

Understanding Nebulizer Utilization by Patients and Healthcare Providers: A COPD Foundation Nebulizer Consortium Survey Study

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Rationale: Inhaled medications are the mainstay of chronic obstructive pulmonary disease (COPD) management. While consensus guidelines for pharmacological management in COPD are well-established, few guidelines exist regarding inhaled medication delivery systems. The COPD Foundation Nebulizer Consortium conducted a cross-sectional survey of patients with COPD and healthcare providers to understand their perceptions and utilization of nebulized medications.

Methods: An online survey was conducted from February 7 through April 9, 2024. Patients completed a 42-question survey, including demographic information, tobacco use, symptoms severity, and the role of nebulizers in their treatment. Healthcare providers responded to a 17-question survey about their clinical experience with nebulized medications.

Results: We analyzed responses from 347 patients and 39 healthcare providers. Among patients, 76.4% (265/347) were ≥65 years old, 72.0% (250/347) were female, 93.4% (324/347) were white, 90.5% (314/347) had a current or former smoking history, 77.6% (263/339) reported at least one exacerbation in the past year, and 70.8% (240/339) used some form of supplemental oxygen. Nebulizer use was reported by 84.1% (292/347) of patients. Among nebulizer users, 94.5% (276/292) used short-acting while only 22.3% (65/292) used long-acting nebulized medications. Patients reported that hand-held inhaler devices were easier to use (69.8%, 171/245), but nebulized therapy led to better symptom control (64.9%, 159/245) and had lower copays (67.8%, 166/245). Among prescribers surveyed, most (82.1%, 32/39) believed nebulizers were preferable for patients experiencing exacerbations. Impediments to wider use of nebulizers included difficulties with insurance coverage (69.2%, 27/39), cost (53.8%, 21/39), and lack of combination nebulized drugs (46.2%, 18/39). Two-thirds of providers thought that nebulizers were underused.

Conclusion: We demonstrate that while patients and providers both perceive nebulizers as preferred in clinical management of COPD, there is discordance between patient and provider perception of nebulizer use on the basis of cost and feasibility of use.

Keywords: chronic obstructive pulmonary disease, COPD exacerbation, nebulizer, nebulized medication

Introduction

Inhaled medications are the mainstay of chronic obstructive pulmonary disease (COPD) management.^{1–3} While consensus guidelines for pharmacological management in COPD are well-established, few guidelines exist regarding inhaled medication delivery systems.^{1–3} Over time, the number and mode of delivery of inhaled medications have increased, now including short and long acting muscarinic antagonists as well as inhaled corticosteroids, offered alone in inhaler or nebulizer formulation or in a variety of combinations via hand-held inhalers.^{4–9} The four most commonly used delivery devices are metered-dose inhalers (MDI), dry powder inhalers (DPI), soft mist inhalers (SMI), and small volume nebulizers (SVN).⁴ Together, MDI, DPI, and SMI will herein be referred to as “hand-held inhalers” and are discussed in comparison to all types of SVNs, which include standard jet, vibrating mesh, breath-enhanced, breath-actuated, and ultrasonic nebulizers.

Overall, studies on inhaled medication delivery systems for COPD management have generally been equivocal, showing no consistent data to support the use of one delivery system over another.^{1,10–13} Prior work has evaluated both patient and provider perspectives on COPD management and inhalation delivery device selection. Patients and providers alike were more knowledgeable about, and paid most attention to, the pharmacologic class of medication, rather than the delivery system.¹⁴ Most patients were prescribed hand-held inhaler devices rather than nebulizers for delivery of both rescue and maintenance medication.^{15,16} Providers generally perceived SVNs to be superior to hand-held inhalers for treatment of a COPD exacerbation, and slightly more than half of patients who had used both types of devices preferred their nebulizer when compared to their hand-held inhalers.^{15,16}

During the COVID-19 pandemic, healthcare institutions and practitioners expressed concerns about infection risk and spread of respiratory viruses with the use of nebulizers, which resulted in further evaluation of nebulization therapy.¹ The COPD Foundation Nebulizer Consortium (CNC), a panel of subject matter experts, recently published their findings.¹⁷ In summary, the CNC concluded that adherence to recommended safety measures should be emphasized and nebulizer use should not be discouraged when clinically indicated.¹⁷ Further work showed that use of a breath-actuated nebulizer with a filtered mouthpiece did not lead to increased dispersion of SARS-CoV-2 RNA compared with an MDI in patients with COVID-19 infection.¹⁸

In part due to the COVID-19 pandemic, the CNC has worked to better understand the hesitancy about the use of nebulizers. Medication delivery remains predominantly via MDIs and DPIs with a significant shift away from SVNs as a delivery method.^{15,16} Meanwhile, new respiratory therapies have been approved via a nebulized delivery method, and updated nebulizer delivery systems, such as vibrating mesh nebulizers, have become commercially available.^{5,8,19–22}

Post-pandemic, we sought to re-evaluate the current state of nebulizer prescribing and utilization patterns. The CNC conducted a cross-sectional survey of patients with COPD and healthcare providers to better understand their perception of nebulizers and their utilization in clinical practice.

Methods

Survey Development

The CNC designed and distributed a survey to patients and healthcare providers regarding nebulizer use for the treatment of COPD. The patient section of this survey was reviewed by patients who were on the CNC. Those questions were written using fifth grade level English, and critical sections like questions about their medications included images of the medicine brand and labeling to facilitate selection. The healthcare provider section was written using college level English and included medical terms.

Survey Format and Administration

The CNC online survey was deemed IRB Exempt under 45 CFR § 46.104(d)(2) by the Western Institutional Review Board (WCG IRB, Puyallup, WA, USA), because this research study only involves a survey and data collection maintained the privacy of its subjects in compliance with the Declaration of Helsinki. Participation was voluntary, and the following language was present on the introduction page of the survey and served as the consent process:

The following short survey is designed to better understand the use of nebulized drugs in COPD. Your responses are completely anonymous and will not be linked back to you. By completing the survey, you acknowledge that the COPD Foundation has your consent to analyze the anonymous responses (data). Your participation in the survey is completely voluntary, and you are not required to share any information that could identify you.

The survey was conducted from February 7 through April 9, 2024.

Participants included adult patients with COPD who resided in the United States and its Territories and healthcare providers recruited through the COPD Foundation's online patient platform, newsletters, and social media channels, and healthcare providers were recruited via COPD Foundation and the CNC provider network. There was no financial incentive for participation. Patients were included if they confirmed they had a diagnosis of COPD, were 18 years of age or older, and lived in the United States. Healthcare providers, such as pulmonologists, primary care practitioners, nurse practitioners and physician assistants were among the respondents asked whether they prescribed nebulizers. Those who prescribed nebulizers and confirmed that they practiced in the United States were asked to complete the rest of the survey.

Survey Content

Patients completed a 42-question survey ([Supplemental Figure 1](#)), including demographic information, smoking status and lifetime cumulative amount in pack-years, and history of lung disease. Specifically, they described their tobacco, marijuana, and vaping use currently and historically. They also described their current level of daily activities, symptoms, and number of COPD exacerbations per year. The final section of the survey included questions about oxygen use, corticosteroid use for exacerbations, and the role of nebulized medication and delivery devices in their treatment regimen.

Healthcare providers were asked 17 questions ([Supplemental Figure 2](#)) about their prescribing experience regarding nebulizers, selected nebulized medications, hand-held inhalers, reasons for selection of medications, and patient characteristics considered when selecting nebulizer devices and medications.

Data Analysis

Data was collected via Survey Monkey [SurveyMonkey Inc.; San Mateo, California, USA; www.surveymonkey.com] using the platform's software. Due to the nature of survey distribution, no formal sample size calculations for this convenience sample were able to be performed. Descriptive analyses of the survey results are provided for both patient and healthcare provider surveys. Comparisons between groups for hand-held inhaler and nebulizer preferences were done using a Chi-square test or Fisher's Exact Test where appropriate. Analyses were conducted using SAS Version 9.4 (SAS Institute, Cary, NC, USA).

Results

We analyzed 347 responses from patients and 39 responses from healthcare providers who met the inclusion criteria. Response rate is not known given that the survey was distributed to an unknown number of participants via the listed online platforms.

Patient Characteristics

Patient characteristics are shown in [Table 1](#). Notably, 76.4% (265/347) were ≥ 65 years old, 72.0% (250/347) were female, 93.4% (324/347) were white. Survey participants reported using a variety of health insurances including private insurance (17.3%, 60/347), Medicare (55.0%, 191/347), and Medicaid (10.4%, 36/347), and they spanned socioeconomic groups with 40.3% (140/347) reporting total household incomes of less than \$40,000 per year. Most of the patients surveyed (80.7%, 280/347) had a pulmonologist involved in their respiratory-related care.

Most patients (82.7%, 287/347) were former tobacco smokers, only 7.8% (27/347) were currently smoking, and 9.5% (33/347) were never-smokers ([Table 1](#)). Among current and former smokers, median [inter-quartile range (IQR)] pack-year history was 49 [35–80]. Only 10.4% (36/347) of patients reported any marijuana use. Among marijuana users,

Table 1 Patient Characteristics

Characteristic	Patient Surveys (N=347)
Age ≥ 65 years	265 (76.4%)
Sex, Female	250 (72.0%)
Race, White	324 (93.4%)
Body Mass Index (kg/m ²)	
< 18.5	47 (13.5%)
18.5 to < 25	99 (28.5%)
25 to < 30	104 (30.0%)
≥ 30	97 (28.0%)
Household income (US dollars)	
< 40,000	140 (40.3%)
40,000–80,000	93 (26.8%)
> 80,000	38 (11.0%)
Prefer not to answer	76 (21.9%)
Health Insurance ^a	
Medicare	191 (55.0%)
Medicare Advantage	106 (30.5%)
Medicare Supplement	69 (19.9%)
PPO/HMO/POS private plan	60 (17.3%)
Medicaid	36 (10.4%)
Disability	8 (2.3%)
No health insurance	1 (<1%)
Region of residence ^b	
South	114 (32.9%)
Northeast	84 (24.2%)
Midwest	84 (24.2%)
West	65 (18.7%)
Smoking status	
Never	33 (9.5%)
Former	287 (82.7%)
Current	27 (7.8%)
Current marijuana use, any type	36 (10.4%)
Via smoking and/or vaping	13 (3.7%)

Notes: ^aHealth insurance was self-selected, and patients chose all that applied. ^bRegions defined by the United States Census Bureau.

31.6% (11/36) used marijuana on a daily basis, with less than half using it in an inhaled form (31.6% or 11/36 smoking and 5.6% or 2/36 vaping).

Patient COPD Disease Severity

Survey participants generally had moderate to severe COPD based on symptoms, exacerbation history, medications, and oxygen use (Table 2). Most patients (96.5%, 327/339) experienced some degree of daily symptoms, indicated by modified Medical Research Council (mMRC) dyspnea score of 1 or more, with 21.5% (73/339) being too short of breath to leave the house (as indicated by a modified Medical Research Council score of 4) and 70.8% (240/339) using supplemental oxygen. The majority (77.6%, 263/339) reported at least one exacerbation that required steroids and/or antibiotics in the past year.

Table 2 Patient Clinical Characteristics

Characteristic	Patient Surveys (N=347)
mMRC ^a	
n	339
0 – Short of breath only with strenuous exercise	12 (3.5%)
1 – Short of breath when hurrying or walking up a slight hill	73 (21.5%)
2 – Walk slower or stop for breath because of dyspnea	99 (29.2%)
3 – Stop for breath after walking 100 yards or after a few minutes	82 (24.2%)
4 – Too short of breath to leave house or breathless when dressing	73 (21.5%)
Exacerbation in the past year	339
n	76 (22.4%)
0	88 (26.0%)
1	175 (51.6%)
2 or more	
Use of supplemental oxygen at home	
n	339
None	99 (29.2%)
Any use	240 (70.8%)
Continuously	147 (43.4%)
Only during the night	52 (15.3%)
Only during exertion	41 (12.1%)
How often do you use a nebulizer?	
n	347
I have not used a nebulizer / No response	55 (15.8%)
Any use	292 (84.2%)
Regular daily, scheduled use	138 (39.8%)
Regular daily, as needed	60 (17.3%)
Occasional use, rescue use	94 (27.1%)

Notes: ^aWorst response was taken where multiple categories were selected. 8 responses were missing for mMRC, exacerbation and oxygen questions.

Abbreviation: mMRC, modified Medical Research Council dyspnea scale.

Patient-Reported Nebulizer Use

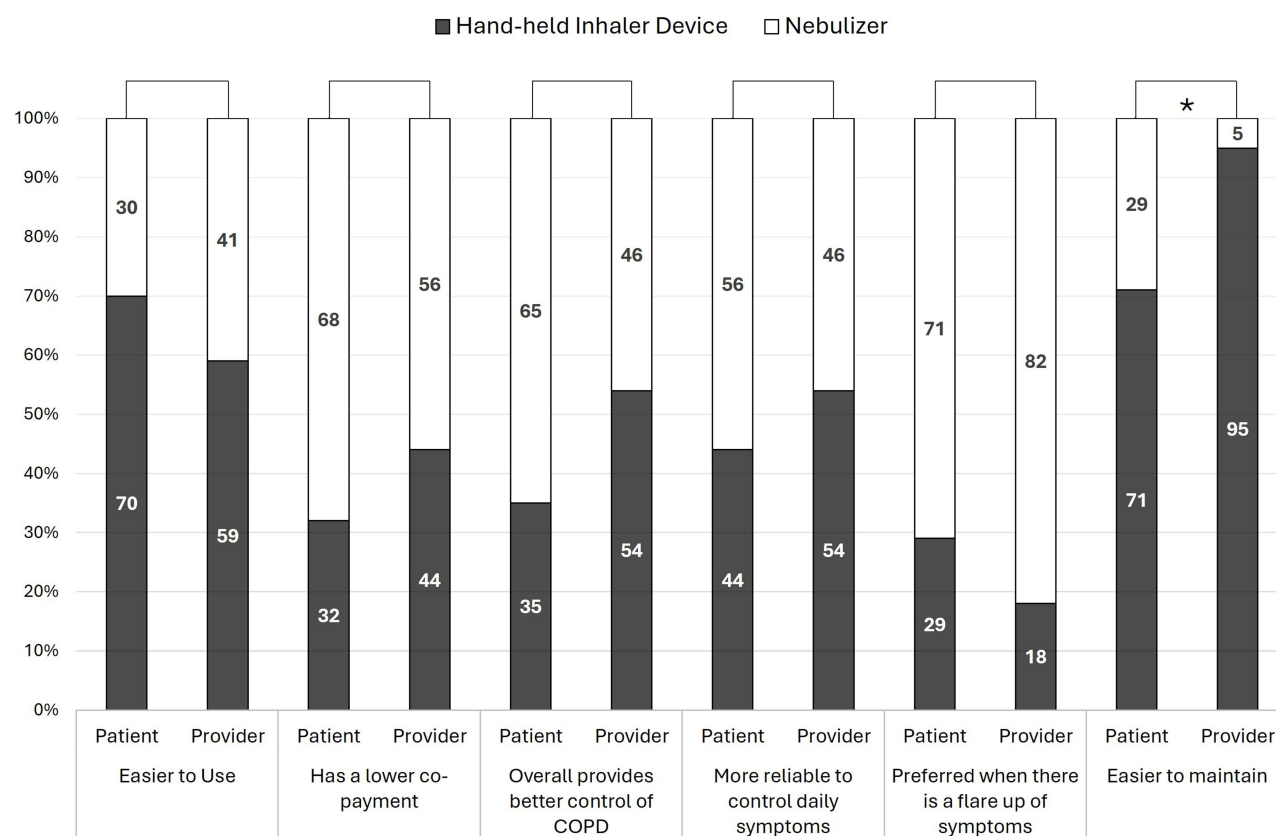
The majority of surveyed patients (84.2%, 292/347) had used nebulizers in the past, with 57.1% (198/347) reporting daily use (Table 2). Jet nebulizers as a category include standard (or continuously operated) models, as well as more advanced types such as breath-enhanced and breath-actuated jet nebulizers. Among nebulizer users, standard jet (36.6%, 107/292) and breath-enhanced nebulizers (29.8%, 87/292) were used more frequently compared to breath-actuated (16.1%, 47/292), vibrating mesh (13.7%, 40/292), and ultrasonic (13.0%, 38/292) nebulizers. Nebulizers were initially prescribed most often by the patient's pulmonologist (53.4%, 156/292), followed by the primary care doctors (20.6%, 60/292). The most common reason patients reported for receiving nebulized therapy included worsening symptoms (40.4%, 118/292) or because other inhaler devices did not work well enough (21.9%, 64/292) (Table 3).

Among patients who reported using nebulizers, most (94.5%, 276/292) used nebulized short-acting bronchodilators, which included beta agonists (63.0%, 184/292), muscarinic antagonists (8.6%, 25/292), or a combination of both (18.2%, 53/292). Only 22.3% (65/292) used one or more long-acting nebulized medications, inhaled corticosteroid most frequently (16.8%, 49/292) followed by long-acting beta agonists (11.0%, 32/292) and long-acting muscarinic antagonists (8.9%, 26/292). Aside from inhaled steroids or bronchodilators, 10.3% (30/292) used other nebulized medications such as hypertonic saline, N-acetylcysteine, or inhaled antibiotics.

Table 3 Nebulizer User Characteristics

Characteristic	Nebulizer Users (N=292)
Who first prescribed the nebulized drug for you?	
My pulmonologist	156 (53.4%)
My primary care physician	60 (20.6%)
Emergency/Urgent care doctor	16 (5.5%)
Nurse practitioner or physician assistant	10 (3.4%)
Other	12 (4.1%)
No response	38 (13.0%)
Why did you start using nebulizers? (select all that apply)	
Worsening of my symptoms	118 (40.4%)
Unsure, it was just prescribed by my doctor	94 (32.2%)
Other inhaler devices did not work well-enough for me	64 (21.9%)
Insurance/copayment reasons	16 (5.5%)
Personal preference	13 (4.4%)
Other	27 (9.2%)

Of the 292 patients who use nebulizers, 254 responded to the nebulizer training question, and of those 21.6% (55/254) reported that no teaching about the use of nebulizers was provided by the prescribing team, and another 24.8% (63/254) stated that teaching, when provided, was perceived to be insufficient. In comparison, teaching about the use of hand-held devices was missing or insufficient in 16.1% (41/254) and 22.4% (57/254) of cases, respectively.



* $p < 0.001$

Figure 1 Patient and provider perception of hand-held inhaler devices versus nebulizers.

245 patients responded to the nebulizer preference questions. Compared to hand-held inhaler devices, surveyed participants reported that nebulized therapy had lower co-pay (67.8%, 166/245) and led to better COPD symptom control (64.9%, 159/245), particularly during an exacerbation (71.0%, 174/245). However, hand-held inhaler devices were perceived as easier to use compared to nebulizers (69.8%, 171/245), and easier to maintain (71.0%, 174/245) (Figure 1). When divided by patient characteristic, patients who were 65 years or older more often found that nebulizers resulted in better daily symptom control compared to younger patients (60% vs 44%, $p = 0.0353$), whereas those who used oxygen or had one or more exacerbation more often preferred nebulizers during an flare up of symptoms (76% vs 56% $p = 0.0017$ and 75% vs 54% $p = 0.0057$ respectively) (Table 4).

Regarding the impact of the recent COVID-19 pandemic on the utilization of nebulized drugs, the majority (55.5%, 141/254) reported never having had a COVID-19 infection. Of the 109 patients who reported having COVID-19, 21.1% (23/109) reported being hospitalized, including three who reported being placed on a ventilator for COVID-19 pneumonia during the pandemic. A COVID-19 diagnosis did not change home nebulizer use for most patients (75.2%, 82/109), although 5.5% (6/109) decreased use or changed from using nebulizers.

Provider Responses

Of the 113 healthcare providers who initiated the survey, 55 did not prescribe nebulizers and 19 did not practice in the United States, leaving 39 that met our inclusion criteria. Of these 39 providers, 84.6% (33/39) were pulmonologists. Most providers prescribed jet nebulizers (89.7%, 35/39), followed by ultrasonic (28.2%, 11/39) and vibrating mesh nebulizers (20.5%, 8/39). A majority of prescribers (71.8%, 28/39) indicated that more than 75% of patients for whom they prescribed nebulizers used short-acting bronchodilators, whereas most indicated that long-acting bronchodilators or inhaled corticosteroids were used in less than 20% of their patients (61.5% or 24/39 and 53.8% or 21/39, respectively). The most frequently reported drugs prescribed for nebulization were albuterol and budesonide (each 92.3%, 36/39), followed by albuterol-ipratropium combined formulation (84.6%, 33/39), ipratropium (71.8%, 28/39), and long-acting medications with less frequency (revefenacin at 61.5% or 24/29; arformoterol at 46.2% or 18/46; formoterol at 35.9% or 14/39; glycopyrrolate at 7.7% or 3/39). Other nebulized therapies that were also prescribed by these providers included hypertonic saline (79.5%, 31/39), inhaled antibiotics (41.0%, 16/39), and N-acetylcysteine (25.6%, 10/39).

Prescribing providers were asked to rank the reasons for initiating nebulized drugs on a 5-point scale from 1 (least important) to 5 (most important), with higher scores indicating greater perceived importance. The most important factor for prescribing nebulizers were reported to be the need for better control of symptoms than provided by hand-held

Table 4 Summary of Preference for Nebulizer Over Hand-Held Inhaler by Patient Demographic Splits

Category	Age 65+ (n=186) ^a	Age < 65 (n=57)	p-value
Easier to Use	55 (30%)	18 (32%)	0.7722
Overall provides better control of COPD	124 (67%)	33 (58%)	0.2256
More reliable to control daily symptoms	111 (60%)	25 (44%)	0.0353
Preferred when there is a flare up of symptoms	136 (73%)	36 (63%)	0.1480
	Uses oxygen (n=182)	No oxygen use (n=63)	p-value
Easier to Use	60 (33%)	14 (22%)	0.1094
Overall provides better control of COPD	121 (66%)	38 (60%)	0.3768
More reliable to control daily symptoms	109 (60%)	29 (46%)	0.0559
Preferred when there is a flare up of symptoms	139 (76%)	35 (56%)	0.0017
	None ^b (n=46)	One or more ^b (n=199)	p-value
Easier to Use	14 (30%)	60 (30%)	0.9698
Overall provides better control of COPD	22 (48%)	137 (69%)	0.0071
More reliable to control daily symptoms	21 (46%)	117 (59%)	0.1053
Preferred when there is a flare up of symptoms	25 (54%)	149 (75%)	0.0057

Notes: ^aTwo patients declined to give their age and are not included in this summary. ^bGroup divided by number of self-reported exacerbations over the preceding year.

Table 5 Factors Considered by Healthcare Providers When Deciding to Prescribe Nebulized Therapy

Factor	Most Important	More Important	Neutral	Less Important	Least Important	Average Rating
Persistence of symptoms despite hand-held inhaler use and failure to control disease with hand-held devices	23	15	1	0	0	4.56
Patient's dexterity and hand-mouth coordination issues	20	18	0	1	0	4.46
Patient's cognitive issues	13	21	4	1	0	4.18
Patient's cognitive skills	11	23	3	2	0	4.10
History of hospitalization(s) for COPD	14	17	6	1	1	4.08
Peak inspiratory flow inadequacy	14	16	6	3	0	4.05
Patient's exacerbation history	11	21	5	1	1	4.03
Patient's severity of obstruction (severity of FEV1 compromise)	8	25	4	2	0	4.00
Patient's preference	10	17	12	0	0	3.95
Comfort	7	18	13	1	0	3.79
Significant comorbidities	7	16	14	2	0	3.72
Patient's age	0	21	15	3	0	3.46
Patient's lifestyle	0	22	13	4	0	3.46
Patient's insurance	6	13	13	3	4	3.36
Socio-economic status	0	8	18	6	7	2.69

Notes: Ratings were scored ranging from "Most important" = 5 to "Least important" = 1. Average rating is the mean numerical score for the 39 provider responses.

inhalers, followed closely by cognitive or dexterity issues (Table 5). Factors such as patient preference, comfort and socio-economic status were rated as less influential, illustrating that prescribers prioritize clinical need and functional limitations over convenience or cost when selecting a delivery device. When asked to compare the benefits of nebulizers versus hand-held inhalers, prescribers reported that nebulizers were preferable to hand-held inhalers for patients experiencing exacerbations (82.1%, 32/39), whereas 53.8% (21/39) believed hand-held inhalers were preferable for routine symptom control (Figure 1).

For hand-held inhaler device versus nebulizer preference, there were no statistical differences between surveyed providers' perception of patient preference and surveyed patients' preference for all surveyed dimensions ($p > 0.17$), with one the exception that more providers believed that hand-held inhaler devices are easier to maintain (94.9% vs 71.0%, $p = 0.0007$) (Figure 1).

Common provider-perceived barriers to wider use of nebulizers among their patients were lack of patient insurance coverage or approvals (69.2%, 27/39), patient cost (53.8%, 21/39), and lack of availability of combination drugs for nebulization (46.2%, 18/39). Two-thirds (26/39) of providers thought that nebulizers were underused. Providers' suggested interventions most likely to increase the use of nebulizers among their patients (in descending order from most to least likely) were availability of combination long-acting drugs (79.5%, 31/39), optimization of insurance coverage and cost (76.9%, 30/39), availability of once-daily drugs (71.8%, 28/39), and improved portability of nebulizers (56.4%, 22/39).

The surveyed prescribers of nebulized pharmacotherapies believed that the COVID-19 pandemic negatively impacted the utilization of nebulizers in hospital settings (48.7%, 19/39) more than in outpatient settings (20.5%, 8/39), but they

generally believed that the use of nebulizers has currently returned to the pre-pandemic degree of use in both hospital (69.2%, 27/39) and outpatient settings (74.4%, 29/39).

Discussion

Nebulized medications are widely used by patients to treat COPD. Despite consensus guidelines and numerous clinical trials that have informed pharmacological management of COPD,^{1–3} as well as the ongoing development of inhaled medications via a variety of delivery systems, there are scant recommendations as to which delivery system to select.^{1,4–8} Similar to prior data,^{15,16} our survey found that both patients and providers believed that nebulizers offer superior symptom control compared to hand-held inhaler devices, particularly during an exacerbation. This effect was best seen in older patients, those who use oxygen, and those who have had an exacerbation in the preceding year, suggesting that nebulizers might be particularly beneficial for these high risk groups. This is in contrast to the published literature that does not demonstrate a consistent objective benefit of nebulizers over hand-held inhalers. Some evidence suggests that these findings may have been confounded by the exclusion of patients who did not demonstrate proper technique with hand-held inhalers and the clinically stable setting in which these studies were performed.^{10–13} With regard to the latter, symptoms or air-trapping during an exacerbation could interfere with the ability of patients to use a hand-held inhaler with proper technique. Interestingly, we found that home nebulizer use and prescribing patterns have not been impacted significantly by the COVID-19 pandemic, despite initial concerns that nebulizers may result in increased infectious risk.

In our study, despite perceived benefits by prescribers and patients, as well as the viewpoint of providers that nebulized medications are underutilized, nearly half of the providers do not prescribe nebulized medications. Also, while 95% of patients in our survey who use nebulizers reported using nebulized short-acting bronchodilators, the use of long-acting nebulized medication, including the long-acting beta-2 agonists, formoterol and aformoterol, the long-acting antimuscarinic bronchodilators, glycopyrrolate and revefenacin, and the inhaled corticosteroid, budesonide, was notably infrequent. With the recent advent of these two nebulized long-acting antimuscarinic bronchodilators,^{19–22} the number of prescriptions for these drugs as controller medications for select patients may increase. Both glycopyrrolate and revefenacin have been approved for use over the last several years, and more recently after the conclusion of our survey, a novel dual phosphodiesterase 3 and 4 inhibitor, ensifentrine, was approved.²³ Moreover, evidence is lacking concerning the efficacy and safety of administering long-acting nebulized controller medications, including long-acting bronchodilators and inhaled corticosteroids, administered in combination, potentially limiting their incorporation into daily practice for the maintenance of moderate to severe COPD.

Our survey explored barriers to nebulizer use by patients and prescription by providers. First, patients consistently reported that nebulizers are considerably more cumbersome to use and maintain compared to hand-held inhalers, although providers seemingly overestimated the degree to which maintenance is prohibitive for patients. Providers also suggested that increased portability and availability of once-daily controller medications would increase nebulizer use. Second, we found that providers perceived that nebulized medications were just as, if not more, expensive than similar drugs available in hand-held inhalers, due to either higher copays or lack of insurance coverage. Interestingly, patients reported the opposite; some nebulized medications were more affordable than hand-held inhalers, often covered through Medicare Part B. Physicians self-reported that the lack of combination long-acting medications as a barrier to more frequent prescribing of nebulized medications. This may help explain why nebulizers are used predominantly for short-acting medications rather than controller medications. Together, this demonstrates discordance between patient and provider perception of nebulizer utility in the clinical management of COPD based on cost and ease of use, but both patient and provider are aligned in the perceived superior symptom control with nebulizer utilization.

There are other possible benefits of nebulizer use as an alternative to hand-held inhalers that exist but are outside the scope of this survey. Several studies have reported the adverse environmental impacts of MDIs.^{24–27} Recommendations have suggested that use of DPIs and SMIs are preferred to MDIs to minimize the use of hydrofluorocarbons and decrease the carbon footprint. Future work could explore the potential environmental benefit of nebulized medications as well.

Our study has several limitations. First, given the survey distribution methods, which included posting on social media, online platforms, and via newsletters, we are not able to calculate a response rate. Similarly, we acknowledge that our cohort may not accurately reflect all COPD patients, particularly those who do not access to online platforms. Our

patients were predominantly female, nearly entirely white, and over half had not been infected by SARS-CoV-2. However, we did have a representative and diverse geographic and socioeconomic sample. Additionally, this is a self-reported patient and provider survey, which carries with it recall bias. Further, patients may not have understood or misremembered details, such as which medications they have been prescribed or which type of nebulizer machine. We did not collect objective patient data to accurately classify their COPD disease severity, although the self-reported questions did give us information to understand that this patient population generally has severe disease. The small sample size limited our ability to perform comparative analysis within the patient or provider cohort. Additionally, not all participants completed the survey in full, leaving some questions with fewer responses than others. Despite these limitations, this is a hypothesis-generating survey to help us understand prescriber and patient utilization practices in the wake of the COVID-19 pandemic.

Conclusions

Our data indicate that nebulized medications are perceived as more affordable and reported to offer better control of chronic disease and acute exacerbation symptoms than hand-held inhalers by both patients and providers alike. However, nebulized therapy is prescribed less often due to several physician-perceived factors, including increased burden to maintain, comparative ease of inhaler maintenance, higher cost of nebulizers, and lack of combination long-acting nebulized therapies. Further work is needed to better understand the relative benefits and drawbacks of nebulized medications, examine the discordance between perceived and objective clinical benefit of nebulized medications, and explore ways of optimizing the identification of patients who will derive the most benefit from their use.

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We would like to acknowledge the additional member of the COPD Foundation.

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