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In Vivo Transfer of Nitric Oxide between a Plasma Protein-bound Reservoir and Low Molecular Weight Thiols

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Abstract

Plasma albumin reacts with nitric oxide (NO) to form the bioactive adduct, S-nitroso-albumin (S-NO-albumin). The limited intracellular access of S-NO-albumin suggests the need for a vascular transfer mechanism of NO from a large plasma S-NO-albumin pool to effect biologic function. To study the role of low molecular weight (LMW) thiols in NO transfer in vivo, we administered intravenous S-NO-albumin (1–300 nmol/kg) to rabbits before and after an intravenous infusion of L-cysteine or N-acetyl-L-cysteine. S-NO-albumin produced dose-dependent hypotension that was significantly augmented by prior infusion of either LMW thiol. LMW thiol infusion significantly accelerated the rate of onset and reduced the duration of action of the hypotension induced by S-NO-albumin. The hemodynamic effects of S-NO-albumin after pretreatment with LMW thiols were mimicked by administration of the corresponding LMW S-nitrosothiol. The transfer of NO from albumin to L-cysteine was directly measured in rabbit plasma using a novel technique that couples high performance liquid chromatography to electrochemical detection. These data demonstrate that NO exchange between plasma protein thiol-bound NO and available LMW thiol pools (transnitrosation) occurs in vivo. (J. Clin. Invest. 1994. 94:1432–1439.)

Key words: nitric oxide • proteins • thiols • endothelium-derived relaxing factor • S-nitrosothiols

Introduction

Since the discovery of endothelium-derived relaxing factor (EDRF)1 by Puchgott and Zawadzki in 1980 (1), much information has become available about the biochemistry and physiology of this endothelial product. Many investigators have found that EDRF shares similar kinetic, biochemical, and physiologic properties with the nitric oxide radical (NO). Both agents cause equipotent relaxations of arterial and venous smooth muscle (2–4), inhibit platelet aggregation (5–8), and activate guanylyl cyclase (3–4). At least two groups have found the half-lives of these two agents to be identical (2, 9).

Despite these extensive similarities, several investigators have reported important differences between EDRF and NO. While some investigators have observed that EDRF and NO both relax nonvascular smooth muscle (10), in some studies EDRF fails to do so consistently (11). Myers et al. (12) found that EDRF was 80-fold more potent than NO in a bioassay system. In addition, Kelm and Schrader (13) reported the half-life of NO in vivo to be ~0.1 s, considerably shorter than the estimated 3–30-s half-life of EDRF ex vivo (9, 12).

Although several of these studies have been criticized for methodologic flaws and the experimental systems used varied, taken together these data have led to the speculation that EDRF bioactivity may not be entirely accounted for by NO alone, but may be evinced by a closely related NO adduct (12, 14–15). One likely candidate class of molecular species that has been shown to react readily with NO under physiologic conditions, without inactivating it, is that of biologic thiols, which form thionitrates or S-nitrosothiols (RSNOs) (14–16). We have recently shown that the sulfhydryl group(s) of plasma proteins can combine with oxides of nitrogen under physiologic conditions to form stable, biologically active NO adducts (17–20). The physiologic relevance of such protein-associated NO has been highlighted by recent evidence that NO circulates in mammalian plasma primarily as S-nitroso-albumin (18) and by the demonstration that S-nitroso-albumin possesses EDRF-like properties in vivo (20).

Although these direct plasma measurements and in vivo experimental data confirm the existence and biologic activity of S-nitroso-albumin, the role of serum albumin in NO metabolism remains imprecisely defined. It is possible that S-nitroso-albumin acts as an extracellular “buffer” of NO from which NO can be transferred to the intracellular milieu by smaller, less diffusion-limited S-nitrosothiol species. To investigate this possibility, we studied the effects of low molecular weight thiols on the systemic hemodynamic responses to S-nitroso-albumin in a rabbit model. Using a novel technique that couples reverse-phase high performance liquid chromatography (HPLC) with electrochemical detection (ECD), we also obtained direct evidence for the biochemical transnitrosation of NO from a plasma albumin pool to a low molecular weight thiol pool in vivo.

Methods

Materials

Sulfanilamide, N-(1-naphthyl)ethylenediamine dihydrochloride, and N-acetyl-L-cysteine were purchased from Aldrich Chemical Co. (Milwau-
Sodium nitrite was obtained from Fisher Scientific (Fairlawn, NJ) and atropine sulfate was obtained from LyphoMed Inc. (Rosemont, IL). Xylazine was purchased from Mobay Corp. (Shawnee, KS). Ketamine hydrochloride was obtained from Aveco Co., Inc. (Fort Dodge, IA) and sodium pentobarbital was purchased from Anthony Products Co. (Arcadia, CA). Hexamethonium bromide, timolol maleate, L-cysteine, bovine serum albumin, and all other reagent-grade chemicals were purchased from Sigma Chemical Co. (St. Louis, MO).

**Synthesis of S-nitroso-serum albumin and S-nitrosothiols**

_S-nitroso-albumin (S-NO-albumin)_ was prepared as previously described (17). Briefly, bovine serum albumin was exposed to an equimolar concentration of sodium nitrite in 1.0 N HCl for 30 min at 25°C and the solution was subsequently neutralized with equal volumes of Tris-buffered saline (TBS, 10 mM Tris, 0.15 M NaCl, pH 7.4) and 1.0 N NaOH. The presence of an S-NO bond was confirmed and quantified by the method of Saville (21), and synthesis of S-nitroso-BSA by this method yielded a stoichiometry of 0.85±0.07 mol S-NO/mol BSA. Previous studies have also confirmed the presence of an S-nitrosothiol using [15N]NMR spectroscopy and chemiluminescence (17).

_S-nitroso-L-cysteine (S-NO-cysteine) and S-nitroso-N-acetyl-L-cysteine (S-NO-N-acetyl-cysteine)_ were synthesized by combining equimolar concentrations of L-cysteine (cysteine) or N-acetyl-L-cysteine (N-acetyl-cysteine) with sodium nitrite in 0.5 N HCl at room temperature. Synthesis of S-NO-cysteine and S-NO-N-acetyl-cysteine by this method produces S-nitrosothiol that is 99% pure and stable at pH 3 (22). These low molecular weight S-nitrosothiols were synthesized on the day of use and maintained at pH 3 until immediately before administration, at which time they were diluted with 10 vol TBS.

Bovine serum albumin (BSA), N-acetyl-L-cysteine, and L-cysteine controls were each produced by acidification and neutralization in the absence of sodium nitrite. Similarly, sodium nitrite was acidified and neutralized in the absence of BSA as another control. L-cysteine and N-acetyl-L-cysteine were dissolved in phosphate-buffered saline (PBS, 10 mM sodium phosphate, 0.15 M NaCl, pH 7.4) and titrated to a pH of 4. All doses used in the experiments described were based on the component of mercuric ion-displaceable NO.

**Measurement of plasma S-nitrosothiols**

Plasma S-NO-cysteine was measured by HPLC using a C18 reverse-phase column coupled to an electrochemical detector with a dual Au/Hg electrode set at both oxidizing (+0.15 V) and reducing (-0.15 V) potentials versus a Ag/AgCl reference electrode (Bioanalytical Systems, West Lafayette, IN). The working electrodes were configured in series with the reducing electrode upstream. Using this configuration, S-nitrosothiols are detected at both electrodes—as S-nitrosothiols at the reducing electrode and as reduced thiol (the product of upstream reduction) at the oxidizing electrode. The mobile phase consisted of 0.1 M monochloroacetic acid, 0.125 M disodium ethylenediaminetetraacetic acid (EDTA), 1.25 mM sodium octyl sulfate, and 1% (vol/vol) acetonitrile, pH 2.8. The flow rate was 1 cc/min. This technique can reliably separate and detect S-nitrosothiols in the nanomolar range. Blood samples were diluted 1:10 with citrate-phosphate-dextrose (CPD) anticoagulant (15.6 mM citric acid, 90 mM sodium citrate, 16 mM NaH₂PO₄, H₂O, and 142 mM dextrose, pH 7.35) and 75% diethylenetriaminepentaacetic acid (DTPA), pH 7.4, then immediately centrifuged at 1,000 g for 10 min at 4°C. Plasma was microfiltered using a Centrifree microfiltration system (Amicon, Beverly, MA) with a molecular weight cutoff of 30,000 at 1,500 g. The resulting filtrate was injected into the HPLC/ECD apparatus and S-nitrosothiols were identified and quantitated by comparison with authentic standards. Biochemical identity of S-nitrosothiol peaks was also confirmed by augmentation of peak height by the addition of authentic (internal) standard to the plasma samples.

**Measurement of plasma thiols**

Plasma thiols were measured by HPLC/ECD. The reduction of plasma thiols was performed by a modification of a previously described technique (23). Briefly, blood samples were diluted 1:10 with citrate-phosphate-dextrose (CPD) anticoagulant and 75 mg% DTPA, pH 7.4, and immediately centrifuged at 1,000 g for 10 min at 4°C. 150 μl of plasma were then mixed with 50 μl 0.1 M potassium borate, pH 9.5, containing 2 mM ethylenediaminetetraacetic acid (EDTA). 20 μl of 100 ml/liter of tri-n-butyl-phosphine was added and the sample was incubated at 4°C for 30 min. The sample was then microfiltered using a Centrifree microfiltration system at 1,500 g. The resulting filtrate was injected into the HPLC/ECD apparatus with both electrodes set at +0.15 V configured in parallel, and thiols were identified and quantitated by comparison with authentic standards.

**Animal preparation**

New Zealand White rabbits (3.5–5.0 kg) were sedated with 0.1 mg/kg atropine sulfate and 5 mg/kg xylazine intramuscularly (IM) followed by 40 mg/kg of ketamine hydrochloride. Anesthesia was maintained by intravenous infusion of sodium pentobarbital at 5–7 mg/kg per h with a Harvard Pump (model 22, Harvard Apparatus Co. Inc., South Natick, MA). Tracheostomies were performed through a midline neck incision, and the trachea was intubated with a specifically designed endotracheal tube. Body temperature was maintained with a homeothermic blanket system (Harvard Apparatus Ltd., Edinbridge, KY, UK).

The femoral vein was cannulated through an inguinal incision with polyethylene tubing (PE-90; Clay Adams, Parsippany, NJ) for intravenous access, and the femoral artery was cannulated with a 5 French high fidelity pressure transducer (model SPC-350; Millar Instruments, Houston, TX). A Doppler flow probe (model T101; Transonic Systems Inc., Ithaca, NY) was placed on the contralateral femoral artery to measure hindlimb blood flow. All measurements were recorded on a Gould physiograph (model 2600; Gould Electronics Inc., Cleveland, OH).

Reflex responses were minimized in rabbits in groups 1–3 with hexamethonium bromide (20 mg/kg intravenously over 20 min), timolol maleate (1 mg/kg intravenous bolus), and atropine sulfate (0.2 mg/kg intravenous bolus). Autonomic blockade was maintained with continuous infusions of hexamethonium bromide (10 mg/kg per h) and timolol maleate (0.06 mg/kg per h).

**Experimental protocols**

**Group 1.** After the establishment of a baseline hemodynamic state, rabbits received intravenous bolus doses of S-NO-albumin (1–300 nmol/kg) in random order of concentration. A new baseline steady-state was established prior to the administration of each dose. After completion of this dosage regimen, rabbits received an intravenous infusion of 613 μmol/kg of either L-cysteine (n = 5) or N-acetyl-L-cysteine (n = 6) over 20 min followed by repeated administration of S-NO-albumin at identical doses.

**Group 2.** Rabbits in this group were prepared as described above and received intravenous bolus doses of S-NO-cysteine (n = 5) or S-NO-N-acetyl-cysteine (n = 6) (1–300 nmol/kg) in random order. These rabbits also served as their own controls, receiving similar doses of L-cysteine or N-acetyl-L-cysteine after the initial dosage regimen was completed.

**Group 3.** Rabbits in this group (n = 5) served as controls for group 1, and received intravenous bolus doses (1–300 nmol/kg) of BSA and sodium nitrite controls in an escalating dosage regimen.

**Group 4.** Rabbits in this group (n = 10) were used for measurement of S-nitrosothiols. A single 300 nmol/kg dose of S-NO-albumin was administered intravenously to each rabbit before and after a 10-min infusion of 400 μmol/kg L-cysteine. The S-NO-albumin used for this group of rabbits was synthesized by combining a 1.5-fold molar excess of bovine serum albumin with sodium nitrite to minimize free sodium nitrite in the synthesized product (≤ 50 μM). Blood samples were drawn for measurement of plasma S-NO-cysteine at baseline, as well as 1 min after administration of S-NO-albumin following pre-infusion of L-cysteine.

**Group 5.** Rabbits in this group (n = 4) served as controls for rabbits in group 4 and received 60 nmol/kg of acidified and neutralized sodium nitrite as an intravenous bolus both before and after a 10-min infusion.
of 400 μmol/kg L-cysteine. This dose was chosen because it reflected at least a 20% increase over the maximal dose of free sodium nitrite in the bolus of S-NO-albumin administered to rabbits in group 4.

Group 6. Rabbits in this group (n = 10) were used for measurement of plasma L-cysteine levels. A single 300 nmol/kg dose of S-NO-albumin was administered intravenously to each rabbit before and after a 10-min infusion of 613 μmol/kg L-cysteine. The S-NO-albumin used for this group of rabbits was prepared as described for rabbits in group 4. Blood samples were drawn for measurement of plasma L-cysteine at baseline, as well as 1 min after administration of S-NO-albumin after pre-treatment with L-cysteine.

**Definition of hemodynamic and time parameters**

EC50 was defined as the dose of S-NO-albumin that caused a half-maximal decrease in mean arterial blood pressure. This parameter was determined for the dose-range tested and does not reflect an actual pharmacologic EC50 per se. Recovery halftime was calculated by measuring the time to recovery of the systolic arterial blood pressure to half its maximal decrease (Fig. 1). This parameter was chosen as a reflection of recovery time because of the asymptotic nature of the restoration in blood pressure after a dose of test agent. EC100, was defined as the dose of S-NO-albumin that effected a recovery time of 100 s. Rate of onset was obtained by calculating the maximal slope of the decline of the systolic arterial blood pressure tracing (Fig. 1). EC100 was derived from the dose of S-NO-albumin that effected a slope of -2 mm Hg/s. Hindlimb resistance was calculated by dividing the nadir of mean arterial pressure by the coincidental peak increase in mean hindlimb blood flow after administration of a dose of test agent and is expressed in Wood units (mm Hg min/liter).

**Statistical analysis**

All data are presented as the mean±SEM. All dose-response curves were analyzed by repeated measures analysis of variance (SAS Institute, Inc., Cary, NC). Dose-response curves were compared among agents by the same method. Comparison of S-NO-cysteine and L-cysteine concentrations at baseline and after administration of S-nitroso-albumin following pre-treatment with L-cysteine was performed using a paired, one-tailed Student’s t test. Statistical significance was accepted if the null hypothesis was rejected with P < 0.05.

**Results**

**Hemodynamics: blood pressure response.** A representative tracing depicting the effect of an intravenous bolus dose of S-NO-albumin on mean arterial pressure is shown in Fig. 1, and dose-response curves for this variable are shown in Fig. 2. S-NO-albumin produced a significant dose-dependent decrease in mean arterial blood pressure (P < 0.0001). Infusion of L-cysteine (Fig. 2 A) enhanced the vascular sensitivity to S-NO-albumin 18-fold (EC50 before cysteine, 62.5 nmol/kg; after cysteine, 3.5 nmol/kg; P = 0.01), while infusion of N-acetyl-L-cysteine was associated with a three-fold enhancement of the vascular sensitivity to S-NO-albumin (EC50 before N-acetyl-cysteine, 62.5 nmol/kg; after N-acetyl-L-cysteine, 20 nmol/kg; P < 0.02).

S-NO-cysteine and S-NO-N-acetyl-cysteine both produced significant dose-dependent reductions in mean arterial blood pressure (P < 0.0001). As shown in Fig. 2 A, S-NO-cysteine was significantly more potent, on a molar basis, than S-NO-albumin (EC50 7 nmol/kg vs. 62.5 nmol/kg; P < 0.005). By contrast, after the infusion of L-cysteine (Fig. 2 A), S-NO-albumin was nearly equipotent to S-NO-cysteine (EC50 3.5 nmol/kg vs. 7 nmol/kg, P < 0.02). As shown in Fig. 2 B, S-NO-N-acetyl-cysteine was fourfold more potent in reducing mean arterial pressure than S-NO-albumin (EC50 15 nmol/kg vs. 62.5 nmol/kg, P < 0.02). After infusion of N-acetyl-L-cysteine, however, S-NO-albumin was equipotent to S-NO-N-acetyl-cysteine (EC50 20 nmol/kg vs. 15 nmol/kg, P = NS). L-cysteine, N-acetyl-L-cysteine, BSA, and sodium nitrite controls had no significant effect on systemic arterial blood pressure over this concentration range (P > 0.001, data not shown).

**Kinetics of the hemodynamic response: recovery half-time.** To quantify the biologic half-life of the NO adducts studied, recovery half-times (as defined in Fig. 1) were determined for each agent. Dose-response curves for this parameter were con-
structed and are shown in Fig. 3. S-NO-cysteine, S-NO-N-acetyl-cysteine, and S-NO-albumin before and after infusion of either biologic low molecular weight thiol all produced dose-dependent increases in the time for recovery of the hypotensive response (P < 0.05 for all dose-response curves). Infusion of L-cysteine produced a 10-fold increase in the EC_{100} for S-NO-albumin (18 nmol/kg vs. 180 nmol/kg, P < 0.01), reflecting a significant reduction in the duration of action of S-NO-albumin (Fig. 3 A). At a dose of 100 nmol/kg (Fig. 1), L-cysteine pretreatment reduced the recovery half-time of S-NO-albumin by 61%. S-NO-cysteine administration produced recovery half-times similar to those for S-NO-albumin after infusion of L-cysteine (P = NS).

In contrast, N-acetyl-L-cysteine pretreatment did not produce a significant reduction in the recovery half-time or increase in EC_{100} for S-NO-albumin (Fig. 3 B, P = NS). In concordance with this finding, the duration of action of S-NO-N-acetyl-cysteine is also similar to that for S-NO-albumin (P = NS). In addition, comparison of Fig. 3, A and B reveals that S-NO-cysteine has a significantly shorter duration of action than S-NO-N-acetyl-cysteine (EC_{100}, 275 nmol/kg vs. 20 nmol/kg, P < 0.03).

**Kinetics of the hemodynamic response: rate of onset.** The rate of onset of the hypotensive response induced by the NO adducts studied reflects the rate of delivery of NO to vascular smooth muscle. This parameter increased in a dose-dependent fashion for both low molecular weight S-nitrosothiols as well as for S-NO-albumin before and after infusion of either low-molecular-weight thiol (Fig. 4; P < 0.02 for all dose-response curves). L-cysteine (Fig. 4 A) accelerated the rate of onset of the hypotensive response of S-NO-albumin twenty-seven-fold (EC_{-2 nmol/kg}, for S-NO-albumin before L-cysteine, 80 nmol/kg; after L-cysteine, 3 nmol/kg; P < 0.003), while N-acetyl-L-cysteine (Fig. 4 B) infusion was associated with a four-fold decrease in the EC_{-2 nmol/kg} (80 nmol/kg vs. 25 nmol/kg, P < 0.02). At a dose of 100 nmol/kg, L-cysteine (Fig. 1) and N-acetyl-L-cysteine infusions significantly accelerated the rate of onset of hypotension by 58 and 49%, respectively.

With regard to the low molecular weight S-nitrosothiols, S-NO-cysteine (Fig. 4 A) and S-NO-N-acetyl-cysteine (Fig. 4 B) were both significantly more potent, on a molar basis, in accelerating the rate of onset of the hypotensive response compared to S-NO-albumin before infusion of thiol. S-NO-cysteine (Fig. 4 A) was eight-fold more potent (EC_{-2 nmol/kg}, 10 nmol/kg vs. 80 nmol/kg, P < 0.04), and S-NO-N-acetyl-cysteine was four-fold more potent than S-NO-albumin (EC_{-2 nmol/kg}, 20 nmol/kg vs. 80 nmol/kg, P < 0.03). S-NO-cysteine and S-NO-N-acetyl-cysteine produced rates of onset that were similar to those produced by S-NO-albumin after pretreatment with the corresponding low molecular weight thiol (P = NS).

**Hindlimb vascular resistance.** Administration of systemic intravenous S-NO-albumin before and after administration of low-molecular-weight thiols produced a significant dose-depen-
Figure 4. The response and S-NO-cysteine, N-acetyl-L-cysteine, respectively of (P dose-dependent potentiate the hypotensive curves). Infusion of the blood flow was markedly increased by prior infusion of (A) L-cysteine (○, P < 0.002) and (B) N-acetyl-L-cysteine (△, P < 0.02). The dose-response curves for S-NO-cysteine (□) and S-NO-N-acetyl-cysteine (△) were not significantly different from the dose-response curves induced by S-NO-albumin after infusion of L-cysteine and N-acetyl-L-cysteine, respectively (P = NS).

Figure 5. The response of hindlimb vascular resistance to S-NO-albumin, S-NO-cysteine, and S-NO-N-acetyl-cysteine. S-NO-albumin (●) produced a dose-dependent decrease in hindlimb vascular resistance that was significantly potentiated by L-cysteine (○, P < 0.001) and enhanced by N-acetyl-L-cysteine (△, P = 0.07). The dose-response curves for S-NO-cysteine (□) and S-NO-N-acetyl-cysteine (△) were not significantly different from those for S-NO-albumin before or after L-cysteine and N-acetyl-L-cysteine, respectively (P = NS).

Figure 6. Measurement of S-nitroso-L-cysteine (S-NO-CYS) concentration in vivo by HPLC coupled to ECD. Administration of 300 nmol/kg of S-nitroso-albumin after pretreatment with 400 μmol/kg cysteine (post) produced a significant 12-fold increase in S-NO-cysteine (S-NO-CYS) concentration compared with baseline (post) (P < 0.05).

The response and S-NO-cysteine, N-acetyl-L-cysteine, respectively of (P dose-dependent potentiate the hypotensive curves). Infusion of the blood flow was markedly increased by prior infusion of (A) L-cysteine (○, P < 0.002) and (B) N-acetyl-L-cysteine (△, P < 0.02). The dose-response curves for S-NO-cysteine (□) and S-NO-N-acetyl-cysteine (△) were not significantly different from the dose-response curves induced by S-NO-albumin after infusion of L-cysteine and N-acetyl-L-cysteine, respectively (P = NS).

of free sodium nitrite before and after L-cysteine infusion had no effect on baseline S-NO-cysteine concentration (data not shown). Mean plasma L-cysteine concentration increased 4.7-fold (n = 10) after infusion of L-cysteine followed by S-NO-albumin (P = 0.0005).

Plasma S-NO-cysteine measurements. The results of S-NO-cysteine measurements in plasma for rabbits in group 4 are shown in Fig. 6, and a representative chromatogram is shown in Fig. 7. Importantly, S-NO-cysteine was detected in 7 of 10 rabbits in the baseline state. The range was 30.1 nM to 873 nM with a mean S-NO-cysteine concentration of 221.1 nM±259.7 nM. Administration of 300 nmol/kg S-NO-albumin following infusion of L-cysteine caused an increased in S-NO-cysteine concentration to 2.75±2.42 μM reflecting an twelve-fold increase over baseline concentration (P = 0.017). Administration of free sodium nitrite before and after L-cysteine infusion had no effect on baseline S-NO-cysteine concentration (data not shown). Mean plasma L-cysteine concentration increased 4.7-fold (n = 10) after infusion of L-cysteine followed by S-NO-albumin (P = 0.0005).
Discussion

Although the origin of NO has been definitively traced to the terminal guanidino group of L-arginine (24), the specific biochemical intermediates that exist between this precursor and the presumed site of NO action, the heme moiety of guanylyl cyclase, are unknown. Given the extreme lability and facile inactivation of NO by heme iron, non-heme iron (25), superoxide anion (26), oxygen (27), and other biochemical species (28–29), it is likely that some intermediate molecular species serves to stabilize NO and/or transfer it from its site of origin (endothelial cell) to target effector sites (e.g., vascular smooth muscle, platelets). Plasma thiols represent likely candidates for this role as they readily form thionitrites (S-nitrosothiols) under physiologic conditions by complexation with NO or by reacting with an endogenous nitrosating species (15–16, 30), are present in abundant quantities in vivo, and confer stability to the NO radical (14). Furthermore, there is ample evidence that low molecular weight S-nitrosothiols (as well as plasma protein thiols) participate in the mechanism of action of nitrateroxidosides (14–15, 31–34), and, by inference, in the mechanism of action of NO.

We have recently shown that the S-nitrosation of albumin at its single free thiol, cysteine 34, forms the bioactive NO adduct S-NO-albumin. S-NO-albumin relaxes vascular smooth muscle ex vivo (17) and in vivo (20, 34), and activates guanylyl cyclase (17). The role of S-NO-albumin, present in the medium range in mammalian plasma (17), in the intermediary metabolism of NO is, however, unclear. With prolonged in vivo hemodynamic effects (20) and limited (if any) intracellular access, S-NO-albumin would be unlikely to serve as an adduct that "presents" NO directly to the intracellular milieu. However, S-NO-albumin may serve as an extracellular buffer or reservoir of relatively stable NO that can be transferred more efficiently to vascular smooth muscle or to the platelet cytosol by low molecular weight, less diffusion-limited thiol species.

NO exchange among thiols has been documented by several investigators, including Oae and coworkers (16) and others (35–37), under non-physiologic conditions. We have recently reported that transnitrosation between S-NO-albumin and low molecular weight thiols occurs rapidly and completely in vitro under physiologic conditions (18). In addition, we have recently found that NO appears to shift transiently from high molecular-weight to low molecular-weight S-nitrosothiol pools in vivo in response to the administration of N\(^\text{O}\)-monomethyl-L-arginine (18).

The hemodynamic data presented here provide direct evidence that transnitrosation occurs in vivo. The hemodynamic responses to S-NO-albumin after infusion of L-cysteine or N-acetyl-L-cysteine were mimicked by administration of the corresponding synthetic S-nitrosothiol. This similarity was particularly striking for N-acetyl-L-cysteine (Fig. 2 B), with nearly identical dose-response curves characterizing the hemodynamic responses to both S-NO-N-acetyl-cysteine as well as S-NO-albumin after infusion of N-acetyl-L-cysteine.

The kinetic data are also consistent with in vivo transnitrosation. Both L-cysteine and N-acetyl-L-cysteine accelerate the rate of onset of the hypotensive response to S-NO-albumin. For both low molecular weight thiols, the dose-response curves for this variable induced by S-NO-albumin after infusion of thiol were not significantly different from the dose-response curves for the corresponding synthetic S-nitrosothiol. These results reflect the more efficient and rapid delivery of NO to the vascular smooth muscle cell by the low molecular weight S-nitrosothiols formed by transnitrosation of NO from S-NO-albumin to the exogenously administered low molecular weight thiol.

The data for recovery half-time also support the hypothesis that transnitrosation occurs in vivo, and highlight the differences in the stability of various low molecular weight S-nitrosothiols. L-cysteine shortens the duration of action of S-NO-albumin, and the dose-response curve for S-NO-albumin after infusion of L-cysteine was nearly identical to that for synthetic S-NO-cysteine. Although N-acetyl-L-cysteine infusion did not shorten the duration of action of S-NO-albumin, this finding was similarly consistent with the occurrence in vivo of transnitrosation as the duration of action of synthetic S-NO-N-acetyl-cysteine was also similar to that for S-NO-albumin.

The hindlimb vascular resistance data also provide important insights into the differences among S-nitrosothiols. While infusion of N-acetyl-L-cysteine produced a significant increase in baseline mean hindlimb blood flow (with no change in hindlimb resistance), infusion of L-cysteine did not (data not shown). This increase in baseline flow induced by N-acetyl-L-cysteine likely results from the formation of an appreciable pool of S-NO-N-acetyl-cysteine by transnitrosation of NO from endogenous and exogenously administered S-NO-albumin to N-acetyl-L-cysteine. Although such an S-nitrosothiol pool is likely formed after infusion of cysteine as well, the duration of action of S-NO-N-acetyl-cysteine is sufficiently prolonged in the hindlimb microcirculation that baseline hindlimb blood flow is significantly increased only by N-acetyl-L-cysteine. This rationale may also explain the lack of a statistically significant decrease in hindlimb resistance induced by S-NO-albumin after administration of N-acetyl-L-cysteine. Repeat administration of S-NO-albumin in the presence of an appreciable, relatively long-acting S-NO-N-acetyl-cysteine plasma pool does little to decrease the already near maximally dilated hindlimb microcirculation.

Taken together, the similarities between the hemodynamic properties of S-NO-albumin after pretreatment with thiol to the hemodynamic profiles of corresponding S-nitrosothiols are convincing evidence for transnitrosation. The HPLC/ECD data provide direct biochemical confirmation of this process. Using this technique, we have detected S-NO-cysteine in plasma in 7 out of 10 rabbits at baseline and markedly increased levels in all
rabbits after administration of S-NO-albumin following preinfusion of l-cysteine. The range of S-NO-cysteine in the basal state varied from low nM to several hundred nM; this range was even more variable (several hundred nM to low μM) after infusion of S-NO-albumin and cysteine. This dramatic variability is not unexpected given the highly reactive nature of S-nitrosothiols in biologic systems and the fact that wide variability in nitrite and nitrate levels, other labile molecular species sensitive to redox conditions, have also been reported (38).

Based on these in vivo data and our in vitro observations, we propose a hypothetical mechanism for the transnitrosation of NO among available thiol species. The cysteine at position 34 of albumin may react with NO\(^+\), nitrosonium, to form S-NO-albumin, the dominant form of NO in plasma. This reaction may be favored because of the anomalously low pKa\(_e\) (< 5) of albumin's cysteine 34, facilitating the formation of thiolate anion (RS\(^-\)) at physiologic pH and redox modulation of NO (39). We postulate that S-NO-albumin may serve as a buffer and reservoir of NO in plasma from which NO can be transferred more efficiently to the intracellular milieu by transnitrosation to low molecular weight thiols forming the corresponding low molecular weight S-nitrosothiol. These latter species may be readily transported across cell membranes by thiol transport mechanisms.

In these experiments, we have facilitated this proposed transnitrosation process by adding a significant excess of low molecular weight thiol to a supplemented S-NO-albumin plasma pool. In the case of l-cysteine, an endogenous biologic thiol present in the micromolar range in mammalian plasma (40), administration of exogenous thiol augments l-cysteine concentrations nearly fivefold and likely alters the equilibrium balance of endogenous NO, l-cysteine, and S-NO-albumin. Administration of l-cysteine without S-NO-albumin supplementation had no hemodynamic effect in these animals; this is likely a consequence of the complex equilibria that exist among the many competing reactions with which l-cysteine is involved in vivo, such as disulfide formation with cyst 34 of albumin. After S-NO-albumin supplementation, however, the demonstrated increase in plasma S-NO-cysteine concentration by exogenous l-cysteine administration results in enhanced delivery of NO to the vasculature. This is manifested by acceleration of the rate of onset, potentiation of the hemodynamic effect, and reduction of the duration of action of exogenously administered S-NO-albumin.

The transnitrosation of NO from albumin to biologic thiols depends on the relative reactivity of the competing high and low molecular weight thiols as well as the relative stability of the S-nitrosothiols formed. Factors that likely contribute to the relative reactivities of the various thiols include the pK\(_a\)s of the individual thiol moieties which influence nucleophilicity, the molecular structure of the individual thiols which influence thiol or thiolate reactivity, and the ambient physiologic conditions of pH and ionic strength (40). The marked difference between the pK\(_a\) of albumin's cysteine 34 (pK\(_a\) < 5) (41) compared with the pK\(_a\) of the low molecular weight biologic thiols (pK\(_a\) = 7–8) (40) is one important factor that should favor the transnitrosation of NO from albumin to low molecular weight thiol under physiologic conditions by facilitating thiol-S-nitrosothiol exchange.

Once transnitrosation has occurred and S-nitrosothiols have been formed, these smaller, less diffusion-limited NO adducts can transport NO more efficiently to the vascular smooth muscle cell or platelet surface. Once at the cell surface, the specific metabolic steps that lead to the activation of intracellular guanylyl cyclase and other target effector sites remain unclear. Possible mechanisms include the direct transport of S-nitrosothiols into the cytosol or, alternatively, cell surface interactions with membrane bound species bearing thiols or heme-containing prosthetic groups (42). This contention draws support from the findings of Malinski and Taha (43), who used a porphyrin-based microsensor to measure single cell NO. These investigators found that a significant NO gradient exists across the vascular smooth muscle cell and platelet membrane, suggesting that transmembrane NO transport is not a diffusion-limited process.

The data presented here strongly support the occurrence of transnitrosation reactions in vivo and a role for these reactions in the transport and metabolism of NO. Alterations in transnitrosation may be an important factor in the pathophysiologic of vascular disease.

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