

Brief Report

Improving Referral Patterns for Bronchoscopic Lung Volume Reduction: A Quality Improvement Initiative

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Abstract

Bronchoscopic lung volume reduction (BLVR) is a minimally invasive treatment option for patients with severe emphysema and hyperinflation refractory to optimal medical care. This therapy is effective in improving functional status and quality of life, underscoring the importance of identifying potential procedure candidates. To our knowledge, scalable strategies to improve the referral of advanced lung disease patients are lacking. This quality improvement project aimed to increase identification and referral for BLVR in a large Veterans Affairs academic medical center. We show implementing case identification within a pulmonary function testing report, in conjunction with provider education, increased referral rates for BLVR. Because of the ubiquity of lung function testing, other advanced lung disease programs may consider adopting this strategy to improve patients' access to timely clinical evaluation and therapy.

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Abbreviations:

BLVR=bronchoscopic lung volume reduction; **COPD**=chronic obstructive pulmonary disease; **DLCO**=diffusing capacity for carbon monoxide; **EMR**=electronic medical record; **FEV₁**=forced expiratory volume in 1 second; **GOLD**=Global initiative for chronic Obstructive Lung Disease; **IQR**=interquartile range; **LVRS**=lung volume reduction surgery; **PCP**=primary care provider; **PFT**=pulmonary function testing (tests); **QI**=quality improvement; **RV**=residual volume; **SD**=standard deviation; **TLC**=total lung volume capacity

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Introduction

The health and economic burden of chronic obstructive pulmonary disease (COPD) is profound, with global prevalence¹ estimated at roughly 10%. Disease prevalence is even higher in certain at-risk populations, with both male and female United States military veterans demonstrating higher rates of COPD compared to non-veterans.² Hyperinflation is one of the many pathophysiologic changes seen in COPD that is known to contribute to patients' symptom burden and is an independent risk factor for mortality.³ Lung volume reduction surgery (LVRS) can reduce the impact of

hyperinflation on pulmonary mechanics in patients with advanced COPD refractory to optimized medical care.⁴ While LVRS was shown to improve several patient-centered outcomes in a large clinical trial, the procedure requires relative surgical fitness and many patients are ineligible.

Bronchoscopic lung volume reduction (BLVR) is a minimally invasive alternative to LVRS for select patients with advanced emphysema and is supported by the Global initiative for chronic Obstructive Lung Disease (GOLD) treatment guidelines.¹ Patient selection for this procedure follows the inclusion and exclusion criteria of the associated clinical trials investigating BLVR.^{5,6} A key step in the evaluation of potential procedure candidates is pulmonary function testing (PFT) to identify severe airflow obstruction, hyperinflation, and gas trapping. Patients with hyperinflated, severely emphysematous type COPD who undergo BLVR show improved lung function, exercise performance, quality of life, and survival.⁵⁻⁷ In a survey of advanced emphysema patients queried about their treatment preferences, most respondents opted for BLVR over continued medical management.⁸

Although many patients may be interested candidates, strategies to improve identification and referral for BLVR are lacking. This quality improvement (QI) project explores the implementation of a referral strategy using PFT reports and provider education to identify BLVR-eligible patients in a large Veterans Affairs academic medical center.

Methods

The Veterans Affairs Northeast Ohio Healthcare System has been offering BLVR at the Louis Stokes Cleveland Department of Veterans Affairs Medical Center location since January 2022. This QI project aimed to identify potentially eligible BLVR patients for referral and evaluation. This QI project was exempt from IRB review. Before our QI initiative, BLVR evaluations primarily relied on word-of-mouth referrals within the pulmonary section. Our approach was to: (1) develop a process to communicate eligibility to ordering providers using PFT reports, and (2) educate providers about BLVR and how to refer patients to our program. We tracked BLVR referrals for 8 months pre- and postintervention to assess effectiveness in improving referral patterns.

In September 2022, we launched a restructured PFT template in the electronic medical record (EMR) to indicate potential BLVR qualification and outline steps for referral. When applicable PFT parameters were met (prebronchodilator forced expiratory volume in 1 second [FEV₁] < 50% and residual volume [RV] > 175%), the following comment would be included in the PFT

report: “The combination of gas trapping and severe obstruction may qualify patients for lung volume reduction. Consider pulmonary referral for evaluation if clinically indicated. Consults -> Pulmonary -> COPD -> BLVR.” The EMR sent an automated notification to the PFT-ordering provider of the completed results for review. To improve referral tracking, a separate BLVR consult order was added to the EMR. An overview of the PFT review and referral process is provided in Figure 1.

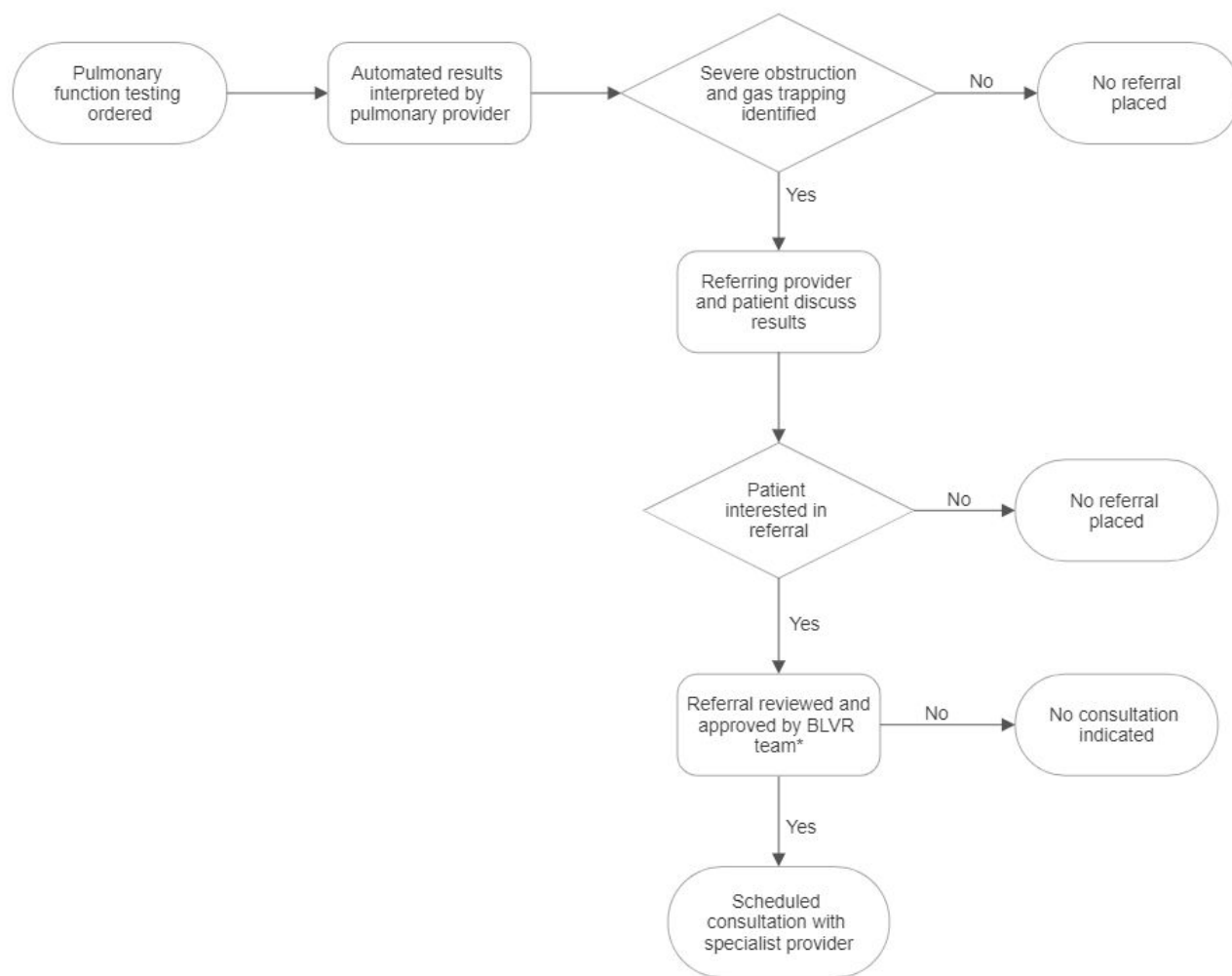
As part of the implementation process, pulmonary and primary care medicine staff were invited to a lecture focused on BLVR eligibility and the referral process specific to our institution. All BLVR evaluation referrals were reviewed by an interventional pulmonary provider and nurse navigator. Upon referral, patient eligibility for BLVR was determined based on previously published inclusion and exclusion criteria.^{5,6} As an internal quality check, our PFT system was queried biweekly to ensure the BLVR referral comment was added by the interpreting pulmonologist when PFT criteria were met.

Data Collection & Analysis

Baseline demographic information including age, gender, smoking status, PFT results, referring provider type, and comorbidities were reviewed. The comorbidity burden was calculated using the Charlson Comorbidity Index score.⁹ Charlson Comorbidity Index estimates the 10-year survival based on patient comorbidities. Comorbid conditions include age (1 point for every decade age 50 years and over), myocardial infarction, congestive heart failure, peripheral vascular disease, cerebrovascular accident or transient ischemic attack, dementia, COPD, connective tissue disease, peptic ulcer disease, liver disease (mild or moderate/severe), diabetes (with end-organ damage or uncomplicated), moderate to severe chronic kidney disease, hemiplegia, localized solid tumor, metastatic solid tumor, leukemia, lymphoma, and AIDS. Summary statistics for the postintervention cohort are presented as mean ± standard deviation (SD), median [interquartile range (IQR)], or n (%), as appropriate. An interrupted time series analysis was used to evaluate the change in referrals relative to the intervention and the time after intervention. The 2-proportion z-test was used to compare eligibility before and after intervention. Data analysis was conducted using R software¹⁰ version 4.1.2.

Results

The postintervention cohort included 118 patients who were identified as potentially BLVR-eligible based on PFT criteria between September 2022 and April 2023 (Table 1). The mean age of the population was 69 years old, of which 97%

Figure 1. An Overview of the Pulmonary Function Test Review and Referral Process

*The BLVR team included a pulmonary provider and nurse navigator. Each referral was reviewed to ensure PFT criteria were met and no obvious contraindications to the procedure were identified in the EMR. Eligible patients were mailed an institution-approved patient education packet and scheduled for consultation.

BLVR= bronchoscopic lung volume reduction; PFT=pulmonary function testing; EMR=electronic medical record

were male. More than 80% were GOLD stage 3 or 4 with a mean postbronchodilator FEV₁ of 38.3%±11.5% and an RV of 216% (196, 246). Most patients were former smokers with a mean Charlson Comorbidity Index of 4.12±1.35. The PFT-ordering providers were predominantly pulmonary (55.1%) and primary care (41.5%) specialists.

Before the intervention, 14 patients were referred to our program between January and August 2022 for BLVR evaluation. Of the 14 BLVR evaluations, 7 patients were candidates for the procedure, a rate of 0.9 eligible referrals per month. After implementation of the restructured PFT template, 31 patients had a referral placed for BLVR, 4 of whom were not sent for evaluation due to a procedure contraindication. A total of 27 patients were evaluated for BLVR, of which 15 were candidates for the procedure, a rate of 1.9 eligible referrals per month (Figure 2). All evaluations for BLVR were referred by either a pulmonary (70%) or primary care (30%) provider. Two of the 12 BLVR-ineligible patients were referred for evaluation of surgical lung volume reduction and lung transplantation.

Overall, the intervention led to a significant increase in the number of BLVR referrals ($\beta=4.6$, $p=0.007$). There was no significant trend (e.g., regression toward the number of referrals before intervention) in the 8 months after intervention ($\beta=-0.61$, $p=0.07$). In addition, there was no significant difference in the percentage of eligible patients before and after the intervention ($p=0.99$).

Discussion

The severe emphysema population continues to grow, making advanced therapeutic options like BLVR an important adjunct treatment strategy.¹¹ Developing a system to identify eligible patients and educate providers about these therapies may significantly impact population morbidity and mortality. After implementing an institution-specific QI initiative, we observed a marked increase in the number of BLVR evaluations, and twice as many eligible referrals, a finding that was sustained in the 8 months of monitoring after the intervention. The positive change in referral patterns was

Table 1. Characteristics of Potential Bronchoscopic Lung Volume Reduction-Eligible Patients Using the Pulmonary Function Test Identification Strategy

Characteristic	Postintervention Cohort
n	118
Age, yr	69.4±8.2
Male	114 (96.6)
Post-BD FEV ₁ , % predicted	38.3±11.5
RV, % predicted	215.5 [196, 246]
TLC, % predicted	118.5 [112, 134]
DLCO, % predicted	44 [35, 59]
GOLD Stage	
2	19 (16.1)
3	70 (59.3)
4	29 (24.6)
Smoking Status	
Current	36 (30.5)
Former	80 (67.8)
Never	2 (1.7)
PFT Ordering Provider	
Pulmonary	65 (55.1)
PCP	49 (41.5)
Surgery	2 (1.7)
Cardiology	1 (0.8)
Oncology	1 (0.8)
Charlson Comorbidity Index	4.12±1.35

Summary statistics for the post-intervention cohort are presented as mean± (standard deviation), median [IQR], or n (%), as appropriate.

post-BD=postbronchodilator; FEV₁=forced expiratory volume in 1 second; RV=residual volume; TLC=total lung capacity; DLCO=diffusing capacity for carbon monoxide; GOLD=Global initiative for chronic Obstructive Lung Disease; PFT=pulmonary function test; PCP=primary care provider; IQR=interquartile range

partly attributed to the efforts of primary care providers who prior to the intervention did not account for any BLVR evaluations. In fact, we were pleasantly surprised that the percentage of eligible referrals was similar before and after implementation, suggesting many primary care specialists, like pulmonary, are providing optimal COPD care. Even for ineligible patients, this was a value-added consultation as they were assessed for optimal COPD therapy and other advanced interventions including surgical lung volume reduction and lung transplantation. Moreover, those yet to complete pulmonary rehabilitation or abstain from tobacco products for at least 3 months are still being longitudinally followed for BLVR eligibility.

There are some limitations to our work. While the number of BLVR referrals increased after project implementation, we cannot definitively attribute this change to restructuring the PFT report alone. It could be argued that increasing awareness and education drove the improvement in referrals; however, educational interventions tend to fade over time, meaning if the improvement in referrals was only due to education, we would expect decreasing effectiveness the longer from the intervention^{12,13} —a pattern not observed in our post-implementation cohort. Therefore, the PFT identification strategy, combined with provider education, is likely what accounts for the sustained

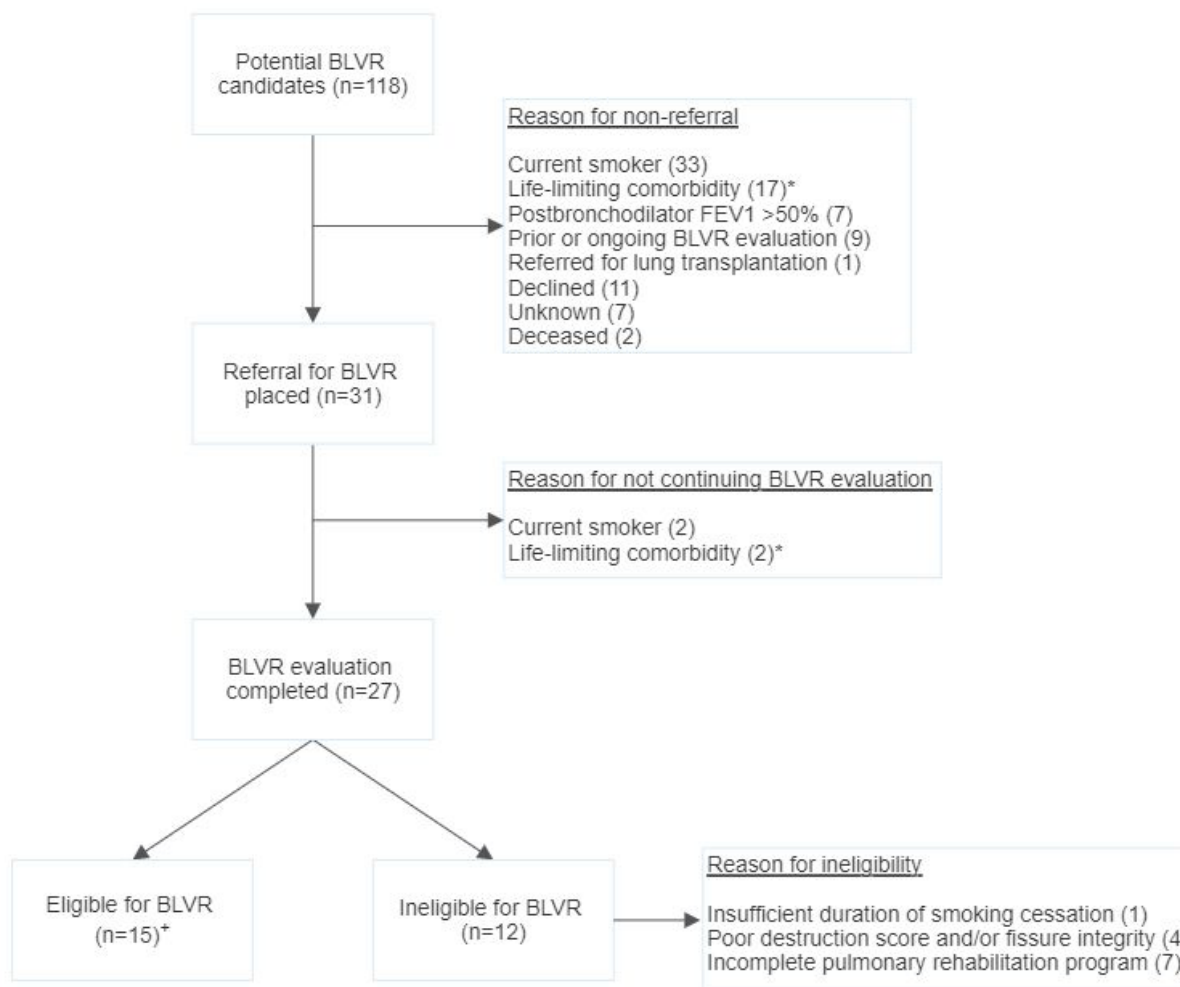
improvement in our referral rates.

A potential drawback of our implementation approach was the inclusion of prebronchodilator, instead of postbronchodilator, FEV₁ values <50%. This led to inaccurately identifying a small proportion with significant postbronchodilator improvement as potential BLVR candidates. It is also worth noting that our cohort was predominantly older males, limiting the generalizability of our results. Finally, although an increase in referral volumes was anticipated, this required close coordination with our referring providers and nurse navigator to ensure timely evaluation. It is foreseeable that with a growing advanced lung failure program, leveraging the support of nurse navigators, coordinators, and advanced practice providers will be instrumental to the program's success.¹⁴

Conclusion

Implementing a restructured PFT template in conjunction with provider education was effective in improving case identification and referral rates for BLVR. Future quality improvement efforts should explore larger-scale VA and non-VA medical center implementation, with careful attention to institutional preferences and cultural norms.

Figure 2. A Flowchart of the Identification, Review, and Referral Process for Bronchoscopic Lung Volume Reduction for the Post-Intervention Cohort



*A life-limiting comorbidity included patients with advanced cardiovascular or other comorbid disease, extrapulmonary end-organ failure, and active or suspected malignancy

+Eligible patients should have met all inclusion criteria including completion of a supervised pulmonary rehabilitation program within the past year (minimum of 10 sessions), abstinence from tobacco products for at least 3 months, and adequate emphysema destruction scores and fissure integrity based on quantitative chest computed tomography.

BLVR= bronchoscopic lung volume reduction; FEV₁=forced expiratory volume in 1 second

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Declaration of Interest

CD has a consulting agreement with Pulmonx. AMM has received payment from LivaNova, Inc., for participation on a data safety monitoring board. The remaining authors have no relevant conflicts of interest to disclose.

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