Asthma and chronic obstructive pulmonary disease (COPD) are 2 of the 3 most likely diagnoses for obstructive airway disorders in a patient with symptoms of shortness of breath. In Canada, obstructive airway disorders are common and are associated with substantial morbidity and mortality. In Alberta, exacerbations of COPD accounted for 85,330 emergency department visits during a 6-year period. In Ontario, researchers projected that 1 in 8 individuals will have asthma by the year 2022, suggesting that obstructive airway disorders will continue to be a considerable burden on the Canadian health care system. Guidelines on obstructive airway disorders, including those from the Global Initiative for Chronic Obstructive Lung Disease and the Global Initiative for Asthma, sponsored by international associations such as the American Thoracic Society, the European Respiratory Society, the Canadian Thoracic Society, and the National Heart, Lung and Blood Institute, have recommended use of pulmonary function testing for the diagnostic workup of patients with suspected obstructive airway disorders. Despite this recommendation, evidence exists that pulmonary function tests are underused, especially in primary care settings.

Authors of recent studies have expressed concern about the accuracy of diagnosis of obstructive airway disorders, including
under- and overdiagnosis of both asthma and COPD.\textsuperscript{8–13} Moreover, it has been reported that among patients receiving treatment in the emergency department for heart failure, 38% had received improper treatment with inhaled β-agonists.\textsuperscript{14} These researchers used a starting point of a “previous physician diagnosis” of obstructive airway disorders in their determination of diagnostic accuracy. Some patients, however, are not specifically informed of their diagnosis (i.e., are prescribed a “trial” of a β-agonist or other inhaled therapy), and such patients would be excluded from these studies.

The implications of poor diagnostic accuracy include patients with severe asthma not receiving optimal management for their condition, patients with COPD misclassified as having asthma and vice versa, increased risk of hospital admission, and subsequent exacerbations. Additionally, patients may be misclassified as having an obstructive airway disorder when they have other causes of dyspnea, such as heart failure or pulmonary hypertension.

Taken together, these studies give rise to concerns about the appropriateness of providing treatment with inhaled lung medications, often before a definitive diagnosis of an obstructive airway disorder. As such, we examined previous diagnosis of obstructive airway disorders and other conditions in patients who received treatment with inhaled medications for shortness of breath in a community setting. We chose to recruit participants through referral from community pharmacies to evaluate a different population from that previously studied.

Methods

Study design, setting and participants
We used a cross-sectional design, recruiting consecutive patients aged 18 years and older from community pharmacies in Edmonton and Saskatoon, Canada, between February 2009 and February 2012. There were no inclusion or exclusion criteria for the selection of pharmacists (other than their interest in participating). Some were drawn from a network of pharmacists in Edmonton and Saskatoon that had previously participated in other health services research studies.

Pharmacists considered eligible patients who had a current prescription (refilled within the past 6 months) for an inhaled medication for shortness of breath symptoms. This included β-agonists (long and short acting), anticholinergics (long and short acting), corticosteroids and combination agents. Patients were identified by either a review of their medication lists generated by each recruiting pharmacy or when patients presented for refills of their inhaled medications. Patients were excluded if they were prescribed inhaled medications for symptoms other than shortness of breath (e.g., for a cough only). Patients were also excluded if they could not communicate in English (unless someone could facilitate translation and interpretation), were pregnant, or were unable to attend the appointment for pulmonary function testing and physical examination.

Study procedures
Patients meeting the study inclusion criteria were approached by the pharmacist (in person or over the phone and using standardized scripts) to obtain verbal consent for the project office to make contact by phone. During a telephone call, trained research personnel from the Epidemiology Coordinating and Research (EPICORE) Centre based at the University of Alberta (www.epicore.ualberta.ca) informed patients about the study procedures and asked for their verbal consent to participate.

Consenting patients were invited to undergo pulmonary function tests at either the University of Alberta or the University of Saskatchewan laboratories. Patients were contacted by telephone, to a maximum of 3 times, to schedule their testing. At the testing session, a research coordinator serving as the lead researcher trainee (or designated laboratory technician) obtained written consent and collected standardized patient information on sociodemographic characteristics, clinical history and appropriate validated disease-specific measures through self-report of the COPD Assessment Test (8-item questionnaire that provides a 0–40 score from less to more severe impact of COPD on a patient’s life), the Asthma Control Questionnaire (7-point scale that provides a 0–6 score from no impairment to maximum impairment for symptoms and rescue use in patients with asthma), functional capacity using the Medical Research Council Dyspnea Scale (5-item scale that provides a 1–5 score from none to almost complete incapacity to breathe in patients with cardiorespiratory conditions) and the New York Heart Association Functional Classification scale (4-item scale that provides a 1–4 score from no to severe limitation in physical activity in patients with heart failure).\textsuperscript{15–18} Patients were also asked to report on their knowledge of their diagnosis of asthma and COPD from their primary care provider (family physician or specialist). Case report forms are shown in Appendix 1 (available at www.cmajopen.ca/content/8/3/E605/suppl/DC1), and the COPD Assessment Test is available at www.catestonline.org.

We performed standard pulmonary function tests, as per recommendations from the European Respiratory Society\textsuperscript{19} and the American Thoracic Society.\textsuperscript{20} This included pre–post bronchodilator spirometry testing. Methacholine challenge testing,\textsuperscript{21} using the tidal breathing method, was performed in all patients who did not show evidence of obstructive airway disorders in the initial pre- and postbronchodilator spirometry. We measured lung volumes, and we excluded pulmonary restriction based on normal total lung capacity.

A blood sample for brain natriuretic peptide measurement was collected in all patients to rule out heart failure or other heart conditions as underlying entities of the shortness of breath. Complete blood work was not examined. The Canadian Cardiovacular Society has recommended a brain natriuretic peptide measurement less than 100 pg/mL as the threshold to differentiate obstructive airway disorders as a possible cause of shortness of breath.\textsuperscript{22} For pulmonary hypertension, we measured diffusing capacity, which, in combination with elevated brain natriuretic peptide, led to a presumptive diagnosis of pulmonary hypertension.

Finally, all patients gave us permission to review their medical records (e.g., to determine the specific condition for
which inhalers were prescribed) and the results of previous diagnostic tests, including pulmonary function tests, chest radiographs, echocardiograms and methacholine challenge testing for the determination of alternative diagnoses. Trained research assistants recorded this information from the provincial electronic health record in Alberta (www.albertanetcare.ca) and the patients’ primary care provider in Saskatchewan using standardized case report forms (Appendix 1).

### Study outcomes

The prevalence of asthma, COPD and nonobstructive airway disorders (heart failure and other conditions) was determined as definite or probable by our physician panel (B.H.R., I.M. and D.V.) (Table 1) through standardized criteria based on international guideline recommendations. A group of 3 expert physicians (1 emergency physician [B.H.R.] and 2 respirologists [L.M. and D.V.]) reviewed and adjudicated any case that remained unclear after the methacholine challenge test. Panelists were blinded to previous diagnoses and resolved any disagreements by consensus. The agreement between diagnoses derived from pulmonary function tests and diagnoses from primary care providers was evaluated.

### Sample size

The sample size for estimating a proportion with a 95% confidence interval of expected width $+/-d$ around the estimate can be calculated as $n = 1.96^2 \times P (1–P)/d^2$, where $P$ is the expected value of the proportion. With $d = 0.05$ and a prevalence of obstructive airway disorders of $P = 0.70$, $n = 323$.

### Statistical analysis

Descriptive analyses were used to report on patient characteristics and prevalence of obstructive airway disorders in shortness of breath. Analyses were performed using SPSS Version 20.0 (IBM). Percentage of baseline characteristics and patient final diagnostic outcomes are presented for categorical variables. For continuous variables, mean and standard deviation or median and interquartile range (IQR) are presented, as appropriate. The agreement between diagnoses derived from pulmonary function tests and diagnoses from primary care provider was evaluated using the $\kappa$ statistic. $\kappa$ statistics were interpreted as follows: poor ($< 0.20$), fair (0.20–0.40), moderate (0.41–0.60), good (0.61–0.80) and excellent (> 0.80).

### Ethics approval

Ethics approval for this study was received from the health research ethics boards at the University of Alberta and University of Saskatchewan.

### Results

The participant flow chart and overall diagnostic results are outlined in Figure 1. Of the 475 eligible patients who were initially screened and consented to participate in the study, 328 completed lung function testing and provided baseline information for diagnostic evaluation. The 147 patients who were lost to follow-up mainly did not present to the pulmonary function tests laboratory.

### Participant characteristics

Twenty-eight pharmacies volunteered to enrol patients using the study protocol. Major characteristics of the study population are summarized in Table 2. Most participants were women (188, 57.3%), white (283, 86.3%) and not single (190, 57.9%). The median age of participants was 50 (IQR 32–64) years, 161 (49.1%) had completed postsecondary education, and 182 (55.5%) were current or former smokers, with a median smoking pack-year of 19 (IQR 10–31) years. Nearly all (319, 97.3%) reported at least 1 comorbid condition with a median of 4 (IQR 3–6) comorbidities per patient. The most frequent reported comorbid conditions were allergies (255, 77.7%), gastroesophageal reflux disease (142, 43.3%) and depression (104, 31.7%). Patients reported an existing or prior history of asthma (184/216, 83.8%) and COPD (34/216, 15.7%).

The median score for the Asthma Control Questionnaire was 1.0 (IQR 0.4–1.7) among 157 of the 181 participants who self-reported being previously diagnosed with asthma; the median score for the COPD Assessment Test was 22 (IQR 17–29) among 160 of the 181 participants who self-reported being previously diagnosed with COPD.

### Table 1: Criteria for study diagnosis of obstructive airway disorders in patients with shortness of breath

| Disease         | Study diagnosis                      | Criteria*4,5 |
|-----------------|--------------------------------------|--------------|
| Asthma          | Definite asthma | Increase in FEV1 > 200 mL and > 12% above prebronchodilator FEV1 OR Positive response to methacholine provocation test: an airway hyperresponsiveness as defined by use of PC20 |
|                 | Probable asthma | Diagnosis by two-thirds consensus of the expert physician panel from review of examination, current respiratory symptoms and prior medical records |
| COPD            | Definite COPD | Postbronchodilator FEV1 < 80% predicted together with an FEV1/FVC < 0.70 |
|                 | Probable COPD | Diagnosis by two-thirds consensus of the expert physician panel from review of examination, current respiratory symptoms and prior medical records |
| Non-OADs        | Probable normal or with other entities | Does not fulfill criteria for definite or probable asthma or COPD diagnoses* |

Note: COPD = chronic obstructive pulmonary disease, FEV1 = forced expiratory volume in 1 second, FVC = forced vital capacity, OAD = obstructive airway disorder, PC20 = the provocative concentration of methacholine that results in a 20% drop in FEV1.

*In patients with a brain natriuretic peptide measurement > 100 pg/mL, a cardiologist consultation was obtained to evaluate the patient for heart failure.
15–30) among 28 of the 34 who self-reported being previously diagnosed with chronic obstructive airway disease. A total of 79 (24.1%) of patients reported moderate (3) to severe (5) scores on the Medical Research Council Dyspnea Scale, and 86 (26.2%) reported New York Heart Association Functional Classification Scale III or IV symptoms. Current symptoms most often reported were day- or nighttime cough (29.9%), fatigue (21.3%), chest tightness (19.5%) and wheezing (19.5%), and sputum production (17.1%).

A total of 216 patients (65.8%) reported ever receiving a diagnosis for their shortness of breath symptoms by their primary care provider, of which 83.8% were diagnosed with asthma and 15.7% with COPD. A total of 39.0% of patients reported seeing a specialist for their shortness of breath symptoms, and 40.8% had previously had pulmonary function tests conducted (Table 2). Most of the patients previously diagnosed with asthma and COPD reported using quick-reliever medication (short-acting β-agonists) and inhaled combination agents (inhaled corticosteroids–long-acting β-agonists) in the last 6 months (Table 3).

Diagnoses

Table 4 shows the participant diagnoses derived from pulmonary function testing, using the criteria outlined in Table 1 and adjudicated by the expert physician panel. Information on previous diagnostic tests (e.g., chest radiographs, echocardiograms, methacholine challenge testing and pulmonary function tests) was available for 275 patients (83.8% of the study population).

Asthma was confirmed in 149 patients (45.4%) and COPD in 97 patients (29.6%). Of the patients who had a prior diagnosis of asthma or COPD from their primary care provider, 54.9% of diagnoses were confirmed for asthma and 71.1% for COPD by adjudicated diagnosis derived from pulmonary function tests (the positive predictive values and negative predictive values are shown in Table 5). Some patients (93; 28.4% of the study sample) had no evidence of obstructive airway diseases either by spirometry data or with further methacholine challenge testing.

Of those patients diagnosed with asthma or COPD by our expert physician panel, 11 had both conditions (asthma–COPD overlap syndrome). Confirmatory diagnosis was elusive in 62 patients (18.9% of the study sample). An additional 8 of the 62 participants could have been classified as having Global Initiative for Chronic Obstructive Lung Disease stage 1 COPD, but we elected to include them in the group of participants with indeterminate cause for shortness of breath. Most of these participants (5/8) had forced expiratory volume in 1 second greater than 95% predicted normal, and, after review, we believed that they did not have clinically significant airflow obstruction.

In Table 5, the κ statistics indicate there was only a poor agreement between previous diagnoses by primary care providers and by our expert physician panel (κ = 0.22 for asthma; κ = 0.28 for COPD). Heart failure and restrictive lung disease were the 2 most common other diagnoses determined for a patient without obstructive airway disorders. Diagnoses could not be determined by our expert

Figure 1: Flow of patients through the study. *By patient report and medical record review. Note: COPD = chronic obstructive pulmonary disease, OAD = obstructive airway disorder, PFT = pulmonary function test, SOB = shortness of breath.
physician panel for 62 patients (18.9% of the study sample) after pulmonary function testing and evaluation of information available in patients’ medical records.

Of the 149 patients diagnosed with asthma by our expert physician panel, 34.9% reported ever having a pulmonary function test performed, whereas for the 97 patients diagnosed with COPD, 50.5% reported ever having a pulmonary function test performed (Figure 1).

Interpretation

In our study of community-dwelling patients who received treatment with inhalers for shortness of breath symptoms and referred by pharmacists, a considerable proportion did not meet the diagnostic criteria for asthma or COPD when evidence-based recommendations were applied. Despite efforts made to obtain detailed information from chart reviews, patient interviews, medication record reviews, standardized assessment tools and additional testing (e.g., pulmonary function tests and methacholine challenge tests), a confirmatory diagnosis was elusive in an important proportion of patients (19%).

Our results are comparable to previous reports that used different means for participant recruitment. A multicentre study conducted in Italy (involving 24 pulmonary or geriatric institutions) investigated to what extent COPD was misdiagnosed in older patients with asthma. Among the 128 patients with asthma included in this study, COPD had been improperly diagnosed in 19.5%, whereas 27.3% of these patients had not been diagnosed previously. Older age and greater degree of disability were factors strongly associated with misdiagnosis. A survey conducted in North America evaluated the sex bias associated with diagnosis of COPD. A random sample of 192 primary care providers (96 American and 96 Canadian) underwent a structured interview after being presented a hypothetical clinical case (with 6 versions differing only in the sex and age of the patients). In this study, following an initial diagnosis without spirometry, the likelihood of diagnosis of COPD increased after conducting spirometry and after receiving a corticosteroid trial for both men (58% v. 74% v. 85%) and women (42% v. 66% v. 79%). The study showed that only 22% of physicians would have requested spirometry at presentation by patients with symptoms consistent with COPD.

Another study conducted in Scotland and the United States compared prior diagnoses of asthma and COPD with study diagnosis based on spirometric results. This study found that among participants with spirometry-based diagnosis of COPD, 52% reported prior diagnosis of asthma without concurrent chronic bronchitis or emphysema diagnosis, 38% reported prior diagnosis of chronic bronchitis or emphysema, and 11% reported no prior diagnosis of COPD.

Similarly, a study carried out in 2 provinces in Canada evaluated primary care practice in patients with COPD compared with recommended care. Among the 1090 patients recruited by the participating physicians, spirometric confirmation of

### Table 2: Characteristics of patients with shortness of breath referred by pharmacists

| Characteristic                                      | No. (%) of patients* |
|----------------------------------------------------|----------------------|
| **Characteristics**                                 | **n = 328**          |
| Age, yr, median (IQR)                              | 50 (32–64)           |
| Weight, kg, mean ± SD                              | 83.3 ± 20.7          |
| Height, cm, mean ± SD                              | 166.4 ± 9.6          |
| Sex, female                                        | 188 (57.3)           |
| Ethnicity                                          |                      |
| White                                              | 283 (86.3)           |
| South Asian                                        | 19 (5.8)             |
| Aboriginal                                         | 12 (3.6)             |
| Other†                                             | 14 (4.3)             |
| Highest level of education                         |                      |
| Completed postsecondary education                  | 161 (49.1)           |
| Some postsecondary education                       | 62 (18.9)            |
| High school diploma                                | 61 (18.6)            |
| Less than grade 12                                 | 44 (13.4)            |
| Married or common-law status                       | 190 (57.9)           |
| **Most common comorbidities**                      |                      |
| Allergies                                          | 255 (77.7)           |
| GERD                                               | 142 (43.3)           |
| Depression                                         | 104 (31.7)           |
| Sinus or nasal polyps                              | 103 (31.4)           |
| Anxiety                                            | 88 (26.8)            |
| Other relevant comorbidities                       |                      |
| Hypertension                                       | 85 (25.9)            |
| Anemia                                             | 42 (12.8)            |
| Coronary artery disease                            | 28 (8.5)             |
| Heart failure                                      | 6 (1.8)              |
| **Smoking history**                                |                      |
| Current                                            | 65 (19.8)            |
| Former                                             | 117 (35.7)           |
| Never                                              | 146 (44.5)           |
| Pack-years for current or former smokers, median (IQR) | 19 (10–31)          |
| **Symptoms**                                       |                      |
| Cough symptoms (day or night)                      | 98 (29.9)            |
| Sputum production                                  | 56 (17.1)            |
| Chest tightness                                    | 64 (19.5)            |
| Wheeze                                             | 64 (19.5)            |
| Other symptoms                                     |                      |
| Peripheral edema                                   | 31 (9.5)             |
| Fatigue                                            | 70 (21.3)            |
| Bilateral ankle edema                              | 34 (10.4)            |
| Fever or flu-like symptoms                         | 18 (5.5)             |
| Currently absent from work or school                | 23 (7.0)             |
| Previously diagnosed by a primary care provider for SOB symptoms | 216 (65.8) |
| Asthma                                             | 181/216 (83.8)       |
| COPD                                               | 34/216 (15.7)        |
| Other                                              | 113/216 (52.3)       |
| Previously seen by a specialist for SOB            | 128 (39.0)           |
| Previously had PFTs conducted for SOB              | 134 (40.8)           |

*Note: COPD = chronic obstructive pulmonary disease, GERD = gastroesophageal reflux disease, IQR = interquartile range, PFT = pulmonary function test, SD = standard deviation, SOB = shortness of breath.
†Included black, Hispanic, East and Southeast Asian, Middle Eastern and other ethnicities not stated in the case report form.
diagnosis was reported by only 56% of them. In addition, pharmacological treatment matching guideline recommendations was identified in only a third of the study population.

A more recent study conducted in Italy assessed the level of asthma misdiagnosis among patients reporting respiratory symptoms to their general practitioner and undergoing treatment with inhaled corticosteroids. From a total of 2090 patients registered in general practitioners’ databases as receiving at least 3 prescriptions of inhaled or nebulized corticosteroids during the 12 months preceding the start of the study, only 47% were diagnosed with asthma.

Finally, a study conducted in 10 Canadian cities used random digital dialing to recruit adults who reported a history of previous physician-diagnosed asthma within the past 5 years to determine whether a current diagnosis of asthma could be ruled out and asthma medications safely stopped. Asthma was ruled out in 33% of 613 participants who underwent home peak flow and symptom monitoring, spirometry and serial bronchial challenge tests. Those participants in whom asthma was ruled out were less likely to have undergone testing for airflow limitation in the community at the time of diagnosis when compared with those in whom asthma diagnosis was confirmed (43.8% v. 55.6%, respectively).

The findings of these studies and the current study suggest that misdiagnosis in community-dwelling patients led to exposure to unnecessary and expensive treatments and, likely, to undesired health outcomes and delayed diagnosis. Importantly, misdiagnosis could also compromise the validity of research studies based on pharmacy dispensations (e.g., pharmacoepidemiology studies that often use 2 dispensations of β-agonists as a proxy for a diagnosis of asthma). In addition, the poor agreement between previous diagnoses by primary care providers and by our expert physician panel could have an important clinical impact on patients’ outcomes when they are exposed to chronic and inappropriate therapies. Educational efforts should be directed to improve the diagnostic approaches of primary care providers and other health practitioners. Moreover, the results suggest that other health professionals, such as pharmacists, should be empowered to direct patients toward optimal diagnosis and management.

**Limitations**

Our sample of 475 represents eligible patients. Unfortunately, we were not able to document the total number of

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**Table 3: Medications used by patients previously diagnosed with asthma and chronic obstructive pulmonary disease, by diagnosis**

| Medication                          | Previous diagnosis of asthma n = 181† | Previous diagnosis of COPD n = 34† | No previous diagnosis of asthma or COPD n = 127 |
|-------------------------------------|--------------------------------------|-----------------------------------|-----------------------------------------------|
| Short-acting β-agonist              | 141 (77.9)                           | 22 (64.7)                         | 83 (65.4)                                     |
| Inhaled corticosteroid             | 45 (24.9)                            | 3 (8.8)                           | 38 (29.9)                                     |
| Inhaled corticosteroid–long-acting β-agonist | 120 (66.3)                         | 30 (88.2)                         | 54 (42.5)                                     |
| Oral corticosteroid                | 6 (3.3)                              | 2 (5.9)                           | 5 (3.9)                                       |
| Long-acting anticholinergic         | 7 (3.9)                              | 13 (38.2)                         | 7 (5.5)                                       |
| Short-acting anticholinergic        | 5 (2.8)                              | 2 (5.9)                           | 6 (4.7)                                       |
| Theophylline                        | 4 (2.2)                              | 1 (2.9)                           | 1 (0.8)                                       |
| Leukotriene antagonist             | 20 (11.0)                            | 2 (5.9)                           | 5 (3.9)                                       |

Note: COPD = chronic obstructive pulmonary disease. *Based on pharmacist report. †A total of 14 patients (4.3% of the study population) reported a previous diagnosis of both asthma and COPD.

**Table 4: Adjudicated diagnoses by expert physician panel for patients with shortness of breath referred by pharmacists**

| Diagnosis by expert physician panel | No. (%) of patients n = 328 |
|-------------------------------------|-----------------------------|
| Asthma*                             | 149 (45.4)                  |
| Definite diagnoses                  | 147/149 (98.7)              |
| Probable diagnoses                  | 2/149 (1.3)                 |
| COPD*                               | 97 (29.6)                   |
| Definite diagnoses                  | 94/97 (96.9)                |
| Probable diagnoses                  | 3/97 (3.1)                  |
| Other                               | 20 (6.1)                    |
| Heart failure                       | 9/20 (45.0)                 |
| Restrictive lung diseases or pulmonary hypertension | 8/20 (40.0) |
| Bronchitis                          | 2/20 (10.0)                 |
| Obesity                             | 1/20 (5.0)                  |
| Unknown origin (indeterminate cause for SOB) | 62 (18.9) |

Note: COPD = chronic obstructive pulmonary disease, SOB = shortness of breath. *A total of 11 patients had both asthma and COPD diagnoses.
patients from which these patients with shortness of breath were screened.

One of the main strengths of our study is the use of standardized diagnostic approaches following the recommendations from international guidelines by an outcome adjudication panel of experts. We collected robust new information, and previous diagnostic tests were available for all but 16% of the study population. Despite these details, there was a substantial proportion of patients (19%) for whom expert clinicians, using accepted standards and protocols, could not provide a diagnosis. This implies that a considerable number of patients are receiving medication for chronic respiratory conditions based on non-evidence-based medical practice; there is variability of airway responsiveness measurements in diseases like asthma; and the accuracy of our estimates could be affected by the completeness of the information available for assessment. Although we did not explore in detail potential sources for diagnostic opportunity bias, we acknowledge that variation in the quality of medical reporting could be an important source of nondifferential bias in our study.

Although the recruitment of community-dwelling patients through volunteering community pharmacists minimizes the selection bias introduced by physician diagnosis and random digit dialing used in previous studies, some cases may have reflected the pharmacist’s clinical concern for confirmation of diagnosis. Our study excluded patients who could not communicate in English, were pregnant or were unable to attend the study appointments. The participating pharmacists may not be representative, given that they volunteered to participate; however, that should not have biased the patient sample. Importantly, many patients with shortness of breath may be untreated; therefore, patients with mild obstructive airway disorders are underrepresented in our study.

Despite multiple efforts to facilitate participants’ attendance to the pulmonary function tests laboratory, 147 (about 30%) participants did not present to the laboratory and were considered lost to follow-up. There were no baseline characteristics for these patients to evaluate the difference with those who attended pulmonary function testing, and therefore the distribution of obstructive airway disorders in these participants could have been different.

### Conclusion

We found that less than half of community-dwelling patients receiving treatment with inhaled medications for shortness of breath and presumed obstructive airway disorders had confirmed asthma, about a quarter had COPD and a quarter had no demonstrable obstructive airway disorders. These findings, coupled with the fact that only about 40% of participants had ever had pulmonary function tests performed, highlights the need to avoid empiric treatment with β-agonists and inhaled corticosteroid agents, increase the use of objective measures of lung function for the diagnosis of obstructive airway disorders, and identify factors associated with patient misdiagnosis.

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### Table 5: Agreement between diagnoses of asthma and chronic obstructive pulmonary disease by primary care providers and those derived from pulmonary function testing

| Prior diagnosis of OAD by PCP | Asthma* | COPD* |
|-----------------------------|---------|-------|
|                             | Yes     | No    | Total | Yes | No | Total |
| Yes                         | 106     | 87    | 193   | 27  | 11 | 38    |
| No                          | 43      | 92    | 135   | 70  | 220| 290   |
| Total                       | 149     | 179   | 328   | 97  | 231| 328   |
| PPV, %                      | 54.9    |       |       | 71.1|    |       |
| NPV, %                      | 68.1    |       |       | 75.9|    |       |
| κ value                     | 0.22    |       |       | 0.28|    |       |
| p value                     | < 0.001 |       |       | < 0.001|  |       |

Note: COPD = chronic obstructive pulmonary disease, NPV = negative predictive value, OAD = obstructive airway disease, PCP = primary care provider, PFT = pulmonary function testing, PPV = positive predictive value.

*A total of 11 patients had both asthma and COPD diagnoses.*
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