mail: montesdeoca.maria@gmail.com.

Am J Respir Crit Care Med Vol 209, Iss 8, pp 928–937, Apr 15, 2024

Copyright © 2024 by the American Thoracic Society

Originally Published in Press as DOI: 10.1164/rccm.202311-2120PP on February 15, 2024

Internet address: www.atsjournals.org

reported by Miravittles and colleagues, who also showed poorer HRQOL and reduced activities of daily living in Spanish individuals with undiagnosed COPD (9).

Exacerbation Events and Healthcare Use

Individuals with previously undiagnosed COPD discovered via population-based screening with spirometry were followed for a median of 6.1 years. Their age- and sex-adjusted hazard ratios for exacerbations and pneumonia were 15.5 (95% CI, 11.0–21.8) and 2.8 (95% CI, 2.4–3.3), respectively, compared with individuals without COPD (7). Other authors reported more physician visits for breathing difficulties in the previous year in individuals with undiagnosed COPD than in healthy control subjects (26% vs. 9%; *P* < 0.001), without differences in emergency department visits or hospitalizations for respiratory illness (6).

The CanCOLD study found that the exacerbation rate in undiagnosed, untreated individuals was about half that of individuals diagnosed with COPD (0.30 exacerbation events/patient-year vs. 0.63/patient-year); however, undiagnosed individuals used health services to treat exacerbation events in an equivalent manner to individuals who had received a previous COPD diagnosis (10). The authors concluded that people with undiagnosed COPD contribute greatly to the healthcare burden of COPD.

Work Absenteeism and Productivity

Gerstein and colleagues reported that individuals with undiagnosed COPD

discovered via case finding had significantly higher work absenteeism (missing work because of a respiratory illness during the previous week) than healthy control subjects (14% vs. 2%, respectively), as well as greater losses in both work productivity and regular daily activities. Interestingly, work absenteeism or productivity was similar between undiagnosed and previously diagnosed patients with COPD (6).

De Sousa Sena and colleagues reported that work absenteeism was higher in subjects with physician-diagnosed COPD than in those with undiagnosed COPD (14.6% vs. 5.7%). However, presenteeism (the act of attending work while sick, resulting in decreased work quality or quantity) and overall work productivity losses were similar among the groups. Individuals with undiagnosed COPD who had a high symptom burden (CAT score ≥ 10) were more likely to have experienced work productivity loss (8).

Mortality

In Denmark, subjects with undiagnosed, symptomatic COPD had age- and sex-adjusted hazard ratios for death of respiratory causes of 4.3 (95% CI, 2.8–6.7) and 2.0 (95% CI, 1.8–2.3) for death of any cause compared with individuals without COPD (7). An analysis of U.S. data demonstrated that among people with undiagnosed COPD, the risk for mortality over 20 years of follow-up was increased (hazard ratio, 1.23; 95% CI, 1.08–1.40) relative to those without COPD (3).

Why Does COPD Go Undiagnosed until Disease Is Far Advanced?

Clinical diagnosis of COPD is often delayed in many patients until disease is far advanced. Indeed, a study using British Columbia’s administrative data records showed that 44% of incident cases of COPD detected from 2001 to 2010 were only first diagnosed with COPD when they had a COPD-related hospitalization (11). An overview of potential patient, healthcare provider, and healthcare system factors that may be associated with underdiagnosis of COPD is shown in Figure 1.

Patient-related Factors

Some patients with COPD may go undetected because of underrecognition and/or underreporting of symptoms. Patients with COPD may adapt their activities to minimize their respiratory symptoms, deciding that their symptoms do not merit a discussion with their physician (12, 13). Chronic breathlessness may be attributed to a normal, inevitable part of aging (14). Furthermore, some patients may be reluctant to discuss their respiratory symptoms because of shame associated with smoking (15). In one sample of Dutch adults, only 34% of participants with undiagnosed obstructive lung disease had previously discussed their symptoms with a physician (16).

Failure to report respiratory symptoms is not the only factor accounting for underdiagnosis. A study evaluating undiagnosed symptomatic obstructive lung

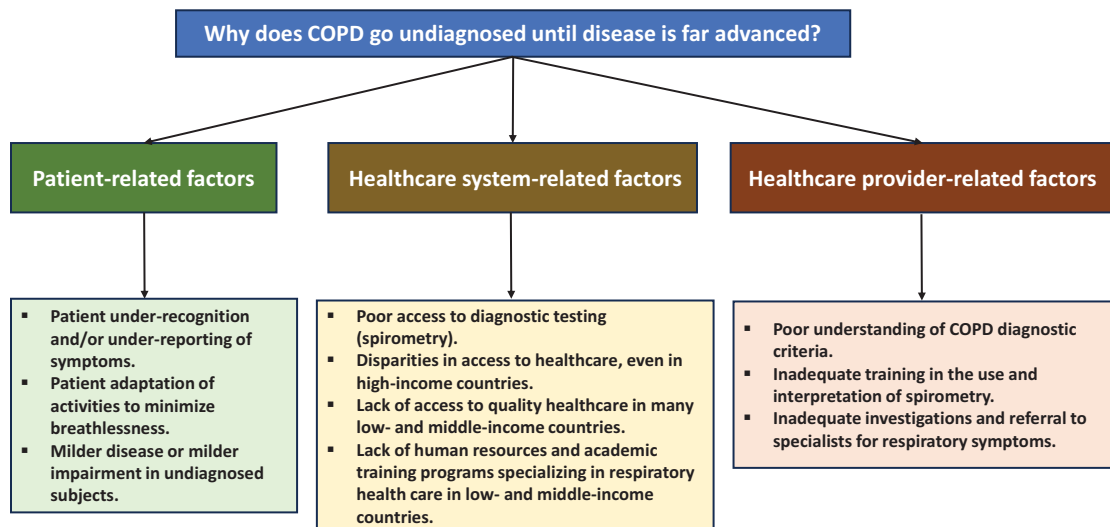


Figure 1. Patient, healthcare provider, and healthcare system factors that may be associated with underdiagnosis of chronic obstructive pulmonary disease (COPD).

disease (detected through active case finding) showed that 69% of undiagnosed participants reported that they had previously discussed their respiratory symptoms with a physician, and 18% had been given an alternative diagnosis, other than obstructive lung disease, to explain their respiratory symptoms (17).

Some underdiagnosed COPD may be attributable to milder disease or milder impairment, especially if undiagnosed COPD is found via population-based screening rather than active case finding of symptomatic individuals. Çolak and colleagues showed that among subjects with undiagnosed COPD found via population-based screening, 71% were symptomatic; however, the majority (73%) had minimal symptoms (modified Medical Research Council dyspnea scale score, <2) (7). Similarly, a Spanish population-based study found that individuals with diagnosed COPD had more severe airflow obstruction, higher cumulative tobacco consumption, and more severely impaired HRQOL than undiagnosed subjects (9).

Healthcare Provider Factors

Underdiagnosed COPD may also be associated with systematic health system deficiencies at the healthcare provider level (18). Primary care practitioners have limited time, lack of financial incentives, and multiple competing priorities (18). The effectiveness of COPD therapies in mild disease may be underestimated, and consequently therapeutic nihilism in primary care may play a role in underdiagnosis (18). Diagnosis may also be limited by physicians' sometimes poor understanding of COPD diagnostic criteria, inadequate training in the use and interpretation of spirometry, poor reimbursement for the test, and poor access to diagnostic spirometry in many communities (19, 20).

Healthcare System Factors

Studies from high-income countries indicate that spirometry is underused. In Sweden, only one-third of patients with physician-diagnosed COPD had their diagnosis confirmed with spirometry (21). A population study of all individuals newly diagnosed with COPD in Ontario, Canada, showed that only 36% had received spirometry (19). Another Canadian study showed that those with undiagnosed symptomatic COPD were less likely to have undergone spirometry or to have been referred to a specialist than subjects with diagnosed COPD, suggesting that

underdiagnosis is associated with inadequate investigations and specialist referral for respiratory symptoms (17). Deficiencies in the provision of respiratory health care may be more acute in LMICs (22). In Latin America, data from the PLATINO (Proyecto Latinoamericano de Investigación en Obstrucción Pulmonar) study revealed that only 16% of subjects with undiagnosed COPD reported ever undergoing spirometry (23).

COPD diagnosis is based on post-bronchodilator (post-BD) spirometry; however, measurement of post-BD airflow adds extra time and complexity to the test, and this may potentially discourage the use of diagnostic spirometry. An analysis of 10,192 tobacco-exposed subjects from the COPDgene (Genetic Epidemiology of COPD) study showed that both pre- and post-BD spirometry were associated with dyspnea, radiographic emphysema, gas trapping, and impaired exercise capacity (24). The predictive value of pre- and post-BD spirometry for exacerbations, change in FEV₁, dyspnea, and exercise capacity was relatively similar, but post-BD measures better predicted mortality. On the basis of this evidence, the latest Global Initiative for Chronic Obstructive Lung Disease (GOLD) document states that if pre-BD spirometry does not show airflow obstruction, then a post-BD measurement is not necessary (25). This may help mitigate a perceived barrier to spirometry. However, if airflow obstruction is found on pre-BD spirometry, then diagnosis of COPD should be confirmed with post-BD measurements.

Many healthcare settings, especially in LMICs, do not have guidelines for the management of COPD (26), and even where guidelines exist, there are significant challenges to their effective implementation (27). In Latin America, patients who did not visit a physician or who only attended a general practitioner had greater risk of undiagnosed COPD than those who saw a specialist (28). The utility of diagnosis may be deprioritized where there are problems with access to affordable interventions, leaving much unmet need (29).

Early Diagnosis of COPD: Screening versus Case Finding

Although no consensus definition exists for "early diagnosis of COPD," it is generally

accepted that early diagnosis happens when an individual undergoes targeted assessment for COPD and the disease is identified before a conventional diagnosis is made by the individual's healthcare professionals (30, 31). Early diagnosis can be potentially achieved by either screening or case finding, and differentiating between these two methodologies is important. The goal of screening is to detect unrecognized COPD in apparently healthy individuals. As such, screening is performed in the general population and involves testing mostly asymptomatic individuals, potentially based on demographic factors (such as age or smoking history). This approach has been shown to be relatively low yield and cost inefficient for detecting unrecognized COPD. For these reasons, the U.S. Preventive Services Task Force (USPSTF) recommends against screening for COPD in asymptomatic individuals (32). In contrast, case finding involves an assessment of an individual's respiratory symptoms and disease risk factors to determine the need for further testing. Case finding is a potential strategy for identifying undiagnosed individuals at high risk of COPD to allow earlier identification of disease and to direct resources to these individuals.

Active versus Opportunistic Case Finding

The objective of targeted case finding is to detect the largest possible number of cases in the community through assessment of at-risk individuals who present with symptoms (33). Case-finding approaches include active or opportunistic case finding, and the two differ on the basis of approach and the target population. Active case finding involves proactively searching for individuals at higher risk of the condition, often based on presence of symptoms. In active case finding, respiratory symptoms and risks for COPD are elicited from the individual via questionnaire, and, on the basis of responses if the individuals exceed a preset threshold, they are targeted to receive spirometry. In contrast, opportunistic case finding involves a passive approach to case identification, and screening of individuals occurs when they present themselves for healthcare services for reasons unrelated to the condition being screened for. Opportunistic case finding does not focus on a specific group or population. Instead, it

captures cases as they arise in the general patient population during routine healthcare encounters.

An active case-finding strategy targeting symptomatic individuals with risk factors for COPD was more effective at detecting new COPD cases than opportunistic case finding conducted in the general patient population during routine healthcare encounters (adjusted odds ratio, 2.34) and was more cost-effective (£333 vs. £376 per case detected, respectively [34]). However, opportunistic case finding conducted in specific target populations (such as smokers being screened for lung cancer) has been shown to be effective and relatively inexpensive (35, 36).

Case-Finding Tools

The optimal strategy for identifying undiagnosed COPD remains an area of controversy. Different tools have been assessed for COPD case finding (questionnaires, handheld devices, or a combination of tools), with wide heterogeneity among them.

Questionnaires

Several COPD risk questionnaires have been developed and validated in different populations. Among them, the COPD Diagnostic Questionnaire (CDQ), the COPD Population Screener (COPD-PS), the Lung Function Questionnaire (LFQ), COPD assessment in primary care to identify undiagnosed respiratory disease and exacerbation risk (CAPTURE), and the PUMA (Prevalence Study and Regular Practice, Diagnosis and Treatment, among General Practitioners in Populations at Risk of COPD in Latin America) study questionnaire (PUMA) are the most widely used (37–53). All include the following elements: age, smoking history, and respiratory symptoms. In addition, some include other items, such as body mass index, allergic history, sex, previous use of spirometry, and history of acute respiratory diseases. Table 1 shows the performance results (area under the curve [AUC]) reported for some questionnaires. The Undiagnosed COPD and Asthma Population Questionnaire (UCAP) detects both undiagnosed COPD and asthma (53). The PUMA and COLA (COPD in LMICs Assessment) questionnaires were developed in LMIC settings and had performance similar to those of other questionnaires when

they were used in LMIC settings (5). The evidence indicates that all diagnostic case-finding questionnaires are valuable tools to discriminate between subjects with or without COPD (37–53).

A meta-analysis aimed at assessing the best practice in COPD case finding in a primary care setting found that the LFQ was a slightly stronger tool to identify high-risk individuals who require diagnostic evaluation for COPD than the CDQ and the COPD-PS questionnaires (54). A study from China showed that the LFQ and the COPD-PS showed the highest diagnostic accuracy and the CAPTURE showed the lowest. Except for CAPTURE, the other five questionnaires demonstrated moderate diagnostic accuracy (55). Another Chinese study in primary care settings showed that PUMA as a screening tool performed better than the CDQ and COPD-PS (AUC, 0.75 vs. 0.66 and 0.61, respectively) (42).

Handheld Airflow Measurement Devices

Measurements such as the peak expiratory flow (PEF) and devices such as microspirometers (COPD-6 and the PiKo-6) have been widely tested. In general, studies have reported that microspirometers have high accuracy in the diagnosis of COPD; therefore, they could be of great diagnostic help in primary care settings with limited resources. Table 1 shows the AUC reported for COPD-6 and PiKo-6 using the FEV₁/FEV₆ ratio.

A meta-analysis showed an overall AUC of microspirometers (including COPD-6 and PiKo-6) of 0.84 (95% CI, 0.80–0.89). In addition, it showed that microspirometers were the most accurate tools for COPD diagnosis compared with diagnostic questionnaires (19). Another meta-analysis reported that the pooled AUC of portable spirometers was 0.91 (0.89–0.94). The accuracy of the PiKo-6 was higher (0.95) than that of COPD-6 (0.91) and PEF (0.82) (43).

Combining Questionnaires with Handheld Airflow Measurement Devices

The use of combined tools has shown better diagnostic accuracy than either tool alone. Table 1 shows results of studies that have assessed the accuracy of combining tools for COPD diagnosis (44, 50, 52, 56).

Soriano and colleagues showed that the combination of COPD-PS and PEF (AUC, 0.76) had greater diagnostic accuracy than

COPD-PS alone (AUC, 0.71) or PEF alone (AUC, 0.66). Case finding for COPD using COPD-PS + PEF led to a 90% reduction in the number of spirometry tests performed (56). Another study showed that the use of the combination of the CAPTURE questionnaire with a PEF device yielded a higher AUC (0.95) than the CAPTURE questionnaire alone (0.93) (50).

Taken together, these findings suggest that the combined use of a questionnaire and a handheld device is a more effective strategy for identifying individuals at increased risk for COPD, although questionnaires used alone are still valuable tools to predict COPD. Other potentially sensitive tools for COPD case finding, such as DL_{CO}, oscillometry, and nitrogen washout, need further research. To date, no prospective study has compared the diagnostic accuracy of these tools with screening questionnaires or handheld flow devices, and the practicality of using such approaches in community settings needs to be assessed.

Potential Target Populations for COPD Case Finding

Because both COPD and lung cancer share similar risk factors, using a single test to simultaneously target early diagnosis of both diseases may be cost-effective. Mets and colleagues reported that low-dose computed tomography (LDCT) scans for lung cancer screening could identify participants with COPD. The diagnostic model showed AUCs of 0.87 for participants with symptoms and 0.78 for those without symptoms (35). In some LDCT lung cancer screening studies, the prevalence of COPD within the study cohorts was between 19% and 57%, and between 49% and 86% of these individuals were undiagnosed (36, 57, 58). Emphysema prevalence rates established by CT scan in those with known and undiagnosed COPD were 73% and 68%, respectively, suggesting that imaging-confirmed emphysema can help find undiagnosed COPD cases in current or former smokers who are at risk for lung cancer (36, 57).

Tang and colleagues evaluated a residual network-based software pipeline for identifying people with COPD using LDCT scans obtained for lung cancer screening on smokers (59). The results showed that using artificial intelligence in case-finding approaches was feasible; the best performing

networks achieved AUCs of 0.88 and 0.89 in three-way cross-validation experiments in two independent cohorts (59).

Multimorbidity is the norm rather than the exception in patients with COPD. The coexistence of COPD with cardiac diseases, including ischemic heart disease or heart failure, is common, and these diseases share similar risk factors, including cigarette smoking, aging, low socioeconomic status, and sedentary behavior (60). The co-occurrence of COPD and cardiovascular diseases has led investigators to recommend a proactive search for these conditions using diagnostic tests that help to differentiate COPD symptoms from those of cardiac diseases (61). Efforts to enhance detection of COPD in patients with cardiovascular disease present an opportunity to explore broader medical collaborations among primary care physicians, pulmonologists, and cardiologists and potentially allow an assessment of the general wellness of their patients.

Taken together, these findings suggest that incorporating COPD case-finding tools and targeting at-risk populations, such as those with cardiac disease, or those being screened for lung cancer is feasible and could identify a significant number of individuals with undiagnosed COPD. Figure 2 shows a possible proposal for COPD case finding.

Cost-Effectiveness of Case Finding

Lambe and colleagues compared the cost-effectiveness of a systematic case-finding program administered every 3 years to ever smokers aged ≥50 years against routine diagnostic process in U.K. primary care (62). The results indicated an incremental cost-effectiveness ratio (ICER) of systematic case finding versus current care of £16,596 per additional quality-adjusted life-year (QALY) gained. The active case-finding strategy was more expensive but more effective, with a greater number of QALYs gained over a lifetime time horizon (62). This ICER of £16,596/QALY is below the National Institute for Health and Care Excellence common threshold to consider an intervention cost-effective (i.e., £30,000/QALY). It is important to highlight that the treatment effect in this model only captured the benefits associated with inhaled medications, so the effect of other interventions that can slow progression of COPD and improve HRQOL, such as

Table 1. Performance of Chronic Obstructive Pulmonary Disease Case-Finding Strategies

Case-Finding Tools	Study	AUC	
Questionnaires			
COPD-PS	Ronaldson <i>et al.</i> (2018) (37)	0.66	
	Spyratos <i>et al.</i> (2017) (38)	0.79	
	Sogbetun <i>et al.</i> (2016) (39)	0.62	
	Llordés <i>et al.</i> (2017) (41)	0.65	
	Miravittles <i>et al.</i> (2012) (40)	0.79	
	Au-Doung <i>et al.</i> (2022) (42)	0.61	
	Zhou <i>et al.</i> (2022) (55)	0.74	
	Fujita <i>et al.</i> (2020) (44)	0.73	
	Ronaldson <i>et al.</i> (2018) (37)	0.67	
	Spyratos <i>et al.</i> (2017) (38)	0.80	
	Stanley <i>et al.</i> (2014) (45)	0.71	
	Frith <i>et al.</i> (2011) (46)	0.72	
	Kotz <i>et al.</i> (2008) (47)	0.65	
	Price <i>et al.</i> (2006) (48)	0.82	
CDQ	Au-Doung <i>et al.</i> (2022) (42)	0.66	
	Zhou <i>et al.</i> (2022)	0.73	
	LFQ	Spyratos <i>et al.</i> (2017) (38)	0.81
	Sogbetun <i>et al.</i> (2016) (39)	0.66	
	Zhou <i>et al.</i> (2022)	0.78	
	Simple PUMA	López Varela <i>et al.</i> (2016) (51)	0.70
		Lopez Varela <i>et al.</i> (2019) (49)	0.70–0.76
		Au-Doung <i>et al.</i> (2022) (42)	0.75
	Weighted PUMA CAPTURE	López Varela <i>et al.</i> (2016) (51)	0.79
		Quezada <i>et al.</i> (2017) (50)	0.93
		Martinez <i>et al.</i> (2017) (52)	0.79
		Zhou <i>et al.</i> (2022)	0.67
	UCAP-Q	Huynh <i>et al.</i> (2022) (53)	0.81–0.82
		Hand devices	
PEF	Ronaldson <i>et al.</i> (2018) (37)	0.77	
	Fujita <i>et al.</i> (2020) (44)	0.82	
	Labor <i>et al.</i> (2016) (73)	0.83	
	Kim <i>et al.</i> (2016) (74)	0.76	
	Llordés <i>et al.</i> (2017) (41)	0.87	
	Represas-Represas <i>et al.</i> (2016) (75)	0.80	
	Thorn <i>et al.</i> (2012) (76)	0.84	
	Miravittles <i>et al.</i> (2012) (40)	0.85	
	van den Bemt <i>et al.</i> (2014) (77)	0.94	
	Frith <i>et al.</i> (2011) (46)	0.85	
	Hidalgo Sierra <i>et al.</i> (2018) (78)	0.86	
	Lin <i>et al.</i> (2021) (79)	0.90	
	Chen <i>et al.</i> (2021) (80)	0.87	
	Spirobank Smart	Combination tools	
Changed CDQ + COPD-6 score		Fujita <i>et al.</i> (2020) (44)	0.87
COPD-PS + PEF		Soriano <i>et al.</i> (2018) (31)	0.76
CAPTURE + PEF		Quezada <i>et al.</i> (2017) (50)	0.95
		Martinez <i>et al.</i> (2017) (52)	0.90
Martinez <i>et al.</i> (2023) (81)	0.81		

Definition of abbreviations: AUC = area under the curve; CDQ = COPD diagnostic questionnaire; COPD-PS = COPD-Population Screener; LFQ = Lung Function Questionnaire; CAPTURE = COPD assessment in primary care to identify undiagnosed respiratory disease and exacerbation risk; PEF = peak expiratory flow; PUMA = PUMA study questionnaire; UCAP-Q = Undiagnosed COPD and Asthma Population Questionnaire.

smoking cessation, pulmonary rehabilitation, and self-management, was not considered. Therefore, it is very likely that the inclusion of nonpharmacologic interventions for COPD could have made systematic case finding even more cost-effective.

In a high-risk population for COPD in China, Qu and colleagues used a modeling approach and found that portable spirometer

screening was cost-saving compared with questionnaire screening and no screening (i.e., usual care), with incremental ICERs of 5,026 and 1,766 renminbi per QALY, respectively (63).

Johnson and colleagues assessed the cost-effectiveness of primary care-based case detection strategies for COPD in Canada using economic models relying on case

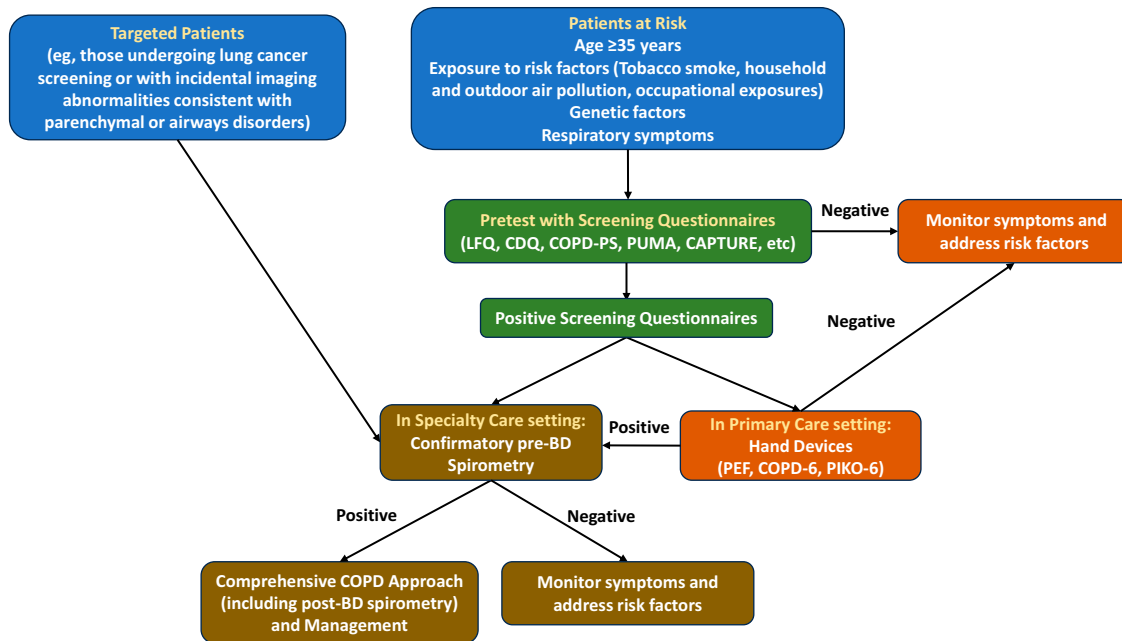


Figure 2. An algorithm for chronic obstructive pulmonary disease case finding. BD = bronchodilator; CAPTURE = COPD assessment in primary care to identify undiagnosed respiratory disease and exacerbation risk; CDQ = COPD Diagnostic Questionnaire; COPD = chronic obstructive pulmonary disease; COPD-PS = COPD Population Screener; LFQ = Lung Function Questionnaire; PEF = peak expiratory flow; PUMA = Prevalence Study and Regular Practice, Diagnosis and Treatment, among General Practitioners in Populations at Risk of COPD in Latin America.

simulations (64). The eligible patients (based on age, smoking history, or symptoms) received the CDQ or screening spirometry at 3- or 5-year intervals during routine visits to a primary care physician. Newly diagnosed patients received treatment for smoking cessation and guideline-based inhaler pharmacotherapy. The most efficient scenario (all patients aged ≥ 40 yr received the CDQ at 3-yr intervals) was associated with an incremental cost of \$287 and incremental effectiveness of 0.015 QALYs per eligible patient over the 20-year time horizon, resulting in an ICER of \$19,632/QALY compared with no case detection (64). Taken together, the results of these three modeling studies are consistent and suggest that both questionnaire-based methods and screening spirometry implemented in routine primary care visits are cost-effective methods for early COPD detection in those at risk by virtue of exposure history and/or symptoms.

Potential Benefits of Case Finding and Early Diagnosis of COPD

A retrospective study of U.K. primary care centers found that only 33% of patients identified with COPD were diagnosed within

5 years after first presentation and were considered early diagnoses. Those defined as receiving early diagnosis had slightly poorer lung function; however, they had a longer time to first exacerbation, decreased exacerbation rates (57 vs. 109 exacerbations/100 person-years), and decreased rate of hospitalizations over 3 years (65). Although this observational study could not establish causality, the study showed an association between early diagnosis of COPD and outcome benefits to both the patients and the healthcare system.

Similar findings were reported in a Swedish study which showed that receiving a COPD diagnosis soon after having visited a doctor for respiratory symptoms was associated with lower risk of exacerbations, fewer comorbidities, and lower costs than delayed diagnosis (66). It is important to note that both the U.K. and Swedish studies evaluated patients who were diagnosed conventionally by their physicians, and early diagnosis was not achieved through case finding.

Seven randomized controlled trials in adult smokers evaluated the effect of providing spirometry results (FEV₁ and/or lung age) in addition to counseling on smoking cessation rates compared with counseling alone (67). Although participants

did not necessarily report respiratory symptoms, these can be considered COPD case-finding studies conducted in an at-risk population. Only two studies showed improved results when smokers were informed of spirometry results (67). Although there is currently only limited evidence to suggest that the use of feedback from spirometry improves smoking quit rates when added to smoking cessation counseling, it needs to be noted that many of the individuals enrolled in these studies did not have COPD on spirometry.

Case Finding and Early Diagnosis of COPD Coupled to Interventions

Two comparative studies have combined active COPD case finding coupled with nonpharmacologic interventions to potentially improve patient health outcomes. One trial conducted in Australia assessed the effectiveness of early intervention by a practice nurse general practitioner team regarding HRQOL and process of care in patients with newly diagnosed COPD discovered via case finding compared with usual care. Among patients invited to a case-finding appointment, 16% attended and 18%

were diagnosed with COPD. Only 15% of patients in the intervention group saw the nurse for COPD care after case finding. There was no between-group difference in SGRQ score at follow-up. The investigators concluded that intervention uptake was low and had no additional beneficial effect over usual care on participants' HRQOL (68).

The second clinical trial was conducted in three LMICs. Individuals with COPD (GOLD grades B–D) were identified through case finding and randomized to a community health worker–supported 1-year self-management intervention or to standard of care. The trial did not find differences in SGRQ scores between the two groups at 1 year (69).

The USPSTF recommends against screening for COPD in asymptomatic adults, mainly because they concluded that there was no evidence to support pharmacological therapy in patients identified by screening (32). In contrast, the USPSTF recommends annual lung cancer screening with LDCT in adults aged 50 to 80 years who have a 20-pack-year smoking history and currently smoke or have quit within the past 15 years. It is interesting that this same approach is not recommended for COPD, which kills more people each year in the United States than lung cancer and shares the same risk factors. Importantly, the evidence indicates that, once diagnosed, COPD has therapies that not only improve lung function, health status, and lung function decline but also decrease exacerbation rates, hospitalizations, and risk of death.

Although no trials have coupled case finding and early pharmacologic treatment of COPD, evidence from randomized clinical trials suggests that treatment of mild and/or moderate COPD can lead to improved clinical outcomes. Two trials (SUMMIT [Study to Understand Mortality and Morbidity in COPD] and UPLIFT [Understanding Potential Long-Term Impacts on Function with Tiotropium]) found that long-acting beta agonists, long-acting muscarinic antagonists, inhaled corticosteroids, or long-acting beta agonists/inhaled corticosteroids reduced exacerbations in persons with symptomatic moderate COPD (70, 71). One trial (UPLIFT) found that tiotropium also reduced

exacerbations in a subgroup analysis of persons with minimal symptoms (i.e., GOLD grade A) (71). However, these trials enrolled individuals with physician-diagnosed COPD, and subjects were not discovered through case finding.

Two ongoing randomized controlled trials couple active case finding with interventions designed to improve health outcomes. The CAPTURE clinical trial is a pragmatic, cluster-randomized trial that explores the impact of using the CAPTURE COPD case-finding tool to identify clinically significant COPD in primary care. The study has randomized 68 practices to either usual care or clinician receipt of patient-level CAPTURE results. One-year follow-up chart reviews and participant surveys will assess the impact of sharing versus not sharing CAPTURE results with clinicians on clinical outcomes, including clinicians' initiation of recommendation-concordant COPD care and changes in CAT respiratory symptom scores in those diagnosed with COPD (72).

The UCAP (Undiagnosed COPD and Asthma in the Population) study is an ongoing Canadian clinical trial being conducted to explore the potential benefits of case finding to discover previously undiagnosed asthma or COPD in symptomatic individuals coupled with an intervention strategy. Participants with previously undiagnosed asthma or COPD identified in the community via case finding have been randomly assigned to one of two trial arms: early treatment with guideline-based therapy provided by a pulmonologist and nurse educator or standard care (6). This trial aims to determine if early intensive treatment for undiagnosed obstructive lung disease reduces healthcare use and improves QOL. A predefined subgroup analysis will explore outcomes in the COPD subgroup, exclusive of those with asthma. The trial has completed enrollment and concluded in January 2024.

Conclusions and Future Research

Undiagnosed COPD is a major global health problem. Most patients diagnosed

with COPD in clinical practice are recognized only when their disease is already relatively advanced and therapy is less effective. As has been shown for lung cancer detection in at-risk subjects, COPD case finding is a potential strategy for identifying individuals with COPD who are still undiagnosed to allow earlier identification of disease and to direct nonpharmacologic and pharmacologic treatments to these individuals.

Newer evidence suggests that active case finding of symptomatic people in the community and opportunistic case finding in people at risk for lung cancer or other comorbidities are effective. Three cost-effectiveness modeling studies from the United Kingdom, China, and Canada suggest that case finding for COPD is cost-effective with ICERs below \$50,000 per QALY gained (62–64). However, these economic models rely on case simulations rather than data derived from actual clinical studies. Although there are theoretical clinical and economic benefits of case finding, the causal benefits of case finding have not been conclusively established, because there are, to date, no randomized controlled trials that have demonstrated that patient outcomes are improved when COPD case-finding strategies are coupled to pharmacological or nonpharmacological interventions.

It is hoped that upcoming results from two ongoing randomized controlled trials will provide hard evidence demonstrating that early diagnosis of COPD achieved via case finding, coupled with pharmacologic and nonpharmacologic treatments for previously undiagnosed COPD, improves patient outcomes (6, 72). Subsequent health economic studies linked to these clinical trials can determine if the strategies are cost-effective.

A nihilistic approach to the growing health problem of COPD across the world has resulted in poor outcomes for patients with COPD. Case finding will provide opportunities for healthcare providers to intervene early and be proactive. ■

Author disclosures are available with the text of this article at www.atsjournals.org.

References

- Centers for Disease Control and Prevention. COPD data and statistics. Accessed September 15, 2023. Available from: <https://www.cdc.gov/copd/data.html>.
- Diab N, Gershon AS, Sin DD, Tan WC, Bourbeau J, Boulet LP, et al. Underdiagnosis and overdiagnosis of chronic obstructive pulmonary disease. *Am J Respir Crit Care Med* 2018;198:1130–1139.
- Martinez CH, Mannino DM, Jaimes FA, Curtis JL, Han MK, Hansel NN, et al. Undiagnosed obstructive lung disease in the United States. Associated factors and long-term mortality. *Ann Am Thorac Soc* 2015;12:1788–1795.
- Lamprecht B, Soriano JB, Studnicka M, Kaiser B, Vanfleteren LE, Gnatiuc 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* 2015;148:971–985.
- Siddharthan T, Pollard SL, Quaderi SA, Rykiel NA, Wosu AC, Alupo P, et al.; GECO Study Investigators. Discriminative accuracy of chronic obstructive pulmonary disease screening instruments in 3 low- and middle-income country settings. *JAMA* 2022;327:151–160.
- Gerstein E, Bierbrier J, Whitmore GA, Vandemheen KL, Bergeron C, Boulet LP, et al. Impact of undiagnosed COPD and asthma on symptoms, quality of life, healthcare utilization and work productivity. *Am J Respir Crit Care Med* 2023;208:1271–1282.
- Çolak Y, Afzal S, Nordestgaard BG, Vestbo J, Lange P. Prognosis of asymptomatic and symptomatic, undiagnosed COPD in the general population in Denmark: a prospective cohort study. *Lancet Respir Med* 2017;5:426–434.
- de Sousa Sena R, Ahmed S, Tan WC, Li PZ, Labonté L, Aaron SD, et al.; CanCOLD Collaborative Research Group and The Canadian Respiratory Research Network. Work productivity loss in mild to moderate COPD: lessons learned from the CanCOLD study. *Eur Respir J* 2017;50:1701154.
- Miravittles M, Soriano JB, García-Río F, Muñoz L, Duran-Tauleria E, Sanchez G, et al. Prevalence of COPD in Spain: impact of undiagnosed COPD on quality of life and daily life activities. *Thorax* 2009;64:863–868.
- Labonté LE, Tan WC, Li PZ, Mancino P, Aaron SD, Benedetti A, et al.; Canadian Respiratory Research Network; CanCOLD Collaborative Research Group. Undiagnosed chronic obstructive pulmonary disease contributes to the burden of health care use. Data from the CanCOLD study. *Am J Respir Crit Care Med* 2016;194:285–298.
- Khakban A, Sin DD, FitzGerald JM, Ng R, Zafari Z, McManus B, et al. Ten-year trends in direct costs of COPD: a population-based study. *Chest* 2015;148:640–646.
- Troosters T, Sciurba F, Battaglia S, Langer D, Valluri SR, Martino L, et al. Physical inactivity in patients with COPD, a controlled multi-center pilot-study. *Respir Med* 2010;104:1005–1011.
- Ekström M. Why treatment efficacy on breathlessness in laboratory but not daily life trials? The importance of standardized exertion. *Curr Opin Support Palliat Care* 2019;13:179–183.
- Ahmadi Z, Sandberg J, Shannon-Honson A, Vandersman Z, Currow DC, Ekström M. Is chronic breathlessness less recognised and treated compared with chronic pain? A case-based randomised controlled trial. *Eur Respir J* 2018;52:1800887.
- Halding AG, Heggdal K, Wahl A. Experiences of self-blame and stigmatisation for self-infliction among individuals living with COPD. *Scand J Caring Sci* 2011;25:100–107.
- van Schayck CP, van Der Heijden FM, van Den Boom G, Tirimanna PR, van Herwaarden CL. Underdiagnosis of asthma: is the doctor or the patient to blame? The DIMCA project. *Thorax* 2000;55:562–565.
- Cherian M, Magner KMA, Whitmore GA, Vandemheen KL, FitzGerald JM, Bergeron C, et al. Patient and physician factors associated with symptomatic undiagnosed asthma or COPD. *Eur Respir J* 2023;61:2201721.
- Han MK, Martinez CH, Au DH, Bourbeau J, Boyd CM, Branson R, et al. Meeting the challenge of COPD care delivery in the USA: a multiprovider perspective. *Lancet Respir Med* 2016;4:473–526.
- Gershon AS, Hwee J, Croxford R, Aaron SD, To T. Patient and physician factors associated with pulmonary function testing for COPD: a population study. *Chest* 2014;145:272–281.
- Summers RH, Sharmeen T, Lippiett K, Gillett K, Astles C, Vu L, et al. A qualitative study of GP, nurse and practice manager views on using targeted case-finding to identify patients with COPD in primary care. *NPJ Prim Care Respir Med* 2017;27:49.
- Arne M, Lisspers K, Ställberg B, Boman G, Hedenström H, Janson C, et al. How often is diagnosis of COPD confirmed with spirometry? *Respir Med* 2010;104:550–556.
- Vázquez-García JC, Salas-Hernández J, Pérez Padilla R, Montes de Oca M. Respiratory health in Latin America: number of specialists and human resources training. *Arch Bronconeumol* 2014;50:34–39.
- Tálamo C, de Oca MM, Halbert R, Perez-Padilla R, Jardim JR, Muiño A, et al.; PLATINO team. Diagnostic labeling of COPD in five Latin American cities. *Chest* 2007;131:60–67.
- Fortis S, Eberlein M, Georgopoulos D, Comellas AP. Predictive value of prebronchodilator and postbronchodilator spirometry for COPD features and outcomes. *BMJ Open Respir Res* 2017;4:e000213.
- Global Initiative for Chronic Obstructive Lung Disease (GOLD). Global Strategy for Prevention, Diagnosis and Management of COPD: 2024 Report. Accessed January 23, 2024. Available from: <https://goldcopd.org/2024-gold-report/>.
- Tabyshova A, Hurst JR, Soriano JB, Checkley W, Wan-Chun Huang E, Trofor AC, et al. Gaps in COPD guidelines of low- and middle-income countries: a systematic scoping review. *Chest* 2021;159:575–584.
- Hurst JR, Buist AS, Gaga M, Gianella GE, Kirenga B, Khoo EM, et al. Challenges in the implementation of chronic obstructive pulmonary disease guidelines in low- and middle-income countries: an official American Thoracic Society workshop report. *Ann Am Thorac Soc* 2021;18:1269–1277.
- Casas Herrera A, Montes de Oca M, López Varela MV, Aguirre C, Schiavi E, Jardim JR; 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* 2016;11:e0152266.
- Florman KEH, Siddharthan T, Pollard SL, Alupo P, Barber JA, Chandyo RK, et al.; Additional GECO Study Investigators. Unmet diagnostic and therapeutic opportunities for chronic obstructive pulmonary disease in low- and middle-income countries. *Am J Respir Crit Care Med* 2023;208:442–450.
- Rennard SI, Drummond MB. Early chronic obstructive pulmonary disease: definition, assessment, and prevention. *Lancet* 2015;385:1778–1788.
- Soriano JB, Polverino F, Cosio BG. What is early COPD and why is it important? *Eur Respir J* 2018;52:1801448.
- Webber EM, Lin JS, Thomas RG. Screening for chronic obstructive pulmonary disease: updated evidence report and systematic review for the US Preventive Services Task Force. *JAMA* 2022;327:1812–1816.
- Ranson JM, Kužma E, Hamilton W, Lang I, Llewellyn DJ. Case-finding in clinical practice: an appropriate strategy for dementia identification? *Alzheimers Dement (N Y)* 2018;4:288–296.
- Jordan RE, Adab P, Sitch A, Enocson A, Blissett D, Jowett S, et al. Targeted case finding for chronic obstructive pulmonary disease versus routine practice in primary care (TargetCOPD): a cluster-randomised controlled trial. *Lancet Respir Med* 2016;4:720–730.
- Mets OM, Buckens CF, Zanen P, Isgum I, van Ginneken B, Prokop M, et al. Identification of chronic obstructive pulmonary disease in lung cancer screening computed tomographic scans. *JAMA* 2011;306:1775–1781.
- Ruparel M, Quaife SL, Dickson JL, Horst C, Tisi S, Hall H, et al. Prevalence, symptom burden, and underdiagnosis of chronic obstructive pulmonary disease in a lung cancer screening cohort. *Ann Am Thorac Soc* 2020;17:869–878.
- Ronaldson SJ, Dyson L, Clark L, Hewitt CE, Torgerson DJ, Cooper BG, et al.; DOC study team. Determining the optimal approach to identifying individuals with chronic obstructive pulmonary disease: the DOC study. *J Eval Clin Pract* 2018;24:487–495.
- Spyratos D, Haidich AB, Chloros D, Michalopoulou D, Sichelidis L. Comparison of three screening questionnaires for chronic obstructive pulmonary disease in the primary care. *Respiration* 2017;93:83–89.
- Sogbetun F, Eschenbacher WL, Welge JA, Panos RJ. A comparison of five surveys that identify individuals at risk for airflow obstruction and chronic obstructive pulmonary disease. *Respir Med* 2016;120:1–9.

40. Miravittles M, Llor C, Calvo E, Diaz S, Díaz-Cuervo H, Gonzalez-Rojas N. Validation of the Spanish version of the Chronic Obstructive Pulmonary Disease-Population Screener (COPD-PS). Its usefulness and that of FEV₁/FEV₆ for the diagnosis of COPD [in Spanish]. *Med Clin (Barc)* 2012;139:522–530.
41. Lordés M, Zurdo E, Jaén Á, Vázquez I, Pastrana L, Miravittles M. Which is the best screening strategy for COPD among smokers in primary care? *COPD* 2017;14:43–51.
42. Au-Doung PLW, Wong CKM, Chan DCC, Chung JWH, Wong SYS, Leung MKW. PUMA screening tool to detect COPD in high-risk patients in Chinese primary care—a validation study. *PLoS One* 2022;17:e0274106.
43. Zhou J, Li X, Wang X, Yu N, Wang W. Accuracy of portable spirometers in the diagnosis of chronic obstructive pulmonary disease: a meta-analysis. *NPJ Prim Care Respir Med* 2022;32:15.
44. Fujita M, Nagashima K, Takahashi S, Suzuki K, Fujisawa T, Hata A. Handheld flow meter improves COPD detectability regardless of using a conventional questionnaire: a split-sample validation study. *Respirology* 2020;25:191–197.
45. Stanley AJ, Hasan I, Crockett AJ, van Schayck OC, Zwar NA. COPD Diagnostic Questionnaire (CDQ) for selecting at-risk patients for spirometry: a cross-sectional study in Australian general practice. *NPJ Prim Care Respir Med* 2014;24:14024.
46. Frith P, Crockett A, Beilby J, Marshall D, Attewell R, Ratnanesan A, et al. Simplified COPD screening: validation of the PiKo-6 in primary care. *Prim Care Respir J* 2011;20:190–198.
47. Kotz D, Nelemans P, van Schayck CP, Wesseling GJ. External validation of a COPD diagnostic questionnaire. *Eur Respir J* 2008;31:298–303.
48. Price DB, Tinkelman DG, Nordyke RJ, Isonaka S, Halbert RJ; COPD Questionnaire Study Group. Scoring system and clinical application of COPD diagnostic questionnaires. *Chest* 2006;129:1531–1539.
49. Lopez Varela MV, Montes de Oca M, Wehrmeister FC, Rodriguez C, Ramirez L, Menezes A. External validation of the PUMA COPD diagnostic questionnaire in a general practice sample and the PLATINO study population. *Int J Chron Obstruct Pulmon Dis* 2019;14:1901–1911.
50. Quezada WA, Whippo BA, Jellen PA, Leidy NK, Mannino DM, Kim KJ, et al.; High-Risk-COPD Screening Study Group. How well does CAPTURE translate? An exploratory analysis of a COPD case-finding method for Spanish-speaking patients. *Chest* 2017;152:761–770.
51. López Varela MV, Montes de Oca M, Rey A, Casas A, Stirbulov R, Di Boscio V; PUMA Team. Development of a simple screening tool for opportunistic COPD case finding in primary care in Latin America: the PUMA study. *Respirology* 2016;21:1227–1234.
52. Martinez FJ, Mannino D, Leidy NK, Malley KG, Bacci ED, Barr RG, et al.; High-Risk-COPD Screening Study Group. A new approach for identifying patients with undiagnosed chronic obstructive pulmonary disease. *Am J Respir Crit Care Med* 2017;195:748–756.
53. Huynh C, Whitmore GA, Vandemheen KL, FitzGerald JM, Bergeron C, Boulet LP, et al. Derivation and validation of the UCAP-Q case-finding questionnaire to detect undiagnosed asthma and COPD. *Eur Respir J* 2022;60:2103243.
54. Schnieders E, Únal E, Winkler V, Dambach P, Louis VR, Horstick O, et al. Performance of alternative COPD case-finding tools: a systematic review and meta-analysis. *Eur Respir Rev* 2021;30:200350.
55. Zhou J, Yu N, Li X, Wang W. Accuracy of six chronic obstructive pulmonary disease screening questionnaires in the Chinese population. *Int J Chron Obstruct Pulmon Dis* 2022;17:317–327.
56. Soriano JB, Molina J, Miravittles M. Combining case-finding methods for COPD in primary care: a large, two-stage design study. *Int J Tuberc Lung Dis* 2018;22:106–111.
57. Balata H, Harvey J, Barber PV, Colligan D, Duerden R, Elton P, et al. Spirometry performed as part of the Manchester community-based lung cancer screening programme detects a high prevalence of airflow obstruction in individuals without a prior diagnosis of COPD. *Thorax* 2020;75:655–660.
58. Undrunas A, Kasprzyk P, Rajca A, Kuziemski K, Rzyman W, Zdrojewski T. Prevalence, symptom burden and under-diagnosis of chronic obstructive pulmonary disease in Polish lung cancer screening population: a cohort observational study. *BMJ Open* 2022;12:e055007.
59. Tang LYW, Coxson HO, Lam S, Leipsic J, Tam RC, Sin DD. Towards large-scale case-finding: training and validation of residual networks for detection of chronic obstructive pulmonary disease using low-dose CT. *Lancet Digit Health* 2020;2:e259–e267.
60. Fabbri LM, Celli BR, Agusti A, Criner GJ, Dransfield MT, Divo M, et al. COPD and multimorbidity: recognising and addressing a syndemic occurrence. *Lancet Respir Med* 2023;11:1020–1034.
61. Natali D, Cloatte G, Hovette P, Cochrane B. Screening for comorbidities in COPD. *Breathe (Sheff)* 2020;16:190315.
62. Lambe T, Adab P, Jordan RE, Sitch A, Enocson A, Jolly K, et al. Model-based evaluation of the long-term cost-effectiveness of systematic case-finding for COPD in primary care. *Thorax* 2019;74:730–739.
63. Qu S, You X, Liu T, Wang L, Yin Z, Liu Y, et al. Cost-effectiveness analysis of COPD screening programs in primary care for high-risk patients in China. *NPJ Prim Care Respir Med* 2021;31:28.
64. Johnson KM, Sadatsafavi M, Adibi A, Lynd L, Harrison M, Tavakoli H, et al. Cost effectiveness of case detection strategies for the early detection of COPD. *Appl Health Econ Health Policy* 2021;19:203–215.
65. Kostikas K, Price D, Gutzwiller FS, Jones B, Loeffroth E, Clemens A, et al. Clinical impact and healthcare resource utilization associated with early versus late COPD diagnosis in patients from UK CPRD database. *Int J Chron Obstruct Pulmon Dis* 2020;15:1729–1738.
66. Larsson K, Janson C, Ställberg B, Lisspers K, Olsson P, Kostikas K, et al. Impact of COPD diagnosis timing on clinical and economic outcomes: the ARCTIC observational cohort study. *Int J Chron Obstruct Pulmon Dis* 2019;14:995–1008.
67. Westerdahl E, Engman KO, Arne M, Larsson M. Spirometry to increase smoking cessation rate: a systematic review. *Tob Induc Dis* 2019;17:31.
68. Zwar NA, Bunker JM, Reddel HK, Dennis SM, Middleton S, van Schayck OC, et al. Early intervention for chronic obstructive pulmonary disease by practice nurse and GP teams: a cluster randomized trial. *Fam Pract* 2016;33:663–670.
69. Pollard SL, Siddharthan T, Hossen S, Rykiel NA, Flores-Flores O, Alupo P, et al.; GECO2 Trial Investigators. Chronic obstructive pulmonary disease self-management in three low- and middle-income countries: a pilot randomized trial. *Am J Respir Crit Care Med* 2023;208:1052–1062.
70. Vestbo J, Anderson JA, Brook RD, Calverley PM, Celli BR, Crim C, et al.; SUMMIT Investigators. Fluticasone furoate and vilanterol and survival in chronic obstructive pulmonary disease with heightened cardiovascular risk (SUMMIT): a double-blind randomised controlled trial. *Lancet* 2016;387:1817–1826.
71. Decramer M, Celli B, Kesten S, Lystig T, Mehra S, Tashkin DP; UPLIFT investigators. Effect of tiotropium on outcomes in patients with moderate chronic obstructive pulmonary disease (UPLIFT): a prespecified subgroup analysis of a randomised controlled trial. *Lancet* 2009;374:1171–1178.
72. Yawn BP, Han M, Make BM, Mannino D, Brown RW, Meldrum C, et al. Protocol summary of the COPD Assessment in Primary Care to Identify Undiagnosed Respiratory Disease and Exacerbation Risk (CAPTURE) validation in primary care study. *Chronic Obstr Pulm Dis (Miami)* 2021;8:60–75.
73. Labor M, Vrbica Ž, Gudelj I, Labor S, Plavec D. Diagnostic accuracy of a pocket screening spirometer in diagnosing chronic obstructive pulmonary disease in general practice: a cross sectional validation study using tertiary care as a reference. *BMC Fam Pract* 2016;1:112.
74. Kim JK, Lee CM, Park JY, Kim JH, Park SH, Jang SH, et al.; Active case finding strategy for chronic obstructive pulmonary disease with handheld spirometry. *Medicine (Baltimore)* 2016;95:e5683.
75. Represas-Represas C, Fernández-Villar A, Ruano-Raviña A, Priegue-Carrera A, Botana-Rial M; study group of “Validity of COPD-6 in non-specialized healthcare settings”. Screening for chronic obstructive pulmonary disease: validity and reliability of a portable device in non-specialized healthcare settings. *PLoS One* 2016;11:e0145571.
76. Thorn J, Tilling B, Lisspers K, Jörgensen L, Stenling A, Stratelis G. Improved prediction of COPD in at-risk patients using lung function

- pre-screening in primary care: a real-life study and cost-effectiveness analysis. *Prim Care Respir J* 2012;21:159–166.
77. van den Bemt L, Wouters BC, Grootens J, Denis J, Poels PJ, Schermer TR. Diagnostic accuracy of pre-bronchodilator FEV1/FEV6 from microspirometry to detect airflow obstruction in primary care: a randomised cross-sectional study. *NPJ Prim Care Respir Med* 2014;24:14033.
 78. Hidalgo Sierra V, Hernandez Mezquita MA, Palomo Cobos L, Garcia Sanchez M, Castellanos RD, Jodra Sanchez S, *et al.*; Usefulness of the Piko-6 portable device for early COPD detection in primary care. *Arch Bronconeumol (Engl Ed)* 2018;54:460–466.
 79. Lin CH, Cheng SL, Wang HC, Hsu WH, Lee KY, Perng DW, *et al.*; Novel app-based portable spirometer for the early detection of COPD. *Diagnostics (Basel)* 2021;11:785.
 80. Chen CZ, Sheu CC, Cheng SL, Wang HC, Lin MC, Hsu WH, *et al.*; Performance and clinical utility of various chronic obstructive pulmonary disease case-finding tools. *Int J Chron Obstruct Pulmon Dis* 2021;16:3405–3415.
 81. Martinez FJ, Han MK, Lopez C, Murray S, Mannino D, Anderson S, *et al.*; Discriminative accuracy of the CAPTURE tool for identifying chronic obstructive pulmonary disease in US primary care settings. *JAMA* 2023;329:490–501.