Reactions of HNO with Heme Proteins: New Routes to HNO—Heme Complexes and Insight into Physiological Effects

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The formation and interconversion of nitrogen oxides has been of interest in numerous contexts for decades. Early studies focused on gas-phase reactions, particularly with regard to industrial and atmospheric environments, and on nitrogen fixation. Additionally, investigation of the coordination chemistry of nitric oxide (NO) with hemoglobin dates back nearly a century. With the discovery in the early 1980s that NO is biosynthesized as a molecular signaling agent, the literature has been focused on the biological effects of nitrogen oxides, but the original concerns remain relevant. For instance, hemoglobin has long been known to react with nitrite, but this reductase activity has recently been considered to be important to produce NO under hypoxic conditions. The association of nitrosyl hydride (HNO; also commonly referred to as nitroxy) with heme proteins can also produce NO by reductive nitrosylation. Furthermore, HNO is considered to be an intermediate in bacterial denitrification, but conclusive identification has been elusive. The authors of this article have approached the bioinorganic chemistry of HNO from different perspectives, which have converged because heme proteins are important biological targets of HNO.

HNO—Heme Adducts

Interest in HNO in the Farmer laboratory originated from efforts to model reductive heme-based catalysis involved in the global nitrogen cycle (Scheme 1). The six-electron reduction of nitrite to ammonia can be driven by a single enzyme, as in assimilatory nitrite reductases, or can occur stepwise via dissimilatory enzymes, which take nitrite to nitric oxide (NO), N₂O, and N₂. Heme nitroxyl intermediates have been postulated in these NO reductions, as indicated by the dotted line in Scheme 1.

A common mechanistic question in enzymatic reduction of nitrogen oxides is whether a nitroxyl intermediate is generated by sequential electron transfer or by a two-electron process such as hydride transfer (Scheme 2). The contention is that a ferrous nitrosyl intermediate (FeII NO) if transiently formed would be stable and difficult to reduce, thus acting as a catalytic dead end. For example, the single-electron reduction of NO–FeIII Mb occurs at ca. −650 mV vs NHE, which is at the edge of the biological reduction range.

An authentic nitroxyl intermediate has been observed during turnover of the fungal NO reductase cytochrome P450nor. In its catalytic cycle, a ferric nitrosyl complex of P450nor is reduced by NADH to generate an intermediate (λmax at 444 nm) that subsequently reacts with NO to give ferric heme and N₂O, which is the nitrogen product of the HNO self-consumption pathway. Ulrich et al. showed that the putative nitroxyl intermediate may be formed from the reaction of NaBH₄ with the ferric nitrosyl adduct, suggesting that nature does indeed bypass the thermodynamically stable FeII NO in forming a reactive nitroxyl intermediate. The electronic structure, basicity, and possible sites of protonation during turnover of this formal [FeII NO] species have been investigated recently.

Corresponding to the free species, Lehner et al. found an energetic

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In a recent series of papers, we have shown that an HNO adduct of ferrous myoglobin (HNO−FeII Mb) can be formed by the reduction of NO−FeII Mb or by the trapping of free HNO by deoxymyoglobin (FeII Mb).15,16 The HNO−FeII Mb adduct is relatively stable, and as a unique model for heme nitroxy intermediates, it has been structurally characterized by 1H NMR,17 resonance Raman, and X-ray absorption spectroscopy/X-ray absorption near-edge structure analysis.18 The 1H NMR spectrum is particularly diagnostic, with characteristic signals for HNO at 14.9 ppm and for the methyl group of Val68 at −2.5 ppm.

In the Farmer laboratory,19 NO−FeII Mb is generally synthesized by the common technique of mixing metmyoglobin (FeII Mb) with nitrite and dithionite (eq 1).20 This reaction also has physiological relevance for the biosynthesis of NO.21 Ulrich’s work9 prompted us to change the reductant from NaNO2 to NaBH4, in the hope of obtaining the HNO−FeII Mb adduct of ferrous myoglobin (FeII Mb).15,16 The HNO−FeII Mb adduct is relatively stable, and as a unique model for heme nitroxy intermediates, it has been structurally characterized by 1H NMR,17 resonance Raman, and X-ray absorption spectroscopy/X-ray absorption near-edge structure analysis.18 The 1H NMR spectrum is particularly diagnostic, with characteristic signals for HNO at 14.9 ppm and for the methyl group of Val68 at −2.5 ppm.

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within a protein pocket will produce two different diastereomers. To examine this hypothesis, a sample of apomyoglobin was reconstituted with the iron complex of 2,4-dimethyldeuteroporphyrin (Scheme 3), and its HNO adduct was prepared by the nitrite/borohydride method. The resulting \(^1\)H NMR spectrum of the low-concentration product solution yields single HNO and valine resonances, consistent with the heme orientation hypothesis. Further 2D NMR studies are underway to better characterize the structural differences of these diastereomeric forms.

To better understand how HNO adducts are formed in the nitrite/borohydride procedure, the three reactants were combined sequentially in several ways (see part S1 in the Supporting Information). HNO−FeII Mb was formed when either met- or deoxymyoglobin was reacted with nitrite and borohydride in any addition order. One plausible reaction of hydride with nitrite is the generation of free HNO, which might then be trapped by FeII Mb (eqs 4 and 5).

\[
\text{HONO} + \text{H}^- \rightarrow \text{HNO} + \text{OH}^- \quad (4)
\]

\[
\text{Fe}^{III}\text{Mb} + \text{HNO} \rightarrow \text{HNO} - \text{Fe}^{III}\text{Mb} \quad (5)
\]

The possible generation of free HNO was examined with the known scavengers nickel(II) tetracyanate (Ni(CN)\(_4\))\(^2-\) and iron(II) N-methyl-N-glucamine-dithiocarbamate (FeII MGD)\(^26\) (eqs 6 and 7). The characteristic 498 nm absorbance band of Ni(CN)\(_3\)NO\(_3\)\(^-\) was observed when Ni(CN)\(_4\))\(^2-\) was reacted with nitrite/borohydride (Figure 2) but was not apparent in the analogous reaction of Ni(CN)\(_4\))\(^2-\) with nitrite/dithionite. Likewise, the reaction of FeII MGD with nitrite/dithionite yielded a product solution whose electron paramagnetic resonance (EPR) spectrum at 77 K matched that of the reported NO−FeII MGD complex (Figure 2). Substitution of dithionite by borohydride produced a spectrum with a similar signal but at less than 5% of the intensity, suggesting that the majority of the product was the diamagnetic NO−FeII MGD species. Both reactions clearly suggest that HNO or an HNO-releasing species is formed in the presence of nitrite/borohydride.

\[
\text{Ni(CN)}_4^{2-} + \text{NaNO}_2/\text{NaBH}_4 \rightarrow \text{NO}^- - \text{Ni(CN)}_3^{3-} \quad (6)
\]

\[
\text{Fe}^{III}\text{MGD} + \text{NaNO}_2 + \text{NaBH}_4 \rightarrow \text{NO}^- - \text{Fe}^{III}\text{MGD} \quad (7)
\]

The versatility of using borohydride to generate HNO adducts in aqueous solutions was investigated with nitroprusside

\[
\text{([Fe(CN)\(_5\))\(^2-\)NO\(^-\) + NaBH}_4 \rightarrow [\text{Fe(CN)\(_5\))\(^2-\)HNO\(^-\)]^2-} \quad (8)
\]

**Acidity/Basicity of HNO−Metal Adducts.** In 1970, the pK\(_a\) of HNO in solution was estimated at 4.7, but more recent work showed the value to be ≈11.5.\(^{27,\text{28}}\) Coordination to a cationic metal ion should lower the pK\(_a\) of HNO, but this has been difficult to characterize in known small-molecule complexes. As mentioned above, Olabe and co-workers reported the generation of an HNO adduct of nitroprusside ([FeII (CN)\(_5\)(HNO)]\(^3-\)) by two sequential reductions of [FeII (CN)\(_5\)]\(^2-\) using dithionite at high pH.\(^{27}\) The adduct is unstable at pH 10; however, lowering the pH to 7 increases the stability such that an HNO resonance can be observed at 20.02 ppm (Figure 3). Analysis of the \(^1\)H NMR signal during pH titration indicated a pK\(_a\) value of 7.7.

For HNO−FeII Mb, a change in the HNO resonance in the \(^1\)H NMR spectrum was not observed from pH 6.5 to 10, suggesting that the pK\(_a\) is well above the range of the well-known acid−alkaline transition for FeII Mb.

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![Figure 2](image-url)
Protein samples at higher pH are unstable over time and unsuitable for NMR studies. However, electronic absorption spectra of dilute samples of HNO–Fe II Mb at high pH do show characterizable changes. As seen in Figure 4, the high-energy band blue shifts at pHs above 11, whereas changes were not evident in the spectra of NO–Fe II Mb or Fe II Mb. This suggests that the pK a of the HNO adduct is above 10 and likely close to 11.

For heme-based oxidoreductases and other metalloproteins, the influence of hydrogen-bonding residues within an active site is often crucial to the mechanism of action. A recent DFT analysis of resonance Raman data on a variety of NO adducts of heme proteins led Xu and Spiro to postulate a differential effect of hydrogen bonding to the nitrogen or oxygen atoms of the coordinated nitrosyl. Whereas hydrogen bonding to the oxygen atom strengthens back-bonding with the metal, hydrogen bonding to the nitrogen atom weakens both the Fe–N and N–O bonds and primes the nitrosyl adduct for reduction to the HNO–Fe II Mb state. Such hydrogen-bonding interactions would likely play an important role in NO x-reducing enzymes, such as several heme-based nitrite reductases and the P450 and binuclear iron nitric oxide reductases.

Evidence for such interactions is observed by analysis of the deuteration exchange in HNO–Fe III Mb, which can be quantified by integration of the HNO peak at 14.9 ppm against that of the methyl group of Val68 at -2.5 ppm in the 1 H NMR spectrum (Figure 5). The rate of H/D exchange of HNO in HNO–Fe III Mb is also quite distinctive. The exchange rate is slow at physiological pH (t 1/2 \sim 5.5 h at pH 8) but increases significantly under more alkaline conditions (t 1/2 \sim 16 or 9 min at pH 9 or 10, respectively). This behavior may be linked to changes in hydrogen-bonding interactions with the distal His64, which hydrogen bonds to the oxygen atom of the HNO adduct.

Previous NMR analysis of HNO–Fe III Mb found few differences between spectra collected at pH 7 or 10; notable exceptions were resonances assigned to N–H protons on the proximal His93, which shifted from 9.32 to 9.68 ppm, and that of the distal His64, which is at 8.11 ppm at pH 8.5 but is not observed at pH 10. Loss of this distal His64 hydrogen-bonding interaction with the HNO adduct would explain the abrupt change in H/D exchange in the same pH region. As illustrated in Scheme 4, the His64 hydrogen-bonding interaction promotes charge buildup on the HNO moiety via back-bonding with the electron-rich Fe II . This may be considered as stabilization of a resonance form with full charge delocalization onto the ligand, i.e., an Fe IV NHOH form for which the nitrogen-bound proton would be more tightly held. A similar doubly protonated form was suggested as an Fe IV NHOH form.

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intermediate in the P450nor cycle.11 Loss of this charge-stabilizing interaction at the oxygen atom may result in a lowering of the pKa of the nitrogen-bound proton and thus an increase in H/D exchange. A reviewer suggests that deprotonation of His64 may open access to the distal pocket, thus facilitating H/D exchange.34

**HNO as an O2 Analogue.** HNO is the simplest analogue of alkynitroso compounds (RNO), which have long been known to bind to ferrous heme proteins.35,36 Mansuy and co-workers were the first to describe the binding of RNO to Mb and Hb,37 as well as to make the analogy of RNO binding to that of O2.38 Although quite rare, a small number of well-characterized organometallic HNO complexes have been identified. Several routes to HNO—metal complexes have been reported, including the direct reduction of a metal nitrosyl, two-electron oxidation of a metal—hydroxylamine adduct,43,44 the insertion of NO into a metal hydride bond,45,46 and the addition of hydride to a metal-bound NO.45–47 Similarly to O2–FeII Mb, all known HNO complexes are low-spin d6 and diamagnetic, and all have a characteristic HNO resonance significantly downfield in the 1H NMR.48

Until this year, only FeIII Mb had been shown to directly complex HNO in solution to form an identifiable HNO complex. Recently,47 other oxygen-binding proteins such as hemoglobin (Hb), leghemoglobin (legHb), and an H3S-binding hemoglobin from the clam L.pectinata were shown to readily trap free HNO in solution to form HNO adducts in good yield as characterized by peaks at ca. 15 ppm in the 1H NMR spectra. Peptidic protons in strong hydrogen bonds may also have downfield resonances, as shown in Figure 6 for the HNO adduct of Hb, which has distinctive downfield resonances at ca. 12–13 ppm because of hydrogen bonds at the α and β subunit interface. Therefore, a key proof is to generate a labeled H15NO adduct because the resulting HNO resonance will be split into a doublet by the 15N nuclear spin.49 15N-labeled samples also allow heteronuclear single quantum coherence (HSQC) spectra to be readily obtained, which provide characterization of the 15N chemical shifts, as demonstrated in Figure 6.

The affinity of the oxygen-binding heme proteins for both HNO and O2 also suggests that HNO might bind and/or inhibit nonheme oxygenases. For instance, HNO precursors inhibit the pigmentation of melanogenic cells,50 which depend on the activity of tyrosinase, an oxygenase that binds O2 between two copper centers.49

**HNO in Mammals.** Interest in HNO in mammalian systems dates to the early 1980s when vasodilation was determined to be actively mediated by an unidentified species50 designated as the endothelium-derived relaxing factor (EDRF).51 The EDRF was subsequently determined to be NO, but the identification process led to comparisons of the effects of NO and HNO donors in vasorelaxant assays. Such experiments were the genesis of the current expanding interest in the pharmacological effects of HNO donors. Consequently, analysis of the aequous chemistry as well as the biochemistry of HNO has arisen in order to identify the chemical origins of the pharmacological effects. The chemical reactions of HNO under physiological conditions and their consequences in mammalian biology are beyond the scope of this review and have been presented recently elsewhere.46,52–54 Other recent research has focused on the identification of mechanisms and markers of HNO biosynthesis and in the production of novel HNO donors with properties tailored for clinical use. Donor compounds are necessary not only for facile or controlled delivery but also because of the self-consumption of HNO via irreversible dehydration of the dimer.55–58

Donors of HNO have also been recently reviewed.59,60

About the same time that the vasoactivity of HNO was observed, Nagasawa and colleagues determined that cyanamide (H2NCN), an alcohol detergent used in Europe, Canada, and Japan for the clinical treatment of chronic
alcoholism, is bioactivated by mitochondrial catalase to produce HNO.\(^{59-61}\)

\[
\text{H}_2\text{NCO} + \text{catalase} \rightarrow \text{HNO} + \text{HCN}
\]

Cyanamide is a potent inhibitor of aldehyde dehydrogenase (AIDH), which catalyzes the conversion of the acetaldehyde generated in the oxidative metabolism of ethanol to acetate. Inhibition of AIDH results in acet-aldehydemia, provoking an unpleasant physiological response and ostensibly leading to alcohol avoidance. The inhibition mechanism involves association of HNO with an active-site thiol.\(^{62}\)

These results provided the first clinical application of an HNO donor and demonstrated that HNO donors could be administered safely and with effect to humans. Consequently, a large series of prodrugs of HNO were developed to elicit this response in vivo (see, for instance, refs 63–68). Additionally, thiols were shown to be major targets of HNO. Subsequently, the interaction of HNO with thiols was shown to lead to reversible (eq 11) and potentially irreversible (eq 12) modifications, depending on the availability of a second thiol to bind to the N-hydroxysulfenamide (RSNHOH) intermediate (eq 10).\(^{69-73}\)

\[
\text{RSH} + \text{HNO} \rightarrow \text{RSNH} \quad \text{(10)}
\]

\[
\text{RSNH} + \text{RSNH} \rightarrow \text{RSSR} + \text{NH}_2\text{OH} \quad \text{(11)}
\]

\[
\text{RSNOH} \rightarrow \text{RS(ОН)NH} \rightarrow \text{RSOOH} \quad \text{(12)}
\]

That protein thiols are able to be modified by HNO donors despite the presence of low-molecular-weight thiols such as glutathione (GSH) has now been shown in a number of systems.\(^{74-78}\) The mechanism by which HNO escapes scavenging by GSH is

\[\text{(eq 10).} 69\]

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\]
not entirely understood but may relate to its hydrophobicity or to the unique properties of protein thiols.

In a somewhat later study, Wink and co-workers demonstrated that an HNO donor (Angeli’s salt) elicited significant cytotoxicity toward lung fibroblasts compared to NO donors.29 This cytotoxicity is dependent upon an aerobic environment, is exacerbated by chemical deple-
tion of cellular GSH, and is induced, in part, by double-
stranded DNA breaks and base oxidation.29 Moreover,
this comparative analysis provided the first indica-
tion that HNO could affect cellular functions by altering
the redox status of the cell in a manner unique from that
of NO. Although HNO is capable of inducing oxidative
stress, it can also act as an antioxidant via facile hydro-
gen-atom donation to oxidizing radical species (akin to
tocopherol) and subsequent generation of NO, which is
an established antioxidant.84 Significantly, the studies
demonstrating the prooxidant effects were performed at
high levels of HNO, whereas the antioxidant properties
were observed at much lower concentrations.

Later collaborative comparisons demonstrated that the
in vitro toxicity of HNO could be replicated in vivo in a
model of ischemia-reperfusion injury in rabbits in con- 
trast to NO, which proved to be protective in the same
model.85 Importantly, a subsequent study showed that
HNO could be protective toward reperfusion injury
if administered prior to the ischemic event.86 Similar
O2-dependent responses to HNO were observed in neu-
ronal channel response.87

Together, these studies led to examination of the reac-
tion of HNO with O2, but the product has yet to be
identified;29–83,88–90 for further discussion of this reac-
tion, see ref 52. Significantly, the autoxidation of HNO
is generally too slow (103 M−1 s−1) to be of kinetic
consequence in many biological systems, particularly at
pharmaceutical levels of HNO.91,92 Furthermore, a num-
er of comparisons of HNO and NO donors have now
appeared in the literature (reviewed in refs 52, 53, 91,
and 93–97) and nearly universally demonstrate that the
physiological properties of HNO and NO are discrete.
Perhaps most importantly, analyses by Wink and col-
leagues led to intensive investigation of the cardiovas-
cular properties of HNO in dogs (reviewed in refs 53
and 93). HNO was found to enhance myocardial con-
tractility even in failing hearts.98,99 As such, HNO donors
may act as a novel class of vasodilators and treatments
for heart failure.100 This discovery substantially increased
interest in HNO and led to an accelerated publication
rate. Ensuing analyses demonstrated that HNO targets
key regulators of normal myocyte contractile function
and increases the sensitivity of myofilaments to calcium in
a thiol-sensitive manner.101–103

During their cardiovascular studies, Paolocci and col-
leagues98 determined that the vascular effects of HNO in
dogs were limited to the venous side of the circulatory
system, unlike NO donors, which are systemic hypotensive
agents. In vascular smooth muscle, NO leads to dilation
by binding to the regulatory ferrous heme of soluble
guanylyl cyclase (sGC), which increases the rate of
conversion of GTP to cGMP. The strong trans effect of
NO induces cleavage of the proximal histidine upon
binding, leading to an activating conformational change
(Scheme 5). The observation that infusion of HNO donors
did not lead to elevated cGMP levels in plasma both indicated
that HNO and NO do not interconvert in blood and renewed
interest in the vasoactive mechanisms of HNO.91

Several HNO donors have been observed to induce
vasorelaxation in vivo or in rodent arterial/aortic ring assays.63,104–106 These results instigated further

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examination of the effects of HNO donors on sGC function, usually by measurement of the cGMP levels or determination of the vasorelaxative potency in the presence of sGC inhibitors. The question of the direct activation of sGC was addressed by Dierks and Burstyn, who exposed partially purified bovine lung sGC to donors of NO, NO\(^+\), and HNO and observed enhanced cGMP formation only upon the introduction of NO. This finding, in addition to earlier examinations of the reaction of HNO with myoglobin and hemoglobin, led to the assumption that HNO does not react with ferrous hemes. Kinetic analysis later suggested that the primary cellular targets for HNO are thiols and oxidized metals, while NO is thought to principally interact with other free radicals and with reduced metals.

To explain the vasoactivity of HNO, the suggestion was made that HNO is converted to NO particularly in the aortic ring assays, for instance, by superoxide dismutase (SOD), metHb, and flavins or by the release of normally sequestered species during tissue preparation. Furthermore, it may be that the adduction of millimolar dithiothreitol (DTT), which is a vital stabilizing agent for sGC because of the oxidative instability of the heme and protein thiols under aerobic conditions, scavenged HNO before it could bind to the heme. This possibility in conjunction with the demonstration by Farmer and co-workers of the thermally stable adduct of HNO with deoxymyoglobin led the Fukuto and Miranda laboratories to further investigate whether HNO can directly enhance the activity of sGC using bovine lung sGC purified in the Burstyn laboratory.

Exposure of sGC to two structurally distinct HNO donors in thiol-free media led to a concentration-dependent increase in cGMP formation. The extent of activation was lower than that from NO but was significantly elevated compared to basal levels. That sGC was not affected by a metal chelator but was modified by DTT clearly indicated that HNO can directly interact with sGC.

Both the heme and the multiple cysteine thiols of sGC are reasonable targets for HNO (Scheme 6). Removal of the heme decreased both HNO- and NO-mediated activity, supporting a direct interaction with the heme for both nitrogen oxides. Surprisingly, HNO did not activate ferric sGC, which was expected to undergo reductive nitrosylation to form the ferrous heme complex (eq 13).

\[ \text{Fe}^{III} + \text{HNO} \rightarrow \text{Fe}^{II}\text{NO} + \text{H}^{+} \]  

Ferric sGC has been previously noted to be substitutionally inert to cyanide. The impact of the reactivity of HNO with the protein thiols was investigated by substitution of the heme with the metal-free porphyrin, which activates the enzyme to a similar extent to the ferrous nitrrosyl complex. In this case, HNO decreased the activity, suggesting that cysteine thiols can function in a negative allosteric fashion when oxidized.

A second recent investigation appeared to confirm that HNO does not affect the sGC activity unless SOD is present. We suggest that, as with other studies that suggest the requirement for oxidation of HNO to NO, buffer components may be lead to unexpected scavenging of HNO. In addition to thiols, a \( N \)-(2-hydroxyethyl)-piperazine-\( N \)\(^{-2}\)-ethanesulfonic acid (HEPES) buffer has been shown previously to scavenge HNO, and related agents such as triethylene (TGA), which was used in the recent activity studies, may have a similar effect.

**Conclusion.** Exposure to HNO is known to affect a variety of thiol-containing proteins. That thermally stable complexes of HNO with heme proteins can now be readily produced suggests the possibility that metalloproteins may also be significant pharmacological targets for HNO. Because investigation of the chemical origin for the differing cardiovascular effects observed for HNO and NO donors in the studies by Paolocci et al. demonstrated that infusion of HNO did not result in a measurable increase in the plasma levels of cGMP, questions remain about the role of HNO in whole organisms compared to in vitro assays or studies using excised tissues, which may have artifactual responses to HNO donors. Furthermore, whether HNO can activate sGC or coordinate to hemoglobin or myoglobin under...
physiological/cellular conditions remains to be determined. The coordination chemistry of HNO should be a fruitful area for both metalloprotein and small-molecule chemistry for the foreseeable future.

Materials and Methods
Horse skeletal muscle myoglobin (95–100%), adult human hemoglobin, sodium nitrite, sodium borohydride, sodium trimethoxyborohydride, and zinc dust were purchased
from Sigma-Aldrich and used as received. Piloty’s acid was purchased from Cayman Chemicals. 4,4′-Dimethyl-2,2′-dipyridyl (DTDP), tetracyanato nickelate,123 and N-methyl-o-glucaminedithiocarbamate26 were prepared and purified following literature procedures. Stock solutions of Piloty’s acid were freshly prepared in deionized water before each experiment. Manipulation of various hemes and their adducts of NO and HNO was performed inside an anaerobic glovebox. Purification of heme adducts was carried out on a preequilibrated Sephadex G25 column in a 50 mM phosphate buffer at either pH 7 or 9.4. All absorption spectra were recorded on a Hewlett-Packard 8453A spectrophotometer. 1H NMR experiments were recorded on a Bruker Avance 600 or Varian 800 MHz spectrometer. The spectra were acquired by direct saturation of the residual water peak during the relaxation delay. Chemical shifts were referenced to the residual water peak at 4.8 ppm. X-band EPR spectra were recorded with a Bruker EMX spectrometer equipped with a standard TE102 (ER 4102ST) or a high-sensitivity ER 4119HS resonator.

UV–Vis Experiments. A sample of HNO–Fe111Mb was concentrated on Centricon YM10. Several microliters were added to 2 mL of the appropriate buffer (100 mM phosphate buffer for pH 7 and 8; 100 mM carbonate buffer for pH 10; the appropriate concentration of NaOH with 50 mM NaCl for pH 11–13). Spectra were collected in a glovebox on a USB2000 spectrophotometer.

H/D Exchange Experiments. For the NMR experiments, HNO–Fe111Mb was prepared using Piloty’s acid, as described above. Aliquots were removed and concentrated on Centricon YM10 using a 100 mM carbonate buffer at pH 10.0, 9.5, and 9.0 and a 100 mM phosphate buffer at pH 8.0. Two dilution concentration cycles were performed, until the pH of the solution in the waste reservoir was at the appropriate pH. The protein solution was concentrated to ~200 μL. To the bottom of a J. Young tube was added 360 μL (60%) of D2O, and to the top compartment was added a total of 240 μL (40%) of a HNO–Fe111Mb solution diluted with a buffer. The solution was mixed, and time-course 1H NMR spectra were collected. The HNO peak was integrated and plotted versus time. For the 95+ % D2O sample, the HNO–Fe111Mb solution was fully exchanged with D2O in a Centricon column.

Reaction of Iron(II) N-Methyl-o-glucaminedithiocarbamate with Nitrite/Dithionite and Nitrite/Borohydride Mixtures. Tetracyanatonickelate (5 mg) was dissolved in 1 mL of a 50 mM carbonate buffer at pH 10 in a 1 cm cuvette. After the addition of sodium nitrite (1 mg), followed by sodium borohydride (2 mg), the solution immediately turned dark purple. The absorbance spectrum was collected with no attempt to isolate the reaction products.

Reduction of Sodium Nitroprusside by Sodium Borohydride. Sodium nitroprusside (10 mg) was dissolved in 1 mL of a 50 mM carbonate buffer. Sodium nitrite and sodium borohydride were then added, and the brick-red-colored solution was then used for studies without further purification.

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Supporting Information Available: Additional experimental details and figures. This material is available free of charge via the Internet at http://pubs.acs.org.