Health care utilization history, GOLD guidelines, and respiratory medication prescriptions in patients with COPD

Background: The relationship between prior health care utilization and respiratory medication prescriptions in an unselected population of patients with COPD is not known.

Methods: We determined the prescribed respiratory medications and respiratory and nonrespiratory health care encounters in 523 Veterans with COPD at the Cincinnati Veterans Affairs Medical Center between 2000 and 2005. Prescribed treatments were compared with the GOLD guidelines and each patient was classified as receiving less medications than recommended in the guidelines (<G), medications according to the guidelines (=G), or more medications than recommended (>G).

Results: Respiratory medications were <G for 54%, =G in 33%, and >G for 14% of the patients studied. For GOLD stages 1 and 2, <G patients had the fewest and >G patients the most prior respiratory encounters during a 12 month period (0.31 ± 0.073 (0.21, 0.47), 0.75 ± 0.5 (0.37, 1.5), 1.1 ± 0.27 (0.74, 1.6) visits/person/year, <G, =G, >G, respectively, mean ± standard error of mean (SEM) (95% confidence limits) 2 degrees of freedom (df) ANOVA P < 0.001 for prescription effect). For GOLD stages 3 and 4, <G was associated with significantly fewer prior respiratory visits than was =G (0.78 ± 0.11 (0.6, 1.0) and 2.4 ± 0.47 (1.9, 3.1) visits/person/year, respectively, P < 0.001). There were no differences in nonrespiratory health care visits for GOLD stages 1 and 2 by prescription level (3.1 ± 0.24 (2.6, 3.5), 3.1 ± 0.46 (2.1, 4.6) and 4.1 ± 0.55 (3.3, 5.1) visits/person/year, <G, =G, >G respectively, 2 df ANOVA P = 0.096) or for GOLD stages 3 and 4 (3.6 ± 0.25 (3.2, 4.1) and 4.0 ± 0.44 (3.3, 4.9) visits/person/year, <G and =G, respectively, P = 0.36).

Conclusions: Respiratory medications prescribed for an unselected population with a broad range of COPD severity complied poorly with the GOLD pharmacologic treatment guidelines but correlated with the number of prior respiratory health care visits.

Keywords: health care, COPD, respiratory visits, GOLD guidelines, prescription

Introduction

Chronic obstructive pulmonary disease (COPD) is a common disease affecting 12–24 million adults in the United States.1,2 COPD is currently the fourth leading cause of mortality and is projected to be the third leading cause of death by 2020.1,2 Outpatient health care encounters, emergency department visits, and hospitalizations prompted by COPD cost more than 32 billion dollars yearly.1,2

Over 50 clinical practice guidelines have been developed for the management of COPD.3,4 The Global Strategy for the Diagnosis and Management of Chronic Obstructive Lung Disease (GOLD), a collaboration of the World Health Organization and the National Heart Lung and Blood Institute of the National Institute of Health, has created evidence-based guidelines with standards for grading evidence, explicit recommendations, and a
well organized implementation group with regular, systematic updates and revisions. Nevertheless, despite international dissemination and intensive promotion, the GOLD guidelines have not been universally adopted and implemented by primary care physicians or pulmonologists.

The reasons for poor adherence with COPD treatment guidelines are not well understood. Potential factors contributing to noncompliance include low utilization of spirometry, unawareness of guideline recommendations, and perceived ineffectiveness. To examine the relationship between prior health care utilization and the prescription of respiratory medications, we determined respiratory health care encounters and the prescription of respiratory medications compared with the GOLD guidelines in an unselected population with COPD.

**Patients and methods**

**Patients**

We reviewed, retrospectively, the medical records of all patients at the Cincinnati Veterans Administration Medical Center (VAMC) with a diagnosis of COPD between June 1, 2000 and June 1, 2005 (Figure 1). To ensure that all participants were followed actively, individuals who did not have at least one health care encounter in the 12 months prior to the study were excluded ($n=189$, Figure 1). Prescribed therapeutic regimens compared with GOLD treatment guidelines are presented in Figure 1. All spirometry was performed in the Cincinnati VAMC Pulmonary Function Laboratory according to American Thoracic Society (ATS) guidelines. For 131 patients, the forced expiratory volume in 1 second ($\text{FEV}_1$) divided by the forced vital capacity ($\text{FVC}$), $\text{FEV}_1/\text{FVC}$, was $\geq 0.70$ and they were classified as clinical COPD (former GOLD stage 0).

The remaining 392 patients were classified into 4 stages (modified GOLD 1, modified GOLD 2, modified GOLD 3, modified GOLD 4) defined by the GOLD guidelines based upon prebronchodilator spirometry.

**Study design**

Each patient’s prescribed medication regimen was compared with the GOLD treatment guidelines. Subjects were classified as less medications than guidelines ($\leq G$) if they were prescribed less than the recommended regimen, according to guidelines ($=G$) if their respiratory medication prescription met the guidelines, and more medications than guidelines ($\geq G$) if they were prescribed more respiratory medications than recommended. The Veterans Affairs medical system electronic medical record was reviewed and every outpatient visit, emergency department encounter, and hospitalization during the preceding 60 months was categorized as respiratory or nonrespiratory based upon the chief complaint and diagnostic coding for the encounter. Examples of respiratory chief complaints included breathlessness, cough, or wheezing.

Because the prescription record could be determined most accurately at the time of the chart review, the primary analysis studied health care encounters during the preceding 12 months. In a secondary analysis, we extended the time frame to the preceding 60 months.

This protocol was approved by the Research and Development Committee of the Cincinnati VAMC and the University of Cincinnati Institutional Review Board. The need for consent was waived.

**Statistics**

The analyses of health care visits were conducted on the counts of encounters over 12 months and 60 months. Based upon the inclusion criteria, all patients were assumed to have been followed actively throughout the 60 month period. Only health care encounters that occurred within the Cincinnati VAMC system were counted and any period with no entry in the electronic medical record was recorded as zero visits for that time span. All visit frequencies were normalized to yearly rates. For many analyses, modified GOLD groups 1 and 2 were pooled, as were groups 3 and 4, because the combined modified GOLD groups shared similar physiologic characteristics and treatment recommendations.

Patients with clinical COPD could be categorized as $G$ or $>G$. Patients in modified GOLD stage 1 and 2 could be classified as $<G$, $=G$, or $>G$, whereas modified GOLD 3 and 4 subjects were grouped as $<G$ or $=G$.

For all analyses in which the number of visits was the dependent variable, confidence limits and $P$-values were based upon Wald tests within Poisson models with the Poisson variance estimates adjusted by the ratio of the model deviance to the model degrees of freedom. The study alpha was $P=0.05$, two-tailed. Results are presented as mean ± SEM (upper and lower 95% confidence limits).

**Results**

**Study population**

1338 patient records were evaluated and 523 patients were studied (Figure 1). The clinical characteristics of the study population are presented in Tables 1 and 2.

**Health care visits**

Hospital, emergency department (ED), outpatient respiratory and nonrespiratory visits over the 12 and 60 month periods
are shown in Figure 2. ED and outpatient visits comprised similar proportions of total respiratory visits (35% and 48%) whereas there were 6-fold more outpatient visits (83%) than ED visits (12%–13%) for nonrespiratory causes. Hospitalizations represented 16%–17% of respiratory visits but only 4%–5% of nonrespiratory encounters. The number of hospital, ED, and outpatient respiratory visits increased with each modified GOLD stage, whereas the number of nonrespiratory visits/person/year, \( P = 0.02 \) (Table 3) This increased total health care utilization was caused by more ED visits and hospitalizations. In the composite group modified GOLD 1 and 2, the prescription level increased as the number of prior respiratory event related encounters rose (2 df ANOVA \( P < 0.0001 \), Table 3). There were no significant differences in total nonrespiratory visits (2 df ANOVA \( P = 0.096 \), Table 3).

In the composite group modified GOLD 3 and 4, those individuals receiving \( > G \) had three times as many total respiratory visits as those who received \( < G, P < 0.0001 \) (Table 3). This increase was due to elevated numbers of office visits as well as ED encounters and hospitalizations. There were no differences in hospital, office, or total nonrespiratory visits, \( P = 0.36 \). Similar relationships between prescribed treatment and respiratory and nonrespiratory encounters were present during the 60 month period (Table 3).

**Discussion**

Respiratory medications prescribed for patients in this unselected population with a broad range of air flow limitation and COPD severity complied poorly with the GOLD guidelines. In the clinical COPD group (former GOLD 0), patients receiving \( > G \) had twice as many prior respiratory visits as patients receiving \( = G \), \( 0.58 \pm 0.19 \text{ versus } 0.27 \pm 0.057 \text{ visits/person/year, } P = 0.02 \). (Table 3) This increased total health care utilization was caused by more ED visits and hospitalizations. In the composite group modified GOLD 1 and 2, the prescription level increased as the number of prior respiratory event related encounters rose (2 df ANOVA \( P < 0.0001 \), Table 3). There were no significant differences in total nonrespiratory visits (2 df ANOVA \( P = 0.096 \), Table 3).

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The COPD Resource Network Needs Assessment Survey found that although 54% of generalists and 94% of pulmonologists were aware of published COPD guidelines, they often under-prescribed recommended pharmacologic treatments. We found that 54% of patients were prescribed less medications than recommended. Although unawareness of COPD guidelines may be one factor contributing to the lack of adherence, independent prescribing habits and absence of apparent effect on patient outcomes may also influence physician utilization of practice guidelines. A history of fewer respiratory encounters correlated with less than recommended prescription of respiratory medications suggesting that providers and possibly patients perceived less need for these treatments. Conversely, our findings indicate that physicians prescribe more than recommended medications for patients who have more frequent respiratory health care encounters. Thus, prior health care visits for respiratory symptoms may influence caregivers to prescribe more respiratory treatments in an effort to alleviate symptoms and improve respiratory health regardless of a patient’s lung function measured by spirometry. Similarly, the Resource Use Study in COPD demonstrated that patients experiencing COPD exacerbations were prescribed significantly more medications than those who did not have respiratory symptoms.

In our unselected population, most health care encounters for COPD were confined to a small fraction of patients with COPD and this population may be disproportionately represented in some studies of respiratory medications. The results of these studies provide, in part, the evidence for spirometry-based COPD management guidelines and may, potentially, bias recommendations toward more treatment. Increased health care visits for respiratory symptoms may influence physicians to prescribe more respiratory treatments in an effort to alleviate symptoms and improve respiratory health regardless of a patient’s lung function measured by spirometry. Similarly, the Resource Use Study in COPD demonstrated that patients experiencing COPD exacerbations were prescribed significantly more medications than those who did not have respiratory symptoms.

Table 2 Distribution of patients, FEV₁, and FEV₁/FVC according to modified GOLD stage

<table>
<thead>
<tr>
<th>Modified GOLD stage</th>
<th>n (%)</th>
<th>FEV₁ (l)</th>
<th>FEV₁/FVC (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Clinical</td>
<td>131 (25%)</td>
<td>2.47 ± 0.71</td>
<td>77.2 ± 5.69</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(2.35 – 2.59)</td>
<td>(76.27 – 78.22)</td>
</tr>
<tr>
<td>COPD</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>28 (5%)</td>
<td>2.66 ± 0.61</td>
<td>65.1 ± 3.17</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(2.44 – 2.88)</td>
<td>(63.89 – 66.25)</td>
</tr>
<tr>
<td>2</td>
<td>134 (25.6%)</td>
<td>2.03 ± 0.50</td>
<td>58.8 ± 6.57</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(1.94 – 2.11)</td>
<td>(57.63 – 59.86)</td>
</tr>
<tr>
<td>3</td>
<td>113 (21.6%)</td>
<td>1.29 ± 0.32</td>
<td>47.9 ± 8.92</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(1.13 – 1.35)</td>
<td>(46.2 – 49.49)</td>
</tr>
<tr>
<td>4</td>
<td>117 (22.4%)</td>
<td>0.89 ± 0.28</td>
<td>42.3 ± 9.11</td>
</tr>
<tr>
<td></td>
<td></td>
<td>(0.84 – 0.94)</td>
<td>(40.63 – 44)</td>
</tr>
</tbody>
</table>

Notes: Results are mean ± SEM (95% confidence intervals).
Abbreviations: FEV₁, forced expiratory volume in 1 second; FVC, forced vital capacity.

Despite extensive publicity and promotion, clinical practice guidelines for the management of COPD were adopted by primary care practitioners and pulmonologists.

The results of these studies provide, in part, the evidence for spirometry-based COPD management guidelines and may, potentially, bias recommendations toward more treatment. Increased health care visits for respiratory symptoms may influence physicians to prescribe more respiratory treatments in an effort to alleviate symptoms and improve respiratory health regardless of a patient’s lung function measured by spirometry. Similarly, the Resource Use Study in COPD demonstrated that patients experiencing COPD exacerbations were prescribed significantly more medications than those who did not have respiratory symptoms.

In our unselected population, most health care encounters for COPD were confined to a small fraction of patients with COPD and this population may be disproportionately represented in some studies of respiratory medications. The results of these studies provide, in part, the evidence for spirometry-based COPD management guidelines and may, potentially, bias recommendations toward more treatment. Increased health care visits for respiratory symptoms may influence physicians to prescribe more respiratory treatments in an effort to alleviate symptoms and improve respiratory health regardless of a patient’s lung function measured by spirometry. Similarly, the Resource Use Study in COPD demonstrated that patients experiencing COPD exacerbations were prescribed significantly more medications than those who did not have respiratory symptoms.

Figure 2 Distribution of health care encounters over 12 and 60 month intervals.
Notes: Results are mean ± SEM. Panel A shows hospital, ED, outpatient, and total respiratory visits. Panel B shows hospital, ED, outpatient, and total nonrespiratory visits.
Abbreviations: Hospital, hospitalizations; ED, emergency department visits; Outpatient, outpatient health care encounters; All, total health care encounters.
utilization may be a distinguishing clinical characteristic that differentiates a phenotypically distinct group of individuals with COPD.\textsuperscript{18} Our results suggest that studies including only individuals with prior respiratory health care encounters may not be applicable to a general, unselected COPD population and may provide a partial explanation for the apparent less than recommended prescription of respiratory medications.\textsuperscript{19}

There is a paucity of available data to demonstrate that COPD treatment guidelines reduce health care utilization by individuals with COPD or affect mortality.\textsuperscript{20} An observational study comparing patients managed in general practices according to the British Thoracic Society guidelines or usual care found no differences in general practitioner or nurse visits, outpatient referrals, or hospitalizations.\textsuperscript{21} Similarly, we did not see fewer prior health care encounters in patients prescribed respiratory medications per guideline recommendations. In fact, for the entire cohort over 12 months, more total respiratory visits occurred in those receiving $G$ than for those who received $<G$ or $>G$ (0.6 $\pm$ 0.071(0.48, 0.75), 1.06 $\pm$ 0.19(0.85, 1.3), and 0.84 $\pm$ 0.17(0.58, 1.2) visits/person/year, for $<G$, $=G$, and $>G$, respectively (2 df ANOVA $P=0.002$). However, this study’s methodology did not allow us to determine whether this effect was caused by prescription of fewer respiratory medications for less symptomatic patients and prescription of increased treatments for more symptomatic patients regardless of their pulmonary function or whether step-wise pharmacologic management strategy...
based upon spirometry does not reduce respiratory health care encounters.

A systematic review to develop clinical practice guidelines for the diagnosis and management of stable COPD revealed insufficient evidence to support using spirometry to guide therapy. Furthermore, spirometry is not an effective guide for the management of patients with COPD and frequent exacerbations. The BODE index, a score based upon FEV1, 6 minute walk test, level of dyspnea, and body mass index, predicts mortality, hospitalization, and COPD exacerbation frequency and severity better than FEV1 alone. Analysis of patients enrolled in the National Emphysema Treatment Trial suggests that multifactorial models incorporating physiologic parameters, breathlessness, prior exacerbations, and comorbidities may prognosticate emergency department visits or hospitalizations. It is not yet known whether these factors or a multivariable index that includes prior respiratory health care visits can be used to guide the pharmacologic management of individuals with COPD.

The frequency of respiratory health care visits by individuals with COPD increased as the FEV1 declined and the modified GOLD stage increased whereas the number of nonrespiratory visits was the same regardless of GOLD stage. Although other investigators have demonstrated that the number of COPD exacerbations increases as the FEV1 decreases, the relationship between spirometry and exacerbation frequency is not consistent. Others have shown that multivariable indices such as the BODE index alone or combined with other parameters predict COPD exacerbations. However, these studies are difficult to compare because the clinical characteristics of the studied populations varied greatly as did the definitions of COPD exacerbation. In our population of unselected patients with a broad range of disease severity, the frequency of respiratory health care encounters increased as FEV1 declined.

In this study, we utilized only the Veterans Affairs medical record system and may have underestimated total health care utilization by omitting any encounters that occurred outside of the Veterans Health care Administration (VHA). Approximately 30%–40% of health care encounters occurred outside the VHA in a multicenter trial evaluating COPD medications based in Veteran Affairs medical centers. We required that all subjects have at least one encounter in the prior year to insure that they were actively receiving care at the Cincinnati VAMC and our results were similar for both the 12 and 60 month time periods suggesting stable respiratory health care utilization within our cohort. Furthermore the data within the electronic medical records available for our review was the same information that was available to the prescribing caregiver so it is representative of the clinical data upon which respiratory medications were prescribed.

Although our results are comparable to findings from other VAMC the findings of this study are restricted to a predominantly male population from a single medical center and will require validation in a larger, more diverse population of individuals with COPD. We used the pre-bronchodilator FEV1 rather than the post-bronchodilator FEV1 to classify patients according to the modified GOLD guidelines because the majority of patients had not undergone bronchodilator testing. We did not assess compliance with the prescribed medication regimen so it is possible that differences in patients’ adherence with their medications within the various GOLD defined groups might have contributed to the different rates of respiratory health care encounters and, possibly, to the prescription of more respiratory medications. Future prospective trials will be necessary to assess these factors.

This study demonstrates that respiratory medications prescribed for an unselected population of individuals with COPD of diverse severity complied poorly with the GOLD treatment guidelines. The majority of patients were prescribed fewer medications than recommended and these patients had fewer prior respiratory health care visits. Prescription of more than the recommended respiratory medications correlated with a history of increased frequency of respiratory health care encounters suggesting that more respiratory medications are prescribed for patients with increased health care utilization regardless of their lung function measured by spirometry. In addition, the majority of respiratory health care encounters occurred in a minor-
<table>
<thead>
<tr>
<th>Health care encounters</th>
<th>Clinical COPD</th>
<th>Modified GOLD 1 + 2</th>
<th>Modified GOLD 3 + 4</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Prescriptions per guidelines</td>
<td>More prescriptions than guidelines</td>
<td>Less prescriptions than guidelines</td>
</tr>
<tr>
<td><strong>Dependent variable</strong></td>
<td><strong>(-G)</strong></td>
<td><strong>(-G)</strong></td>
<td><strong>P value</strong></td>
</tr>
<tr>
<td>Respiratory Visits in 12 months visits/person/year</td>
<td>Hospital</td>
<td>0.032 ± 0.018</td>
<td>0.14 ± 0.071</td>
</tr>
<tr>
<td>ED</td>
<td>0.053 ± 0.023</td>
<td>0.22 ± 0.11</td>
<td>0.0004</td>
</tr>
<tr>
<td>Non-respiratory ED</td>
<td>0.19 ± 0.05</td>
<td>0.22 ± 0.09</td>
<td>0.67</td>
</tr>
<tr>
<td>All</td>
<td>0.27 ± 0.057</td>
<td>0.58 ± 0.19</td>
<td>0.016</td>
</tr>
<tr>
<td>Respiratory Visits in 60 months visits/person/year</td>
<td>Hospital</td>
<td>0.025 ± 0.0091</td>
<td>0.056 ± 0.02</td>
</tr>
<tr>
<td>ED</td>
<td>0.061 ± 0.014</td>
<td>0.17 ± 0.055</td>
<td>0.034</td>
</tr>
<tr>
<td>Non-respiratory ED</td>
<td>0.19 ± 0.014</td>
<td>0.27 ± 0.042</td>
<td>0.13</td>
</tr>
<tr>
<td>All</td>
<td>0.28 ± 0.03</td>
<td>0.49 ± 0.084</td>
<td>0.0032</td>
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<tr>
<td>Non-respiratory Visits in 12 months visits/person/year</td>
<td>Hospital</td>
<td>0.14 ± 0.046</td>
<td>0.33 ± 0.12</td>
</tr>
<tr>
<td>ED</td>
<td>0.42 ± 0.1</td>
<td>0.67 ± 0.2</td>
<td>0.15</td>
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<tr>
<td>Non-respiratory ED</td>
<td>3.0 ± 0.19</td>
<td>3.6 ± 0.4</td>
<td>0.078</td>
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<tr>
<td>All</td>
<td>3.5 ± 0.26</td>
<td>4.6 ± 0.56</td>
<td>0.028</td>
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<tr>
<td>Non-respiratory Visits in 60 months visits/person/year</td>
<td>Hospital</td>
<td>0.099 ± 0.023</td>
<td>0.18 ± 0.047</td>
</tr>
<tr>
<td>ED</td>
<td>0.37 ± 0.059</td>
<td>0.57 ± 0.12</td>
<td>0.077</td>
</tr>
<tr>
<td>Non-respiratory ED</td>
<td>2.5 ± 0.15</td>
<td>3.2 ± 0.25</td>
<td>0.034</td>
</tr>
<tr>
<td>All</td>
<td>3 ± 0.2</td>
<td>3.9 ± 0.32</td>
<td>0.014</td>
</tr>
</tbody>
</table>

Notes: Data are mean ± SEM (95% confidence intervals).
*ANOVA comparing prescription level; **2dfANOVA comparing prescription level.
Abbreviations: Hospital, hospitalization; ED, emergency department; Outpatient, outpatient visit; All, total health care encounters.

**Table 3: Respiratory and nonrespiratory health care encounters over the 12 and 60 month periods based upon modified GOLD stage and prescription level**
ity of patients and these phenotypically distinct patients may be over-represented in some studies of respiratory medications that provide, in part, the evidence for spirometry-based COPD treatment guidelines, potentially biasing recommendations toward increased prescription of respiratory medications. Prior respiratory-related health care encounters may provide a measure of a patient’s respiratory symptoms and may be an important factor influencing caregiver prescription of respiratory medications. In an unselected population with COPD, both a history of fewer respiratory health care encounters in an individual patient and guidelines based upon studies of patients with increased respiratory health care utilization may provide potential explanations for the prescription of less than recommended respiratory medications compared with the spirometry-based GOLD guidelines.

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References

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