Aim: Angiotensin-converting enzyme (ACE) inhibitors and beta blockers were shown to be effective in prevention of atrial fibrillation (AF) recurrences after electrical cardioversion, by interfering with atrial structural and electrical remodeling. In this study, we aimed to compare the effects of ACE inhibitor cilazapril and beta blocker metoprolol after cardioversion of AF in terms of sinus rhythm maintenance.

Patients and Methods: The study population comprised 120 patients with persistent atrial fibrillation (>7 days) who underwent successful cardioversion. Randomization into 3 groups was done after successful cardioversion. Group I (n=41) was treated with amiodarone, group II (n=41) was treated with amiodarone plus metoprolol, and group III (n=38) was treated with amiodarone plus cilazapril. The primary end-point of the study was the recurrence rates between groups at one year and with those who recurred, the time to recurrence of AF.

Results: After 1 year follow-up, maintenance of sinus rhythm were similar among groups (Kaplan-Meier analysis, 24%, 37% and 26% respectively; log rank=0.3). There was no difference among groups with respect to time to recurrence of AF.

Conclusion: Patients treated with amiodarone only, amiodarone plus metoprolol and amiodarone plus cilazapril had similar rates of recurrence of AF.

Key Words: Atrial fibrillation, cardioversion, angiotensin-converting enzyme inhibitors, beta blockers

Amaç: Atriyal fibrilasyon (AF) elektriksel kardiyoversiyonu sonrasında anjiyotensin dönüştürücü enzim (ACE) inhibitörleri ve beta blokerlerin atriyal yapısal ve elektriksel remodeling'i azaltmaya surette AF nükslerini azalttığı gösterilmiştir. Bu çalışmada, kardiyoversiyon sonrasında ACE inhibitörü cilazapril ve beta bloker metoprolol'un sinüs ritmi idamesindeki etkileri karşılaştırdık.

Hastalar ve Yöntem: Çalışmaya persistan AF'si olup başarılı kardiyoversiyon yapılan 120 hasta alındı. Kardiyoversiyon sonrasında hastalar 3 gruba randomize edildi. Grup I' e (n=41) amiodaron, Grup II' ye (n=41) amiodaron+metoprolol, Grup III' e (n=38) amiodaron+cilazapril verildi. Çalışmanın primer sonlanım noktası AF nüks oranları ve AF nüksü görülenlerde nükse kadar geçen zaman idi.

Bulgular: Bir yıllık takip sonunda gruplar arasında sinüs ritmi idamesi arasında anlamlı fark saptanmadı (Kaplan Meier analizi, sırasıyla %24, %37 and %26; log rank=0.3). AF nüks zamandan açı- sıldan da gruplar arasında fark gözlemedi.

Sonuç: Amiodaron, amiodaron+metoprolol veya amiodaron+cilazapril tedavilerinin AF nüksü üzerine etkileri benzer bulunmuştur.

Anahtar Kelimeler: Atriyal fibrilasyon, kardiyoversiyon, anjiyotensin dönüştürücü enzim inhibitörleri, beta blokerler

Atrial fibrillation (AF) is a common arrhythmia in clinical practice, and may result in serious complications (1). Direct current cardioversion of persistent AF is the most effective treatment for the restoration of sinus rhythm but it may be hampered by high recurrence rates (2). One of the accepted mechanisms of AF recurrence is electrical and
structural remodelling caused by changes in the refractory period of the atrial muscle and atrial fibrosis with intraatrial conduction disturbances (3). Renin-angiotensin system (RAS) leads to cardiac fibrosis via increased angiotensin II and aldosterone levels in a variety of cardiac disorders (4). Three-fold increase in angiotensin-converting enzyme (ACE) expression occurs in chronic persistent AF (5), and several studies reported the relationship between increased angiotensin II levels and arrhythmogenic atrial electrical and structural remodelling that could be reversed by blockade of the RAS (6-9).

Abnormal autonomic control is another mechanism that was suggested to increase AF recurrence. Increased adrenergic and reduced vagal stimulation contribute to atrial electrical remodeling by facilitating intracellular calcium overload (10). Intracellular calcium lowering drugs (beta blockers and calcium channel blockers) (11) and beta blockers (12, 13) were found to be effective in reduction of AF recurrences after cardioversion.

Up to our knowledge, there is no prospective study comparing the effect of ACE inhibitors and beta blockers on AF recurrence rates after electrical cardioversion. In the present study, we aimed to investigate whether the ACE inhibitor cilazapril or the beta blocker metoprolol add benefit on the antiarrhythmic drug amiodarone and to compare the effects of these drugs in terms of maintaining sinus rhythm after electrical cardioversion of AF.

Methods

The study population comprised 120 patients with persistent atrial fibrillation (>7 days) who underwent successful cardioversion between December 2002 and February 2005. During this time period a total of 205 patients referred to our clinic for electrical cardioversion were screened and 85 of them were excluded. The reasons were: Presence of thrombus in the left atrium (14 patients), unsuccessful cardioversion (26 patients), unwilling to participate (5 patients) and presence of one of the exclusion criteria listed below (40 patients).

Exclusion criteria included a left atrium size > 6 cm, acute coronary syndrome within 6 weeks, known thyroid, hepatic or pulmonary disease, heart surgery within 6 weeks, contraindications to treatment with amiodarone, beta-blockers or ACE-inhibitors, heart failure (NYHA class III-IV), rheumatic valve disease, paroxysmal atrial fibrillation, known allergy to amiodarone, beta-blockers or ACE-inhibitors.

Written informed consent was obtained from all patients before they entered the study. All patients clinical history, physical examination, TSH measurements and transthoracic echocardiograms were done and scheduled for transoesophageal echocardiography (TEE). Patients were asked to withdraw ACE inhibitors and/or beta-blockers 1 week before TEE. Those with hypertension, amiodarone was initiated. After TEE demonstrated no visible thrombus in the left atrium, cardioversion was done in the same day. Patients who were not previously anticoagulated started both warfarin and heparin infusion. The infusion was stopped when the international normalized ration (INR) >2. Cardioversion was performed with a biphasic defibrillator (HeartStart XL, Philips). Successful cardioversion was defined as sinus rhythm recovery lasting as least 1 minute after the shock. Anticoagulation was continued for at least 4 weeks after electrical cardioversion.

Randomization into 3 groups was done after successful cardioversion: Group I: Amiodarone (Cor-darone, Sanofi-Synthelabo) only group (900mg IV infusion just after the shock for 24 hours, followed by oral amiodarone 200mg three times a day for one week, two times a day for subsequent week and thereafter daily 200mg maintenance dose); Group II: Amiodarone plus metoprolol (Be-lod, Astra-Zeneca) (50mg/day which could be increased to 100mg/day) and Group III: Amiodarone plus cilazapril (Inhibace, Roche) (5mg/day which could be increased to 10mg/day in hypertensive patients). The drugs were given in an open-label fashion. Patients were not allowed to use ACE-inhibitors, angiotensin receptor blockers or beta-blockers other than the assigned treatment. If the patient had high blood pressure amlodipin was initiated.

Patients were examined at 1, 3, 6 and 12 months, and at any time the patient complained of palpitations or any other symptoms. Standard 12-lead ECG and inquiry about any recurrence of palpitation was done at each visit. The cardiologist who assessed the outcome was blinded for the patient’s group assignment.

The primary end-point of the study was the comparison of recurrence rates between groups at one year and with those who recurred, the time to recurrence of AF.

Statistical Analysis:

Statistical analyses were performed
Comparison of Angiotensin Converting Enzyme Inhibitors and Beta Blockers in Prevention of Atrial Fibrillation Recurrences After Electrical Cardioversion

using SPSS 10.0 (version 10.0 for Windows, SPSS Inc., Chicago, Illinois). Data are expressed as numbers and percentages for discrete variables and as means ± SD for continuous variables. Comparisons between groups were performed by univariate analysis with the one-way ANOVA or Kruskal-Wallis analysis of variance test for the continuous variables and by using chi-square test for the other parameters. Estimates of the proportion of patients remaining in sinus rhythm over time were constructed using the method of Kaplan–Meier and compared with log rank test. Results with a p value less than 0.05 were considered significant.

Results

The study population consisted 120 patients with persistent atrial fibrillation who underwent successful cardioversion (mean age: 62 ± 12 years, 50 were male). The baseline demographic and clinical characteristics of the groups were given in Table 1. As noted, groups were similar, whereas diabetes mellitus prevalence and pulmonary artery pressure were tended to be lower in group I (Table 1). The mean duration of AF before randomization was 19 months, with no differences among the groups. As shown in table 1, the groups were similar with regard to all concomitant medications.

At the 1st month follow-up visit, 10 patients had a recurrence of AF (4 patients in group I, 3 patients in group II, and 3 patients in group III, p=0.9). The proportion of patients with sinus rhythm at 3rd and 6th months follow-up were also similar (Table 2). After a median follow-up period of 12 months, Kaplan-Meier analysis (Figure 1) showed that maintenance of sinus rhythm between groups did not differ (Kaplan-Meier analysis, Figure 1: Comparison of recurrence times among groups.

Table 1: Characteristics of the patients

<table>
<thead>
<tr>
<th></th>
<th>Group I n=41</th>
<th>Group II n=41</th>
<th>Group III n=38</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, years</td>
<td>62±13</td>
<td>63±11</td>
<td>61±11</td>
<td>0.8</td>
</tr>
<tr>
<td>Male, n (%)</td>
<td>16 (39)</td>
<td>16 (39)</td>
<td>18 (47)</td>
<td>0.7</td>
</tr>
<tr>
<td>Underlying heart disease, n (%)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Coronary artery disease</td>
<td>20 (48)</td>
<td>12 (29)</td>
<td>10 (26)</td>
<td>0.1</td>
</tr>
<tr>
<td>Mitral valve disease</td>
<td>13 (32)</td>
<td>13 (32)</td>
<td>11 (29)</td>
<td></td>
</tr>
<tr>
<td>Hypertension</td>
<td>4 (10)</td>
<td>12 (29)</td>
<td>7 (18)</td>
<td></td>
</tr>
<tr>
<td>Dilated CMP</td>
<td>2 (5)</td>
<td>2 (5)</td>
<td>7 (18)</td>
<td></td>
</tr>
<tr>
<td>Other</td>
<td>2 (5)</td>
<td>2 (5)</td>
<td>3 (9)</td>
<td></td>
</tr>
<tr>
<td>Body mass index, kg/m²</td>
<td>25±3.4</td>
<td>27±4.3</td>
<td>27±3.2</td>
<td>0.7</td>
</tr>
<tr>
<td>Diabetes mellitus, n (%)</td>
<td>5 (12)</td>
<td>11 (27)</td>
<td>10 (26)</td>
<td>0.2</td>
</tr>
<tr>
<td>Smoking, n (%)</td>
<td>6 (15)</td>
<td>2 (5)</td>
<td>6 (16)</td>
<td>0.2</td>
</tr>
<tr>
<td>Left atrial diameter, mm</td>
<td>5±0.6</td>
<td>4.9±0.7</td>
<td>4.7±0.8</td>
<td>0.1</td>
</tr>
<tr>
<td>PAP, mmHg, n (%)</td>
<td>35±5</td>
<td>46±16</td>
<td>43±8</td>
<td>0.3</td>
</tr>
<tr>
<td>Ejection fraction, %</td>
<td>51±12</td>
<td>55±11</td>
<td>49±14</td>
<td>0.08</td>
</tr>
<tr>
<td>Duration of AF, months, n (%)</td>
<td>18±8</td>
<td>21±10</td>
<td>19±11</td>
<td>0.3</td>
</tr>
<tr>
<td>Concomitant medications, n (%)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Digoxin</td>
<td>5 (12)</td>
<td>1 (3)</td>
<td>4 (11)</td>
<td>0.3</td>
</tr>
<tr>
<td>CCB</td>
<td>6 (15)</td>
<td>4 (10)</td>
<td>1 (3)</td>
<td>0.2</td>
</tr>
<tr>
<td>Diuretics</td>
<td>2 (5)</td>
<td>1 (3)</td>
<td>1 (3)</td>
<td>0.8</td>
</tr>
<tr>
<td>Anticoagulant drugs</td>
<td>41 (100)</td>
<td>41 (100)</td>
<td>38 (100)</td>
<td>1</td>
</tr>
<tr>
<td>Aspirin</td>
<td>20 (48)</td>
<td>12 (29)</td>
<td>12 (32)</td>
<td>0.1</td>
</tr>
<tr>
<td>Number of shocks</td>
<td>2.4±0.5</td>
<td>1.7±0.8</td>
<td>1.7±0.9</td>
<td>0.3</td>
</tr>
</tbody>
</table>

AF, atrial fibrillation; CCB, calcium channel blockers; CMP, cardiomyopathy; PAP, pulmonary arterial pressure.
76%, 63% and 74% respectively; log rank=0.3).

Adverse clinical events resulted in discontinuation in 4 patients (10%) treated with amiodarone, 8 patients (20%) treated with amiodarone plus metoprolol and 5 (13%) patients treated with amiodarone plus cilazapril (Table 2). Three patients discontinued cilazapril because of dry cough, one patient because of elevated potassium levels. Discontinuation of amiodarone occurred in 4 patients in group I, 3 patients in group II and 1 patient in group III. The reasons were: thyroid pathologies (4 patients), gastrointestinal events (2 patients) and elevation of liver enzymes (2 patients). Five patient discontinued metoprolol because of symptomatic bradycardia in group II. None of the patients died during the study period and no thrombo-embolic events occurred.

Discussion

The results of the present study show that adding cilazapril or metoprolol to amiodarone does not decrease the recurrence rates of AF after electrical cardioversion. Amiodarone and combination of these drugs were well tolerated by the patients.

Most of the AF recurrences are thought to be due to atrial electrical and structural remodeling that are partly mediated by RAS and autonomic nervous system (3, 10). AF leads to development of atrial fibrosis, which has been suggested to be responsible for electrophysiological changes such as atrial conduction delay or reduced atrial effective refractory period (14).

Experimental animal studies showed the critical role of angiotensin II in both types of atrial remodeling in which histological and electrophysiological properties were reversed by ACE inhibition and angiotensin II blockade (6, 7). These findings were supported by clinical studies in which blockade of RAS with ACE inhibitor enalapril (8) or angiotensin II receptor antagonists irbesartan (9) as an adjunct to amiodarone facilitated sinus rhythm maintenance after cardioversion. Our results are in contrast with those findings. The present and previous studies differ in some aspects. First, in the previous studies (8, 9) the drugs were prescribed 4 weeks before, whereas our patients began to take the drugs in the day of electrical cardioversion. In persistent AF, the effects of ACE inhibitors are thought to be mostly on structural, rather than electrical remodeling (15), which may need a sufficient period of time to exert their beneficial effects. However, we believe that one-year follow-up period in our study is long enough to observe the results of possible structural influences of cilazapril. Secondly, in the study of Madrid et al. (9), there was a trend towards to higher beta blocker use in the irbesartan+amiodarone group than the amiodarone only group. So, co-administration of angiotensin II antagonists and beta blockers might have exerted synergistic effect on RAS and sympathetic nervous system, rather than RAS blockade only. Third, we used a different kind of ACE inhibitor, cilazapril. Studies investigating the effect of cilazapril on AF recurrence is limited. Recently, Li et al. demonstrated the supressive effect of cilazapril on atrial structural remodeling and the incidence of AF in dogs paced with high atrial rates (16). However, up to our knowledge, evidence lacks about the effect of cilazapril on AF recurrences in humans. Finally, atrial angiotensin II concentrations were found to be increased before plasma levels rised, suggesting in situ cardiac tissue synthesis as the source of atrial angiotensin II increases in an experimental model of atrial fibrillation in dogs (17). So, differences in the capability of blocking tissue RAS among ACE inhibitors may be responsible for the different results.

The other finding in the present study is the absence of any benefit of metoprolol on sinus rhythm maintenance. Van Noord et al. demonstrated that the beneficial effects of beta blockers in preventing AF relapse after cardioversion were
more pronounced in the setting of hypertension rather than lone AF (18). The possible mechanism was suggested to be the lengthening of diastole, thereby decreasing atrial stretch related arrhythmogenicity. In our study, the spectrum of underlying heart diseases were similar among groups. Van Noord et al. also underscored the importance of initiating beta blocker therapy before the scheduled cardioversion, especially in terms of preventing earlier relapses (18). This was further pronounced by the study of Workman et al., in which atrial electrophysiological changes such as prolongation of atrial action potential duration and effective refractory period were suggested to be consistent with a long-term adaptive response, a type of “pharmacological remodeling”, that appears as a result of long-term beta blockade (19). In the present study, beta blockers were not initiated before the scheduled cardioversion. In fact, others demonstrated that preventive effect of metoprolol from AF recurrence still persevered despite initiation on the day of cardioversion (12). However, none of the studies above (12, 18) used the combination of amiodarone and beta blocker together. So, amiodarone might have masked the effects of metoprolol on AF recurrences, in the present study.

In the present study neither cilazapril nor metoprolol were superior to each other. In the literature, the studies comparing ACE inhibitors and beta blockers after electrical cardioversion of AF is limited. In a large hypertension trial, blockade of RAS by angiotensin antagonist losartan had superior effects over the beta blocker atenolol in terms of reducing new onset AF (20). However, this study enrolled only the patients with left ventricular hypertrophy, a group of patients with more advanced hemodynamic abnormalities. As well, beta blockers are less effective in reducing hypertrophy and may be possibly less effective in preventing AF (21). The absence of any benefit of ACE inhibitors over beta blockers in terms of reducing new-onset AF in two other large hypertension trials (22, 23) partly supports our results. In fact, the two drugs act somewhat in a parallel manner. The possible mechanisms other than reversal of atrial remodeling, by which ACE inhibitors may exert antiarrhythmic effect include the decrease of wall stress, improvement of left ventricular function, decrease of left ventricular end-diastolic pressure and left atrial pressure, and modulation of ion currents and refractoriness (24). Beta blockers have also modulatory actions on ion channels (19), and they may be antiarrhythmic by lengthening diastole and enhancing ventricular filling, thereby ameliorating atrial-stretch (18). ACE inhibitors have beta-blocking properties, and beta blockers depress renin activity. Furthermore, structural and electrical remodeling, the two entities which ACE inhibitors and beta blockers are proposed to act on, are interrelated, in such a way that increases in atrial pressure have been shown to produce electrical remodeling, and prolonged rapid atrial rates can cause atrial dilatation (25).

This study has some limitations. First, the beginning time (at the time of cardioversion) of the drugs may be relatively late. This might effect the comparison of the recurrence rates in short-term, because longer time may be needed to exert their effects. However, our relatively long follow-up (one year) is thought to be enough to eliminate this problem at least in long-term. Secondly, we did not perform electrophysiological study or histological analysis for ethical reasons. These were performed previously in humans (26) and animals (6, 7). Finally, we did not evaluate possible paroxysmal AF attacks in our patients, so we might have missed asymptomatic episodes. However, our aim was to investigate recurrence of persistent AF, not paroxysmal AF. We examined the patients at definite intervals or at any time they had symptoms to evaluate if there were any recurrences of persistent AF. So, we think that our methodology was sufficient for the aim of the study.

In conclusion, we could neither demonstrate any beneficial effect of concomitant use of cilazapril or metoprolol with amiodarone on cardioversion outcome in persistent AF. Our results may be important for that ACE inhibitors and beta blockers as an adjunct to amiodarone were compared with respect to their effect on sinus rhythm maintenance after electrical cardioversion of AF. Larger prospective, randomized, controlled trials with different drug combinations (eg. comparison of ACE inhibitors and beta blockers without amiodarone) are warranted.

REFERENCES

4. Mehta PK, Griending KK. Angiotensin II


