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Association of Exercise-Based Cardiac Rehabilitation with Progression of Paroxysmal to Sustained Atrial Fibrillation

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Abstract: Progression of atrial fibrillation (AF) is associated with worsened prognosis for cardiovascular events and mortality. Exercise-based-cardiac rehabilitation programmes have shown preliminary promise for primary and secondary prevention of AF. Yet, such interventions are typically reserved for patients with acute coronary syndrome or undergoing revascularization. Using a retrospective cohort design, the present study investigated the association of exercise-based cardiac rehabilitation on the progression of paroxysmal to sustained AF, compared to propensity-matched controls. Patients with a diagnosis of paroxysmal AF were compared between those with and without an electronic medical record of exercise-based cardiac rehabilitation within 6-months of diagnosis. Using cox regression models, we ascertained odds of 2-year incidence for AF progression. This cohort of 9808 patients with paroxysmal AF demonstrated that exercise-based cardiac rehabilitation was associated with 26% lower odds of AF progression (odds ratio 0.74, 95% CI 0.66–0.83) compared to propensity-matched controls. This beneficial effect seemed to vary across patient subgroups. In conclusion, findings revealed that exercise-based cardiac rehabilitation was associated with significantly lower odds of progression from paroxysmal to sustained AF at 2-years follow-up compared to propensity-matched controls.

Keywords: atrial fibrillation; cardiac rehabilitation; secondary prevention; disease progression; rehabilitation

1. Introduction

Atrial fibrillation (AF) can present as one of three main clinical subtypes: paroxysmal AF (PAF; episodes of arrhythmia that terminate spontaneously), persistent AF (episodes that continue for >7 days and are not self-terminating), and permanent AF (ongoing long-term episodes). AF typically progresses from PAF to sustained AF (SAF; persistent or permanent AF), with a worsened prognosis for death, heart failure, and stroke during the peri-progression period [1]. Although excessive levels of vigorous exercise may be associated with increased incident AF, guideline levels of physical activity and exercise training have been shown to elicit potent AF-specific protection [2]. Indeed, 1500 metabolic equivalents (METs)-mins/week were associated with reduced risk of incident AF in men and even exceeding...
these recommendations (5000 MET-mins/week) were AF-protective in females [3]. Further, exercise-based cardiac rehabilitation (CR), in association with optimized drug therapy, can improve sinus rhythm maintenance in patients with SAF [4]. However, little is known about the impact of CR and exercise programmes on disease progression in those with PAF. Therefore, we investigated the association between exercise-based CR and the progression of PAF to SAF, compared to matched PAF patients without exercise-based CR.

2. Materials and Methods

This retrospective observational study was conducted on October 29, 2020 with anonymized data provided by TriNetX, a global federated health research network with access to electronic medical records (EMR) from participating academic medical centres, specialty physician practices, and community hospitals, predominantly in the United States. AF was identified from International Classification of Diseases, Tenth Revision, Clinical Modification (ICD-10-CM) codes in patient EMRs: I48.0 (PAF), I48.1 and I48.21 (SAF). Restrictions on the search dates were used so that all included patients had an EMR of PAF recorded at least 2 years ago (to allow for follow-up). Exercise-based CR was identified from ICD-10-CM codes Z71.82 (Exercise counselling), Healthcare Common Procedure Coding System (HCPCS) code S9472 (CR program, non-physician provider, per diem), or Current Procedural Terminology (CPT) code 1013171 (Physician or other qualified health care professional services for outpatient CR). Correspondingly, these exercise-based CR codes were excluded in the matched controls. The exercise-based CR cohort were aged ≥18 years with exercise-based CR recorded in EMRs within 6 months of PAF diagnosis. Controls were aged ≥18 years with PAF and no EMR history of exercise-based CR. At the time of the search, 39 participating healthcare organizations had data available for patients who met the study inclusion criteria.

Baseline characteristics were compared using chi-squared tests or independent-sample t-tests. Current CR provision is typically reserved for cardiovascular patients following an acute coronary syndrome, those undergoing a revascularization procedure (coronary artery bypass graft or planned percutaneous coronary intervention), and patients with heart failure. Thus, propensity score matching was used to control for these differences in the two cohorts. Using logistic regression, exercise-based CR patients were 1:1 propensity score-matched with controls for age at PAF diagnosis, sex, race, hypertensive diseases, ischaemic heart diseases, heart failure, cerebrovascular diseases, diabetes mellitus, chronic kidney disease, cardiovascular procedures (including cardiography, echocardiography, catheterization, cardiac devices, and electrophysiological procedures), and cardiovascular medications (including beta-blockers, antiarrhythmics, diuretics, antilipemic agents, antianginals, calcium channel blockers, and ACE inhibitors). Following propensity score matching, logistic regression produced odds ratios (OR) with 95% confidence intervals (CIs) for 2-year incidence of SAF (persistent/permanent AF), comparing exercise-based CR with non-CR controls. Additional sub-analyses (following propensity score matching) were conducted to produce ORs with 95% CIs to explore the effect of population subgroups (age, sex, and comorbidities) on the odds of AF progression between the exercise-based CR cohort and controls. Statistical significance was set at \( p < 0.05 \).

3. Results

In total, 379,966 patients with PAF from 39 US healthcare organizations met the inclusion criteria for the control group and 4905 patients with PAF met the inclusion criteria for the exercise-based CR cohort. Compared to controls, the exercise-based CR cohort were younger, had a lower proportion of females, had a higher proportion of White and Black or African American, and had a higher proportion of comorbidities (Table 1). Following propensity score matching, there were 4904 patients in each cohort (\( n = 9808 \) in total), which were overall, well balanced for age, race, sex, included comorbidities, cardiovascular procedures, and cardiovascular medications (Table 1).
Table 1. Baseline characteristics % (n) * of the AF populations with and without CR and exercise before and after propensity score matching.

|                      | Initial Populations | Propensity Score Matched Populations |  p-Value |  p-Value |
|----------------------|---------------------|--------------------------------------|----------|----------|
|                      | AF without CR       | AF with CR                           |          |          |
|                      | (n = 379,966)       | (n = 4905)                           |          |          |
| Age (years) at diagnosis; mean (SD) | 70.6 (12.5)         | 69.6 (11.2)                          | <0.001   |          |
| Female               | 45.4 (172,581)      | 32.4 (1587)                          | <0.001   |          |
| Race                 |                     |                                      |          |          |
|                      | White               | 84.4 (320,653)                       | 87.1 (4272) | <0.001  |          |
|                      | Black or African American | 7.4 (28,052) | 8.5 (416) | 0.004 |          |
|                      | Asian               | 1.3 (4937)                           | 1.5 (72) | 0.303 |          |
|                      | Unknown             | 6.8 (25,731)                         | 2.8 (138) | <0.001  |          |
| Hypertensive diseases | 46.8 (177,861)      | 83.4 (4090)                          | <0.001   |          |
|                      | Ischaemic heart diseases | 23.9 (91,001) | 77.4 (3795) | <0.001  |          |
|                      | Cerebrovascular diseases | 10.1 (38,352) | 24.1 (1180) | <0.001  |          |
|                      | Diabetes Mellitus | 18 (68,336)                          | 37.3 (1830) | <0.001  |          |
|                      | Chronic Kidney Disease | 12.2 (46,533) | 30.5 (1497) | <0.001  |          |
|                      | Cardiovascular Procedures b | 38.7 (147,131) | 89.1 (4372) | <0.001  |          |
|                      | Catheter ablation  | 0.5 (1959)                           | 1.6 (79) | <0.001 |          |
|                      | Cardiovascular Medications c | 57 (216,644) | 93.5 (4588) | <0.001  |          |
|                      | Most common antiarrhythmic drugs | | | | |
|                      | Lidocaine          | 18.7 (70,923)                       | 64.3 (3153) | <0.001 |          |
|                      | Amiodarone         | 8.4 (32,044)                        | 45.4 (2226) | <0.001  |          |
|                      | Adenosine          | 1.0 (3755)                          | 8.1 (396) | <0.001 |          |
|                      | Dofetilide         | 1.0 (3860)                          | <0.1 (95) | <0.001 |          |

* Values are % (n) unless otherwise stated. Baseline characteristics were compared using a chi-squared test for categorical variables and an independent-sample t-test for continuous variables. a Data are taken from structured fields in the electronic medical record systems of the participating healthcare organizations, therefore, there may be regional or country-specific differences in how race categories are defined. b Cardiovascular procedures include cardiography, echocardiography, catheterization, cardiac devices, electrophysiological procedures. c Cardiovascular medications include beta-blockers, antiarrhythmics, diuretics, antilipemic agents, antianginals, calcium channel blockers, Angiotensin-converting-enzyme (ACE) inhibitors. AF; atrial fibrillation, CR; cardiac rehabilitation and exercise programmes, SD; standard deviation.
Using the propensity score matched cohort, progression from PAF to SAF at 2-year follow-up from PAF diagnosis was proportionally lower with 19.3% \( (n = 617 \text{ of } 3197 \text{ patients}) \) in the exercise-based CR cohort compared to 24.5% \( (n = 909 \text{ of } 3716 \text{ patients}) \) in the matched controls \( (\text{OR} 0.74, 95\% \text{ CI: } 0.66–0.83) \).

Subgroup analyses comparing exercise-based CR with controls demonstrated lower odds of SAF progression in patients aged <75 years \( (\text{OR} 0.66, 95\% \text{ CI: } 0.56–0.78) \); male \( (\text{OR} 0.71, 95\% \text{ CI: } 0.62–0.83) \); patients without chronic kidney disease \( (\text{OR} 0.66, 95\% \text{ CI: } 0.56–0.79) \); patients with heart failure \( (\text{OR} 0.81, 95\% \text{ CI: } 0.71–0.93) \) and patients without heart failure \( (\text{OR} 0.72, 95\% \text{ CI: } 0.56–0.93) \); and patients without a history of stroke \( (\text{OR} 0.73, 95\% \text{ CI: } 0.64–0.83) \). Patients aged \( \geq 75 \text{ years} \) \( (\text{OR} 0.92, 95\% \text{ CI: } 0.77–1.09) \); female \( (0.97, 95\% \text{ CI: } 0.81–1.13) \); and with a history of stroke \( (\text{OR} 0.97, 95\% \text{ CI: } 0.73–1.29) \) did not demonstrate a significant association with reduced odds of SAF progression.

4. Limitations

A number of limitations are noteworthy. First, the data were collected from health care organization EMR databases and some co-morbidities may be underreported. Indeed, recording of ICD codes in administrative datasets may vary by factors such as age, number of comorbidities, severity of illness, length of hospitalization, and whether in-hospital death occurred. In particular, an EMR of exercise-based CR does not provide information as to whether a patient attended, the intervention type and dose, or intervention adherence. We could also not determine the influence of attending different healthcare organizations due to data privacy restrictions. Further, the current AF classification scheme supported by the American Heart Association (AHA), American College of Cardiology (ACC), and the European Society of Cardiology (ESC) relies on duration of rhythm and spontaneous conversion with patients considered to have paroxysmal, persistent, or permanent AF. However, the accuracy and reliability of this classification scheme, especially in EMRs, is unknown. Second, the data were from multiple healthcare organizations in the United States, which may not be representative of the wider population, thus the generalizability of the results beyond this cohort is unclear. Third, residual confounding may have impacted our results, including lifestyle factors and socioeconomic status, which were not available from EMRs. Fourth, it was not possible to factor for multiple comparisons in the subgroup analyses within the online database. Finally, subsequent randomized controlled trials are needed to further investigate the causal impact of exercise-based CR on AF-specific outcomes.

5. Conclusions

The present study of 9808 patients with PAF demonstrates that exercise-based CR was associated with 26% lower odds of progression to SAF at 2-years, compared to propensity score matched controls. Since progression from PAF to SAF is associated with a further increased risk in mortality and morbidity, our observation of delayed progression has clinical value. Despite the promise of exercise as a primary and secondary therapy for AF, CR is typically not prescribed for patients with AF. Our findings provide support for the inclusion of patients with AF in exercise-based CR programmes for AF-specific protection.

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Conflicts of Interest: Benjamin J. R. Buckley has received funding from Pfizer. Stephanie L. Harrison has received funding from Bristol-Myers Squibb (BMS). Elnara Fazio-Eynullayeva and Paula Underhill are employees of TriNetX LLC. Deirdre A. Lane has received investigator-initiated educational grants from Bristol-Myers Squibb (BMS), has been a speaker for Boehringer Ingelheim, and BMS/Pfizer and has consulted for BMS, Boehringer Ingelheim, and Daiichi-Sankyo. Gregory Y. H. Lip: consultant for Bayer/Janssen, BMS/Pfizer, Medtronic, Boehringer Ingelheim, Novartis, Verseon and Daiichi-Sankyo and speaker for Bayer, BMS/Pfizer, Medtronic, Boehringer Ingelheim, and Daiichi-Sankyo. No fees are directly received personally.

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