Repeat Catheter Ablation of Long-standing Persistent Atrial Fibrillation in Patients with a Total Atrial Fibrillation Duration of More Than 2 Years: Effects of the CHA2DS2-VASc Score and Estimated Glomerular Filtration Rate on the Outcomes

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Abstract

Objective Little is known about the outcome of repeat catheter ablation of long-standing persistent atrial fibrillation (AF) in patients with a total AF duration of more than 2 years. The main objective of this study was to explore the results and factors affecting the clinical success rate of these repeat procedures.

Methods We enrolled 99 patients with a total AF duration of more than 2 years and recurrent atrial arrhythmias after the initial catheter ablation of long-standing persistent AF. The enrolled patients were divided into two groups named the AF-recurrence group (50 patients) and the atrial tachycardia (AT)-recurrence group (49 patients) and all underwent a strict follow-up. The quality of life (QOL) and AF-related symptom classification were assessed at baseline and at 24 months post re-ablation.

Results After a mean follow-up of 31 months, 30 (30.3%) patients were free from arrhythmia recurrence, and the success rate in the AT-recurrence group was higher than that in the AF-recurrence group (32.7% vs. 28.0%, p=0.614). A Cox regression analysis revealed a CHA2DS2-VASc score ≥3 to be a predictor of recurrence. AF recurrent patients with an abnormal renal function were more prone to undergo a failed procedure. However, an abnormal renal function had no effect on the outcome of the repeat procedure for patients with AT recurrence. At the 24-month follow-up, patients maintaining sinus rhythm (SR) had a significantly improved QOL and AF-related symptoms.

Conclusion The success rate of repeat procedures for long-standing persistent AF and a total AF duration of more than 2 years is poor for patients with a CHA2DS2-VASc score ≥3. An impaired renal function has an unfavorable effect on the outcome for patients with AF recurrence. For patients maintaining SR, both the QOL and AF symptomatology improve significantly.

Key words: atrial fibrillation, atrial tachycardia, repeat catheter ablation, CHA2DS2-VASc score, estimated glomerular filtration rate, quality of life

Figure 1. Study design flowchart and the type of recurrent atrial arrhythmias prior to the repeat ablation procedure.

ity of achieving a durable sinus rhythm (SR). Previous studies have reported various predictors of recurrent AF after the index procedure, such as the CHADS2 score, left atrial (LA) dimension (3), LA size (4), AF cycle length, AF termination (5), persistent AF duration (6), valvular heart disease, non-ischemic dilated cardiomyopathy (7) and so on. Furthermore, patients with a total AF duration of more than 2 years were almost 3 times more likely to relapse after the ablation (6). However, there is a paucity of data predicting a favorable outcome after repeat procedures for patients with long-standing persistent AF and a total AF duration of more than 2 years (8).

In the present study, we aimed to investigate the long-term outcomes after repeat procedures for patients with long-standing persistent AF and a total AF duration of more than 2 years. Moreover, we intended to respectively identify the clinical factors that could predict the redo ablation outcome for patients with AF recurrence and atrial tachycardia (AT) (including atrial flutter) recurrence. Additionally, to analyze the effectiveness of catheter ablation, we evaluated the QOL and AF-related symptoms of the enrolled patients at baseline and again at 24 months post re-ablation.

Materials and Methods

Participants

A total of 424 consecutive patients with long-standing persistent AF and a total AF duration of more than 2 years undergoing the initial procedure of catheter ablation at Shanghai Chest Hospital Affiliated to Shanghai Jiao Tong University were enrolled from January 2008 to December 2011. Out of the 424 patients, 249 had recurrence after the initial procedure, and eventually 99 patients received repeat procedures in our center (Fig. 1). According to the type of recurrent atrial arrhythmia prior to the repeat procedure, we...
divided the enrolled patients into two groups named the AF-recurrence group (50 patients) and the AT-recurrence group (49 patients). The indications for undergoing repeat procedures were: (1) an age between 18 and 80 years; (2) symptomatic and drug-refractory atrial arrhythmia after the initial procedure; (3) time since the initial procedure: a minimum of 3 months; (4) signed informed consent. Patients with insufficient data, previous catheter ablation or intraoperative ablation for AF at other institutions, left atrial thrombus on transesophageal echocardiography were excluded. The study protocol was reviewed and approved by the Institutional Ethics Committee of Shanghai Chest Hospital. Among the patients with recurrent atrial arrhythmias, there was no significant difference in the baseline characteristics between the patients undergoing repeat procedures and those not undergoing repeat procedures (Table 1).

**Definitions**

Procedural termination refers to atrial arrhythmias terminated and converted to SR during the repeat procedure. The first 3 months after the initial procedure were considered to be a blanking period. After the blanking period, any over-30-second episode of AF and AT documented by electrocardiogram or Holter monitoring was considered to signify recurrence. CHADS2 assigns scores as follows: congestive heart failure, hypertension, age >75 years and diabetes mellitus count for one point each, and previous stroke or transient ischemic attack counts as two points. Compared to the CHADS2 score, CHA2DS2-VASc includes three additional factors: female gender, age 65-75 years, and vascular events. Each additional factor counts as one point, while an age older than 75 years was upgraded to two points. The R2CHADS2 score was obtained by incorporating the components of the CHADS2 score and awarding 2 points for renal dysfunction, defined as an estimated glomerular filtration rate (eGFR) <60 mL/min/1.73 m² (9). The four-variable Modification of Diet in Renal Disease (MDRD) study equation was used to calculate the eGFR from serum creatinine: 

\[ \text{eGFR (mL/min/1.73 m²)} = \frac{186.3 \times \text{[serum creatinine (mg/dL)]}^{-1.154} \times \text{[age (years)]}^{-0.203} \times (0.742 \text{ if a woman})}{10} \]

An abnormal renal function refers to the patients with eGFR <90 mL/min/1.73 m².

**Pre-ablation management**

Before the scheduled procedure, all patients underwent transthoracic echocardiography and transesophageal echocardiography to evaluate the cardiac function and exclude any intracardiac thrombi. All antiarrhythmic drugs (AAD), with the exception of amiodarone, were discontinued for at least five half-lives during the periprocedural period. Patients received effective anticoagulation with warfarin for at least 1 month, with a therapeutic international normalized ratio (INR) ranging from 2.0 to 3.0. Warfarin was discontinued 3 days before the procedure, and full-dose weight-adjusted (1.5 mg/kg daily) subcutaneous low molecular weight heparin was used to bridge the procedure until oral anticoagulation was achieved following ablation.

**Electrophysiological study and ablation procedure**

All 424 long-standing persistent AF patients with a total AF duration of more than 2 years followed the step-wise approach, including, in a progressive fashion, circumferential pulmonary vein isolation (CPVI), complex fractionated atrial electrograms (CFAEs) ablation followed by linear ablation as necessary to achieve the endpoint of AF termination. The methods and endpoints of ablation were described in detail in our previous study (11). During the repeat procedure, firstly, pulmonary vein (PV) isolation was reconfirmed with a circular mapping catheter at each PV, and CPVI was used to eliminate a recovery of the PV potentials. Secondly, for

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**Table 1. Clinical Characteristics of Patients with Recurrent Atrial Arrhythmias Undergoing Redo Procedure and Those Not Undergoing Redo Procedure.**

<table>
<thead>
<tr>
<th></th>
<th>Redo (n=99)</th>
<th>Not redo (n=76)</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>62.4±9.5</td>
<td>62.6±9.6</td>
<td>0.796</td>
</tr>
<tr>
<td>Male n(%)</td>
<td>69 (69.7%)</td>
<td>48 (63.2%)</td>
<td>0.686</td>
</tr>
<tr>
<td>Echocardiographic parameters</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>LA dimension (mm)</td>
<td>45.0±5.0</td>
<td>45.2±4.7</td>
<td>0.428</td>
</tr>
<tr>
<td>LVEF (%)</td>
<td>60.2±4.9%</td>
<td>60.4±5.5%</td>
<td>0.373</td>
</tr>
<tr>
<td>Comorbidities n (%)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hypertension</td>
<td>50 (50.5%)</td>
<td>40 (52.6%)</td>
<td>0.877</td>
</tr>
<tr>
<td>Diabetes mellitus</td>
<td>11 (11.1%)</td>
<td>10 (13.2%)</td>
<td>0.867</td>
</tr>
<tr>
<td>Coronary artery disease</td>
<td>5 (5.1%)</td>
<td>5 (6.6%)</td>
<td>0.996</td>
</tr>
<tr>
<td>HCM</td>
<td>3 (3.0%)</td>
<td>2 (2.6%)</td>
<td>0.687</td>
</tr>
<tr>
<td>DCM</td>
<td>1 (1.0%)</td>
<td>0 (0.0%)</td>
<td>0.459</td>
</tr>
<tr>
<td>COPD</td>
<td>13 (13.1%)</td>
<td>10 (13.2%)</td>
<td>0.866</td>
</tr>
<tr>
<td>Metabolic syndrome n (%)</td>
<td>40 (40.4%)</td>
<td>27 (35.5%)</td>
<td>0.458</td>
</tr>
<tr>
<td>Body mass index (kg/m²)</td>
<td>23.6±2.89</td>
<td>22.4±3.0</td>
<td>0.369</td>
</tr>
</tbody>
</table>

Data are expressed as mean±standard deviation, or percentage.

LA: left atrial; LVEF: left ventricular ejection fraction; HCM: hypertrophic cardiomyopathy; DCM: dilated cardiomyopathy; COPD: chronic obstructive pulmonary disease.
AF recurrence, the ablation of CFAEs was employed, and after restoring SR by ablation or cardioversion, the bidirectional blocks of the mitral isthmus (MI) line, the roof line, and the cavotricuspid isthmus (CTI) line were verified using the above-mentioned method (11). For AT, the suspected macroreentrant AT was mapped using CARTO three dimensional mapping, and ablated at the corresponding isthmus. If a focal AT was suspected, it was mapped and ablated at the site of earliest activation. After restoring SR, PV isolation and the complete bidirectional block of the MI, roof line and CTI were verified and consolidated as required.

**Post-ablation management**

All patients were administered low molecular weight heparin for 3-5 days after the procedure, and also took warfarin for at least 3 months, achieving the target intensity INR of 2.0-3.0. AAD (Normally, the patients take a daily oral maintenance dose of either 200-400 mg amiodarone or 150 mg propafenone three times a day. If the patients cannot tolerate such a dose, then we reduced the dose circumstances of each individual patient.) was administered for 3 months to prevent any early recurrence of atrial arrhythmia. After the blanking period, warfarin and AAD therapy were encouraged for patients who did not show a good response after the repeat procedure. The second redo ablation procedure was performed in the case of a recurrent drug-refractory atrial arrhythmia.

**Follow-up**

All patients were scheduled for regular follow-up appointments in the outpatient clinic at 1, 3, 6, and 12 months after the repeat procedure and then every 6 months. The routine investigations included transthoracic echocardiogram, 12-lead surface electrocardiogram and 24-hour Holter monitoring. Monthly telephone inquiry by the referring physicians was performed to evaluate the severity of symptoms. Clinical success was defined as free from atrial arrhythmia recurrence without taking any AAD before the last follow-up. Furthermore, cardiac computed tomography was performed at 3 months after the procedure to evaluate the presence of PV stenosis.

**Quality of life assessment and EHRA atrial fibrillation-related symptoms classification**

The Medical Outcomes Study Short-Form 36 Health Survey (SF-36) and the European Heart Rhythm Association (EHRA) class were used to assess the QOL and AF-related symptoms classification at baseline (refers to the condition when the repeat procedure was performed) and at 24 months post re-ablation (12, 13).

All patients receiving the repeat procedures subsequently completed the SF-36 questionnaire. The SF-36 assesses health-related QOL on 8 domains pertaining to physical health (4 domains) and mental health (4 domains), namely physical functioning, role limitations due to physical health, bodily pain, general health, vitality, social functioning, role limitations due to emotional problem, and mental health, respectively. For each subscale, the scores were transformed to a scale ranging from 0 to 100, with lower scores representing a lower QOL. EHRA assigns patients with AF into 4 EHRA classes based on symptoms and their effect on daily activity, namely no symptoms, mild symptoms, severe symptoms and disabling symptoms (Table 2) (13).

**Statistical analysis**

Continuous variables were expressed as the mean±standard deviation or median and interquartile ranges (25th and 75th percentile) when appropriate; while categorical variables were represented by frequencies and relative percentages. Between-group comparisons were performed using the t-test or Chi-square test. Univariate and multivariate predictors were identified using Cox proportional hazards regression models. Candidate variables with p values <0.10 according to univariate regression analyses were entered into a multivariate model. Relative risks were expressed as hazard ratios (HRs) with corresponding 95% confidence intervals (CIs). The optimal cut-off value of the eGFR for predicting AF recurrence was identified using the receiver operation characteristic (ROC) curve. Event-free survival was estimated by a Kaplan-Meier survival analysis with the log-rank test. A p value <0.05 was considered to be statistically significant. All statistical analyses were performed using the SPSS 17.0 software program.

**Results**

**Patient characteristics**

In this single-center cohort study, after the exclusion process, we eventually included 99 patients with recurrent atrial arrhythmias and a total AF duration of more than 2 years who underwent a repeat procedure in our center. Fifty out of the 99 patients had AF recurrence. Among the remaining 49 patients, AT (including atrial flutter) was the cause of recur-
rence (Fig. 1). The baseline characteristics of patients undergoing the repeat procedure are summarized in Table 3. Patients with AF recurrence had a longer total AF duration (p=0.022) and a longer duration of persistent AF (p=0.020) than those with AT recurrence (Table 3).

**Antiarhythmic drugs administration during follow-up**

After the blanking period, for patients with SR maintenance, AAD was no longer administered during follow-up. For patients who failed in the repeat procedure, AAD was firstly encouraged. After the blanking period, a total of 43 patients required ongoing AAD therapy for recurrent atrial arrhythmias. Amiodarone was predominantly used in 22 patients, propafenone was administered in 11 patients, and a combination of AAD therapy was used in 10 patients (amiodarone plus propafenone in 6 patients, and propafenone plus verapamil in 4 patients). Due to drug intolerance or severe side effect, 23 patients discontinued AAD administration and underwent a third procedure.

**Predictors of recurrence after the repeat procedure**

To analyze the predictors of recurrence after the repeat procedure, both univariate and multivariate analyses were respectively performed for AT recurrent patients (Table 4) and AF recurrent patients (Table 5). For patients with AT recurrence, a CHA2DS2-VASc score ≥3 (HR 3.720, 95% CI: 1.367-10.124, p=0.010) was identified to be a significant predictor of recurrence after the repeat procedure according to a multivariate Cox proportional hazard analysis (Table 4), while for patients with AF recurrence, a CHA2DS2-VASc score ≥3 (HR 2.707, 95% CI: 1.151-6.363, p=0.022) as well as an abnormal renal function (HR 2.821, 95% CI: 1.277-6.232, p=0.010) was independently associated with procedural failure (Table 5).

Derived from the ROC curve, eGFR <86 mL/min/1.73 m² predicted recurrence significantly in the Kaplan-Meier analysis (log-rank probability value =0.001) for AF recurrent patients after the repeat procedure, with a sensitivity of 63.9% and specificity of 92.9% (area under the ROC curve =0.812; Fig. 2).

**Long-term results after the repeat procedure and corresponding arrhythmia-free survival**

Among the 99 patients who underwent the repeat procedure, 16 out of the 49 AT recurrent patients (32.7%) were free from atrial arrhythmias; however, only 14 out of the 50 AF recurrent patients (28.0%) remained SR during the long-
Regarding the findings of the Kaplan-Meier analysis, freedom from asymptomatic or symptomatic arrhythmia for patients with AT/AF recurrence was present in Fig. 3.
Quality of life and EHRA class

Because the period of follow-up was not identical for each patient, we consistently choose 24 months post re-ablation to judge the postoperative QOL and AF-related symptom classification. According to the follow-up results, we then divided enrolled patients into two groups named non-responders and responders. Non-responders are patients who demonstrated treatment failure for the repeat procedure and had recurrent atrial arrhythmia, while responders are patients who achieved and maintained SR after the redo procedure. As portrayed in Fig. 4, there was no significant difference between the responders and non-responders at QOL baseline. At 24 months post re-ablation, compared with baseline, all eight subscales of QOL showed a significant improvement in the patients maintaining SR (responder), while only two out of the eight subscales of QOL showed an improvement in the patients with recurrent atrial arrhythmias (non-responder). The change in proportion of each ERHA level demonstrated that maintaining SR after the repeat procedure successfully reduced the AF-related symptoms.

Complications

No atrial-esophageal fistula, significant PV stenosis and procedure-related deaths occurred. The complications were as follows: cardiac tamponade (n=2, both of the patients were treated conservatively with percutaneous pericardiocentesis), arteriovenous femoral fistulae (n=1), and femoral hematoma (n=2).

Discussion

Principal study findings

We studied the outcome of a repeat procedure for long-standing persistent AF with an AF duration of more than 2 years and analyzed the predictors of recurrence after the repeat procedure. The main findings were as follows. (1) After a mean follow-up of 31 months, the overall success rates for redo AF ablation were low (30.3%). Compared with AT recurrent patients, patients with AF recurrence had a lower success rate (32.7% vs. 28.0%, p=0.614). (2) A CHA2DS2-VASc score ≥3 was a predictor of recurrence after the repeat procedure. (3) AF recurrent patients with eGFR <86 mL/min/1.73 m² after undergoing the initial procedure of long-standing persistent AF were more prone to undergo a failed repeat ablation compared to those with eGFR ≥86 mL/min/1.73 m². However, an abnormal renal function had no influ-
The majority of prior studies showed arrhythmia-free survival rates after single and multiple procedures. McCready et al. investigated 191 persistent AF patients who received catheter ablation from 2003 to 2009 and demonstrated the single procedure success rate to be 32.0% after a mean follow-up of 13.5±9.4 months, while the overall success rate was 64.0% after a mean follow-up of 13.0±8.9 months (4). Rostock et al. analyzed the outcome in 395 persistent AF patients and showed that with a single procedure, the recurrence rate was 73.0% (5). The recent study performed by
patients receiving a stepwise catheter ablation from 2006 to 2008, noted a success rate of 28.4%, 47.7% and 51.1% after the first, second and multiple procedures, respectively (3). Further, Tilz et al. describes that during 5-year follow-up, single and multiple ablation procedure success was 20% and 45%, respectively, for patients with long-standing persistent atrial fibrillation. For patients with a total AF duration of more than 2 years, the outcomes were unfavorable (6).

Considering the reasons for the difference between these outcomes, diverse populations, various types of ablation strategies, the definition of the recurrence, different types of recurrent arrhythmias after catheter ablation, and the accuracy of follow-up may together explain this discrepancy. In comparison with previous studies, we focused on the outcomes of patients with a total AF duration more than 2 years after the repeat procedure. Furthermore, all of the enrolled patients suffered from long-standing persistent AF, and they underwent the same ablation strategy in the initial procedure. During the repeat procedure, all patients reached current ablation endpoints, which meant pulmonary vein isolation, bidirectional block of lines, or the disappearance of CFAEs. The study excluded the effects of incomplete pulmonary vein isolation, insufficient ablation of CFAEs, and unblocked lines on the outcome of a repeat procedure. In addition, we pointed out that the recurrence rate after the repeat procedure for patients with long-standing persistent AF and a total AF duration more than 2 years was as high as 69.7%, especially for patients with AF recurrence (72.0%). Accordingly, due to the high recurrence rate, the decision to perform a repeat ablation for long-standing persistent AF should be made cautiously by physicians.

**CHA2DS2-VASc score and recurrence after catheter ablation during a long-term follow-up**

In the present study, we showed a CHA2DS2-VASc score ≥3 to be associated with a poor outcome after the repeat procedure for patients with AF recurrence as well as AT recurrence. Recently, a series of studies have demonstrated the CHA2DS2-VASc score to be a significant predictor of adverse events in patients undergoing the catheter ablation (14). It also serves as a combined risk predictor of first cardiovascular hospitalization for patients with AF and a risk factor of recurrence after catheter ablation (3, 15, 16). Furthermore, Park et al. proved that the electroanatomical remodeling of the left atrium (LA) was related to the CHA2DS2-VASc score in patients with non-valvular AF (17). Park and coworkers found the LA volume to be significantly higher in patients with high CHA2DS2-VASc score. A higher LA volume was closely related to a lower endocardial voltage of LA, which may help to classify the severity of the tissue pathology underlying AF before and after the catheter ablation procedure (18). The high CHA2DS2-VASc score, as a predictor of AF recurrence, implies that it does have something to do with a larger LA volume and a lower LA voltage, and plays a role in the electroanatomical remodeling of patients with AF (19).

**Estimated glomerular filtration rate and recurrence after catheter ablation during a long-term follow-up**

Recently published studies analyzed the association of an impaired renal function with post-ablation recurrence of AF (20-23). Chao et al. proved that even a mildly decreased eGFR was associated with an arrhythmogenic LA substrate and a high recurrence rate of catheter ablation in patients with paroxysmal AF (23). Tokuda et al. certified that low eGFR independently influenced the outcome of catheter ablation for paroxysmal AF (22). Our study found that for patients with AF recurrence prior to the repeat procedure, eGFR <86 mL/min/1.73 m² was a risk factor of recurrence for the redo procedure, which reconfirmed these results. Nevertheless, for patients with AT recurrence, an abnormal renal function did not help to predict the outcome of re-ablation.

Regarding the strong association between arrhythmia recurrence and a low eGFR after the repeat procedure, one possible explanation may be the correlation between an impaired renal function and established LA remodeling as already described (23). The renin-angiotensin-aldosterone system (RAAS) plays a key role in the relationship between renal disease and cardiac disease. There is strong evidence that RAAS is involved in the genesis of AF (24). When the renal function is impaired, RAAS is activated, and the secretion of angiotensin II is known to increase. Further, angiotensin II increases the atrial pressure, promotes atrial fibrosis, and modulates ion channels (25). All of these effects are involved in atrial structural and electrical remodeling, thus triggering AF. Considering that the abnormal renal function has a different effect on the outcome of the patients with AF recurrence and AT recurrence, it is likely that for patients with AT recurrence prior to the repeat procedure, atrial structural and electrical remodeling were not so serious as for patients with AF recurrence prior to the repeat procedure.

In addition, emerging evidence suggests inflammation and oxidative stress play an important role in the pathophysiology of both AF and renal dysfunction (26). The proinflammatory and profibrotic state in the patients with moderate to severe renal impairment is more prone to result in a recurrence after the repeat procedure of AF. In brief, the patients with a low eGFR after the procedure were associated with advanced atrial remodeling, including structural and electrical changes, which were shown to result in recurrence after the repeat ablation (23).

**Quality of life and EHRA class**

Although AF is seldom immediately life threatening, it imposes a substantial QOL burden on many patients (27). Previous reports have demonstrated that for patients with persistent AF, the restoration and maintenance of SR after catheter ablation was associated with an improvement in QOL measures (28, 29). In our study, a significant improve-
ment was observed in eight subscales of the SF-36 questionnaire for patients maintaining SR after the repeat procedure at 24 months post re-ablation; while only two out of the eight subscales of QOL improved in patients with recurrent atrial arrhythmias. The change in the proportion of each ERHA level indicated that maintaining SR after a repeat procedure could well reduce the AF-related symptoms. Improved QOL scores denoted the importance of the long-term maintenance of SR in patients with persistent AF.

**Study limitations**

There are several limitations associated with the present study. Firstly, this study was performed in a single-center with a relatively small sample size. Therefore, larger studies are required to confirm the validity of our observed results. Secondly, 11 patients (6.0% of 182 recurrent patients) dropped out from the follow-up because they underwent ablation at other institutions. Exclusion of these patients may have influenced the results of this study. Thirdly, the routine follow-up investigations were limited by reliance on 12-lead surface electrocardiogram and 24-hour Holter monitoring. These assessments are less precise than continuous cardiac rhythm monitoring devices. It was likely that some asymptomatic recurrence of arrhythmias was undetected. This inevitably underestimated the recurrence rate. The patients who had undetected recurrence could thus have been included in the success group, perhaps some important predictors of the repeat procedure may be masked. Fourthly, renal disease and atrial fibrillation share common risk factors, including diabetes mellitus, hypertension and abnormal left ventricular systolic function, the extent to which eGFR is a direct causal link to AF or merely a marker of systemic disease is not clear. Finally, because the follow-up periods were not identical for each patient, the QOL was only assessed at baseline and at 24 months post re-ablation.

**Conclusion**

As a whole, the success rate of all patients with long-standing persistent AF and a total AF duration of more than 2 years undergoing repeat ablation is poor with the corollary that patients with a CHA2DS2-VASc score ≥3. An impaired renal function in AF recurrent patients is a significant risk factor for a failed redo procedure. However, for those who do achieve durable SR, the QOL and AF symptomatology improve significantly.

The authors state that they have no Conflict of Interest (COI).

**Acknowledgement**

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**References**


15. Naccarelli GV, Panaccio MP, Cummins G, Tu N. CHADS2 and CHA2DS2-VASc risk factors to predict first cardiovascular hospitalization among atrial fibrillation/atrial flutter patients. Am J Car-


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