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Meta-Analysis of the Usefulness of Catheter Ablation of Atrial Fibrillation in Patients with Heart Failure with Preserved Ejection Fraction

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Conflicts of Interest:

Dr. Hsu reports receiving honoraria from Medtronic, Abbott, Boston Scientific, Biotronik, Zoll Medical, Biosense-Webster, Janssen Pharmaceuticals, and Bristol-Myers Squibb, research grants from Biotronik and Biosense-Webster, and has equity interest in Acutus Medical and Vektor Medical.

Dr Krummen reports owning equity in Vektor Medical Inc for unrelated work. Dr. Ho reports receiving a research grant from the American Heart Association (AHA 19CDA34760021), National Institutes of Health (NIH 1KL2TR001444), owns equity in Vektor Medical Inc and fellowship support from Medtronic, Abbott, Boston Scientific, and Biotronik.

Dr. Feld reports receiving consulting fees from Acutus Medical, Inc., Vektor Medical, Inc., and Altathera Pharmaceuticals, Inc., fellowship stipend support from Medtronic, Inc., Boston Scientific, Inc., Biotronik, Inc., Biosense Webster, Inc., and St. Jude Medical, Inc., is co-founder of Perminova, Inc., and has received stock or stock-options from Acutus Medical, Inc., and Perminova, Inc.

ABSTRACT

Catheter ablation improves clinical outcomes in atrial fibrillation (AF) patients with heart failure (HF) with reduced ejection fraction (HFrEF). However, the role of catheter ablation in HF with a preserved ejection fraction (HFpEF) is less clear. We performed a literature search and systematic review of studies that compared AF recurrence at one year after catheter ablation of AF in patients with HFpEF versus those with HFrEF. Risk ratio (RR; where a RR<1.0 favors the HFpEF group) and mean difference (MD; where MD<0 favors the HFpEF group) 95% confidence intervals were measured for dichotomous and continuous variables, respectively. Six studies with a total of 1,505 patients were included, of which 764 (51%) had HFpEF and 741 (49%) had HFrEF. Patients with HFpEF experienced similar recurrence of AF one year after ablation on or off antiarrhythmic drugs compared to those with HFrEF (RR 1.01; 95% CI 0.76, 1.35). Fluoroscopy time was significantly shorter in the HFpEF group (MD -5.42; 95% CI -8.51, -2.34), but there was no significant difference in procedure time (MD 1.74; 95% CI -11.89, 15.37) or periprocedural adverse events between groups (RR 0.84; 95% CI 0.54,1.32). There was no significant difference in hospitalizations between groups (MD 1.18; 95% CI 0.90, 1.55), but HFpEF patients experienced significantly less mortality (MD 0.41; 95% CI 0.18, 0.94). In conclusion, based on the results of this meta-analysis, catheter ablation of AF in patients with HFpEF appears as safe and efficacious in maintaining sinus rhythm as in those with HFrEF.

Key words: catheter ablation, atrial fibrillation, heart failure, diastolic dysfunction, hospitalizations, mortality

INTRODUCTION

Atrial fibrillation (AF) and heart failure (HF) have become increasingly prevalent and frequently co-occur, resulting in increased morbidity and mortality relative to either disease alone.¹⁻⁵ Although there are evidence-based guidelines for both diseases,^{6,7} it is less clear how to manage patients in whom both AF and HF are present. While more data has emerged to guide the management of patients with AF and heart failure with reduced ejection fraction (HFrEF), including updated guidelines advocating for ablation of AF in patients with HFrEF, data on patients with heart failure with preserved ejection fraction (HFpEF) were relatively sparse until recently.^{7,8} Given that AF has been shown to incur greater morbidity and mortality in patients with HFpEF relative to those with AF and HFrEF and in those with AF and no HF, understanding how to best manage AF in patients with HFpEF is of particular importance.⁹⁻¹⁴ The purpose of our current study was to perform a systematic review of the literature and meta-analysis to determine the effect of catheter ablation of AF on rates of recurrent AF, fluoroscopy and procedure times, and rates of periprocedural adverse events, hospitalizations and mortality in patients with HFpEF compared to those with HFrEF, in order to determine any difference in benefits or risks between these groups.

METHODS

We searched PubMed, Clinicaltrials.gov, Medline, Google scholar and the Cochrane Central Register of Clinical Trials (Cochrane Library, Issue 09, 2017). This was assessed up to March 2020. No language restriction was applied. The reference list of all eligible studies was also reviewed. Search terms included (*Catheter Ablation*) AND (*Atrial Fibrillation*) AND (*Heart Failure with Preserved Ejection Fraction* OR *Diastolic Dysfunction*).

Studies were selected by two independent reviewers. The PRISMA statement for reporting systemic reviews and meta-analyses was applied to the methods for this study.¹⁵ The studies had to fulfill the following criteria to be considered in the analysis: 1) Studies were required to evaluate outcomes in patients with HFpEF and HFrEF (HF with a moderately reduced ejection fraction (EF 40-49%) was included as part of the HFrEF group); 2) Studies were required to report the rates of recurrent AF; 3) Studies were required to have a minimum follow up of 12 months; 4) Studies were required to have been published in a peer-reviewed scientific journal.

We aimed to compare rates of recurrent AF, fluoroscopy and procedure times, and rates of periprocedural adverse events, hospitalizations and mortality in patients with HFpEF compared to those with HFrEF, from baseline procedure to follow up.

Two authors (O.M.A. and F.L.) independently performed the literature search and extracted data from eligible studies. Outcomes

were extracted from original manuscripts and supplementary data. Information was gathered using a standardized protocol and reporting forms. Disagreements were resolved by consensus. Two reviewers (O.M.A. and F.L.) independently assessed the quality items and discrepancies were resolved by consensus or involvement of a third reviewer (J.C.H), if necessary.

Two authors (O.M.A. and F.L.) independently assessed the risk of bias of the included trials using standard criteria defined in the Cochrane Handbook for Systematic Reviews of Interventions. Discrepancies were resolved by discussion or adjudication by a third author (J.C.H.).

Data were summarized across treatment arms using the Mantel-Haenszel risk ratio (RR), where a RR < 1.0 favored the HFpEF group, and inverse variance mean difference (MD), where a MD < 0 favored the HFpEF group. Heterogeneity of effects was evaluated using the Higgins I-squared (I²) statistic. Random effects models for analyses were used with high heterogeneity (defined as $I^2 > 25\%$), otherwise fixed effects models of DerSimonian and Laird were used. Funnel plot analyses were used to address publication bias. The statistical analysis was performed by the Review Manager (RevMan). Version 5.3. Cochrane Copenhagen: The Nordic Centre, The Cochrane Collaboration, 2014. Descriptive statistics are presented as means and

standard deviations (SD) for continuous variables or number of cases (n) and percentages (%) for dichotomous and categorical variables.

RESULTS

An initial search resulted in 325 abstracts, of which 93 were duplicates and 214 were excluded based on titles and abstracts (Figure 1). We included six studies in our final analysis; three retrospective¹⁶⁻¹⁸ prospective¹⁹⁻²¹ observational studies. and three Baseline demographics and characteristics of the six studies are summarized in Table 1 and 2. All of the included studies were observational. The majority of HFpEF and HFrEF patients in all of the included studies had persistent AF and were on antiarrhythmic drugs and beta-blockers, with the exception that only 37% of HFrEF patients in the study by Eietl et al. were on antiarrhythmic drugs. We included a total of 1,505 patients. Among these, 764 (51%) patients had HFpEF and 741 (49%) had HFrEF. The risk of bias is summarized in Table 3. While all the studies accounted for major comorbidities when making comparisons and had adequate follow-up, the majority of studies did not control for antiarrhythmic drug use. The majority of the studies used a 3-month blanking period.^{16-18,21}

There was no difference in the risk of recurrent AF in patients with HFpEF and HFrEF (RR 1.01; 95% CI 0.76, 1.35) one year after ablation (Figure 2). Although fluoroscopy time was significantly shorter

in the HFpEF group (MD -5.42; 95% CI -8.51, -2.34), there was no significant difference in procedure times (MD 1.74; 95% CI -11.89, 15.37) or peri-procedural adverse events between groups (RR 0.84; 95% CI 0.54,1.32) (Figure 3). Additionally, while there was no significant difference in hospitalizations between groups (MD 1.18; 95% CI 0.90, 1.55), HFpEF patients experienced significantly less mortality (MD 0.41; 95% CI 0.18, 0.94) (Figure 4). Funnel plot analysis of the included studies showed no evidence of publication bias on any of the reported outcomes (Figures 2-4). Furthermore, in a sensitivity analysis where only prospective studies were included, the results were similar.

DISCUSSION

This is the first systematic review and meta-analysis of studies comparing outcomes during and after catheter ablation of AF in patients with HFpEF versus those with HFrEF. The results of this metaanalysis show that there are no significant differences in rates of recurrence of AF one year after catheter ablation between patients with HFpEF and HFrEF. Fluoroscopy time was significantly shorter in the HFpEF group, but there were no significant differences in procedure time or periprocedural adverse events between groups. While there was no significant difference in hospitalizations, HFpEF patients had significantly less mortality over follow-up. These findings should encourage larger, randomized control trials to be done specifically in patients with HFpEF, to establish a benefit of catheter ablation in this population of patients.

In addition to the findings reported in this meta-analysis, there are data that suggest ablation of AF in patients with HFpEF portends improvements in quality of life, short-term hospitalizations and longterm AF recurrence. Although Black-Meier et al.¹⁶ found no significant difference in quality of life scores pre- and post-ablation, both Cha et al. and Ichijo et al. reported significant improvements in quality of life in patients with HFpEF post-ablation.^{17,19} Elkaryoni et al. found that there was no significant difference in the relative reduction in hospitalization rates 120 days before and after index admission for catheter ablation among HFpEF patients (28.5%) and those with HFrEF (25.2%).²² Fukui et al. also showed that catheter ablation significantly reduced HF hospitalizations over a mean follow-up of 720 ±377 days in patients with HFpEF when compared to conventional pharmacotherapy.²³ Similar to the data presented here, Jayanna et al. found, in a subgroup analysis, that there was no difference in AF recurrence 3 months and 1 year post-ablation between patients with HFpEF and HFrEF, but this data was not included in the analyses as the numbers of patients with HFpEF and HFrEF were not explicitly stated.²⁴ Much of the AF recurrence data presented at one year remained to be true over longer follow-up, with two other studies that had extended

follow-up out to 5 years showing no difference in recurrence between patients with HFpEF and HFrEF.¹⁹ These data suggest that patients with AF and HFpEF do just as well, if not better, than those with AF and HFrEF after catheter ablation, which should encourage larger studies to evaluate this patient population, especially since they are largely lacking from current guidelines.⁷

Atrial remodeling, AF and HFpEF share similar risk factors, which in part explains the increased prevalence of one disease in the presence of the other ²⁵⁻²⁸ AF can lead to HFpEF predominantly through hemodynamic effects and left ventricular fibrosis. The loss of atrial systole, loss of atrio-ventricular synchrony and decreased filling time seen in AF decreases cardiac output and results in a series of neurohormonal changes. The excess sympathetic tone and renin activity result in an increase in central venous pressure and the rise in plasma norepinephrine and subsequent arteriolar vasoconstriction increases the afterload.^{29,30} Additionally, there are data to suggest that the burden of AF is important in the development of fibrosis.³¹ Conversely, HFpEF can also promote AF through several mechanisms including atrial fibrosis,^{32,33} left atrial mechanical dysfunction.^{9,34} changes in calcium handling,^{35,36} and electrical remodeling.³⁷ The shared risk factors and pathophysiology may account for the increased burden and mortality of AF in the HFpEF population relative to those with HFrEF.^{9,38} Due to the pathophysiology of AF in patients with

HFpEF, the importance of our meta-analysis findings in aggregate show that catheter ablation of AF in patients with HFpEF is important and may be as effective at improving clinical outcomes as has been shown in patients with HFrEF.

The current systematic review and meta-analysis has several important limitations that should be acknowledged. First, all of the included studies were observational and, with the exception of the study by Eitel et al., were single-center, which significantly restricts the generalizability of the results. Second, there were different study protocols, with both retrospective and prospective studies included and various lesion sets employed. Third, each study had different protocols to monitor for arrhythmia recurrence, but all met the standard of consensus guidelines.^{39,40} Fourth, follow-up was only analyzed out to 12 months post-ablation. However, as mentioned above, reported results were similar in the studies that extended follow-up out for several years. In conclusion, based on the results of this meta-analysis, catheter ablation of AF in patients with HFpEF appears as safe and efficacious in maintaining sinus rhythm as in those with HFrEF.

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Figure 1. Selection of studies

Figure 2. Forrest plots and funnel plots for the comparative analysis of risk of recurrent atrial fibrillation in patients with heart failure with preserved ejection fraction compared to those who have heart failure with reduced ejection fraction at one year after catheter ablation.

Figure 3: Forrest plots and funnel plots for the comparative analysis of A) fluoroscopy time, B) procedure time, and C) peri-procedural adverse events in patients with heart failure with preserved ejection fraction compared to those with heart failure with reduced ejection fraction. Peri-procedural adverse events varied by study, including access site/vascular complications, cardiac perforation/tamponade, stroke/transient ischemic attack, pericarditis, acute heart failure, pulmonary vein stenosis, phrenic nerve injury, esophageal atrial fistula, air embolism and prolonged hospitalization.

Figure 4: Forrest plots and funnel plots for the comparative analysis of A) hospitalizations and B) mortality in patients with heart failure with preserved ejection fraction compared to those who have heart failure with reduced ejection fraction.