Detailed mechanistic investigation into the S-nitrosation of cysteamine

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Abstract

The nitrosation of cysteamine (H₂NCH₂CH₂SH) to produce cysteamine-S-nitrosothiol (CANO) was studied in slightly acidic medium by using nitrous acid prepared in situ. The stoichiometry of the reaction was H₂NCH₂CH₂SH + HNO₂ → H₂NCH₂CH₂SNO + H₂O. On prolonged standing, the nitrosothiol decomposed quantitatively to yield the disulfide, cystamine: 2H₂NCH₂CH₂SNO → H₂NCH₂CH₂S–SCH₂CH₂NH₂ + 2NO. NO₂ and N₂O₃ are not the primary nitrosating agents, since their precursor (NO) was not detected during the nitrosation process. The reaction is first
order in nitrous acid, thus implicating it as the major nitrosating agent in mildly acidic pH conditions. Acid catalyzes nitrosation after nitrous acid has saturated, implicating the protonated nitrous acid species, the nitrosonium cation (NO$^+$) as a contributing nitrosating species in highly acidic environments. The acid catalysis at constant nitrous acid concentrations suggests that the nitrosonium cation nitrosates at a much higher rate than nitrous acid. Bimolecular rate constants for the nitrosation of cysteamine by nitrous acid and by the nitrosonium cation were deduced to be $17.9 \pm 1.5 \text{ (mol/L)}^{-1} \text{s}^{-1}$ and $6.7 \times 10^4 \text{ (mol/L)}^{-1} \text{s}^{-1}$, respectively. Both Cu(I) and Cu(II) ions were effective catalysts for the formation and decomposition of the cysteamine nitrosothiol. Cu(II) ions could catalyze the nitrosation of cysteamine in neutral conditions, whereas Cu(I) could only catalyze in acidic conditions. Transnitrosation kinetics of CANO with glutathione showed the formation of cystamine and the mixed disulfide with no formation of oxidized glutathione (GSSG). The nitrosation reaction was satisfactorily simulated by a simple reaction scheme involving eight reactions.

**Keywords**
nitric oxide; nitrosation; thiols; cysteamine; kinetics

**Introduction**

Nitric oxide (NO) is a simple gaseous molecule with fascinating biological functions.$^1$ It is one of the most versatile bioactive molecules ever identified. Unlike other free radicals, it is relatively stable, but becomes unstable in biological systems due to its reaction with oxygen, superoxide anion (O$_2^{−}$), and haem proteins.$^2$ Nitrite (NO$_2^{−}$) is the major end product of its reaction with oxygen. Nitrite can also release NO on acidification and by the reductase activity of the enzyme, xanthine oxidoreductase, at pH conditions lower than 6.0.$^3$–$^5$

Nitric oxide has been the focus of many research groups in recent years, probably because of the realization that it is synthesized by mammalian cells and can act both as a physiological messenger and as a cytotoxic agent.$^6$ It is an essential physiological signaling molecule necessary for cell-to-cell communication and it is also involved in the regulation of gene expression and modification of sexual and aggressive behavior.$^7$,$^8$ It is actively involved in smooth muscle relaxation due to its ability to activate guanosine 3′,5′-cyclic monophosphate (cGMP) in smooth muscle tissue.$^9$,$^{10}$ The role of nitric oxide in the physiological system at any given time significantly depends on its concentration and its environment. It is known to perform most of its beneficial roles, such as vasorelaxation$^{11}$,$^{12}$ and neurotransmission,$^{13}$,$^{14}$ at low concentrations, on the order of 5–10 nmol/L.$^{15}$ At this concentration range it cannot effectively compete with superoxide dismutase (SOD) for O$_2^{−}$.

NO also readily reacts with O$_2^{−}$ to produce the highly damaging and genotoxic oxidant (peroxynitrite, ONOO$^−$):

$$\text{NO} + \text{O}_2^{−} \rightarrow \text{ONOO}^{−} \quad [R1]$$
The bimolecular rate constant for rxn. [R1] is \(6.7 \times 10^9\) (mol/L\(^{-1}\) s\(^{-1}\)). Superoxide anion radical concentrations in the physiological environment are in the order of 4–10 μmol/L. When NO concentrations are in the micromolar range, rxn. [R1], which produces the deleterious peroxynitrite, becomes competitive with the deactivating reaction that removes superoxide anion radical.

Excessive production of nitric oxide has thus been linked to so many cytotoxic activities such as septic shock and liver injury. It is necessary to have processes that can reduce NO concentrations to beneficial levels in the physiological environment. Such processes would include its reaction with thiols to form S-nitrosothiols (RSNOs). For example, RSNOs have been shown to improve end-organ recovery in models of ischemia–reperfusion injury in the heart and liver. Biological nitrosothiol formation is a post-translational modification of biological thiols that is subsequently linked to all those functions that are attributed to nitric oxide. Possible roles of S-nitrosation include the regulation of apoptosis through glyceraldehyde-3-phosphate dehydrogenase modification as well as involvement in the pathogenesis of Parkinson’s disease. There are several studies on nitrosothiol formation by Williams, but these studies dwell on formations and characterizations and not so much on the mechanisms and kinetics. Despite the potential benefits of the relevance of the mechanistic basis of nitrosations (both S- and N-nitrosations), very few studies have been directed at elucidating these mechanisms.

Cysteamine (2-aminoethanethiol) is a very important physiologically active β-aminothiol. It is produced in vivo by degradation of coenzyme A. Coenzyme A is a cofactor in many enzymatic pathways, which include fatty acid oxidation, heme synthesis, synthesis of the neurotransmitter acetylcholine, and amino acid catabolism. Cysteamine is medically known as Cystagon and can be used as oral therapy for the prevention of hypothyroidism and as a growth enhancer in patients with nephropathic cystinosis. Cysteamine and its disulfide (cystamine) can also be used in topical eye drops to dissolve corneal cystine crystals. It has also been shown to be an effective molecule against hepatotoxicity caused by excessive production of nitric oxide.

Although a series of studies have been performed on the metabolism of this thiol and its metabolites with respect to oxidation by reactive oxygen species, nothing has been done on the elucidation of its mechanism of nitrosation by NO and (or) NO-derived metabolites (N\(_2\)O\(_3\) and NO\(^+\)). Its nitrosation is well-known and has been reported in several manuscripts, but the exact mechanistic details are still unknown. We report herein on the detailed kinetics and mechanism of the nitrosation of cysteamine. The nitrosating agent is produced in situ, by premixing H\(^+\) and NO\(_2\) and utilizing a double-mixing stopped-flow spectrophotometer. This manuscript also reports on detailed product analysis as well as on transnitrosation kinetics. The most prevalent thiol-based antioxidant in the physiological environment is glutathione. It is envisaged that, in any discussion of S-nitrosation, glutathione would be a major player.
Experimental

Materials

Cysteamine hydrochloride (CA, 2-aminoethanethiol hydrochloride) 98%, cystamine, cysteine (99+%), sodium perchlorate (Acros), glutathione, sodium nitrite, sodium chloride, perchloric acid (72%), ferrous sulfate (Fisher Scientific), cuprous chloride, and cupric chloride (GSF chemicals) were used as purchased. Stock solutions of CA and nitrite were prepared just before use. Reagent solutions were prepared with distilled and deionized water from a Barnstead Sybron Corporation water purification unit. Inductively coupled plasma mass spectrometry (ICP–MS) was used to show that our aqueous reaction medium did not contain enough transition-metal ions to affect the overall reaction kinetics and mechanism. The highest metal ion concentration was cadmium at 1.5 ppb followed by lead at 0.43 ppb. Copper was virtually undetectable.

Methods

All experiments were carried out at 25.0 ± 0.1 °C and at a constant ionic strength of 1.0 mol/L (NaClO₄). The progress of the nitrosation reaction was monitored spectrophotometrically by following the absorbance of S-nitrosocysteamine (CANO) at its experimentally determined absorption peak of 545 nm, where an absorptivity coefficient of 16.0 ± 0.1 (mol/L)⁻¹ cm⁻¹ was evaluated. Although CANO has a second absorption peak at 333 nm, with a much higher absorptivity coefficient of 536.0 (mol/L)⁻¹ cm⁻¹, its maximum absorption at 545 nm was used owing to the lack of interference from the absorption peaks of nitrogen species (NO₂⁻, NO⁺, and HNO₂) at this absorption wavelength (see Fig. 1).

Product identification and verification was achieved by complementary techniques: UV–vis spectrophotometry, HPLC, quadrupole time-of-flight mass spectrometry (TOF-MS), and electron paramagnetic resonance (EPR) spectrometry. Kinetics measurements for fast reactions were performed on a Hi-Tech Scientific double-mixing SF61-DX2 stopped-flow spectrophotometer. For slower reactions, a PerkinElmer Lambda 25 UV–vis spectrophotometer was used.

Stoichiometric determinations were performed by mixing excess perchloric acid and nitrite with a fixed amount of CA such that CA was the limiting reagent. The reaction was allowed to proceed to completion while rapidly scanning the reacting solution between 200 and 700 nm. The concentration of CANO was deduced from its maximum absorption at 545 nm.

HPLC technique—The HPLC system utilized a Shimadzu Prominance (Columbia, Maryland) SPD-M10A VP diode array detector, dual LC 600 pumps, LPT-6B LC–PC interface controller, SIL-10AD auto injector, and Class-VP chromatography data system software. All samples were loaded on a reversed phase Discovery 5 μm C18 column (Supelco, Bellefonte, Pennsylvania). They were run isocractically at 0.15% trifluoroacetic acid (TFA) / H₂O to 10% methanol / H₂O, and CANO was detected at 545 nm (for selective scans) with maximum spectral scan mode employed to detect all eluents. A flow rate of 1 mL/min was maintained. All solutions for HPLC analysis were made with Milli-Q Millipore purified water and filtered with Whatman Polypropylene 0.45 μm pore-size filter devices.
before injection (10 μL) into the column using the auto injector. To curb the interaction of the protonated amines on the analytes with the silanol groups on the stationary phase (which was causing tailing of peaks), the sodium salt of 1-octanesulfonic acid (0.005 mol/L) was incorporated into the aqueous mobile phase as an HPLC ion-pairing agent. This was sufficient to neutralize the protonated amines and produce good resolution while eliminating peak tailing.

**TOF-MS**—Mass spectra were acquired on a Micromass QTOF-II (Waters Corporation, Millford, MA) quadrupole time-of-flight mass spectrometer. Analytes were dissolved in 50:50 acetonitrile / water – 1% formic acid mixture, and analyte ions were generated by positive-mode electrospray ionization (ESI) at a capillary voltage of 2.8 kV and a flow rate of 5 μL/min. The source block was maintained at 80 °C and the nitrogen desolvation gas was maintained at 150 °C and at a flow rate of 400 L/h. Data were visualized and analyzed with the Micromass MassLynx 4.0 software suite for Windows XP (Water Corporation, Millford, Massachusetts).

**EPR Measurements**—A Bruker e-scan EPR spectrometer (X-band) was used for the detection of radicals and to confirm the release of NO upon the decomposition of CANO. Detection of NO was through the use of the NO-specific spin trap, aci-nitroethane. The following settings were used for a typical run: microwave power, 19.91 mW; modulation amplitude, 1.41 G; receiver gain, 448; time constant, 10.24 ms; sweep width, 100 G; and frequency, 9.78 GHz. All measurements were taken at room temperature.

**Results**

The major reaction studied was a simple nitrosation of cysteamine through in situ production of nitrous acid as the nitrosating agent. The yield of CANO varied, based on the order of mixing reagents. The highest yield, which gave quantitative formation of CANO, was obtained by premixing nitrite with acid and incubating the solution for approximately 1 s before subsequently reacting this nitrous acid with cysteamine in a third reagent stream of the double-mixing stopped-flow spectrophotometer. The value of the absorptivity coefficient of CANO at 545 nm adopted for this study of 16.0 mol/L$^{-1}$ cm$^{-1}$ was derived from this mixing order. Premixing CA with acid before the addition of nitrite gave a lower yield of CANO. The available protons are reduced by the protonation of CA.

\[ H_2NCH_2CH_2SH + H^+ \leftrightarrow [H_2NCH_2CH_2SH]^+ \]  \[ R2 \]

This subsequently reduces the amount of the immediately available nitrosating agent, HNO$_2$.

**Stoichiometry and product analysis**

The stoichiometry, in conditions of excess acid and nitrite over CA, was strictly 1:1.

\[ H_2NCH_2CH_2SH + HNO_2 \rightarrow H_2NCH_2CH_2S + NO + H_2O \]  \[ R3 \]
Stoichiometry rxn. [R3] is attained within a minute of mixing the reagents at high acid concentrations, and a little longer in lower acid concentrations. High acid concentrations rapidly form the product (CANO), which also rapidly decomposes. A resolvable mass spectrum is taken at low acid concentrations, and this generally will show both the reagent (CA) and the single product (CANO). This mass spectrum is shown in Fig. 2A. There are no other peaks that can be discerned from the reaction mixture, apart from those belonging to the reagent and the product. Incubation of the product of nitrosation overnight shows a final conversion of the CANO into the disulfide, cystamine (H$_2$NCH$_2$CH$_2$S–SCH$_2$CH$_2$NH$_2$, Fig. 2B).

\[ 2H_2NCH_2CH_2S + NO \rightarrow H_2NCH_2CH_2S – SCH_2CH_2NH_2 + 2NO \]  

Nitric oxide can be trapped and observed by EPR techniques through the use of a nitroethane trap.\textsuperscript{44} The aci-anion of this trap, formed in basic environments, generates a long-lived spin adduct with NO, which is detectable and quantifiable by its distinct EPR spectrum pattern and g value.

\[ \text{CH}_3\text{CH} = \text{NO}_2^- + \text{NO} + \text{OH}^- \rightarrow [\text{CH}_3\text{C'(NO)(NO}_2^-]^2^- + H_2O \]

This adduct ([CH$_3$C(NO)(NO$_2$)]$^{2-}$) is EPR-active. No peaks were obtained in the EPR spectrum when the nitroethane trap was utilized during the formation of CANO, indicating that NO is not released during the S-nitrosation and that it is not a major nitrosation agent in this mechanism. Upon use of the trap, however, during the decomposition of CANO, a strong EPR spectrum indicating the presence of NO was obtained, proving stoichiometry [R4]. Figure 3 shows a series of EPR spectra that proves the presence of NO in the decomposition of CANO. The first three spectra show that, on their own, nitroethane, CA, and NO$_2^-$ cannot generate any active peaks that can be attributed to NO. In excess concentrations of the trapping reagent, heights of the spectral peaks can be used, after a standardization, as a quantitative measure of the available NO. NO has a very poor solubility in water, and so the technique is limited only to low NO concentrations.

**Reaction kinetics**

The reaction has a very simple dependence on cysteamine concentrations (see Figs. 4A and 4B). There is a linear first-order dependence on the rate of nitrosation with CA concentrations with an intercept kinetically indistinguishable from zero. The reaction kinetics are more reproducible in the format utilized in Fig. 4 in which concentrations of acid and nitrite are equal and in overwhelming excess over CA concentrations. The effect of nitrite is also simple and first-order (see Figs. 5) when nitrite concentrations do not exceed the initial acid concentrations. Effectively, under these conditions, all N(III) species in the reaction environment are in the form of protonated HNO$_2$. Acid effects are much more complex and are dependent on the ratio of initial acid to nitrite concentrations. The nitrosation response to acid differs, depending on whether initial proton (H$_3$O$^+$) concentrations exceed initial nitrite concentrations. The most important parameter affecting the rate of nitrosation is the concentration of HNO$_2$. A set of kinetics traces that encompass
a proton concentration variation, which ranges from concentrations that are lower than nitrite to those that exceed, was performed. A simple plot of added acid to the rate of nitrosation shown in Fig. 6A clearly displays a discontinuity at the point where acid exceeds nitrite concentrations. By utilizing the dissociation constant of HNO₂, the final concentrations of H₃O⁺, NO₂⁻, and HNO₂ can be calculated for each set of initial concentrations. These data are shown in Table 1 for the input concentrations used for the set of kinetics experiments. Close examination of the data in Table 1 shows that the kinetics of the reaction are vastly different depending on whether initial acid concentrations exceed initial nitrite concentrations. When nitrite is in excess (the first four readings), there is a sharp nonlinear increase in the rate of nitrosation with nitrous acid concentrations (Fig. 6B), and a saturation in rate with respect to the actual proton concentrations despite the recalculation of the concentrations undertaken in Table 1. Figure 6B would strongly indicate second-order kinetics in nitrous acid, and a plot of nitrosation rate versus nitrous acid to the second power is linear (plot not shown). Table 1 shows that, in the series of data plotted for Fig. 6B, nitrous acid concentrations and excess proton concentrations increase while nitrite concentrations decrease, and the final observed rate of nitrosation is derived from these three species concentrations and not isolated to nitrous acid. At acid concentrations that exceed initial nitrite concentrations, there is a linear dependence in rate on acid concentrations (Fig. 6C). Note that a saturation would have been achieved in nitrous acid concentrations as would be expected in this format in which the nitrous acid saturates at 50 mmol/L. The increase in nitrosation rate with an increase in acid after saturation of HNO₂ indicates that other nitrosants apart from HNO₂ exist in the reaction environment.

**Effect of copper**

Copper has been implicated in many reactions involving organosulfur compounds. Cu(I) is known to be much more active than Cu(II). It is easier to assay for Cu(II) and the addition of Cu(II) to organosulfur reactions involves its initial conversion to the more active Cu(I). Figure 7A shows that addition of Cu(II) rapidly increases the nitrosation reaction. Cu(II) ions, however, also catalyze the decomposition of CANO. A plot of the rate of nitrosation versus Cu(II) concentrations shows the sigmoidal rate increase followed by an abrupt saturation, which is typical of catalytic mechanisms (figure not shown). No nitrosation occurs in the presence of nitrite without acid (which is necessary to generate the HNO₂). Figure 7B, however, shows that, even in the absence of acid, Cu(II) ions can still effect the nitrosation of CA. Cu(I) ions, however, could not effect nitrosation in the absence of acid. Figure 7C shows the catalytic effect of Cu(II) ions on the decomposition of CANO. CANO was prepared in situ from two feed streams of the double-mixing stopped flow ensemble. Cu²⁺ ions derived from cupric chloride were next added from a third feed stream after complete production of CANO. All feed streams were buffered at pH 7.4. The two control experiments (traces a and b) show the slow decomposition expected from CANO solutions at pH 7.4. Trace b was treated with EDTA to sequester all metal ions in the reaction solution. This trace is identical to the one without EDTA (trace a), confirming the absence of metal ions in reagent solution, which can affect the decomposition kinetics. Addition of micromolar quantities of Cu(II) ions effects a big acceleration in the rate of decomposition. Higher Cu(II) concentrations give a strong autocatalytic decay of the nitrosothiol (trace f in Fig. 7C). The catalytic effect of Cu(II) ions on the decomposition can
also be evaluated through the rate of production of nitric oxide, which is known to be the major product, together with the disulfide, of nitrosothiol decompositions. Figure 7D shows EPR spectra using the nitroethane trap, of the enhanced production of NO with addition of Cu(II) ions. All three spectra were taken after the same time lapse from CANO incubation, and thus the peak heights of the NO-induced spectra can be correlated with the extent of reaction.

**Transnitrosation**

The most abundant thiol in the physiological environment is glutathione (GSH). A pertinent reaction to study would be the interaction of any nitrosothiol formed in the physiological environment with glutathione. There is a crucial question about the lifetimes of the nitrosothiols formed. If thiols are carriers of NO from the point of production to the point of its usage, the nitrosothiols formed should be stable enough and have a long enough lifetime to enable them to perform this activity. Spectrophotometric study of transnitrosation is rendered complex by the fact that many nitrosothiols seem to absorb around the same wavelength and with similar absorptivity coefficients. Figure 8A shows a series of superimposed spectra of four well-known nitrosothiols, which show nearly identical spectra. Figure 8B shows the spectrophotometric characterization of possible transnitrosation in mixtures of CANO and GSH. GSNO has a higher absorptivity coefficient than CANO at the same wavelength, and this explains the initial increase in absorbance on addition of GSH to CANO. LC–MS data have shown that the reaction mixture can transiently (before attainment of the equilibrium mixture) contain any of at least six components at varying times and concentrations: CANO, S-nitrosoglutathione (GSNO), GSH, CA–CA (cystamine), GSSG (oxidized glutathione), and CA–GS (mixed disulfide). The rate of transnitrosation will be difficult to evaluate owing to the presence of the mixed disulfide. Figure 8B shows that, relatively, CANO is more stable than GSNO because of the rapid rate of consumption of GSNO upon its formation as compared with the much slower CANO decomposition with or without EDTA. Figure 9 shows a GC–MS spectrum of mixtures of CANO and GSH. This involved a 3:5 ratio of CANO to GSH, which is also trace f in Fig. 8B. The remarkable aspect of this spectrum is the absence of the GSSG, whose precursor is GSNO. Thus, GSNO, when formed, rapidly decomposes by reacting with CA to form the mixed disulfide and releasing NO. The cystamine formed and observed in the spectrum is derived from the normal decomposition of CANO (see rxn. [R10], vide infra).

**Mechanism**

The overall rate of nitrosation is dependent on the rate of formation and accumulation of the nitrosating agents. A number of possibilities have been suggested, and these include HNO₂, NO, NO₂, N₂O₃, and the nitrosonium cation, *NO. Some nitrosants cannot exist under certain conditions; for example, the *NO cannot exist at high pH conditions and NO₂ cannot be produced in strictly anaerobic conditions. Figure 10 shows two nitrosation reactions at equivalent conditions except that one was run in solutions that had been degassed with argon and immediately capped. Aerobic environments can form NO₂ from NO and subsequently N₂O₃ will also be formed.
Both NO$_2$ and N$_2$O$_3$ are potent nitrosating agents.\textsuperscript{29} There is a slightly higher rate of nitrosation in aerobic environments when compared with those in anaerobic environments. This is a reflection of the low dissolved oxygen concentrations available in aqueous solutions at approximately $2 \times 10^{-4}$ mol/L. Thus, although formation for these nitrosants is favored, their viability is hampered by the low concentrations of oxygen as well as NO, which is only produced during the decomposition of the nitrosothiol. NO was not detected during the formation of CANO. In the aqueous environment utilized for these experiments, reaction of aqueous N$_2$O$_3$ to nitrous acid would be overwhelmingly dominant. This is a rapid equilibrium reaction that favors the weak acid, HNO$_2$.

\[ N_2O_3 \text{(aq)} + H_2O \text{(l)} \rightleftharpoons 2\text{HNO}_2 \text{(aq)}; \quad K_H \] \textsuperscript{[R8]}

Early studies by Bunton and Steadman\textsuperscript{51} had erroneously estimated a value for $K_H^{-1} = 0.20$ (mol/L)$^{-1}$, which would indicate appreciable amounts of N$_2$O$_3$(aq) over nitrous acid. A more accurate evaluation by Markovits et al.\textsuperscript{52} gave a more realistic value of $K_H^{-1} = 3.03 \times 10^{-3}$ (SD 0.23 $\times 10^{-3}$) (mol/L)$^{-1}$. According to Turney,\textsuperscript{53} appreciable amounts of N$_2$O$_3$ can only exist in highly acidic environments, in the region of 5 mol/L perchloric acid. Such conditions are not applicable for the physiological environment. Most nitrosation studies in aerobic NO environments neglect the contribution to nitrosation by NO$_2$ and only concentrate on NO and N$_2$O$_3$. NO$_2$ concentration will always be negligibly low because of rxn. [R6], which is two orders of magnitude greater than the rate of nitrosation by NO$_2$ ($1.1 \times 10^9$ (mol/L)$^{-1}$ s$^{-1}$ vs $2.0 \times 10^7$ (mol/L)$^{-1}$ s$^{-1}$, respectively)\textsuperscript{54,55} is hampered by the low concentrations of oxygen available. In the presence of NO, it is more unstable with respect to N$_2$O$_3$.

All nitrosation kinetics show first-order dependence in both thiol and nitrite with a complex dependence on acid. Data in Figs. 4 and 5 show this strong first-order dependence on CA and on nitrite. In conditions in which acid concentrations are less than initial nitrite concentrations, the effect of nitrite is more difficult to interpret, since a change in its concentrations concomitantly induces a change in both acid and HNO$_2$ concentrations, and thus its sole effect cannot be isolated. First-order kinetics in nitrite strongly suggest that the major nitrosant at the beginning of the reaction is HNO$_2$, since there is a direct correlation between nitrite concentrations and HNO$_2$, especially when there is consistently excess acid over nitrite. After performing calculations for the relevant reactive species in the reaction medium (Table 1), the plot of the initial rate of nitrosation versus nitrous acid concentrations shows a parabolic dependence, which erroneously suggests second-order kinetics in HNO$_2$ (Fig. 6B) for the region in which initial nitrite concentrations exceed acid. As mentioned in the Results section, the data in Table 1 show that HNO$_2$ is not changing in isolation. Both nitrite and acid are also concomitantly changing, and so the effect observed in rate is not solely attributable to changes in HNO$_2$ concentrations. When acid is in excess over nitrite,
the amount of HNO₂ saturates, but as can be seen in Fig. 6C, the rate keeps increasing linearly with an increase in acid, indicating the presence of multiple nitrosating agents apart from HNO₂. The simplest rate law that can explain this nitrosation rate involves +NO and HNO₂. +NO can be formed from the protonation of HNO₂:

\[
\text{HNO}_2 + H_3O^+ \rightarrow H(OH)N = \text{O}^+ + H_2O; \quad K_b \ [R9]
\]

\((H(OH))N=O^+\) is a hydrated nitrosonium cation, \(+\text{NO} + H_2O\).

The overall rate of reaction, using these two nitrosants gives

\[
\text{Rate} = \left( \frac{[N(\text{III})]}{[H^+] + K_a} \right) [CA] \left( k_1 + k_2K_b \left[ H^+ \right] \right) \ [1]
\]

where \([N(\text{III})]_T\) is the total concentrations of nitrous acid and nitrite as calculated in Table 1 (columns 4 and 5) and \([H^+]\) is the calculated acid concentration in column 6 of Table 1. \(K_a\) is the acid dissociation constant of nitrous acid, \(K_b\) is the equilibrium for the protonation of nitrous acid (rxn. [R9]), \(k_1\) is the bimolecular rate constant for the nitrosation of CA by HNO₂, and \(k_2\) is the bimolecular rate constant for the nitrosation by +NO. \(K_a\) was taken as \(5.62 \times 10^{-4}\) (mol/L)⁻¹ and, thus, at high acid concentrations, a plot of Rate/[N(III)]ₜ [CA] vs [H⁺] should give a straight line with slope \(k_2K_b\) and intercept \(k_1\). This type of plot only applies to conditions of excess acid over nitrite (Fig. 6C). One can derive values for \(k_1\) and product \(k_2K_b\) from these series of plots. A value of \(k_1 = 17.9 \pm 1.5\) (mol/L)⁻¹ s⁻¹ was derived from a series of several complementary experiments. \(k_2\) was deduced as \(6.70 \times 10^4\) (mol/L)⁻¹ s⁻¹ after assuming a \(K_b\) value of \(3.16 \times 10^{-2}\) (mol/L)⁻¹.

**Decomposition of CANO**

RSNOs differ greatly in their stabilities, and most of them have never been isolated in solid form. Homolysis of the S–N bond occurs both thermally and photochemically, but, generally, these processes are very slow at room temperature and in the absence of radiation of the appropriate wavelength. In vivo decomposition of RSNOs is likely to proceed through other pathways apart from these two. The presence of trace amounts of Cu(II) or Cu(I) can efficiently catalyze RSNOs to disulfide and nitric oxide (see Figs. 7A–7D).

\[
2\text{RSNO} \rightarrow \text{RSSR} + 2\text{NO} \ [R10]
\]

The characterization of the S–N bond is crucial in determining the stability of RSNOs, and, unfortunately, very little work has been performed in this area. The little crystallographic data available on isolable RSNOs indicate that the S–N bond is weak, sterically unhindered, elongated at between 0.17 and 0.189 nm, and exists as either the cis (syn) or trans (anti) conformers. These conformers are nearly isoenergetic but separated by a relatively high activation barrier of ~11 kcal mol⁻¹ (1 cal = 4.184 J). CANO is a primary S-nitrosothiol, with primary and secondary nitrosothiols known to be kinetically unstable to decomposition when compared with tertiary nitrosothiols. S-Nitroso-N-acetylpenicillamine is the most well-known stable tertiary nitrosothiol. The mechanism of decomposition of nitrosothiols was believed to proceed through the homolytic cleavage of this weak S–N bond. Computed
S–N bond dissociation energies for nitrosothiols show very little variation, however, between primary, secondary, and tertiary nitrosothiols, resulting in very similar homolysis rates of reaction at elevated temperatures for all nitrosothiols. The computed activation parameters for thermal homolysis to occur are prohibitively high for this to be a dominant pathway under normal biological conditions. For example, assuming an approximate S–N bond energy of 31 kcal, a rough calculation will indicate that if decomposition was to proceed solely through homolysis, the half-life of a typical nitrosothiol would be in the region of years instead of the minutes to hours that are observed. This analysis does not support the work reported by Roy et al. in which they established homolysis as the major decomposition pathway. They also managed to trap the thyl radical, RS·. Close examination of their reaction conditions show high trap concentrations (0.2 mol/L DMPO) and nitrosothiol (0.1 mol/L). Such conditions would catch even very low concentrations of thyl radical, but would not necessarily indicate that this was the dominant pathway and whether it was derived solely from homolytic cleavage of the S–N bond. Nitrosothiol decomposition rates in the absence of metal ions are known to be mostly zero order and this is inconsistent with an S–N bond homolysis-driven decomposition, which would have been expected to be first order. In the presence of excess thiol, however, decomposition of nitrosothiols proceeds via a first-order decay process. The absence of first-order decay kinetics in the decomposition of CANO also rules out homolysis as a dominant pathway. For a series of nitrosothiols, the homolysis-dependent decomposition rates are not dependent on the bulkiness of the substituent groups attached to the nitrosothiols, which seems to enhance the calculations that showed no discernible difference in the S–N bond energies irrespective of whether the nitrosothiol was primary, secondary, or tertiary. In aerated solutions, the decay was autocatalytic, thus implicating N₂O₃ as the autocatalytic propagating species (see rxns. [R6] and [R7]). Nitrosothiol decompositions were much slower in solutions in which NO was not allowed to escape, implicating NO as a possible retardant of nitrosothiol decomposition. This can be explained by the recombination of the thyl radical with NO in the reaction cage after initial cleavage. Elimination of NO from the medium by bubbling air forces rxn. [R11] to the right.

\[ RSNO \leftrightarrow RS^* + NO \]  \[ R11 \]

No mechanism has yet been proposed for the implication of N₂O₃ in the autocatalytic mechanism. One can assume that N₂O₃ should assist in the cleavage of the S–N bond, thus accelerating the rate of homolytic decomposition of RSNOs.

In aerated solutions, NO then undergoes rxns. [R6] and [R7] to yield N₂O₃.

\[ RSNO + N_2O_3 \rightarrow RSN(NO_2)^* + NO \]  \[ R12 \]

The incorporation of the NO₂⁻ group in the nitrosothiol would weaken the S–N bond, leading to its easy cleavage and regeneration of the N₂O₃ molecule, which will further catalyze the decomposition.
RSN (NO₂) O• → RS• + N₂O₃  \[\text{[R13]}\]

The sequence of rxns. [R11]–[R13] will deliver autocatalysis in N₂O₃, since rxn. [R13] produces N₂O₃, which recycles back into rxn. [R12].

Recent results have suggested a novel pathway for the decomposition of nitrosothiols that involves an initial internal rearrangement to form an N-nitrosation product.\(^6\) Typical nitrosothiol decomposition products after such an internal rearrangement would include thirranes and 2-hydroxy-mercaptans. This mechanism has been used successfully to explain the production of a small but significant quantity of 2-hydroxy-3-mercaptopropionic acid (apart from the dominant product cystine) in the decomposition of S-nitrosocysteine. The GC–MS spectrum shown in Fig. 2, however, gives a dominant signal for dimeric cystamine, with very little evidence of any other significant products, indicating very little initial N-transnitrosation.

The copper-catalyzed decomposition of nitrosothiols has been well-documented.\(^6\)–\(^8\) The varying rates of nitrosothiol decompositions reported by different research groups can be traced possibly to adventitious metal ion catalysis from the water used in preparing reagent solutions. Even in our case, with the maximum metal ion concentrations of 0.43 ppb in Pb\(^{2+}\), small differences can be observed in EDTA-laced reaction solutions and those that are not (see traces a and b in Fig. 7C). Nitrosothiol decompositions have also implicated other metal ions and not just specifically copper.\(^6\) Although Cu(I) ions are used to catalyze nitrosothiol decomposition, the active reagent is Cu(I). In this catalytic mechanism, Cu redox cycles between the +2 and +1 oxidation states. Only trace amounts of Cu(I) are needed to effect the catalysis, and these can be generated from the reduction of Cu(II) ions from minute amounts of thiolate anions that exist in equilibrium with the nitrosothiol.

RSNO ↔ RS⁻ + NO⁺  \[\text{[R14]}\]

2Cu\(^{2+}\) + 2RS⁻ → RSSR + 2Cu\(^+\)  \[\text{[R15]}\]

Cu\(^+\) + RSNO → [RS (Cu) NO]⁺  \[\text{[R16]}\]

[RS (Cu) NO]⁺ → RS⁻ + NO + Cu\(^{2+}\)  \[\text{[R17]}\]

The cascade of rxns. [R14]–[R17] continues until the nitrosothiol is depleted and converted to the disulfide. The complex formed in rxn. [R16] has been justified through density functional theory studies that report that Cu(I) binds more strongly to the S center than to the N center of the nitrosothiol.\(^6\) This preferential binding weakens and lengthens the S–N bond, thus facilitating its cleavage. It is difficult to extrapolate this proposed mechanism into the physiological environment, but some experimental results have shown that protein-bound Cu\(^{2+}\) catalyzed nitric oxide generation from nitrosothiols, but not with the same
efficiency as the free hydrated ion. Further studies are needed to evaluate the efficiency of this catalysis, especially in copper-based enzymes.

**Effect of Cu(II) on the nitrosation of cysteamine (Fig. 7B)**

The fact that Cu(II) ions can assist in nitrosating CA from nitrite in the absence of acid while Cu(I) ions are unable to effect this can be explained by recognizing that nitrosation is an electrophilic process, where NO effectively attaches as the (formally) positive charge, while eliminating a positively charged proton. Cu(II), in this case, acts by complexing to the nitrite and forming a nitrosating species that will release the Cu(II) ions after nitrosation.

Further experiments on the effect of nitrous acid—Since nitrous acid is the major nitrosation species, another series of experiments were undertaken in which the effect of nitrous acid could be unambiguously determined. For the data shown in Fig. 11A, nitrite and acid were kept at equimolar concentrations and varied with that constraint. Final proton and nitrite concentrations after dissociation are also equal for electroneutrality (see Table 2). Thus, if our simple mechanism is plausible, a plot of nitrous acid versus initial nitrosation rate should be linear with the expected slight tailing at either extreme of the nitrous acid concentrations owing to nonlinear generation of the free acid derived from the quadratic equation. Using the same rate law as eq. [1], a value of $k_1$ can be derived and compared with values obtained from the Fig. 6C plot. Kinetics constants derived from the Fig. 11B plot are not expected to be as accurate as those from Fig. 6C, but should deliver values close to those from high acid environments. For the case of nitrous acid as the nitrosating agent (too low a concentration of excess $H_3O^+$ to effect equilibrium of rxn. [R9]), data derived from Fig. 11B delivered an upper limit rate constant for $k_1$ that was higher than that derived from Fig. 6C-type data, but well within the error expected for such a crude extrapolation. Other sets of kinetics data were derived in which CA concentrations were varied while keeping the nitrous acid concentrations constant. Varying nitrous acid concentrations were used and, for each, a CA dependence was plotted (data not shown). These were straight lines through the origin and with the slope determined by the concentration of nitrous acid and the free protons existing in solution after the dissociation of nitrous acid. For these kinetics data, eq. [1] had to be evaluated completely, since $K_a \approx [H_3O^+]$ and no valid approximation could be made. The values of $k_1$ and $k_2K_b$ were utilized in eq. [1] to recalculate initial rates and slopes with respect to CA concentrations. These values were utilized to check the accuracy of the kinetics constants derived from high-acid-dependent experiments. It was noted that, within experimental error, these values checked and were generally robust.

**Simulations**

A small set of simple reactions was utilized to simulate the nitrosation kinetics. This set is shown in Table 3. At low acid concentrations, the nitrosation is dominated by $HNO_2$ as the nitrosant, and as acid concentrations are increased, both nitrous acid and the nitrosonium
cation are involved in the nitrosation. Even though Table 3 includes nitrosation by \(N_2O_3\), the lack of oxygen (approximately kept constant at \(2.0 \times 10^{-4}\) mol/L) and its most favored and rapid reaction to nitrous acid made the nitrosation by \(N_2O_3\) insignificant. \(N_2O_3\) is the dehydrated form of HNO\(_2\) (rxn. [R8]), while reaction conditions were in an overwhelming excess of water. Its insignificant concentration ensured that the simulations were insensitive to value of the bimolecular nitrosation constant for \(N_2O_3\) \((k_{M8})\). Figure 12B clearly shows that the concentration of \(N_2O_3\) never rises to a level where it would be significant. The rate constants adopted for rxn. [M1] (Table 3) were not crucial for as long as they were fast (in both directions) and were connected by \(K_a\). Increasing the forward and reverse rate constants while maintaining \(K_a\) only slowed the simulations, but did not alter the simulation results. Reaction [M2] (Table 3) was also a rapid hydration reaction that strongly favored HNO\(_2\), thus rapidly depleting any \(N_2O_3\) formed or merely keeping these concentrations depressed should the tandem rxns. [R6] + [R7] have been added to the mechanism. These were not utilized for this simulation because EPR experiments had not detected NO at the beginning of the reaction and had only detected NO during the decomposition of the nitrosothiol. There are no reliable thermodynamics parameters for rxn. [M3] (Table 3), since this would be heavily dependent on the pH of the medium. For the purposes of this simulation and some of our previous estimates, the equilibrium constant for rxn. [M3] was set at \(3.16 \times 10^{-2}\) (mol/L\(^{-1}\)). Kinetics constants for rxns. [M4] and [M5] (Table 3) were derived from this study. In the absence of significant concentrations of \(N_2O_3\), the simulations were insensitive to the kinetics constants utilized for rxns. [M6] and [M8] (Table 3). Reaction [M7] (Table 3) was assumed to be slow and, in the absence of Cu(II) ions or other adventitious ions, this is indeed so (see Figs. 7A and 7C). The simulations are shown in Fig. 12A, which gives a reasonably good fit from such a crude mechanism. Figure 12B is provided to show the concentration variation of some of the species that could not be experimentally determined. Of note is that \(N_2O_3\) concentrations are depressed throughout the lifetime of the nitrosation. Even though \(\cdot NO\) is also depressed in concentration, this can be explained by its high rate of nitrosation as soon as it is formed and an equilibrium that favors its hydration back to HNO\(_2\) once the thiol is depleted.

### Conclusion

This detailed kinetics study has shown that, even though there are several possible nitrosating agents in solution, in the physiological environment, with its slightly basic pH and low oxygen concentrations, the major nitrosant is HNO\(_2\). Furthermore, the lifetime of CANO is not long enough to be able to be effectively used as a carrier of NO. GSH, which has been touted as a possible carrier of NO, appears to have a shorter lifetime than CANO. However, because of its overwhelming concentration in the physiological environment it might still be a major carrier of NO when it is compared with other thiols. Protein thiols, it appears, would be the best carriers by either intra- or inter-molecular transfer of NO through protein nitrosothionylation through exposed and accessible cysteine groups.\(^{69}\)

### Acknowledgements

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report are those of the authors and do not necessarily represent the views of the National Institute for Occupational Safety and Health.

References


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Fig. 1.
UV–vis spectral scan of reactants 0.01 mol/L cysteamine hydrochloride (CA, a), 0.001 mol/L NaNO$_2^-$ (b), 0.001 mol/L HNO$_2$ (c), and product 0.001 mol/L S-nitrosocysteamine (CANO, d).
Fig. 2.
Mass spectrometry analysis of the product of nitrosation of cysteamine hydrochloride (CA) showing (A) formation of S-nitrosocysteamine (CANO), $m/z = 107.9999$; (B) product of decomposition of CANO after 24 h. Cystamine, a disulfide of CA was produced with a strong peak, $m/z = 153.0608$. 

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Fig. 3.
Electron paramagnetic resonance (EPR) spectra of the NO radical generated during the decomposition of S-nitrosocysteamine (CANO) using 0.5 mol/L nitroethane (NE) in 1.0 mol/L NaOH solution as the spin trap. (A) 0.5 mol/L NE, (B) 0.01 mol/L NO$_2^-$, (C) 0.04 mol/L cysteamine hydrochloride (CA), (D) [CANO] = 0.01 mol/L, (E) [CANO] = 0.02 mol/L, (F) [CANO] = 0.03 mol/L, and (G) [CANO] = 0.04 mol/L.
Fig. 4.
(A) Absorbance traces showing the effect of varying cysteamine hydrochloride (CA) concentrations. The reaction displays first-order kinetics in CA, $[\text{NO}_2^-]_0 = 0.10 \text{ mol/L}$; $[\text{H}^+]_0 = 0.10 \text{ mol/L}$; and (a) $[\text{CA}]_0 = 0.005 \text{ mol/L}$; (b) $[\text{CA}]_0 = 0.006 \text{ mol/L}$; (c) $[\text{CA}]_0 = 0.007 \text{ mol/L}$; (d) $[\text{CA}]_0 = 0.008 \text{ mol/L}$; (e) $[\text{CA}]_0 = 0.009 \text{ mol/L}$; and (f) $[\text{CA}]_0 = 0.010 \text{ mol/L}$. (B) Initial rate plot of the data in Fig. 4A. The plot shows the strong first-order dependence of the rate of formation of S-nitrosocysteamine (CANO) on CA, with an intercept kinetically indistinguishable from zero.
Fig. 5.
Initial rate plot from the data derived from varying nitrite ion concentrations in excess acid over nitrite. \([\text{CA}]_0 = 0.10 \text{ mol/L}; [\text{H}^+]_0 = 0.10 \text{ mol/L};\) and \([\text{NO}_2^-]_0\) was varied from 0.005 to 0.010 mol/L. The plot shows strong first-order dependence on initial rate of formation of S-nitrosocysteamine (CANO) on nitrite. CA, cysteamine hydrochloride.
Fig. 6.

(A) Plot of the raw kinetics data for the effect of added acid on the rate of nitrosation. \([\text{CA}]_0 = 0.05 \text{ mol/L}; [\text{NO}_2^-]_0 = 0.05 \text{ mol/L}; \) and \([\text{H}^+]_0\) was varied between 0.01 and 0.08 mol/L. For the first four data points, \([\text{NO}_2^-]_0 > [\text{H}^+]_0\). The linear portion of the plot represents excess proton concentrations over nitrite. Acid concentrations plotted here are those experimentally added and not calculated. (B) Plot of initial rate versus nitrous acid concentrations of the experimental data in Fig. 6A and calculated in Table 1. This plot involves the region at which initial nitrite concentrations exceed acid concentrations. (C) Plot showing the linear dependence of the rate of formation of S-nitrosocysteamine (CANO) on \([\text{H}_3\text{O}^+]\) in high acid concentrations, where initial acid concentrations exceed nitrite concentrations. Nitrous acid concentrations are nearly constant in this range (see Table 1).
Fig. 7.

(A) Absorbance traces showing the effect of varying Cu$^{2+}$ concentrations on the rate of formation of CANO. There is a progressive increase in the rate of S-nitrosocysteamine (CANO) formation with an increase in Cu$^{2+}$ concentration. $[CA]_0 = [NO_2^-]_0 = [H^-]_0 = 0.05$ mol/L; and (a) $[Cu^{2+}]_0 = 0.00$ mol/L, (b) $[Cu^{2+}]_0 = 5$ μmol/L, (c) $[Cu^{2+}]_0 = 15$ μmol/L, (d) $[Cu^{2+}]_0 = 100$ μmol/L, (e) $[Cu^{2+}]_0 = 1$ mmol/L, and (f) $[Cu^{2+}]_0 = 2.5$ mmol/L. CA, cysteamine hydrochloride. (B) Absorbance traces showing the formation of CANO via a reaction of CA, nitrite, and copper(II) without acid. The amount of CANO formed increases with an increase in Cu$^{2+}$ concentration. No formation of CANO was observed with Cu$^+$. The result shows the catalytic effect of Cu$^{2+}$ on the rate of formation of S-nitrosothiol (RSNO). $[CA]_0 = [NO_2^-]_0 = [H^-]_0 = 0.05$ mol/L; and (a) $[Cu^{2+}] = 0.0025$ mol/L, (b) $[Cu^{2+}] = 0.0030$ mol/L, (c) $[Cu^{2+}] = 0.0035$ mol/L, (d) $[Cu^{2+}] = 0.0040$ mol/L, (e) $[Cu^{2+}] = 0.0045$ mol/L, and (f) $[Cu^{2+}] = 0.0050$ mol/L. (C) Effect of Cu$^{2+}$ on the stability of CANO in a pH 7.4 phosphate buffer. This reaction involves the reduction of Cu$^{2+}$ to Cu$^+$, which causes the decomposition of CANO. $[CANO]_0 = 0.028$ mol/L; (a) $[Cu^{2+}]_0 = 0.00$ mol/L (pure CANO), (b) $[Cu^{2+}]_0 = 0.00$ mol/L (pure CANO in 0.00001 mol/L EDTA), (c) $[Cu^{2+}]_0 = 5$ μmol/L, (d) $[Cu^{2+}]_0 = 10$ μmol/L, (e) $[Cu^{2+}]_0 = 20$ μmol/L, and (f) $[Cu^{2+}]_0 = 30$ μmol/L. (D) Electron paramagnetic resonance (EPR) spectra of the NO radical showing the catalytic effect of Cu$^{2+}$ on the rate of decomposition of CANO. The intensity of the spectra increases with an increase in Cu$^{2+}$ concentration. $[CANO]_0 = 0.01$ mol/L; (a) $[Cu^{2+}]_0 = 0.00$, (b) $[Cu^{2+}]_0 = 1.0 \times 10^{-4}$ mol/L, and (c) $[Cu^{2+}]_0 = 2.0 \times 10^{-4}$ mol/L.
Fig. 8.
(A) Superimposed spectra of four well-known nitrosothiols (a) CysNO, (b) S-nitrosocysteamine (CANO), (c) S-nitrosoglutathione (GSNO), and (d) nitrosothiol of penicillamine (SNAP). Three of them nearly have the same $\varepsilon$ at 545 nm. The observed separation in the absorbances of these three at 545 nm is due to staggering of the time lag before acquiring the spectrum. (B) The effect of glutathione (GSH) on the stability of CANO in a pH 7.4 phosphate buffer. All experimental traces have $[CA]_0 = 0.03$ mol/L, $[NO_2^-]_0 = 0.03$ mol/L, and $[H^+]_0 = 0.05$ mol/L. (a) $[GSH] = 0$, (b) $[GSH] = 0$, (c) $[GSH] = 0.01$ mol/L, (d) $[GSH] = 0.02$ mol/L, (e) $[GSH] = 0.03$ mol/L, and (f) $[GSH] = 0.05$ mol/L. Traces a and b are controls. Trace a has no EDTA. Trace b is the same as trace a with 10 $\mu$mol/L of EDTA. There is no significant difference between traces a and b, indicating that the water used for preparing reagent solutions did not contain enough trace metal ions to affect the kinetics. Ten microlitres of EDTA was added to
Fig. 9.
A GC–MS spectrum of a 3:5 ratio of $S$-nitrosocysteamine (CANO) to glutathione (GSH); trace $f$ in Fig. 8B. Final products were predominantly the mixed disulfide and cysteamine with no evidence of oxidized glutathione (GSSG). CA, cysteamine hydrochloride.
Fig. 10.
Evaluating the effect of oxygen on the rate of nitrosation of cysteamine. The solutions purged with argon (trace a) give a very slightly lower rate of nitrosation, indicating a small and insignificant contribution of nitrosation by $N_2O_3$. $[CA]_0 = 0.05 \text{ mol/L}; [H^+]_0 = 0.08 \text{ mol/L}$; and $[NO_2^-]_0 = 0.07 \text{ mol/L}$. CA, cysteamine hydrochloride.
Fig. 11.

(A) Evaluation of unambiguous nitrous acid dependence by employing \([H^+]_0 = [NO_2^-]_0\) and varying both at the same equimolar concentrations. \([CA]_0\) fixed at 0.25 mol/L. (a) \([H^+]_0 = [NO_2^-]_0 = 0.005\) mol/L, (b) \([H^+]_0 = [NO_2^-]_0 = 0.010\) mol/L, (c) \([H^+]_0 = [NO_2^-]_0 = 0.015\) mol/L, (d) \([H^+]_0 = [NO_2^-]_0 = 0.020\) mol/L, (e) \([H^+]_0 = [NO_2^-]_0 = 0.025\) mol/L, (f) \([H^+]_0 = [NO_2^-]_0 = 0.030\) mol/L, and (g) \([H^+]_0 = [NO_2^-]_0 = 0.035\) mol/L. (B) Plot of initial rate of nitrosation versus initial nitrous acid concentrations as calculated in Table 2 and derived from the data in Fig. 11A. This plot is linear, as opposed to Fig. 6C.
Fig. 12.
(A) Simulation of the short set of reactions shown in Table 3. Solid lines are experimental data and the symbols indicate simulations. The mechanism is dominated by the value of the bimolecular rate constant for the nitrosation of cysteamine hydrochloride (CA) by HNO₂ ($k_1$). [CA]₀ = [NO₂⁻]₀ = 0.05 mol/L. (B) Results from the modeling that delivered simulations of the two traces shown in Fig. 12A. These simulations show the concentration variations of those species that could not be experimentally monitored. Neither N₂O₃ nor NO⁺ rise to any significant concentration levels.
Table 1
Input species concentrations and final reagent concentrations for acid dependence experiments (units of mol/L).

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**Table 2**

Input species concentration and final reagent concentrations for nitrous acid dependence experiments with $[\text{H}^+]_0 = [\text{NO}_2^-]_0$.

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Table 3

The set of reactions used for simulating the nitrosation of cysteamine. RSH, cysteamine.

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<th>Reaction</th>
<th>$k_f$</th>
<th>$k_r$</th>
</tr>
</thead>
<tbody>
<tr>
<td>M1</td>
<td>$\text{H}^+ + \text{NO}_2^- \rightleftharpoons \text{HNO}_2$</td>
<td>$1.10 \times 10^9$</td>
<td>$6.18 \times 10^5$</td>
</tr>
<tr>
<td>M2</td>
<td>$2\text{HNO}_2 \rightleftharpoons \text{N}_2\text{O}_3 + \text{H}_2\text{O}$</td>
<td>$1.0 \times 10^7$</td>
<td>$1.0 \times 10^8$</td>
</tr>
<tr>
<td>M3</td>
<td>$\text{HNO}_2 + \text{H}^+ \rightleftharpoons \text{N}=\text{O} + \text{H}_2\text{O}$</td>
<td>$3.16 \times 10^5$</td>
<td>$1.00 \times 10^7$</td>
</tr>
<tr>
<td>M4</td>
<td>RSH + HNO$_2$ $\rightarrow$ RSNO + H$_2$O</td>
<td>17.9</td>
<td>0</td>
</tr>
<tr>
<td>M5</td>
<td>RSH + N=O $\rightleftharpoons$ RSNO + H$^+$</td>
<td>$6.7 \times 10^6$</td>
<td>$1.0 \times 10^{-3}$</td>
</tr>
<tr>
<td>M6</td>
<td>N$_2$O$_3$ $\rightleftharpoons$ NO + NO$_2$</td>
<td>$8.0 \times 10^9$</td>
<td>$6.0 \times 10^8$</td>
</tr>
<tr>
<td>M7</td>
<td>2RSNO $\rightleftharpoons$ RSSR + 2NO</td>
<td>$4.0 \times 10^{-3}$</td>
<td>0</td>
</tr>
<tr>
<td>M8</td>
<td>RSH + N$_2$O$_3$ $\rightarrow$ RSNO + HNO$_2$</td>
<td>$5.0 \times 10^3$</td>
<td>0</td>
</tr>
</tbody>
</table>

Note: $k_f$, forward rate constant; $k_r$, reverse rate constant.