Effects of voluntary exercise on blood pressure, angiotensin II, aldosterone, and renal function in two-kidney, one-clip hypertensive rats

Brian M Waldman1,2  
Robert A Augustyniak1–3  
Haiping Chen1,2  
Noreen F Rossi1,2,4

1Department of Internal Medicine,  
2Department of Physiology,  
Wayne State University School of Medicine, Detroit, MI,  
3Department of Biomedical Sciences, Edward Via College of Osteopathic Medicine-Carolinas, Spartanburg, SC,  
4Department of Internal Medicine, John D Dingell Veterans Administration Medical Center, Detroit, MI, USA

Abstract: Spontaneous dynamic exercise promotes sympathoinhibition and decreases arterial pressure in two-kidney, one-clip (2K-1C) hypertensive rats. Renal sympathetic nerves stimulate renin secretion and increase renal tubular sodium reabsorption. We hypothesized that daily voluntary wheel running exercise by 2K-1C rats will decrease mean arterial pressure (MAP), plasma angiotensin II (Ang II), and aldosterone as well as normalize urinary sodium and potassium excretion independent of changes in glomerular filtration rate (GFR). Five-week-old male Sprague Dawley rats underwent sham clipping (Sham) or right renal artery clipping (2K-1C). Rats were randomized to standard caging (SED) or cages with running wheels (EX). After 12 weeks, rats were assigned to either collection of aortic blood for measurement of Ang II and aldosterone or assessment of inulin clearances and excretory function. Running distances were comparable in both EX groups. MAP was lower in 2K-1C EX vs 2K-1C SED rats (P<0.05). Plasma Ang II and aldosterone were significantly higher in 2K-1C SED rats and decreased in 2K-1C EX rats to levels similar to Sham SED or Sham EX rats. clipped kidney weights were significantly lower in both 2K-1C groups, but GFR and urine flow rates were no different from right and left kidneys among the four groups. Total and fractional sodium excretion rates from the unclipped kidney of 2K-1C SED rats were higher vs either Sham group (P<0.05). Values in 2K-1C EX rats were similar to the Sham groups. Potassium excretion paralleled sodium excretion. These studies show that voluntary dynamic exercise in 2K-1C rats decreases plasma Ang II and aldosterone, which contribute to the lower arterial pressure without deleterious effects on GFR. The effects on sodium excretion underscore the impact of pressure natriuresis despite elevated plasma Ang II and aldosterone in sedentary 2K-1C rats. In contrast, potassium excretion is primarily regulated by circulating aldosterone and distal sodium delivery.

Keywords: glomerular filtration rate, Goldblatt kidney, sodium excretion, potassium excretion, sympathetic nervous system

Introduction

Regular dynamic exercise has been shown to decrease systemic arterial blood pressure, and is accompanied by diminishing efferent sympathetic nerve activity,1,2 in humans3,4 and in several, but not all, models of hypertension in rats.5–8 In spontaneously hypertensive rats (SHR), daily volitional exercise decreases blood pressure and plasma renin activity, whereas results in stroke-prone SHR have been less consistent.5,7 In Dahl salt-sensitive rats, a model of kidney-mediated hypertension, regular dynamic exercise does not reduce systolic pressure despite decreasing the response to ganglionic blockade consistent with diminished sympathetic outflow.4 In two-kidney, one-clip (2K-1C) hypertensive rats, the type of dynamic exercise influences the hemodynamic...
response. Although neither treadmill exercise nor forced swimming exercise decreases blood pressure, voluntary wheel running exercise significantly decreases systemic arterial pressure, lowers plasma angiotensin II (Ang II) levels, and modulates renal sympathetic nerve activity in this model. The effect of exercise on both blood pressure and sympathetic activity is dependent on the severity of 2K-1C hypertension, with reductions occurring in moderate but not severe hypertension.

Improvement in structural parameters, such as collagen deposition and inflammatory infiltration, have been reported in the kidneys of exercised 2K-1C rats, but scant information exists about the effects of exercise on parameters of renal function in this model. Notably, efferent renal sympathetic nerves influence three major renal functions: renal blood flow, renin secretion, and renal tubular sodium reabsorption. Evidence from studies with renal denervation indicate that efferent renal sympathetic nerve activity exerts little or no influence on effective renal blood flow rate in either normal or hypertensive rats. In contrast, renal sympathetic nerve activation increases renin secretion leading to activation of the renin–angiotensin–aldosterone cascade. Ang II and aldosterone actions on renal tubules are known to increase renal tubular sodium reabsorption. Moreover, direct action of norepinephrine released from nerve endings onto adrenergic receptors on the tubules enhances renal sodium reabsorption independent of changes in glomerular filtration rate (GFR). Renal artery clipping and telemetry transmitter placement

Given the foregoing observations, we hypothesize that the decrease in arterial pressure seen with 12 weeks of daily wheel running exercise by moderately hypertensive 2K-1C rats is due, at least in part, to decreases in plasma Ang II and aldosterone concentrations and normalization of urinary sodium and potassium excretion independent of changes in GFR. Methods

Male Sprague Dawley rats (Harlan Sprague Dawley, Indianapolis, IN, USA) used in all the studies were housed under controlled conditions (21°C–23°C; lights on, 07:00–19:00 hours) and were permitted free access to water and standard rat chow. The rats were cared for in accordance with the principles of the National Institutes of Health Guide for the Care and Use of Laboratory Animals. All protocols were approved by the Wayne State University Institutional Animal Care and Use Committee.

Renal artery clipping and telemetry transmitter placement

Five-week-old rats were anesthetized with combined ketamine 80 mg/kg and xylazine 5 mg/kg intraperitoneally (ip). A silver clip (0.2 mm) was surgically placed around the right renal artery (2K-1C group) via a flank incision as we have previously reported. Rats in the sham-clipped (Sham) groups underwent identical surgery but were not clipped. The flank incision was closed. A venral midline incision was made in the neck and a radiotelemetry transmitter (TA11PA-C40; Data Sciences Intl) was implanted into each rat by exposing the left carotid artery, temporarily occluding the proximal end, and inserting the gel-filled catheter attached to the transmitter device into the artery with a 21-gauge needle. The catheter was advanced into the aorta and secured with medical adhesive and the transmitter was tunneled subcutaneously and sutured to the muscles between the scapulae. The incision was closed with surgical staples. At the end of surgery, each rat received butorphanol tartrate 0.075 mg subcutaneously for analgesia.

Exercise and sedentary protocols

Five days after surgery, Sham or 2K-1C rats were further assigned randomly to either sedentary (SED) or exercise
(EX) groups resulting in four groups: Sham SED, Sham EX, 2K-1C SED, and 2K-1C EX. Roughly, 15% additional rats were placed into each EX group to account for rats that would fail to exercise within 2 SDs of the overall mean distance and would require a priori removal from the protocol. SED rats were left in their individual home cages for the duration of the experiment. EX rats were placed in cages identical to the SED rats but with access to running wheels (1.1 m per revolution) equipped with an optical sensor to detect and record total revolutions (Lafayette Instruments). Running distances were averaged over 24-hour periods for each animal. Rats were allowed free access to the wheels 24 h/day, 7 days/week for 12 weeks. Any rat running <1000 m/day for 2 weeks was excluded from analysis a priori. To avoid acute exercise-induced effects, the running wheel was closed off to the EX rats for 24 hours before undergoing further testing.

Protocol 1: plasma Ang II and aldosterone measurements

After the sedentary and exercise protocols were completed, the running wheels were closed off to the exercise groups for 24 hours. Mean arterial pressure (MAP) and heart rate were assessed by telemetry in the home cage over the next 24 hours. Then, a random subset of rats from each of the four groups (Sham SED, Sham EX, 2K-1C SED, and 2K-1C EX) was chosen to assess plasma Ang II and aldosterone. This approach sought to minimize the impact of surgical stress, fluid infusion, and blood drawing from the clearance experiments that may have independently influenced the hormone levels. Each rat was anesthetized with pentobarbital sodium (50 mg/kg ip). The aorta was exposed via midline abdominal incision. A ligature was placed on distal aorta just above the bifurcation of the iliac arteries. A suture was loosely placed around the proximal aorta between the superior mesenteric and renal arteries. The aorta was rapidly cannulated and 2 mL of blood was collected for subsequent analysis of Ang II and aldosterone. Then, the rat was euthanized and the kidneys were harvested and weighed.

Protocol 2: renal clearance and excretion

A separate set of rats from each of the four groups underwent assessment of GFR and urinary sodium and potassium excretion rates using standard clearance methodology we have previously reported.22 As in Protocol 1, the running wheels were closed off to the exercised groups for 24 hours, and then hemodynamic data were recorded during the next 24 hours. Each rat was anesthetized with pentobarbital sodium (50 mg/kg ip). A 23-gauge polyethylene catheter was inserted into the right femoral artery for blood pressure monitoring and for blood sampling and into the right femoral vein for infusion. Arterial pressure was monitored continuously during the clearance periods by connecting the arterial catheter to a pressure transducer (Gould P23 XL) coupled to an amplifier (Digi0Med BPA-200). The heart rate was derived from the beat-to-beat arterial pressure pulse and averaged over 1-second intervals using custom programmed data acquisition software (Dasylab; Biotech Products). To determine GFR, a bolus of 2.0 mL/kg body weight of 8 mg/mL fluorescein isothiocyanate (FITC)-inulin (Sigma-Aldrich Co.) in 0.9% saline was infused followed by continuous infusion of 4 mg/mL at 0.055 mL/min to maintain a steady-state plasma inulin concentration and to account for fluid loss.23 The right and left ureters were cannulated with 27-gauge polyethylene catheters for urine sampling from each kidney. After a 30-minute stabilization period, urine was collected for two consecutive 15-minute time periods, and an arterial blood sample (0.2 mL) was collected from the femoral arterial catheter at the midpoint of each collection to assess plasma inulin, sodium, and potassium concentrations. Rats were then euthanized with an overdose of sodium pentobarbital (200 mg/kg intravenously). The kidneys were collected and weighed.

Analytical measurements and calculations

Plasma and urinary FITC-inulin concentrations were quantitated using a Synergy H1 microplate reader (BioTek Instruments) at Ex/Em 485/535. Plasma and urinary sodium and potassium concentrations were determined with flame photometry with internal lithium standardization (Cole Parmer Instruments). Urine volume was assessed gravimetrically. GFR as well as urinary excretion rates of sodium and potassium and fractional excretion of sodium and potassium were assessed using standard formulae. Inulin clearance values were expressed after correction for body weight (mL/min/kg BW) as well as for kidney weight (mL/min/g KW).

Plasma samples for Ang II were processed as described previously24 with minimal modification. Briefly, 1 mL of plasma was extracted twice with 90% methanol. The extracts were combined, taken to dryness under nitrogen, and stored overnight at ~20°C. The extracts were reconstituted in 0.5 mL assay buffer consisting of 50 mM Na phosphate, 1 mM ethylenediaminetetraacetic acid, 0.25 mM thimerosal, and 0.25% peptidase-free human serum albumin. All samples were assayed in duplicate using 125I-labeled Ang II (Perkin-Elmer, Billerica, MA, USA) as the tracer and anti-Ang II antibody (Peninsula Laboratories, San Carlos, CA, USA) at final dilution of 1:660,000. Nonspecific binding was 2.1%, the lower
limit of detection was 0.5 fmol/tube, and 50% binding was 15.2 fmol/tube. Plasma aldosterone was assessed directly on plasma with 125I-aldosterone as the tracer in the solid phase Coat-a-Count radioimmunoassay system (Siemens).

Statistical analysis
Values are presented as mean ± SE. All comparisons among groups were made by one-way analysis of variance with the Tukey–Kramer post hoc analysis for multiple comparisons. Comparisons within the same group of rats were analyzed by paired t-test. A P-value <0.05 was taken as significant.

Results
Running distances by the Sham EX and 2K-1C EX rats over the 12 weeks were comparable for Protocol 1 (Figure 1, upper panel) and Protocol 2 (Figure 1, lower panel). The 2K-1C rats in Protocol 2 tended to run greater distances early in the experiment but distances became similar after week 5. One 2K-1C rat in Protocol 2 consistently ran <800 m/day for 3 weeks and was excluded from the study by design.

Figure 1 Running distances in the rats in Protocols 1 and 2.
Notes: Average running distances over the course of 12 weeks of voluntary wheel running for Sham EX (n=12) and 2K-1C EX (n=7) rats in Table 1 (Protocol 1; upper panel) and for Sham EX (n=7) and 2K-1C EX (n=7) rats in Table 2 (Protocol 2; lower panel). Values are mean ± SE; *P<0.05 vs Sham.
Abbreviations: 2K-1C, two-kidney, one-clip; Sham, sham clipped; EX, exercise.

Protocol 1
Parameters for rats in Protocol 1 are provided in Table 1. Body weights of Sham and 2K-1C rats were similar at the beginning of Protocol 1. At the end of the 12 weeks, body weights of both EX groups were similar to each other but were significantly less than their respective SED groups. This occurred despite equivalent caloric intake (average ~14 g chow/day/rat in all groups). Kidney weights of the right (clipped) kidney were significantly lower than those of the left kidney in both 2K-1C groups. Diurnal variation during the brief 24-hour period of telemetric data collection appeared to be preserved and the differences among the groups during daytime and nighttime MAP were similar (data not shown). MAP assessed by telemetry and averaged over the entire 24-hour period was significantly lower in the 2K-1C EX group compared with the 2K-1C SED group. Exercise training did not result in a difference in MAP in the Sham EX rats but this group displayed a significant bradycardic effect compared with Sham SED. The 2K-1C SED rats exhibited significantly elevated levels of plasma Ang II and aldosterone compared with either Sham SED or Sham EX rats. Plasma Ang II and aldosterone in the 2K-1C EX rats decreased significantly compared with 2K-1C SED rats and were not significantly different from those in either of the Sham groups.

Protocol 2
Average body weights at the end of 12 weeks in Protocol 2 did not differ significantly among the four groups (Table 2). Initial weights were also similar among the groups (data not shown). MAP assessed by telemetry was higher in the 2K-1C SED group (189.5±8.2 mmHg) compared with either Sham SED (132.0±2.8 mmHg) or Sham EX (133.9±2.7 mmHg) groups (P<0.01). MAP in the 2K-1C EX rats was lower in the 2K-1C EX group (162.7±6.7 mmHg) compared with the 2K-1C SED rats (P<0.05), but remained higher than in either of the Sham groups (P<0.05). Similar to measurements in conscious rats acquired by telemetry, MAP in 2K-1C SED rats after anesthesia for clearance studies was lower (P<0.05 vs telemetry) than in the conscious state but remained significantly higher than MAP in any of the other groups (Table 2). Notably, the decrease in MAP with anesthesia was less evident in the Sham groups and the 2K-1C EX group and did not achieve significance (~1–5 mmHg; P>0.05).

As in Protocol 1, the Sham EX group displayed a significantly lower heart rate (372±10 bpm) compared with the Sham SED group (425±11 bpm, P<0.05). The 2K-1C EX group had a lower heart rate (384±6 bpm) compared with 2K-1C SED group (402±10 bpm), but this did not achieve significance (P>0.05).
The clipped kidneys exhibited significantly lower kidney weights. GFRs did not differ among the groups. When GFR was expressed per body weight, there was no difference between left and right kidneys. When GFR was expressed per kidney weight, in light of the differences in kidney weights, the clipped kidney of the 2K-1C SED group was significantly higher than that of the nonclipped, left kidney.

Plasma sodium concentrations were similar among the groups. Plasma potassium levels were significantly lower in the 2K-1C SED rats compared with values in the three other groups (Table 2). Figure 2 shows that the left unclipped kidneys in the 2K-1C SED group displayed higher sodium excretion rates than the left kidneys of either of the Sham groups. Sodium excretion by the unclipped kidney of the 2K-1C EX group was significantly lower than that of the 2K-1C SED group and did not differ from that of either of the Sham groups. Sodium excretion by the clipped kidneys did not differ among any of the groups. Fractional sodium excretion exhibited the same pattern among the groups (Figure 3).

Potassium excretion rates from the unclipped kidneys were higher in the 2K-1C SED rats than either Sham SED or Sham EX rats (Figure 2). Potassium excretion was also higher from the unclipped kidneys in the 2K-1C SED rats compared with Sham EX rats. The 2K-1C EX group displayed potassium excretion rates from both left and right kidneys that were significantly lower than the corresponding kidneys of the 2K-1C SED group. Potassium excretion from right or left kidneys did not differ from the corresponding kidneys in either of the Sham groups. Likewise, fractional excretion of potassium from the left unclipped kidney was significantly higher in the 2K-1C SED group compared with either Sham group and was significantly lower from the left kidney of the 2K-1C EX group (Figure 3).

**Discussion**

As the prevalence of renovascular hypertension rises, knowledge about the impact of regular voluntary dynamic exercise on blood pressure and renal function in this condition becomes vital. The present study reports several findings in the 2K-1C rat, a model of unilateral renal artery stenosis. First, daily exercise for 12 weeks decreased arterial pressure in 2K-1C hypertensive rats. Second, with daily exercise in 2K-1C rats, elevated plasma Ang II and aldosterone levels returned to levels not different from those observed in the sham-clipped rats. Third, GFR did not differ between sedentary sham and 2K-1C groups and, further, was not altered by exercise in either group. Fourth, both absolute and fractional urinary sodium and potassium excretion values from the nonclipped kidneys of sedentary 2K-1C rats was significantly elevated compared with that of sham-clipped rats. Notably, urinary sodium and potassium excretion by the 2K-1C exercised rats was not different from that of sham-clipped normotensive rats.

**Hemodynamic response to exercise training**

Our data indicate that voluntary wheel running exercise in 2K-1C rats decreased arterial pressure when measured by telemetry in conscious freely moving rats or by direct arterial pressure recording in pentobarbital-anesthetized rats. The arterial pressure–lowering effect of exercise training was most evident in the hypertensive 2K-1C group rather than the sham-clipped normotensive rats. The blood pressure–lowering effect of exercise training in 2K-1C hypertension has been observed by some investigators, but not others. These discrepancies may be related to the type of exercise regimen, the technique used for assessing blood pressure, or the presence, type, or depth of anesthesia. For example, an 8- to 12-week forced treadmill regimen did not decrease arterial pressure. Forced swimming exercise over a 4-week period reduced arterial pressure to an extent similar to that seen in the present study. However, Maia et al showed that additional weeks of forced swimming resulted in no difference between exercise and sedentary 2K-1C groups.
**Table 2** Body weight, kidney weight, mean arterial pressure, plasma sodium and potassium concentrations, and renal function data in anesthetized sham-clipped and two-kidney, one-clip hypertensive rats after 12-week sedentary or exercise conditions (Protocol 2)

<table>
<thead>
<tr>
<th>Groups</th>
<th>n</th>
<th>BW (g) ± SE</th>
<th>MAP (mmHg)</th>
<th>Plasma (Na⁺) (mM)</th>
<th>Urine flow (μL/min/g) KW</th>
<th>GFR (mL/min/g) Left</th>
<th>GFR (mL/min/g) Right</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sham SED</td>
<td>6</td>
<td>416±20</td>
<td>131±0.10</td>
<td>146±6.0.6</td>
<td>1.37±0.11</td>
<td>146±6.0.6</td>
<td>1.37±0.11</td>
</tr>
<tr>
<td>Sham EX</td>
<td>7</td>
<td>403±8</td>
<td>135±2</td>
<td>146±6.0.6</td>
<td>1.43±0.06</td>
<td>146±6.0.6</td>
<td>1.43±0.06</td>
</tr>
<tr>
<td>2K-1C SED</td>
<td>6</td>
<td>428±9</td>
<td>170±4³</td>
<td>146±6.0.6</td>
<td>1.78±0.10</td>
<td>146±6.0.6</td>
<td>1.78±0.10</td>
</tr>
<tr>
<td>2K-1C EX</td>
<td>7</td>
<td>398±18</td>
<td>158±3</td>
<td>145±2.0.8</td>
<td>1.19±0.06</td>
<td>145±2.0.8</td>
<td>1.19±0.06</td>
</tr>
</tbody>
</table>

Notes: Values are mean ± SE. Right kidneys clipped. *P* < 0.05 vs Sham SED. **P** < 0.05 vs Sham EX. ***P*** < 0.05 vs Sham SED; °P < 0.05 vs sham kidney; #P < 0.05 vs 2K-1C SED.

Abbreviations: BW, body weight; KW, kidney weight; MAP, mean arterial pressure; SED, sedentary; 2K-1C, two-kidney, one-clip.

Plasma Ang II and aldosterone levels

Plasma Ang II and aldosterone levels in both groups of sham-clipped rats and sedentary 2K-1C rats were elevated similar to values reported previously. Daily exercise clearly decreased plasma Ang II and aldosterone levels to concentrations similar to those in sham-clipped animals. Taken together with evidence indicating that daily exercise attenuates renal sympathetic nerve activity and that sympathetic inputs to the kidney regulate activity of the renin–angiotensin–aldosterone system, it is possible that exercise-induced augmentation of parasympathetic control of heart rate may be less effective in 2K-1C than sham-clipped normotensive rats or that renal sympathetic tone may decrease to a greater extent than cardiac sympathetic activity. It should be noted that the study was not powered to evaluate changes in heart rate, which tends to have greater variability.
that sympathoinhibition due to exercise can attenuate activation of the renin–angiotensin–aldosterone system in 2K-1C hypertension. As circulating Ang II and aldosterone promote renal sympathoexcitation, the decline in both of these hormones may, in turn, further contribute to the sympathoinhibitory effect of regular exercise and lowering of arterial pressure in this model.

Glomerular filtration rate

In the present study, GFR did not differ significantly between sedentary sham-clipped and 2K-1C groups. This has been observed in some, but not all, earlier studies of the 2K-1C model and may have depended on the size of the clip, the age of the rat at the time of clipping, and the time after clipping at which renal function was assessed.

In addition, GFR remained unchanged in exercised 2K-1C rats despite lowering arterial pressure and Ang II. Our previous studies have shown that voluntary wheel running exercise inhibits renal sympathetic nerve activity. A possible explanation is that the combined effects of sympathoinhibition and lower Ang II result in a greater decrease in afferent vs efferent arteriolar resistance thereby...
maintaining intraglomerular pressure and GFR. These data are consistent with existing studies demonstrating that pharmacologic inhibition of Ang II actions alone does not decrease glomerular filtration.\textsuperscript{42,45,46} In addition, glomerular filtrate rate remains unchanged when Ang II is blocked after acute renal denervation in 2K-1C rats.\textsuperscript{43} Thus, regular dynamic exercise does not exert a detrimental effect on GFR in moderately hypertensive 2K-1C rats. The small but significant increase in GFR from the clipped kidney of sedentary 2K-1C when expressed per kidney weight may be due to a more complex interplay of the sympathetic nervous system and both systemic and intrarenal renin–angiotensin systems. While the elevated systemic Ang II levels may serve to preserve GFR via preferential efferent arteriolar vasoconstriction, intrarenal Ang II has been known to modulate tubuloglomerular feedback favoring afferent vasoconstriction.\textsuperscript{47} Intrarenal Ang II was high in the clipped kidney 1 week after renal artery clipping; however, after 9 weeks, renal cortical Ang II decline levels were no different from those in sham-clipped rats.\textsuperscript{20} Thus, the timing after clipping may also contribute substantially to the impact on GFR. It is plausible that efferent arteriolar vasoconstriction in response to systemic Ang II, which is threefold higher in the sedentary 2K-1C group, serves to maintain GFR at 12 weeks. Studies designed to assess these complex interactions would be required.

Figure 3 Fractional excretion of sodium and potassium.

Notes: Fractional excretion of sodium (upper panel) and potassium (lower panel) for left and right kidneys in the four groups. Values are mean ± SE; n as in Table 2. *P<0.01 vs Sham SED; \$P<0.05 vs Sham EX; †P<0.001 vs left kidney; and ‡P<0.05 vs 2K-1C SED.

Abbreviations: 2K-1C EX, two-kidney, one-clip exercise; 2K-1C SED, two-kidney, one-clip sedentary; Sham EX, sham-clipped exercise; Sham SED, sham-clipped sedentary.
Excretory function
The higher urinary sodium excretion rate by the unclipped kidney but not the clipped kidney in the sedentary 2K-1C group reflects the substantial influence that renal perfusion pressure exerts via pressure natriuresis. Our findings are consistent with studies showing that acutely elevated levels of Ang II are antinatriuretic when renal perfusion pressure is prevented from increasing but natriuretic when renal perfusion pressure is permitted to rise. A similar excretory pattern has been observed in 2K-1C rats when renal perfusion pressure is controlled. This would not, however, fully explain how urinary sodium excretion was virtually identical to that in the sham- clipped groups although arterial pressure was still significantly elevated in the exercise-trained 2K-1C group. Thus, only a component of the decrease in sodium excretion can be attributed to a decrease in renal perfusion pressure.

Together with the known inhibition of renal sympathetic activity, the observed decline in plasma Ang II would have predicted an increase in urinary sodium excretion with regular exercise in the 2K-1C group. Consistent with our findings, renal sympathetic innervation is paradoxically permissive for sodium excretion in response to exogenous Ang II in the canine split bladder model. Despite a decrease of plasma Ang II to values similar to sham-clipped animals, denervation of the clipped kidney results in a profound decrease in renal sympathetic nerve activity but no change in intrarenal Ang II to the unclipped kidney. Although renal tissue Ang II was not measured in the current study, it is possible that persistent tubular effects by intrarenal Ang II contributed to the decrease in sodium excretion by the exercise-trained 2K-1C rats. Furthermore, as these short-term clearance experiments were performed at the end of the 12-week exercise regimen at a time when the animals were likely in electrolyte balance, we cannot exclude the possibility that a greater or lesser amount of sodium was excreted earlier in the course by the exercised 2K-1C rats. In contrast to the results with sodium excretion, the impact on urinary potassium excretion appears more straightforward. The elevated plasma aldosterone levels in the sedentary 2K-1C rats contributed to the greater urinary potassium excretion by the unclipped kidney despite lower plasma potassium levels. Increased sodium delivery to the distal nephron segments also potentially increased potassium secretion despite a counter-regulatory effect by Ang II. By decreasing aldosterone, daily exercise attenuated urinary potassium losses and restored plasma potassium levels. In addition, exercise-induced sympathoinhibition may have further contributed to the lower urinary potassium excretion in the exercised 2K-1C rats, thereby further supporting the dual action of diminished activation of the renin-angiotensin-aldosterone system and sympathetic nervous system.

Summary and perspectives
In summary, these studies support the concept that voluntary dynamic exercise lowers systemic arterial pressure and decreases plasma Ang II and aldosterone levels in the 2K-1C model of renovascular hypertension without deleterious effects on GFR. The differences among the groups in sodium excretion underscore the impact of pressure natriuresis despite elevated plasma Ang II and aldosterone in the sedentary 2K-1C rats. In contrast, potassium excretion appears to be primarily regulated by ambient aldosterone levels and distal sodium delivery. Importantly, atherosclerotic renal artery stenosis is increasing in prevalence, particularly in high-risk populations, and carries a high risk of cardiovascular morbidity and mortality. In addition to the beneficial effects on arterial pressure, regular dynamic exercise may be of benefit via diminished circulating Ang II and aldosterone, both of which contribute to deleterious cardiac and vascular remodeling. Existing studies in humans evaluated renal function during exercise and suggested a detrimental effect in renovascular hypertension. To our knowledge, follow-up investigations of the impact of regular exercise on renal function have not occurred. Our findings strongly support the possibility of beneficial long-term effects and need for further studies on the cardiovascular and renal impact of regular dynamic exercise in humans with renovascular hypertension.

Acknowledgment
This study was supported by a Merit Award from the Department of Veterans Affairs to NFR.

Disclosure
The authors report no conflicts of interest in this work.

References


Exercise and renal function in 2K-1C rats


