

Safety and Efficacy of Switching Anticoagulation to Aspirin Three Months after Successful Radiofrequency Catheter Ablation of Atrial Fibrillation

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Received: August 6, 2013

Revised: October 23, 2013

Accepted: November 22, 2013

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The authors have no financial conflicts of interest.

Purpose: Although current guidelines recommend continuing the same antithrombotic strategy regardless of rhythm control after radiofrequency catheter ablation (RFCA) of atrial fibrillation (AF), anticoagulation has a risk of major bleeding. We evaluated the safety of switching warfarin to aspirin in patients with successful AF ablation. **Materials and Methods:** Among 721 patients who underwent RFCA of AF, 608 patients (age, 57.3±10.9 years; 77.0% male, 75.5% paroxysmal AF) who had no evidence of AF recurrence at 3 months post-RFCA were included. We compared the thromboembolic and hemorrhagic events in patients for whom warfarin was switched to aspirin (ASA group; n=296) and patients who were kept on warfarin therapy (W group; n=312). **Results:** There were no significant differences in CHA₂DS₂-VASc or HAS-BLED scores between the groups. In 30 patients in the ASA group and 37 patients in W group, AF recurred and warfarin was restarted or maintained during the 18.0±12.2 months of follow-up. There were no significant differences in thromboembolic (0.3% vs. 1.0%, *p*=0.342) and major bleeding incidences (0.7% vs. 0.6%, *p*=0.958) between ASA and W groups during the follow-up period. In the 259 patients with a CHA₂DS₂-VASc score ≥2, there were no significant differences in thromboembolism (0.8% and 2.2%, *p*=0.380) or major bleeding incidences (0.8% and 1.4%, *p*=0.640) between ASA and W groups. **Conclusion:** Switching warfarin to aspirin 3 months after successful RFCA of AF could be as safe and efficacious as long-term anticoagulation even in patients with CHA₂DS₂-VASc score ≥2. However, strict rhythm monitoring cannot be overemphasized.

Key Words: Anticoagulation, aspirin, atrial fibrillation, catheter ablation

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INTRODUCTION

Atrial fibrillation (AF) is significantly associated with the risk of strokes,^{1,2} and given this association, antithrombotic therapy is essential in the management of AF. CHADS₂ or CHA₂DS₂-VASc scores have been utilized for risk stratification of ischemic stroke in patients with AF,^{3,4} and anticoagulation is recommended for

the patients with CHADS₂ score ≥ 1 or CHA₂DS₂-VASc score ≥ 1 .^{5,6} Radiofrequency catheter ablation (RFCA) has been known to be as effective as or more effective than antiarrhythmic drugs (AAD) at controlling rhythm in AF patients.⁷⁻⁹ According to the 2012 HRS/EHRA/ECAS Expert Consensus Statement, AAD-resistant symptomatic paroxysmal AF (PAF) and persistent AF (PeAF) are class I and IIa indications for RFCA, respectively.¹⁰ However, the same guidelines recommend that anticoagulation be continued after RFCA of AF based on CHADS₂ or CHA₂DS₂-VASc score.¹⁰ Although warfarin is effective for the prevention of stroke in patients with AF, using warfarin is difficult given the various dose-effect relationships as well as potential drug-drug or drug-food interactions. Moreover, warfarin introduces an annual risk of major bleeding of 3.1–3.7%.¹¹⁻¹³ Of the various bleeding risk stratification schemas, the HAS-BLED score is recommended by the current guidelines for prediction of bleeding risk in patients who take warfarin due to AF,^{5,14} and special attention should be paid to patients with high HAS-BLED score. Given the risks and benefits of warfarin, controversies remain regarding long-term use of warfarin after successful RFCA of AF. Therefore, we hypothesized that switching warfarin to aspirin (ASA group) may have an equivalent efficacy and safety in the prevention of ischemic stroke, compared with continuous warfarin administration (W group) in patients with no evidence of AF recurrence after RFCA during standard rhythm monitoring.

MATERIALS AND METHODS

Study population

The study protocol was approved by the Institutional Review Board of Severance Cardiovascular Hospital, Yonsei University Health System and Asan Medical Center, and adhered to the Declaration of Helsinki. This study included 721 patients who were prospectively registered in a database of AF catheter ablation between March 2009 and December 2011 in two centers. Clinical outcomes, thromboembolic events, and hemorrhagic complications were compared between the ASA group and the W group in the registry database. Among these 721 patients, there were 608 patients (age, 57.3 \pm 10.9 years; 77.0%, male; 75.5%, PAF; 24.5%, PeAF) who had no evidence of AF recurrence on Holter monitoring 3 months after RFCA in this study. All patients provided written informed consent, and all were AAD-resistant or intolerant patients. The exclusion criteria were as

follows: 1) permanent AF refractory to electrical cardioversion; 2) left atrium (LA) size >55 mm as measured by echocardiogram; 3) critical coronary artery stenosis ($>75\%$ of luminal diameter); 4) severe rheumatic mitral valvular disease; and 5) prior RFCA of AF. Three-dimensional (3D) spiral computerized tomography (CT) scans (64 Channel, Light Speed Volume CT, Philips, Brilliance 63, the Netherlands) were performed to visually define pulmonary vein (PV) anatomy. The presence of an LA thrombus was excluded by transesophageal echocardiography. All AADs were discontinued for a period corresponding to at least five half-lives. In total, 141 patients (19.6%) were taking amiodarone, which was discontinued for at least four weeks prior to the procedure. Anticoagulation therapy was maintained before RFCA for all patients.

Electrophysiologic mapping and radiofrequency catheter ablation

Intracardiac electrograms were recorded using the Prucka CardioLabTM electrophysiology system (General Electric Health Care System Inc., Milwaukee, WI, USA) or Workmate (St. Jude Medical Inc., Minnetonka, MN, USA). RFCA was performed in all patients using 3D electroanatomical mapping (NavX; St. Jude Medical Inc., Minnetonka, MN, USA or Carto; Johnson & Johnson Inc., Diamond Bar, CA, USA) merged with 3D spiral CT. We used an open irrigated-tip catheter (Celsius or Thermocool, Johnson & Johnson Inc.; Diamond Bar, CA, USA; irrigation flow rate 20 to 30 mL/min; 25–35 W; 47°C) to deliver radiofrequency energy for ablation (Stockert generator, Biosense Webster Inc.; Diamond Bar, CA, USA). All patients initially underwent circumferential PV isolation (CPVI) and bi-directional block of the cavotricuspid isthmus. For the patients with PAF, we added linear ablations and complex fractionated atrial electrogram (CFAE) ablation guided by 3D-CFAE-CL map¹⁵ in a stepwise approach in patients without AF termination during RFCA. For PeAF, we conducted CPVI, cavotricuspid isthmus block, roof line, posterior inferior line, and anterior line¹⁶ as a routine lesion set, and CFAE ablation was added based on the operator's decision. If AF persisted beyond the aforementioned ablation protocols for PAF or PeAF, we stopped the procedure after internal cardioversion. The endpoint of our procedure was no immediate recurrence of AF after cardioversion with isoproterenol infusion (5–20 μ g/min). If there were non-PV foci under isoproterenol, we ablated them all. During RFCA, activated clotting time was maintained between 350 to 400 seconds.

Post-RFCA follow-up

All patients were followed up at the outpatient clinic after 1, 3, 6, 9, and 12 months post-RFCA and then every 6 months. Electrocardiography (ECG) was performed at every visit and whenever the patients complained of palpitation. A Holter monitoring (24- or 48-hour) and/or an event recorder were performed at 3, 6, 12, 18, and 24 months after RFCA and whenever patients complained of palpitation following the 2012 HRS/EHRA/ECAS Expert Consensus Statement guidelines.¹⁰ We defined recurrence of AF as any episode of AF or atrial tachycardia of at least 30 sec in duration.¹⁷ If any ECG-documented AF episode occurred within the three-month blanking period during follow-up, patients were then diagnosed as an early recurrence, and any AF recurrence thereafter was diagnosed as clinical recurrence.¹⁷ In case of early recurrence and clinical recurrence, AADs were prescribed.

Comparisons of ASA switching and continuous warfarin during follow-up

A total of 608 patients without evidence of AF recurrence on ECG and Holter monitoring at 3 months after RFCA were included in this study. We compared the thromboembolic events and major bleeding complications in 296 patients for whom warfarin was changed to 100 mg of aspirin (ASA group) and 312 patients for whom anticoagulation was maintained (W group) after confirming no recurrence in the 3rd-month post-RFCA Holter monitoring (Fig. 1). The primary endpoints were thromboembolism including ischemic stroke/transient ischemic attack (TIA) and major bleeding after 3 months from RFCA. Stroke was defined as symptomatic ischemic cerebral infarction with apparent brain lesion in imaging studies. TIA was defined as a transient episode of neurologic dysfunction confirmed by a neurologist without brain lesion in imaging studies, with spontaneous symptomatic recovery within 24 hours. Major bleeding events were defined as any type of hemorrhage re-

quiring blood transfusion or intervention, and bleeding with reduction of hemoglobin levels by ≥ 4.0 g/dL.¹⁸ We compared incidences, hazard ratio and event-free survival of thromboembolic events (including stroke, TIA, and other thromboembolism) and major bleeding events between ASA and W groups depending on CHA₂DS₂-VASc score.

Statistical analyses

The results are expressed as mean \pm standard deviation. The Student's t-test, χ^2 test were used for comparison of incidences of primary endpoints between the groups. Univariate and multivariate Cox regression analyses were used for comparison of hazard ratio (HR) between the groups. The assumption was assessed by log-minus-log-survival function and found that the proportion hazards assumption was reasonable. Parameters of a p value ≤ 0.1 by univariate analysis were included for multivariate analysis. HRs and 95% confidence intervals (CIs) were calculated. The Kaplan-Meier method was used for analyzing event-free survival. A p value < 0.05 was considered significant. The data were analyzed using the Statistical Package for the Social Sciences version 20.0 (IBM Inc., Armonk, NY, USA).

RESULTS

Patient characteristics

Patient demographic and clinical characteristics in the ASA and W groups at baseline were shown (Table 1). There were no significant differences in clinical profiles, LA size, and left ventricular function. CHADS₂, CHA₂DS₂-VASc, and HAS-BLED scores were not significantly different between two groups. However, past history of stroke/TIA was more frequent in the W group (14.5%) than in the ASA group (7.1%, $p=0.006$). The time in the therapeutic range of international normalized ratio (INR) for the W group was 44.2% during the follow-up period. In sixty-seven patients (6.1%), AF re-

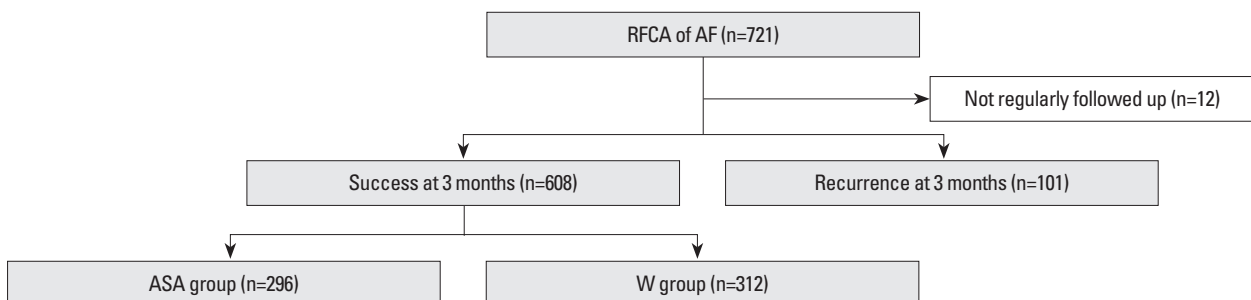


Fig. 1. Flow diagram and numbers of patients. AF, atrial fibrillation; RFCA, radiofrequency catheter ablation.

Table 1. Baseline Characteristics of All Patients

	ASA group (n=296)	W group (n=312)	p value
Age (yrs)	56.9±11.7	57.6±10.1	0.453
Male sex (%)	77.7	76.3	0.677
PAF (%)	75.7	75.3	0.919
CHADS ₂ score	0.91±0.98	1.04±1.04	0.089
CHA ₂ DS ₂ -VASc score	1.45±1.34	1.55±1.36	0.335
HAS-BLED score	1.37±0.83	1.45±1.02	0.268
LA diameter (mm)	41.2±6.0	42.9±7.2	0.002
Ejection fraction (%)	63.1±8.7	61.8±8.6	0.066
Prior aspirin use (%)	21.3	26.6	0.125
Comorbidities (%)			
Hypertension	53.4	53.2	0.966
Diabetes	13.9	16.0	0.453
Stroke/TIA	7.1	14.5	0.006
Chronic kidney disease	1.0	1.3	0.756
Heart failure	4.1	2.6	0.303

LA, left atrium; PAF, paroxysmal atrial fibrillation; TIA, transient ischemic attack.

Table 2. Patients with Primary Endpoints

Group	Age/sex	AF type	CHA ₂ DS ₂ -VASc score	HAS-BLED score	Primary endpoints	Time to event (month)	INR at event
ASA	60/F	Paroxysmal	3	1	Stroke; occipital	7	0.91
ASA	73/M	Paroxysmal	5	4	Pseudoaneurysm rupture	3.1	0.95
ASA	49/M	Paroxysmal	0	1	Epistaxis	3.3	1.12
W	42/M	Paroxysmal	2	1	Stroke; frontal	25	4.82
W	62/M	Persistent	2	1	TIA	6	1.19
W	71/F	Paroxysmal	4	1	TIA	28	2.61
W	62/F	Paroxysmal	3	3	Intramuscular hematoma	4	6.22
W	69/F	Paroxysmal	4	3	Gingival bleeding	4	2.28

AF, atrial fibrillation; INR, international normalized ratio; TIA, transient ischemic attack.

curred after 3 months of post-RFCA. After recurrence of AF, warfarin was restarted according to CHA₂DS₂-VASc score.

Hemorrhagic and thromboembolic risks were not different between ASA and W groups

During the 18.0±12.2 months of follow-up, stroke occurred in a patient in the ASA group (0.3%). Stroke and TIA occurred in one and two patients in the W group, respectively (1.0%, $p=0.342$). All patients who experienced a stroke recovered without sequelae. In all patients with thromboembolic complications, heart rhythm was sinus rhythm at the time of stroke or TIA. Major bleeding events occurred in two patients in ASA and W groups (0.7% and 0.6%, respectively; $p=0.958$). The detailed information about events in each patient was shown (Table 2). There were no thromboembolic or major bleeding complications in patients with AF recurrence.

In univariate and multivariate Cox regression analyses,

CHA₂DS₂-VASc score (HR, 3.80; 95% CI, 1.13–12.71; $p=0.031$) was an independent risk factor for thromboembolic events, and HAS-BLED score (HR, 3.90; 95% CI, 1.13–13.45; $p=0.031$) was an independent risk factor for major bleeding events (Table 3). By Kaplan-Meier analysis, there were no significant differences of event-free survival from the composite primary endpoint (thromboembolic and major bleeding events) between the groups ($p=0.968$) (Fig. 2A).

Comparisons in patients with CHA₂DS₂-VASc score ≥2

Among the 608 included patients, 259 patients (42.6%) had a CHA₂DS₂-VASc score ≥2 (ASA group, 121; W group, 138). Table 4 shows comparison of clinical characteristics of the patients with CHA₂DS₂-VASc score ≥2 in the ASA and W groups. The patients in the W group were younger (62.9±9.0 years vs. 65.5±8.3 years, $p=0.019$) and more likely to have had a history of stroke/TIA (32.6% vs. 17.3%, $p=0.008$) than the patients in the ASA group. In patients with CHA₂DS₂-

Table 3. Clinical Factors Related to Thromboembolic or Major Bleeding Events by Multivariate Cox Regression Analysis

	Thromboembolism		Bleeding	
	HR (95% CI)	<i>p</i> value	HR (95% CI)	<i>p</i> value
Age	0.96 (0.85–1.08)	0.452	0.96 (0.85–1.08)	0.466
Male sex	0.83 (0.10–6.90)	0.865	0.23 (0.03–1.50)	0.123
CHA ₂ DS ₂ -VASc score	3.80 (1.13–12.71)	0.031	1.23 (0.44–3.41)	0.691
HAS-BLED score	0.24 (0.04–1.42)	0.115	3.90 (1.13–13.45)	0.031
Warfarin duration	0.89 (0.65–12.16)	0.931	0.86 (0.16–4.61)	0.863

CI, confidence interval; HR, hazard ratio.

Table 4. Baseline Characteristics of Patients with CHA₂DS₂-VASc Score ≥2

	ASA group (n=121)	W group (n=138)	<i>p</i> value
Age (yrs)	65.5±8.3	62.9±9.0	0.019
Male sex (%)	64.5	60.9	0.551
PAF (%)	72.7	73.2	0.934
CHADS ₂ score	1.69±0.99	1.89±0.92	0.098
CHA ₂ DS ₂ -VASc score	2.78±1.00	2.82±0.98	0.734
HAS-BLED score	1.89±0.83	1.96±1.13	0.601
LA diameter (mm)	42.5±6.1	43.5±6.5	0.216
Ejection fraction (%)	63.1±10.0	61.9±8.9	0.345
Prior aspirin use (%)	23.5	29.7	0.287
Comorbidities (%)			
Hypertension	83.5	76.8	0.182
Diabetes	29.8	33.3	0.536
Stroke/TIA	17.3	32.6	0.008
Chronic kidney disease	2.5	1.4	0.548
Heart failure	9.1	5.1	0.205

LA, left atrium; PAF, paroxysmal atrial fibrillation; TIA, transient ischemic attack.

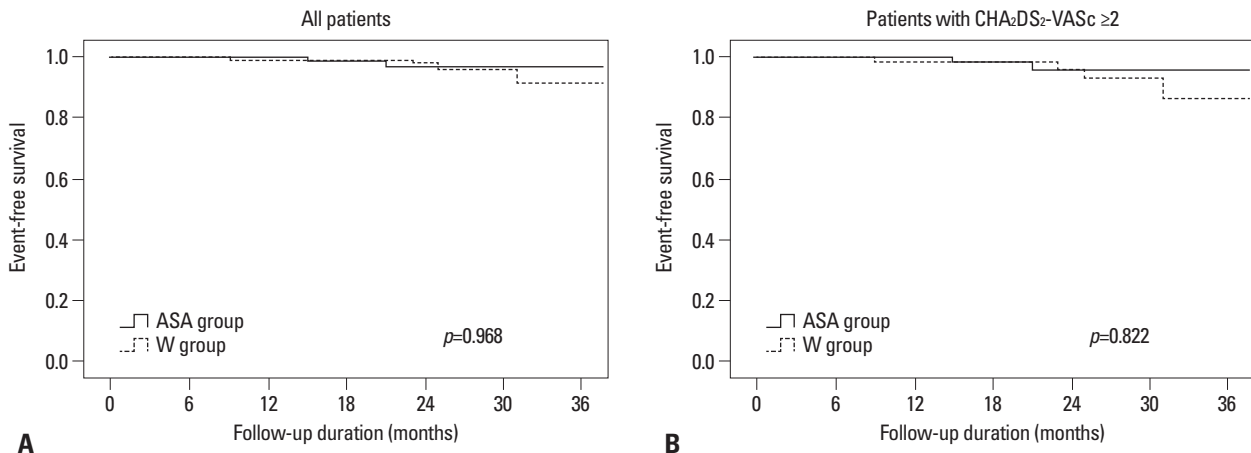


Fig. 2. Thromboembolic and major bleeding event-free survival by Kaplan-Meier method in all patients (A) and the patients with CHA₂DS₂-VASc score ≥2 (B) in ASA (solid line) and W (dotted line) groups.

VASc score ≥2, thromboembolic events occurred in one patient in the ASA group (0.8%) and three patients in the W group (2.2%, *p*=0.380), respectively during follow-up period. Major bleeding events occurred in one and two patients in the ASA (0.8%) and W groups (1.4%, *p*=0.640), respectively. By Kaplan-Meier analysis, there were no significant differences of event-free survival from the composite pri-

mary endpoint in the patients with CHA₂DS₂-VASc score ≥2 between the groups (*p*=0.822) (Fig. 2B).

DISCUSSION

This is a retrospective observational study that evaluated the

risk and benefit of switching anticoagulation to aspirin after successful RFCA of AF, based on CHADS₂, CHA₂DS₂-VASc, HAS-BLED scores, and recommended a standard rhythm monitoring follow-up strategy. The present study suggested that switching warfarin to aspirin 3 months after RFCA could be as safe and efficacious as continuous anticoagulation even in patients with a CHA₂DS₂-VASc score ≥ 2 . However, strict and continuous rhythm monitoring and maintenance of optimal INR should be emphasized to achieve these results. We also proved that CHA₂DS₂-VASc and HAS-BLED scores were very valuable parameters in predicting ischemic stroke and hemorrhagic complications, even after RFCA of AF.

Risk and benefits of anticoagulation in patients with non-valvular AF

Although warfarin can significantly reduce stroke risk by 64% in patients with non-valvular AF,¹⁹ it still has an annual 1.6–2.4% risk of stroke or systemic embolism and an annual 3.1–3.4% risk of major hemorrhagic complications.^{11–13} Moreover, the annual risk of fatal hemorrhagic stroke is between 0.7–0.8%.^{11–13} Therefore, the 2010 ESC guidelines recommended caution, regular review, and correction of reversible bleeding risk factors for patients with a HAS-BLED score ≥ 3 .⁵ In this study, 2 out of 721 patients experienced intracranial and spinal cord hemorrhages during anticoagulation, but they were not included in this analysis because these complications occurred within 3 months after RFCA. Although aspirin has a lower potential risk of major bleeding than warfarin, it reduces ischemic stroke risk by no more than 22% as compared with the 64% risk reduction achieved by anticoagulation.¹⁹ Therefore, aspirin can be considered in patients with a very low AF burden and a high HAS-BLED score. However, a previous study reported that the risk of intracranial hemorrhage was similar in patients taking aspirin to those taking warfarin, and that aspirin increases bleeding risk significantly when administered simultaneously with warfarin.²⁰ In a large cohort study from Denmark including over 132000 patients with AF, warfarin was superior for stroke prevention and similar for bleeding risk compared to aspirin.²¹ Aspirin has rather higher bleeding risk than warfarin in Japan Atrial Fibrillation Stroke Trial (JAST).²² Discrepancies of current study from these two studies were that we included the patients with significantly reduced AF burden by catheter ablation, and those with young and low CHADS₂ score. Generally, the risk of bleeding (HAS-BLED score) increases parallel with the risk of

stroke (CHADS₂ score), such as hypertension, old age, or previous stroke. In spite of ethnical similarity, major bleeding rates were 1.6% in JAST and 0.7% in current study. 2010 EHRA-EAPCI Consensus Documents recommended using warfarin monotherapy in patients with AF and percutaneous coronary intervention who are stable longer than 1 year.²³ However, whether warfarin monotherapy is safe enough to prevent an annual 2.8% risk of very late stent thrombosis in patients with drug-eluting stents²⁴ and major hemorrhagic complications remains unclear.

Anticoagulation issue after rhythm control of AF

The main reason for continuing oral anticoagulation after successful RFCA is the concern for asymptomatic AF recurrence.¹⁰ Previous studies have shown a 0.5 to 7% risk of stroke or thromboembolic events after RFCA of AF,^{25–27} and current guidelines suggest anticoagulation should be maintained even after successful RFCA of AF according to CHADS₂ or CHA₂DS₂-VASc score.¹⁰ However, there have been efforts to stop anticoagulation after potentially successful RFCA of AF because of the bleeding risks and inconvenience of warfarin. There have been several studies demonstrating incidence of thromboembolic events decrease after successful RFCA of AF.^{28–30} Oral, et al.²⁵ showed that, among 755 successfully ablated AF patients, discontinuation of anticoagulation therapy was safe in patients younger than 65 years without a history of stroke. Bunch, et al.³¹ demonstrated the safety of 325 mg aspirin without warfarin in 690 patients with CHADS₂ score 0 to 1, and Saad, et al.³² reported that no thromboembolic morbidity without anticoagulation existed in 327 patients with CHADS₂ score ≤ 3 . However, in a prior nonrandomized study of 3355 patients who underwent CPVI, suspension of oral anticoagulation therapy after successful RFCA was beneficial in patients at moderate to high risk of thromboembolism.³⁰ Therefore the decision to anticoagulate after successful RFCA should be based on assessment of the risk-benefit ratio, and switching anticoagulation to aspirin can be considered in patients with high risk of hemorrhage and no recurrence of AF. There were two reasons that we grouped the patients based on anticoagulation therapy 3 months after RFCA. First, the current guidelines recommend that warfarin should be continued for 2 months after RFCA. Second, post-procedural 3 months are considered as blanking period and not considered as clinical recurrence.³³

However, strict rhythm monitoring cannot be overemphasized. In this study, heart rhythm was sinus rhythm at

the time of stroke or TIA in patients with thromboembolic complications. We might miss asymptomatic AF recurrence or non-cardioembolic source of stroke in these patients. Careful monitoring for rhythm is important in patients with high CHA₂DS₂-VASc or high HAS-BLED scores and patients at high risk of AF recurrence, such as those with high LA volume, PeAF, or long ablation time.³⁴ In this study, we suggest that a regular rhythm monitoring strategy based on the current guidelines was acceptable for making a clinical decision of switching warfarin to aspirin. We also implied that CHA₂DS₂-VASc and HAS-BLED scores were still independent predictors for stroke/TIA and major hemorrhage after successful catheter ablation of AF.

Study limitations

This study was designed as a retrospective observational study, but not as a randomized study. Although CHADS₂ and CHA₂DS₂-VASc scores were not significantly different between two groups, past history of stroke or TIA were higher in W group than in ASA group. It may be because the physicians tended to continue warfarin in patients with a history of stroke or TIA. Although we tried to follow the consensus guidelines for AF rhythm monitoring strictly, all silent recurrence of AF could not be detected on surface ECG or Holter monitoring. The time in therapeutic range was relatively low (44.2%) in the W group compared to previous large scale clinical trials, and it could be one of factors for no significant differences in thromboembolic complication between W group and ASA group. Although recovery of LA contractile function is an important factor for prediction of thromboembolic events, we did not analyze LA contractile function in the current study.

In conclusion, after successful RFCA of AF, switching warfarin to aspirin 3 months after RFCA could be as safe and efficacious as long-term anticoagulation, even in patients with CHA₂DS₂-VASc score ≥ 2 . Strict heart rhythm monitoring in the aspirin group and careful INR monitoring in the anticoagulation group are important in patients after RFCA of AF. Future large randomized trials are warranted.

ACKNOWLEDGEMENTS

This work was supported by a grant (A085136) from the Korea Health 21 R&D Project, Ministry of Health and Welfare, and a grant (2010-0010537) from the Basic Science Research Program of the National Research Foundation of

Korea under the Ministry of Education, Science and Technology of the Republic of Korea.

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