

# Angiotensin II receptor blockers in the prevention of complications from atrial fibrillation

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**Abstract:** Atrial fibrillation (AF) is the most common form of cardiac arrhythmia and is associated with an increased risk of cardiovascular morbidity and mortality, especially due to ischemic stroke. The occurrence of AF leads to atrial electrical and structural remodeling. The renin-angiotensin system appears to play a role in the development of atrial arrhythmias by its involvement in both of these processes. Large-scale hypertension trials and heart failure trials have indicated the potential value of angiotensin II receptor blockers in the treatment of AF.

**Keywords:** angiotensin II receptor blockers, atrial fibrillation, cardiac arrhythmia

## Introduction

Atrial fibrillation (AF) is the most common clinically significant cardiac arrhythmia. Overall, it affects >1% of the population and is common in the elderly, with a prevalence of almost 10% in individuals aged over 75 years.<sup>1</sup> Currently, approximately 2.3 million people in the United States are diagnosed with AF, and this number is expected to rise to 5.6 million by 2050.<sup>2-4</sup>

A 38-year follow-up review of data from the Framingham Study has indicated that men have a 1.5-fold greater risk of developing AF than women and that hypertension and diabetes are significant independent predictors of AF, after adjustment for age and other predisposing conditions. Hypertension was shown to be the most common, independent, and potentially modifiable risk factor for AF,<sup>5,6</sup> primarily because of its high prevalence in the population.<sup>5</sup> The cardiac conditions that pose the greatest risk for development of AF include heart failure, myocardial infarction, and valvular heart disease (especially mitral valve disease). In addition, other cardiac conditions that have been associated with the occurrence of AF include myocarditis, hypertrophic cardiomyopathy, congenital heart disease, pericarditis, hypertensive cardiovascular disease, left ventricular hypertrophy, and coronary artery disease.<sup>5,7</sup> Obesity was not associated with the incidence of AF in the review of Framingham data.<sup>5</sup> However, it has been proposed as a risk factor for the development of AF in relation to associated left atrial dilation.<sup>8</sup>

The development of AF has been associated with a 1.5- to 1.9-fold greater mortality risk, after adjusting for preexisting cardiovascular conditions, in the original cohort of the Framingham Study.<sup>9</sup> Much of the morbidity and mortality associated with AF is due to thromboembolic complications resulting in ischemic stroke, especially in the elderly.<sup>10</sup> Ischemic stroke incidence among patients with nonvalvular AF averages 5% per year and is approximately 2 to 7 times greater than in individuals without AF.<sup>11</sup>

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The occurrence of AF leads to atrial electrical and structural remodeling, which both contribute to the maintenance and recurrence of the arrhythmia.<sup>12–18</sup> Patients who initially present with paroxysmal AF often progress to longer, non-self-terminating bouts,<sup>19</sup> and enlarged atria are a major risk factor for its recurrence.<sup>20,21</sup>

Current treatment strategies for AF include the prevention of thromboembolism and either rate-control or rhythm-control strategies, with pharmacotherapy or left atrial ablation as first- or second-line therapy options, respectively.<sup>8</sup> These approaches may produce similar outcomes, but rate control may be preferable because agents used to maintain sinus rhythm can have proarrhythmic effects. However, both approaches have important limitations, including proarrhythmic and negative dromotropic effects, respectively.<sup>8,22,23</sup> In view of this, a preventive strategy may be a more attractive option.<sup>7</sup> The angiotensin-converting enzyme (ACE) inhibitors have been shown to reduce the incidence of AF postmyocardial infarction and in patients with left ventricular dysfunction.<sup>24,25</sup> In addition, in recent years, large-scale hypertension trials (Losartan Intervention For Endpoint reduction in hypertension [LIFE] and Valsartan Antihypertensive Long-term Use Evaluation [VALUE]) and heart failure trials (Candesartan in Heart Failure – Assessment of Mortality and Morbidity [CHARM] and Valsartan Heart Failure Trial [Val-HeFT]) have indicated the potential value of angiotensin II receptor blockers (ARBs) in the treatment of AF.<sup>26–30</sup> The aim of this review is to examine the mechanisms of action and clinical efficacy of ARBs in the prevention of AF.

## Preventive mechanisms of ARBs in AF

The occurrence of uncomplicated AF leads mainly to electrical remodeling due to the high atrial rate (indicated by shortening of refractoriness), whereas structural remodeling develops when AF leads to a hemodynamic burden on the atria.<sup>31</sup> Cellular electrophysiological studies have indicated that marked reductions occur in the densities of the L-type voltage-gated calcium ( $\text{Ca}^{2+}$ ) current,  $I(\text{Ca}_L)$ , the transient outward potassium ( $\text{K}^+$ ) current,  $I(\text{TO})$ , and the ultra-rapid delayed rectifier  $\text{K}^+$  current,  $I(\text{Kur})$  in atrial myocytes from patients with chronic AF. Similar changes in currents have been noted in myocytes from a canine model of AF.<sup>32</sup> In both human and canine AF, the reduction in  $I(\text{Ca}_L)$  may explain the observed decrease in action potential duration and effective refractory period that are characteristic electrophysiological features of the remodeled atria.<sup>32</sup> Thus, it has

been suggested that calcium overload and perturbations in calcium handling play prominent roles in AF-induced atrial remodeling in humans.<sup>32</sup>

In addition, angiotensin II exerts its physiologic effects by activating AT1 and AT2 receptor subtypes. Goette et al noted that the occurrence of AF was associated with down regulation of AT1 and upregulation of AT2.<sup>33</sup> However, recently von Lewinski et al reported data that suggests that the major arrhythmic effects occur secondary to stimulation of the AT1 receptor and thus blockade of the AT1 receptor may be antiarrhythmic.<sup>34</sup>

In contrast, structural remodeling is associated with fibrosis: this is characterized by increased deposition of connective tissue in the atria. The histological substrate of atrial biopsies in patients with lone AF has suggested that the probability of AF increases as the degree of fibrosis increases, and further electrophysiologic modification ultimately leads to self-perpetuation of AF.<sup>31,35–39</sup>

The renin-angiotensin-aldosterone system (RAAS) appears to play a role in the development of arrhythmias by its involvement in both forms of remodeling. For example, angiotensin II has been shown to play a critical role in cardiac remodeling via the promotion of cardiac myocyte hypertrophy and cardiac fibroblast interstitial fibrotic changes associated with left ventricular hypertrophy, postmyocardial infarction remodeling, and congestive heart failure (CHF).<sup>40</sup> In particular, angiotensin II has been shown to mediate cardiac myocyte hypertrophy directly via induction of immediate early genes through a mitogen-activated protein kinase (MAPK)-dependent pathway. In addition, it has also been shown to mediate cardiac hypertrophy indirectly via stimulation of norepinephrine release from cardiac nerve endings and endothelin from endothelial cells and through multiple effects on cardiac fibroblasts.<sup>40–45</sup>

Electrical remodeling is brought about by the shortening of the atrial effective refractory period (AERP) and of the action potential duration.<sup>46–49</sup> The inhibitory effects of an ARB (candesartan) and an ACE inhibitor (captopril) on atrial electrical remodeling induced by rapid pacing have been examined in a canine model of AF.<sup>50</sup> The AERP was measured before, during, and after rapid atrial pacing; infusions of saline, ARB, ACE inhibitor, or angiotensin II were initiated 30 minutes before rapid pacing and continued throughout the study. In the saline and angiotensin II groups, the AERP was significantly shortened during rapid atrial pacing and the rate adaptation of the AERP was lost. In contrast, in the ARB and ACE inhibitor groups, shortening of the AERP after rapid pacing was completely inhibited and the rate adaptation of

the AERP was preserved, which indicated that endogenous angiotensin II may be involved in the mechanism of atrial electrical remodeling.<sup>50</sup>

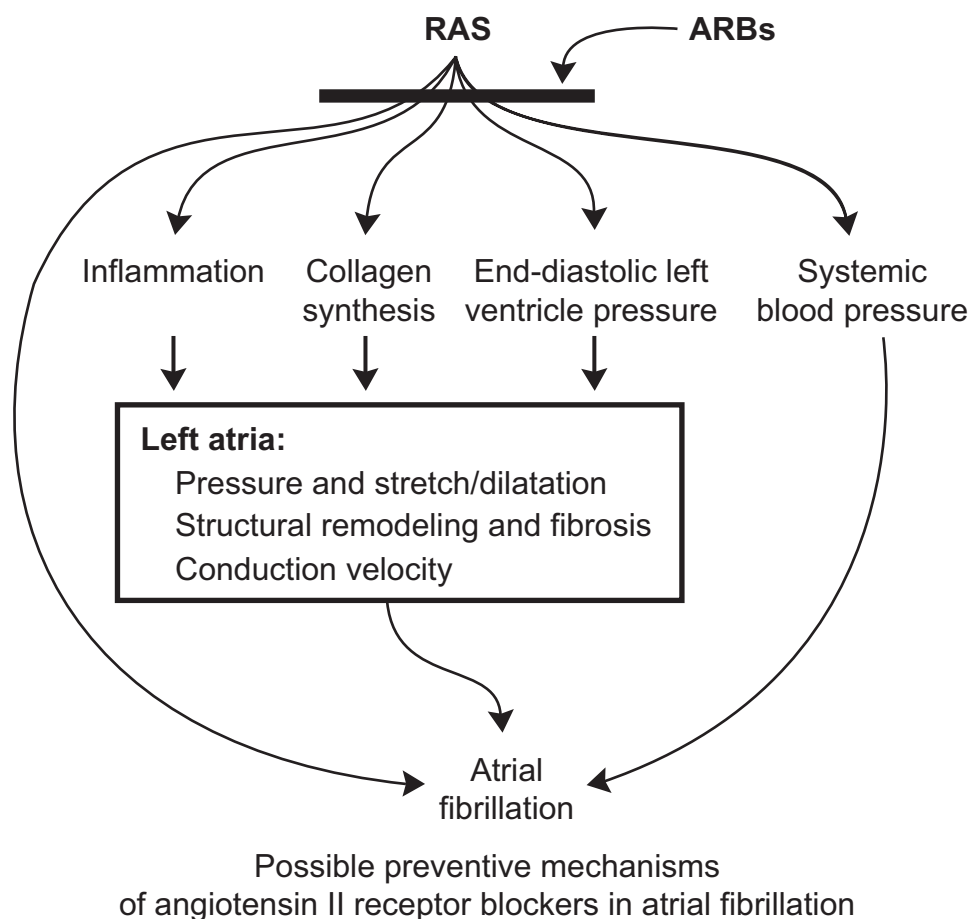
In addition to the reduction of blood pressure per se, there are a range of other potential mechanisms by which inhibition of the RAAS may reduce AF (Figure 1).<sup>51</sup> For example, blockade of the RAAS may prevent left atrial dilatation, atrial fibrosis, dysfunction, and slowing of conduction velocity<sup>44,50,52,53</sup> and, thus, may be particularly effective in patients with left ventricular dysfunction/heart failure.<sup>54</sup> Inhibition of the RAAS has also been shown to reduce inflammation and oxidative stress, and to modulate sympathetic nerve activity.<sup>55</sup> The use of agents that block the RAAS has been shown to increase the efficacy of electrical cardioversion of AF,<sup>56–59</sup> and some studies have also indicated that blockade of the RAAS by ARBs may have direct antiarrhythmic properties.<sup>60,61</sup> Thus, inhibition of the RAAS may lead to improvements in the management of AF.

## Clinical trials with ARBs in AF

A range of clinical and experimental studies have shown that the use of ACE inhibitors or ARBs in CHF reduces the occurrence of AF and AF vulnerability via the reduction of atrial structural remodeling and fibrosis.<sup>44,53,54,62–66</sup>

## ARBs in patients with heart failure

A secondary analysis of the results of the CHARM program, which included 7601 patients with symptomatic CHF, has demonstrated that, in addition to significant reductions in cardiovascular deaths and hospital admissions for heart failure with reduced ejection fraction, the ARB candesartan significantly reduced the development of new AF compared with placebo. Moreover, the benefits of treatment with an ARB were accrued regardless of treatment at baseline (which included treatment with ACE inhibitors) and in a wide spectrum of patients with CHF, including those with preserved as well as reduced left ventricular systolic



**Figure 1** Possible preventive mechanisms of ARBs in atrial fibrillation. Reproduced with permission from Aksnes T, Flaa A, Strand A, et al. Prevention of new-onset atrial fibrillation and its predictors with angiotensin II receptor blockers in the treatment of hypertension and heart failure. *J Hypertens*. 2007;25:15–23.<sup>51</sup> Copyright © Wolters Kluwer Health.

**Abbreviations:** ARBs, angiotensin II receptor blockers; RAS, renin-angiotensin-aldosterone system.

function (Table 1).<sup>67,68</sup> In Val-HeFT, the occurrence of AF was associated with worse outcomes and was evaluated on the basis of adverse event reports in patients with heart failure treated with valsartan vs placebo on top of optimal heart failure therapy, which included ACE inhibitors and beta-blockers.<sup>69,70</sup> Treatment with valsartan significantly reduced the combined endpoint of mortality and morbidity and improved clinical signs and symptoms in patients with heart failure, and also significantly reduced new-onset AF and reduced the relative risk of developing AF by 37% compared with placebo, when added to prescribed therapy. However, when ECG recordings from patients with sinus rhythm at enrolment were evaluated at 4, 12, and 24 months and at study end, the presence of at least 1 episode of AF showed no statistically significant difference between valsartan-treated patients and the placebo group (3.9% and 4.5%, respectively;  $P = 0.15$ ) (Table 1).<sup>69,70</sup>

## ARBs in the treatment of AF after cardioversion

The efficacy of treatment with irbesartan in maintaining sinus rhythm after cardioversion has been evaluated in patients with persistent AF.<sup>56</sup> Patients were divided into 2 groups: group 1 was treated with amiodarone, group 2 with amiodarone plus irbesartan, and both groups underwent electrical cardioversion after 3 weeks of amiodarone administration. All patients started amiodarone after at least 3 weeks of anticoagulation to achieve an international normalized ratio greater than 2, and the primary endpoint of the study was the length of time to the first recurrence of AF. After 2 months of follow-up, the amiodarone plus irbesartan group had significantly fewer recurrences of AF (63.2%) than the amiodarone-only group (84.8%,  $P = 0.008$ ), and had a greater probability of maintaining sinus rhythm (79.5% vs 55.9%,  $P = 0.007$ ).<sup>56</sup> There was

also a trend toward a lower number of shocks and lower electrical threshold, but this did not reach statistical significance.<sup>56</sup> Furthermore, in a subsequent study, the combination of irbesartan plus amiodarone decreased the rate of AF recurrence in a dose-dependent manner in lone AF patients.<sup>57</sup>

## ARBs in the prevention of new AF

In the Study on Cognition and Prognosis in the Elderly (SCOPE), there was a marked reduction in nonfatal stroke compared with placebo in elderly patients with hypertension who received treatment with an ARB, which may have been due to a reduction in AF (Table 2).<sup>71</sup>

Similarly, the results of the LIFE study indicated that ARB-based treatment produced a 25% reduction in fatal and nonfatal stroke compared with atenolol-based treatment in hypertensive patients with left ventricular hypertrophy. This was attributable in part to a 45% lower rate of stroke (24.1 vs 46.5 strokes per 1000 patient-years of follow-up) on ARB treatment in patients with a history of AF,<sup>29</sup> but may have also been a reflection of benefits from a reduced incidence of new-onset AF. In a subsequent subanalysis of ECG data from patients without AF by history at baseline, new-onset AF occurred in 150 patients randomized to losartan vs 221 to atenolol (6.8 vs 10.1 per 1000 person-years; relative risk [RR] 0.67,  $P < 0.001$ ) despite similar levels of blood pressure reduction (Table 2).<sup>28</sup> Patients who received losartan tended to stay in sinus rhythm longer (mean, 1809 vs 1709 days from baseline,  $P = 0.057$ ) than those who received atenolol. Patients with new-onset AF had 2-, 3-, and 5-fold increased rates, respectively, of cardiovascular events, stroke, and hospitalization for heart failure, but there were fewer composite endpoints and strokes in patients who developed new-onset AF in the losartan arm compared to the atenolol treatment arm of the study.<sup>28</sup>

**Table 1** The effects of ARBs on AF and outcomes in clinical trials in patients with left ventricular dysfunction/congestive heart failure

Study	Design/follow-up	N	Interventions	AF-related endpoints
CHARM <sup>67,68</sup>	MC, R, DB, PL Mean, 37.7 months	7601	Candesartan (Can) PL	Incidence of AF 392/6379 (6.15%) of patients with no AF at baseline developed AF during follow-up Can 5.55% vs PL 6.74% (OR 0.802, $P = 0.039$ )
Val-HeFT substudy <sup>69,70</sup>	MC, R, DB, PL Mean, 23 months	4395	Valsartan (Val) PL	Incidence of AF 287/4395 (6.35%) of patients with sinus rhythm at baseline had AF as an adverse event during follow-up Val 5.12% vs PL 7.95% ( $P = 0.0002$ )

**Note:** Trial acronyms are expanded in the text.

**Abbreviations:** AF, atrial fibrillation; ARB, angiotensin II receptor blocker; DB, double-blind; MC, multicenter; OR, odds ratio; PL, placebo-controlled; R, randomized.

**Table 2** Studies of the effectiveness of ARBs in the prevention of AF (new onset and recurrent)

Study	Design/follow-up	N	Interventions	AF-related endpoints
<b>New-onset AF</b>				
SCOPE <sup>71</sup>	MC, R, DB, hypertension (elderly)	4964	Candesartan (Can) PL	Incidence of nonfatal stroke Can 7.4 vs PL 10.3/1000 patient-years (risk reduction 27.8%, $P = 0.04$ )
	Mean, 3.7 years			
LIFE <sup>28</sup>	MC, R, DB, hypertension/LVH	8851 <sup>a</sup>	Losartan (Los)	Incidence of AF
	Mean, 4.8 years		Atenolol (At)	Los 6.8 vs At 10.1/1000 patient-years (RR 0.67, $P < 0.001$ )
				Maintenance of sinus rhythm Los 1809 ± 225 days vs At 1709 ± 254 days ( $P = 0.057$ )
VALUE <sup>27,30</sup>	Retrospective analysis of MC, R, DB study (hypertension)	15,245	Valsartan (Val) Amlodipine (Aml)	Incidence of new-onset AF Val 3.7% vs Aml 4.3% ( $P = 0.044$ ) Rate of persistent AF Val 1.4% vs Aml 2.0% ( $P = 0.005$ )
	Mean, 4.2 years			
ONTARGET <sup>72</sup>	MC, R, DB, patients at high risk of vascular events	25,620	Telmisartan (Tel) Ramipril (Ram)	Incidence of new-onset AF Tel 6.7% vs Ram 6.9% vs Ram ± Tel 6.5% (all $P = NS$ between treatments; Tel vs Ram RR 0.97)
	Median, 56 months		Ram ± Tel combination therapy	
<b>Recurrent AF</b>				
Fogari et al <sup>73</sup>	R, open-label, hypertension/type 2 diabetes/AF	296	Valsartan (Val) Atenolol (At)	Incidence of recurrent AF Val ± Aml 20.3% vs At + Aml 34.1% ( $P < 0.01$ )
	1 year		(± Amlodipine [Aml])	
Fogari et al <sup>74</sup>	R, DB, hypertension/AF	369	Valsartan (Val) Ramipril (Ram) Amlodipine (Aml)	Incidence of recurrent AF Val 16.1% vs Ram 27.9% vs Aml 47.4% ( $P < 0.01$ Val vs Aml and $P < 0.05$ Val vs Ram)
	1 year			
GISSI-AF <sup>75</sup>	MC, R, DB, PL, AF	1442	Valsartan (V) PL	Incidence of recurrent AF Val 51.4% vs PL 52.1% (HR 0.99), but trend favored Val in patients with CHF and/or LV dysfunction (HR 0.81)
	Median, 1 year			
CAPRAF <sup>76,77</sup>	R, DB, PL, AF	171	Candesartan (Can) PL	Incidence of recurrent AF Can 71% vs PL 65% ( $P = 0.20$ ) in patients with persistent AF who underwent ECV
	6 months			

**Note:** <sup>a</sup>No AF at baseline.

**Abbreviations:** AF, atrial fibrillation; ARB, angiotensin II receptor blocker; CHF, congestive heart failure; DB, double-blind; ECV, electrical cardioversion; HR, hazard ratio; LVH, left ventricular hypertrophy; MC, multicenter; PL, placebo; R, randomized; RR, relative risk. Trial acronyms are expanded in the text.

In the VALUE trial, new-onset AF was a secondary prespecified endpoint and ECG recordings were obtained every year and centrally analyzed. During the study, the incidence of at least 1 documented occurrence of new-onset AF was significantly lower in the valsartan treatment group (3.7%) than in the amlodipine treatment group (4.3%, odds ratio 0.84) (Table 2). The incidence of persistent AF was also significantly lower with valsartan than with amlodipine

(odds ratio 0.68, Table 2), and the effects of valsartan on the incidence of AF remained significant even when potential confounding covariates (age, history of coronary artery disease, left ventricular hypertrophy) were taken into account.<sup>27,30</sup>

In addition, the noninferiority of ARBs compared to ACE inhibitors in the prevention of new-onset AF was demonstrated in the ONTARGET study (Ongoing Telmisartan



Alone and in Combination With Ramipril Global Endpoint Trial) in patients with vascular disease or high-risk diabetes (Table 2).<sup>72</sup> In this study, which compared the effectiveness of telmisartan and ramipril in reducing cardiovascular death, myocardial infarction, stroke, and hospitalization for heart failure in patients at risk, the diagnosis of new-onset AF was a secondary endpoint.<sup>72</sup> Telmisartan treatment produced a greater reduction in blood pressure than ramipril and, after a median follow-up of 56 months, the primary outcome of death from cardiovascular causes, myocardial infarction, stroke, or hospitalization for heart failure had occurred in 1412 patients in the ramipril group (16.5%) vs 1423 patients in the telmisartan group (16.7%, RR 1.01). However, the combination of the 2 drugs was associated with a greater frequency of adverse events without an increase in benefit.<sup>72</sup>

## ARBs in the prevention of recurrent AF

Patients with hypertension and type 2 diabetes have an elevated risk of cardiovascular events and typically require combination antihypertensive therapy to achieve goal blood pressure levels. In view of this, Fogari et al<sup>73</sup> compared the effectiveness of valsartan + amlodipine and atenolol + amlodipine combinations on the prevention of AF recurrence in hypertensive patients with well-controlled type 2 diabetes in sinus rhythm who had at least 2 ECG-documented episodes of AF in the previous 6 months. Significantly fewer patients who were followed up for 1 year had recurrent AF with an ARB in addition to antiarrhythmic agents than with atenolol (Table 2).<sup>73</sup> Overall, despite similar levels of blood pressure reduction, the valsartan + amlodipine combination was more effective in preventing AF in patients treated with amiodarone or propafenone.<sup>73</sup> In addition, ARB therapy with valsartan led to significantly fewer recurrences of AF after 12 weeks of treatment than amlodipine, and significantly less recurrence than both amlodipine and ramipril after 1 year in patients with mild hypertension and a history of AF (Table 2).<sup>74</sup> Although blood pressure lowering could be a beneficial mechanism of action for the reduction of AF with both types of RAAS inhibitors, there were no statistically significant differences in blood pressure between the 3 treatment groups in this study. This suggested that both ARBs and ACE inhibitors may exert an antiarrhythmic effect beyond their effects on blood pressure.<sup>74</sup> However, no statistically significant reduction in the recurrence of AF was noted with valsartan treatment compared with placebo in the recent GISSI-AF study (*Gruppo Italiano per lo Studio della Sopravvivenza nell'Infarto Miocardico*: Use of Valsartan and Angiotensin II AT1-Receptor Blocker in the Prevention of

Atrial Fibrillation Recurrence).<sup>75</sup> This may have been due to issues relating to the study design, broad selection criteria, and/or the patients' relatively low-risk clinical conditions (and possible consequent low level of RAAS activation), and optimized background therapy. Moreover, the use of candesartan had no influence on echocardiographic variables or on the recurrence rate of AF after cardioversion in the 6-month Candesartan in the Prevention of Relapsing Atrial Fibrillation (CAPRAF) study.<sup>76,77</sup>

## Ongoing trials of ARBs in AF

It is hoped that a range of ongoing and planned future studies will help to clarify the possible beneficial effects of treatment with ARBs in terms of prevention of AF complications and prevention of AF recurrence (Table 3). The Atrial Fibrillation Clopidogrel Trial with Irbesartan for Prevention of Vascular Events (ACTIVE I) trial will randomize 9000 patients with a history of AF to receive irbesartan or placebo with a planned mean follow-up of 3 years. ACTIVE I is part of the ACTIVE trial program investigating the effect of irbesartan on the occurrence of stroke, myocardial infarction or vascular death, hospitalization for heart failure (primary endpoints) and recurrence of AF, development of cardiac structural remodeling, microalbuminuria, cognitive function, and quality of life (secondary endpoints).<sup>78</sup> In addition, the Angiotensin II Antagonist in Paroxysmal Atrial Fibrillation (ANTIPAF) trial will compare the effect of olmesartan and placebo in patients with paroxysmal AF,<sup>79</sup> and the aim of the planned I-PACE trial is to investigate the effects of irbesartan in hypertensive patients with pacemakers who have recurrent atrial high-rate episodes (Irbesartan for the Prevention of Atrial Arrhythmias and Cardiac Electrical Remodeling in Patients With Hypertension and Permanent Pacemakers).<sup>80</sup>

## Conclusions and clinical recommendations

The prevention of AF is particularly challenging because of its high incidence.<sup>1,3,81</sup> Because ARBs block the actions of angiotensin II, it seems likely that they interfere with structural and electrical remodeling and consequently provide benefits in terms of prevention of new-onset AF and recurrence of AF. So far, the body of evidence supports that ARBs may be useful in combination with other antiarrhythmic agents and not as sole antiarrhythmic therapy in suppressing AF recurrences. However, while the possibility of hemodynamic benefits and direct antiarrhythmic effects cannot be excluded, it is hoped that the results of ongoing prospective studies, such as ACTIVE I and I-PACE, will clarify

**Table 3** Summary of ongoing prospective trials of ARBs in patients with AF

Study	Design/follow-up	N	Interventions	AF-related endpoints
ACTIVE I <sup>78</sup>	Partial factorial, DB, PL, AF Mean, 3 years	9000	Irbesartan (Irb) PL Open clopidogrel plus ASA or oral anticoagulation	Incidence of recurrent AF
ANTIPAF <sup>79</sup>	MC, R, DB, PL, AF 1 year	422 <sup>a</sup>	Olmesartan (Ol) PL	Incidence of pAF (Percentage of days with documented episodes of pAF on daily transtelephonic tele-ECG recordings)
I-PACE <sup>80</sup>	MC, R, DB, PL, hypertensive patients with permanent pacemakers and risk factors for developing AF	200	Irbesartan (Irb) PL	Time to recurrent AHRE

**Note:** <sup>a</sup>Patients stratified according to beta-blocker use.

**Abbreviations:** AF, atrial fibrillation; ARB, angiotensin II receptor blocker; AHRE, atrial high-rate episodes; ASA, aspirin; DB, double-blind; ECG, electrocardiograph; MC, multicenter; pAF, paroxysmal AF; PL, placebo; R, randomized.

the benefits of ARB treatment in relation to AF. Although valsartan was ineffective in suppressing AF in GISSI-AF, this study and the results of ACTIVE I may be limited by the high percentage of patients who had background ACE inhibitor therapy. The results of ACTIVE I may support the use of ARBs related to meaningful endpoints and not surrogates such as AF recurrence. Although it is not currently possible to specifically recommend the use of ARBs for prevention of AF in routine clinical practice, ARBs are well tolerated and may be considered for use in patients with AF and coexisting clinical conditions such as hypertension, coronary heart disease, CHF, and type 2 diabetes.

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