Hemodynamic and renal effects of acute and progressive nitric oxide synthesis inhibition in anesthetized dogs

ALEIX CASES, JOHN HAAS, JOHN C. BURNETT, AND JUAN CARLOS ROMERO
Department of Physiology, Mayo Foundation, Rochester, Minnesota 55905

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However, changes of renal blood flow (RBF) relative to any other vascular bed and systemic hemodynamics have not been evaluated. Hence, our interest in knowing if the acute inhibition of NO produced in the dog, by progressively increasing doses of NO-synthesis inhibition in anesthetized dogs. The NO-synthesis inhibition in anesthetized dogs. The NO-synthesis inhibition in anesthetized dogs. NO synthesis inhibition induces a progressive antidiuretic and antinatriuretic effect, which is partially offset by the increase in blood pressure.

N\textsuperscript{\textit{G}}-nitro-L-arginine methyl ester; regional blood flows; systemic hemodynamics; renal function; urinary sodium excretion

IT HAS BEEN WIDELY DEMONSTRATED that the synthesis of nitric oxide (NO) by endothelial cells plays a critical role in the regulation of the main determinants of blood pressure control: vascular tone (vascular resistance) and tubular sodium reabsorption (volemia). Inhibition of NO synthesis results in an increase in blood pressure due to an increase in peripheral vascular resistance and an antinatriuretic effect (18, 19, 21, 26, 28).

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HEMODYNAMIC AND RENAL EFFECTS OF NITRIC OXIDE INHIBITION IN DOGS

The right external jugular vein was cannulated with a 7F Swan-Ganz catheter (CritiCath, Ohmeda, Oxnard, CA) and advanced into the pulmonary artery for measuring cardiac output (CO) and cardiac filling pressures. The right femoral artery was cannulated to collect peripheral arterial blood samples and for continuous measurement of MAP with the use of a pressure transducer (Pd23 ID, Statham, Hato Rey, PR) connected to a chart recorder (2600, Gould, Cleveland, OH). The femoral vein was cannulated for infusion of vehicle or N®-nitro-l-arginine methyl ester (1 ml/min) and 2% inulin solution at a rate of 1 ml/min with additional anesthetic as necessary.

Transonic flow probes (Transonic #3R, Transonic Systems, Ithaca, NY) were placed through a left flank incision on the left renal, superior mesenteric, and left iliac arteries in segments proximal to the aorta. The left ureter was cannulated with PE-200 tubing for urine collection.

After completion of the surgical procedure, dogs were allowed to stabilize for 60 min without intervention. At the end of this period, a 15-min baseline clearance period was obtained. After this baseline period, vehicle or increasing doses of N®-nitro-l-arginine methyl ester (0.1, 1, 10, and 50 μg·kg⁻¹·min⁻¹) were sequentially intravenously administered during 45-min periods. During the last 15 min of each period, a clearance period was obtained. During the clearance periods, serum and urine samples were collected to measure electrolytes, inulin concentration, hemodynamic parameters, and CO.

At the end of the stabilization period and at the end of N®-nitro-l-arginine methyl ester infusion, blood samples were also obtained for measuring plasma endothelin (ET) and plasma renin activity (PRA). Plasma and urinary sodium were measured by flame photometry (IL943, Instrumentation Laboratory, Lexington, MA). GFR was calculated by measuring the clearance of inulin. Plasma and urine inulin were measured by a colorimetric method (8). Plasma ET and PRA were measured by radioimmunoassay with the use of commercial RIA kits.

CO was measured by a thermodilution technique with a CO computer (model 9520A, Edwards Laboratories, Santa Ana, CA) and considered as the average of three measurements. Calculated hemodynamic parameters included: systemic vascular resistances (SVR) = MAP – right arterial pressure/CO. Pulmonary vascular resistances are equal to pulmonary artery pressure minus pulmonary capillary wedge pressure divided by CO.

Animals were euthanized at the end of each experiment without emerging from anesthesia by an intravenous injection of 200 meq potassium chloride.

Statistics. Data are expressed as means ± SE. The analysis of the data included a repeated-measures ANOVA and a Scheffe’s test to assess differences in the means between periods. To assess differences between control dogs and the N®-nitro-l-arginine methyl ester group, ANOVA was performed for each dependent variable, using dog, time, and drug terms as the independent variables. The drug effects were then added to the residuals of this ANOVA model, which created a data set adjusted for dog and time. Two-sample comparisons (t-tests with the Bonferroni correction) were then performed on the adjusted data to determine the significance of N®-nitro-l-arginine methyl ester. Differences between baseline and the last infusion period in hormonal variables were assessed by means of a paired t-test. A value of P < 0.05 was considered as statistically significant.

RESULTS

As can be seen in Fig. 1, MAP increased in a dose-dependent manner during the infusion of N®-nitro-l-arginine methyl ester [from 125.4 ± 3.7 to 127.7 ± 4.1, 130.1 ± 4.3, 135.8 ± 4.2*, and 144.9 ± 4.91* (*P < 0.05 vs. baseline)]. However, these increments did not achieve statistical significance with respect to the baseline period or the control group until the doses of 10 and 50 μg·kg⁻¹·min⁻¹ were administered. At this point, the overall increase of MAP was 16% over the basal period. MAP of the control group did not experience any significant change throughout the experiment [from 129.3 ± 1.85 to 132 ± 2.7, 129.4 ± 3.9, 127.4 ± 4.8, and 126.1 ± 4.4 mmHg, P not significant (NS)].

CO also decreased in a progressive dose-dependent manner in the experimental group [from 3.13 ± 0.14 to 2.95 ± 0.16, 2.74 ± 0.16*, 2.33 ± 0.14*, and 1.98 ± 0.15* l/min (P < 0.05 vs. baseline)], which became statistically significant during the infusion of 1 μg·kg⁻¹·min⁻¹, before MAP was significantly in-

Fig. 1. Response of mean arterial pressure (MAP), cardiac output (CO), and pulmonary arterial pressure (PAP) after progressive doses (0.1, 1.0, 10, 50 μg·kg⁻¹·min⁻¹) of N®-nitro-l-arginine methyl ester (L-NAME) and saline in a time control group. *P < 0.05 vs. baseline; and #P < 0.05 vs. control.
creased (Fig. 1). The fall in CO was already significant during the infusion of 1 μg·kg⁻¹·min⁻¹ compared with the control group, in which the CO remained unaltered until the last period (period 4) when it decreased by ∼10% [from 3.24 ± 0.27 to 3.2 ± 0.28, 3.1 ± 0.23, 3.03 ± 0.23, and 2.92 ± 0.23 dyn·s·cm⁻5 (*P < 0.05 vs. baseline)]. SVR of the experimental group changed in a parallel fashion with respect to MAP [from 40.43 ± 1.9 to 43.65 ± 1.9, 47.7 ± 1.97, 58.1 ± 2.38, and 73.93 ± 4.78 dyn·s·cm⁻5 (*P < 0.05 vs. baseline)]. The increments in SVR became significant during the last two periods (10 μg and 50 μg·kg⁻¹·min⁻¹) with respect to both the baseline period and the control group. SVR in the control group remained unchanged throughout the experiment (from 41.6 ± 3.7 to 42.9 ± 3.4, 42.8 ± 2.28, 43.03 ± 2.17, and 44.2 ± 2.35 dyn·s·cm⁻5).

In the experimental group, there was a continuous and progressive increase in pulmonary vascular resistance [from 3.23 ± 0.34 to 3.4 ± 0.34, 3.72 ± 0.33, 3.98 ± 0.38, and 4.2 ± 0.37 dyn·s·cm⁻5 (*P < 0.05 vs. baseline)], which was reflected in parallel elevations of pulmonary arterial pressure (PAP) [from 14.6 ± 0.41 to 15.4 ± 0.51, 17.9 ± 0.71, 20.8 ± 0.98, and 23.2 ± 0.98 mmHg (*P < 0.05 vs. baseline)] (8.6 mmHg at the end of the study) (Fig. 1). Both of these parameters were significantly increased by the 1-μg·kg⁻¹·min⁻¹ dose. Pulmonary vascular resistances increased in the last period by 2.45-fold, which is greater than the increase recorded in SVR (1.82-fold). In the control group, pulmonary vascular resistance [from 3.23 ± 0.34 to 3.4 ± 0.34, 3.72 ± 0.33, 3.98 ± 0.38, and 4.2 ± 0.37 (*P < 0.05 vs. baseline)] (30%) and pulmonary artery pressure [from 13.2 ± 0.33 to 13.6 ± 0.4, 14.25 ± 0.35, 14.9 ± 0.23, and 15.4 ± 0.29 dyn·s·cm⁻5 (*P < 0.05 vs. baseline)] also experienced a mild increase from the second infusion (∆2.2 mmHg, P < 0.05). The differences in mean pulmonary pressure and pulmonary vascular resistances between the experimental and control groups reached significance during the last two infusion periods.

As it is shown in Fig. 2, in the experimental group, mesenteric blood flow (MBF) [from 203.8 ± 21.1 to 206.1 ± 21.5, 206.8 ± 20.1, 188.1 ± 17.4, and 153.8 ± 16.8 ml/min (*P < 0.05 vs. baseline)] and RBF [from 185 ± 24.5 to 184.3 ± 23.9, 178.3 ± 22.2, 161 ± 17.3, and 145 ± 16.4 ml/min (*P < 0.05 vs. baseline)] both are decreased by approximately the same proportion, 25 and 22%, respectively. These decrements occur only during the highest dose of N⁵-nitro-L-arginine methyl ester infusion. This contrasted with the decrease in iliac blood flow (IBF) [from 105.7 ± 8.7 to 98.29 ± 7.8, 94 ± 8.7, 81.43 ± 9.8, and 60.3 ± 7.7 ml/min (*P < 0.05 vs. baseline)], which occurred earlier and was more marked. In the control group, MBF (from 225.4 ± 13.5 to 222.3 ± 13.0, 227.4 ± 13.4, 233.7 ± 12.8, and 232.9 ± 15.6 ml/min) and IBF (142.6 ± 10, 141.4 ± 9.7, 142 ± 12.3, 132.9 ± 10.7, and 130.9 ± 9.9 ml/min) did not significantly change, whereas RBF [from 159.4 ± 6.21 to 158.9 ± 8.35, 177.7 ± 13.4, 187.7 ± 13.4, and 185.1 ± 13.6 ml/min (*P < 0.05 vs. baseline)] underwent a progressive elevation of 16%, which became statistically significant from period 3. Furthermore, between the experimental and time control groups, there were significant differences in IBF from the dose of 1 μg·kg⁻¹·min⁻¹ and in the MBF and RBF from the dose of 10 μg·kg⁻¹·min⁻¹.

As shown in Fig. 3, GFR did not change significantly in either group (N⁵-nitro-L-arginine methyl ester group from 44.45 ± 5.16 to 38.5 ± 5.1 ml/min and control group from 31.14 ± 1.5 to 33.6 ± 2.2 ml/min at the last infusion period, NS). Urine volume (UVol) (from 0.18 ± 0.05 to 0.21 ± 0.05, 0.29 ± 0.07, 0.29 ± 0.07, and 0.25 ± 0.06, NS), urine sodium excretion (UNa) (from 45.93 ± 16.5 to 55.3 ± 14.5, 69.1 ± 16.1, 62.73 ± 14.5, and 39.9 ± 11.6 μeq/min), and fractional excretion of sodium (FeNa) (1.11 ± 0.43, 1.3 ± 0.34, 1.62 ± 0.37, 1.67 ± 0.39, and 1.03 ± 0.25, NS) did not exhibit any significant change in the experimental group. These last three parameters showed a marked tendency toward elevation in the control group, which

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Fig. 2. Response of renal, iliac, and mesenteric blood flows after progressive doses (0.1, 1.0, 10, 50 μg·kg⁻¹·min⁻¹) of L-NAME and saline in a time control group. *P < 0.05 vs. baseline; #P < 0.05 vs. control.
became significant during the last two periods for UNa [from 38.12 ± 9.1, 37.58 ± 9.22, 48.25 ± 13.42, 60.47 ± 11.35*, and 70.11 ± 12.7* μeq/min (*P < 0.05 vs. baseline)] and FeNa [from 1.06 ± 0.23 to 1.07 ± 0.24, 1.19 ± 0.23, 1.57 ± 0.29*, and 1.85 ± 0.32* % (*P < 0.05 vs. baseline)] and during the last period for UVol [from 0.17 ± 0.03 to 0.17 ± 0.03, 0.19 ± 0.03, 0.25 ± 0.05, and 0.31 ± 0.06* ml/min (*P < 0.05 vs. baseline)]. The differences between the two groups in these three parameters became significant during the last infusion period.

PRA decreased in the N\textsuperscript{G}-nitro-L-arginine methyl ester group (from 3.01 ± 0.99 to 1.84 ± 0.77 ng·ml\(^{-1}·\text{h}^{-1}\), *P < 0.05), but it did not change in the vehicle group (from 5.05 ± 0.81 to 4.48 ± 0.87 ng·ml\(^{-1}·\text{h}^{-1}\), NS). Plasma ET levels increased after \textit{N}\textsuperscript{G}-nitro-L-arginine methyl ester infusion (from 19.35 ± 0.75 to 25.4 ± 1.4 pg/ml, *P < 0.05), but not in the control group (from 22.1 ± 2.9 to 23.1 ± 2.9, NS).

**DISCUSSION**

**Changes in systemic and pulmonary circulation.** This study shows that basal release of NO regulates blood pressure and modulates the vasculature of systemic and pulmonary circulation. In systemic circulation, however, the sensitivity of different vascular beds in response to NO synthesis inhibition is not homogeneous. Such findings agree to that reported by many other investigators (9, 10, 25, 33, 34, 38). The IBF exhibited dependency on the synthesis of NO, which was greater than that exhibited by other vascular beds. Similar results have been observed by some authors (9, 10), but not by others (33, 38). In fact, the iliac vasculature developed an early vasoconstriction at doses of \textit{N}\textsuperscript{G}-nitro-L-arginine methyl ester (1 μg·kg\(^{-1}·\text{min}^{-1}\)) that was not seen in the renal or mesenteric vascular beds. At maximal doses of \textit{N}\textsuperscript{G}-nitro-L-arginine methyl ester (50 μg·kg\(^{-1}·\text{min}^{-1}\)), the reduction of IBF was almost twofold higher than that produced in RBF and MBF. This effect is important, because the role of NO in the control of skeletal muscle blood flow has been linked to exercise performance (29) and to the amount of glucose uptake that, when decreased, could lead to situations comparable to those observed during insulin resistance (1). It should be pointed out that the mesenteric vasculature was the least dependent on changes in NO synthesis, because it experienced a 25% decrease only at the highest doses of \textit{N}\textsuperscript{G}-nitro-L-arginine methyl ester. Renal vasculature exhibited an intermediate response between the iliac and the mesenteric, considering that RBF increased by 16% in time control animals.

Our results also showed that inhibition of NO synthesis has a major impact on pulmonary circulation, which is an observation previously reported by others (7, 14, 23, 38). In our study, the increase in pulmonary vascular resistance (+146%) and in PAP (+59%) was greater compared with the changes observed in SVR (+83%) and systemic blood pressure (+19.5%), at a dose of 50 μg·kg\(^{-1}·\text{min}^{-1}\). Previous studies reported similar results (7, 38), but this is not a universal finding (23).

**Changes in renal hemodynamics and sodium excretion.** In previous studies, intrarenal infusion of NO synthesis inhibitors induced potent renal vasoconstriction and antinatriuresis (19). However, Baylis et al. (2) showed that inhibition of NO synthesis causes a dose-dependent increase in arterial blood pressure and sodium excretion in both conscious and anesthetized rats. In the experimental group of animals, RBF did not decrease significantly until the last period. However, these decrements in blood flow became more apparent if one compares it with the steady increments observed in the control group, which became significant from the third period. This renal vasoconstriction was not accompanied by any change in GFR, thus suggesting that the increase in resistance was taking place simultaneously in glomerular afferent and efferen-

![Graph](https://via.placeholder.com/150)
ent arterioles. This assumption is supported by the finding of other investigators, who have reported similar findings in dogs (5, 23). Schnackenberg et al. (31) and Juncos et al. (13) observed that inhibition of NO renders the afferent preglomerular arteries susceptible to be constricted by doses of ANG II that produce only efferent vasoconstriction under normal circumstances when NO synthesis is not inhibited.

UNa and FeNa in the experimental group showed a tendency to increase in proportion to the increase in MAP, except during the last two periods in which UNa and FeNa declined toward baseline levels. This antinatriuretic effect occurred despite the marked increase in blood pressure. Furthermore, the decrease of sodium excretion is more noticeable when compared with the natriuresis observed in the control group of dogs, which blood pressure did not change, suggesting that N\textsuperscript{G}-nitro-L-arginine methyl ester induced a dose-dependent antinatriuretic effect that was able to completely blunt pressure natriuresis at the highest dose administered in this study. This phenomenon has been reported by others in rats and dogs (19, 26, 27). The importance of NO in the regulation of sodium excretion is underscored by the demonstration that NO synthesis inhibition blunts both pressure-induced (20) and volume-induced natriuresis (17).

A decrease in PRA was observed at the end of the experiment in the N\textsuperscript{G}-nitro-L-arginine methyl ester group, whereas no changes were observed in the control group. Studies on the effects of NO on renin release have yielded disparate results. In early in vitro studies, NO synthesis inhibition increased renin release (3, 39). However, more recent studies indicate a more complex interaction between NO and renin secretion (16, 36). Although several authors reported decreases in plasma renin levels during systemic NO synthesis inhibition (12), other authors observed no changes (5) or even increases (32). Systemic inhibition of NO synthesis is also associated with increases in blood pressure, stimulation of sympathetic nerve activity (35), and increases in other vasoactive factors (4) that can have different directional effects on renin release. Schnackenberg et al. (30) recently demonstrated, in an elegant study, that intrarenal infusion of N\textsuperscript{G}-nitro-L-arginine methyl ester did not alter renal perfusion pressure or GFR and stimulates renin release in the dog and that the macula densa is important in mediating this effect.

An increase in plasma ET was observed in the N\textsuperscript{G}-nitro-L-arginine methyl ester group but not in the control group, in agreement with previous in vivo and in vitro studies (4, 23). The increase in the plasma levels of this vasoconstrictor peptide may partly account for the hemodynamic, renal, and renin-angiotensin system changes observed during NO synthesis inhibition. Recent studies in rats indicate that ET is partially involved in the hemodynamic changes during NO inhibition, but its role in the renal effects in this situation is controversial (6, 22, 24, 37).

It should be noted that in the control dogs, the administration of 2 ml of saline per minute produced a mild volume expansion reflected in an increase in RBF, UVol, and UNa in the last two experimental periods. However, these changes were not sufficient to alter MAP, CO, IBF, MBF, GFR, PRA, and plasma ET. For these reasons, we believe that the vasodilator actions do not preclude the overall conclusions about the vasoconstrictor effects exerted by NO on different vascular territories.

In summary, this study shows that in anesthetized dogs, the progressive inhibition of NO synthesis with doses of N\textsuperscript{G}-nitro-L-arginine methyl ester that range from 0.1 to 50 µg·kg\textsuperscript{-1}·min\textsuperscript{-1} produces a marked elevation of pulmonary vascular resistance and pulmonary blood pressure, which is almost doubled from that observed in SVR. Systemic vascular territories, on the other hand, do not respond in a uniform manner. The iliac vascular bed is by far the most sensitive, whereas the renal vascular territories supplied by the mesenteric artery do not seem to be significantly affected. Renal vascular territory exhibits an intermediate response. The volume of blood passing through skeletal muscle may be important, because the progressive fall induced by the administration of N\textsuperscript{G}-nitro-L-arginine methyl ester correlated with the CO. Finally, inhibition of NO produces a decrease in RBF with no changes in GFR. Under these conditions, pressure natriuresis is not manifested because of the opposing effects of NO inhibition on sodium excretion.

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REFERENCES


