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# Dinitrosyl Iron Complexes with Cysteine. Kinetics Studies of the Formation and Reactions of DNICs in Aqueous Solution

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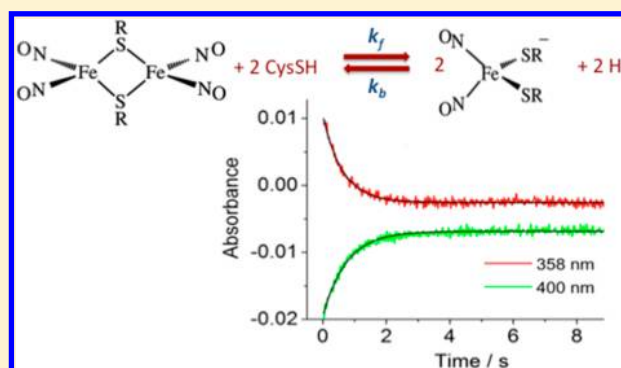
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## S Supporting Information

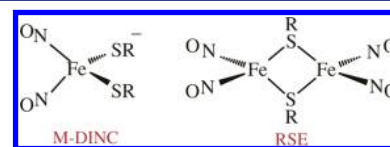
**ABSTRACT:** Kinetics studies provide mechanistic insight regarding the formation of dinitrosyl iron complexes (DNICs) now viewed as playing important roles in the mammalian chemical biology of the ubiquitous bioregulator nitric oxide (NO). Reactions in deaerated aqueous solutions containing FeSO<sub>4</sub>, cysteine (CysSH), and NO demonstrate that both the rates and the outcomes are markedly pH dependent. The dinuclear DNIC Fe<sub>2</sub>(μ-CysS)<sub>2</sub>(NO)<sub>4</sub>, a Roussin's red salt ester (Cys-RSE), is formed at pH 5.0 as well as at lower concentrations of cysteine in neutral pH solutions. The mononuclear DNIC Fe(NO)<sub>2</sub>(CysS)<sub>2</sub><sup>-</sup> (Cys-DNIC) is produced from the same three components at pH 10.0 and at higher cysteine concentrations at neutral pH. The kinetics studies suggest that both Cys-RSE and Cys-DNIC are formed via a common intermediate Fe(NO)(CysS)<sub>2</sub><sup>-</sup>. Cys-DNIC and Cys-RSE interconvert, and the rates of this process depend on the cysteine concentration and on the pH. Flash photolysis of the Cys-RSE formed from Fe(II)/NO/cysteine mixtures in anaerobic pH 5.0 solution led to reversible NO dissociation and a rapid, second-order back reaction with a rate constant  $k_{\text{NO}} = 6.9 \times 10^7 \text{ M}^{-1} \text{ s}^{-1}$ . In contrast, photolysis of the mononuclear-DNIC species Cys-DNIC formed from Fe(II)/NO/cysteine mixtures in anaerobic pH 10.0 solution did not labilize NO but instead apparently led to release of the CysS<sup>•</sup> radical. These studies illustrate the complicated reaction dynamics interconnecting the DNIC species and offer a mechanistic model for the key steps leading to these non-heme iron nitrosyl complexes.



## INTRODUCTION

Nitric oxide (nitrogen monoxide, NO) is an bioregulator that has important roles in mammalian physiological functions such as vasodilation, inflammation, neuronal transmission, and immune system response.<sup>1</sup> Other NO derivatives, such as S-nitrosothiols (RSNO) and N-nitrosoamines (R<sub>2</sub>NNO), heme and non-heme iron nitrosyl complexes as well as the oxidation products, NO<sub>2</sub><sup>-</sup>, ONOO<sup>-</sup>, and NO<sub>3</sub><sup>-</sup> also have physiological presence, and their specific roles remain the subjects of continuing studies. The dinitrosyl iron complexes (DNICs) comprise one class of such species. DNICs are four-coordinate Fe(NO)<sub>2</sub>L<sub>2</sub><sup>n±</sup> complexes thought to be formed from the chelatable iron pool, thiol-containing ligands (for example, glutathione, cysteine, or protein thiol), and endogenous or exogenous NO.<sup>2</sup> Although DNICs were first discovered some decades ago,<sup>3–6</sup> there has been a recent upsurge of interest in the biological or pathological pathways to which these species contribute.<sup>2,7–15</sup>

Previous studies have demonstrated reversible transformations between the Roussin's red salt ester analogs Fe<sub>2</sub>(μ-RS)<sub>2</sub>(NO)<sub>4</sub>,<sup>6</sup> which are binuclear DNICs, and mononuclear species [Fe(NO)<sub>2</sub>L<sub>2</sub>]<sup>n±</sup> (Figure 1). In aqueous media, these



**Figure 1.** Generic formulas for a mononuclear dinitrosyl iron complex (M-DNIC) (left) and for the binuclear DNIC, a Roussin's red salt ester, (RSE) (right). In the present case, the thiolate RS<sup>-</sup> is the cysteinate anion, and the M-DNIC Fe(NO)<sub>2</sub>(CysS)<sub>2</sub><sup>-</sup> is designated as Cys-DNIC and the RSE Fe<sub>2</sub>(μ-Cys)<sub>2</sub>(NO)<sub>4</sub> is designated as Cys-RSE.

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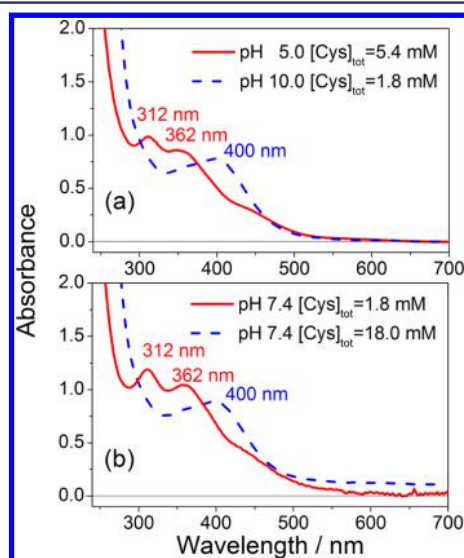
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equilibria are dependent on both pH and thiol ligand concentration.<sup>9</sup> The dinuclear species is the dominant form at lower pH, while the mononuclear complex predominates at higher pH. However, there is relatively little quantitative information regarding the dynamics of the formation and interconversion of these DNICs.<sup>14</sup> The present stopped-flow and flash photolysis kinetics study was initiated to address this issue.

## RESULTS AND DISCUSSION

The present studies use aqueous ferrous sulfate as the iron source, purified NO gas, and L-cysteine (CysSH) in order to model the formation of biological DNICs. In each case the solutions were deaerated in order to avoid the autoxidation of NO. In discussing the reactions of these species in aqueous solutions, it is important to remember that this system involves a number of dynamic pH-dependent equilibria involving various iron complexes as well as that between CysSH and its conjugate base CysS<sup>-</sup>. In this context, kinetics experimental conditions are reported in terms of the following variables:  $[\text{Fe}]_{\text{tot}}$  = the total concentration of iron(II) introduced to the system;  $[\text{Cys}]_{\text{tot}}$  = the total concentration of L-cysteine introduced,  $\sim([\text{CysSH}] + [\text{CysS}^-])$  since  $[\text{Cys}]_{\text{tot}}$  is generally  $\gg [\text{Fe}]_{\text{tot}}$ ; and  $[\text{NO}]_{\text{tot}}$  = the solution nitric oxide concentration, based on the total amount of NO introduced to the system and the partition coefficient between the gas and liquid phases.

**Spectra of Fe/NO/CysSH Solutions at Various pH.** The earlier study by Vanin and co-workers<sup>9</sup> reported that the binuclear DNIC  $\text{Fe}_2(\mu\text{-CysS})_2(\text{NO})_4$  (Cys-RSE) is the dominant iron-containing species in such solutions at low pH. This was confirmed by recording the optical spectrum of a deaerated solution containing Fe(II) (0.18 mM), NO (0.93 mM), and CysSH (5.4 mM) in pH 5.0 citrate buffer. This spectrum (Figure 2, top) displayed the two absorption bands

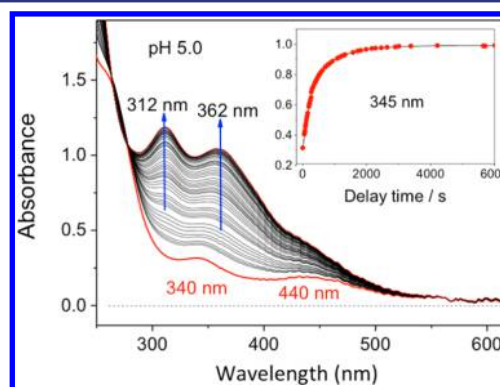


**Figure 2.** Top: Absorption spectra of Cys-RSE (red line) formed from Fe(II) ( $[\text{Fe}]_{\text{tot}} = 0.18$  mM) and NO ( $[\text{NO}]_{\text{tot}} = 0.93$  mM) with CysSH ( $[\text{Cys}]_{\text{tot}} = 5.4$  mM) at pH 5.0 citrate buffer and Cys-DNIC (dashed blue) formed from the mixture of Fe(II) (0.18 mM) and NO (0.93 mM) with CysSH (1.8 mM) at pH 10.0 borate buffer. Bottom: Spectra at pH 7.4 (HEPES buffer) with CysSH at 1.8 mM (red) and at 18.0 mM (dashed blue). In all cases the solutions were deaerated.

(312 and 362 nm) with a shoulder (434 nm) reported<sup>9a</sup> for Cys-RSE and characteristic of that for other RSEs.<sup>16,17</sup> In contrast, a deaerated pH 10.0 borate buffer solution prepared from Fe(II) (0.18 mM), NO (0.93 mM), and CysSH (1.8 mM) displayed a spectrum (Figure 2, top) with a single band centered at 400 nm as reported<sup>9a</sup> for Cys-DNIC. In pH 7.4 HEPES buffer, a spectrum analogous to that of Cys-RSE was seen for  $[\text{CysSH}]_{\text{tot}} = 1.8$  mM, while one close to that of Cys-DNIC was seen for  $[\text{CysSH}]_{\text{tot}} = 18.0$  mM under otherwise analogous conditions (Figure 2, bottom). None of these features are apparent in the spectra of analogous deaerated solutions of Fe<sup>2+</sup> alone, of cysteine alone, or of Fe<sup>2+</sup> and CysSH together in pH 5 citrate buffer, in pH 7.4 HEPES buffer or in pH 10 borate buffer (Supporting Information (SI) Figure S-1).

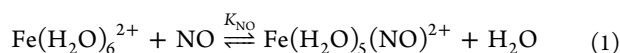
The EPR spectrum recorded for a solution prepared from ferrous sulfate ( $[\text{Fe}]_{\text{tot}} = 0.18$  mM), CysSH ( $[\text{Cys}]_{\text{tot}} = 20$  mM), and NO ( $[\text{NO}]_{\text{tot}} = 1.86$  mM) in deaerated pH 7.4 HEPES buffer (100 mM) indicated the presence of a paramagnetic species with  $g_{\text{ave}} = 2.03$  (SI Figure S-2). This was essentially identical to the EPR spectrum reported by Vanin et al.<sup>9c</sup> for an analogous solution and attributed by these workers to the mononuclear DNIC formed from cysteinate, that is, Cys-DNIC. Similar EPR spectral signatures have been noted by others<sup>17,18</sup> and are considered to be diagnostic of thiolate coordinated DNIC anions of the type  $\text{Fe}^{\text{I}}(\text{NO})_2(\text{RS})_2^-$ . When a less than stoichiometric NO concentration was used ( $[\text{NO}]_{\text{tot}} = 0.093$  mM), the EPR spectrum was still dominated by the signal for Cys-DNIC with  $g = 2.03$ . However, a weaker signal with  $g = 2.04$  was also evident (SI Figure S-3) similar to one that Vanin et al.<sup>9c</sup> have attributed to a triplet mononitrosyl iron complex, presumably  $\text{Fe}(\text{NO})(\text{CysS})_2^-$  in the present case.<sup>18,19</sup> Notably, DFT computations suggest that the mononitrosyl iron intermediate  $\text{Fe}^{\text{I}}(\text{NO})(\text{CysS})(\text{H}_2\text{O})_2$  intermediate also has an  $S = 1$  electronic configuration (see below).

**Kinetics Studies at pH 5.0: Formation of Cys-RSE.** At this pH, the overall reaction to form Cys-RSE from a solution containing Fe(II), NO, and CysSH was quite slow, so optical spectral changes were monitored using a conventional UV-vis spectrophotometer (Figure 3). A solution of Fe(II) (0.18 mM) and CysSH (5.4 mM) was prepared in a deaerated Schlenk cuvette,<sup>20</sup> and it is notable that the resulting spectrum did not display obvious differences from the spectra of solutions of Fe(II) and of CysSH alone, suggesting that there was little



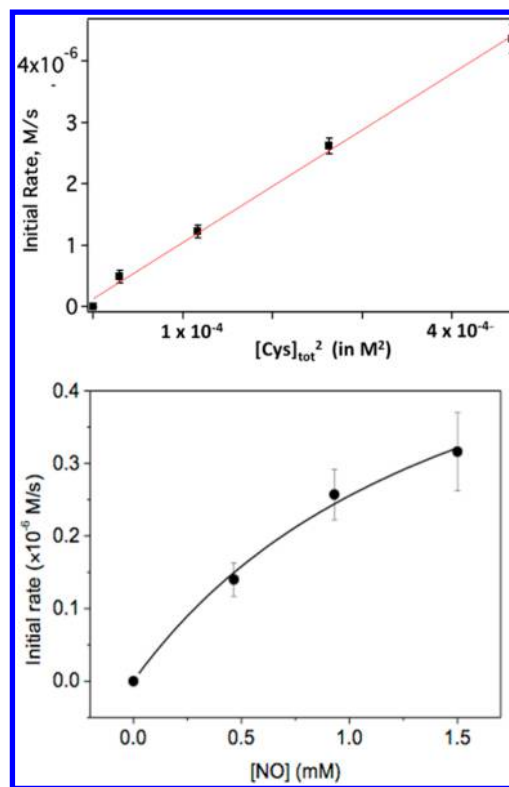
**Figure 3.** Spectral changes seen in pH 5.0 (100 mM citrate buffer) upon adding NO to a deaerated solution of Fe(II) and CysSH to give the final concentrations  $[\text{Fe}]_{\text{tot}} = 0.18$  mM,  $[\text{CysSH}]_{\text{tot}} = 5.4$  mM, and  $[\text{NO}]_{\text{tot}} = 1.86$  mM. The inset shows the temporal absorbance at  $\lambda_{\text{mon}} = 345$  nm.

complex formation between  $\text{Fe}^{2+}$  and CysSH under these conditions. However, immediately upon adding NO ( $P_{\text{NO}} = 1$  atm) and shaking the solution to give a NO concentration of 1.86 mM (based upon the solubility of NO in water),<sup>21</sup> the spectrum displayed two bands with maxima at 340 and 440 nm (Figure 3). These bands are ascribed to rapid formation of  $\text{Fe}(\text{H}_2\text{O})_5(\text{NO})^{2+}$  ( $\text{FeNO}^{2+}$ , eq 1), which was shown by Wannet et al. to be formed from aqueous Fe(II) and NO with a relatively small equilibrium constant  $K_{\text{NO}}$  ( $1.15 \times 10^3 \text{ M}^{-1}$  in pH 5.0, acetate buffer, 23 °C)<sup>22</sup> but with a large second-order rate constant  $k_{\text{NO}} = 1.4 \times 10^6 \text{ M}^{-1} \text{ s}^{-1}$ . The solution spectrum then continued to evolve over a period of minutes to give a final, stable spectrum that can be ascribed to that of Cys-RSE (Figures 2 and 3).



The absorbance changes of the type shown in Figure 3 can be fit to an exponential function to obtain  $k_{\text{obs}}$  values indicating the reaction to be first order in the iron substrate. However, at lower values of  $P_{\text{NO}}$  in the cuvette, a feature characteristic of systems depending on gas/liquid transport was apparent. The gas phase volume in the cuvette was about 8 times that of the liquid phase, and therefore, given NO's low aqueous solubility ( $1.86 \text{ mM atm}^{-1}$  at 298 K),<sup>21</sup> the vast majority of the nitric oxide present in the system for any experiment is in the gas phase. Thus, at lower  $P_{\text{NO}}$ , the solution phase NO concentration is no longer in such excess over  $[\text{Fe}]_{\text{tot}}$  that the  $[\text{NO}]$  remains effectively constant throughout the experiment, unless adequate provision is made to ensure that the gas and liquid phases remain in equilibrium. For a slow reaction, this can be done by periodically shaking the cell, under such conditions, the temporal absorbance changes are exponential (see SI Figure S-4), and this leads to the conclusion that the reaction being observed is first order in  $[\text{Fe}(\text{II})]$ . However, for the faster reactions, an alternative approach is to measure the initial rates, since these are established for the initial reaction conditions. The latter approach was taken for most of the experiments described in this section.

Figure 4 illustrates the results of kinetics experiments in pH 5.0 solution systematically studied as functions of the cysteine and NO concentrations. When  $[\text{Cys}]_{\text{tot}}$  was varied over the range 0–27.0 mM with  $[\text{NO}]_{\text{tot}}$  (0.93 mM) and  $[\text{Fe}]_{\text{tot}}$  (0.18 mM) held constant, the initial rates proved to be a linear function of  $[\text{Cys}]_{\text{tot}}$  squared. Notably, the kinetics behaviors of these relatively slow reactions were essentially indistinguishable whether the Fe(II) and the NO were premixed and the CysSH was then added or the Fe(II) and the CysSH were premixed and the NO was then introduced. This behavior is consistent with the view that, under these conditions, the reactions before the rate-limiting step are rapidly established pre-equilibria. Figure 4 (bottom) illustrates the effect of  $[\text{NO}]_{\text{tot}}$  on the initial rates over the range 0–1.5 mM with  $[\text{Cys}]_{\text{tot}}$  (5.4 mM) and  $[\text{Fe}]_{\text{tot}}$  (0.18 mM) held constant. Notably the latter plot is nonlinear, leveling off at higher values of  $[\text{NO}]$ , an observation consistent with an equilibrium such as that described by eq 1. The curve fit shown in Figure 4 (bottom) is according to eq 2, with the values  $C_1 = (1.19 \pm 0.24) \times 10^2 \text{ M}^{-3} \text{ L}^{-1} \text{ s}^{-1}$  and  $C_2 = (6.2 \pm 2.0) \times 10^2 \text{ M}^{-1}$ . Notably, the value for  $C_2$  falls within the range of values (440–1150  $\text{M}^{-1}$ ) reported for the equilibrium constant ( $K_{\text{NO}}$ ) for eq 1 under similar conditions.<sup>22b</sup>



**Figure 4.** Initial rates as functions of  $[\text{Cys}]_{\text{tot}}$  and of  $[\text{NO}]_{\text{tot}}$  from mixing Fe(II), CysSH, and NO in pH 5.0 (100 mM citrate buffer),  $T = 298 \text{ K}$ . Top: Initial rates vs  $[\text{Cys}]_{\text{tot}}^2$  ( $[\text{Fe}]_{\text{tot}} = 0.18 \text{ mM}$ ;  $[\text{NO}] = 0.93 \text{ mM}$ ). Bottom: Initial rates vs  $[\text{NO}]$  ( $[\text{Fe}]_{\text{tot}} = 0.18 \text{ mM}$ ;  $[\text{NO}]_{\text{tot}} = 0, 0.5, 1.0, \text{ and } 1.5 \text{ mM}$ ;  $[\text{Cys}]_{\text{tot}} = 5.4 \text{ mM}$ ). The curve is a fit of the data according to form of eq 2, see text.

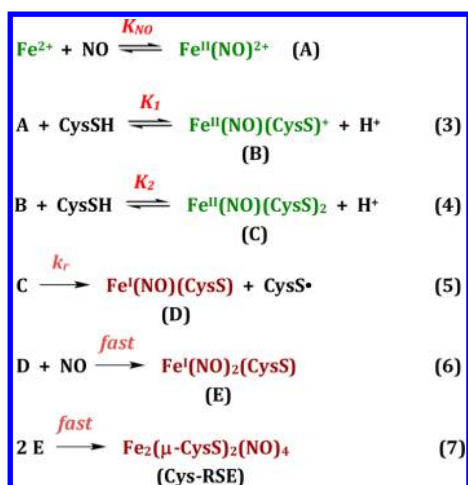
$$\text{rate} = C_1 C_2 [\text{Cys}]_{\text{tot}}^2 [\text{Fe}]_{\text{tot}} [\text{NO}] / (1 + C_2 [\text{NO}]) \quad (2)$$

The sequence of processes represented by Cys-RSE formation from a solution mixture of Fe(II), CysSH and NO must be very intricate, requiring the substitution of several components into the coordination sphere of the iron, the reduction of Fe(II) to Fe(I), and the assembly of two iron/cysteinate/nitrosyl units into the dimeric structure. In this context, the first point to take into consideration is that, given the relative lability of iron(II) complexes,<sup>22,23</sup> it is very unlikely that the slow formation of Cys-RSE at pH 5.0 seen in Figure 3 is rate limited by simple substitution into the coordination sphere of the  $\text{Fe}^{2+}$  ion. This leads us to propose that reduction of Fe(II) is the rate-limiting process. A scenario consistent with the rate law described by eq 2 is outlined in Scheme 1.

In Scheme 1, the rate-limiting step (eq 5) is proposed to be a spontaneous reduction of the Fe(II) center of  $\text{Fe}(\text{NO})(\text{Cys})_2$  (C, the remaining coordination sites being occupied by waters) to give the Fe(I) intermediate D plus a cysteinyl radical  $\text{CysS}^\bullet$ . C would be formed reversibly by reaction of the ferrous center with NO and two cysteines (eq 1 and eqs 3 and 4 in Scheme 1) by relatively fast substitutions for the coordinated waters of the Fe(II) center. The equilibria represented by eqs 3 and 4 are certainly very pH dependent, but the data under consideration here were recorded at a single pH (5.0). Thus, at this relatively low pH, the overall concentration of C would be small, and the overall rate ( $k_r[\text{C}]$ ) would be relatively slow. As seen below, the reactions are much faster at higher pH. Once D is formed, it is likely that it would rapidly capture another NO from the



Scheme 1. Proposed Sequence of Reactions Leading to DNIC Products



solution to give a dinitrosyl species E, which is a likely precursor to either Cys-RSE or Cys-DNIC.

An alternative, essentially kinetically equivalent, rate-limiting step at pH 5.0 would be for B to undergo outer-sphere reduction by a CysS<sup>-</sup> anion to give the same products. However, the stopped flow experiments at higher pH showing a two-stage process, one fast and dependent on [Cys]<sub>tot</sub><sup>2</sup> and pH and the second being much slower and pH independent (see below), would appear to argue against that alternative. In the presence of excess NO, either reduction pathway should lead to formation of the S-nitroso adduct CysSNO, since CysS<sup>•</sup> radical would be trapped by the excess NO. Another essentially kinetically equivalent mechanism at pH 5.0 would be for CysS<sup>-</sup> to undergo nucleophilic attack on the coordinated NO of B to give a Fe<sup>I</sup>(CysS<sup>-</sup>) complex plus CysS-NO. Similar reductive reactions have been noted for Fe(III) and Cu(II) complexes.<sup>24</sup> However, this pathway to Fe(I) intermediates would not be consistent with the kinetics data acquired at higher pH (see below).

The reactions outlined by Scheme 1 would give the rate law:

$$\text{rate} = k_r K_1 K_2 [\text{H}^+]^{-2} K_{\text{NO}} [\text{Cys}]_{\text{tot}}^2 [\text{NO}] [\text{Fe}]_{\text{tot}} / (1 + K_{\text{NO}} [\text{NO}]) \quad (8)$$

This has the same form as eq 2, if  $C_1 = k_r K_1 K_2 [\text{H}^+]^{-2}$  and  $C_2 = K_{\text{NO}}$  and predicts that the overall transformation will be markedly accelerated at higher pH, consistent with the results of the experiments described below. However, it should be noted that this relatively “simple” form of the rate law would be applicable only under the mildly acidic conditions used in the above experiments where  $\text{pH} \ll \text{p}K_a(\text{CysSH})$  (see SI Scheme S-1)

**DFT Studies of eq 5.** Figure 5 illustrates the result of DFT computations of the optimized structures of the proposed species C and D in a dielectric continuum (Scheme 1). All optimizations were performed at unrestricted B3PLYP/DGDZVP level without symmetry constraints using polarizable continuum model (Gaussian 2009) and were starting from “octahedral” geometry with water molecules occupying any coordination sites not taken by NO or the cysteine thiolate anion. (Coordinates for DFT calculated structures of intermediates C and D are listed in SI Table S-1.) Notably, the optimized structure of species C is pentacoordinate

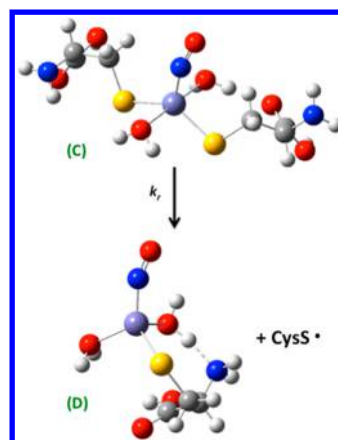
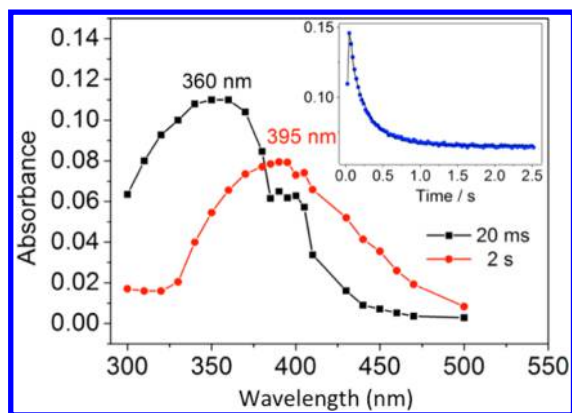


Figure 5. DFT computed structures for the rate-limiting step proposed in Scheme 1.

(pseudo square pyramidal) with a quartet ground state, while the optimized structure of D is tetra-coordinate (pseudo tetrahedral) with a triplet ground state. A number of other configurations (including structures with S<sub>2</sub>O bidentate cysteinate ligands) were also investigated, but those shown in Figure 5 proved to be the most energetically likely. The calculated energy difference between the reactants and products of eq 5 (Scheme 1) in a dielectric continuum was +14 kcal mol<sup>-1</sup>, indicating that this reaction is a thermodynamically reasonable explanation of the kinetics described. Notably, trapping of the CysS<sup>•</sup> by NO is calculated to be exothermic by 37 kcal mol<sup>-1</sup>, thus making the overall process quite favorable.

**Kinetics Studies at pH 10.0: Formation of Cys-DNIC, Fe(NO)<sub>2</sub>(CysS)<sub>2</sub><sup>-</sup>.** When the analogous experiment was performed in pH 10.0 borate buffer (100 mM), the product spectrum displayed a band centered at ~395 nm indicating the formation of Cys-DNIC (Figure 2). The product formation was too fast to monitor by conventional spectroscopy, so the stopped-flow spectroscopic technique was used to follow the temporal absorbance changes at 300–500 nm. When a solution containing FeSO<sub>4</sub> (0.36 mM) and excess CysSH (3.6 mM) in nanopure water was rapidly mixed with an equal volume of a solution containing NO (1.86 mM) in pH 10.0 borate buffer (200 mM), two steps were observed. First, there was a rapid increase of absorption at 360 nm over the course of a few milliseconds indicating the formation of a transient intermediate we designate as “X”, and this was followed by a much slower decrease over a period of a few seconds (Figure 6 inset). The absorbance changes at 360 nm could be fit to a two-stage kinetics model for sequential reactions with exponential functions giving rate constants  $k_{\text{obs}}(\text{a}) = 105 \pm 2 \text{ s}^{-1}$  for generation of the first observable intermediate and  $k_{\text{obs}}(\text{b}) = 4.7 \pm 0.1 \text{ s}^{-1}$  for formation of Cys-DNIC in subsequent steps. These results are summarized in SI Table S-2.

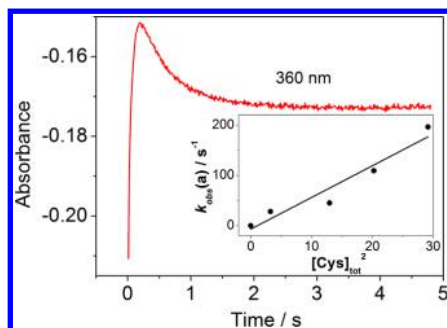
The kinetics data obtained at different observation wavelengths allow one to construct the spectrum of the intermediate(s) formed after the initial reaction. Figure 6 displays the result of using this point-by-point method to obtain the transient spectra at 20 ms and at 2 s from kinetic curves obtained at different wavelengths. The spectrum recorded at 20 ms displays a band centered at 360 nm ascribed to the intermediate “X”. The spectrum recorded at 2 s displays a band centered at 395 nm, which is consistent with the final spectrum (Figure 2), indicating the rapid formation of Cys-DNIC under these conditions. If the initial steps in the reactions of Fe(II),



**Figure 6.** Transient difference spectra recorded at 20 ms (black squares) and at 2 s (red circles) using the point-by-point method with the stopped-flow spectrophotometer when a solution of Fe(II) nanopure water was mixed with a solution of CysSH and NO in pH 10.0 borate buffer,  $T = 298$  K (concentrations after mixing are  $[\text{Fe}]_{\text{tot}} = 0.18$  mM,  $[\text{CysSH}]_{\text{tot}} = 1.8$  mM, and  $[\text{NO}]_{\text{tot}} = 0.93$  mM, 100 mM buffer). Before mixing the solution in the stopped-flow optical path was the solution of Fe(II) in nanopure water. Inset: Typical absorbance changes at 360 nm showing a fast absorbance increase to give an intermediate X, followed by slower reaction to form Cys-DNIC.

CysSH, and NO at both pH 5 and 10 are described by Scheme 1, then a logical identity of X would be as  $\text{Fe}^{\text{II}}(\text{NO})(\text{CysS})_2$ , that is, C.

**Kinetics Studies at pH 7.4.** Similar temporal absorbance changes at 360 nm were also observed at pH 7.4 when an analogous solution of Fe(II) and CysSH was mixed in the stopped flow spectrophotometer with a NO/HEPES buffer solution (Figure 7). Again sequential changes were observed

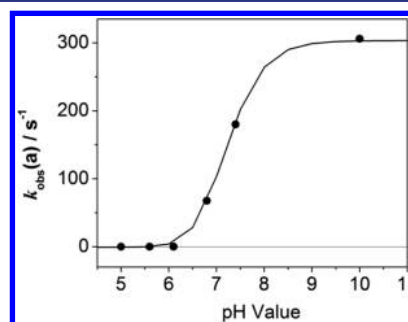


**Figure 7.** Temporal absorbance changes at 360 nm for the reaction of Fe(II) (0.18 mM), CysSH (1.8 mM), and NO (0.93 mM) in pH 7.4 HEPES buffer (100 mM),  $T = 298$  K. Inset: Plot of rate constants  $k_{\text{obs}}(\text{a})$  vs  $[\text{Cys}]_{\text{tot}}^2$  for the reaction of Fe(II) (0.18 mM), CysSH (1.8, 3.6, 4.5, and 5.4 mM), and NO (0.93 mM); slope =  $6.3 (\pm 0.9) \times 10^6 \text{ M}^{-2} \text{ s}^{-1}$ .

with the final spectrum being consistent with the formation of Cys-DNIC. For the same initial concentrations of Fe(II), CysSH, and NO, the first step leading to an intermediate proved to be considerably slower ( $k_{\text{obs}}(\text{a}) = 28 (\pm 1) \text{ s}^{-1}$ ) than at pH 10, while the second step displayed a comparable rate constant ( $k_{\text{obs}}(\text{b}) = 4.8 (\pm 1.5) \text{ s}^{-1}$ ). Plots of  $k_{\text{obs}}(\text{a})$  vs  $[\text{Cys}]_{\text{tot}}^2$  proved to be linear with a slope of  $6.3 (\pm 0.9) \times 10^6 \text{ M}^{-2} \text{ s}^{-1}$  (Figure 7). In contrast,  $k_{\text{obs}}(\text{b})$  appears to be independent of the cysteine concentration. This behavior is consistent with the

$k_{\text{obs}}(\text{a})$  step reflecting formation of the intermediate C, as illustrated in Scheme 1.

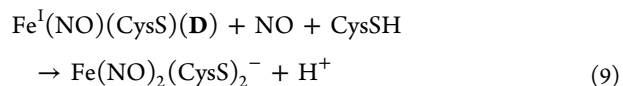
**pH Dependence.** As noted above, whether Cys-DNIC or Cys-RSE is the predominant DNIC formed in Fe/NO/CysSH solutions is dependent both on the solution pH and on  $[\text{Cys}]_{\text{tot}}$ . In order to elucidate the pH effect on the reaction dynamics, we probed the temporal absorbance changes at 360 nm as a function of pH. This experiment was carried out by stopped-flow mixing of a solution of Fe(II) and CysSH in nanopure water with an equal volume of a NO solution in various buffers at pH 5.0, 5.6, 6.1, 6.8, 7.4, and 10.0 under anaerobic conditions at 298 K. After mixing, the initial concentrations were  $[\text{Fe}(\text{II})]_{\text{tot}} = 0.18$  mM,  $[\text{CysSH}]_{\text{tot}} = 5.4$  mM,  $[\text{NO}]_{\text{tot}} = 0.93$  mM, and  $[\text{buffer}] = 100$  mM. Temporal spectral changes similar to those seen in Figure 7 were observed, but as noted above, the rates of the first step proved to be markedly pH dependent. A plot of the  $k_{\text{obs}}(\text{a})$  values against pH for the first spectral change (that we have attributed to formation of C is shown in Figure 8 (SI Table S-3), although



**Figure 8.** Plot of the rate constants  $k_{\text{obs}}(\text{a})$  (●) in  $\text{s}^{-1}$  vs pH values for the reaction of Fe(II) (0.36 mM) and CysSH (10.8 mM) in nanopure water with equal volume of NO solution (1.86 mM) at pH 5.0, 5.6, 6.8, 7.4, and 10.0 buffer (200 mM) at 298 K. Reactant concentrations after mixing were 0.18 mM (Fe(II)), 5.4 mM (CysSH), and 0.93 mM (NO). The line is drawn for illustrative purposes only.

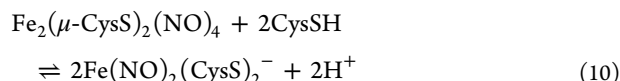
it should be noted that the rate constants given for the lower two pH values are for the overall reaction, since the two stages could not be independently observed. The rates increase markedly as the pH increases, suggesting that the cysteinate anion is the much more reactive form, although the curve drawn through this data suggests a  $\text{pK}_a$  of  $\sim 7.2$ , perhaps indicating enhanced acidity of coordinated CysSH, given that the  $\text{pK}_a$  of CysSH is reported as 8.3.<sup>25</sup>

In contrast, the rate constant  $k_{\text{obs}}(\text{b})$  for the second decay stage, proved to be pH independent (Table S-2). These results are consistent with the first stage of the reaction being the formation of  $\text{Fe}^{\text{II}}(\text{NO})(\text{CysS})_2$  (C). The second-order dependence of  $k_{\text{obs}}(\text{a})$  on  $[\text{CysSH}]_{\text{tot}}$  would imply that eq 4 is rate-limiting in the sequence of steps leading to the intermediate C (Scheme 1). Accordingly, the second stage seen in the stopped-flow experiments would be the spontaneous reduction of C to D followed by rapid reaction of D with CysSH and NO to give Cys-DNIC at higher pH (eq 9).

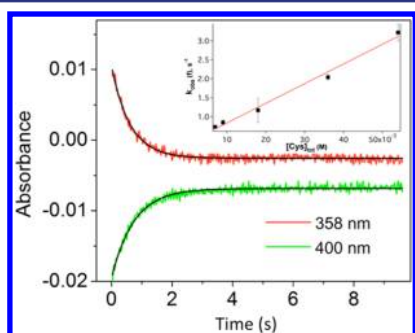


**Interconversion between Cys-RSE and Cys-DNIC at pH 7.4. Cysteine Concentration Dependence.** The above data showed that Cys-RSE is formed in a pH 7.4 HEPES buffer

solution containing Fe(II) (0.18 mM), NO (0.93 mM), and CysSH (1.8 mM). However, at higher [CysSH] (18.0 mM) Cys-DNIC is formed (Figure 2) implying that the two species are readily interconverted (eq 10). Similar interconversions have been reported by Tinberg et al.<sup>26</sup> and by Lu et al.<sup>11a</sup> for the addition of excessive thiolates to RSEs.



The rates of this interconversion were probed by the stopped-flow mixing of a solution containing mostly Cys-RSE in pH 7.4 HEPES buffer ([Cys] = 0.36 mM, [NO] = 1.86 mM) with an equal volume of a pH 7.4 solution containing a large excess of cysteine (36 mM) (final concentrations: [Fe]<sub>tot</sub> = 0.18 mM; [Cys]<sub>tot</sub> = 18.2 mM; [NO]<sub>tot</sub> = 0.93 mM; [buffer] = 100 mM). Under these conditions, the form of the iron complexes shifts almost entirely to Cys-DNIC. The exponential absorbance decrease at 358 and the corresponding increase at 400 nm are shown in Figure 9. Plotting the  $k_{\text{obs}}(f)$  values



**Figure 9.** Temporal absorbance changes at 358 nm (red) and 400 nm (green) for the reaction resulting from the stopped-flow mixing of equal volumes of one solution predominately Cys-RSE in pH 7.4 HEPES buffer solution (100 mM) containing Fe (0.36 mM), CysSH (0.36 mM) and NO (1.86 mM) and a pH 7.4 of CysSH (36.0 mM) at  $T = 298$  K. The solid lines display the fits of these data to single exponential functions to give the  $k_{\text{obs}}(f)$  values  $1.34 \pm 0.03$  and  $1.62 \pm 0.03$   $\text{s}^{-1}$ , respectively. Inset: Plot of rate constants  $k_{\text{obs}}$  ( $\text{s}^{-1}$ ) monitored at 400 nm vs [Cys]<sub>tot</sub> giving the slope  $k_2 = 52 \pm 3.0$   $\text{M}^{-1} \text{s}^{-1}$ .

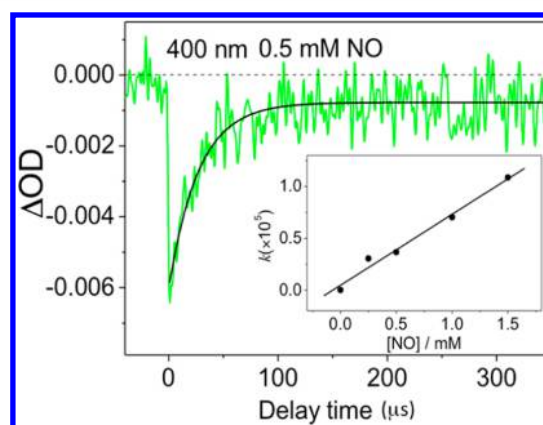
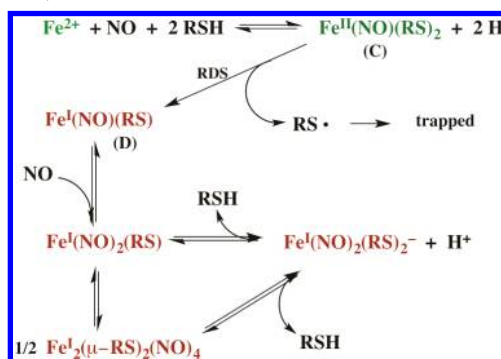
determined in this manner vs [Cys]<sub>tot</sub> for different cysteine concentrations proved to be linear with a slope of  $52 (\pm 3) \text{M}^{-1} \text{s}^{-1}$  and an intercept of  $0.03 (\pm 0.01) \text{s}^{-1}$  (see inset of Figure 9). Thus, the rate-limiting step in the conversion of the dinuclear complex Cys-RSE to the mononuclear Cys-DNIC appears to involve the attack of CysSH (or more likely CysS<sup>-</sup>) on one of the two iron centers in a step leading to one equivalent of Cys-DNIC plus the transient species Fe<sup>I</sup>(CysS)(NO)<sub>2</sub> (E). Rapid reaction of the latter with another CysSH or CysS<sup>-</sup> should give a second Cys-DNIC.

Based on the above kinetics data, we propose the molecular pathways for the formation of Cys-RSE and Cys-DNIC and their interconversion as summarized in Scheme 2.

#### Flash Photolysis Kinetics of Cys-RSE at pH 5.0.

Nanosecond kinetic flash photolysis ( $\lambda_{\text{ex}}$  355 nm) was used to investigate the dynamics of the processes resulting from irradiating solutions of Cys-RSE formed in pH 5.0 citrate buffer by mixing FeSO<sub>4</sub> (0.05 mM), cysteine (2.0 mM), and NO (various concentrations: 0.25, 0.50, 1.00, or 1.50 mM). Figure 10 illustrates one such experiment. At the 400 nm monitoring

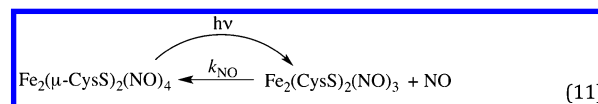
#### Scheme 2. Proposed Scheme for the Formation of Fe(NO)<sub>2</sub>(CysS)<sub>2</sub><sup>-</sup> (Cys-DNIC) and Fe<sub>2</sub>(μ-CysS)<sub>2</sub>(NO)<sub>4</sub> (Cys-RSE) and Their Interconversion



**Figure 10.** Temporal absorbance changes at 400 nm following 355 nm laser photolysis of the Fe/CysSH/NO species (Cys-RSE) formed from a mixture of FeSO<sub>4</sub> (0.05 mM), cysteine (2.0 mM), NO (0.50 mM) in pH 5.0 citrate buffer at 298 K. Inset: Plot of  $k_{\text{obs}}$  vs [NO] for the relaxation of transients formed from the flash photolysis. The slope  $k_{\text{NO}}$  is  $6.9 \times 10^7 \text{M}^{-1} \text{s}^{-1}$ .

wavelength, flash-induced transient bleaching was immediately apparent, and this was followed by relaxation over several hundred  $\mu\text{s}$  back to an absorbance close to that of the original solution. However, the relaxation was only ~91%, complete, and a small fraction of this bleaching does not relax within 1 ms. Absorption spectra recorded before and after the flash experiment (SI Figure S-5) indicated the formation of some long-lived species. Notably control flash photolysis reactions indicated that cysteine itself shows no photochemistry under these conditions or at pH 10.0 (Figure S-6).

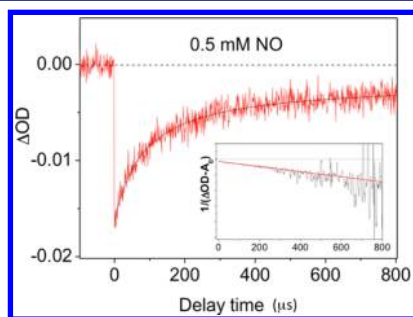
The temporal decay of the transient bleaching upon flash photolysis of Cys-RSE at pH 5.0 could be fit to single exponential functions (Figure 10), to give the calculated  $k_{\text{obs}}$  values for different NO concentrations (Table S-4). A plot of these  $k_{\text{obs}}$  values vs [NO] is linear with an approximately zero intercept (Figure 10 inset). These observations are consistent with photoinduced NO dissociation from Cys-RSE, followed by a back reaction of the resulting intermediate with NO (eq 11). Similar reactions have been observed previously for other





RSEs  $\text{Fe}_2(\mu\text{-RS})_2(\text{NO})_4$  ( $\text{R} = -\text{CH}_2\text{CH}_2\text{OH}$  and  $-\text{CH}_2\text{CH}_2\text{SO}_3^-$ )<sup>16</sup> as well as for Roussin's red salt anion  $\text{Fe}_2(\mu\text{-S})_2(\text{NO})_4$ .<sup>27</sup> The slope of this line ( $6.9 \times 10^7 \text{ s}^{-1} \text{ M}^{-1}$ ) would be the second-order rate constant  $k_{\text{NO}}$  value for this relaxation process. Changing  $[\text{Cys}]_{\text{tot}}$  had little effect on  $k_{\text{obs}}$  and this observation argues against the possibility that the transient bleaching results from dissociation of a cysteinyl moiety.

**Flash Photolysis of Cys-DNIC at pH 10.0.** Similar nanosecond flash photolysis ( $\lambda_{\text{ex}}$  355 nm) was used to investigate the photochemistry of Cys-DNIC formed from mixing  $\text{FeSO}_4$  (0.05 mM) with cysteine (2.0 mM) and NO (0.05, 0.10, 0.25, 0.50, 0.75, 1.00, or 1.25 mM) in pH 10.0 borate buffer. Again flash photolysis leads to an initial bleach followed by the partial relaxation back to the original absorbance (for example, Figure 11). However, the latter

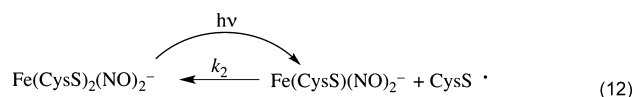


**Figure 11.** Temporal absorbance changes at 400 nm following 355 nm laser photolysis of the DNIC species formed from a mixture of  $\text{FeSO}_4$  (0.05 mM), cysteine (20.0 mM), NO (0.50 mM) in pH 10.0 borate buffer at 298 K. Inset: the linear plot of the  $1/(\Delta\text{OD} - A_0)$  vs time in  $\mu\text{s}$ .

process is slower than that seen at the lower pH under otherwise analogous conditions. Furthermore, the temporal absorbance decay curves could not be fit by a simple exponential, but instead fit well with a second-order reaction function giving rate constant  $k_2 \sim 10^9 \text{ M}^{-1} \text{ s}^{-1}$  ( $\epsilon_{400 \text{ nm}}(\text{Cys-DNIC}) = 6.0 \times 10^3 \text{ L mol}^{-1} \text{ cm}^{-1}$ , Table S-5). Although there was some scatter ( $\pm 20\%$ ) in the  $k_2$  values so determined, a 25-fold increase in  $[\text{NO}]$  from 0.05 mM to 1.25 mM had no effect (SI Figure S-7). Thus, unlike the transient spectral changes at pH 5 attributed to NO dissociation and recombination to regenerate Cys-RSE (eq 10), the transient bleaching seen at pH 10 is not due to NO dissociation. Furthermore, varying  $[\text{Cys}]_{\text{tot}}$  from 2.0 to 20 mM led, not to a linear increase in  $k_2$ , but instead to a modest, systematic decrease in the relaxation rate constant from  $6.6 \times 10^9$  to  $4.4 \times 10^9 \text{ M}^{-1} \text{ s}^{-1}$  (SI Table S-6, Figure S-8). Thus, CysS<sup>-</sup> photodissociation and recombination can also be excluded as being responsible for transient bleaching under these conditions.

One possible scenario to explain these spectral changes would involve the photochemical dissociation not of the cysteinyl anion, but instead a cysteine radical CysS<sup>•</sup> via homolytic cleavage of a Fe–S bond, and the corresponding recombination of the intermediate species (eq 12). The modest decrease in  $k_2$  with increasing  $[\text{Cys}]_{\text{tot}}$  (Table S-6) may be due to the reversible adduct formation between CysS<sup>•</sup> and CysS<sup>-</sup>.<sup>28</sup>

Notably, the relaxation process seen for the flash photolysis studies in pH 10 solution appeared not to be fully reversible during the time frame of the experiment (Figures 11 and S-6). Furthermore, long-term repetitive photolysis with the pulsed



laser (355 nm) led to net change observed as in the solution spectrum, most prominently decreased absorbance over the 340–500 nm range (SI Figure S-9), when the solution was photolyzed with magnetic stirring in a Schlenk photolysis cell.<sup>20</sup> Possible explanations for this irreversibility would be dimerization of the CysS<sup>•</sup> radical to (CysS)<sub>2</sub> or trapping of this species by NO to form the S-nitrosothiol CysSNO. Surprisingly, when this solution was shaken manually, the spectrum of the photolyzed sample nearly returns to that seen before photolysis. We see no obvious explanation for this phenomenon other than the possibility that exposure of the solution to longer wavelength room light decomposed S-nitrosothiols to the RS<sup>•</sup> radical plus NO,<sup>29,30</sup> thereby providing a pathway for restoration of Cys-DNIC.

## SUMMARY

Dinitrosyl iron complexes, especially those involving thiols, have been argued to be the most common form of nitric oxide equivalents in mammalian physiology.<sup>10</sup> DNICs are formed by the reaction of  $\text{Fe}^{2+}$  in the chelatable (labile) iron pool with NO and biological thiols, and these reactions are generally argued to be quite rapid.<sup>7,12</sup> While the majority of these are likely to be protein bound,<sup>31</sup> more mobile forms such as Cys-DNIC and its glutathione analog may play important roles in the transport of this functionality through the cell.<sup>12</sup> Various reactivities have been attributed to DNICs, including a role in the formation of another biological reservoir of NO, the S-nitrosothiols.<sup>7,32</sup> It is clear that there is a marked interplay between all these species in mammalian biology.

The investigations described here have used temporal changes in the optical spectra to outline the dynamics of DNIC formation from solutions of iron(II), the thiol cysteine, and nitric oxide in aqueous solutions of various pH values. As observed previously by Vanin and co-workers<sup>9</sup> the dinuclear Roussin's red ester Cys-RSE is formed at low pH, while the mononuclear complex Cys-DNIC is formed in alkaline media. The nature of the latter species was confirmed by EPR experiments. Notably all these reactions were carried out under deaerated conditions. At the near neutral pH 7.4, Cys-RSE is formed at relatively low total cysteine concentrations, but the transformation to Cys-DNIC occurs readily when additional CysSH is introduced. The kinetics of DNIC formation at 298 K are also strongly pH dependent and demonstrate a second-order dependence on the total cysteine concentration  $[\text{Cys}]_{\text{tot}}$ . The rapid biological formation of DNICs can thus be attributed to the high concentration of thiols in cells (especially glutathione at 0.5–10 mM).<sup>33</sup> Notably, the studies here were at 25 °C, so one might expect larger values of key rate constants such as the  $k_r$  for eq 5 (Scheme 1) at physiological temperatures.

Based on these data we have proposed a mechanism for DNIC formation from components that, regardless of the eventual product under these conditions, proceeds through pH-dependent formation of a common mononuclear intermediate  $\text{Fe}^{\text{II}}(\text{CysS})_2(\text{NO})(\text{H}_2\text{O})_x$ . Spontaneous decomposition of this species, presumably accompanied by formation of the radical CysS<sup>•</sup>, would give the Fe(I) precursors of both Cys-RSE and Cys-DNIC (Scheme 2). That such precursors are quite labile is



illustrated by flash photolysis studies that demonstrated that reaction of the unsaturated dinuclear complex  $\text{Fe}_2(\mu\text{-CysS})_2(\text{NO})_3$  with NO (eq 10) proceeds with a second-order rate constant  $k_{\text{NO}}$  of  $6.9 \times 10^7 \text{ M}^{-1} \text{ s}^{-1}$ . Interestingly, flash photolysis of Cys-DNIC did not labilize NO but instead apparently leads to a reversible photoredox reaction.

Notably, in pH 7.4 solution, the dinuclear DNIC Cys-RSE undergoes a reaction first-order in added CysSH to form Cys-DNIC, further demonstrating the labilities of these systems. By such studies, we have begun the task of mapping the dynamics landscape defining the key transformations interlinking these labile and biologically relevant metal centers.

## EXPERIMENTAL SECTION

**Materials.** L-Cysteine ( $\geq 98.0\%$ ) and iron(II) sulfate heptahydrate ( $>99.0\%$ ) were purchased from Sigma. Nitric oxide gas (99.5%) from the tank (PraxAir) was purified to remove  $\text{NO}_2$  and  $\text{N}_2\text{O}_3$  by passing through a stainless steel column containing Ascarite II (Sigma).<sup>34</sup>

Solutions at pH 5.0 and 5.6 were prepared in citrate buffer (citrate tribasic dehydrate,  $\geq 99.0\%$ ), those at pH 6.8 and 7.4 were prepared in HEPES ( $\geq 99.5\%$ ) buffer, and those at pH 9.0 and 10.0 are prepared in borate buffer (sodium borate decahydrate,  $\geq 99.0\%$ ). Schlenk quartz cuvettes<sup>20</sup> were used to prepare anaerobic solutions for recording the absorption spectra of Cys-RSE and Cys-DNIC and for the kinetics studies on microsecond time scale using laser photolysis. The initial buffered solutions (5 mL) in the cuvette contained the desired concentrations of cysteine and of iron(II) sulfate. These were deoxygenated by the use of the freeze–pump–thaw (3 $\times$ ) method. Measured amounts of NO gas were then added to the cell (either by using a vacuum line and a monometer or using a gastight syringe depending on the experiment) to give the desired NO concentrations calculated according to the partition coefficient between the liquid and gas phases.<sup>21</sup>

**Computations.** The structures of iron cysteine nitrosyl complexes were optimized using unrestricted approach with B3LYP functional and the full-electron, DFT-optimized, double  $\zeta$  + valence polarization basis set equivalent to the DZVP basis set used in DGauss software (DGDZVP), inside the Gaussian 09 package. All calculations were performed without symmetry constraints. The starting geometries around iron were either pseudotetrahedral or pseudo-octahedral in which all coordination sites were occupied by NO, cysteinate ion, and the remainder by water molecules. The lowest energy spin states were determined in vacuum, and then these molecules were optimized again using the polarized continuum model (PCM, solvent = water) also at UB3LYP/DGDZVP theory level.

**Instrumental Methods.** Optical absorbance measurements were recorded using a Shimadzu dual beam UV-2401 PC spectrophotometer in 1.00 cm path-length quartz cells. Stopped-flow kinetics studies for reactions completed on the time frame of ms to s were measured using an Applied Photophysics model SX-18 MV spectrophotometer. Oxygen was removed from the stopped-flow mechanism by vacuum pumping for 1 h. Deaerated solutions were prepared in an inert atmosphere box or by vacuum line Schlenk techniques and transferred to the stopped-flow system using gastight Hamilton syringes with Luer locks. The temperature of the reaction system was controlled with a thermostated, circulating water bath. The results of the stopped-flow kinetics data were evaluated using the SX-18 software.

The laser flash photolysis apparatus and technique have previously been described in detail.<sup>35</sup> Briefly, solution samples in Schlenk quartz cells were photolyzed using the third harmonic (excitation wavelength  $\lambda_{\text{ex}}$  355 nm) output of a Spectra-Physics INDI-HG Nd:YAG pulse laser. The laser excitation energy was  $\sim 10$  mJ/pulse with a pulse duration of  $\sim 10$  ns. The probe beam, perpendicular to the excitation beam, was the output of a 300 W xenon lamp, passed through an IR filter and a SPEX (1681 Spectrometer) monochromator (to select the probe wavelength) prior to the sample combined with a second SPEX (1681 Spectrometer) monochromator (to filter scattered laser pulse

light) prior to a 1P28 photomultiplier tube (PMT). Signals from the PMT were recorded on a LeCroy digital oscilloscope (LT342), and each kinetic trace was an average of 50 traces. The kinetics was analyzed using Matlab and Origin software.

The EPR spectra were recorded using a Bruker EMX Plus EPR spectrometer, operating in the X-band frequencies, with a field modulation of 100 kHz at room temperature. The microwave power and the amplitude modulation were kept at 20 mW and 0.05 or 0.2 mT. During the spectrum recording, sweep time and time constant were 15.36 s and 5.12 ms, respectively. The samples were measured in a 1 mm capillary tube in the absence of oxygen.

## ASSOCIATED CONTENT

### Supporting Information

One scheme, six tables, and nine figures provide additional documentation of the studies described in this manuscript. This material is available free of charge via the Internet at <http://pubs.acs.org/>.

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### Notes

The authors declare no competing financial interest.

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- (18) (a) Vanin and coworkers have assigned the DNIC displaying the EPR  $g_{\text{ave}} = 2.03$  signature as being  $[\text{Fe}^{\text{I}}(\text{NO}^+)_2\text{L}_2]^{n+}$  (or formally  $[\text{Fe}^{\text{III}}(\text{NO})_2\text{L}_2]^{n+}$ ) in character ( $\{\text{Fe}(\text{NO})_2\}^7$  in Enemark/Feltham notation, ref 18b). However, others have demonstrated with crystallographically characterized complexes (refs 13b and 18c) that this  $g_{\text{ave}} = 2.03$  signature defines a  $\{\text{Fe}(\text{NO})_2\}^9$  species, such as the  $[\text{Fe}^{\text{I}}(\text{NO})_2(\text{CysS})_2]^-$  anion that we have assigned as the mononuclear DNIC in this case. Notably, the Roussin's red salt esters are also clearly  $\text{Fe}^{\text{I}}(\text{NO})_2$  in character, although they are diamagnetic owing to coupling between the two iron centers. (b) Enemark, J. H.; Feltham, R. D. *Coord. Chem. Rev.* **1974**, *13*, 339–406. (c) Tsou, C.-C.; Tsai, F. T.; Chen, H.-Y.; Hsu, I.-J.; Liaw, W. F. *Inorg. Chem.* **2013**, *52*, 1631–1639.
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- (20) (a) A Schlenck cuvette has the optical cell fused to a small all-glass apparatus designed for attaching to vacuum lines for preparing deaerated solutions of reactive gases in equilibrium with a gas phase volume of ~20 mL. A diagram of this apparatus is shown in the Supporting Information of ref 17b. (b) Rimmer, R. D.; Richter, H.; Ford, P. C. *Inorg. Chem.* **2010**, *49*, 1180–1185.
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