An Unprecedented NADPH Domain Conformation in Lysine Monooxygenase NbtG Provides Insights into Uncoupling of Oxygen Consumption from Substrate Hydroxylation*

Claudia Binda 1, Reeder M. Robinson 1, Julia S. Martin del Campo 2, Nicholas D. Keul 3, Pedro J. Rodriguez 5, Howard H. Robinson 4, Andrea Mattevi 3, and Pablo Sobrado 4

From the 4Department of Biology and Biotechnology, University of Pavia, Pavia 27100, Italy, 5Department of Biochemistry, Virginia Tech, Blacksburg, Virginia 24061, and 6Biology Department, Brookhaven National Laboratory, Upton, New York 11973

**Background:** Flavin-dependent lysine monooxygenases are involved in siderophore biosynthesis and are promising bacterial drug targets.

**Results:** Biochemical and structural characterization of lysine monooxygenase from *Nocardia farcinica* (NbtG) is presented.

**Conclusion:** An unprecedented domain conformation blocks the proper binding of NAD(P)H in the active site, which explains the high level of uncoupling observed in NbtG.

**Significance:** The structural and biochemical data should aid in drug design.

*N*-Hydroxylating monooxygenases are involved in the biosynthesis of iron-chelating hydroxamate-containing siderophores that play a role in microbial virulence. These flavoenzymes catalyze the NADPH- and oxygen-dependent hydroxylation of amines such as those found on the side chains of lysine and ornithine. In this work we report the biochemical and structural characterization of *Nocardia farcinica* Lys monooxygenase (NbtG), which has similar biochemical properties to mycobacterial homologs. NbtG is also active on D-Lys, although it binds l-Lys with a higher affinity. Differently from the ornithine monooxygenases PvdA, SidA, and KtzI, NbtG can use both NADH and NADPH and is highly uncoupled, producing more superoxide and hydrogen peroxide than hydroxylated Lys. The crystal structure of NbtG solved at 2.4 Å resolution revealed an unexpected protein conformation with a 30° rotation of the NAD(P)H domain with respect to the flavin adenine dinucleotide (FAD) domain that precludes binding of the nicotinamide cofactor. This “occluded” structure may explain the biochemical properties of NbtG, specifically with regard to the substantial uncoupling and limited stabilization of the C4a-hydroperoxoflavin intermediate. Biological implications of these findings are discussed.

Flavin-dependent *N*-hydroxylating monooxygenases (NMOs) are involved in the biosynthesis of hydroxamate-containing siderophores. These enzymes catalyze the NADPH- and oxygen-dependent hydroxylation of amine groups on the side chains of l-ornithine (l-Orn), l-lysine (l-Lys), and some primary aliphatic diamines (1–6). Many of these NMOs are linked to virulence in pathogenic bacteria. For instance, gene deletion experiments have shown that the l-Orn monooxygenases SidA from *Aspergillus fumigatus* and PvdA from *Pseudomonas aeruginosa* as well as the l-Lys monooxygenase MbtG from *Mycobacterium tuberculosis* are essential for functional siderophore biosynthesis and virulence (7–9). It is possible that these enzymes would make promising drug targets for various infectious diseases because of their requirement for the production of virulence-conferring siderophores and the lack of any human homologs.

Substantial mechanistic and structural data has been obtained in recent years on the l-Orn monooxygenases SidA, PvdA, and, more recently, on *Kutzneria* sp. 744 l-Orn monooxygenase (Ktzl) (10–12). These enzymes are very selective for l-Orn and are highly coupled, producing little or no hydrogen peroxide during catalysis (1, 13) as they stabilize the C4a-hydroperoxoflavin (FAD$_{O2H}$) intermediate long enough to allow effective hydroxylation of l-Orn (14–16). A different strategy for oxygen consumption regulation was found in Ktzl where flavin movement from an “out” to “in” conformation controls the oxidized/reduced state of the cofactor, similarly to what happens in Class A monooxygenases such as p-hydroxybenzoate hydroxylase. Despite this difference, the structures of SidA, PvdA and Ktzl all share a similar overall fold to other Class B flavin monooxygenases and a similar binding conformation for NADP$^+$, which is important for stabilization of the FAD$_{O2H}$ (6, 10, 17, 18).

Comparison of the structures of SidA in complex with l-Orn and l-Lys revealed the mechanism of substrate selectivity where the Ne of l-Orn is in a favorable position for hydroxylation by the distal oxygen of the FAD$_{O2H}$. In contrast, l-Lys protrudes farther into the active site in a suboptimal position, which causes breakdown of the FAD$_{O2H}$ resulting in produc-
tion of high levels of hydrogen peroxide and low levels of hydroxylated l-Lys (10).

In contrast to the significant mechanistic and structural data of l-Orn monooxygenases, relatively little is known about l-Lys monooxygenases. IucD from *Escherichia coli* was the first l-Lys monooxygenase characterized. IucD shares similar mechanistic features to SidA and PvdA, as this enzyme has been shown to be highly coupled and specific for NADPH and l-Lys (3, 19). Recently, characterization of the l-Lys monooxygenase MbsG from *M. smegmatis* showed that this enzyme can use both NADH and NADPH and is highly uncoupled, making more superoxide and hydrogen peroxide than hydroxylated Lys (20, 21). Also, unlike other NMOs where substrate binding accelerates turnover, l-Lys binding decreases the activity of MbsG ~2-fold by regulating the reaction with NAD(P)H (21). These results suggest that there is a higher degree of mechanistic variability among NMOs than previously thought.

The biosynthesis of the siderophore nocobactin in *Nocardia farcinica* requires functioning of NbtG, a l-Lys monooxygenase that shares >50% of sequence identity with MbsG (Scheme 1) (22). The *Nocardia* genus is comprised of parasitic bacteria, which can infect both plants and animals, including humans, and has a high mortality rate due to antibiotic resistance (22).

Here, we present a thorough biochemical and structural characterization of NbtG. The enzyme is an effective l-Lys monooxygenase that is also able to hydroxylate d-Lys. NbtG is unable to stabilize the FAD_{OOH} intermediate, which results in production of hydrogen peroxide and superoxide. The structure of NbtG was solved by x-ray crystallography and the results show an unexpected and unprecedented protein conformation that can rationalize the biochemical properties of NbtG, specifically with regard to the substantial uncoupling and limited stabilization of the FAD_{OOH}.

**EXPERIMENTAL PROCEDURES**

*Materials*—The synthetic gene encoding for NbtG from *N. farcinica* IFM 10152 was obtained from GenScript (Piscataway, NJ). Pmel and SgfI were from Promega (Madison, WI). *E. coli* TOP10 and BL21-T1R chemically competent cells were from Invitrogen. pVP56K was obtained from the Center for Eukaryotic Structural Genomics (University of Wisconsin, Madison, WI). DNA sequencing was performed at the DNA sequencing facility of the Virginia Bioinformatics Institute (Blacksburg, VA). Protein purification was performed on an ÄKTA Prime Plus FPLC (GE Healthcare). L-Lys, D-Lys, NADPH, NADP+, buffers, and salts were from Fisher. Amino acid analysis was performed using an Acquity Uitraformance Liquid Chromatography system (Waters, Milford, MA). Oxygen consumption studies were performed on a Hansatech Oxygraph (Norfolk, UK). WST-1, used for superoxide quantification, was from Dojindo Molecular Technologies, Inc. (Rockville, MD). Alcohol dehydrogenase from *Thermoanaerobium brockii* was obtained from Sigma. Glucose oxidase was from MP Biomedical (Solon, OH). Rapid-reaction studies were performed on an Applied Photophysics SX20 stopped-flow spectrophotometer (Leatherhead, UK) housed in a Coy Glove Box (Grass Lake, MI).

*Cloning*—To subclone the NbtG gene into the expression vector pVP56K, the pUC57:nbtG plasmid was incubated with Pmel and SgfI restriction endonucleases at 37 °C for 2 h. After digestion, the restriction endonucleases were inactivated by
incubation at 65 °C for 30 min. The digested gene was then ligated into pVP56K, which was previously treated with PmeI and SfiI. The ligation reaction was then transformed into E. coli TOP10 cells. Five colonies were grown for plasmid purification and sequenced to determine proper ligation of nbtG into pVP56K.

**Site-directed Mutagenesis**—Mutagenesis was performed using the QuikChange (Agilent Technologies) method following the manufacturer’s instructions. The wild-type NbtG gene, subcloned into the pVP56K plasmid, was used as the template. The mutations were confirmed by DNA sequencing at the Virginia Bioinformatics Core Sequencing Facility.

**Protein Expression and Purification**—A single colony of E. coli BL21-T1R transformed with pVP56K-nbtG was used to inoculate a 50-ml overnight LB culture flask (supplemented with 50 μg/ml kanamycin). After overnight incubation at 37 °C, 6 flasks containing 1 liter each of autoinduction media (23) (supplemented with 50 μg/ml kanamycin) were each inoculated with 10 ml of the overnight culture and grown at 37 °C with agitation at 250 rpm. When the optical density at 600 nm reached a value of ~3, the temperature was lowered to 18 °C. After overnight incubation, the cells were harvested by centrifugation, and the cell pellet was stored at −80 °C. All subsequent steps were performed at 4 °C. Cell paste (50 g) was resuspended in 150 ml of buffer A (25 mM HEPES, 300 mM NaCl, 25 mM imidazole, 5% glycerol, pH 7.5) containing 1 mM phenylmethanesulfonyl fluoride, 100 μg/ml FAD, and 25 μg/ml each of DNase, RNase, and lysozyme. This solution was stirred for 30 min. Cells were lysed by sonication using a Fisher Scientific Sonic Dismembrator Model 500 at 70% amplitude for 10 min (5 s pulses with 15 s delays). Next, the lysate was centrifuged at 35,000 × g for 1 h. The resulting supernatant was loaded at 5 ml/min onto 3 in-tandem 5-ml HisTrap FF crude columns (GE Healthcare) previously equilibrated with buffer A. After loading, the column was washed with buffer A followed by 100 ml of buffer A containing 75 mM imidazole. Bound His8-MBP-NbtG was eluted with 150 ml of buffer A containing 500 mM imidazole. Fractions were analyzed by sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE), and those containing His8-MBP-NbtG were combined. To liberate NbtG from the His8-MBP, 10 mg of tobacco etch virus protease was added, and the solution was then transferred to a 4,000 molecular weight cutoff dialysis bag and dialyzed in 2 liters of buffer A. The next day, the solution was loaded at 3 ml/min onto 3 in-tandem 5-ml HisTrap FF crude columns previously equilibrated with buffer A. In this step His8-MBP, His8-tobacco etch virus, and any uncleaved His8-MBP-NbtG remain bound to the column, whereas NbtG elutes in the flow-through. SDS-PAGE analysis of the flow-through showed highly pure NbtG; however, small amounts of His8-MBP-NbtG were still present. To remove the fusion from free NbtG, the sample was loaded at 3 ml/min onto 2 in-tandem 5-ml MBPTrap HP columns equilibrated in buffer B (25 mM HEPES, 500 mM NaCl, pH 7.5). The flow-through contained NbtG and was determined to be >95% pure by SDS-PAGE. The sample (~20 ml) was dialyzed for 3 h in 1 liter of storage buffer (100 mM sodium phosphate, 50 mM NaCl, pH 7.5) then dialyzed in 1 liter of fresh storage buffer overnight. Dialyzed NbtG was concentrated to ~300 μM, aliquoted (in ~60 μl drops), and frozen in liquid nitrogen before storage at −80 °C. NbtG K184A seleno-l-methionine was produced using the metabolic inhibition method (24). Both seleno-l-methionine-labeled NbtG K184A and unlabeled enzymes were purified as described above.

**Characterization of the Flavin Cofactor**—A solution of NbtG (25 μM) was heated at 95 °C for 10 min. After incubation on ice for 5 min, the denatured protein was precipitated by centrifugation at 15,000 × g for 10 min. The supernatant containing the flavin cofactor was used for mass spectroscopy analysis as previously described (20, 21). The extinction coefficient of the bound FAD to NbtG was determined by the addition of 5% SDS and incubation at room temperature for 5 min. The spectrum of the liberated FAD was recorded, and an extinction coefficient at 452 nm of 12,600 M⁻¹ cm⁻¹ was calculated using the extinction coefficient for free FAD of 11,300 M⁻¹ cm⁻¹.

**NADP⁺ Binding**—The binding of NADP⁺ to oxidized NbtG was monitored by recording the change in the flavin spectra at increasing concentrations of NADP⁺. Each solution consisted of 100 μl of 100 mM sodium phosphate, pH 7.5, NbtG (60 μM) and NADP⁺ (0–10 mM). The solutions were incubated on ice for 10 min before data collection. The spectra of NADP⁺ alone were subtracted from spectra of NADP⁺-NbtG at each NADP⁺ concentration. The absorbance differences showed the largest decrease at 395 nm. An extinction coefficient of 1,300 M⁻¹ cm⁻¹ for this change in absorbance at 395 nm was determined and used to calculate the concentrations of NbtG-NADP⁺ complex.

**Oxygen Consumption Assay**—Oxygen consumption was monitored in 1-ml reactions of 100 mM sodium phosphate, pH 7.5. NAD(P)H was varied between 0.1 and 5 mM. L-Lys was varied between 0 and 5 mM, and D-Lys was varied between 0 and 100 mM. For assays where NAD(P)H was varied, no Lys, 2 mM L-Lys, or 10 mM D-Lys were present. In assays where Lys was varied, NAD(P)H was held constant at 2 mM. Assays were initiated with 1 μM NbtG and monitored on a Hansatech Oxygraph system at 25 °C under constant stirring.

**Product Formation Assay**—To determine if L-Lys was hydroxylated, NbtG (8 μM) was mixed with 5 mM NADPH and 1 mM L-Lys in 100 μl of 50 mM Tris-Cl, 100 mM NaCl, pH 8.0. At 0.5, 1, 2, and 3 min, the reaction was quenched with 200 μl of 100% acetonitrile and centrifuged at 13,000 rpm for 1 min. The supernatant was withdrawn and combined with 25 μl of 200 mM borate, pH 8.0. L-Lys was derivatized by adding 3.4 μl of FMOCl (150 mM) to 130 μl of the quenched reaction. This was allowed to incubate for 5 min at 25 °C. To remove any remaining derivatizing reagent, 158 μl of 1-aminoadamantane (53 mM) was added and incubated at room temperature for an additional 15 min. The sample was analyzed by reverse phase ultraperformance liquid chromatography by loading 2 μl onto a C18 AcQuity column (2.1 × 100 mm) equilibrated in 70% eluent A (0.1% trifluoroacetic acid in water) at 0.5 min/ml. The samples were eluted with a linear gradient from 30 to 100% eluent B (0.1% trifluoroacetic acid in acetonitrile) at 0.5 min/ml, and samples were monitored at 263 nm. The elution profiles of derivatized L-Lys and L-Lys-OH were consistent with previously published data (25).

Hydroxylated Lys was quantified by a variation of the Csaky iodine oxidation assay as previously described (20, 26). Each
reaction consisted of 104 μl of 100 mM sodium phosphate, pH 7.5. NAD(P)H was varied between 0.1 and 10 mM, l-Lys was varied between 0.1 and 5 mM, and D-Lys was varied between 0 and 10 mM. In assays where NAD(P)H was varied, 2 mM l-Lys, or 10 mM D-Lys were used. For reactions with varying concentrations of d/L-Lys, 2 mM NAD(P)H was used. Reactions were initiated with 2 μM NbtG and allowed to proceed at 25 °C for 10 min while shaking at 750 rpm.

**Hydrogen Peroxide Formation Assay**—Hydrogen peroxide formed during the reaction was measured using the Pierce hydrogen peroxide detection kit as described previously (20, 21). Each reaction consisted of 104 μl of 100 mM sodium phosphate, pH 7.5. NAD(P)H was varied between 0.1 and 10 mM in the absence of d/L-Lys or with l-Lys or D-Lys held constant at 2 mM and 10 mM, respectively. Reactions were initiated with 2 μM NbtG and allowed to proceed at 25 °C for 10 min under constant orbital shaking at 750 rpm.

**Superoxide Formation Assay**—Superoxide formed during the reaction was measured by coupling the reaction with WST-1 as described previously (20, 21). Each reaction consisted of 200 μl of 100 mM sodium phosphate, pH 7.5. NAD(P)H was varied between 0.1 and 10 mM in the absence of d/L-Lys or with l-Lys or D-Lys held constant at 2 and 10 mM, respectively. Reactions were initiated with 2 μM NbtG and allowed to proceed at 25 °C for 10 min while shaking at 750 rpm.

**Flavin Oxidation**—Flavin reduction was monitored at 24 °C in 100 mM sodium phosphate, pH 7.5, in a stopped-flow spectrophotometer in single mixing mode. The stopped-flow spectrophotometer system and buffers were made anaerobic as described previously (21). Oxidized NbtG (15 μM after mixing) was mixed with various concentrations of NADPH (0.025–5 mM) after mixing). The reaction was monitored with a photodiode array spectrophotometer until flavin reduction was complete. This was repeated with l-Lys or D-Lys at 2 mM and 10 mM (after mixing), respectively.

**Flavin Reduction**—Flavin reduction was monitored at 25 °C in 100 mM sodium phosphate, pH 7.5, in a stopped-flow spectrophotometer in single mixing mode. The stopped-flow spectrophotometer system and buffers were made anaerobic as described previously (21). Oxidized NbtG (15 μM after mixing) was mixed with various concentrations of NADPH (0.025–5 mM after mixing). The reaction was monitored with a photodiode array spectrophotometer until flavin reduction was complete. This was repeated with l-Lys or D-Lys at 2 mM and 10 mM (after mixing), respectively.

**Crystallization and Structure Determination**—Purified samples of NbtG (6 mg/ml) in 25 mM HEPES/NaOH, pH 7.5, 500 mM NaCl, and 5 mM NADP+ in the presence of either l-Lys or D-Lys (various concentrations ranging from 5 to 100 mM) were used for crystallization screenings. As no crystals were obtained in these initial trials, surface mutagenesis experiments were carried out. Using the surface entropy reduction prediction server (SERp Server) (29), Lys-184, Lys-200, and Lys-202 were selected and mutated to Ala. After initial screening, promising conditions were found only with the K184A variant. As shown by the crystal structure, Lys-184 is on a loop, which is fully exposed to the solvent at >20 Å distance from the flavin ring. Moreover, the kinetic properties were very similar to those of the wild type (data not shown). Yellow crystals of the K184A NbtG variant grew in 3–4 days at 20 °C using the hanging drop vapor diffusion method by mixing 1 μl of protein solution with an equal volume of a reservoir solution containing 10% w/v PEG 6,000, 5% v/v 2,4-methylpentanediol, and 0.1 M HEPES/NaOH, pH 7.5. Optimization was achieved by the micro-seeding technique using a seed stock generated by vortexing crystals for 90 s in a seed bead tube. Crystallization experiments were then performed by adding 0.3 μl of a 100-fold diluted seed stock solution into a drop with a 2:1 ratio of protein to reservoir solution. Similar procedures were followed to crystallize a seleno-l-methionine-labeled sample of K184A NbtG.

For x-ray data collection, NbtG crystals were soaked in a cryo solution containing 16% w/v PEG 6,000, 14% v/v 2,4-methylpentanediol, and 0.1 M HEPES/NaOH, pH 7.5, 5 mM NADP+, d/L-Lys (at the concentration used for crystallization), 20% v/v glycerol, and flash-cooled in liquid nitrogen. Many crystals were

**Structure and Mechanism of Lysine Monooxygenase NbtG**
screened for x-ray diffraction at the beamlines ID23-EH1 (European Synchrotron Radiation Facility, Grenoble, France), X06SA (Swiss Light Source, Villigen, Switzerland), and X29 (National Synchrotron Light Source, Upton, NY). Data processing and scaling were performed using MOSFLM (30) and programs of the CCP4 package (31). Final statistics of the collected datasets are reported in Table 1. The NbtG structure was solved by single-wavelength anomalous diffraction on the selenium-edge using the program SHELX (32). Because the crystals were highly isomorphous, it was possible to merge two datasets at 2.7 and 2.9 Å resolution to increase redundancy. The eight selenium sites (including that of the N-terminal methionine, which is left in the recombinant protein after His-tag cleavage) were identified and used for phasing with the program PHENIX (33). The quality of the initial electron density map was poor (figure of merit = 0.27), but a significant improvement was obtained after PHENIX density modification (averaging among the 4 molecules present in the asymmetric unit). This allowed us to perform autobuilding, which yielded good statistics (model-map CC = 62%, R1 = 36%, free R1 = 41%), although a large portion of the protein needed to be manually traced by COOT (34). Refinement was carried out using a native data set at 2.4 Å resolution with the program REFMAC5 (35). Figures were produced by CCP4mg (36).

**RESULTS**

**Purification of Wild-type NbtG and Variants**—Wild-type NbtG and the K184A isoform used for crystallography were expressed as a fusion to His8-MBP in pVP56K. This was critical to obtaining soluble active enzymes, as expression with only a His8 tag produced insoluble protein. Overall, a yield of ~30 mg of purified NbtG was obtained per liter of growth media with a flavin incorporation of ~75%. High resolution MALDI-TOF of the extracted flavin from the active site of NbtG yielded an m/z value of 784.15, which is consistent with FAD (data not shown). The flavin spectrum of NbtG displayed characteristic absorptions at 380 nm and 452 nm.

![Extinction Coefficient](chart.png)

**TABLE 2**

Activity of NbtG monitored by oxygen consumption

| Parameter          | No Lys | L-Lys | D-Lys |
|--------------------|--------|-------|-------|
| kcat (s⁻¹)         | 0.58 ± 0.02 | 1.08 ± 0.01 | 2.9 ± 0.1 |
| KNADPH (mM)        | 0.20 ± 0.02 | 0.14 ± 0.01 | 0.51 ± 0.06 |
| kcat/KNADPH (M⁻¹s⁻¹) | 2900 ± 300 | 7700 ± 500 | 5600 ± 700 |
| NADH kcat (s⁻¹)    | 0.32 ± 0.01 | 1.29 ± 0.04 | 2.5 ± 0.1 |
| KNADH (mM)         | 0.36 ± 0.05 | 0.40 ± 0.04 | 1.6 ± 0.2 |
| kcat/KNADH (M⁻¹s⁻¹) | 900 ± 100 | 3300 ± 400 | 1500 ± 200 |

**TABLE 1**

Data collection and refinement statistics for the NbtG structure

| SeMet, seleno-L-methionine. | SeMet, seleno-L-methionine (crystal 1) | SeMet, seleno-L-methionine (crystal 2) | Native |
|-----------------------------|----------------------------------------|----------------------------------------|-------|
| PDB code                    | P2₁                                   | P2₁                                    | P2₁   |
| Space group                 |                                       |                                       |       |
| a (Å)                       | 80.3                                  | 80.4                                   | 80.6  |
| b (Å)                       | 121.5                                 | 122.1                                  | 121.9 |
| c (Å)                       | 92.2                                  | 92.6                                   | 91.9  |
| β (°)                       | 97.7                                  | 97.6                                   | 97.7  |
| Resolution (Å)              | 2.7                                   | 2.9                                    | 2.4   |
| Rsym (%)                    | 10.1 (96.8)                           | 10.8 (99.5)                            | 7.6 (98.4) |
| CC1/2 (%)                   | 0.997 (0.423)                         | 0.997 (0.627)                         | 0.999 (0.563) |
| Completeness (%)            | 85.4 (79.9)                           | 100.0 (99.9)                          | 99.2 (98.5) |
| Anomalous completeness (%)  | 82.3 (78.3)                           | 99.6 (99.0)                           |       |
| Redundancy                  | 6.1 (6.4)                             | 7.1 (7.1)                              | 6.5 (6.3) |
| Anomalous redundancy        | 3.0 (3.1)                             | 3.6 (3.6)                              |       |
| Unique reflections          | 40862                                 | 39427                                  | 68,260 |
| I/σI (cryst)                | 13.4 (1.4)                            | 14.2 (1.8)                             | 12.3 (1.0) |
| Protein/FAD atoms (N)       | 11,431/53 (N × 4)                     |                                       |       |
| Water atoms (N)             | 239                                   |                                       |       |
| Average B value (Å²)        |                                       |                                       | 43.5  |
| Rcryst (%)                  |                                       |                                       | 20.7  |
| Rfree (%)                   |                                       |                                       | 28.2  |
| Root mean square bond length (Å) | 0.013                             |                                       |       |
| Root mean square bond angles (°) | 1.7                                  |                                       |       |

* Rsym = Σ[Ii−(I)/ΣIi], where Ii is the intensity of ith observation, and I is the mean intensity of the reflection.

* A cut-off criterion for resolution limits was applied on the basis of the mean intensity correlation coefficient of half-samples of each dataset (CC1/2) (45).

* d Rcryst = Σ|Fc-Fe|/ΣFc, where Fe and Fc are the observed and calculated structure factor amplitudes, respectively. Rcryst and Rfree were calculated using the working and test sets, respectively.

* Values in parentheses are for reflections in the highest resolution shell.
zyme selectivity, in contrast to other Class B flavin monooxygenases, which are all highly selective for NADPH (1, 37–39).

**Lysine Hydroxylation—**NbtG displayed significant activity as measured by the oxygen consumption assay. To determine if hydroxylation of L-Lys was taking place, NbtG was reacted with NADPH in excess and 1 mM L-Lys, and the reaction was monitored by ultraperformance liquid chromatography after derivatization of the amino acid as described under “Experimental Procedures.” The results clearly show that NbtG is able to produce L-Lys-OH (Fig. 2). To quantitate the amount of L-Lys-OH produced, the Csaky iodine oxidation assay was performed. The reaction was initially done using NADPH as the reducing substrate (Table 3). Under fully coupled conditions, one equivalent of hydroxylated d/l-Lys should be produced for every 1 eq of oxygen consumed. Very similar \( k_{cat} \) values were determined for hydroxylation of d/l-Lys; however, the values were \( \sim 3–7 \)-fold lower than the \( k_{cat} \) values obtained in the oxygen consumption assay. These results suggest that NbtG is highly uncoupled, producing reactive oxygen species in addition to hydroxylated d/l-Lys. Substrate inhibition was observed when NADPH was varied in the presence of saturating levels of either d- or l-Lys, with a lower \( K_m^{(NADPH)} \) value with l-Lys. The \( k_{cat}/K_m \) value for l-Lys was \( \sim 6 \)-fold higher than for d-Lys, which originates mainly from a lower \( K_m \) value for l-Lys. Weak

**FIGURE 2. Production of hydroxylated lysine (L-Lys-OH) by NbtG.** A, scheme of the derivatization reaction of L-Lys with FMOC-Cl. FMOC-Cl can react with the amino groups of either L-Lys or L-Lys-OH. B, ultraperformance liquid chromatography chromatogram (263 nm) showing the elution of L-Lys and L-Lys-OH derivatized with FMOC-Cl. The inset shows a zoomed-in view of the L-Lys peak decreasing and the L-Lys-OH peak increasing between 0 and 3 min.
TABLE 3
Activity of NbtG monitored by Lys hydroxylation

Conditions were 100 mM sodium phosphate, pH 7.5, 25°C. The final concentration of L-Lys was 2 mM and of D-Lys was 10 mM. Uncoupling was calculated using the expression (1 - $k_{cat(O2)} / k_{cat(N-OH)}$) × 100, where $k_{cat(N-OH)}$ is the $k_{cat}$ calculated measuring hydroxylated D/L-Lys, and $k_{cat(O2)}$ is $k_{cat}$ from the oxygen consumption assay.

| Parameter | L-Lys | D-Lys | NADPH | NADH |
|-----------|-------|-------|-------|-------|
| $k_{cat}$ (s$^{-1}$) | 0.32 ± 0.02 | 0.41 ± 0.03 | 0.62 ± 0.08 | 0.31 ± 0.02 |
| $K_{M,NADPH}$ (mM) | 0.7 ± 0.2 | 0.5 ± 0.1 | 0.6 ± 0.1 | 3.8 ± 2.4 |
| $K_{M,NADPH}$ (mM) | 2.4 ± 0.1 | 2.6 ± 0.1 | 5.7 ± 1.7 | 2.1 ± 1.5 |
| $k_{cat}/K_{M,NADPH}$ (m$^{-1}$ s$^{-1}$) | 460 ± 130 | 520 ± 120 | 1100 ± 300 | 80 ± 50 |
| $K_{M,Lys}$ (mM) | 0.35 ± 0.05 | 2.6 ± 0.3 | 0.9 ± 0.3 | 4.2 ± 0.6 |
| $K_{M,D-Lys}$ (mM) | 15 ± 3 | | 6.3 ± 2.5 | |
| $k_{cat}/K_{M,D-Lys}$ (m$^{-1}$ s$^{-1}$) | 910 ± 140 | 100 ± 20 | 580 ± 230 | 70 ± 10 |
| Uncoupling (%) | 70 | 86 | 52 | 92 |

Substrate inhibition was observed with L-Lys, which is a common feature observed in other NMOs. With D-Lys, no substrate inhibition was observed, most likely because of the lower affinity. Very similar results were observed when NADH was used as the coenzyme (Table 3).

TABLE 4
Oxygen reactive species produced during oxidation of NbtG

Conditions were 100 mM sodium phosphate, pH 7.5, 25°C. The final concentration of L-Lys was 2 mM and of D-Lys 10 mM.

| Assay | No Lys | 2 mM L-Lys | 10 mM D-Lys |
|-------|--------|------------|------------|
| Hydrogen peroxide | | | |
| NADPH | 0.3 ± 0.1 | 0.20 ± 0.04 | 0.40 ± 0.05 |
| NADH | 0.27 ± 0.02 | 0.32 ± 0.05 | 0.40 ± 0.04 |
| Superoxide | | | |
| NADPH | 0.30 ± 0.02 | 0.30 ± 0.02 | 0.23 ± 0.01 |
| NADH | 0.22 ± 0.02 | 0.170 ± 0.004 | 0.34 ± 0.02 |

With enzymic assays highlighting such a substantial uncoupling, we next asked the question as to the type of product generated, as flavoenzymes can produce both hydrogen peroxide and superoxide upon reacting with oxygen. The kinetic analysis indicated that the oxidase activity of NbtG partly produces superoxide and hydrogen peroxide, which were detected using both NADPH and NADH either in the absence or presence of D/L-Lys (Table 4). In summary, NbtG is rather inefficient in Lys hydroxylation because flavin reoxidation can inefficiently occur through the direct generation of hydrogen peroxide and/or superoxide. This finding warranted a more in-depth kinetic and structural analysis to understand the features underlying such a functional property.

**Flavin Oxidation**—Flavin oxidation was monitored as a function of molecular oxygen concentration. NbtG oxidation occurs in a single step process without the stabilization of a FAD$\text{O}OH$ intermediate. In the absence of D/L-Lys, a rate of 3,100 ± 200 m$^{-1}$ s$^{-1}$ was determined. In the presence of L-Lys, this value increased to 3,900 ± 100 m$^{-1}$ s$^{-1}$, whereas the rate was 5,400 ± 400 m$^{-1}$ s$^{-1}$ in the presence of D-Lys (Fig. 4). The effect of excess NADP$^+$ on flavin oxidation was also determined. The addition of 0.5–10 mM NADP$^+$ did not change the rate of flavin oxidation, nor was there evidence of the spectra for the FAD$\text{O}OH$ intermediate (Fig. 4C).

**NADP$^+$ Binding**—It has been determined that NADP$^+$ is essential for stabilization of the FAD$\text{O}OH$ intermediate in members of Class B flavin-dependent monooxygenases. In PvdA and SidA, NADP$^+$ has similar affinity to NADPH. Using the spectral perturbation methods, a $K_p$ value of 2.0 ± 0.5 mM was determined for NADP$^+$ binding to oxidized NbtG (Fig. 5). Thus, for NbtG, the $K_p$ for NADP$^+$ is ~10-fold higher than the $K_p$ for NADH (Table 5).

**Three-dimensional Structural Analysis of NbtG**—The crystal structure of NbtG was solved by single-wavelength anomalous diffraction at the selenium edge and refined at 2.4 Å resolution (Table 1). The four NbtG molecules present in the asymmetric unit form a compact tetramer with the same arrangement...
observed in the L-Orn monooxygenases SidA (10) and PvdA (11) (their monomeric and dimeric structures, respectively, form a tetramer through application of crystallographic symmetry operators); in KtzI a similar but not identical tetramer was observed (12). The NbtG subunit folds into a two-domain topology similar to that of other NMOs: the FAD binding domain (residues 1–178 and 337–429), including the non-covalently bound FAD cofactor (Fig. 6B), and the NADPH binding domain (residues 179–336). Nearly all 429 residues of the protein could be modeled for each of the four polypeptide chains present in the asymmetric unit except for monomer D, in which 1/3 of the molecule (i.e. most of the NADP+ domain) lacks clear electron density (Fig. 6A).

The structures of the homologous SidA (10) and PvdA (11) NMOs were elucidated in complex with L-Orn and NADP+H, whereas despite extensive screening of NbtG crystals grown in the presence of either NADP+H/NAD+H and/or D/L-Lys, no electron density for any ligand except for the FAD cofactor could be detected. NbtG shares about 20% sequence identity and the same domain topology with SidA, PvdA, and KtzI. However, comparison of NbtG structure with those of SidA, PvdA, and KtzI reveals a large change in domain orientation. More specifically, the program DynDom (40) indicates that the NADPH domain of NbtG is rotated by 33° with respect to the same domain of SidA using the superimposed FAD domains as reference. Such a rotation occurs through an axis passing through the residues (175–178 and 335–337 in NbtG) linking the two protein domains (Fig. 7A). This finding provides a rationale for the unsuccessful attempts to obtain the NADP+ and D/L-Lys bound to NbtG in this crystalline form.

The domain rotation found in the NbtG structure makes the nicotinamide binding site in front of the FAD ring become occluded by an α-helical segment, preventing NADP+ binding (Figs. 7B and 8). Likewise, structural comparison shows that the residues responsible for binding the L-Orn backbone group in SidA are conserved in NbtG (Lys-64, Asn-252, Ser-402), but their relative positions are modified (Fig. 8). In essence, the domain rotation featured by the NbtG structure drastically alters the active site geometry and accessibility. Differences in the NAD(P)+ binding site may also explain the lack of NAD+...
NADP\(^+\) specificity found in NbtG. In SidA the position of the NADP\(^+\) adenine-ribose moiety is stabilized by a network of interactions that also involve the phosphate. In particular, the ribose 3’-OH is H-bonded to both the side chain and the peptide nitrogen of Ser-254, whereas the phosphate interacts with Ser-325 and with Arg-279 (37). In NbtG, Arg-279 is substituted by Pro-238, and Ser-325 is substituted by Asp-277, whose longer side chain is likely to interact with the ribose rather than with the phosphate. All of these NbtG residues are also conserved in MbsG, which is consistent with the similar cofactor specificity properties of the mycobacterial enzyme.

**Steady-state Kinetic Properties of P238R**—To determine if replacing Pro at position 238 to Arg resulted in a higher affinity and specificity for NADPH, the NbtG P238R variant was created. The steady-state kinetic parameters were determined by monitoring oxygen consumption and lysine hydroxylation. Oxygen consumption data show that in the presence of L-Lys the \(k_{\text{cat}}\) value is only ~2-fold lower, and the \(K_{\text{NADPH}}\) value is ~5-fold higher than wild type; however, no activity was observed with NADH (Table 6). Similar results were obtained when lysine hydroxylation was monitored. The levels of uncoupling are very similar to those calculated for the wild-type enzyme.

**Steady-state Kinetic Properties of K64A**—Although the structure of NbtG did not contain electron density for Lys, the amino acids involved in L-Orn binding in SidA, PvdA, and KtzI are conserved in NbtG (Fig. 8). We probed the role of K64 in L- and D-Lys binding by mutating it to Ala. The steady-state lysine hydroxylation data clearly show that the major effect is on the \(K_m\) value for L- and D-Lys. The \(K_{\text{L-Lys}}\) value increased ~40-fold, whereas the \(K_{\text{D-Lys}}\) value increased ~15–20-fold. In contrast, the \(k_{\text{cat}}\) values decreased only 3–13-fold (Table 7). The uncoupling values for K64A are similar to the values obtained for the wild-type enzyme (Table 7). These data indicate that K64A is primarily involved in Lys binding in NbtG as predicted from the three-dimensional structure.

**DISCUSSION**

NMOs represent a class of flavin monooxygenases involved in the biosynthesis of hydroxamate-containing siderophores whose iron-chelating properties are responsible for the virulence of microbes that secrete them. The L-Lys monooxygenase NbtG from *N. farcinica* catalyzes the hydroxylation of L-Lys in the pathway leading to the production of the siderophore noco-

![Flavin oxidation](image)

The data were fit to Equation 4. B, \(k_{\text{obs}}\) values as a function of \(\text{O}_2\) without Lys (open circles), with 2 mM L-Lys (solid circles), and with 10 mM D-Lys (open squares, dashed line). C, spectra of NbtG flavin oxidation with 3 mM NADP\(^+\). The spectra of flavin oxidation were subtracted from the spectrum of 3 mM NADP\(^+\) alone. The inset shows the deconvoluted flavin spectra. The data were fit to a two-step process to determine if a transient C4a-hydroperoxyflavin is present. Pro-K (Applied Photophysics) was used to deconvolute the flavin spectra.
binding of NADPH, which reduces the flavin cofactor that, in turn, reacts with molecular oxygen generating the FAD$_{OOH}$ intermediate capable of donating the distal oxygen to the substrate.

We first studied the NbtG reaction by probing the enzyme cofactor specificity. In contrast to L-Orn monooxygenases, which show a clear preference for NADPH, NbtG can also use NADH as a reducing agent, although at a lower efficiency. The same was observed for MbsG where, nevertheless, the $K_m$ values for both cofactors are higher than those measured with NbtG: for the mycobacterial enzyme $K_{NADPH}$ and $K_{NADH}$ are 12.0 and 7.0 mM, respectively, whereas values of 0.20 and 0.36 mM were determined with NbtG (Table 2) (21). In either case the presence of the L-Lys substrate does not significantly

**FIGURE 5.** NADP$^+$ binding. A, flavin spectra changes as a function of increasing concentrations of NADP$^+$ (0–10 mM). The spectral changes show an increase in absorbance at 340 nm and a decrease in absorbance at 395 nm. B, spectra differences after subtracting the spectrum of NbtG with 0 mM NADP$^+$. C, determination of the $K_D$ value of NADP$^+$. The concentration of the NbtG-NADP$^+$ complex was plotted as a function of NADP$^+$ to determine a $K_D$ value.

**FIGURE 6.** Crystal structure of NbtG. A, ribbon diagram of the crystallographic tetramer of NbtG observed in the asymmetric unit. Monomers A, B, C, and D of the tetramer are represented in green, light gray, dark gray, and black, respectively. The FAD cofactor is drawn as sticks for each monomer, with carbon, oxygen, nitrogen, and phosphorus atoms colored in yellow, red, blue, and magenta, respectively. For the sake of clarity, flexible regions that lack electron density are indicated only for monomer A as green dashed lines (residues 143–149, 158–160, 241–247; the C-terminal 414–429 residues could not be included in the final model). B, close-up view (same orientation as A) of FAD bound to monomer A with its refined weighted 2$F_o$ – $F_c$ electron density contoured at 1.2 $\sigma$.

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Structure and Mechanism of Lysine Monooxygenase NbtG

![Comparison of NbtG (green) and SidA (orange) active sites. For the sake of clarity, only the flavin ring of NbtG is drawn. Equivalent residues surrounding the flavin site of the two enzymes and the position of the L-Lys substrate bound to the NbtG active site are drawn (dashed lines indicate hydrogen bonds, which stabilize substrate binding). Loop 276–279 of NbtG and loop 321–324 of SidA are shown to highlight the different conformation.](image)

![Comparison of SidA (PDB code 4b64; NADP⁺ complex) and NbtG. A, the large change in the orientations of the NADPH binding domains is visualized by superimposing NbtG (green) and SidA (orange) structures using their respective FAD binding domains (root mean square deviation = 2.5 Å for the equivalent Cα atoms). The shaded arrow outlines the additional 33° rotation, which is needed for the optimal superposition of the NADP⁺ binding domains (highlighted as bold worms). B, close-up view of the flavin site in NbtG (green) and SidA (orange). Because of the domain rotation, the nicotinamide-ribose site in NbtG is physically occluded by an α-helix (residues 215–229) of the NADP⁺ binding domain.](image)

![Figure 7: Comparison of SidA (PDB code 4b64; NADP⁺ complex) and NbtG. A, the large change in the orientations of the NADPH binding domains is visualized by superimposing NbtG (green) and SidA (orange) structures using their respective FAD binding domains (root mean square deviation = 2.5 Å for the equivalent Cα atoms). The shaded arrow outlines the additional 33° rotation, which is needed for the optimal superposition of the NADP⁺ binding domains (highlighted as bold worms). B, close-up view of the flavin site in NbtG (green) and SidA (orange). Because of the domain rotation, the nicotinamide-ribose site in NbtG is physically occluded by an α-helix (residues 215–229) of the NADP⁺ binding domain.](image)

![Figure 8: Comparison of NbtG (green) and SidA (orange) active sites. For the sake of clarity, only the flavin ring of NbtG is drawn. Equivalent residues surrounding the flavin site of the two enzymes and the position of the L-Lys substrate bound to the NbtG active site are drawn (dashed lines indicate hydrogen bonds, which stabilize substrate binding). Loop 276–279 of NbtG and loop 321–324 of SidA are shown to highlight the different conformation.](image)

![TABLE 7 Steady-state kinetic parameters for NbtG K64A monitored by Lys hydroxylation](table)

| Parameter | NADPH | NADH |
|-----------|-------|------|
| L-Lys   | D-Lys | L-Lys | D-Lys |
| $k_{cat}$ (s⁻¹) | 0.34 ± 0.01 | 0.130 ± 0.004 | 0.180 ± 0.005 | 0.021 ± 0.001 |
| $K_{cat}$ (mM) | 13 ± 1 | 49 ± 4 | 40 ± 3 | 62 ± 5 |
| $k_{cat}/K_{cat} (M^{-1} s^{-1})$ | 27 ± 3 | 2.7 ± 0.2 | 4.4 ± 0.4 | 3.8 ± 0.03 |
| Uncoupling, % | 78 | 89 | 69 | 94 |

![TABLE 6 Steady-state kinetic parameters for NbtG P238R](table)

| Parameter | O₂ consumption | L-Lys hydroxylation |
|-----------|----------------|---------------------|
| $k_{cat}$ (s⁻¹) | 0.60 ± 0.04 | 0.146 ± 0.009 |
| $K_{cat}$ (mM) | 0.8 ± 0.2 | 0.7 ± 0.2 |
| $k_{cat}/K_{cat} (M^{-1} s^{-1})$ | 790 ± 170 | 210 ± 60 |
| $K_{cat}$ (mM) | 3.6 ± 0.4 | Uncoupling (%) |
| | 76 | |

binding (Arg-279 and Ser-325 side chains) do not form in NbtG (and neither in MbsG) because these residues are not conserved, and therefore, either NADH or NADPH would indifferently bind to these Lys monooxygenases. Substitution of Pro to an Arg at position 238 (the corresponding position for Arg-279 in SidA) converted NbtG into a NADP⁺-specific monooxygenase but increased the $K_{cat}$ value for NADPH. Furthermore, the P238R enzyme was as uncoupled as wild-type NbtG. These results suggest that the mode of binding of NADP(H) in NbtG is different from what is observed in SidA and PvdA.

The kinetics of the NbtG reaction were investigated by measuring the O₂ consumption rate under steady-state conditions by using an Oxygraph. A $k_{cat}$ value of 0.58 s⁻¹ was determined for NADPH, and a ~2-fold increase was obtained in the presence of L-Lys, with this substrate effect being even more pronounced with NADH. We also probed the substrate stereospecificity of NbtG, and we found that the enzyme is also active on D-Lys, although it binds L-Lys with a higher affinity. The results of the K64A variant support a conserved amino acid binding site among members of the NMO group of enzymes.

We then investigated the overall reaction leading to substrate hydroxylation by isotope effect analysis and by quantifying the final product using the Csaky assay. Flavin reduction occurs through two phases, one being ~10-fold faster than the other.

alter the affinity for the coenzymes. A comparative analysis of NbtG and SidA structures shows that the interactions in SidA that stabilize the NADPH 2'-OH-phosphate during
and both dependent on NADPH concentration. Although hydride transfer has been shown to be the rate-limiting step in SidA, isotope effect experiments showed that in NbtG this step is only partially rate-limiting. The large solvent kinetic isotope effect suggests that a proton exchangeable step is the slow step in the reaction. Similar to MbsG, we found that the $k_{\text{cat}}$ of d/L-Lys hydroxylation is significantly lower than that of O$_2$ consumption even in the presence of the substrate, and production of superoxide and hydrogen peroxide was also detected using both NADPH and NADH (Table 4). This means that NbtG undergoes a number of unproductive catalytic cycles with an uncoupling level of 70% for l-Lys hydroxylation from O$_2$ consumption using NADPH as the reducing agent and 52% in the case of NADH (Table 3). Even more pronounced uncoupling effects were detected with n-Lys as the substrate.

NbtG represents the first crystal structure of a l-Lys monooxygenase (Fig. 6A). Structural comparison with l-Orn monooxygenases revealed that although the NADPH domain is individually very similar to those of SidA and PvdA, in NbtG it is differently oriented by about 30° with respect to the FAD domain. Importantly, such a domain rotation alters the NADPH binding site, which becomes physically inaccessible to the dinucleotide cofactor. Although domain rotations have been found among members of this family of flavoenzymes (41) (including NMOs, Baeyer-Villiger enzymes, and flavin-containing monooxygenases), such a large change so drastically affecting the active center has not been observed thus far. The characteristic mechanistic feature of this family of enzymes is stabilization of the FADOOH intermediate by NADP$^+$, which remains bound throughout the catalytic reaction ensuring a high degree of coupling. This effect is absent in NbtG as even high NADP$^+$ concentrations do not promote FADOOH stabilization. Consistently, the enzyme exhibits a rather loose binding of NADP$^+$ as demonstrated by the measured $K_D$ value of 2 mM, which indicates a much weaker affinity compared with that observed in the family members (typical values in the 2–100 μM range) (11, 12, 15–17, 39, 42, 43). Collectively, these findings indicate that in NbtG the catalytic uncoupling of O$_2$ consumption from substrate hydroxylation is related to the pronounced stability of an NADP-domain conformation, which occludes the NADP(H) binding site. After reduction of the flavin by NADPH, the enzyme might quickly release NADP$^+$ and adopt the occluded conformation observed in the crystal structure. Along with this, flavin reoxidation will occur without any stable formation of the FADOOH whose stability requires bound NADP$^+$ (16, 44). Alternatively, the FADOOH might form but will quickly decay upon release of NADP$^+$.

These findings raise the intriguing question as to the biological significance of these functional and structural features. The possibility exists that NbtG and the homologous mycobacterial MbsG are uncoupled because they dually function as both Lys hydroxylases and NADPH oxidases, with this second activity possibly being involved in ROS generation and signaling. Alternatively, these properties might be instrumental in enzyme regulation by yet-unknown factors that control the oxidase versus the hydroxylase activities, for instance through stabilization of the functionally distinct enzyme conformations.

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