Endothelin-1 (ET-1) is a potent vasoconstrictor peptide produced and released by the vascular endothelium. In humans, the vascular actions of ET-1 are mediated by 2 distinct ET receptor subtypes: ET_A receptors located exclusively on vascular smooth muscle and ET_B receptors located on both the vascular smooth muscle and endothelial surfaces. In combination with the endothelial vasodilator nitric oxide, ET-1 plays a central role in the regulation of vascular tone. ET-1 system activity and elevations in BP. Thus, clear links have been established between ET-1 system activity and elevations in BP.

**Clinical Trial Registration**—URL: http://www.clinicaltrials.gov. Unique identifier: NCT01395329.

(Hypertension. 2016;67:1196-1204. DOI: 10.1161/HYPERTENSIONAHA.115.06979.)

**Key Words:** blood pressure ■ endothelin-1 ■ metoprolol ■ nebivolol ■ vasoconstriction
Nebivolol, a third generation β-blocker with high selectivity for β₁-adrenergic receptors, has proven to be effective in treating elevated BP. A distinguishing feature of nebivolol from other β-blockers is its hemodynamic profile, specifically the unique ability to enhance both basal and stimulated nitric oxide release resulting in peripheral vasodilation, improved endothelial function and increased myocardial compliance. Cockcroft et al demonstrated that the vasodilatory effects of nebivolol were attenuated by the infusion of the nitric oxide synthase inhibitor N⁶-monomethyl-L-arginine, indicating that nebivolol-induced improvement in vasodilator function is mediated, in part, by increased nitric oxide bioavailability. However, the favorable vascular effects of nebivolol that contribute to its BP-lowering action may not be limited to nitric oxide. Indeed, there are in vitro data to suggest that nebivolol suppresses endothelial ET-1 production, but there is currently no in vivo clinical evidence that treatment with nebivolol reduces ET-1–mediated vasoconstrictor tone.

Accordingly, we tested the hypothesis that chronic nebivolol treatment will reduce ET-1–mediated vasoconstrictor tone in adult humans with elevated BP. Moreover, reducing ET-1 vasoconstrictor activity contributes to the improvement in endothelial vasodilator function associated with nebivolol treatment. To address this hypothesis, we used a 3-month randomized, double-blind, placebo controlled study to determine the effects of nebivolol, compared with metoprolol and placebo, on ET-1 vasoconstrictor tone in adults with suboptimal BP.

**Methods**

**Subjects**

Forty-two middle-aged adults with elevated BP (systolic BP ≥130 or diastolic BP ≥85 mm Hg) participated in a 3-month, double-blind, randomized, placebo controlled trial: 14 received nebivolol (8 men/6 women; 5 mg per day; Forest Laboratories, Inc, Jersey City, NJ); 14 received metoprolol succinate (9 men/5 women; 100 mg per day; AstraZeneca LP, Wilmington, DE); and 14 received placebo (9 men/5 women; 1 gelatin capsule per day; Forest Laboratories, Inc). The doses of nebivolol and metoprolol were chosen to elicit similar reductions in BP. Resting BP was determined by the average of ≥2 seated BP readings from 2 separate visits per American Heart Association guidelines. All subjects were free of overt coronary and metabolic disease as assessed by medical history, physical examination, fasting blood chemistries, and electrocardiograms and BP at rest and during incremental exercise performed to exhaustion. In addition, all subjects presented with a resting heart rate more than 50 bpm. None of the subjects smoked, were taking medications (including vitamins), or performed regular physical exercise for at least 1 year before the start of the study. All of the women were at least 1-year postmenopausal and had never taken or had discontinued the use of hormone replacement therapy at least 1 year before the start of the study. After baseline testing, subjects were randomly assigned to 1 of the 3 experimental groups. Before participation, all of the subjects had the research study and its potential risks and benefits explained fully before providing written informed consent according to the guidelines of the University of Colorado at Boulder. The study was approved by the Institutional Review Board of University of Colorado, Boulder.

**Measurements**

**Blood Pressure**

Resting BP measurements were performed in the sitting position on at least 2 separate days at least 1 week apart. Subjects were instructed not to

### Table 1. Selected Subject Characteristics

<table>
<thead>
<tr>
<th>Variable</th>
<th>Nebivolol Before</th>
<th>Nebivolol After</th>
<th>Metoprolol Before</th>
<th>Metoprolol After</th>
<th>Placebo Before</th>
<th>Placebo After</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sex, M/F</td>
<td>8/6</td>
<td>8/6</td>
<td>9/5</td>
<td>9/5</td>
<td>9/5</td>
<td>9/5</td>
</tr>
<tr>
<td>Age, y</td>
<td>57±1</td>
<td>57±1</td>
<td>55±1</td>
<td>55±1</td>
<td>56±1</td>
<td>56±1</td>
</tr>
<tr>
<td>Body mass, kg</td>
<td>77.7±2.7</td>
<td>78.4±2.5</td>
<td>88.7±4.8</td>
<td>90.7±4.7</td>
<td>88.7±5.1</td>
<td>89.1±5.2</td>
</tr>
<tr>
<td>BMI, kg/m²</td>
<td>26.6±0.7</td>
<td>26.9±0.7</td>
<td>29.3±1.4</td>
<td>30.1±1.5</td>
<td>28.6±1.2</td>
<td>28.8±1.2</td>
</tr>
<tr>
<td>Body fat, %</td>
<td>33.6±1.9</td>
<td>34.6±1.9</td>
<td>34.6±2.1</td>
<td>35.7±1.9</td>
<td>35.5±2.6</td>
<td>35.9±2.6</td>
</tr>
<tr>
<td>Waist circumference, cm</td>
<td>88.3±2.4</td>
<td>89.0±2.4</td>
<td>95.9±4.0</td>
<td>97.6±3.8</td>
<td>94.2±2.9</td>
<td>94.5±2.7</td>
</tr>
<tr>
<td>VO₂ max, mL/kg per min</td>
<td>28.6±2.2</td>
<td>28.0±1.9</td>
<td>27.5±1.9</td>
<td>27.3±1.8</td>
<td>26.5±1.9</td>
<td>25.5±1.5</td>
</tr>
<tr>
<td>Total cholesterol, mmol/L</td>
<td>5.2±0.2</td>
<td>4.7±0.2</td>
<td>5.1±0.2</td>
<td>4.5±0.2</td>
<td>5.3±0.1</td>
<td>4.9±0.2</td>
</tr>
<tr>
<td>LDL cholesterol, mmol/L</td>
<td>3.1±0.1</td>
<td>2.6±0.1*</td>
<td>3.0±0.2</td>
<td>2.8±0.2</td>
<td>3.6±0.1</td>
<td>3.2±0.3</td>
</tr>
<tr>
<td>HDL cholesterol, mmol/L</td>
<td>1.4±0.1</td>
<td>1.1±0.1</td>
<td>1.3±0.1</td>
<td>1.1±0.1</td>
<td>1.2±0.1</td>
<td>1.1±0.1</td>
</tr>
<tr>
<td>Triglycerides, mmol/L</td>
<td>1.5±0.3</td>
<td>1.2±0.4</td>
<td>1.7±0.2</td>
<td>1.4±0.2</td>
<td>1.3±0.1</td>
<td>1.1±0.1</td>
</tr>
<tr>
<td>Glucose, mmol/L</td>
<td>4.9±0.2</td>
<td>4.9±0.2</td>
<td>5.1±0.1</td>
<td>5.0±0.1</td>
<td>5.2±0.1</td>
<td>5.0±0.1</td>
</tr>
<tr>
<td>Insulin, pmol/L</td>
<td>62.3±5.6</td>
<td>57.2±6.6</td>
<td>61.0±7.7</td>
<td>67.9±10.1</td>
<td>74.6±7.8</td>
<td>69.0±7.4</td>
</tr>
</tbody>
</table>

Values are presented as mean±SEM. BMI indicates body mass index; F, female; HDL, high-density lipoprotein; LDL, low-density lipoprotein; M, male; and VO₂ max, maximal oxygen consumption.

* P<0.05 vs before intervention.
Table 2. Subject Heart Rate and Blood Pressure

<table>
<thead>
<tr>
<th>Variable</th>
<th>Nebulivolol</th>
<th>Metoprolol</th>
<th>Placebo</th>
</tr>
</thead>
<tbody>
<tr>
<td>Heart rate, bpm</td>
<td>Before</td>
<td>After</td>
<td>Before</td>
</tr>
<tr>
<td></td>
<td>66±1</td>
<td>59±2*</td>
<td>70±2</td>
</tr>
<tr>
<td>Systolic BP, mmHg</td>
<td>144±2</td>
<td>126±2*</td>
<td>140±2</td>
</tr>
<tr>
<td>Diastolic BP, mmHg</td>
<td>89±1</td>
<td>77±1*</td>
<td>90±2</td>
</tr>
<tr>
<td>MAP, mmHg</td>
<td>108±1</td>
<td>94±2*</td>
<td>105±2</td>
</tr>
</tbody>
</table>

Values are presented as mean±SEM. BP indicates blood pressure; and MAP, mean arterial pressure.

*P<0.05 vs before intervention.

ingest caffeine-containing beverages before all BP measurements. The recordings were made under quiet, comfortable ambient (≈24°C) laboratory conditions. To avoid the possibility of investigator bias, measurements were made with a semiautomated device (Dinamap, Critikon, FL) that uses an oscillometric technique over the brachial artery. Recordings were made in triplicate in the upright sitting position. All measurements conformed to American Heart Association guidelines as established by the Council for High Blood Pressure Research.

Body Composition
Body mass was measured to the nearest 0.1 kg using a medical beam balance (Detecto, Webb City, MO). Percent body fat was determined by dual energy x-ray absorptiometry (Lunar Radiation Corporation, Madison, WI). Body mass index was calculated as weight (kg) divided by height (m) squared. Minimal waist circumference was measured according to previously published guidelines.

Metabolic Measurements
Fasting plasma lipid, lipoprotein, glucose, and insulin concentrations were determined using standard techniques by the clinical laboratory affiliated with the Clinical Translational Research Center at the University of Colorado at Boulder.

Intra-arterial Infusion Studies
All studies were performed between 7:00 AM and 10:00 AM after a 10-hour overnight fast in a temperature-controlled room. Under strict aseptic conditions a 5-cm, 20-gauge catheter was inserted into the brachial artery of the nondominant arm under local anesthesia (1% lidocaine). Heart rate and arterial BP were continuously measured throughout the infusion protocol. FBF at rest and in response to each pharmacological agent was measured in both the experimental (non-dominant) and the contralateral (dominant) forearm using strain-gauge venous occlusion plethysmography (D.E. Hokanson, Bellevue, WA) as previously described by our laboratory. Baseline FBF was measured for 5 minutes and for 5 minutes before each drug infusion thereafter. After the measurement of resting blood flow, FBF was assessed in response to infusions of acetylcholine (IOLAB Pharmaceuticals, Duluth, GA) at 4.0, 8.0, and 16.0 μg per 100 mL of tissue per minute and sodium nitroprusside (SNP; Nitroprusside, Abbott Laboratories, Abbott Park, IL) at 1.0, 2.0, and 4.0 μg per 100 mL of tissue per minute. Each dose of acetylcholine and SNP was infused for ≈5 minutes, and sufficient time (≈20 minutes) was allowed for FBF to return to resting levels between each vasoactive agent. To avoid an order effect, the sequence of acetylcholine and SNP administration was randomized. After the initial infusion of acetylcholine and SNP and allowing FBF to return to baseline (≈20 minutes), BQ-123 (Clinalfa, AG, Bubendorf, Switzerland), a selective ET<sub>B</sub> receptor antagonist, was infused at a rate of 100 nmol/min for 60 minutes. FBF was measured every 10 minutes throughout the infusion period. The selected dose of BQ-123 has been shown to completely inhibit the vasoconstrictor effect of ET-1 in the human forearm of healthy adults. After 60 minutes of BQ-123 infusion, the FBF response to nonselective ET-1 receptor blockade was assessed by the coadministration of BQ-123 and BQ-788 (Clinalfa, AG) for an additional 60 minutes. BQ-788, a specific antagonist of ET<sub>B</sub> receptors, was infused at a rate of 50 nmol/min, a dose demonstrated to effectively inhibit ET<sub>B</sub> receptors. Thereafter, the infusion of BQ-123 and BQ-788 was continued at the same dose, and FBF was reassessed during coadministration of acetylcholine as performed earlier.

Statistical Analysis
Differences in subject baseline characteristics were determined by between-groups ANOVA. Differences in FBF responses to acetylcholine, SNP, BQ-123, BQ-123+BQ-788, and BQ-123/BQ-788+acetylcholine involving both main effects and interactions (group×intervention) were determined by repeated-measures ANOVA. Post hoc comparisons were performed using the Tukey procedure. There were no significant sex interactions; therefore, the data were pooled and presented together. All data are expressed as mean±SEM. Statistical significance was set a priori at P<0.05.

Results
Selected subject characteristics are presented in Table 1. There were no differences in age and anthropometric, metabolic, or

Figure 1. Forearm blood flow (FBF) responses to BQ-123 before and after 3 months of nebulivolol (A), metoprolol (B), and placebo (C) intervention. Values are presented as mean±SEM. *P<0.05 refers to the difference in the FBF responses to selective endothelin A blockade before vs after intervention.
hemodynamic variables between the groups. Table 2 shows the BP responses among the groups. There were no differences in resting BP between the nebivolol, metoprolol, and placebo groups. Both nebivolol and metoprolol treatment resulted in similar and significant reductions in systolic (≈10%), diastolic (≈15%), and mean arterial (≈15%) BPs. There were no significant changes in BP in the placebo group.

Before randomization to nebivolol/metoprolol/placebo, FBF responses to selective ET\textsubscript{A} receptor blockade with BQ-123 were similarly and significantly elevated (≈30%) from baseline in all 3 groups. Nebivolol, but not metoprolol or placebo, therapy resulted in a marked (≈25%; \(P<0.05\)) reduction in FBF response to BQ-123 (Figure 1). The vasodilator response to BQ-123 was almost identical before and after

Figure 2. Forearm blood flow (FBF) responses to BQ-123 (100 nmol/min) alone and BQ-123 combined with BQ-788 (50 nmol/min) before and after 3 months of nebivolol (A), metoprolol (B), and placebo (C) interventions. Values are presented as means±SEM. *\(P<0.05\) refers to the difference in the FBF responses to selective endothelin A (ET\textsubscript{A}) blockade before vs after intervention. †\(P<0.05\) refers to the difference in the FBF response to nonselective ET\textsubscript{A/B} blockade before vs after intervention.

Figure 3. Forearm blood flow responses to acetylcholine (graphs, top) and sodium nitroprusside (graphs, bottom) before and after 3 months of nebivolol (A), metoprolol (B), and placebo (C) interventions. Values are presented as means±SEM. *\(P<0.05\) vs before intervention.
either metoprolol or placebo treatment. The FBF responses to nonselective ET_A receptor blockade with BQ-123 and BQ-788 were similar among the groups before treatment. There was a significant increase (≈35%) in FBF beyond that of ET_A receptor blockade in each group (Figure 2). However, after 3 months of treatment, only nebivolol therapy significantly reduced (≈40%) the FBF response to nonselective ET_A receptor blockade (Figure 2). Neither metoprolol therapy nor placebo significantly altered the FBF responses to BQ-123/BQ-788 infusion (Figure 2). Across the groups, the FBF response to BQ-123/BQ-788 after each intervention was significantly greater in the nebivolol treatment group.

FBF responses to the endothelium-dependent vasodilator acetylcholine were not significantly different between the 3 groups before intervention (nebivolol: from 5.1±0.3 to 13.3±0.8 mL per 100 mL of tissue per min; metoprolol: from 5.3±0.4 to 13.9±1.1 mL of 100 mL of tissue per min; and placebo: from 4.9±0.2 to 13.0±0.2 mL per 100 mL of tissue per min). Nebivolol treatment resulted in a significant increase (≈20%) in the vasodilator response to acetylcholine (from 4.9±0.4 to 16.4±0.6 mL per 100 mL of tissue per min; Figure 3). In stark contrast, there was no change in the FBF responses to acetylcholine in either the metoprolol (from 5.7±0.5 to 14.1±1.1 mL per 100 mL of tissue per min) or placebo (from 5.1±0.3 to 13.8±0.7 mL per 100 mL of tissue per min) groups after each respective intervention. Across the groups, the FBF response to acetylcholine after each intervention was significantly greater in the nebivolol treatment group. The FBF responses to SNP were not affected by each intervention (Figure 3). The coinfusion of acetylcholine with nonselective ET_A receptor blockade (BQ-123+BQ-788) resulted in significantly greater (≈30%) vasodilator responses in all 3 groups before each intervention (Figure 4). However, after nebivolol therapy (Figure 4), but not metoprolol (Figure 5) or placebo (Figure 6), FBF response to acetylcholine was not significantly increased by the coinfusion of BQ-123+BQ-788 (Figure 4). In both the metoprolol and placebo groups, nonselective ET_A receptor blockade augmented acetylcholine-mediated vasodilation to similar extent when compared with before each intervention.

Figure 4. Forearm blood flow (FBF) responses (top) and total FBF (bottom) to acetycholine in the absence or presence of nonselective endothelin A/B blockade (BQ-123+BQ-788) before and after nebivolol intervention. Values are presented as mean±SEM. *P<0.05 vs saline.
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Discussion

The BP-lowering effects of nebivolol are well established.\textsuperscript{13,27,28} The seminal and novel finding of the present study, however, is that in addition to, and independent of, lowering BP, nebivolol markedly and favorably affects ET-1 system activity. Indeed, the results reported herein demonstrate that (1) chronic nebivolol, but not metoprolol, therapy reduces ET-1–mediated vasoconstrictor tone in adults with elevated BP and (2) reductions in ET-1 vasoconstriction underlie nebivolol-induced improvements in endothelium-dependent vasodilation. Diminished ET-1–mediated vasoconstrictor tone may represent an important endovascular pleiotropic effect of nebivolol, contributing to its favorable effect on overall cardiovascular risk.\textsuperscript{29}

In this study, there was a similar and significant (≈30%) increase in FBF responses to selective ETA receptor blockade in all 3 groups before treatment. In addition, nonselective ETA/B receptor blockade resulted in a further increase (≈35%) in FBF in all the groups. These findings are fully consistent with previous studies establishing enhanced ET-1–mediated vasoconstrictor tone in adults with BP in both the hypertensive\textsuperscript{12} and the prehypertensive\textsuperscript{11} range. For example, Cardillo et al\textsuperscript{1,12} demonstrated almost identical increases (35%–55%) in FBF to selective and nonselective ET receptor blockade, to that reported herein, in adults with essential hypertension compared with marginal, nonsignificant changes in FBF in normotensive adults. Thus, we are confident that ET-1–mediated vasoconstrictor tone was abnormally high in our subjects with elevated BP without a direct comparison with a normotensive control group.

In vitro, nebivolol has been shown to blunt endothelial production, and in turn release, of ET-1.\textsuperscript{30} The results of this study complement and significantly extend these findings by demonstrating that nebivolol reduces ET-1–mediated vasoconstrictor tone in vivo in adults with elevated BP. After 3 months of nebivolol therapy, there was a marked reduction (≈25%) in the vasodilator response to both selective ET\textsubscript{A} and nonselective ET\textsubscript{A/B} receptor blockade. Of note, the nebivolol-induced reduction in ET-1 vasoconstrictor tone was independent of concomitant reductions in BP. Indeed, BP was equally and significantly reduced in adults randomized to either the nebivolol or metoprolol treatment groups. However, metoprolol therapy had no effect on the vascular responses to

Figure 5. Forearm blood flow (FBF) responses (top) and total FBF (bottom) to acetycholine in the absence or presence of nonselective endothelin A/B blockade (BQ-123+BQ-788) before and after metoprolol intervention. Values are presented as mean±SEM. *P<0.05 vs saline.
either selective or nonselective ET-1 receptor blockade despite its BP-lowering effect. Moreover, there were no significant changes in body composition or cardiometabolic risk factors in response to nebivolol (or metoprolol) treatment. Collectively, this provides further evidence for a direct clinical effect of nebivolol on the ET-1 system. Several mechanisms may underlie this unique feature of nebivolol. Most notably, nebivolol has been shown to ameliorate prepro–ET-1 mRNA production in human coronary endothelial cells. Prepro–ET-1 is the peptide transcribed from prepro–ET-1 mRNA that is post-translationally modified to ET-1. Reduction in prepro–ET-1 would ultimately lead to less ET-1 formation. Other contributing factors may include greater nitric oxide bioavailability and reduced oxidative mediators. Nebivolol increases nitric oxide bioavailability by enhancing endothelial nitric oxide synthase activity through calcium- and noncalcium-dependent pathways. Nitric oxide, in turn, has a potent inhibitory influence on ET-1 at the level of transcription and endothelin-converting enzyme activity. About oxidative stress, nebivolol has been shown to block nicotinamide adenine dinucleotide phosphate oxidase, a known activator of the ET-1 system.

Concurrent with the nebivolol-induced reduction in ET-1–mediated vasoconstrictor tone, we also demonstrate that the nebivolol-induced improvement in endothelium-dependent vasodilation is due, at least in part, to the reduction in ET-1 vasoconstriction. It is important to note that before intervention, the FBF responses to acetylcholine in all 3 groups were similar to that previously reported in prehypertensive and hypertensive adults, supporting diminished endothelium-dependent vasodilation in our study population. Consistent with previous studies, we demonstrate that chronic nebivolol therapy significantly improves acetylcholine-mediated endothelium-dependent vasodilation in adults with elevated BP. In stark contrast, there was no effect of metoprolol therapy (or placebo) on endothelial vasodilator function. It has been suggested that the nebivolol-induced improvement in endothelium-dependent vasodilation is largely because of an increase in nitric oxide bioavailability. A seminal finding of this study is that reduced ET-1 vasoconstrictor tone seems to be a primary contributor to improved endothelial vasodilator function. Indeed, before intervention, the coinfusion of nonselective ET\textsubscript{A} receptor blockade resulted in a significant increase (≈35%) in

![Figure 6. Forearm blood flow (FBF) responses (top) and total FBF (bottom) to acetycholine in the absence or presence of nonselective endothelin A/B blockade (BQ-123+BQ-788) before and after placebo intervention. Values are presented as mean±SEM. *P<0.05 vs saline.](http://hyper.ahajournals.org/doi/abs/10.1161/HYPERTENSIONAHA.116.11202)
acetylcholine-stimulated endothelial vasodilation in all 3 treatment groups. After the 3-month intervention period, this effect was unchanged in the metoprolol and placebo groups; however, in the nebivolol group, nonselective ET<sub>AB</sub> receptor antagonism no longer enhanced the FBF responses to acetylcholine. Although we did not assess whether the nebivolol-induced improvement in acetylcholine-mediated vasodilation was nitric oxide dependent, it is plausible that the previously reported increase in nitric oxide bioavailability with nebivolol is due, in part, to an uncoupling of ET-1–mediated nitric oxide inhibition. Moreover, relieving ET-1–mediated nitric oxide inhibition would allow nitric oxide, and other endothelium-derived relaxing factors, to act without opposition and dilate the vessel appropriately in response to stimulation. Thus, the unique vasomotor properties of nebivolol seem to involve both vasodilator and vasoconstrictor factors. To the best of our knowledge, this is the first study to assess the involvement of the ET-1 system in nebivolol-induced improvements in endothelial vasodilator function. There are a few experimental considerations about the present study that deserve to be mentioned. First, given the extended half-life of ET receptor antagonists, our study design did not involve the singular administration of the selective ET<sub>B</sub> receptor antagonist BQ-788, and therefore, we cannot comment on the effects of nebivolol (or metoprolol) on the independent vascular actions of the ET<sub>B</sub> receptor. Second, we did not measure circulating plasma levels of ET-1 in the present study. ET-1 produced by the endothelium is predominantly (>80%) released abuminally toward the vascular smooth muscle<sup>42</sup>; thus, the pathophysiologic significance of circulating ET-1 levels can be variable.<sup>43</sup> Circulating plasma concentrations of the peptide may not necessarily reflect local vascular production but rather spillover into, and clearance from, the bloodstream.<sup>42</sup> However, elevations in plasma ET-1 concentrations have been linked with ET receptor activity.<sup>44</sup> Third, consistent with previous studies, we infused BQ-123 for 60 minutes before the coinfusion with BQ-788<sup>12</sup>; the time course for the slow-onset vasodilation with BQ-123 has been shown to maximize by 60 minutes in some studies<sup>45</sup>–<sup>47</sup> and 90 minutes in another study.<sup>48</sup> As a result, we cannot rule out the possibility that further increase in FBF noted in the groups in response to the addition of BQ-788 to BQ-123 may involve some residual effects of BQ-123.

**Perspectives**

In conclusion, the results of this study indicate that nebivolol, but not metoprolol, treatment reduces ET-1–mediated vasoconstrictor tone in adult humans with elevated BP. Moreover, nebivolol-induced reduction in ET-1–mediated vasoconstrictor tone seems to be an important factor underlying the favorable effects of this β-blocker on endothelial vasodilation. Importantly, the direct effect of nebivolol on ET-1 system activity is in addition to, and independent of, its established BP-lowering effects and may be a key factor contributing to the improvement in endothelial health and reduction in cardiovascular morbidity and mortality<sup>49</sup> associated with chronic nebivolol treatment.

**Acknowledgments**

We thank all the subjects who participated in the study, as well as the clinical staff at the Clinical Translational Research Center, University of Colorado-Boulder for their assistance.

**Sources of Funding**

This study was supported by funding from Forest Research Institute, Inc (BYS-MD-57) and National Institute of Health award UL1TR001082.

**Disclosures**

None.

**References**


Chronic Nebivolol Treatment Suppresses Endothelin-1–Mediated Vasoconstrictor Tone in Adults With Elevated Blood Pressure
Kyle J. Diehl, Brian L. Stauffer, Caitlin A. Dow, Tyler D. Bammert, Danielle L. Brunjes, Jared J. Greiner and Christopher A. DeSouza

Hypertension. 2016;67:1196-1204; originally published online April 25, 2016; doi: 10.1161/HYPERTENSIONAHA.115.06979

Hypertension is published by the American Heart Association, 7272 Greenville Avenue, Dallas, TX 75231
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Print ISSN: 0194-911X. Online ISSN: 1524-4563

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