The Effects Of Ranolazine On Paroxysmal Atrial Fibrillation In Patients With Coronary Artery Disease: A Preliminary Observational Study

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Abstract

The impact of ranolazine, an anti-ischemic agent with antiarrhythmic properties, on paroxysmal atrial fibrillation (PAF) in patients with coronary artery disease (CAD) remains unclear. Pacing devices can be useful tools for disclosing even asymptomatic PAF. Purpose of this study is to assess the effect of ranolazine on atrial fibrillation (AF), in patients with CAD, PAF and a dual-chamber pacemaker.

We studied 74 patients with CAD, PAF, and sick sinus syndrome or atrio-ventricular block, treated with pacemakers capable to detect PAF episodes. The total time in AF, AF burden, and the number of PAF episodes within the last 6 months before enrolment in the study, mean AF duration per episode, and the QTc interval were initially assessed. Subsequently, patients were randomized into additional treatment with ranolazine (375 mg twice daily) or placebo. Following six months of treatment, all parameters were reassessed and compared to those before treatment.

Ranolazine was associated with shorter total AF duration (81.56±45.24 hours versus 68.71±34.84 hours, p=0.002), decreased AF burden (1.89±1.05% versus 1.59±0.81%, p=0.002), and shortened mean AF duration (1.15±0.41 hours versus 0.92±0.35 hours, p=0.01). In the placebo group no such differences were observed. In both groups, no significant differences in the number of PAF episodes and QTc duration were observed.

We conclude that in patients with CAD and PAF, ranolazine reduces the total time in AF, AF burden, and mean AF duration. These findings may imply additional antiarrhythmic properties of ranolazine on atrial myocardium and might indicate the necessity of its use in ischemic patients with PAF.

Introduction

Ranolazine is an antianginal and anti-ischemic drug with antiarrhythmic properties, as shown in experimental and clinical studies.⁴⁻⁸ In atrial myocytes, ranolazine exerts its antiarrhythmic effect mainly by inhibiting IKr, early INA, and late INA currents in a selective, use-dependent way.⁹,¹⁰ There are reports suggesting that ranolazine prevents the induction of atrial fibrillation (AF) and terminates the episodes of the arrhythmia.⁶,¹¹⁻¹３ However, in clinical practice, a considerable proportion of AF episodes are asymptomatic and this poses difficulties in the detection of the arrhythmic events and the evaluation of antiarrhythmic medications.³⁴,¹⁵

The efficacy of ranolazine in patients with ischemic heart disease and paroxysmal AF (PAF) has not been completely studied. Since implantable pacemakers can store significant information about high atrial rate episodes and rhythm control in patients with AF, they can be useful tools in evaluating the antiarrhythmic potency of ranolazine on both symptomatic and asymptomatic episodes of AF.¹⁶⁻⁻²¹ Therefore, in a prospective, placebo-controlled, randomized study, we used diagnostic data derived from the interrogation of pacing devices in order to assess the antiarrhythmic effect of ranolazine in patients with coronary artery disease (CAD) and PAF, who had been treated with a permanent dual-chamber pacemaker, due to sick sinus syndrome (SSS) or atrio-ventricular conduction block (AVB).

Material and Methods

Patients

Among patients routinely followed in the Pacemaker Clinic within...
an 18-month period, 460 individuals were initially assessed, and patients eligible for enrollment in the study were those with known CAD and myocardial ischemia, who had been implanted a dual-chamber pacemaker due to SSS or AVB. The pacemakers should be implanted at least six months before recruitment in the study and should be able to store and recall atrial high-rate episodes.

The diagnosis of CAD was based on patients’ history, documented in the past by typical symptoms, electrocardiographic recordings, markers of myocardial necrosis, coronary angiography and interventions for cardiac revascularization, when indicated. Exercise stress test or myocardial scintigraphy revealed myocardial ischemia. SSS and atrio-ventricular conduction abnormalities were diagnosed by symptoms (syncope, presyncope or inappropriate fatigue) combined with 12-lead electrocardiogram, 24-hour Holter recordings and electrophysiological study, if necessary. PAF was defined according to the Task Force for management of Atrial Fibrillation of the European Society of Cardiology, and documented by Holter recordings and the interrogation of the pacing devices.22

Exclusion criteria were recent myocardial infarction, unstable angina, candidacy for revascularization, congestive heart failure, renal failure (creatinine clearance < 30 ml/min), hepatic impairment, and age above eighty-five years. Additionally, patients under antiarrhythmic medication were not included in the study, in order to avoid drug interactions that would make the evaluation of the effect of ranolazine difficult.

Patients received the optimal hemodynamically tolerated treatment, and were randomly assigned to adjunctive treatment with ranolazine or placebo (control group), in an 1:1 order. Each patient was given a unique 3-digit numeric code upon enrollment, and the “Random Allocation Software” for parallel group randomized trial was used to produce a simple 1:1 randomization of the predefined candidacy code numbers.

The following pacemaker models had been implanted in the studied population: 1) Adapta DR, Medtronic, Minneapolis, USA, 2) Symphony DR, Sorin, Milan, Italy, 3) Insignia Ultra DR, Boston Scientific, Phoenix, Arizona, USA, 4) T60 DR and C70 DR, Vitatron, Maastricht, Netherlands, and 5) Victory XL DR and Identity XL DR, St. Jude Medical, St. Paul, Minneapolis, USA. The devices had automatic mode switch features and could provide information concerning the number of AF episodes (mode switches), the duration of each episode, and the total time in AF (AF burden) between two sequential interrogations. Additionally, AF burden was automatically calculated and expressed as percentage (%). Initial Evaluation and Assessed Parameters

Clinical examination, twelve-lead electrocardiogram along with assessment of the corrected QT-interval (QTc), evaluation of renal function, and transthoracic echocardiographic study were performed at baseline. Left ventricular end-diastolic and end-systolic diameter (LVEDD and LVESD, respectively) were measured, and left ventricular ejection fraction (EF) was estimated according to the modified Simpson’s method. The anteroposterior diameter of the left atrium (LA) was measured at the parasternal short axis. Demographics, clinical characteristics and medication of the studied population are presented in the Table. The total time in AF, AF burden, and the number of AF episodes during the last six months before enrolment in the study were recalled during the interrogation of the pacemakers at the time of recruitment (baseline). Furthermore, AF duration per episode (mean AF duration) was calculated.

QTc was the Q-T interval corrected for heart rate and expressed in msec. It was calculated by the duration of the QT interval divided by the square root of the R-R interval, according to the Bazett’s formula. AF burden was presented as percentage (%), and defined as the total time in AF (time in mode-switch) during follow-up, divided by the time of follow-up (six months) and multiplied by 100.

AF duration per episode (mean AF duration) was calculated by dividing the total time in AF by the number of AF episodes.

Randomized Treatment and Follow-Up

In the ranolazine group, ranolazine was administered at a dose of 375 mg twice daily, as an adjunctive therapy on top of conventional medication for CAD, according to current indications and the rules of the Hellenic National drug Organization.23 This dose scheme

| Table 1: Demographics, clinical characteristics, history of revascularization and medication of the studied population, presented in groups, according to randomization. |
|-----------------|-----------------|-----------------|
| Ranolazine group | Control group | p value |
| (n=36) | (n=36) |
| Age (yrs) | 73.8 ± 6.8 | 73.9 ± 4.5 | 0.726 |
| Male gender (%) | 25 (69.4%) | 22 (61.1%) | 0.621 |
| SSS | 21 (58.3%) | 23 (63.9%) | 0.809 |
| AVB | 15 (41.7%) | 13 (36.1%) | 0.809 |
| Prior revascularization | 18 (50.0%) | 15 (41.7%) | 0.637 |
| CABG | 5 (13.9%) | 4 (11.1%) | 1.000 |
| PCI | 13 (36.1%) | 11 (30.6%) | 0.803 |
| Angina | 21 (58.9%) | 23 (63.9%) | 0.809 |
| Positive stress test/scintigraphy | 36 (100%) | 36 (100%) | 1.000 |
| Echocardiography | | | |
| EF (%) | 54.6 ± 6.3 | 54.3 ± 7.6 | 0.943 |
| LVEDD (mm) | 51.4 ± 2.7 | 50.7 ± 2.8 | 0.233 |
| LVESD (mm) | 32.4 ± 2.2 | 32.1 ± 2.2 | 0.446 |
| LA diameter (mm) | 42.9 ± 2.7 | 42.6 ± 3.1 | 0.687 |
| QTc (msec) | 436 ± 10 | 437 ± 9 | 0.857 |
| AF statistics at baseline | | | |
| total time in AF (hrs) | 81.56 ± 45.24 | 80.28 ± 43.198 | 0.903 |
| AF burden (%) | 1.89 ± 1.05 | 1.86 ± 1.00 | 0.903 |
| number of AF episodes | 77.3 ± 36.1 | 71.56 ± 30.92 | 0.468 |
| mean duration (hrs) | 1.05 ± 0.41 | 1.12 ± 0.43 | 0.486 |
| Comorbidities | | | |
| hypertension | 28 (77.8%) | 29 (80.6%) | 1.000 |
| dyslipidemia | 24 (66.7%) | 26 (72.2%) | 0.798 |
| diabetes mellitus | 8 (22.2%) | 3 (8.3%) | 0.189 |
| thyroid disease | 2 (5.6%) | 3 (8.3%) | 1.000 |
| Medication | | | |
| nitrates | 30 (83.3%) | 33 (91.6%) | 0.631 |
| β-blocker | 36 (100.0%) | 35 (97.2%) | 1.000 |
| ACE inhibitor/ARB | 29 (80.6%) | 25 (69.4%) | 0.415 |
| calcium channel blocker | 12 (33.3%) | 14 (38.9%) | 0.806 |
| dihydropyridine | 9 (25.0%) | 12 (33.3%) | 0.605 |
| non-dihydropyridine | 3 (8.3%) | 2 (5.6%) | 1.000 |
| statin | 34 (94.4%) | 36 (100%) | 0.798 |
| diuretics | 13 (36.1%) | 16 (44.4%) | 0.631 |

Comparisons of continuous variables between groups were performed using Student’s t-test for independent samples, preceded by Levene’s test for equality of variances, or Mann–Whitney U non-parametric test, as appropriate. Comparisons of continuous variables within the same group prior to and following six months of treatment were performed using paired Student’s t-test or Wilcoxon signed-rank test for related samples, as appropriate. Categorical data were summarized in contingency tables and between groups comparisons were performed using the Fisher’s exact test.

All statistical significance tests were two-sided and the null hypothesis was the presumption of equality of means. Differences between and within groups were considered to be statistically significant if the null hypothesis could be rejected with >95% level of confidence (p-value <0.05).

Results

Baseline Characteristics

Out of the 460 patients’ pool, 77 met the eligibility criteria for the study. Seventy four of them provided written informed consent and were initially enrolled. Two patients were excluded from the study: one from the ranolazine group developed unstable angina and another from the control group had to change his treatment due to unsatisfactory control of his blood pressure (Figure 1). Demographics, clinical characteristics, history, medication of the remaining 72 studied patients and comparison between groups are presented in the Table.

During the initial evaluation, patients treated with ranolazine and controls were comparable regarding the aforementioned parameters (Table 1).

Treatment and Follow-Up

Ranolazine was overall well tolerated and no serious side effects were reported. Initially, five patients complained about constipation, abdominal discomfort or dizziness within the first two weeks of ranolazine administration. However, these symptoms were attenuated...
later on and patients continued treatment for the entire follow-up period.

Within group comparisons showed a statistically significant decrease of the total time of AF with ranolazine, from 81.56 ± 45.24 hours before treatment to 68.71 ± 34.84 hours during therapy (p=0.002), as presented in Figure 2. AF burden decreased too (Figure 3), from 1.89 ± 1.05 % to 1.59 ± 0.81 % (p=0.002). The number of AF episodes under treatment with ranolazine was not significantly lower compared to that before treatment (75.53 ± 33.13 episodes versus 77.33 ± 36.12 episodes, respectively, p=0.61), as depicted in Figure 4. However, the mean AF duration decreased from 1.05 ± 0.41 hours before treatment to 0.92 ± 0.35 hours (p=0.01) with ranolazine (Figure 5). The duration of QTc interval following ranolazine administration was comparable with that before treatment (436 ± 10 msec and 438 ± 10 msec, respectively, p=0.25).

In the control group, no significant changes were observed between the six-month period before enrolment in the study and during follow-up regarding the total time in AF (80.28 ± 43.20 hours versus 81.40 ± 42.54 hours, p=0.57), AF burden (1.86 ± 1.00 % versus 1.88 ± 0.99 %, p=0.57), mean time in AF per episode (1.12 ± 0.43 hours versus 1.11 ± 0.40 hours, p=0.53), the number of AF episodes (71.56 ± 30.92 episodes versus 72.39 ± 26.99 episodes, p=0.66), and QTc duration (437 ± 9 msec versus 437 ± 10 msec, p=0.60).

The comparison between groups confirmed the significant difference in the mean duration of AF episodes (p=0.034). Furthermore, when the changes in all the aforementioned parameters were compared between groups, a statistically significant difference favoring ranolazine was found (Table 2).

### Discussion
In the present study, the effect of adjunctive treatment with ranolazine on AF burden was evaluated in patients with ischemic heart disease and PAF. Implanted dual-chamber pacing devices were used for heart rhythm monitoring, detection of asymptomatic PAF, and providing information regarding the number and duration of AF episodes. At the selected dose-scheme, ranolazine was associated with a statistically significant decrease in the total time in AF, AF burden, and the duration of each AF episode.

The studied patients were clinically stable during follow-up, so that the progression of ischemia would not be involved in the results. Patients with highly symptomatic AF necessitating antiarrhythmic medication were excluded, in order i) to avoid interactions between the antiarrhythmic agents and ranolazine, and ii) to evaluate the pure antiarrhythmic effect of ranolazine. These restrictions resulted in a highly selected sample, and give advantages to the study.

### Total AF Burden
Experimental and clinical studies suggest an antifibrillatory impact of ranolazine on atrial myocardium. It has been reported that ranolazine is more effective than lidocaine in terminating AF in perfused canine right atrial preparations, mainly through the development of a rate-dependent postrepolarization refractoriness.\(^\text{11}\) In canine isolated atrial preparations, ranolazine was comparable to propafenone in terminating acetylcholine-mediated persistent AF.\(^\text{24}\) Besides, ranolazine suppressed triggers of AF originating from the sleeves of pulmonary veins.\(^\text{25}\) Additionally, it has been proposed that ranolazine may prevent AF by suppressing cellular calcium overload and delayed afterdepolarizations that can trigger atrial arrhythmias.\(^\text{26}\) The role of the late phase 3 EAD-induced trigger activity has been

### Table 2: The differences in total time in AF, AF burden, number of AF episodes, and mean duration of each episode between baseline and during follow-up are compared between the two groups.

<table>
<thead>
<tr>
<th></th>
<th>Ranolazine group (n=36)</th>
<th>Control group (n=36)</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Difference in total time in AF (hrs)</td>
<td>-12.85 ± 22.94</td>
<td>1.13 ± 11.77</td>
<td>0.002</td>
</tr>
<tr>
<td>Difference in AF burden (%)</td>
<td>-0.30 ± 0.53</td>
<td>0.03 ± 0.27</td>
<td>0.002</td>
</tr>
<tr>
<td>Difference in the number of AF episodes</td>
<td>-1.81 ± 12.55</td>
<td>0.83 ± 11.28</td>
<td>0.528</td>
</tr>
<tr>
<td>Difference in mean duration (hrs)</td>
<td>-0.13 ± 0.30</td>
<td>-0.01 ± 0.19</td>
<td>0.041</td>
</tr>
</tbody>
</table>

hrs: hours
shown to be involved in the reinduction immediately after an episode of atrial fibrillation and the prolongation of the total duration of the arrhythmia.\textsuperscript{17}

In human right atrial appendages obtained from patients undergoing heart surgery, ranolazine was found to suppress calcium- or isoprenaline-induced premature atrial contractions that might initiate or perpetuate AF.\textsuperscript{28} Furthermore, there are clinical observations suggesting that oral ranolazine could be used in a “pill in the pocket” approach to convert PAF, and that it could facilitate electrical cardioversion in cardioversion-resistant cases.\textsuperscript{12,29}

In accordance with these studies, we found that treatment with ranolazine at the aforementioned dose-scheme was associated with decreased AF burden during follow-up and shorter AF duration per episode. These observations indicate the earlier termination of AF episodes and could be attributed to the use-dependent properties of ranolazine, the increase of atrial refractoriness and the suppression of premature contractions in the atrium or pulmonary veins that can not only induce but also perpetuate AF. The reduction of AF burden and AF duration per episode may have an even greater clinical importance in patients with higher AF burden and more frequent AF relapse.\textsuperscript{30} Finally, an additional anti-ischemic effect and the improvement in diastolic dysfunction might also be involved in this beneficial effect, since ischemia has been implicated in the pathophysiology of AF.\textsuperscript{31-35}

Dual-chamber pacemakers have been used as tools for the evaluation of the efficacy of antiarrhythmic agents before. In a recent study, bjudiodarone, a novel chemical analogue of amiodarone, administered at a dose of 200 mg daily in patients with PAF and a dual-chamber pacemaker, decrease in AF burden by 14% approximately.\textsuperscript{36} Our results are comparable with those of bjudiodarone, since ranolazine was associated with a decrease in total time in AF and AF burden by 16%. The antiarrhythmic effect of ranolazine was observed despite the relatively low initial load of AF in the studied population. Although this effect may have a small impact on the quality of life among asymptomatic patients or those with short AF duration, it is important in terms of pathophysiology and deserves further clinical investigation.

Number of AF Episodes

Treatment with ranolazine was not associated with a significant decrease in the number of AF episodes. This can be explained if the following aspects are taken under consideration: i) Some episodes of AF might have been aborted by the beneficial effect of atrial pacing. In patients with SSS, atrial pacing reduces the incidence of AF, especially when ventricular pacing is minimized.\textsuperscript{37,38} Besides, it has been reported that atrial pacing diminishes the arrhythmogenic dispersion of atrial refractoriness associated with bradycardia.\textsuperscript{19} Furthermore, atrial pacing may prevent the induction of AF by suppressing premature atrial contractions.\textsuperscript{39,41} ii) Ranolazine exhibits rate-dependent antiarrhythmic properties.\textsuperscript{3,11,42} This may result in an increased effect of the drug during each PAF episode compared to that in sinus rhythm. iii) Higher doses of ranolazine, if well tolerated, may be more effective in reducing AF recurrence. iv) The studied population was mainly characterized by short and sporadic episodes of PAF. This might be an additional reason for not observing any significant difference in the number of AF episodes with ranolazine. The confirmation of this antiarrhythmic effect in patients with high AF burden is a challenge for the future and it could introduce new indications for the use of ranolazine.
and individuals with higher AF burden, more frequent and longer AF episodes.

The potential antiarrhythmic effect of pacing cannot be ruled out. However, this beneficial impact was equally distributed in the controls and the ranolazine-group.

The number of patients enrolled in the study is relatively small, due to the specific inclusion criteria and the indications for ranolazine administration. Therefore, they were not separately studied in different groups according to a predominantly atrial pacing (sinus node dysfunction) versus ventricular pacing (impaired atrioventricular conduction). A potential correlation between the ischemic burden and the effect of ranolazine was not assessed in the study. However, in the within-group statistical analysis, each patient was the control of himself and the antiarrhythmic effect was evaluated on the same ischemic substrate, and with the same pacing mode before and following treatment.

Conclusions:

In a selected population of patients with CAD, PAF, and a dual-chamber pacemaker, ranolazine may have an antiarrhythmic effect, indicated by a shorter total time in AF, a reduced AF burden, and decreased mean AF duration, as documented by the pacing devices. These observations may imply a beneficial effect of ranolazine administration in patients with CAD and PAF, necessitating further clinical confirmation.

References:


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Original Research


