



NOX4-dependent hydrogen peroxide overproduction in human atrial fibrillation and HL-1 atrial cells: relationship to hypertension

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Background/Objectives: Atrial fibrillation (AF) is the most common type of cardiac arrhythmia with patients dying frequently of stroke. In view of the unclear etiologies of AF and a potential role of oxidative stress, the present study examined cardiac reactive oxygen species production and NADPH oxidase (NOX) expression in AF patients. **Methods and Results:** Patients with AF were older than those without (58.8 ± 11.7 vs. 47.8 ± 19.2 , $p = 0.047$). Whereas total $O_2^{\bullet-}$ production (determined by electron spin resonance) was similar in patients with and without AF, H_2O_2 production was more than doubled in AF patients (149.8 ± 26.28 vs. 66.9 ± 7.14 pmol/mg/min, $p = 0.0055$), which correlated well with a doubling in NOX isoform 4 (NOX4) expression. AF patients with co-existing hypertension had three-fold higher H_2O_2 production compared to those without (239.0 ± 125.1 vs. 83.6 ± 51.3 pmol/mg/min, $p = 0.003$). Treatment of HL-1 atrial cells with angiotensin II, a known modulator of atrial structural remodeling, resulted in upregulation of NOX4 and H_2O_2 production, further implicating a potential role of NOX4 in atrial remodeling. **Conclusion:** Our data represent the first implication that NOX4-derived H_2O_2 may play an important role in the etiologies of AF.

Keywords: atrial fibrillation, hydrogen peroxide, NADPH oxidase, NOX4, hypertension, HL-1 cells, angiotensin II

INTRODUCTION

Atrial fibrillation (AF) is the most common type of cardiac arrhythmia that is rapidly developing into an epidemic (Savelieva and Camm, 2008). It is associated with markedly increased risk of stroke, due to dislocation of thrombi that are most often originated from left atrial appendage (LAA). Although AF is clearly associated with aging, and cardiovascular conditions such as hypertension, mitral valve stenosis, and heart failure, molecular mechanisms underlying its etiology have remained elusive. Recent advances seem to suggest that AF is initiated by atrial structural and electrical remodeling, and that AF itself further augments these responses to perpetuate AF (Savelieva and Camm, 2008; Chou and Chen, 2009).

The known risk factors for AF have all been linked to oxidative stress, and inflammatory factors such as angiotensin II (Ang II) (Kannel et al., 1998; Nakashima et al., 2000; Kim et al., 2003, 2005). Ang II is a major vasoconstrictive hormone and its production is elevated in patients with hypertension. It is also involved in modulating cardiac hypertrophy and remodeling. Of note, hypertension is considered the most prevalent risk factor for AF. Atrial structural remodeling can be induced by Ang II, resulting in atrial fibrosis and possibly initiation of AF (Zografos and Katritsis, 2010; Tan and Zimetbaum, 2011). A meta-analysis of 7 clinical trials involving a

total of 24,849 patients receiving ACEIs and/or ARBs for treatment of hypertension, ischemic heart disease, heart failure, or diabetes mellitus demonstrated that ACEIs/ARBs significantly reduces the risk of AF or its recurrence by 28% (Healey et al., 2005). Importantly, Ang II is an extremely potent activator of NADPH oxidase (NOX) in the vascular cells (Cai et al., 2003; Choi et al., 2008; Lassegue and Griendling, 2010). NOX-dependent reactive oxygen species (ROS) production leads to nitric oxide (NO^{\bullet}) inactivation and endothelial dysfunction, which underlie pathogenesis of vascular diseases such as hypertension and atherosclerosis (Cai and Harrison, 2000; Cai et al., 2003; Cai, 2005a,b). We hypothesize that NOX-derived ROS might be important in mediating Ang II-dependent development of AF. Some interesting data exist to show that hydrogen peroxide increases Kv1.5 current amplitude at voltages corresponding to the action potential repolarization phase and accelerate Kv1.5 channel opening. These changes can reduce the action potential duration, leading to a shortening of the atrial effective refractory period (Caouette et al., 2003).

To test this hypothesis in human AF, cardiac tissues from heart transplant patients were examined for ROS production and expression of NOX isoforms. Interestingly, we found a specific upregulation of NOX4 that was correlated with H_2O_2 overproduction in AF patients. None of the other NOX isoforms were however

regulated in AF. AF patients with co-existing hypertension had more than 3-fold increase in H₂O₂ production compared to those without. In atrial HL-1 cells, Ang II treatment led to increased AT1 receptor expression, upregulation of NOX4 and increased H₂O₂ production. These human and cell culture data together implicate that there might be a novel role of NOX4-derived H₂O₂ in mediating Ang II-dependent AF development.

MATERIALS AND METHODS

PATIENT POPULATION

Left atrial appendage tissues isolated from randomly recruited patients undergoing cardiac transplant surgeries were subjected to electron spin resonance (ESR) determination of O₂^{•-} production, Amplex red determination of H₂O₂ production and RT-PCR analysis of NOX gene expression. The study was carried out in a double-blinded fashion. Clinical information was gathered upon completion of laboratory analyses. IRB approval was obtained for these studies.

ELECTRON SPIN RESONANCE DETECTION OF SUPEROXIDE RADICAL

The specific O₂^{•-} spin trap CMH (0.5 mmol/L; Noxygen) solution was prepared freshly in nitrogen gas bubbled Krebs/HEPES buffer containing diethyldithiocarbamic acid (DETC; 5 μmol/l; Sigma) and deferoxamine (25 μmol/l; Sigma). The LAA homogenates (in 50 mmol/L Tris-HCl, pH 7.4, 0.1 mmol/L EDTA, 0.1 mmol/L EGTA, and 1% (v/v) of protease inhibitor cocktail from Sigma) were then mixed with the spin trap solution and loaded into glass capillary (Fisher Scientific) for analysis of O₂^{•-} production using ESR spectrometer (Bruker), following a similar procedure as we previously published for cultured endothelial cells (Chalupsky and Cai, 2005; Gao et al., 2009a,b) intact mouse aortas (Chalupsky and Cai, 2005; Oak and Cai, 2007; Gao et al., 2009a), and tissue homogenates (Lu et al., 2009). The ESR settings used were as follows: bio-field 3,410, field sweep 100G, microwave frequency 9.73 GHz, microwave power 13.26 mW, modulation amplitude 9.82G and 512 points of resolution.

AMPLEX-RED ASSAY OF HYDROGEN PEROXIDE PRODUCTION

The LAA lysates [in 20 mmol/L Tris-HCl, pH 7.4, 150 mmol/L NaCl, 1 mmol/L EGTA, 1 mmol/L EDTA, 1% (v/v) Triton X-100, 2.5 mmol/L sodium pyrophosphate, 1 mmol/L Na₃VO₄, 1 mmol/L β-glycerolphosphate, 1 mmol/L PMSF, 1% (v/v) of protease inhibitor cocktail from Sigma] were analyzed for H₂O₂ production using a fluorometric horseradish peroxidase assay (Amplex-Red assay; Molecular Probes). Fluorescence was measured (excitation 530 nm and emission 590 nm) after 1 h of incubation at 37°C in the dark against background fluorescence of buffer. Polyethylene glycol-conjugated catalase (500 U/ml, Sigma) inhibitable fraction reflects specific H₂O₂ signal. The rate of H₂O₂ production was presented as picomoles per milligram protein per minute after calculation, according to a standard curve generated using fresh H₂O₂ in reaction buffer.

RT-PCR ANALYSIS OF NOX mRNA EXPRESSION

Total RNA was extracted from the LAA using TRIzol (Invitrogen) according to the manufacturer's instructions. Reverse transcription was performed in standard fashion with iScript cDNA synthesis Kit (Bio-Rad). The primer sequences used for PCR reactions

are summarized in **Table 1**. The PCRs were carried out using an iCycler (Bio-Rad) with the protocol of [(95°C/2 min) (95°C/25 s, 57°C/5 s, 68°C 5 min) × 35 (72°C/10 min)]. The PCR mix contained GAPDH primers to generate GAPDH amplicon that served as an internal control.

HL-1 CELL CULTURE AND STIMULATION BY ANG II

The HL-1 cells were obtained as generous gifts from Dr. William Claycomb from Louisiana State University that have been extensively characterized to maintain its differentiated cardiac phenotype using microscopic, genetic, immunohistochemical, electrophysiological, and pharmacological techniques (Claycomb et al., 1998; White et al., 2004). These beating cells were cultured following exact protocols as established by the Claycomb group and found to keep their phenotype during the 24 h exposure period to Ang II. The confluent cells were kept quiescent in media containing 0.01% FBS overnight and then stimulated with Ang II (100 nmol/L) for 24 h prior to analysis of NOX4 protein expression by Western blot (primary antibody from Abcam) and H₂O₂ production by Amplex Red assay. NOX4 mRNA expression from identically treated cells was determined by real-time RT-PCR.

STATISTICAL ANALYSIS

All data are presented as mean ± SEM. Differences in O₂^{•-} production, H₂O₂ production, and mRNA expression of different NOX isoforms between AF and non-AF groups were compared using *t* tests. ANCOVA were performed to examine the correlation between H₂O₂ and NOX4 mRNA expression in patients with and without AF. Statistical significance was set as *p* < 0.05.

RESULTS

PATIENT CHARACTERISTICS

Patient characteristics are summarized in **Table 2**. Eighteen patients with AF were older than those without (*n* = 17; *p* = 0.047). The gender distribution was not different. Both groups had similar incidence of co-existing cardiovascular conditions including hypertension, diabetes, and coronary artery disease. The percentages of patients receiving treatment of ACEI, AT1RB, calcium blocker or beta blocker were also similar between the two groups (**Table 2**). Of note, there was no significant difference in ejection fraction between the AF and non-AF groups. Whereas

Table 1 | Primer sequences for RT-PCR.

Gene		Sequence (5'→3')
NOX1	Forward	CAATCTCTCCTGGAATGGCATCCT
	Reverse	CCTGCTGCTCGGATATGAATGGAGAA
NOX2	Forward	AAGGCTTCAGGTCCACAGAGGAAA
	Reverse	AGACTTTGTATGGACGGCCAACT
NOX3	Forward	ACCGTGGAGGAGGCAATTAGACAA
	Reverse	CAGGTTGAAGAAATGCGCCACGAT
NOX4	Forward	AGCAGAGCCTCAGCATCTGTCTCT
	Reverse	TGGTTCTCCTGCTTGGAACTTCT
NOX5	Forward	CCTCCTCATGTTTCATCTGCTCCAGTT
	Reverse	AGGAGGTAGGACAGGTGAGTCCAATA

Table 2 | Patient characteristics.

Characteristic	Non-AF	AF	p Value
Age	47.8 ± 19.2	58.8 ± 11.7	0.047
Gender (%male)	70%	61%	0.568
Hypertension	59%	39%	0.251
Diabetes	29%	28%	0.917
CAD	41%	33%	0.643
Family history of heart disease	65%	44%	0.409
ACEI Therapy	53%	44%	0.627
AT1RB Therapy	29%	17%	0.384
Ca ⁺⁺ blocker therapy	0%	11%	0.166
Beta blocker therapy	76%	55%	0.203
SBP (mmHg)	107.1 ± 18.3	120.8 ± 13.9	0.033
DBP (mmHg)	66.6 ± 12.2	73.8 ± 8.3	0.074
Ejection fraction (%) by ECHO)	20.82 ± 2.3	22.67 ± 2.5	0.586
TC (mg/dL)	158.6 ± 33.6	158.0 ± 44.1	0.967
TG (mg/dL)	137.8 ± 57.2	144.5 ± 53.6	0.266

CAD, coronary artery disease; ACEI, angiotensin converting enzyme inhibitor; AT1RB, angiotensin II type-1 (AT1) receptor blocker; SBP, systolic blood pressure; DBP, diastolic blood pressure; TC, total cholesterol; TG, triglycerides.

diastolic blood pressure, total cholesterol and triglyceride levels were not different in the AF vs. non-AF groups, systolic blood pressure (SBP) was higher in the AF patients (Table 2). ACEI-treated AF group had modestly higher SBP than those of non-AF group receiving ACEI (Table 3).

REACTIVE OXYGEN SPECIES PRODUCTION

Among biologically relevant and abundant ROS, O₂^{•-}, and H₂O₂ appear to be most important in redox signaling. Whereas O₂^{•-} predominantly induces endothelial dysfunction by rapidly inactivating nitric oxide (NO[•]), H₂O₂ influences different aspects of endothelial cell function via complex mechanisms. We hypothesize that O₂^{•-} and H₂O₂ also have differential roles in the regulation of cardiac functions. Both species were quantitatively and specifically determined by ESR and Amplex Red assay respectively. Whereas total O₂^{•-} production was not different between AF and non-AF groups (Figure 1A), we found that H₂O₂ production was more than doubled in AF patients compared to those without (149.8 ± 26.28 vs. 66.9 ± 7.14 pmol/mg/min, *p* = 0.0055, Figure 1B).

H₂O₂ PRODUCTION IN AF PATIENTS WITH CO-EXISTING HYPERTENSION

Interestingly, AF patients with co-existing hypertension had approximately 3-fold higher H₂O₂ production compared to those without (239.0 ± 125.1 vs. 83.6 ± 51.3 pmol/mg/min, *p* = 0.003, Table 4). Older AF patients (>50) also had higher tissue levels of H₂O₂ compared to those younger AF patients (<50; 165 ± 118.8 vs. 95.0 ± 64.9 pmol/mg/min), although yet reaching statistical significance. In contrast, hypertension or age had no effect on H₂O₂ levels in patients without AF (Table 4).

EXPRESSION OF DIFFERENT NOX ISOFORMS AND ITS RELATIONSHIP TO H₂O₂

NOX4 is known to either produce H₂O₂ directly, or produce O₂^{•-} in subcellular organelles leaving membrane permeable H₂O₂ the sole detectable species (Martyn et al., 2006; Lambeth et al., 2007; Serrander et al., 2007; Chen et al., 2008; Block et al., 2009; Lassegue and Griendling, 2010). Other NOX isoforms however, have been shown to produce O₂^{•-} directly (Choi et al., 2008). Intriguingly, NOX4 mRNA expression was found significantly upregulated in AF patients (Figure 2A), which correlated well with a marked increase in H₂O₂ production (Figure 2B). The expression of NOX1 and NOX2 however, was not different in patients with and without AF (Figures 3A,B). NOX3 and NOX5 were not detectable in LAA. We did not find any correlation between ejection fractions and NOX4/H₂O₂ levels in both groups. This indicates that severity of heart failure is not linked to NOX4-H₂O₂ axis as AF does.

NOX4 EXPRESSION AND H₂O₂ PRODUCTION IN ANG II-STIMULATED HL-1 ATRIAL CELLS

In additional experiments, we examined whether Ang II activates NOX4 in HL-1 atrial cells that have been well characterized to maintain a differentiated, adult cardiomyocyte and atrial cell-like phenotype. In our hand we have also been able to reproduce the culture condition and observe the beating HL-1 cells following the published Claycomb conditions. Of note, the Ang II receptor AT1 was abundantly expressed in these cells, and it was upregulated by Ang II (Figures 4A,C). Ang II stimulation resulted in upregulation of NOX4 protein expression and an associated increase in H₂O₂ production (Figures 4A,B,D). The NOX4 mRNA expression, determined by real-time RT-PCR, was also found upregulated by Ang II in HL-1 cells (Figure 4E).

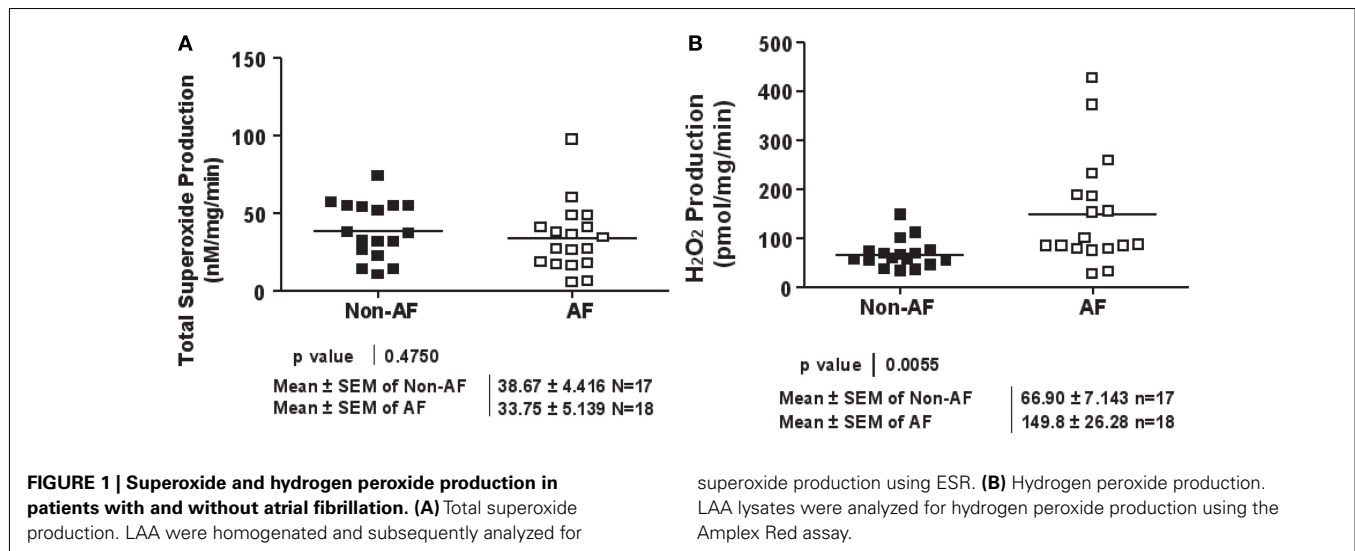
DISCUSSION

In the present study, we have discovered that NOX4-derived H₂O₂ production is markedly increased in the LAA tissues of AF patients. Hydrogen peroxide production in LAA was 3-fold higher in AF patients with co-existing hypertension than those without. Hypertension is an established risk factor for AF. Recent studies have indicated that H₂O₂ regulates electrical properties of isolated rabbit pulmonary vein and atrial cells (Lin et al., 2010). Together with the previously characterized role of H₂O₂ in myocardial fibrosis, these data suggest that NOX4-derived H₂O₂ may play an important role in the development of AF. Indeed, in additional studies utilizing HL-1 atrial cells, we found that Ang II stimulation led to NOX4 upregulation and increased H₂O₂ production, further implicating a potential role of NOX4-derived H₂O₂ in atrial structural remodeling that is considered an important initiating process of AF.

Aging and hypertension are the best-characterized, potential causal factors for AF. Of significant interest, we found that AF patients with co-existing hypertension had 3-folds higher H₂O₂ production compared to those without AF. AF patients that are older (>50) also had apparently higher H₂O₂ production. These observations may further suggest that augmented local cardiac production of H₂O₂ is involved in arrhythmogenesis, besides its proinflammatory and prothrombotic effects. On the other hand,

Table 3 | Blood pressure control in patient subgroups.

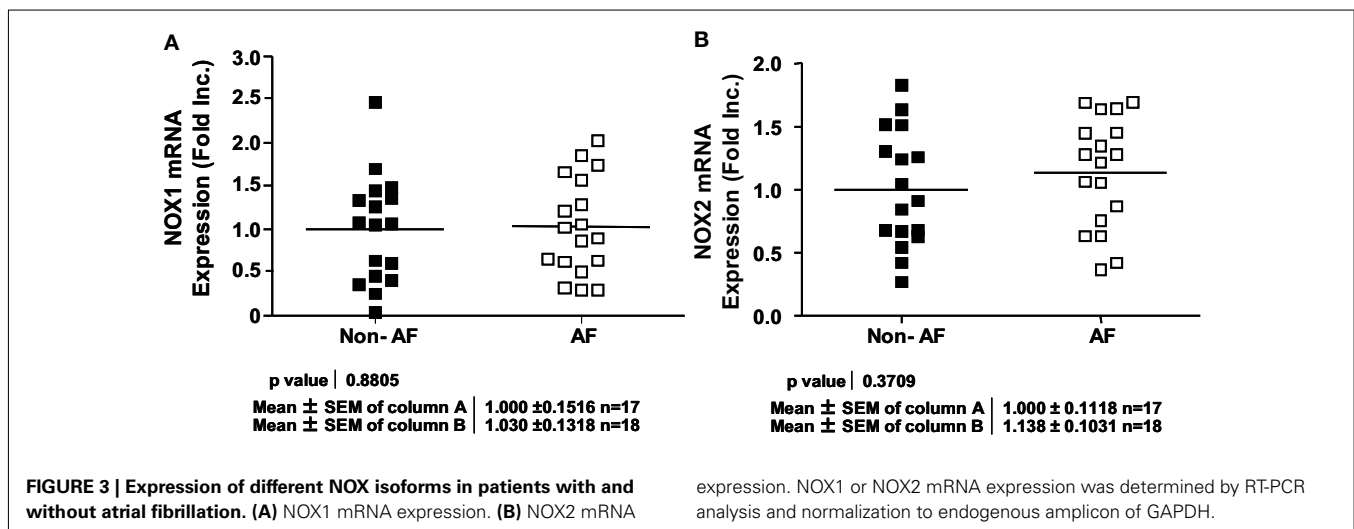
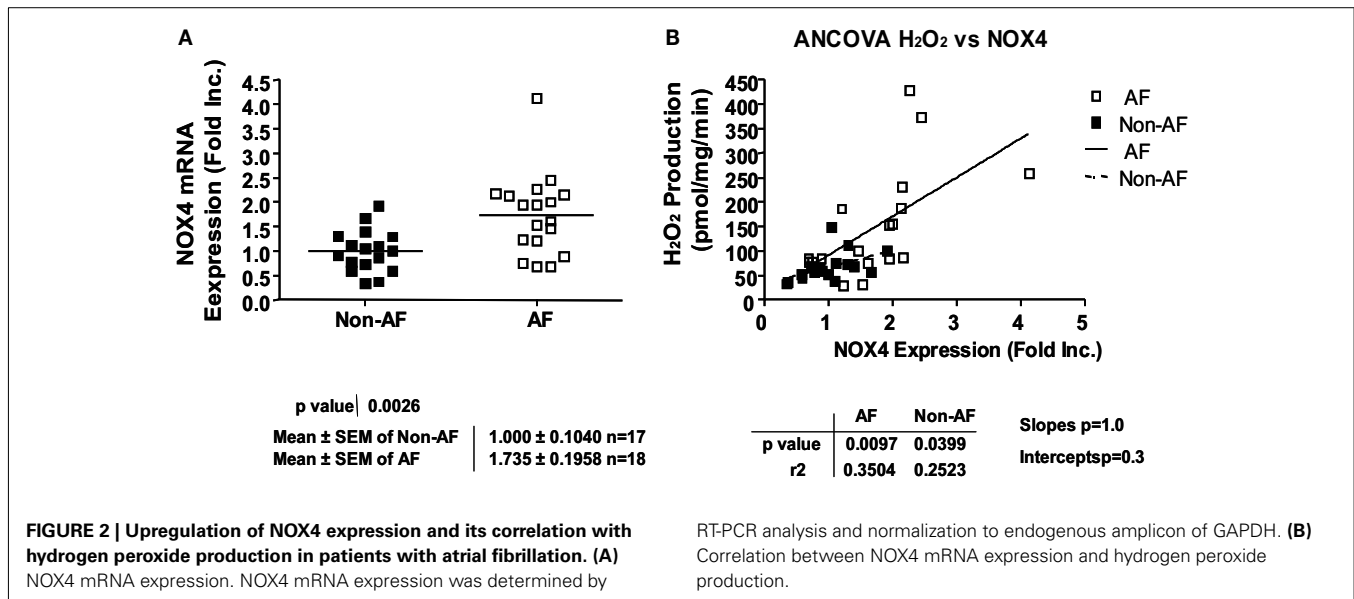
Characteristic	ACEI			Non-ACEI		
	Non-AF (n = 7)	AF (n = 8)	p Value	Non-AF (n = 8)	AF (n = 12)	p Value
SBP	104.1 ± 16.2	119.1 ± 12.4	0.024	110.6 ± 19.4	123 ± 16.8	0.236
DBP	68.6 ± 11.0	74.5 ± 9.4	0.236	70.2 ± 10.3	76.4 ± 4.4	0.233
Characteristic	AT1RB			Non-AT1RB		
	Non-AF (n = 12)	AF (n = 4)	p Value	Non-AF (n = 13)	AF (n = 6)	p Value
SBP	120.8 ± 16.6	131 ± 26.9	0.530	101.7 ± 14.3	119.1 ± 11.9	0.003
DBP	75.1 ± 5.2	78.5 ± 3.5	0.444	67.8 ± 11.4	74.3 ± 7.9	0.120

**Table 4 | Hydrogen peroxide levels in patient subgroups.**

Characteristic	Non-AF			AF		
	Absence	Presence	p Value	Absence	Presence	P Value
Hypertension	58.1 ± 25.7 (n = 6)	71.7 ± 31.4 (n = 11)	0.379	83.6 ± 51.3 (n = 11)	239.0 ± 125.1 (n = 7)	0.003
Diabetes	77.4 ± 33.7 (n = 12)	51.8 ± 12.5 (n = 5)	0.076	151.8 ± 122.7 (n = 13)	144.8 ± 87.3 (n = 5)	0.909
CAD	55.6 ± 13.9 (n = 10)	71.6 ± 33.3 (n = 7)	0.321	155.0 ± 123.8 (n = 12)	139.4 ± 91.6 (n = 6)	0.789
Gender	Male	Female	0.828	Male	Female	0.544
	65.9 ± 32.2 (n = 13)	69.8 ± 21.6 (n = 4)		136.6 ± 117.9 (n = 11)	170.6 ± 105.9 (n = 7)	
Age	<50	>50	0.070	<50	>50	0.278
	53.3 ± 14.6 (n = 8)	79.0 ± 34.5 (n = 9)		95.0 ± 64.9 (n = 4)	165.5 ± 118.8 (n = 14)	

AF may have made SBP control more difficult (ACEI-AF group had higher SBP than ACEI-non-AF group), which is likely related to the substantially higher H₂O₂ production in hypertensive AF patients. Although H₂O₂ can mediate compensatory relaxation under certain conditions such as in the apolipoprotein E deficient atherosclerotic mice (Landmesser et al., 2003), excessive H₂O₂ is involved in vascular remodeling which could in turn result in stiffer blood vessels to deteriorate hypertension.

We found that LAA O₂^{•-} production was not different in patients with and without AF. We also found that xanthine oxidase derived O₂^{•-} was not different between the two groups (data not shown). Xanthine oxidase has been previously shown to play an important role in cardiac O₂^{•-} production in patients with heart failure. However its role in AF patients has never been previously tested. It is interesting to speculate whereas xanthine oxidase and/or other NOX isoforms is more important for O₂^{•-}

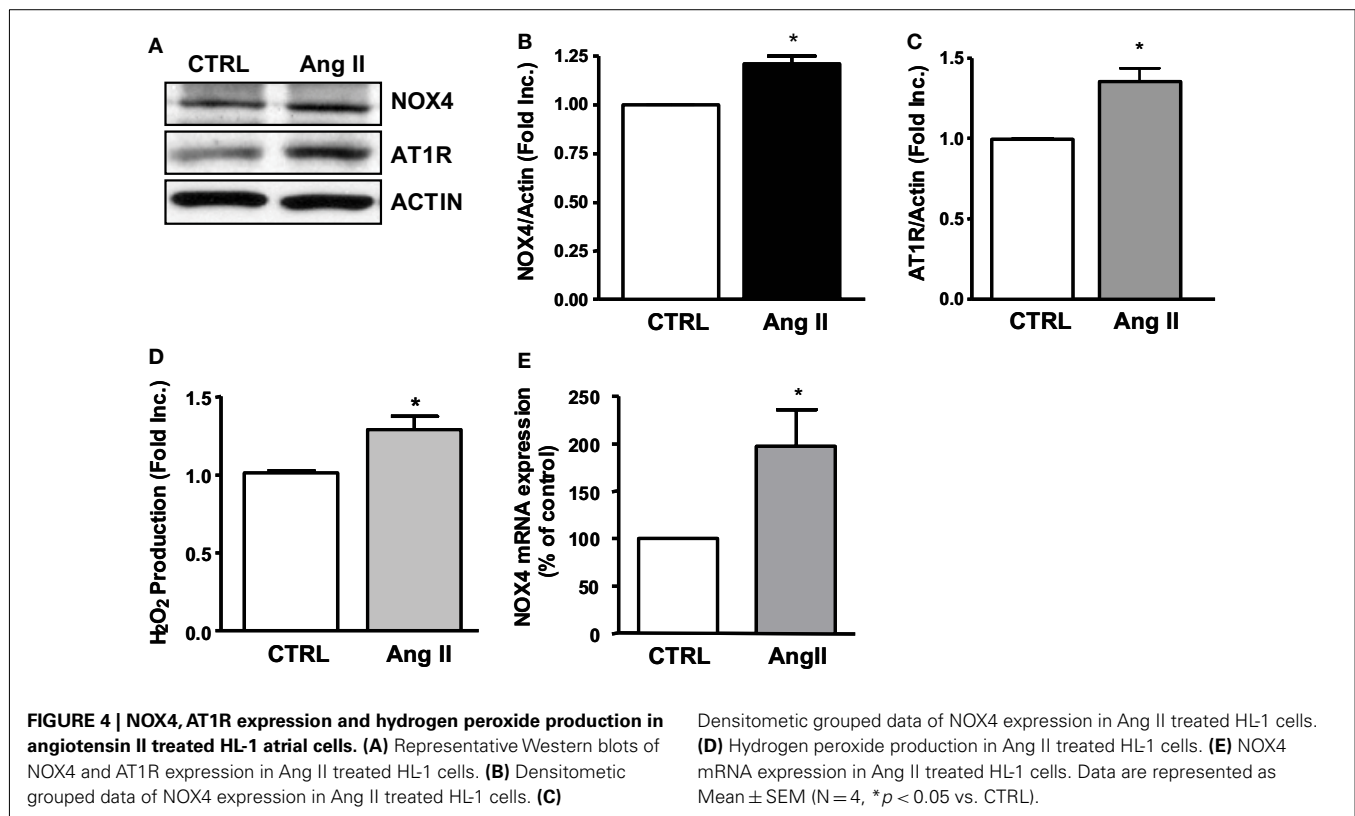


production in heart failure patients, NOX4-derived H₂O₂ production is more important in the condition of AF. Of course, this hypothesis needs to be further investigated using larger patient population. Potential regulations of ROS scavenging enzymes such as superoxide dismutases and glutathione peroxidases, under the environment of NOX4-dependent H₂O₂ overproduction, would also be interesting to be further investigated.

Our data revealed an innovative role of NOX4 in H₂O₂ production in AF patients. Previously NOX1 and NOX4 were found undetectable in RAA (Kim et al., 2005). In human LAA, however, we were able to detect abundant NOX1, NOX2, and NOX4 mRNA expression. Only NOX4 mRNA expression was upregulated by AF, which correlated well with overproduction of H₂O₂ in AF patients. Accumulating evidence have characterized the relatively newly recognized NOX4 and its ROS-producing activity in various cell types such as endothelial cells (Ago et al., 2004, 2005), vascular smooth muscle (Lassegue et al., 2001; Wingler et al., 2001; Touyz et al., 2002), and kidney cells (Shiose et al., 2001; Wingler

et al., 2001; Chabrashvili et al., 2002). Functionally, NOX4 has been implicated in cell differentiation (Clemus et al., 2007; Ismail et al., 2009; Schroder et al., 2009; Xiao et al., 2009), cell cycle transition (Sturrock et al., 2007), transcriptional activation (Bonello et al., 2007), and myofibroblast activation and fibrogenic responses (Hecker et al., 2009). In the heart, NOX4 has been shown to be involved in hypertrophy induced cardiac apoptosis and mitochondrial dysfunction (Ago et al., 2010), and pressure overload-induced heart failure (Byrne et al., 2003; Kuroda et al., 2010). Our data add to this rapidly growing knowledge of NOX4 functions, demonstrating a potential role of NOX4 in enhancing H₂O₂ production in patients with AF, which might be involved in the pathogenesis of AF. Of note, recent studies have demonstrated an important role of myocardial fibrosis in AF (Everett and Olgin, 2007). Therefore, H₂O₂ may also contribute to AF via augmentation of myocardial fibrosis as recently reported (Morita et al., 2009).

HL-1 cardiomyocytes, derived from atrial cardiac muscle cells from the AT1 mouse atrial cardiomyocyte tumor lineage,



maintains the differentiated adult cardiomyocyte phenotype (Claycomb et al., 1998). This cell has been proved to possess the organized sarcomere, ion channels, functional receptors, and intracellular signaling proteins so that it contracts spontaneously, divides, generates action potential, and responds to agonists (White et al., 2004). Therefore, HL-1 cardiomyocyte has been established as the useful tool to study the cellular and molecular mechanisms underlying cardiac pathophysiology (Rodriguez-Sinovas et al., 2005; Brady et al., 2006; Barba et al., 2009). In particular, expression of renin-angiotensin system components and Ang II regulation of L-type calcium channel have been characterized in HL-1 cells (Tsai et al., 2007, 2008). Moreover, transcriptional profile regulated by rapid electrical stimulation in HL-1 cells was similar to those from human AF (Mace et al., 2009). Therefore, we employed HL-1 cells as the cellular model to investigate whether NOX4 is modulated by Ang II, which has been shown to modulate atrial remodeling and fibrillation. We found AT1R was expressed in HL-1 cells and further upregulated by Ang II treatment. Most importantly, exposure of HL-1 cells to Ang II resulted in increased NOX4 mRNA and protein expression, without affecting NOX1

and NOX2 mRNA levels. This was accompanied by a significant elevation in H₂O₂ production.

In summary, our data have uncovered a NOX4-mediated overproduction of H₂O₂ in patients with AF. AF patients with hypertension had markedly higher LAA H₂O₂ levels compared to those without AF. In addition, H₂O₂ has been recently reported to regulate electrical properties of pulmonary vein and atrial cells, where the initiation of ectopic beats is believed to occur to induce AF. It is also known to promote myocardial fibrosis. The modulator of atrial remodeling and fibrosis, Ang II, also significantly upregulated NOX4 expression and H₂O₂ production in atrial HL-1 cells. Taken together, these data suggest that NOX4 may play an important role in the pathogenesis of AF.

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