Research Article

Acupuncture Attenuates Renal Sympathetic Activity and Blood Pressure via Beta-Adrenergic Receptors in Spontaneously Hypertensive Rats

Jing-Wen Yang,1 Yang Ye,2 Xue-Rui Wang,1 Fang Li,1 Ling-Yong Xiao,2 Guang-Xia Shi,1 and Cun-Zhi Liu1

1Acupuncture and Moxibustion Department, Beijing Hospital of Traditional Chinese Medicine Affiliated to Capital Medical University, 23 Meishuguanhou Street, Dongcheng District, Beijing 100010, China
2Beijing University of Chinese Medicine, 11 Beisanhuan East Road, Chaoyang District, Beijing 100029, China

Correspondence should be addressed to Cun-Zhi Liu; lcz623780@126.com

Received 8 August 2016; Accepted 11 January 2017; Published 8 February 2017

Academic Editor: Preston E. Garraghty

Copyright © 2017 Jing-Wen Yang et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

The sympathetic nervous system, via epinephrine and norepinephrine, regulates β-adrenergic receptor (β-AR) expression, and renal sympathetic activation causes sustained increases in blood pressure by enhanced renin release. In this study, we aim to investigate the effect and underlying mechanism of acupuncture at Taichong (LR3) on renal sympathetic activity in spontaneously hypertensive rats. Unanesthetized rats were subjected to daily acupuncture for 2 weeks. Mean blood pressure (MBP) and heart rate variability (HRV) were monitored at days 0, 7, and 14 by radiotelemetry. After euthanasia on the 14th day, blood and the kidneys were collected and subject to the following analyses. Epinephrine and norepinephrine were detected by ELISA. The expression of β-ARs was studied by western blotting and PCR. The renin content was analyzed by radioimmunoassay. 14-day acupuncture significantly attenuates the increase of MBP. The HRV indices, the standard deviation of all normal NN intervals (SDNN), and the ratio of the low-frequency component to the high-frequency component (LF/HF) were improved following acupuncture. Renal sympathetic activation induced upregulation of epinephrine, norepinephrine, and renin content were attenuated by acupuncture. In addition, acupuncture decreased β1-AR expression and improved β2-AR expression. These results indicated that acupuncture relieves the increased MBP via the regulation of renal sympathetic activity and β-ARs.

1. Introduction

Enhanced sympathetic nervous system activity is linked to the development and maintenance of hypertension [1–3]. The extensive sympathetic innervation of the kidney occupies a special place in the development of hypertension. Calaresu and Ciriello [4, 5] showed that renal afferent nerves project directly into a number of areas in the central nervous system, such as the lateral tegmental fields, the paramedial reticular nucleus, the dorsal vagal complex of the medulla, and the lateral hypothalamic area, contributing to arterial blood pressure regulation. Renal efferent sympathetic activity participates in renin release, sodium retention, and reduced renal blood flow, which contribute to the development of hypertension [6–8]. In animal models, Kumagai and his colleagues [9] found that stimulation of afferent sympathetic nerve increased systemic sympathetic nerves activity and caused vasoconstriction. Moreover, the attenuation of either the efferent or afferent renal nerves theoretically contributes to lowering blood pressure (BP) [8]. The renal sympathetic nervous system regulates BP mainly through catecholamines (epinephrine and norepinephrine) binding to β-adrenergic receptors (β-ARs). Moreover, β1-ARs in the kidney may activate renin release [10] and, consequently, result in the renin-angiotensin-aldosterone system (RAAS) activation and angiotensin II formation followed by vasoconstriction [11]. Therefore, the renal sympathetic nervous system and β-ARs are critical for regulating peripheral
resistance and play important roles in the development of hypertension.

Acupuncture has been recommended as a complementary therapy for hypertension. In 1996, 64 acupuncture indications were declared by World Health Organization in Milan conference including hypertension [12]. Numerous animal and clinical studies have reported the efficacy of acupuncture in treating hypertension [13–15]. Although there is some disagreement among the reports, the majority of them indicate that acupuncture causes a significant decrease in BP [13]. However, the underlying mechanism through which acupuncture lowers BP remains to be elucidated. Acupuncture could affect the sympathetic nervous system. Knardahl et al. found that acupuncture produces moderate hypalgesia in humans paralleled by a regulation of muscle sympathetic nerve activity [16]. Song et al. suggested that acupuncture pretreatment improved the survival rate in rats with lethal endotoxemia, which involves the activation of the autonomic nervous system [17]. Research has shown that electroacupuncture modulates the renal sympathetic nerve activity in chronic kidney disease rats [18]. Although electroacupuncture modulates the renal sympathetic nerve activity in chronic kidney disease rats [18]. Although electroacupuncture modulates the renal sympathetic nerve activity in chronic kidney disease rats [18]. Although electroacupuncture modulates the renal sympathetic nerve activity in chronic kidney disease rats [18]. Although electroacupuncture modulates the renal sympathetic nerve activity in chronic kidney disease rats [18]. Although electroacupuncture modulates the renal sympathetic nerve activity in chronic kidney disease rats [18]. Although electroacupuncture modulates the renal sympathetic nerve activity in chronic kidney disease rats [18]. Although electroacupuncture modulates the renal sympathetic nerve activity in chronic kidney disease rats [18]. Although electroacupuncture modulates the renal sympathetic nerve activity in chronic kidney disease rats [18]. Although electroacupuncture modulates the renal sympathetic nerve activity in chronic kidney disease rats [18].

The present study was conducted to examine the hypothesis that the activation of the renal sympathetic nervous system in SHRs is mediated by acupuncture. Furthermore, we assessed if this renal nerve-mediated effect contributes to the downregulation of β1-AR and upregulation of β2-AR, leading to the decreased BP in SHRs.

2. Materials and Methods

2.1. Experimental Animal. Male SHR and WKY rats (12 weeks), weighing 260–300 g, were purchased from Vital River Laboratory Animal Technology Co. Ltd (Beijing China). The animals were housed in cages at 22 ± 2 °C and humidity of 40 ± 5% under a 12-hour light/dark cycle and received standard diet and water ad libitum. The experimental procedures were in accordance with the Guidelines for the Institutional Animal Care and Use Committee of China Academy of Chinese Medical Sciences (Beijing, China).

2.2. Animal Grouping and Acupuncture Stimulation. The rats were randomly divided into 4 groups, the control group (WKY), the model group (SHR), the acupuncture stimulation group at LR3 (located between the first and second metatarsal bones on the dorsum of the foot) (Acu), and nonacupoint acupuncture stimulation (Non-Acu), with 10 rats in each group. Sterilized disposable stainless steel needles (0.3 mm × 40 mm, Hwato, China) were inserted 5 mm deep at LR3. For the Non-Acu group, the rats received similar treatment as Acu group but the acupuncture site was at a sham acupoint (on the bilateral hypochondrium, 10 mm above iliac crest) to replace LR3. The rats in the WKY and SHR groups were given the same time and same level catching-grasping stimulus without acupuncture stimulation. Acupuncture proceeded for 30 seconds each time, once daily for a period of 2 weeks (1 day rest after 6 days of treatment).

2.3. Radiotelemetry. We measured BP and HRV by radiotelemetry combined with fast Fourier transform analysis of BP and HRV. DSCF-FS01 HRV devices or DSCF-FS02 BP devices (Softron, USA) were implanted in rats to follow BP and HRV changes continuously over time. The telemetric techniques and the techniques employed to analyze the autonomic nervous system are described in detail elsewhere [19]. Briefly, we anesthetized the rats with intraperitoneal injection of pentobarbital sodium (Sigma, USA). Then, The telemeter was placed in the abdominal cavity, the blood pressure catheter tip was inserted into the descending abdominal aorta, and the electrode leads approximated the Lead II orientation as previously described [20]. The rats recovered for 7 days before baseline BP and HRV values were recorded. By this time, the rats had regained their circadian BP and HRV rhythm; and surgery and anesthesia-induced changes had abated.

Data were sampled every 5 min for 10 sec continuously day and night and stored on a hard disk. BP and HRV were recorded using the SP 2006 Lan software. Continuous beat-by-beat values of BP and HRV were recorded during morning hours. Measurements to analyze the autonomic control of BP and HR and HRV were performed between 8:00 and 10:00 AM.

2.4. ELISA. NE and E contents in the plasma and kidney were measured by enzyme-linked immunosorbent assay (ELISA) kit (R&D Systems, USA) according to the manufacturer’s instruction. Absorbance in each well was measured using microplate reader (Thermo Fisher, USA) at 450 nm. Concentrations of NE and E in the plasma and renal were determined by interpolation from the standard curve.

2.5. Real-Time Quantitation PCR. Total RNA was extracted from the kidney using Trizol Reagent (Invitrogen) according to the manufacturer’s instructions. Quantification of the relative amounts of mRNA was performed using the Rotor-Gene6000 1.7 version software (Corbett research). In brief, 2 μL of the cDNA was mixed with primers, 100 nM of probe, and 1x TaqMan Universal Master Mix in each reaction. Predeveloped assay reagents were used and were purchased from Applied Biosystems. Sequence for the primers and TaqMan probes for the rat β1-AR, β2-AR, and renin are shown in Table 1. Samples were tested in triplicate, and differences of threshold cycles between target genes and house-keeping gene (18S rRNA) were calculated. The relative mRNA abundance in the treatment groups was calculated using \( 2^{-\Delta\Delta CT} \) method using the control group as the calibrator according to the manufacturer’s user manual. The value of relative mRNA quantity for control group is 1 with arbitrary units.

2.6. Western Blot Analysis. Total protein in each sample was measured by the BCA assay (Bio-Rad, USA) using bovine serum albumin as the standard protein. A fixed amount of protein (20 μg) from each sample was fractionated by 10% SDS-polyacrylamide gel electrophoresis (SDS-PAGE) and transferred to a polyvinylidene difluoride (PVDF) membrane.
(Pall, USA). Membranes were incubated in 5% milk in TBS for 1 h in room temperature. For identification of β-ARs, membranes were exposed to primary antibodies (anti-β₁-AR, 1: 250 dilution, or anti-β₂-AR, 1: 500 dilution, abcam, USA). After washing, the membranes were incubated with secondary antibodies (HRP-conjugated anti-rabbit antibody, 1: 5000 dilution, KBL, USA). The bound antibodies were detected using the enhanced chemiluminescent reagent (GE Health Care, USA). Data are presented as the β-ARs to actin ratio and then expressed as fold-change compared to WKY group.

2.7. Radioimmunoassay. For determination of PRC, 50 µL of blood was collected into EDTA-containing 75-µL microhematocrit tubes from conscious rats by tail vein puncture. For determination of renal renin content, samples of kidney cortex were dissected under the microscopy, frozen in liquid nitrogen, and stored at −80°C until assay. For renin analysis, tissue was weighed, homogenized with two 30-second pulses in a 100-fold excess of homogenization buffer (5% [vol/vol] glycerol, 0.1 mmol/L of PMSF, 10 mmol/L of EDTA, and 0.1 mmol/L of 4-[2-aminomethyl]benzenesulfonyl fluoride) using a Polytron homogenizer (Kinematica), and centrifuged at 4°C at 140,000g for 5 minutes. Supernatants were incubated with saturating concentrations of rat renin substrate, and angiotensin I generation was assayed by radioimmunoassay (DiaSorin).

2.8. Statistical Analysis. Data analysis was performed with SPSS software version 16.0. All values were expressed as the mean ± standard error (SEM). Comparison between the treatment group and the control group was performed by one-way ANOVA test followed by a post hoc least significant difference multiple comparison test. A P value < 0.05 was considered statistically significant.

3. Results

3.1. Effect of Acupuncture on BP in SHRs. To confirm the efficacy of acupuncture, BP levels were measured in all groups by the telemetry method after two weeks of acupuncture treatment (Figure 1(a)). Before acupuncture, the mean blood pressure (MBP) of SHR, Acu, and Non-Acu group were insignificantly different (P > 0.05), but significantly higher than in the WKY rats. After 7 days of acupuncture, the MBP was significantly decreased compared with SHR group and Non-Acu group (P < 0.01), and this decrease was sustained throughout the treatment period (days 7-14). Although the MBP of Acu group was significantly decreased, it was still higher than the WKY group (P < 0.01) over the experimental period. These results suggested that acupuncture could lower BP but is unable to reduce it to normal levels.

3.2. Effect of Acupuncture on the Balance of Autonomic Nervous in SHRs. To investigate whether the attenuation of BP in SHRs by the treatment with acupuncture is associated with beneficial effects of the automatic nervous system, we used radiotelemetry method to examine the heart rate variability (HRV) in all groups at days 0, 7, and 14 (Figures 1(b)–1(d)). The values of standard deviation of all normal NN intervals (SDNN) and root mean square of differences between adjacent NN intervals (rMSSD) show acupuncture effects on HRV by mean variation of the short-term period. Before acupuncture, the SDNN of the WKY group was higher than the other groups, and SDNN was slightly improved in the Acu group after 7 days of treatment without significance (P > 0.05). In the 14 days of acupuncture, the SDNN was significantly increased compared with the SHR group and Non-Acu group (P < 0.01). However, there was no significant change in rMSSD among the four groups (P > 0.05).

The ratio of the low-frequency component to the high-frequency component (LF/HF) shows acupuncture effects on autonomic regulation. LF/HF of the SHR group, Acu group, and Non-Acu group were significantly higher than that of the WKY group at days 0 and 7 (P > 0.05). On the 14th day, acupuncture significantly reduced the ratio of LF/HF compared with the SHR group and Non-Acu group (P < 0.01).

3.3. Effect of Acupuncture on the Contents of Catecholamine in SHRs. The catecholamines, including norepinephrine (NE) and epinephrine (E), acting as neurotransmitters, play important roles in the sympathetic control of arterial BP. Therefore, we tested the contents of NE and E in plasma and renal tissues by ELISA (Figure 2). At the end of the study, NE and E levels induced by sympathetic nervous system activation were significantly higher in plasma and renal tissues from SHR than in those from WKY rats. Acupuncture at LR3 but not the nonacupoint significantly decreased NE and E concentrations in SHR group, and there was no significant difference between WKY group and Acu group.

3.4. Effect of Acupuncture on β-Adrenergic Receptor in SHRs. Both NE and E act at G protein-coupled receptors of the adrenergic receptor family to mediate sympathetic effects. Within this main classification, there are several subtypes α₁A, α₁B, α₁D, α₂A, α₂B, α₂C, β₁, β₂, and β₃. In this study we focused on two of these receptors, β₁ and β₂. Both of these receptor subtypes have been strongly implicated in cardiovascular control. To determine whether the antihypertensive
3.5. Effect of Acupuncture on Plasma Renin Concentration, Renal Renin Content, and Renin mRNA Expression in SHRs. The juxtaglomerular (JG) cells in the media of renal afferent arterioles are the major sites of synthesis of the aspartic protease renin, the rate-limiting enzyme in the formation of angiotensin II. The JG cells are in contact with sympathetic nerve varicosities and express postjunctional β-ARs. Activation of β-ARs directly increases renin secretion. To address the role of acupuncture in the regulation of renin, we used radioimmunoassay to detect the concentration of renin. As shown in Figure 4, plasma renin concentration (PRC) increased significantly in SHR group and Non-Acu group and decreased significantly after acupuncture at LR3. Renal renin content was significantly higher in SHR rats compared with WKY rats. There was a significant reduction in renin content in the Acu group, whereas the decrease in renin content in Non-Acu group did not reach significance. Renin mRNA was markedly higher in SHR rats compared with WKY rats. Renin mRNA was significantly altered by the effect of acupuncture is associated with beneficial outcome in β-ARs, we detected the mRNA and protein levels of β-ARs after acupuncture treatment by PCR and Western blot, respectively (Figure 3). Compared to WKY rats, SHR rats had significantly increased β1-AR protein levels and decreased β2-AR protein levels. After acupuncture, β1-AR protein levels were significantly decreased compared to SHR group. The signal ratio of β1-AR protein to actin was reduced by about 16.5% (1.01 versus 1.21). In contrast, β1-AR protein levels were similar between SHR group and Non-Acu group. The trends of the β-ARs mRNA levels were consistent with its protein levels.
acupuncture at LR3, whereas acupuncture at nonacupoint caused nonsignificant reductions of renin mRNA.

4. Discussion

In the present study, we show that acupuncture at LR3 reduced MBP and regulated HRV in SHR. And the therapeutic effects were associated with renal sympathetic nervous system, which can be manifested by the significant reductions in the levels of E and NE (an indirect marker of sympathetic activity) and the content of renin (indicative of increased renal sympathetic activity) in the plasma and renal by acupuncture. In addition, acupuncture decreased $\beta_1$-AR expression and increased $\beta_2$-AR expression, which means acupuncture has bidirectional regulation effects on $\beta$-ARs.

Autonomic imbalance with increased sympathetic activity has been strongly implicated in the pathophysiology of hypertension. Among the different available noninvasive techniques for assessing the autonomic status, HRV has emerged as a simple, noninvasive method to evaluate the sympathovagal balance. Our findings of depressed SDNN in SHR rats suggested a homeostatic disruption in sympathovagal balance. However, because rMSSD is generally associated with vagal pathways, our results with unchanged rMSSD suggest an increase in sympathetic tone in SHR rats. This result is consistent with Schroeder's finding that individuals with low SDNN at baseline were at an increased risk of developing hypertension over 9 years of follow-up [21]. An analysis of LF/HF ratios rather than single components is considered by many investigators to better reflect the activity of the sympathovagal balance. Neto et al. showed a significant correlation between reduction BP levels and lower LF/HF ratio [22]. In our study, the higher LF/HF ratio demonstrated that SHRs have excessive sympathetic activation and autonomic imbalance. After acupuncture treatment, SDNN increased and LF/HF ratio decreased, but the rMSSD did not change. These results indicated that acupuncture could regulate the balance of the autonomic nervous system mainly through reducing the sympathetic activity.

Evidence collected throughout the years has shown that alterations in sympathetic cardiovascular control participate in the development, maintenance, and progression of hypertension. Renal sympathetic nerves induce high BP via E and NE regulates $\beta$-ARs expression. Of the two main $\beta$-AR subtypes ($\beta_1$ and $\beta_2$), $\beta_1$-AR signaling has been linked to cardiotoxicity. In contrast, $\beta_2$-AR activates signaling pathways.
involved in cardioprotection. Gui et al. found that \( \beta_2 \)-AR has a vasodilatation effect, and Raf kinase inhibitory protein could correct impaired \( \beta_2 \)-AR to treat hypertension [23]. Thus, some have proposed that the \( \beta_1 \)-AR is the “cardiotoxic subtype” whereas the \( \beta_2 \)-AR is the “cardioprotective subtype.” The two receptors regulate the peripheral resistance which reflects the balance between vasoconstrictor and vasodilator mechanisms. Based on this, we examined the effect of acupuncture on E, NE, and \( \beta \)-ARs. Here we observed that the contents of E and NE of SHR rats were significantly higher compared with the WKY rats. This gives direct evidence that the sympathetic nervous system is hyperreactive in the SHR and is in general agreement with previous studies [24, 25]. Then acupuncture at LR3, not nonacupoint, could decrease the contents of E and NE. In addition, acupuncture significantly decreased the expression of \( \beta_1 \)-AR and increased \( \beta_2 \)-AR expression in SHR. Therefore, the antihypertensive effect of acupuncture in SHR might be associated with the balance of \( \beta_1/\beta_2 \)-AR. \( \beta \)-blockers therapy plays a major role in the treatment of cardiovascular diseases. To date, 14 \( \beta \)-blockers have received Federal Drug Administration approval for oral use in patients having systemic hypertension [26]. By receptor selective classification, \( \beta \)-blockers can be divided into three categories, namely, nonselective b-blocker, selective b1-receptor blocker, and b1-receptor blocker with additional a1-receptor blocking activity. Among them, the nonselective b-blocker is used most widely. In general, \( \beta \)-blockers are well tolerated, but serious side effects may occur, which are usually associated with the \( \beta_2 \)-AR antagonistic activity (e.g., increase in peripheral vascular resistance, worsening of asthma symptoms) [27]. Wong et al. suggested that \( \beta_1 \) selective blockers lowered BP by a greater magnitude compared to dual receptor beta-blockers [28]. Acupuncture is reported to have potential for treating hypertension with
Figure 4: Effects of acupuncture on renin content (a and b) and renin mRNA expression (c). Data are presented as mean ± SEM (n = 10 rats). ## indicate P < 0.01, compared with the WKY group; ** indicate P < 0.01, compared with SHR group; & indicate P < 0.05, compared with Non-Acu group.

fewer side effects [29]. Moreover, Zhang et al. reported that acupuncture may serve as an alternative for hypertensive patients, especially for those who cannot tolerate the side effects of antihypertensive drugs [30]. Combined with our experimental results, where acupuncture could decrease β1-AR expression and increase β2-AR expression, we consider that the low side effects of acupuncture may be related to the increased expression of β2 receptors, which needs to be further investigated.

The RAAS is an important mechanism in regulating BP. For that reason, an alteration in any molecule that composes the RAAS contributes to developing hypertension [31]. Classically, the sympathetic nervous system rapidly activates the RAAS through β1-AR mediated release of renin [32]. Therefore, inhibition of renin may exert complete inhibition of the RAAS, leading to a decrease of BP. Hisa et al. found that a β-AR blocker could reduce renal nerve stimulation-induced renin release. In our experiment, we found a significant reduction of renin content after acupuncture. Consistent with our results, Lohmeier et al. demonstrated that renal denervation decreased plasma renin content and abolished the hypertension [33]. However, the effect of acupuncture on RAAS in SHRs needs further study.

The renal sympathetic nervous system plays an active role in the modulation of BP, and its overactivation can lead to hypertension [34]. The Symplicity HTN-1 and HTN-2 studies proposed renal denervation as an effective and safe approach to treat patients with hypertension and were followed by substantial enthusiasm [35, 36]. In contrast with Symplicity HTN-1 and HTN-2, the announcement that Symplicity HTN-3 failed to meet its primary efficacy endpoint and put an abrupt stop to these overoptimistic expectations shows that renal denervation is not ready for clinical dissemination [37]. However, some researchers considered that the execution of the study was hampered by operational weaknesses and drug adherence [38, 39]. Despite the disappointing outcome of Symplicity HTN-3, the renal sympathetic system is still a potential target for treating hypertension, and our results indicate that acupuncture could reduce the renal sympathetic nervous activity and lower BP without related risk.

In conclusion, our results provide evidence that acupuncture at LR3 can significantly relieve the increased MBP through reducing renal sympathetic nervous activity. The effect of acupuncture on renal sympathetic nervous activity is evidenced by the improved HRV and the reduction of NE and E contents. The decreased β1-AR and increased β2-AR
expression we observed may also be involved in the beneficial effect of acupuncture on hypertension.

**Competing Interests**

The authors declare that they have no conflict of interests with the contents of this article.

**Authors’ Contributions**

Jing-Wen Yang conducted experiments on the role of renin, analyzed the results, and wrote most of the paper. Xue-Rui Wang and Fang Li conducted the radiotelemetry. Yang Ye and Ling-Yong Xiao conducted experiments searching for β-ARs function. Guang-Xia Shi conducted the ELISA of epinephrine and norepinephrine content. Cun-Zhi Liu conceived the idea for the project and wrote the paper with Jing-Wen Yang.

**Acknowledgments**

This work was supported in whole or partly by National Natural Science Foundation of China (Grant no. 81473501), Beijing Municipal Administration of Hospitals Clinical Medicine Development of Special Funding Support (code: ZYLYX201412), and National Basic Research Program of China (Grant no. 2014CB543203).

**References**

[1] P. A. Smith, L. N. Graham, A. F. Mackintosh, J. B. Stoker, and D. A. S. G. Mary, “Relationship between central sympathetic activity and stages of high hypertension,” *American Journal of Hypertension*, vol. 17, no. 3, pp. 217–222, 2004.

[2] S. C. Malpas, “Sympathetic nervous system overactivity and its role in the development of cardiovascular disease,” *Physiological Reviews*, vol. 90, no. 2, pp. 513–557, 2010.

[3] A. E. Simms, J. F. R. Paton, A. E. Pickering, and A. M. Allen, “Amplified respiratory-sympathetic coupling in the spontaneously hypertensive rat: does it contribute to hypertension?” *The Journal of Physiology*, vol. 587, no. 3, pp. 597–610, 2009.

[4] F. R. Calaresu and J. Ciriello, “Renal afferent nerves affect discharge rate of medullary and hypothalamic single units in the cat,” *Journal of the Autonomic Nervous System*, vol. 3, no. 2–4, pp. 311–320, 1981.

[5] J. Ciriello and F. R. Calaresu, “Central projections of afferent renal fibers in the rat: an anterograde transport study of horseradish peroxidase,” *Journal of the Autonomic Nervous System*, vol. 8, no. 3, pp. 273–285, 1983.

[6] H. Krum, P. Sobotka, F. Mahfoud, M. Böhm, M. Esler, and M. Schlachter, “Device-based antihypertensive therapy: therapeutic modulation of the autonomic nervous system,” *Circulation*, vol. 123, no. 2, pp. 209–215, 2011.

[7] G. F. DiBona, “Neural control of the kidney: past, present, and future,” *Hypertension*, vol. 41, no. 3, pp. 621–624, 2003.

[8] G. F. DiBona, “Physiology in perspective: The Wisdom of the Body. Neural control of the kidney,” *American Journal of Physiology—Regulatory Integrative and Comparative Physiology*, vol. 289, no. 3, pp. R633–R641, 2005.

[9] H. Kumagai, N. Oshima, T. Matsuura et al., “Importance of rostral ventrolateral medulla neurons in determining efferent sympathetic nerve activity and blood pressure,” *Hypertension Research*, vol. 35, no. 2, pp. 132–141, 2012.

[10] T. K. Keeton and W. B. Campbell, “The pharmacologic alteration of renin release,” *Pharmacological Reviews*, vol. 32, pp. 81–227, 1980.

[11] H. Kubista and S. Boehm, “Molecular mechanisms underlying the modulation of exocytotic noradrenaline release via presynaptic receptors,” *Pharmacology and Therapeutics*, vol. 112, no. 1, pp. 213–242, 2006.

[12] J.-Y. Wang, H. Li, C.-M. Ma, J.-L. Wang, X.-S. Lai, and S.-F. Zhou, “MicroRNA profiling response to acupuncture therapy in spontaneously hypertensive rats,” *Evidence-Based Complementary and Alternative Medicine*, vol. 2015, Article ID 204367, 9 pages, 2015.

[13] C. Yin, B. Seo, H.-J. Park et al., “Acupuncture, a promising adjunctive therapy for essential hypertension: a double-blind, randomized, controlled trial,” *Neurological Research*, vol. 29, no. 1, pp. S98–S103, 2007.

[14] J.-M. Park, A.-S. Shin, S.-U. Park, I.-S. Sohn, W.-S. Jung, and S.-K. Moon, “The acute effect of acupuncture on endothelial dysfunction in patients with hypertension: a pilot, randomized, double-blind, placebo-controlled crossover trial,” *Journal of Alternative and Complementary Medicine*, vol. 16, no. 8, pp. 883–888, 2010.

[15] L. A. Kalish, B. Buczynski, P. Connell et al., “Stop Hypertension with the Acupuncture Research Program (SHARP): clinical trial design and screening results,” *Controlled Clinical Trials*, vol. 25, no. 1, pp. 76–103, 2004.

[16] S. Knardahl, M. Elam, B. Olausson, and B. G. Wallin, “Sympathetic nerve activity after acupuncture in humans,” *Pain*, vol. 75, no. 1, pp. 19–25, 1998.

[17] J.-G. Song, H.-H. Li, Y.-F. Cao et al., “Electroacupuncture improves survival in rats with lethal endotoxemia via the autonomic nervous system,” *Anesthesiology*, vol. 116, no. 2, pp. 406–414, 2012.

[18] J. C. Paterno, C. T. Bergamaschi, R. R. Campos et al., “Electroacupuncture and moxibustion decrease renal sympathetic nerve activity and retard progression of renal disease in rats,” *Kidney and Blood Pressure Research*, vol. 35, no. 5, pp. 355–364, 2012.

[19] V. Gross, J. Tank, M. Obst et al., “Autonomic nervous system and blood pressure regulation in RGS2-deficient mice,” *American Journal of Physiology—Regulatory Integrative and Comparative Physiology*, vol. 288, no. 5, pp. R134–R142, 2005.

[20] L. B. Wichers, J. P. Nolan, D. W. Winsett et al., “Effects of instilled combustion-derived particles in spontaneously hypertensive rats. Part I: cardiovascular responses,” *Inhalation Toxicology*, vol. 16, no. 6–7, pp. 391–405, 2004.

[21] E. B. Schroeder, D. Liao, L. E. Chambless, R. J. Prineas, G. W. Evans, and G. Heiss, “Hypertension, blood pressure, and heart rate variability: the atherosclerosis risk in communities (ARIC) study,” *Hypertension*, vol. 42, no. 6, pp. 1106–1111, 2003.

[22] O. B. Neto, D. T. R. S. Abate, M. Marocolo Júnior et al., “Exercise training improves cardiovascular autonomic activity and attenuates renal damage in spontaneously hypertensive rats,” *Journal of Sports Science and Medicine*, vol. 12, no. 1, pp. 52–59, 2013.

[23] Y.-J. Gui, C.-X. Liao, Q. Liu, Y. Guo, and D.-Y. Xu, “RKIP corrects impaired beta (2)-adrenergic receptor vasodilatation in hypertension by downregulation of GRK2,” *International Journal of Cardiology*, vol. 207, pp. 359–360, 2016.
S. D. Stocker, S. M. Lang, S. S. Simmonds, M. M. Wenner, and W. B. Farquhar, “Cerebrospinal fluid hypernatremia elevates sympathetic nerve activity and blood pressure via the rostral ventrolateral medulla,” *Hypertension*, vol. 66, no. 6, pp. 1184–1190, 2015.

N. Marina, R. Ang, A. Machhada et al., “Brainstem hypoxia contributes to the development of hypertension in the spontaneously hypertensive rat,” *Hypertension*, vol. 65, no. 4, pp. 775–783, 2015.

W. H. Frishman, “Beta-adrenergic receptor blockers in hypertension: alive and well,” *Progress in Cardiovascular Diseases*, vol. 59, no. 3, pp. 247–252, 2016.

G. W. Wong, H. N. Boyda, and J. M. Wright, “Blood pressure lowering efficacy of beta-1 selective beta blockers for primary hypertension,” *Cochrane Database of Systematic Reviews*, vol. 2016, no. 3, Article ID CD007451, 2016.

F. A. Flachskampf, J. Gallasch, O. Gefeller et al., “Randomized trial of acupuncture to lower blood pressure,” *Circulation*, vol. 115, no. 24, pp. 3121–3129, 2007.

L. Zhang, P. Shen, and S. Wang, “Acupuncture treatment for hypertension: a case study,” *Acupuncture in Medicine*, vol. 32, no. 1, pp. 73–76, 2014.

L. Te Riet, J. H. M. van Esch, A. J. M. Roks, A. H. van den Meiracker, and A. H. J. Danser, “Hypertension: renin-angiotensin-aldosterone system alterations,” *Circulation Research*, vol. 116, no. 6, pp. 960–975, 2015.

G. Z. Kalil and W. G. Haynes, “Sympathetic nervous system in obesity-related hypertension: mechanisms and clinical implications,” *Hypertension Research*, vol. 35, no. 1, pp. 4–16, 2012.

T. E. Lohmeier, R. Iliescu, B. Liu, J. R. Henegar, C. Maric-Bilkan, and E. D. Irwin, “Systemic and renal-specific sympathoinhibition in obesity hypertension,” *Hypertension*, vol. 59, no. 2, pp. 331–338, 2012.

K. Kario, D. L. Bhatt, S. Brar, S. A. Cohen, M. Faby, and G. L. Bakris, “Effect of catheter-based renal denervation on morning and nocturnal blood pressure: insights from SYMPLICITY HTN-3 and SYMPLICITY HTN-Japan,” *Hypertension*, vol. 66, no. 6, pp. 1130–1137, 2015.

H. T. N. I. Symplicity, “Catheter-based renal sympathetic denervation for resistant hypertension: durability of blood pressure reduction out to 24 months,” *Hypertension*, vol. 57, pp. 911–917, 2011.

SYMPLICITY HTN-2 Investigators, M. D. Edler, H. Krum et al., “Renal sympathetic denervation in patients with treatment-resistant hypertension (The Symlicity HTN-2 Trial): a randomised controlled trial,” *The Lancet*, vol. 376, no. 9736, pp. 1903–1909, 2010.

D. L. Bhatt, D. E. Kandzari, W. W. O’Neill et al., “A controlled trial of renal denervation for resistant hypertension,” *New England Journal of Medicine*, vol. 370, no. 15, pp. 1393–1401, 2014.

R. E. Schmieder, “Renal denervation—a valid treatment option despite SYMPLICITY HTN-3?” *Nature Reviews Cardiology*, vol. 11, no. 11, article no. 638, 2014.

R. E. Schmieder, “Hypertension: how should data from SYMPLICITY HTN-3 be interpreted?” *Nature Reviews Cardiology*, vol. 11, no. 7, pp. 375–376, 2014.