

Oxygenase Domain of *Drosophila melanogaster* Nitric Oxide Synthase: Unique Kinetic Parameters Enable a More Efficient NO Release[†]

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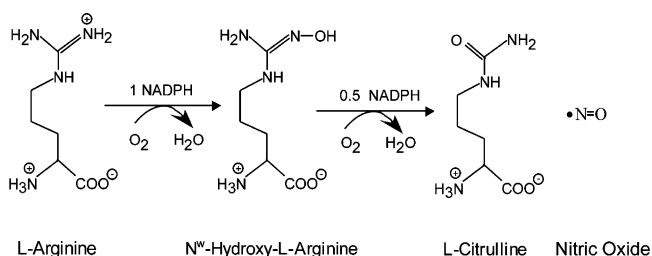
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ABSTRACT: Although nitric oxide (NO) is important for cell signaling and nonspecific immunity in the fruit fly *Drosophila melanogaster*, little is known about its single NO synthase (dNOS). We expressed the oxygenase domain of dNOS (dNOSoxy), characterized its spectroscopic, kinetic, and catalytic properties, and interpreted them in light of a global kinetic model for NO synthesis. Single turnover reactions with ferrous dNOSoxy showed it could convert Arg to *N*^ω-hydroxy-L-arginine (NOHA), or NOHA to citrulline and NO, when it was given 6*R*-tetrahydrobiopterin and O₂. The dNOSoxy catalyzed Arg hydroxylation and NOHA oxidation at rates that matched or exceeded the rates catalyzed by the three mammalian NOSoxy enzymes. Consecutive heme-dioxy, ferric heme–NO, and ferric heme species were observed in the NOHA reaction of dNOSoxy, indicating that its catalytic mechanism is the same as in the mammalian NOS. However, NO dissociation from dNOSoxy was 4 to 9 times faster than that from the mammalian NOS enzymes. In contrast, the dNOSoxy ferrous heme–NO complex was relatively unreactive toward O₂ and in this way was equivalent to the mammalian neuronal NOS. Our data show that dNOSoxy has unique settings for the kinetic parameters that determine its NO synthesis. Computer simulations reveal that these unique settings should enable dNOS to be a more efficient and active NO synthase than the mammalian NOS enzymes, which may allow it to function more broadly in cell signaling and immune functions in the fruit fly.

Nitric oxide synthase enzymes (NOS, EC 1.14.13.39) convert L-arginine to nitric oxide (NO) and citrulline by catalyzing a stepwise, NADPH- and O₂-dependent reaction that forms *N*^ω-hydroxy-L-arginine (NOHA¹) as an intermediate (Scheme 1) (1–3). Three NOS isozymes are expressed in mammals (neuronal or nNOS, inducible or iNOS, and endothelial or eNOS), and they differ with regard to their expression, catalysis, and cellular localization (2, 4–7). All NOSs comprise attached heme-containing (oxygenase) and

Scheme 1



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¹ Abbreviations: CaM, human calmodulin; EPPS, 4-(2-hydroxyethyl)-1-piperazinepropanesulfonic acid; NOS, nitric oxide synthase; nNOS, rat neuronal NOS; NOSoxy, the oxygenase domain of a given NOS; NOHA, *N*^ω-hydroxy-L-arginine.

flavin-containing (reductase) domains that are connected by a central calmodulin (CaM) binding sequence (8–13). Because NOS oxygenase and reductase domains can fold and function independently, they have been overexpressed and subjected to detailed structural, kinetic, and regulatory studies (14–16), along with some NOS oxygenase-like enzymes that are present in bacteria (17–19).

The insect *Drosophila melanogaster* expresses a single NOS enzyme (dNOS) that participates in the developmental, behavioral, and host defense pathways of the fruit fly (20–24). However, besides its generation of NO in a Ca²⁺- and CaM-dependent manner (25, 26), similar to the mammalian eNOS and nNOS enzymes, little is known regarding the catalytic properties of dNOS. A more detailed characterization stands to broaden our understanding of NOS structure–function relationships and could help to clarify NOS functions in the fruit fly. To address this, we characterized

the catalytic and kinetic properties of the dNOS oxygenase domain (dNOSoxy) after its overexpression in *E. coli*. Our results show that dNOSoxy is most similar to the mammalian nNOS but has kinetic features that distinguish it from the mammalian NOS enzymes. These findings, along with results in the accompanying article on the reductase domain of dNOS, suggest that dNOS evolved to support a more efficient and active NO release than do the mammalian enzymes.

EXPERIMENTAL PROCEDURES

Materials. All reagents and materials were obtained from Sigma, Alexis Biochemicals, and Amersham Biosciences, unless noted otherwise.

dNOSoxy Construct Generation. The pCWori vector containing the 5' hexa-His tagged dNOS full-length construct (4050 bp) within NdeI and HindIII restriction sites, as described in (26), was used to generate a 5' hexa-His tagged construct of the dNOS oxygenase domain and the adjoining CaM binding sequence (dNOSoxy). This construct ends at Ser683. To create the stop codon after Ser683, the following primers were used to amplify a fragment of dNOS: 5'-ACG-GTGGCGCTGAAGATGCAACTGG-3' and 5'-GATAAGCTTT-TACGATTTGCCAGTTTCGGTGGC-3'. The second primer contains the stop codon in bold followed by the underlined HindIII restriction site. This PCR product was subcloned into a PCR-4 vector (Invitrogen) and was digested with BsiWI (located downstream from the forward primer) and HindIII (both are unique sites) and cloned into the pCWori plasmid containing dNOS after it had been digested with the same enzymes. Positive clones of dNOSoxy were identified by restriction analysis.

Expression and Purification of dNOSoxy. The dNOSoxy protein was expressed in recombinant *E. coli* Rosetta cells using the pCWori plasmid as done with the mammalian NOSoxy proteins (34). Transformed bacteria were grown at 37 °C in 3 L of terrific broth supplemented with 100 mg/L ampicillin. Protein expression was induced when the cultures reached an OD₆₀₀ of 0.8 to 1 by adding 1 mM isopropyl- β -D-thiogalactoside. After further growth at room temperature for 24 h, the cells were harvested and resuspended in buffer A (40 mM EPPS, 150 mM NaCl, and 10% glycerol) containing 1 mM L-Arg, 10 μ M H₄B, 0.5 mg/mL each of leupeptin and pepstatin, 1 mg/mL lysozyme, and 1 mM phenylmethylsulfonyl fluoride. Cells were lysed by freeze-thawing three times in liquid nitrogen followed by sonication for five 45-s pulses with a 1-min rest on ice between pulses. The cell lysate was centrifuged at 4 °C for 30 min, and the cell-free supernatant was precipitated by adding 50% (w/v) ammonium sulfate. The precipitant was centrifuged at 4 °C for 30 min at 16,000 rpm, and was then resuspended in buffer A containing 5 mM β -mercaptoethanol, 1 mM L-Arg, and 10 μ M H₄B. The resuspended solution was loaded onto a Ni-NTA-Sepharose CL-4B column that had been charged with 50 mM NiSO₄ and equilibrated with buffer A containing 5 mM β -mercaptoethanol, 1 mM L-Arg, and 10 μ M H₄B. The column was washed with 5 volumes of equilibration buffer and 5 volumes with equilibration buffer containing 10 mM imidazole. The dNOSoxy protein was eluted with 100 mM imidazole in buffer A. The eluted protein sample was dialyzed against buffer B (40 mM EPPS and 10% glycerol) and then run onto a Q-Sepharose column equili-

brated with buffer B containing 1 mM L-Arg and 10 μ M H₄B. The bound protein was washed extensively with the equilibration buffer containing increasing concentrations of NaCl, and the bound protein was eluted with Buffer B containing 200 mM NaCl plus L-Arg and H₄B. The resulting protein sample was concentrated, dialyzed to remove the imidazole, and finally stored frozen in aliquots at -80 °C.

UV-visible spectra of diluted protein samples were recorded on an Hitachi U3110 spectrophotometer in the absence or presence of 20 μ M H₄B and 1 mM Arg. Enzyme concentration was quantified using the absorption of the ferrous-CO adduct at 444 nm and an extinction coefficient of 74 mM⁻¹cm⁻¹ ($A_{444} - A_{500}$).

Preparation of Ferrous dNOSoxy. Concentrated dNOSoxy was placed in a cuvette and made anaerobic by several cycles of evacuation and purging with deoxygenated N₂. An anaerobic solution that contained 50 mM EPPS at pH 7.5, H₄B, and L-Arg or NOHA was then added. The enzyme was reduced by the sequential addition of a dithionite solution from which the concentration was standardized against ferricyanide. The heme reduction was monitored spectrophotometrically. The reduced enzyme solution was transferred into a syringe in a stopped-flow instrument using a gastight syringe.

Stopped-Flow Single Catalytic Turnover Reactions. Rapid-Scanning stopped-flow experiments were performed at 10 °C using a HI-TECH SF-61 instrument equipped with a HI-TECH MG-6000 rapid-scanning diode-array detector, as described previously (30, 34, 36). An anaerobic solution containing 50 mM EPPS (pH 7.5), 10 μ M ferrous dNOSoxy, 10 μ M H₄B, and either 1 mM L-Arg or 0.5 mM NOHA was rapidly mixed with an equal volume of air-saturated buffer (50 mM EPPS at pH 7.5). Ninety-six scans from 350 to 700 nm were collected within 0.0018 to 2 s after each mixing. Sequential spectral data were fit to different reaction models using the Specfit global analysis program (provided by Hi-Tech Ltd.), which can calculate the number of different enzyme species, their spectra, and their concentrations versus time during the single turnover reactions. Data from six to eight sequential reactions were averaged to obtain the final traces.

Product Stoichiometry Analyses. Amino acids in aliquots taken from single turnover reactions were derivatized with *o*-phthalaldehyde and then analyzed by reverse-phase HPLC with fluorescence detection (27). Samples were filtered through an Amicon Centricon device (10,000 MW cutoff) prior to derivatization. Samples were injected onto a Hewlett-Packard ODS-Hypersil column that was equilibrated with 5 mM ammonium acetate at pH 6.0 and 20% methanol and then subjected to an increasing gradient of methanol. Retention times and concentrations of amino acids were calculated on the basis of NOHA and citrulline standard solutions.

NO Binding Kinetics of Ferric NOSoxy Enzymes. The experiments were performed as previously described (51). An anaerobic solution containing 2 μ M of either dNOSoxy or nNOSoxy, 1 mM Arg, 50 μ M H₄B, 0.5 mM EDTA in 40 mM EPPS at pH 7.6 was rapidly mixed at 10 °C in the stopped-flow instrument with a buffered solution containing different concentrations of NO. The NO solutions were made by diluting a buffered, NO-saturated solution in anaerobic buffer and assuming a NO concentration of 1.9 mM for a

saturated solution at 25 °C. Six to eight sequential reactions were run at each NO concentration and then averaged to obtain the final traces. Data were fit to a single or double exponential equation.

Ferrous Heme-NO Oxidation Rate. A solution of 8 μM ferric dNOSoxy in a 40 mM EPPS buffer (pH 7.6), containing 1 mM Arg, 50 μM H₄B, and 0.5 mM EDTA, was made anaerobic in a sealed cuvette and reduced by adding a small amount of anaerobic sodium dithionite solution at 10 °C. To minimize adding excess dithionite, titration of the ferric enzyme was stopped as soon as the heme became reduced, as judged by a shift in the Soret absorbance and the disappearance of absorbance at 650 nm. NO was subsequently added from a NO-saturated solution. The ferrous heme-NO enzyme was transferred to the stopped-flow instrument and mixed at 10 °C with buffer containing various O₂ concentrations and 40 mM EPPS (pH 7.6), 1 mM Arg, 50 μM H₄B, and 0.5 mM EDTA. The kinetics of ferrous heme-NO oxidation was determined from the absorbance loss at 438 nm (ferrous heme-NO decay) and buildup at 396 nm (ferric heme buildup). Traces at both wavelengths were averaged from 6 to 10 individual reactions and were fit to a single-exponential function using software provided by the instrument manufacturer.

Computer Simulation of the Global Kinetic Model for NOS Catalysis. Simulations of the model in Figure 6 were done using kinetic values determined here or previously at 10 °C and by using Mathcad software as previously described in detail (31, 44, 50). In the current simulations, the following parameter values were utilized: for dNOS, $k_{\text{cat}1}$ 28 s⁻¹, $k_{\text{cat}2}$ 48 s⁻¹, k_d 18 s⁻¹, and k_{ox} 0.2 s⁻¹; for nNOS (38), $k_{\text{cat}1}$ 18 s⁻¹, $k_{\text{cat}2}$ 29 s⁻¹, k_d 5 s⁻¹, and k_{ox} 0.2 s⁻¹. The heme reduction rate (k_r) was varied as described in the text, and we assumed a constant concentration of O₂ (140 μM) and NADPH (40 μM) and no buildup of NO in solution.

RESULTS AND DISCUSSION

Expression, Purification, and Spectral Properties. The oxygenase domain of the *Drosophila melanogaster* NO synthase (dNOSoxy) was overexpressed in *E. coli* BL21 Rosetta cells. The dNOSoxy protein had a His₆ tag at its N-terminus to enable purification by nickel-nitriloacetic acid sepharose column chromatography. The average yield of purified dNOSoxy was 3 to 4 mg/L of culture, and the final protein was approximately 90% homogeneous as judged by the SDS-PAGE analysis (data not shown). The apparent molecular mass of the dNOSoxy was 76 kDa, which is in good agreement with the predicted size of the polypeptide based on the gene sequence.

Purification of dNOSoxy was done in the presence of L-Arg and H₄B. The UV-visible spectrum of dNOSoxy showed a Soret peak at 396 nm (Figure 1), typical of mammalian NOS enzymes containing bound Arg and H₄B (27, 28) and of five-coordinate, high-spin heme-thiolate proteins in general. Reduction of dNOSoxy with dithionite caused the Soret band to shift to 414 nm (not shown), and subsequent addition of CO shifted the Soret band to 445 nm (Figure 1), confirming an intact heme-thiolate bond in dNOSoxy.

Single Catalytic Turnover Reactions of dNOSoxy. We utilized rapid-scanning stopped flow spectroscopy to docu-

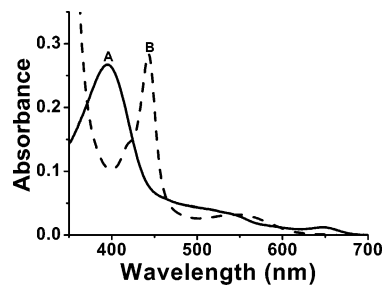


FIGURE 1: UV-visible spectra of purified dNOSoxy. Scan A is ferric dNOSoxy in the presence of L-Arg and H₄B, while scan B is the ferrous heme-CO complex.

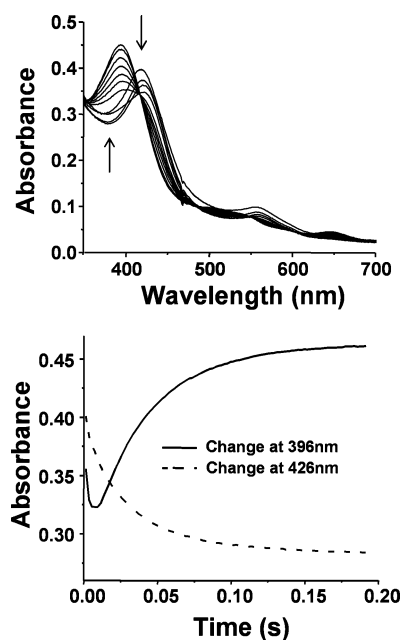


FIGURE 2: Kinetics of heme transitions during Arg oxidation by dNOSoxy under single turnover conditions. Anaerobic ferrous dNOSoxy containing Arg and H₄B was mixed at 10 °C with air-saturated buffer in the stopped flow instrument to start the reaction. The upper panel contains spectral traces collected during the reaction and show the direction of absorbance change. The lower panel shows the absorbance change over time at two wavelengths. Data are representative of four independent experiments.

ment heme transitions during the Arg hydroxylation and NOHA oxidation reactions catalyzed by dNOSoxy. This approach has been quite valuable for understanding the reaction mechanism and kinetics of the mammalian NOS enzymes (29–31). Dithionite-reduced (ferrous) dNOSoxy that contained H₄B and either Arg or NOHA was rapidly mixed at 10 °C with an O₂-saturated buffer to initiate the reaction.

For the Arg single turnover reaction, prior work with mammalian NOSoxy enzymes showed that a heme-dioxy intermediate is formed upon mixing reduced enzyme with O₂-containing buffer, and then this species converts to ferric NOSoxy concurrent with Arg hydroxylation to form NOHA (30–32). However, in the Arg hydroxylation reaction of dNOSoxy, we observed no clear buildup of a heme-dioxy intermediate prior to the formation of ferric dNOSoxy (Figure 2). Global analysis of the spectral data also failed to indicate a clear buildup of a heme-dioxy intermediate in the reaction. This contrasts with the three mammalian NOSoxy enzymes, which typically build up a detectable heme-dioxy intermediate with characteristic heme Soret absorbance near 427 nm

during the Arg hydroxylation reaction (29–32). Despite this, the shifting isosbestic points in Figure 2A and the biphasic absorbance change at 396 nm in Figure 2B do suggest the formation of an intermediate in the reaction. Because the maximally shifted spectrum is centered at 420 nm in Figure 2A, it is possible that the dNOSoxy heme-dioxy intermediate formed in the Arg reaction is the 420 nm species that is observed for mammalian NOS enzymes under some conditions (52). In any case, we still could estimate the rate of Arg hydroxylation catalyzed by dNOSoxy from the concurrent loss of absorbance at 426 nm and the gain of absorbance at 396 nm (equivalent to the rate of ferric dNOSoxy formation) (Figure 2). The rates estimated at the two wavelengths were similar (29 s^{-1} and 26 s^{-1} , respectively). These rates are in the high range of rates obtained for the heme-dioxy to ferric heme transition that is associated with Arg hydroxylation by the mammalian NOSoxy enzymes, which range from 6 to 24 s^{-1} under identical reaction conditions (37). Thus, dNOSoxy appears to catalyze Arg hydroxylation as fast or faster than most mammalian NOS enzymes and is most similar to nNOS in this respect.

Amino acid product analysis confirmed that dNOSoxy generated NOHA from Arg in the single turnover reactions (0.3 NOHA per heme). This product yield is within the range reported for Arg hydroxylation by mammalian NOSoxy enzymes under similar reaction conditions (0.2 to 0.8 NOHA per heme) (32, 34, 35).

For the NOHA single turnover reaction of dNOSoxy, global analysis of the rapid scanning data indicated four consecutive heme species, which, on the basis of previous work with mammalian NOSoxy enzymes (36, 37), could be identified as the starting ferrous dNOSoxy, a transient heme-dioxy intermediate, a ferric heme–NO intermediate, and the ferric dNOSoxy product. Thus, three observable transitions occurred during the NOHA oxidation reaction, where k_1 , k_2 , and k_3 are the calculated rates for the transitions:

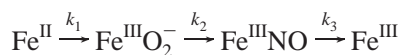


Figure 3 illustrates the UV–visible spectrum of the four enzyme species and the time course of their consecutive appearance and/or disappearance. The Soret absorbance peaks for the dNOSoxy Fe^{II} , $\text{Fe}^{\text{III}}\text{O}_2^-$, $\text{Fe}^{\text{III}}\text{NO}$, and Fe^{III} heme species were 415, 424, 438, and 396 nm, respectively, which closely match the absorbance positions of these same species in mammalian NOSoxy enzymes when they catalyze NOHA oxidation (34, 36). Why a 428 nm ferrous-dioxy species was clearly observable in the NOHA reaction but not in the Arg reaction is puzzling and deserves further investigation. Our NOHA single turnover data imply that dNOS has a similar if not identical mechanism of catalysis compared to the mammalian enzymes. Amino acid product analysis confirmed that dNOSoxy generated citrulline from NOHA in the single turnover reactions (0.4 NOHA per heme), consistent with our observation of a buildup of a ferric heme–NO product complex during the reaction.

The calculated rates for the heme transitions observed during NOHA oxidation by dNOSoxy are listed in Table 1, along with values obtained for NOHA reactions catalyzed by the three mammalian NOSoxy enzymes and by one bacterial NOSoxy protein under identical or nearly identical

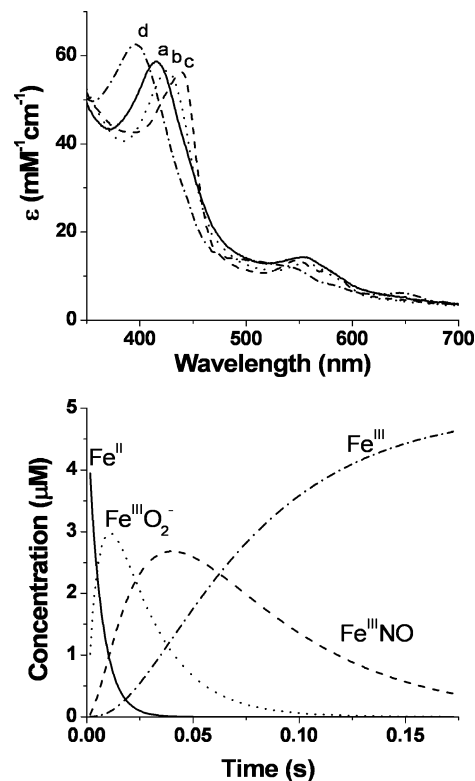


FIGURE 3: UV–visible spectra of heme species detected and their transition kinetics during NOHA oxidation by dNOSoxy under single turnover conditions. Anaerobic ferrous dNOSoxy containing NOHA and H_4B was mixed at 10°C with air-saturated buffer in the stopped flow instrument to start the reaction. The upper panel contains spectra of four species derived from global analysis of the data. These in order of appearance are the ferrous (a, solid line), heme-dioxy (b, dotted line), ferric heme–NO (c, dashed line), and ferric (d, dash-dotted line) forms of dNOSoxy. The lower panel shows the kinetics of disappearance or formation of each spectral species. Line patterns for each species are the same as those in the upper panel. Data are an average of 6 to 10 reactions and are representative of four independent experiments.

reaction conditions (17, 34, 36, 38). The comparison shows that dNOSoxy has a rate of product formation as measured by the k_2 transition (also designated k_{cat}) that is as fast or faster than that in the mammalian and bacterial NOS enzymes (30, 36). The greatest difference, however, appears to be in the k_3 transition, which is the macroscopic dissociation rate (k_d) of the ferric heme–NO product complex that forms at the end of each catalytic cycle in all NOS enzymes examined to date (31). This k_d is important because it helps to determine what percentage of the NO synthesized by a NOS enzyme is actually released into solution (39). Remarkably, the k_d values derived from our NOHA single turnover experiments indicate that NO dissociation from ferric dNOSoxy is 3 to 8 times faster than NO dissociation in the mammalian NOS enzymes and is 90 times faster than that in the bacterial NOS (Table 1).

To confirm whether dNOSoxy has a faster NO dissociation than the mammalian enzymes, we utilized a different stopped flow method (51) to compare the NO binding kinetics of ferric dNOSoxy and nNOSoxy. Each enzyme was mixed under pseudo-first-order conditions with solutions that contained different concentrations of NO, and the observed NO binding rates were plotted versus the initial NO concentration (Figure 4). A slight deviation from a single-exponential fit was observed (particularly for the dNOSoxy

Table 1: Observed Rates of Heme Transitions during NOHA Single Turnover Reactions Catalyzed by Various NOSoxy Enzymes^a

| transformations | dNOSoxy ^b | iNOSoxy ^c | nNOSoxy ^d | eNOSoxy ^e | bsNOS ^f |
|--|----------------------|----------------------|----------------------|----------------------|--------------------|
| heme-dioxy formation (k_1) ^g | 157 ± 15 | 47 ± 4 | 129 ± 5 | NA | 29 ± 1 |
| heme-dioxy disappearance or Fe ^{III} -NO formation (k_2, k_{cat}) | 48 ± 3 | 37 ± 1 | 29 ± 1 | 18 | 7 ± 1 |
| Fe ^{III} -NO disappearance or ferric heme formation (k_3, k_d) | 18 ± 2 | 2.3 ± 0.1 | 5 ± 1 | 3 | 0.2 ± 0.1 |

^a The rates were determined at 10 °C and are the mean ± SD. ^b This work. ^c Wei et al., ref 36. ^d Panda et al., ref 38. ^e Unpublished observations. ^f Adak et al., ref 17. ^g Rates were recorded in a half O₂-saturated reaction except for iNOSoxy and bsNOSoxy, which were recorded in a half air-saturated reaction. NA, not available.

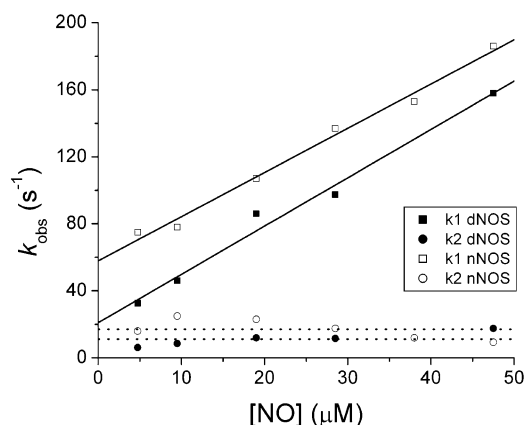


FIGURE 4: Kinetics of NO binding to dNOSoxy and nNOSoxy. An anaerobic solution of either ferric enzyme was mixed with NO containing solution in a stopped-flow spectrophotometer at 10 °C and the ferric heme-NO complex formation was monitored at 438 nm. The absorbance traces were fit to a biexponential equation, and the observed rates are plotted vs the NO concentration with the least-squares lines of best fit as indicated. The data are representative of two independent experiments.

reactions), and therefore, for the sake of comparison both the nNOSoxy and dNOSoxy data were fit to a biexponential equation. The slower phase (k_2) contributed 30% or less of the total absorbance change in all cases and appeared to be independent of NO concentration (Figure 4), whereas the fast phase (k_1) was NO concentration-dependent. The y axis intercepts of the k_1 rate data provide an estimate of the NO k_d for each ferric enzyme and are 21.0 ± 5.5 and $57.9 \pm 3.8 \text{ s}^{-1}$ for dNOSoxy and nNOSoxy, respectively. These data are consistent with our NOHA single turnover experiments and confirm that ferric dNOSoxy has a faster NO k_d than the mammalian NOS enzymes.

In bacterial NOSoxy, we know that a Ile/Val substitution near the entrance to the heme pocket is partly responsible for its slow NO dissociation from the ferric heme (39). However, a protein sequence comparison (Figure S1, Supporting Information) reveals no obvious differences in the residues near the heme pockets of dNOSoxy and mammalian NOSoxy enzymes. This implies that the general dimensions and composition of the dNOSoxy heme pocket are similar to those in the mammalian NOSoxy enzymes (28, 39–41). Thus, faster NO dissociation from the dNOSoxy ferric heme could be due to other more global effects.

A second kinetic parameter that helps determine NOS catalytic behavior is the oxidation rate of the enzyme ferrous heme-NO complex. This complex forms whenever the ferric heme-NO product complex becomes reduced during steady-state turnover in the full-length NOS enzymes (42–44). We studied this reaction in dNOSoxy by mixing the ferrous

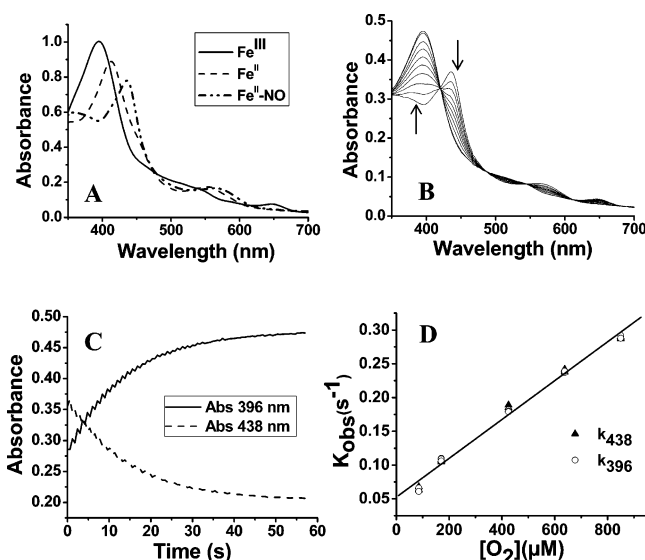


FIGURE 5: Oxidation of the dNOSoxy ferrous heme-NO complex. Panel A shows the UV-visible spectra of the ferric, ferrous, and ferrous-NO forms of dNOSoxy. Panel B contains spectral traces collected within 58 s after mixing the dNOSoxy ferrous heme-NO complex with air-saturated buffer in the stopped flow instrument at 10 °C. Arrows indicate the direction of absorbance change with time. Panel C shows the absorbance changes at 438 and 496 nm vs time during the reaction of the dNOSoxy ferrous heme-NO complex with air-saturated buffer. Panel D plots the k_{obs} values calculated from the absorbance changes at either wavelength for reactions that were performed at the different indicated concentrations of O₂. The least-squares line of best fit is indicated. Data are the average of 6 to 10 individual reactions and are representative of three similar experiments.

heme-NO complex with oxygenated buffer at 10 °C in a stopped-flow spectrometer equipped with a diode-array detector. Figure 5, panel A shows the spectrum of authentic ferric, ferrous, and ferrous-NO forms of dNOSoxy. Panel B contains a group of consecutive spectral scans collected during a typical reaction. The several isobestic points indicate that a monophasic transition occurs between the starting ferrous heme-NO complex of dNOSoxy and the ending ferric enzyme, as is also reported for the reaction catalyzed by mammalian NOSoxy enzymes (38, 44). The rates of heme-NO complex disappearance and ferric enzyme formation were determined from the absorbance changes at 438 nm and at 396 nm, respectively, and were best fit as monophasic transitions (Figure 5C). At the half air-saturated condition used here (O₂ concentration ~140 μM), the oxidation rate of the dNOSoxy ferrous heme-NO complex (designated k_{ox}) was 0.093 s^{-1} at 10 °C. This rate is similar to that of the ferrous heme-NO complex of nNOSoxy at the same O₂ concentration and temperature (0.1 to 0.2 s⁻¹) but is 6 and 13 times slower than the k_{ox} rates measured for

the ferrous heme–NO complexes of eNOSoxy and iNOSoxy, respectively (44, 45).

We went on to measure k_{ox} rates of dNOSoxy at various O_2 concentrations. The rates we obtained are plotted in Figure 5D. The slope gives a second-order rate constant for the reaction of $290 M^{-1} s^{-1}$. This value is quite similar to that obtained for mammalian nNOSoxy, whose k_{ox} second-order rate constant is $220 M^{-1} s^{-1}$ at $10^\circ C$ (unpublished data). We conclude the ferrous heme–NO complex of dNOSr is relatively un-reactive toward O_2 and in this way is similar to nNOSoxy.

Implications for the NO Synthesis Activity of dNOS. Our results provide two (k_d , k_{ox}) of the three kinetic parameters that allow one to model and understand the catalytic behavior of any given NOS (31, 44). The third variable is the heme reduction rate (k_r), which cannot be obtained using the isolated oxygenase or reductase domains of dNOS. However, even without a measured k_r value, we can still make reasonable predictions about the catalytic behavior of dNOS. Our data show that dNOS has a relatively fast k_{cat} and a slow k_{ox} like nNOS. It also has a reductase domain whose electron-transfer properties and regulation (see the accompanying article) are similar to those of the nNOS reductase domain (46, 47, 48). However, there is an important difference regarding the k_d values of ferric dNOS and nNOS that will distinguish their catalytic behaviors. For example, in nNOS, the rates of heme reduction (k_r , 3 to $4 s^{-1}$) and NO dissociation (k_d , $5 s^{-1}$) are similar (34, 43). This causes its ferric heme–NO product complex to partition in about a 2:3 ratio between futile and productive cycles during catalysis (Figure 6A). Because nNOS has a relatively slow k_{ox} ($0.2 s^{-1}$), a significant proportion of enzyme molecules accumulate as the ferrous heme–NO complex during steady-state NO synthesis (31, 44). This in turn compromises the observed NO synthesis activity of nNOS and greatly increases its apparent K_mO_2 , enabling NOS to have a very gradual and near linear oxygen–NO synthesis activity response across the entire physiologic O_2 concentration range (31, 49, 50). In comparison, dNOS has a 5-times faster NO dissociation (k_d is $18 s^{-1}$) that will predispose its ferric heme–NO product complex to release NO rather than become reduced to the ferrous heme–NO complex (Figure 6A). This means that at the heme reduction rate measured for nNOS (approximately $4 s^{-1}$), the k_d of dNOS would allow four enzyme molecules to release NO for every one molecule that enters the futile cycle and therefore should enable dNOS to have a greater NO synthesis activity than nNOS.

How the different k_d values will impact the catalytic behavior of dNOS and nNOS can be determined by computer simulation of the global kinetic model in Figure 6A (44, 50) using the kinetic values we have measured for dNOSoxy and nNOS (44, 50). We performed such simulations using the measured k_d values of 18 and $5 s^{-1}$ for dNOS and nNOS, respectively, a common k_{ox} value of $0.2 s^{-1}$, and heme reduction rates (k_r) ranging from 0.3 to $12 s^{-1}$, which cover and exceed the known k_r values for NOS enzymes. Our simulations indicate that a smaller proportion of dNOS would exist as a ferrous heme–NO complex during the steady state compared to that of nNOS (Figure 6B). Moreover, dNOS would be up to 3.5 times more active than nNOS over the range of heme reduction rates (Figure 6C). Thus, a unique blend of kinetic parameters is likely to enable dNOS to be

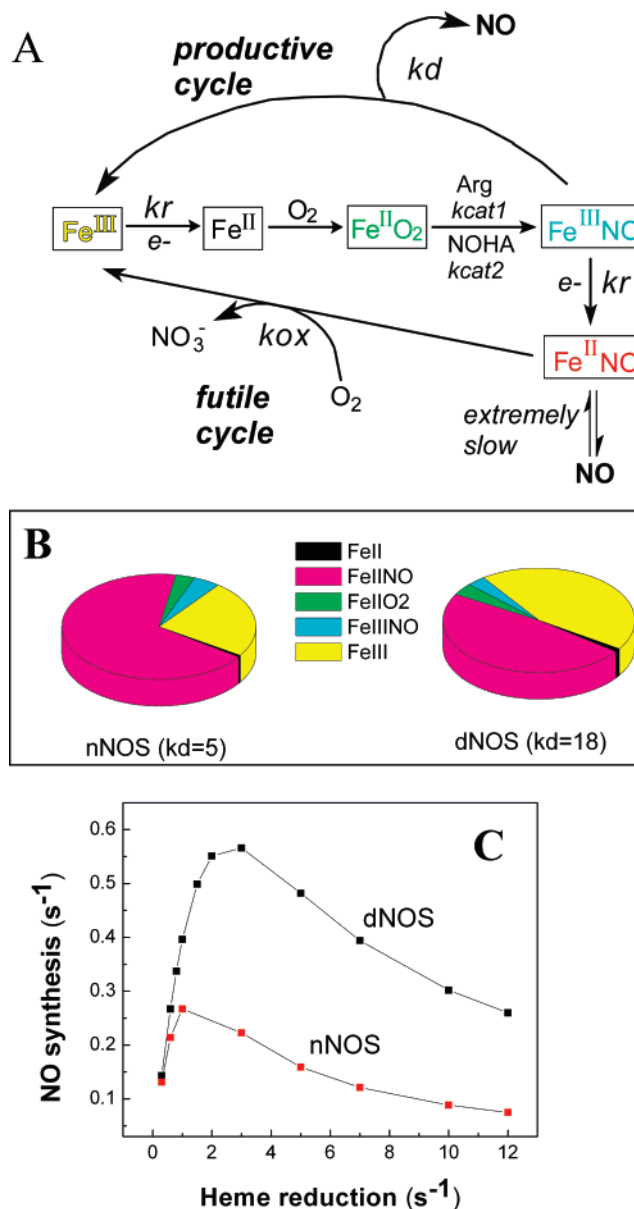


FIGURE 6: Global kinetic model for NOS catalysis and results derived from model simulations. Panel A shows that ferric enzyme reduction (kr) is rate-limiting for the biosynthetic reactions (central linear portion). $kcat1$ and $kcat2$ are the conversion rates of the $Fe^{II}O_2$ species to products in the Arg and NOHA reactions, respectively. The ferric heme–NO product complex ($Fe^{III}NO$) can either release NO (kd) in the productive cycle or become reduced (kr') to a ferrous heme–NO complex ($Fe^{II}NO$), which reacts with O_2 (kox) to generate nitrate and ferric enzyme in the futile cycle. Panel B shows the calculated dNOS and nNOS enzyme distributions during steady-state NO synthesis at a common heme reduction rate of $3 s^{-1}$. Panel C plots the steady-state NO release rates as a function of kr for dNOS and nNOS. Activities were calculated by simulation of the global kinetic model using parameters derived from dNOSoxy and nNOS. See Experimental Procedures for details.

a more active and efficient NO synthase than mammalian nNOS, while still allowing other important catalytic behaviors of the two enzymes (Ca^{2+} concentration–activity response, K_mO_2 for NO synthesis) to remain similar. That dNOS would have a higher NO release capacity plus the regulatory mechanisms of a constitutive mammalian NOS (nNOS) makes some sense because *Drosophila* evolved to express a single NOS for both its cell signaling and host

defense, while mammals utilize a different NOS for each purpose.

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SUPPORTING INFORMATION AVAILABLE

A sequence alignment of the *Drosophila* NOS oxygenase domain and other NOS sequences. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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