Impact of Chronic Obstructive Pulmonary Disease on Atrial Fibrillation Ablation Outcomes According to the National Readmission Database

AHMED M. MARAEY, MD,1,2 MUHAMMAD HAISUM MAQSOOD, MD,3 MAHMOUD KHALIL, MD, MSc,3,4 AHMED HASHIM, MBChB,5 AHMED M. ELZANATY, MD, MPH,6 HADEER R. ELSHARNOBY, MBChB,7 EMAN ELSHEIKH, MBChB,4 LAMIAA ELBATANONY, MBChB, MSc,4 KENNETH ONG, MD,8 and PAUL CHACKO, MD6

1Department of Internal Medicine, CHI St. Alexius Health, Bismarck, ND, USA
2Department of Internal Medicine, University of North Dakota, Bismarck, ND, USA
3Department of Internal Medicine, Lincoln Medical Center, Bronx, NY, USA
4Department of Cardiology, Tanta University Faculty of Medicine, Tanta, Egypt
5Ain Shams University, Cairo, Egypt
6Department of Cardiovascular Medicine, University of Toledo, Toledo, OH, USA
7Tanta University Faculty of Medicine, Tanta, Egypt
8Department of Cardiovascular Medicine, Lincoln Medical Center, Bronx, NY, USA

ABSTRACT. Chronic obstructive pulmonary disease (COPD) is a risk factor for the development of atrial fibrillation (AF). There is a paucity of contemporary data studying the association between COPD and outcomes of AF ablation. The objective of this study was to investigate the impact of COPD on AF ablation outcomes using a large nationwide database. This study was a retrospective analysis of the National Readmission Database for the years 2016–2018 and included patients admitted with a diagnosis of AF who underwent catheter ablation. Admissions were stratified according to COPD diagnosis using International Classification of Diseases, 10th Revision, Clinical Modification codes. Multivariate, linear, Cox, and logistic regressions were performed to study the impact of COPD on AF ablation. A total of 18,224 admissions (mean age, 68 years; standard deviation, 10 years) were included, of whom 3,494 (19%) had a diagnosis of COPD. The COPD group was older (72 ± 8 vs. 67 ± 11 years, P < .001) and more likely to have congestive heart failure (73% vs. 44%, P < .001) and renal failure (31% vs. 17%, P < .001). COPD was associated with an increased risk of readmission (adjusted hazard ratio [aHR], 1.40; 95% confidence interval [CI], 1.26–1.56; P < .001) and all-cause in-hospital mortality (adjusted odds ratio, 2.83; 95% CI, 1.74–4.60; P < .001). However, COPD was not associated with an increased risk of readmission due to recurrent AF (aHR, 0.97; 95% CI, 0.75–1.27; P = .844) or the need for re-ablation (aHR, 0.85; 95% CI, 0.44–1.65; P = .639), respectively. In conclusion, COPD was not associated with an increased risk of recurrent AF after ablation despite higher peri-procedural risks. The present study contributes to a better understanding of this high-risk subgroup of patients undergoing AF ablation.

KEYWORDS. Atrial fibrillation, catheter ablation, COPD, mortality, outcome, readmission.
Introduction

Chronic obstructive pulmonary disease (COPD) is a disease characterized by airflow limitations due to small airway disease (obstructive bronchiolitis) and parenchymal destruction (emphysema), leading to expiratory flow limitation and hyperinflation. COPD continues to pose a growing contribution to the overall prevalence of non-communicable diseases. It was estimated that the global prevalence of COPD was 11.7% by 2010. In addition, COPD is usually associated with other comorbidities, especially atrial fibrillation (AF), the most common sustained arrhythmia affecting 33.5 million people globally. COPD is present in about 23% of patients diagnosed with AF. AF development is directly associated with the frequency of COPD exacerbation. The co-presence of both COPD and AF represents an extra challenge that requires tailoring any medical interventions to minimize any potential collateral damage.

In this study, we have investigated the potential effect of COPD on the outcomes of AF ablation.

Methods

Data source

This was a retrospective cohort study performed using the Agency for Healthcare Research and Quality’s Healthcare Cost and Utilization Project (HCUP) Nationwide Readmission Database (NRD) for the years 2016–2018. The NRD is the largest publicly available all-payer inpatient health care readmission database in the United States. Data in the NRD are drawn from HCUP state inpatient databases containing verified patient linkage numbers that can be used to track a person across hospitals within a state while adhering to strict privacy guidelines. It provides longitudinal information about patients’ initial hospitalization and subsequent readmissions in a calendar year. It contains a weighted sample of hospitalizations in the United States, and this can be used to directly derive national estimates of various hospitalizations. National estimates were produced using sampling weights provided by the sponsor. All values presented are weighted estimates.

Study population

Our study population included admissions with a diagnosis of AF who underwent catheter ablation. To define the admissions with AF who underwent catheter ablation, we used the International Classification of Diseases, 10th Revision (ICD-10), Clinical Modification, codes for AF who underwent catheter ablation (ICD-10 Procedure Coding System codes [02573ZZ, 025S3ZZ, 025T3ZZ, 02583ZZ]).

The authors report no conflicts of interest for the published content. No funding information was provided. The main findings of this study were presented in abstract format at the 2022 American College of Cardiology meeting.

Manuscript received February 8, 2022. Final version accepted February 22, 2022.

Address correspondence to: Ahmed Maraeey, MD, 900 E. Broadway Ave, Bismarck, ND, 58501, USA. Email: maraeeyahmed@gmail.com.

(Supplementary Table 1). Admissions with a diagnosis of other cardiac arrhythmias for which an ablation procedure may be required were excluded. Arrhythmias excluded were supraventricular tachycardia, ventricular tachycardia, atrial flutter, ventricular premature beats, pre-excitation syndrome, atrioventricular (AV) nodal tachycardia, and atrial tachycardia. Also, to avoid inclusion of patients undergoing AV junction ablation, we excluded admissions with codes indicating prior or current implantation of a pacemaker or an implantable cardioverter-defibrillator. This approach for the identification of admissions with AF and catheter ablation has been previously described and used by other investigators. In NRD, patient identifiers cannot be linked across years. Hence, index admissions on July 1 or beyond were excluded to allow 6 months of follow-up. Other exclusion criteria were admissions with age <18 years and those with missing data.

In accordance with the HCUP data use agreement, we excluded reporting any variable containing a small number of observations (≤10) that could pose a risk of person identification or data privacy violation. This study was exempted from approval by the institutional review board because of the anonymized and de-identified data in the NRD.

Study outcomes

Primary outcomes were 180-day readmission with recurrent AF, AF re-ablation, and all-cause readmissions after index admissions for AF ablation.

Secondary outcomes were in-hospital mortality, iatrogenic cardiac complication, postprocedural bleeding, stroke, transient ischemic attack (TIA), acute myocardial infarction, cardiac tamponade, acute kidney injury (AKI), and acute heart failure. Index admission was defined as the first admission with the primary diagnosis of AF without prior admission in a 180-day period. A readmission was defined as the first readmission within 180 days of admission.

Statistical analysis

Continuous variables were compared using Student’s t test, and categorical variables were compared using the chi-squared test. Multivariate regression analyses were used to study the association between COPD and AF ablation outcomes and to adjust for confounders: age; sex; median income; primary insurance; hospital bed size; hospital teaching status; and chronic comorbidities such as hypertension, diabetes mellitus, congestive heart failure, valvular heart disease, peripheral vascular disease, obesity, liver disease, history of myocardial infarction, history of percutaneous coronary intervention, and history of coronary artery bypass graft. Logistic regression was used for dichotomous in-hospital outcomes, and linear regression was used for continuous outcomes. Cox proportional hazards regression analysis was used in readmission outcomes. In-hospital mortality admissions were excluded from readmission analysis. Continuous data were presented as mean and standard deviation (SD) values.
All $P$ values were 2-sided, with 0.05 as the threshold for statistical significance. Data analysis was performed using STATA 17 (StataCorp, College Station, TX, USA).

**Results**

**Baseline characteristics**

Our cohort included 18,224 AF ablation admissions (mean age, 68 ± 10 years) between January and June of 2016–2018, of whom 3,494 (19%) admissions had a diagnosis of COPD. The mean age was different between the COPD group and the non-COPD group (72 ± 8 and 67 ± 11 years, $P < .001$, respectively). COPD admissions were likely to be women (46% vs. 43%, $P = .02$). Medicare was the primary insurance in the cohort; however, COPD admissions were more likely to have Medicare insurance (81% vs. 60%, $P < .001$). AF ablation was less likely to be done in a teaching hospital among the COPD group (80% vs. 95%, $P < .001$).

COPD admissions were more likely to have congestive heart failure (73% vs. 44%, $P < .001$), peripheral vascular disorder (21% vs. 12%, $P < .001$), renal failure (31% vs. 17%, $P < .001$), chronic respiratory failure (18% vs. 1%, $P < .001$), and hypothyroidism (20% vs. 16%, $P < .001$). Other baseline characteristics stratified according to the presence or absence of COPD are included in Table 1.

**Outcomes**

A total of 5,209 readmissions within 180 days occurred after AF ablation, of whom 1,505 (29%) carried a diagnosis of COPD. COPD was associated with an increased risk of readmission after adjustment (adjusted hazard ratio (aHR), 1.40; 95% confidence interval [CI], 1.26–1.56; $P < .001$).

Supplementary Table 1: International Classification of Diseases, Tenth Edition, Clinical Modification (ICD-10-CM) Codes Used to Identify Key Variables

| Atrial Fibrillation          | I48, I48.0, I48.1, I48.11, I48.19, I48.2, I48.20, I48.21, I48.91 |
| Catheter ablation            | 02573ZZ, 02553ZZ, 02573ZZ, 02583ZZ |
| Chronic obstructive pulmonary disease | J44, J44.0, J44.1, J44.9, J41, J41.0, J41.1, J41.8, J42, J43.0, J43.1, J43.2, J43.8, J43.9 |
| Premature ventricular complexes | I49.3 |
| Supraventricular tachycardia  | I47.1 |
| Ventricular premature depolarization | I49.3 |
| Pre-excitation syndrome       | I45.6 |
| Presence of pacemaker         | 295.0 |
| Paroxysmal tachycardia        | I47.9 |
| Atrial flutter                | I489.2, I48.3, I48.4 |
| Postprocedural hemorrhage     | I97.610, I97.618, I97.620, I97.638, J95.830, K91.841, K91.871 |
| Iatrogenic cardiac complications | I97.790, I97.88, I97.89 |
| Hemopericardium               | I31.2 |
| Cardiac tamponade             | I31.4 |
| Acute myocardial infarction   | I21.4, I21.9, I21.3, I21.09, I21.19, I21.02, I21.21, I21.01, I21.11, I21.29 |
| Uncomplicated hypertension    | I10.x |
| Complicated hypertension      | I11.x-I13.x, I15.x |
| Valvular heart disease        | A52.0, I05.x - I08.x, I09.1, I09.8, I34.x - I39.x, Q23.0 - Q23.3, Z95.2 - Z95.4 |
| Pulmonary circulation disorders | I26.x, I27.x, I28.0, I28.8, I28.9 |
| Peripheral vascular disorders | I70.x, I71.x, I72.x, I73.1, I73.8, I73.9, I77.1, I77.7, I79.0, I79.1, I79.8, I79.2, K55.1, K55.8, K55.9, Z95.8, Z95.9 |
| Uncomplicated diabetes mellitus | E08.9, E09.9, E10.9, E11.9, E13.9 |
| Complicated diabetes mellitus | E08.2-E08.8, E09.x, E10.2 - E10.8, E11.2 - E11.8, E12.2 - E12.8, E13.2 - E13.8 |
| Hypothyroidism                | E00.x - E03.x |
| Obesity                       | E66.x, Z68.3, Z68.4, Z68.5 |
| Liver disease                 | B18.x, I85.x, K70.x, K71.1, K71.3 - K71.5, K71.7, K72.x - K74.x, K75.4, K75.8, K76.0, K76.2 - K76.9, Z94.4 |
| Obesity                       | E66.x, Z68.3, Z68.4, Z68.5 |
| Deficiency anemia             | D501, D50.8, D50.9, D51.x - D53.x, D63.1, D63.8 |
| Alcohol abuse                 | F10, E52, G62.1, I42.6, K29.2, K70.0, K70.3, K70.9, T51.x, Z71.4 |
| Drug abuse                    | F11.x - F16.x, F18.x, F19.x, Z71.5 |
| Prior myocardial infarction   | I25.2 |
| Prior percutaneous coronary intervention | Z98.61, Z95.5 |
| Prior coronary artery bypass grafting | Z95.1 |
| Dyslipidemia                  | E78.5 |
an increased risk of readmission due to recurrent AF (aHR, 0.97; 95% CI, 0.75–1.27; \( P = .844 \)) or the need for re-ablation (aHR, 0.85; 95% CI, 0.44–1.65; \( P = .639 \)) (Figure 1).

In-hospital mortality occurred in 168 (1%) admissions. COPD was associated with an increased in-hospital mortality. The adjusted odds ratio (aOR) was 2.83 (95% CI, 1.74–4.60; \( P < .001 \)).

In terms of periprocedural complications, COPD was not associated with an increased odds of postprocedural hemorrhage (aOR, 0.62; 95% CI, 0.36–1.06; \( P = .083 \)).

### Table 1: Baseline Characteristics Stratified According to the Presence or Absence of COPD

| Characteristics                                    | AF Ablation                  | \( P \) Value |
|----------------------------------------------------|------------------------------|---------------|
| No COPD (n = 14,523)                                | COPD (n = 3,432)             |               |
| Female sex                                         | 6,318 (43%)                  | 1,881 (46%)   | .018          |
| Mean age in years (SD)                             | 72 (±8)                      | 67 (±11)      | <.001         |
| Median household income quartile for zip code per percentile |                              |               |
| <25<sup>th</sup>                                   | 3,089 (21%)                  | 944 (28%)     | <.001         |
| 25<sup>th</sup>–50<sup>th</sup>                     | 3,687 (25%)                  | 987 (29%)     |              |
| 50<sup>th</sup>–75<sup>th</sup>                     | 3,735 (26%)                  | 877 (26%)     |              |
| >75<sup>th</sup>                                   | 4,011 (28%)                  | 623 (18%)     |              |
| Insurance                                          |                              |               |
| Medicare                                           | 8,854 (60%)                  | 2,815 (81%)   |              |
| Medicaid                                           | 623 (4%)                     | 187 (5%)      |              |
| Private                                            | 4,804 (33%)                  | 408 (12%)     |              |
| Self-pay                                           | 109 (1%)                     | 22 (1%)       |              |
| Other/no charge                                    | 334 (2%)                     | 60 (2%)       |              |
| Hospital bed size                                  |                              |               |
| Small                                              | 899 (6%)                     | 263 (8%)      | .196         |
| Medium                                             | 3,956 (27%)                  | 972 (25%)     |              |
| Large                                              | 9,875 (67%)                  | 2,259 (65%)   |              |
| Teaching hospital                                  | 12,493 (85%)                 | 2,802 (80%)   | <.001        |
| Charlson Comorbidity Index score                   |                              |               |
| \( \leq 1 \)                                       | 8,926 (60%)                  | 445 (13%)     |              |
| 2                                                  | 2,304 (16%)                  | 850 (24%)     |              |
| \( \geq 3 \)                                       | 3,500 (24%)                  | 2,198 (63%)   |              |
| Comorbidities                                      |                              |               |
| Congestive heart failure                           | 6,492 (44%)                  | 2,563 (73%)   | <.001        |
| Valvular heart disease                             | 3,257 (22%)                  | 950 (27%)     | <.001        |
| Peripheral vascular disease                        | 1,803 (12%)                  | 748 (21%)     | <.001        |
| Pulmonary circulatory disorder                     | 1,110 (8%)                   | 590 (17%)     | <.001        |
| Uncomplicated hypertension                         | 6,617 (45%)                  | 1,147 (33%)   | <.001        |
| Complicated hypertension                           | 4,570 (31%)                  | 1,916 (55%)   | <.001        |
| History of myocardial infarction                   | 1,052 (7%)                   | 462 (13%)     | <.001        |
| History of PCI                                     | 1,327 (9%)                   | 503 (14%)     | <.001        |
| History of CABG                                     | 927 (6%)                     | 370 (11%)     | <.001        |
| History of stroke                                  | 1,109 (8%)                   | 327 (9%)      | .014         |
| Renal failure                                      | 2,571 (17%)                  | 1,081 (31%)   | <.001        |
| Obesity                                            | 3,484 (24%)                  | 990 (28%)     | <.001        |
| Hypothyroidism                                     | 2,338 (16%)                  | 707 (20%)     | <.001        |
| Dyslipidemia                                       | 6,553 (44%)                  | 1,750 (50%)   | <.001        |
| Diabetes without complications                     | 2,289 (16%)                  | 610 (17%)     | .045         |
| Diabetes with complications                        | 1,456 (10%)                  | 682 (20%)     | <.001        |
| Liver disease                                      | 349 (2%)                     | 156 (4%)      | <.001        |
| Alcohol abuse                                      | 327 (2%)                     | 167 (5%)      | <.001        |
| Drug abuse                                         | 132 (1%)                     | 67 (2%)       | <.001        |

**Abbreviations:** AF, atrial fibrillation; CABG, coronary artery bypass graft; COPD, chronic obstructive pulmonary disease; PCI, percutaneous coronary intervention; SD, standard deviation.
Figure 1: Kaplan–Meier estimates comparing time to first all-cause readmission and first atrial fibrillation readmission in the chronic obstructive pulmonary disorder (COPD) group versus non-COPD group. Abbreviations: AF, atrial fibrillation; COPD, chronic obstructive pulmonary disease.

Figure 2: Forest plot of main study outcomes. Abbreviation: TIA, transient ischemic attack.

Discussion

This nationally representative analysis of 18,224 AF admissions who underwent ablation, using NRD data from the years 2016–2018, found no difference in 180-day readmission due to AF or re-ablation despite the higher risk of all-cause readmissions and some periprocedural complications, such as acute heart failure, in-hospital mortality, and AKI, in the COPD group compared to the non-COPD group. Other periprocedural complications, such as acute myocardial infarction, iatrogenic cardiac complications, postprocedural hemorrhage, cardiac tamponade and hemopericardium, and stroke/TIA, were not different between the 2 groups.

It is well known that COPD has been associated with an increased incidence of AF, 11–13 which is attributed to increased arrhythmogenicity from hypoxia, 14 hypercapnia, 15 pulmonary hypertension, 16 ventricular
### Table 2: Study Outcomes

| Outcome                                    | No COPD (n = 14,523) | COPD (n = 3,432) | P Value |
|--------------------------------------------|----------------------|------------------|---------|
| In-hospital mortality (n)                  | 88 (1%)              | 80 (2%)          | <.001   |
| Adjusted odds ratio                        | Reference            | 2.83 (95% CI, 1.74–4.60) | <.001   |
| Acute kidney injury (n)                    | 1,711 (12%)          | 809 (23%)        | <.001   |
| Adjusted odds ratio                        | Reference            | 1.24 (95% CI, 1.03–1.49) | .022    |
| Acute myocardial infarction                | 250 (2%)             | 99 (3%)          | .002    |
| Adjusted odds ratio                        | Reference            | 1.14 (95% CI, 0.8–1.6) | .47     |
| Iatrogenic cardiac complication            | 164 (1%)             | 36 (1%)          | .78     |
| Adjusted odds ratio                        | Reference            | 0.97 (95% CI, 0.54–1.76) | .93     |
| Postprocedural hemorrhage                  | 230 (2%)             | 34 (1%)          | .062    |
| Adjusted odds ratio                        | Reference            | 0.62 (95% CI, 0.36–1.06) | .083    |
| Cardiac tamponade and hemopericardium      | 219 (1%)             | 28 (1%)          | .022    |
| Adjusted odds ratio                        | Reference            | 0.66 (95% CI, 0.38–1.13) | .131    |
| Stroke or TIA                              | 147 (1%)             | 47 (1%)          | .204    |
| Adjusted odds ratio                        | Reference            | 1.27 (95% CI, 0.77–2.11) | .35     |
| Acute heart failure                        | 2,735 (19%)          | 1,466 (42%)      | <.001   |
| Adjusted odds ratio                        | Reference            | 1.45 (95% CI, 1.24–1.70) | <.001   |
| 180-day all-cause readmission             | 3,700 (25%)          | 1,505 (44%)      | <.001   |
| Adjusted hazard ratio                      | Reference            | 1.40 (95% CI, 1.26–1.56) | <.001   |
| Readmission due to atrial fibrillation     | 627 (4%)             | 119 (3%)         | .12     |
| Adjusted hazard ratio                      | Reference            | 0.97 (95% CI, 0.75–1.27) | .844    |
| Need for repeat ablation                   | 131 (1%)             | 20 (1%)          | .18     |
| Adjusted hazard ratio                      | Reference            | 0.85 (95% CI, 0.44–1.65) | .639    |

**Abbreviations:** AF, atrial fibrillation; CI, confidence interval; COPD, chronic obstructive pulmonary disease; NRD, Nationwide Readmission Database.

![Figure 3: Study summary. Abbreviations: AF, atrial fibrillation; COPD, chronic obstructive pulmonary disease; NRD, Nationwide Readmission Database.](image-url)
remodeling,17 inflammation,18 and respiratory drug use.19 At a biochemical level, the forementioned stresses are hypothesized to induce the production of hypoxia-inducible factor 1α, vascular endothelial growth factors, and matrix metalloproteinase-9 through a cascade of reactions that ultimately leads to atrial fibrosis.14,20–22

In 2011, a retrospective study on AF ablation from South Korea from 2000–2009 sparked a huge interest in AF ablation in chronic lung disease patients.23 They did not only find out that chronic lung disease changes the pulmonary vein anatomy and makes it more arrhythmogenic but also reported that non-pulmonary vein foci-related AF was more common in chronic lung disease patients. In 2015, a review by Goudis on the relationship between COPD and AF highlighted the need for studies on AF ablation outcomes in COPD patients.24 Similarly, the abovementioned gap was reinforced in recent reviews on AF, COPD, and pulmonary hypertension.25,26

So, this nationally representative cohort of American adult patients is the first large study to evaluate the outcomes of AF ablation in patients with COPD. This is also the first study of its kind, which evaluated 180-day readmission and other procedure-related complications of AF ablation.

A national cohort analysis of years 2010–2014, which evaluated causes and predictors of readmission (30-day and 90-day readmission) post-ablation, found that patients with chronic pulmonary disease have a higher risk of readmission and COPD is an independent predictor of readmission.10 The results of that analysis align with our analysis, which showed a significantly higher risk of readmission in COPD patients.

AF has been challenging to control in patients with COPD. Non-COPD status was one of the independent predictors of successful electrical cardioversion at 1 year of follow-up in the Euro Heart Survey (P = .003).27 A prospective study by Gu et al. on 550 consecutive patients with symptomatic and medication-refractory AF with first catheter ablation found a significantly greater recurrence of atrial tachyarrhythmias in patients with COPD versus patients without COPD (44% vs. 28.6%, P = .016) with a mean follow-up of 31.4 months.28 In contrast, our analysis with a shorter follow-up duration (6 months) did not find differences between the 2 arms (COPD vs. non-COPD) in readmission with AF or the need for repeat ablation.

Our study demonstrated that post-ablation in-hospital mortality in patients was higher in the COPD group (2% vs. 1%, P < .001). Our results are comparable to those of a multicenter analysis of AF patients from 13 hospitals in China from 2008–2011. The study found that the prevalence of COPD was significantly higher in the mortality group than the survival group (21.4% vs. 10.1%, P ≤ .01).26 In the present study, we observed a higher risk of acute congestive heart failure and AKI in the COPD group.

New-onset congestive heart failure was reported following AF catheter ablation by Tan et al.30 Congestive heart failure was found to be an independent predictor of worsening renal function in a prospective study done by Kawai et al. on 791 non-dialysis patients who underwent AF ablation.31 Due to the lack of patient-level data in the administrative database, such as the degree of ejection fraction reduction and the level of kidney impairment, we could not further investigate the increased risk of acute heart failure and AKI in COPD patients. We hypothesize that the difference was likely due to the higher prevalence of chronic heart failure and chronic kidney disease in the COPD arm and the strong correlation between COPD and congestive heart failure.

COPD has been associated with a greater risk of arteriovascular events, including ischemic stroke, cerebrovascular events, TIs, and myocardial infarctions.32–34 Similarly, COPD patients are more likely to have peripheral vascular diseases and atherosclerosis.35 The present study did not find a difference in outcomes of in-hospital occurrence of acute myocardial infarction and ischemic stroke/TIA between the 2 groups segregated based on COPD status. However, our study evaluated these outcomes if they happened during the index admission. It would be interesting to see future studies on post-ablation vascular complications in the 2 groups.

Limitations

This study should be interpreted in light of some important limitations. First, it is an observational study subject to confounding bias despite statistical adjustment. Second, the severity of COPD and other comorbidities, anti-arrhythmic medication use, and medication compliance could not be assessed due to the administrative nature of the database, which is also subject to coding errors. Third, a real-world all-payer, nationally representative NRD cohort could not include all admissions in the United States, so it might over- or underestimate the overall difference in outcomes. Additionally, AF recurrence without admission was not captured. Fourth, the follow-up period was relatively short as admissions can be missed. Longer-term follow-up will be an area for future research.

Conclusion

Despite being a risk factor for the development of AF, COPD was not associated with an increased risk of AF recurrence after the ablation procedure. However, it was associated with increased AF ablation periprocedural complications. The present study contributes to a better understanding of this high-risk subgroup of patients undergoing AF ablation. The development of a nationally valid risk stratification model and improving post-hospital care have the potential to offset some of the risks inherent to ablating AF in COPD patients.
Impact of COPD on AF Ablation Outcomes

References

1. Dugani S, Gaziano TA. 25 by 25: achieving global reduction in cardiovascular mortality. Curr Cardiol Rep. 2016;18(1):10.
2. Adeloye D, Chua S, Lee C, et al. Global and regional estimates of COPD prevalence: systematic review and meta-analysis. J Glob Health. 2015;5(2):020415.
3. Negewo NA, McDonald VM, Gibson PG. Comorbidity in chronic obstructive pulmonary disease. Respir Investig. 2015;53(6):249–258.
4. Chugh SS, Havmoeller R, Narayan K, et al. Worldwide epidemiology of atrial fibrillation: a Global Burden of Disease 2010 Study. Circulation. 2014;129(8):837–847.
5. Buist AS, McBurnie MA, Vollmer WM, et al. International variation in the prevalence of COPD (the BOLD Study): a population-based prevalence study. Lancet. 2007;370(9589):741–750.
6. Proietti M, Laroche C, Drozd M, et al. Impact of chronic obstructive pulmonary disease on prognosis in atrial fibrillation: a report from the EURObservational Research Programme Pilot Survey on Atrial Fibrillation (EORP-AF) General Registry. Am Heart J. 2016;181:83–91.
7. Rodriguez-Manero M, Lopez-Pardo E, Cordero A, et al. A prospective study of the clinical outcomes and prognosis associated with comorbid COPD in the atrial fibrillation population. Int J Chron Obstruct Pulmon Dis. 2019;14:371–380.
8. Grymonprez M, Vakaet V, Kavousi M, et al. Chronic obstructive pulmonary disease and the development of atrial fibrillation. Int J Cardiol. 2019;276:118–124.
9. HCUP Nationwide Readmission Database (N RD), Healthcare Cost and Utilization Project (HCUP). Rockville, MD. Agency for Healthcare Research and Quality. Available at: https://www.hcupus.ahrq.gov/nrdoverview.jsp. Accessed February 7, 2022.
10. Arora S, Lahewala S, Tripathi B, et al. Causes and predictors of readmission in patients with atrial fibrillation undergoing catheter ablation: a national population-based cohort study. J Am Heart Assoc. 2018;7(12):e009294.
11. Johnson LS, Juhlin T, Engström G, Nilsson PM. Reduced forced expiratory volume is associated with increased incidence of atrial fibrillation: the Malmo Preventive Project. Europace. 2014;16(2):182–188.
12. Li J, Agarwal SK, Alonso A, et al. Airflow obstruction, lung function, and incidence of atrial fibrillation: the Atherosclerosis Risk in Communities (ARIC) study. Circulation. 2014;129(9):971–980.
13. Konecny T, Park JY, Somers KR, et al. Relation of chronic obstructive pulmonary disease to atrial and ventricular arrhythmias. Am J Cardiol. 2014;114(2):272–277.
14. Xu Y, Sharma D, Du E, Liu Y. The role of Toll-like receptor 2 and hypoxia-induced transcription factor-1α in the atrial structural remodeling of non-valvular atrial fibrillation. Int J Cardiol. 2013;168(3):2940–2941.
15. Terzano C, Romani S, Conti V, Paone G, Oriolo F, Vitarelli A. Atrial fibrillation in the acute, hypercapnic exacerbations of COPD. Eur Respir Rev. 2014;23(139):2908–2917.
16. Wanamaker B, Cassino T, McLaughlin V, Oral H, Latchamsetty R, Siontis KC. Atrial arrhythmias in pulmonary hypertension: pathogenesis, prognosis and management. Arrhythm Electrophysiol Rev. 2018;7(1):43–48.
17. Eweda I, Hamada G. Concordance between Doppler and pulsed-wave Doppler tissue imaging in estimation of the degree of left ventricular dysfunction and correlating it to the degree of chronic obstructive pulmonary disease. J Saudi Heart Assoc. 2016;28(1):15–21.
18. Harada M, Van Wagoner DR, Nattel S. Role of inflammation in atrial fibrillation pathophysiology and management. Circ J. 2015;79(3):495–502.
19. Wood-Baker R, Cochrane B, Naughton MT. Cardiovascular mortality and morbidity in chronic obstructive pulmonary disease: the impact of bronchodilator treatment. Intern Med. 2010;49(2):94–101.
20. Lammers WJ, Kirchhof C, Bonke FI, Allessie MA. Vulnerability of rabbit atrium to reentry by hypoxia. Role of inhomogeneity in conduction and wavelength. Am J Physiol. 1992;262(1 Pt 2):H47–H55.
21. Su F, Zhang W, Chen Y, Ma L, Zhang H, Wang F. Significance of hypoxia-inducible factor-1α expression with atrial fibrosis in rats induced with isoproterenol. Exp Ther Med. 2014;8(6):1677–1682.
22. Ogi H, Nakano Y, Niida S, et al. Is structural remodeling of fibrillated atria the consequence of tissue hypoxia? Circ J. 2010;74(9):1815–1821.
23. Roh SY, Choi JJ, Lee JY, et al. Catheter ablation of atrial fibrillation in patients with chronic lung disease. Circ Arrhythm Electrophysiol. 2011;4(6):815–822.
24. Goudis CA. Chronic obstructive pulmonary disease and atrial fibrillation: an unknown relationship. J Cardiol. 2017;69(5):699–705.
25. Valhadtpour CA, Luebert JJ, Palevsky HJ. Atrial arrhythmias in chronic lung disease-associated pulmonary hypertension. Palm Circ. 2020;10(1):1–13.
26. Simons SO, Elliott A, Sastry M, et al. Chronic obstructive pulmonary disease and atrial fibrillation: an interdisciplinary perspective. Eur Heart J. 2021;42(5):532–540.
27. Pisters R, Nieuwlaat R, Frins MH, et al. Clinical correlates of immediate success and outcome at 1-year follow-up of real-world cardioversion of atrial fibrillation: the Euro Heart Survey. Europace. 2012;14(5):666–674.
28. Gu J, Liu X, Tan H, et al. Impact of chronic obstructive pulmonary disease on procedural outcomes and quality of life in patients with atrial fibrillation undergoing catheter ablation. J Cardiovasc Electrophysiol. 2013;24(2):148–154.
29. Wang J, Yang YM, Zhu J, et al. [Analysis of risk factors for all cause-mortality in Chinese emergency atrial fibrillation patients]. Zhonghua Yi Xue Za Zhi. 2013;93(36):2871–2875.
30. Tan HW, Wang XH, Shi HF, et al. Congestive heart failure after extensive catheter ablation for atrial fibrillation: prevalence, characterization, and outcome. J Cardiovasc Electrophysiol. 2011;22(6):632–637.
31. Kawai T, Shizuta S, Aizawa T, et al. Renal function and outcomes in atrial fibrillation patients after catheter ablation. PLoS One. 2020;15(11):e0241449.
32. Nadeem R, Sharief A, Tanna S, Sidhu H, Molnar J, Nadeem A. Potential augmentation of the risk of ischemic cerebrovascular accident by chronic obstructive pulmonary disease in patients with atrial fibrillation. J Stroke Cerebrovasc Dis. 2015;24(8):1893–1896.
33. Donaldson GC, Hurst JR, Smith CJ, Hubbard RB, Wedzicha JA. Increased risk of myocardial infarction and stroke following exacerbation of COPD. Chest. 2013;143(5):1091–1097.
34. Fearn J, Rodrigues LC, Smith CJ, Hubbard RB, Gibson JE. Prevalence of major comorbidities in subjects with COPD and incidence of myocardial infarction and stroke: a comprehensive analysis using data from primary care. Thorax. 2010;65(11):956–962.
35. Angeli F, Reboldi G, Trapasso M, Aita A, Ambrosio G, Verdecchia P. Detrimental impact of chronic obstructive pulmonary disease in atrial fibrillation: new insights from umbria atrial fibrillation registry. Medicina (Kaunas). 2019;55(7):358.