

Structure and Function of NzeB, a Versatile C–C and C–N Bond-Forming Diketopiperazine Dimerase

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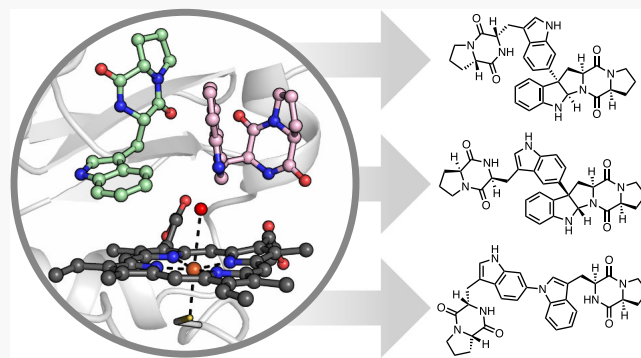


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ABSTRACT: The dimeric diketopiperazine (DKPs) alkaloids are a diverse family of natural products (NPs) whose unique structural architectures and biological activities have inspired the development of new synthetic methodologies to access these molecules. However, catalyst-controlled methods that enable the selective formation of constitutional and stereoisomeric dimers from a single monomer are lacking. To resolve this long-standing synthetic challenge, we sought to characterize the biosynthetic enzymes that assemble these NPs for application in biocatalytic syntheses. Genome mining enabled identification of the cytochrome P450, NzeB (*Streptomyces* sp. NRRL F-5053), which catalyzes both intermolecular carbon–carbon (C–C) and carbon–nitrogen (C–N) bond formation. To identify the molecular basis for the flexible site-selectivity, stereoselectivity, and chemoselectivity of NzeB, we obtained high-resolution crystal structures (1.5 Å) of the protein in complex with native and non-native substrates. This, to our knowledge, represents the first crystal structure of an oxidase catalyzing direct, intermolecular C–H amination. Site-directed mutagenesis was utilized to assess the role individual active-site residues play in guiding selective DKP dimerization. Finally, computational approaches were employed to evaluate plausible mechanisms regarding NzeB function and its ability to catalyze both C–C and C–N bond formation. These results provide a structural and computational rationale for the catalytic versatility of NzeB, as well as new insights into variables that control selectivity of CYP450 diketopiperazine dimerases.



INTRODUCTION

The tryptophan-linked dimeric diketopiperazine (DKP) alkaloids are a family of over 50 natural products (NPs) isolated from bacteria, fungi, and plants.^{1–4} Their dimerization modes encompass an array of stereo- and regiochemical connectivities, and the potent biological activity of these NPs elicited the development and application of effective methods for their total synthesis.^{3–5} In particular, biomimetic strategies enabled access to C₂-symmetrical dimers, such as (–)-ditryptophenalanine (**3**, Figure 1), wherein two identical monomers are coupled at a late stage, resulting in highly convergent and efficient syntheses.^{6,7} However, the chemical synthesis of structurally challenging non-symmetrical DKP dimers such as (+)-naseaezines A (**1**) and B (**2**),^{8,9} (–)-naseaezine C (**4**),^{10,11} and the related alkaloid (–)-aspergilazine A (**5**)^{10,12} continues to inspire the development of new synthetic routes.^{4,9,13–16} The synthesis of structurally complex non-symmetrical dimeric DKPs requires independent synthesis of appropriately substituted precursors to allow selective formation of a key C–C or C–N bond that adjoins the two subunits.^{9,17,18} Seeking to develop an efficient approach toward DKP dimer assembly, we envisioned a process wherein the

site- and stereochemical outcome of dimerization would be dictated by the biocatalyst and a single DKP monomer could be dimerized selectively to a variety of site- and stereoisomeric products.

While dozens of fungal dimeric diketopiperazine natural products have been reported since the isolation of **2**,^{1–4,6} fungal biosynthetic gene clusters (BGCs) responsible for dimerization of DKPs have only recently been identified and have been limited to biosynthesis of C₂-symmetrical dimers.¹⁹ Characterization of these BGCs identified cytochromes P450 that mediate the oxidative convergence of two DKP monomers to generate C₂-symmetrical dimers, validating long-standing hypotheses proposed in isolation, characterization, and total syntheses of these molecules.¹⁹ While the majority of DKP dimers have been isolated from fungi, marine actinomycetes

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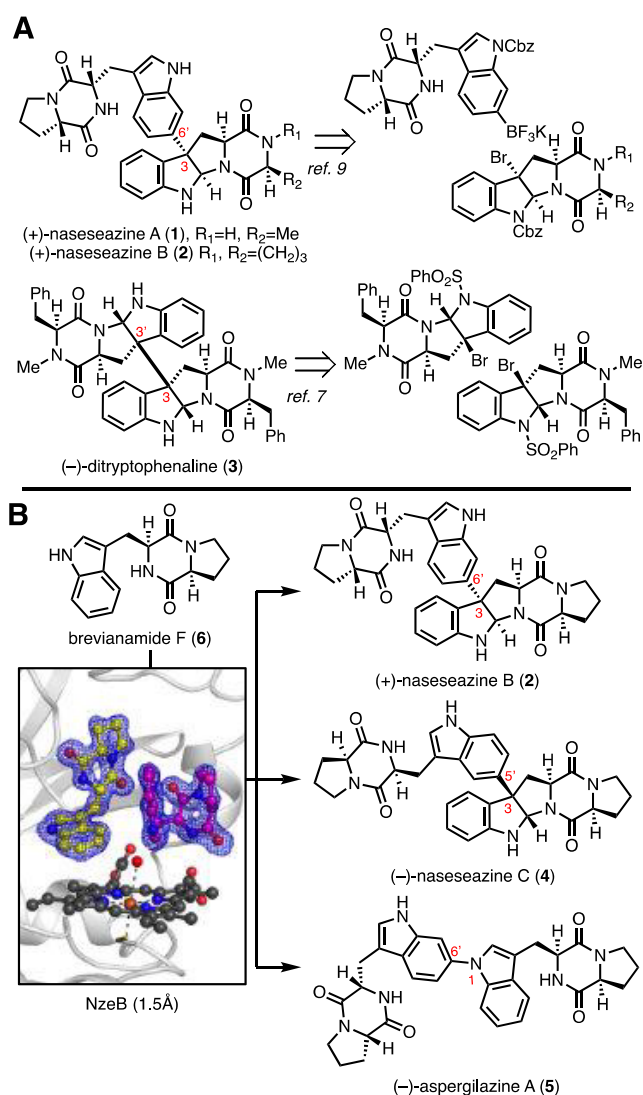


Figure 1. (A) Representative examples of substrate-controlled, biomimetic total syntheses of dimeric diketopiperazine natural products. (B) Current work describing the molecular basis for catalyst-controlled NzeB-mediated assembly of isomeric dimers.

have recently emerged as a new source for non-symmetrical dimeric DKP NPs. Notably, bacterial DKP dimers display coupling patterns distinct from their fungal counterparts, with the exception of **4**, a rare example of a secondary metabolite isolated from both fungal and bacterial sources.^{8,10,11,20} Contemporaneous efforts to identify DKP dimer BGCs from bacteria have led to the recent identification of gene clusters for dimers **4** (*nasc*)²¹ (*Streptomyces* sp. CMB-MQ030) and the (+)-naseaezines A (**1**) and B (**2**) (*nas*) and (-)-aspergilazine A (**5**) gene clusters (*Streptomyces* sp. NRRL-S1868).²² While functional characterization of these BGCs has provided insight into the biogenesis of these NPs, the molecular basis for the selective formation of non-symmetrical dimers by these P450s has remained elusive.

Herein, we report the identification of NzeB, a versatile P450 that catalyzes both C–C and C–N bond formation through bioinformatics and sequence similarity network (SSN) analysis. Structural and functional characterization of NzeB provides the first insights into the molecular basis of selectivity displayed by the DKP dimerizing class of P450s. Quantum

mechanics (QM) calculations were performed to elucidate plausible mechanisms for NzeB-catalyzed C–C and C–N dimerization.

Identification of the *nzn* Gene Cluster. Previous studies by Tian et al. identified a gene cluster (*nasc*) from *Streptomyces* sp. CMB-MQ030 composed of a cyclodipeptide synthase (CDPS), *nascA*, and P450, *nascB*, which mediates assembly of a cryptic DKP dimer, (-)-naseaezine C (**4**).²¹ Our independent sequencing and annotation of the *Streptomyces* sp. CMB-MQ030 genome (Supporting Information, Supplementary Table 1) identified an additional operon, *nzn*, also composed of a CDPS and P450, each with high sequence homology to *nasc* counterparts (NznA/NascB = 61%, NznB/NascB = 68%). Given the high sequence homology to the *nasc* gene cluster, we hypothesized that P450 NznB catalyzes formation of the site- and stereoisomeric (+)-naseaezines A (**1**) and B (**2**), the original dimeric DKP NPs isolated from this strain. To assess NznB as the (+)-naseaezines A and B synthase as well as evaluate relative catalytic activities, both recombinant proteins (NznB and NascB) were subjected to dimerization conditions with predicted native substrate breviramide F (**6**). Homodimerization reactions demonstrated that NznB catalyzes conversion of **6** into **2**, albeit in modest total turnover number (TTN) of 145, relative to the conversion of **6** to **4** by NascB, >500 TTN (Figure 2).²³

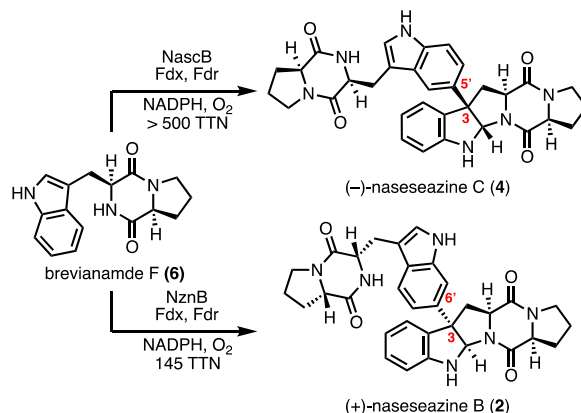
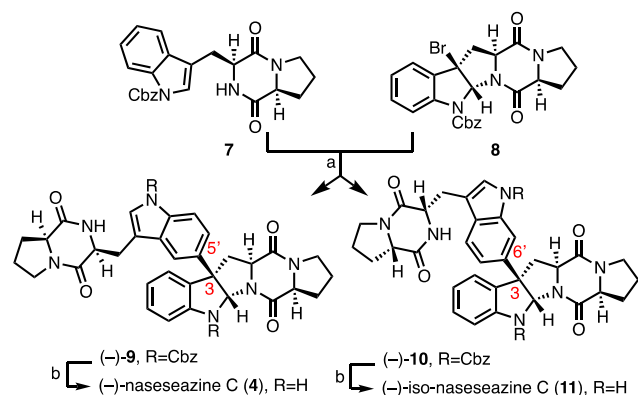


Figure 2. Products (-)-naseaezine C (**4**) and (+)-naseaezine B (**2**) formed from homodimerizations of **6** catalyzed by NascB and NznB, respectively.

To confirm the structure of (-)-naseaezine C (**4**), as well as resolve a discrepancy in the literature regarding its connectivity,^{10,11} the first total synthesis of (-)-naseaezine C (**4**) was performed, using the undirected Friedel–Crafts methodology described in the first-generation total synthesis of (+)-naseaezines A and B (Scheme 1).⁹ This route provided dimers with connectivities at both C5 and C6, encompassing the constitutional isomers proposed in both isolation reports. Detailed analysis of key HMBC correlations confirmed both the stereochemical configuration of the dimerization axis and connectivity of (-)-naseaezine C (**4**) at C5, as well as its C6 constitutional isomer, which we name (-)-*iso*-naseaezine C (**11**) based on previous precedent.⁹

Discovery and Functional Characterization of NzeB. Given the divergent site- and stereoselectivity displayed by NascB and NznB, we reasoned that comparative sequence analysis may uncover regions conferring selectivity to the dimerization cascade. Therefore, we sought to identify

Scheme 1. Total Synthesis of (–)-Naseseazine C (4) and (–)-iso-Naseseazine C (11)^a



^aReagents and conditions: (a) AgSbF₆, EtNO₂, (–)-9:(–)-10 = 1:1.4, 40%. (b) H₂/Pd/C, EtOH, 80% for (–)-4 and 84% for (–)-11.

additional DKP dimerases through SSN analysis.²⁴ Using NascB and NznB as *in silico* probes, along with known DKP functionalizing P450s from the thaxtomin (12),²⁵ pulcherri-
minic acid (13),²⁶ guanitryptmycin (14),^{27–30} bicyclomycin (15),^{31,32} and mycocyclusin (16)^{33,34} BGCs, we mined the NCBI database to identify candidate DKP dimerases. The resulting sequences were assembled into a SSN, which revealed a series of nodes clustered with NascB and NznB. From these data we identified a BGC, *nze*, consisting of a CDPS (NzeA, NCBI Accession #: WP_078874796.1) and P450 (NzeB, NCBI Accession #: WP_030888003.1) pair with high sequence homology to the *nasc* and *nzn* operons (Figure 3).

Heterologous expression of CDPS, NzeA, and analysis of its product profile identified 6 as the sole DKP metabolite (Supporting Information, Supplementary Figure S8). Subsequent NzeB-catalyzed homodimerization of 6 resulted in formation of three dimeric products: (–)-naseseazine C (4) as the major product and two minor products, (+)-naseseazine B (2) and unknown dimer 5 (Figure 4). Scale up and NMR characterization revealed 5 to be C–N-linked dimer (–)-aspergilazine A. This product distribution exemplifies the versatility of NzeB, as it is able to form site- and stereoisomeric Csp³–Csp²-linked dimeric products 2 and 4, as well as display variant chemoselectivity in the formation of Csp²–Nsp²-linked 5 (Figure 1b).^{10,12,14,15} The multifunctional nature of NzeB enables assembly of all currently known *Streptomyces*-derived DKP dimer scaffolds from a single biocatalyst.²⁰

Crystal Structure of NzeB. Given the catalytic flexibility displayed by NzeB, we sought to identify the molecular basis for this versatility using X-ray crystallography. We obtained high-resolution (1.5 Å) crystal structures of NzeB both in the presence and in the absence of native substrate 6 (Figure 5 and Supporting Information, Supplementary Figure 21). NzeB shares highest structural homology with the DKP functionalizing cytochrome P450, CYP121, which catalyzes oxidative intramolecular C–C bond formation to generate mycocyclusin (30% sequence ID, 2.4 Å RMSD, 45.8 Z-score).³⁵ NzeB adopts the prototypical triangular prism P450 fold, with only minor structural rearrangement between the ligand-bound and ligand-free forms (0.45 Å RMSD). Notably, both ligand-free and substrate-bound structures of NzeB display electron density for the conserved axial water 3 Å from the heme-iron.

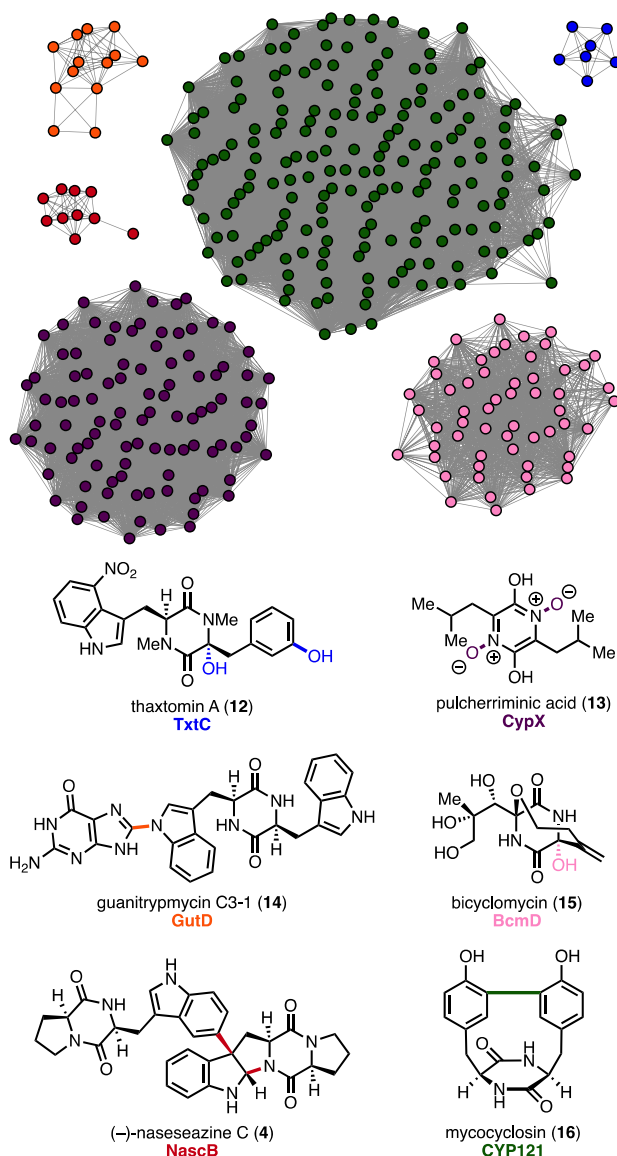


Figure 3. Sequence similarity network with clusters representing characterized DKP functionalizing P450s (clusters of unknown function and orphan nodes omitted for clarity).

The active site of the ligand-bound crystal structure of NzeB contains two units of 6, with each substrate occupying a discrete pocket in the active site. The DKP core and tryptophan side chain of the first substrate (Figure 5, light pink) are “sandwiched” by Gln68 and Phe391, with the DKP adopting a concave architecture. The concave pose of this substrate is suggestive of the polycyclic pyrroloindoline scaffold in the C–C-linked dimeric products 2 and 4, and as such we named this site the “cyclization site”. The proline side chain of the second substrate (Figure 5, mint green) is encapsulated by predominately hydrophobic residues (Gln75, Glu76, Phe177, Phe390, Phe391) and binds in an extended pose. This extended conformation positions the indole atoms that undergo C–H functionalization to form the dimeric axis (C5 for 4 and C6 for 2 and 5) proximal to the iron center of the heme cofactor; therefore, we named this the “dimerization site”.

Inspection of this complex revealed close packing of the substrate proline side chain in the dimerization site compared

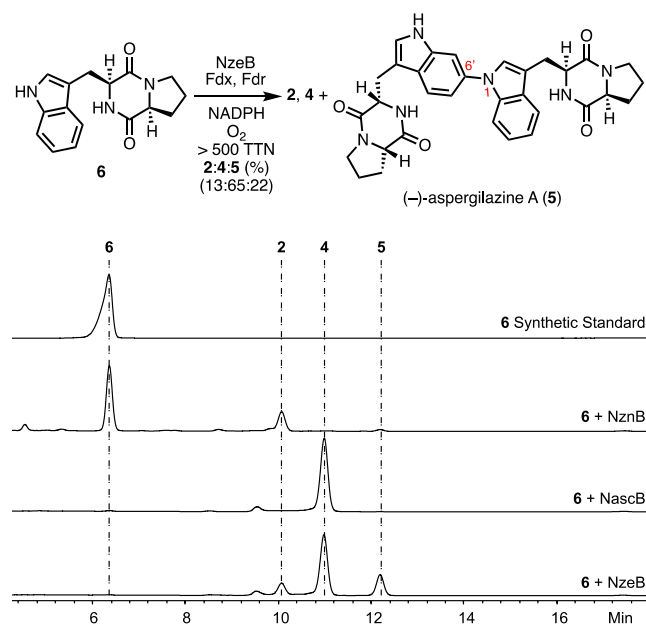


Figure 4. Products formed and HPLC traces (absorbance measured at 280 nm) from homodimerization of **6** by NzeB.

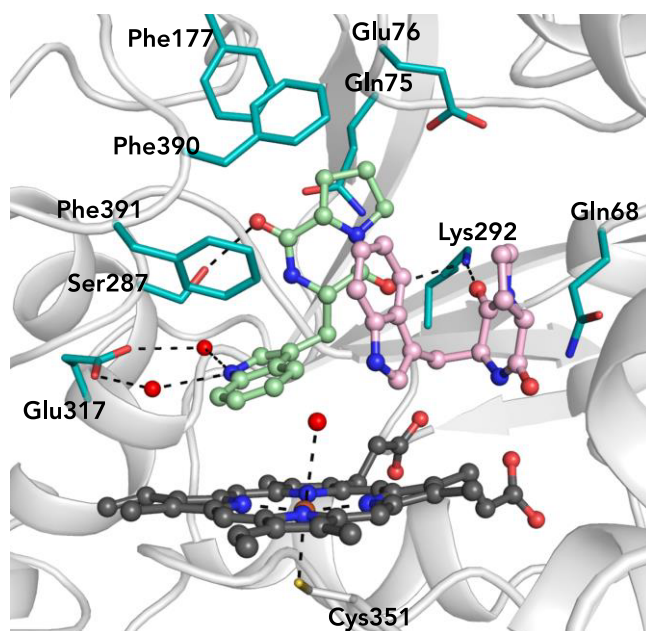


Figure 5. Active site of NzeB with concave substrate (light pink) bound in the “cyclization site” and extended substrate (mint green) bound in the “dimerization site”.

to the relatively open cavity surrounding the proline side chain in the cyclization site (Supporting Information, Supplementary Figure 22).

We reasoned that the more open cyclization site could selectively accommodate non-native substrates with bulkier side chains. We hypothesized that reactions involving heterodimerization of the larger substrate and the native DKP **6** would show pyrroloindoline ring formation in the larger substrate due to selective binding in the putative cyclization site.

To confirm the roles of the cyclization and dimerization sites, non-native DKP substrates with bulky side chains in place

of proline were synthesized and screened for their heterodimerization capacity with native substrate **6** (Supporting Information, Supplementary Figure 12). Although a variety of these substrates were capable of forming heterodimers with **6**, only heterodimerizations with *cyclo*(L-Trp-L-Trp) (**17**) resulted in the exclusive formation of a single heterodimeric product (Figure 6). Suppression of homodimeric product formation in

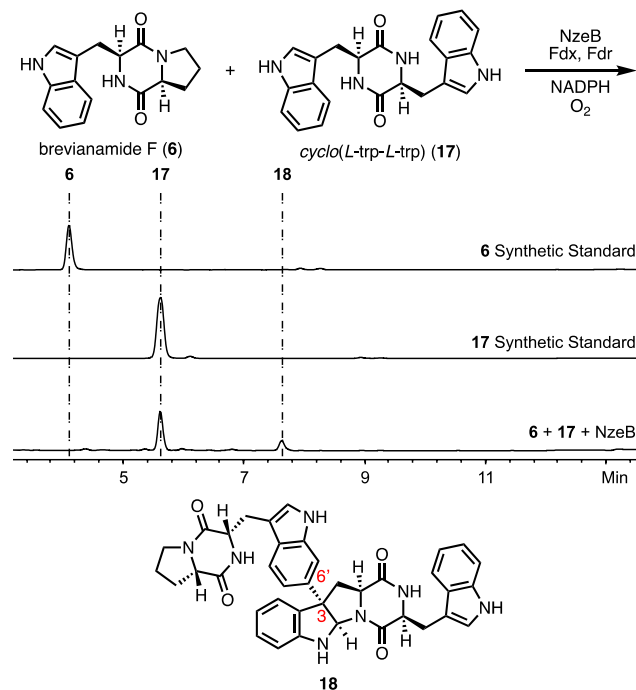


Figure 6. Heterodimerization reaction of NzeB with substrates **6** and **17**, HPLC traces (absorbance measured at 280 nm) from heterodimerization reactions, and structure of heterodimer **18**.

the presence of **17** and formation of a single heterodimer suggested that each substrate bound selectively and preferentially to a single DKP binding site.

This biocatalytic union of substrates **6** and **17** was performed on preparative scale and the structure of product **18** was determined. This result suggests that binding of **17** is limited to the cyclization site. We obtained a crystal structure of a mixed-ligand complex of NzeB with **17** binding in the cyclization site and **6** in the dimerization site, confirming our assignment of the functional roles of each of these active site pockets (Figure 7).

Site-Directed Mutagenesis of NzeB. To identify key residues that control chemoselectivity in the NzeB-catalyzed dimerization cascade, we examined the conservation of amino acid residues in both the cyclization and dimerization sites between NzeB, NascB, and NznB. As (–)-naseaezine C (**4**) is the major product of the NzeB and the exclusive product of NascB, we began our analysis by comparing the active sites of these two homologues. A single residue variation exists in the active sites of NascB and NzeB, (NascB: Ala 287, NzeB: Ser 287). In the ligand-bound structures of NzeB, Ser287 forms a hydrogen bond with the proline carbonyl of the substrate in the dimerization site (Figure 5). NzeB variant NzeB_{S287A} was generated and homodimerization reactions of **6** with this mutant protein resulted in exclusive formation of (–)-naseaezine C (**4**) (Figure 8). Functionally, this mutation completely

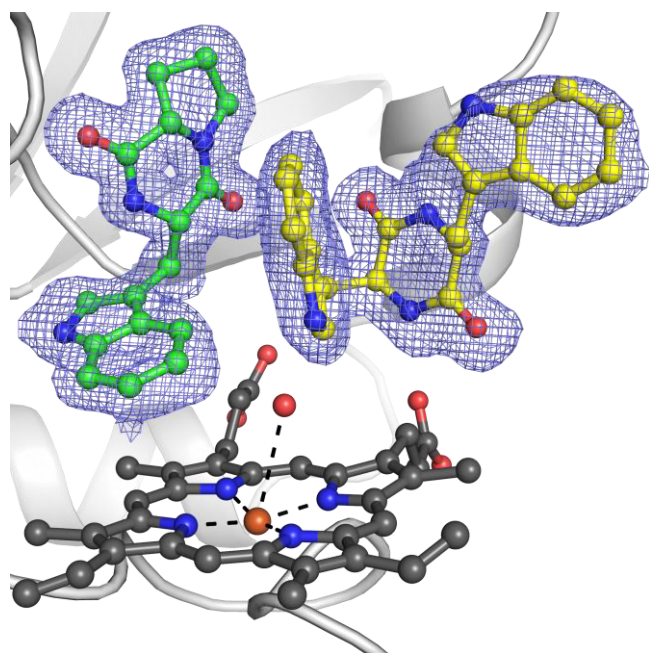


Figure 7. Crystal structure of NzeB heterocomplex with diketopiperazine 17 (yellow) bound in the cyclization site and diketopiperazine 6 (green) bound in the dimerization site, with omit map ($F_o - F_c$) displayed as blue mesh.

reconstituted the specificity of NascB and pinpointed Ser287 as a key determinant of dimerase selectivity.

To examine the generality of remodeling active sites to recapitulate selectivity, we compared the active site residues of (+)-naseasezine B (2) synthase, NznB, to NzeB. This revealed a single active site variation (NznB: Met68, NzeB: Gln68), located in the cyclization site. To assess the consequence of this variation, NzeB_{Q68M} was constructed and subjected to homodimerization reactions with 6. Rather than exclusive formation of 2, homodimerization reactions of 6 with NzeB_{Q68M} generated dimers 2, 4 and 5, as well as a new product, 11 (Figure 8). Scale up and characterization of this product revealed its identity as the C6 constitutional isomer of 2, (–)-iso-naseasezine C (11), based on comparison to synthetic standard (Scheme 1).⁹

To further profile the role of active site residues in controlling selectivity in the C–C/C–N bond-forming cascade, we performed alanine scanning mutagenesis on residues within 5 Å of either substrate. The resulting variants (S287A, Q75A, E76A, Q75A/E76A, F177A, E317A, F390A, F391A, and F390A/F391A) were profiled for catalytic activity and selectivity against native substrate 6 (Supporting Information, Supplementary Figure 13). All functional variants from this experiment displayed the same product profile as NzeB_{Q68M}, formation of 2, 4, 5 and 11, and differed only in the relative distributions of dimers. Given the increased formation of 11 across a variety of active site variants at various positions within the NzeB active site, we hypothesize that 11 results from loss of selectivity during the dimerization cascade.

NzeB Dimerization Mechanism and Quantum Mechanics Calculations. The first biomimetic total synthesis of the (+)-naseasezines proposed two alternative biosynthetic mechanisms for DKP dimer formation.⁹ Both mechanisms involved oxidative formation of a pyrroloindoline radical, but diverged on formation of the intermolecular C–C bond. The

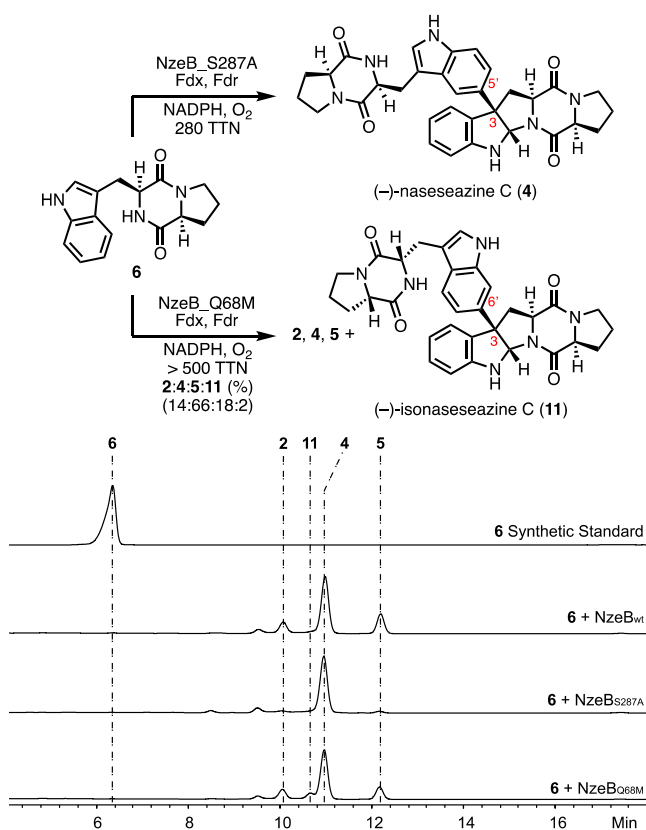


Figure 8. Homodimerization reactions of NzeB variants with 6 and corresponding HPLC traces. Minor peaks (around 9.5 min) are unrelated to either DKP substrate or dimeric products of the reaction.

first mechanism proposed directed pyrroloindoline radical coupling to a second DKP followed by oxidative re-aromatization to afford the non-symmetrical DKP scaffold. The second involved oxidation of the pyrroloindoline radical to a tertiary, benzylic cation followed by electrophilic aromatic substitution (EAS) to form the C–C bond. The reaction was completed by re-aromatization via deprotonation of the resulting Wheland intermediate to forge the dimeric axis. To shed light on the viability of these proposed mechanisms for intermolecular C–C bond formation, as well as the intermolecular C–N bond formation to give 5 (catalyzed by NzeB), we turned to QM calculations.

Given the proximity and orientation of the substrate in the cyclization site relative to the heme iron, we propose that the dimerization cascade is initiated by the oxidation of the substrate occupying the cyclization site. The mechanistic pathways (Figure 9) compare two possible modes for oxidative formation of the cyclized pyrroloindoline radical: hydrogen abstraction of (1) the proximal indole N–H bond or (2) the diketopiperazine N–H bond starting from separated brevianamide F (6) and P450 iron-oxo (Compound I). While indole N–H abstraction has a lower free energy barrier (path 1, 11.5 kcal/mol) than DKP N–H abstraction (path 2, 20.3 kcal/mol), both pathways are energetically plausible and subsequent steps have equal or higher free energy barriers. Furthermore, the diketopiperazyl radical (R-II) can cyclize directly to the pyrroloindoline radical (R-III) with a free energy barrier of 20.7 kcal/mol, while cyclization of the indolyl radical (R-I) would need to be accompanied by the net movement of a hydrogen atom from the DKP nitrogen to the indole nitrogen.

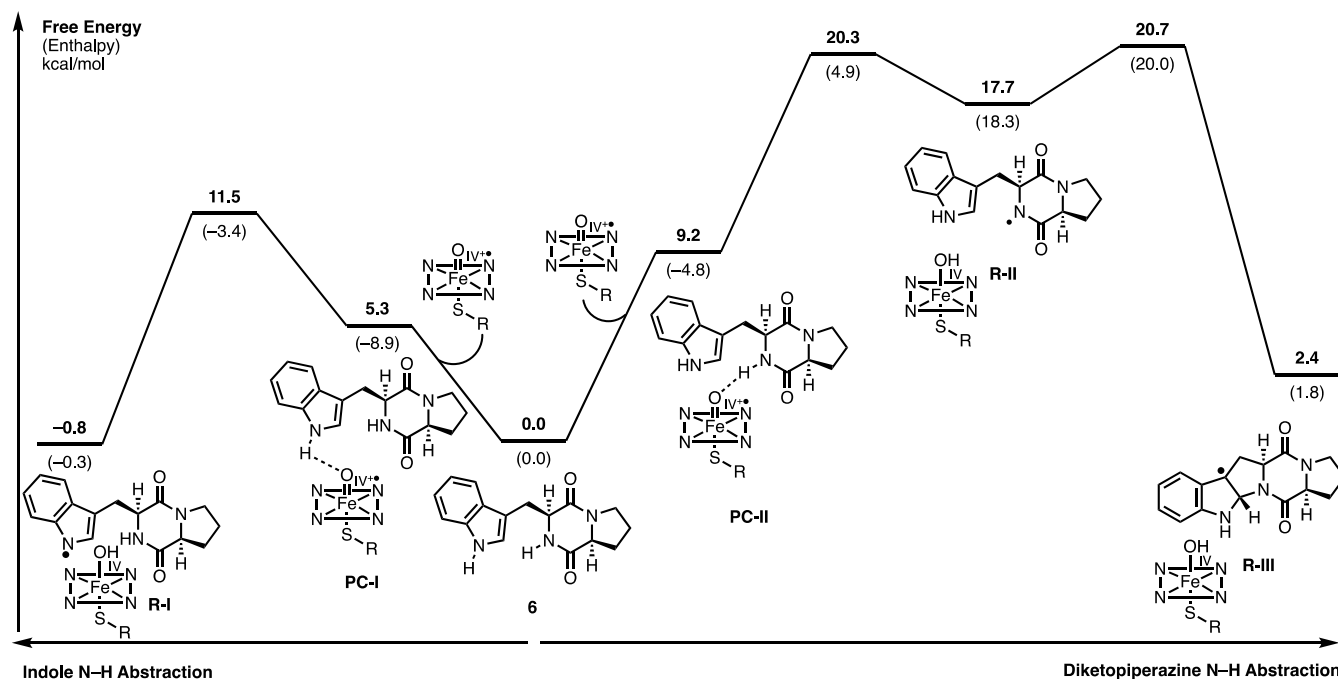


Figure 9. Computed mechanistic pathways for the formation of radical cyclized pyrroloindoline intermediate (**R-III**) via indole N–H abstraction (left) and DKP N–H abstraction (right).

A unique feature of the active site of NzeB is the hydrogen bonding of the DKP N–H to a heme propionate. We considered the heme propionate as a possible hydrogen relay to enable cyclization of indolyl radical (**R-I**); however, calculations using acetate as a surrogate for the heme propionate gave a large free energy barrier (30.8 kcal/mol). Furthermore, there is an overall absence of participant residues (both side chain and backbone atoms) in the active site to catalyze this proton transfer. Single-electron oxidation of indole to give a radical cation was also considered (Supporting Information, Supplementary Figure 23); however, given the high-energy radical cation intermediate (41.2 kcal/mol), we did not to pursue this pathway further. In both depicted N–H abstraction pathways, a significant contribution to the overall free energy barrier is the formation of a pre-complex (**PC**) between the heme iron-oxo species and the substrate N–H bond to be abstracted (Figure 9, **PC-I** and **PC-II**). Compared to separated starting materials (diketopiperazine **6** and the heme iron-oxo), the free energy for the formation of the **PC** for indole N–H abstraction (**PC-I**) and the **PC** for DKP N–H abstraction (**PC-II**) are 5.3 and 9.2 kcal/mol, respectively. However, as evidenced by the crystal structure of NzeB, both DKP substrates are oriented to facilitate **PC** formation, substantially lowering the free energy barrier for the entire transformation.

Following cyclization to the pyrroloindoline radical (Figure 10, **R-III**), formation of the C–C bond with a second unit of **6** (modeled as 3-methylindole) may occur through either a radical-mediated, or a cationic Friedel–Crafts mechanism. The depicted energetic landscape of both dimerization mechanisms led to formation of (–)-naseaezine **C** (**4**) (Figure 10); however, analogous computations of the pathway leading to the formation of (+)-naseaezine **B** (**2**) (Supporting Information), were qualitatively similar and led to the same conclusions. In the radical-mediated mechanism, C–C bond formation is the rate-determining step with a free energy

barrier of 24.3 kcal/mol, while in the cationic mechanism the C–C bond formation barrier of 20.3 kcal/mol falls slightly below the preceding cyclization. Both mechanisms remain viable given that these computations are performed on a model system in the absence of active site residues, and that the computed barriers are relatively close. However, NzeB lacks any side chains, backbone amides, or water molecules at an appropriate location in the active site to act as a general base for deprotonation of Wheland intermediate (**C-IV**) generated in the cationic Friedel–Crafts pathway. Furthermore, alanine mutants of distal acidic/basic residues in the active site (Q78, Q75, E76, E317), which could act to re-aromatize the Wheland intermediate (**C-IV**) are still catalytically competent for dimerization (Supporting Information, Supplementary Figure S23). However, re-aromatization of intermediate (**R-IV**) via H-atom abstraction by the iron(IV)-hydroxo (Compound II) is likely to be facile given the proximity of C5 and C6 to the heme cofactor. Alternatively, localization of C5 and C6 to the heme may also allow a single electron oxidation of intermediate **R-IV** to intermediate **C-IV** by Compound II followed by deprotonation and re-aromatization of **C-IV** by the resulting iron(III)-hydroxo species.

In addition to catalyzing intermolecular C–C bond formation, NzeB also catalyzes the intermolecular C–N bond formation to generate (–)-aspergilazine **A** (**5**). Given the viability of both a radical and cationic dimerization mechanism for NzeB-catalyzed C–C bond formation, we also compared the energetic landscapes of radical and cationic pathways for C–N bond formation starting from the uncyclized indolyl radical (Figure 11). In the radical-mediated pathway, the rate-limiting C–N bond formation has a thermodynamically plausible free energy barrier of 21.7 kcal/mol. In contrast, a potential cationic pathway would require oxidizing the indolyl radical (**R-V**) to a cation (**C-V**), which is thermodynamically demanding (45.5 kcal/mol). Given that the oxidation needed to access a cationic pathway is so highly

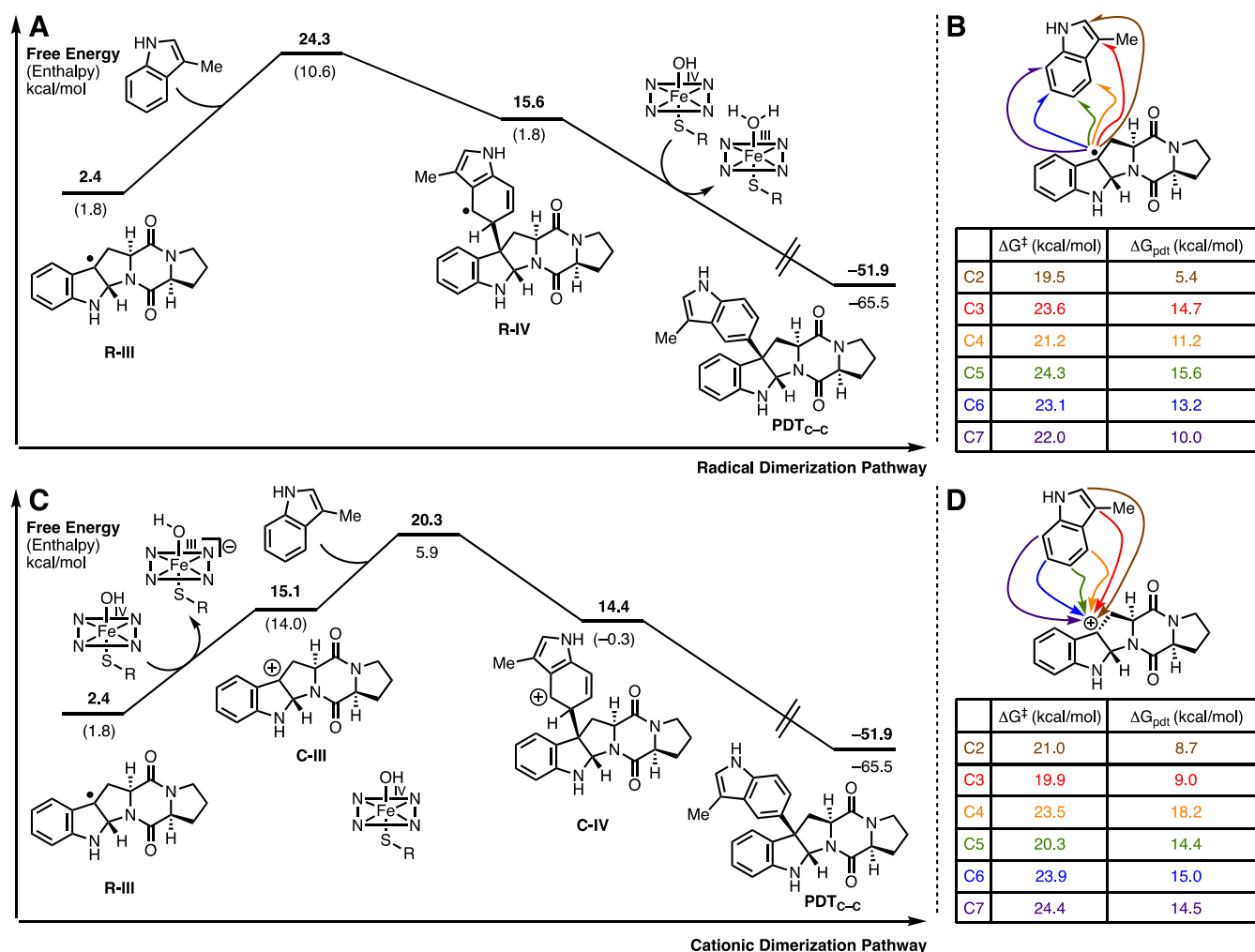


Figure 10. (A) Computed radical dimerization pathways to generate (–)-naseaeazine C model $\text{PDT}_{\text{C-C}}$. (B) Transition state and product free energies for both radical C–C bond formation at indole positions C2–C7 for $\text{PDT}_{\text{C-C}}$. (C) Computed cationic dimerization pathways to generate $\text{PDT}_{\text{C-C}}$. (D) Transition state and product free energies for cationic C–C bond formation at indole positions C2–C7 for $\text{PDT}_{\text{C-C}}$.

unfavorable, QM computations clearly support a radical-mediated pathway in the case of C–N bond formation.

The C–C bond-forming cascade catalyzed by NzeB results in the formation of two C–C-linked dimers **2** and **4**, as well as the C–N bond-linked dimer **5**. Computations were performed to examine the innate selectivity for cation and radical mechanisms for intermolecular C–C bond formation (Figure 10B,D), and analogous computations were performed for radical C–N bond formation (Figure 11B). For all computations relating to C–C and C–N bond formation, regardless of radical or cation mechanism, the observed dimerization site selectivity displayed by NzeB does not coincide with either the transition states or products that are lowest in free energy. Thus, these computations suggest that DKP-dimerization is catalyst-controlled rather than driven by the innate reactivity of the DKP substrates, with NzeB guiding the selective intermolecular C–C and intramolecular C–N bond formation.

DISCUSSION

Bacterial biosynthesis of the tryptophan-linked dimeric DKP alkaloids reveals Nature's ability to build molecular complexity from simple and abundant building blocks captured from primary metabolism. The production of DKP dimers from

both fungal and bacterial origin indicates the biological utility of this NP scaffold. Characterized fungal systems utilize non-ribosomal peptide synthetases (NRPS) to generate DKP monomers, while bacterial pathways generate the same monomers via CDPSs. Characterization of CDPS-associated gene clusters has decrypted the biosynthetic pathways for known NPs such as bicyclomycin,^{31,32} albonoursin,³⁶ and the drimentines.³⁷ Furthermore, these investigations have also identified new classes of NPs as well as unique DKP tailoring reactions including dehydrogenation,³⁶ C–H hydroxylation,^{31,32} C-, O-, and N-methylation,³⁸ prenylation and cyclization of terpenes,³⁷ and oxidative coupling with purine nucleobases.^{27–30} P450-mediated diketopiperazine dimerization further expands the repertoire of tailoring events found in CDPS-associated pathways.

The isolation of isomeric DKP dimers from a single organism is relatively common, which suggested two biosynthetic postulates: (1) a single dimerase with low selectivity enabled the assembly of isomeric products, or (2) a discrete dimerase was required for each coupling mode. Our results demonstrate there is no general axiom for bacterial biosynthesis of DKP dimers. P450s NascB and NznB from *Streptomyces* sp. CMB-MQ030 each catalyze a site- and stereoselective dimerization of **6** resulting in (–)-naseaeazine

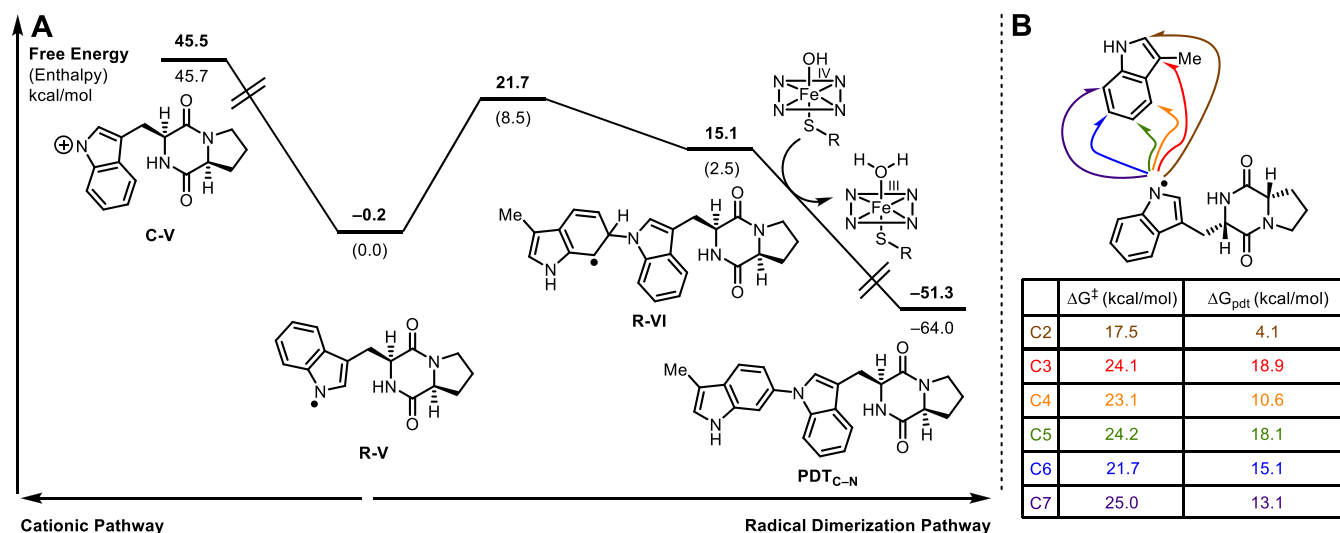


Figure 11. (A) Computed dimerization pathways for the formation of (–)-aspergilazine A (**5**): radical dimerization pathway and demonstration that a cationic pathway is thermodynamically unfavorable. (B) Transition state and product free energies for radical C–N bond formation at indole positions C2–C7 for (–)-aspergilazine A (**5**) model $\text{PDT}_{\text{C-N}}$.

C (**4**) and (+)-naseezazine B (**2**), respectively. By contrast, NzeB catalyzes both known bacterial C–C bond-forming dimerization modes generating **2** and **4**, while also displaying chemodivergent reactivity in the formation of C–N bond-linked dimer, (–)-aspergilazine A (**5**). While “multifunctional” bacterial P450s are well known,³⁹ this typically refers to the ability for a P450 to perform multiple monooxygenase reactions, such as hydroxylations at multiple sites,⁴⁰ hydroxylation and epoxidation activity,⁴¹ hydroxylation with continued reaction to higher oxidation states, or a combination of these processes. NzeB differs from these enzymes in its ability to mediate formation of site- and stereoisomeric C–C-linked products **1** and **2**, and also catalyzes direct $\text{Csp}^2\text{--H}$ amination in the formation of **5**. Recently, multiple reports have disclosed engineered enzymes capable of performing inter- and intramolecular C–H amination.^{42,43} However, this transformation is relatively rare in nature with few examples in the literature,^{44,45} and to our knowledge, NzeB represents the first structurally characterized, wild-type P450 to catalyze intermolecular $\text{Csp}^2\text{--H}$ amination.

We were able to obtain multiple high-resolution crystal structures, including NzeB ligand free, and in complex with select native and non-native substrates. Our biosynthetic hypotheses for the C–C bonded dimers **2** and **4** were supported by direct observation of the NzeB–DKP complex, with the enzyme possessing two distinct binding sites to pre-organize substrates for diastereoselective cyclization as well as site-selective C–C bond formation. Synthesis and screening of non-native substrates for their selective heterodimerization enabled validation of the cyclization and dimerization sites. Moreover, we were able to capture a complex of NzeB with sterically differentiated ligands, brevianamide F (**6**) and *cyclo*(L-Trp–L-Trp) (**17**), which unambiguously confirmed our assignment of roles to each DKP binding site. The connectivity and configuration of the dimerization axis of heterodimeric product **18** is identical to that of the NzeB minor product (+)-naseezazine B (**2**) and is consistent with the binding mode of the two monomeric precursors (Figure 7). We hypothesize this is due to the markedly larger side chain of **17** which occupies the additional channel in the cyclization site that the

native substrates may occupy during catalysis in order for both substrate and enzyme to undergo the conformational changes necessary to access either the (–)-naseezazine C or (–)-aspergilazine A scaffolds. Notably, all other structurally characterized DKP functionalizing CYP450s (CYP121,³⁴ CypX,²⁶ and TxtC²⁵) bind DKPs in the analogous position of the active site, which corresponds to the cyclization site of NzeB. The dimerization site of NzeB is a previously unobserved pocket for DKPs in P450s and accounts for an approximate doubling of the active site volume of NzeB (832 Å³) relative to other DKP binding CYP450s.⁴⁶ Previous efforts in the characterization of NascB demonstrated that in heterodimerization reactions the bulky substrates undergo cyclization to form pyrroloindoline functionality.²¹ Our structural investigations of NzeB and assignment of the role of the dimerization site provides a structural basis for this phenomenon. Furthermore, identifying the side chains in the dimerization site that encapsulate the proline moiety of its substrate paves the way to reshaping this binding pocket. Engineering DKP dimerases to enable coupling of DKPs with larger side chains may enable formation of products similar to those isolated from fungi.^{47–49}

Mutagenesis of the active site of NzeB provided further insight into how NzeB guides the site- and stereoselective formation of C–C and C–N bonds. Comparison of the active site residues of NzeB to NascB and NznB, identified that each possessed a single, active site variation (NzeB: Ser287, NascB: Ala287). To examine the role of this active site variation in selectivity the NzeB_{S287A} variant was generated and homo-dimerization reactions with native substrate **6**, NzeB_{S287A} exclusively generates (–)-naseezazine C (**4**) completely reconstituting the selectivity of NascB with a single substitution. Given this result, we reasoned that through homologous substitution of the active site variation between NznB and NzeB, located in the cyclization site (NzeB: Gln68 vs NznB: Met68) would provide access to a surrogate of NznB. The prospect of engineering the dimerization selectivity of NzeB to recapitulate that of NznB is particularly attractive, as wildtype NznB has low TTN relative to NascB and NzeB (Figure 2). However, counter to the results from NzeB_{S287A}

dimerization reactions, NzeB_{Q68M} did not recapitulate the observed NznB selectivity but led to the formation of previously unobserved dimer (–)-iso-naseaezine C (**11**).

These results demonstrate that the selectivity of a given DKP dimerase cannot be reconstituted simply by interchanging active site residues, and therefore the selectivity of NzeB and its homologues must also be determined by residues outside the active site motif. Thus, we propose that the observed crystal structure represents one of many catalytically relevant conformations, and that the active site must undergo conformational remodeling during catalysis, with the selectivity of the dimerization reaction being influenced by a combination of residues both inside and outside of the active site. Notably, the overall sequence conservation between NzeB and NznB (69% identity) is lower compared to that of NzeB and NascB (87% identical), and conformational changes during catalysis could affect these regions of low conservation present in NznB, which are absent in NascB. This could result in a shift into or near the active site that modulates the selectivity of the dimerization reaction. Support for active site remodeling is best represented by the discrepancy between the conformation of substrates in the ligand bound crystal structure of NzeB and the observed product distribution from the dimerization reaction. In the ligand-bound structure of NzeB, both substrates are apparently pre-organized toward formation of (+)-naseaezine B (**2**), despite this dimer being a minor product. To access conformations that would lead to the major product, DKP dimer **4**, the orientation of substrate in the cyclization site would require inversion from a concave to convex posture. Furthermore, in spin-shift binding assays of NzeB,⁵⁰ substrate **6** was able to displace the water molecules axial to the heme to induce a low- to high-spin transition of the heme iron indicative of direct binding to the active site (Supporting Information, Supplementary Figure 10). However, this axial water is present in both the ligand-bound and ligand-free crystal structures at high-occupancy, suggesting that solution-phase dynamics provide access to currently unobserved conformations. In addition, it is well-precedented that binding of electron transport proteins alters the architecture of cytochromes P450 during catalysis.⁵¹ Given these factors we propose residues and regions distal to the active site are influencing selectivity and catalysis, and efforts to identify them in bacterial DKP dimerases are currently underway.

Finally, QM computations provided new insights into NzeB's ability to catalyze formation of C–C and C–N bonds. Relative to C–O bond-forming P450s, far fewer C–C bond-forming P450s have been structurally characterized; examples include both intramolecular such as StaP (PDB: 2Z3U)⁵² and CYP121 (PDB: 3G5H),³⁴ as well as intermolecular such as CYP158A1/CYP158A2 (PDB: 1T93, 2D09),^{53,54} and recently the intramolecular C–N bond-forming indolactam cytochrome P450 (PDB: 6J82).⁵⁵ A common feature among the characterized intra- and intermolecular C–C/C–N bond-forming P450s is that one of the reactive atoms participating in C–C/C–N coupling is proximal to the heme iron while the second reactive atom is distal, and oxidation of this distal atom is facilitated by a basic amino acid residue or water via proton-coupled electron transfer (PCET).^{56,57} However, unique to the structure of NzeB, reactive atoms from both ligands are positioned within 4 Å of the axial water ligand, indicating the possibility for direct oxidation of both ligands catalyzed by the heme without intervention of neighboring residues or solvent (Figure 5).

Regardless of mechanism, they indicate that a major role of NzeB is to orient substrates for reaction with the heme iron oxo species. By facilitating the formation of a pre-complex, NzeB can reduce the free energy barrier for the overall transformation by up to 5–10 kcal/mol. Previous investigations into the mechanism for C–C bond formation excluded a cationic mechanism due to reactivity of electron-poor substrates.²¹ However, the utilization of electronically differentiated substrates for mechanistic studies such as Hammett plots is well represented in the literature, including enzymatic electrophilic aromatic substitution (EAS)-type reactions.⁵⁸ Our computations demonstrate that both radical-mediated and cationic pathways are thermodynamically plausible for the formation of C–C bonds, with rate-determining transition states of comparable free energies. While there are residues present in the active site that could be positioned for deprotonation of Wheland intermediates, their distance appears too great from the appropriate carbon atoms, and mutation of these residues to alanine does not prevent dimerization. While either N–H abstraction from the indole or DKP could initiate the dimerization cascade, abstraction of the indole N–H would require multiple proton transfer steps in order to allow for cyclization of the pyrroloindoline. The active site of NzeB has a general lack of basic residues as well as active site waters that could act as proton shuttles. Furthermore, distal residues that could be implicated in proton transfer were mutagenized to alanine, and the resulting variants were still catalytically competent repudiating their side chains' participation in proton transfer during catalysis. Taken together we propose that in the case of a C–C-linked dimer, the dimerization cascade is likely initiated by abstraction of the DKP N–H. Given this lack of residues or active site water molecules, which would also be necessary to deprotonate Wheland intermediates generated from a cationic dimerization pathway, and the localization of reactive atoms from both substrates proximal to the heme iron, we propose that C–C bond-forming dimerization modes likely also proceed via a radical-mediated mechanism. However, re-aromatization of the C–C bond-formed radical intermediate may occur through either concerted HAT or stepwise PCET by Compound II. In contrast, computations relating to formation of the intermolecular C–N bond in (–)-aspergilazine A (**5**) demonstrate that the cationic pathway is thermodynamically prohibitive and the radical-mediated mechanism is strongly favored. To examine the site selectivity of NzeB we also explored computationally the innate selectivity for C–C and C–N bond-forming dimerizations. These computations demonstrate that regardless of cationic or radical mechanism, NzeB overrides the innate selectivity for C–C and C–N bond formation and the connectivity of the product dimers **2**, **4**, and **5** are the result of a catalyst-controlled dimerization, rather than the reactivity of the DKP substrates themselves. Further analysis is required to unambiguously exclude other mechanisms for C–C and C–N bond formation and is currently underway.

CONCLUSION

We identified and characterized the chemically versatile cytochrome P450, NzeB, capable of generating both (+)- and (–)-naseaezine frameworks as well as the non-cyclized Csp²–Nsp²-linked (–)-aspergilazine dimeric scaffold. To gain insight into the molecular basis for its versatility, we obtained the first high-resolution crystal structure of a DKP dimerization

P450 in complex with select native and non-native substrates. This structural information revealed a previously unrecognized DKP binding site compared to other reported crystal structures of DKP functionalizing P450s.^{25,26,34} NzeB is distinguished from the flavin dimerase in its relatively close binding of DKP substrates proximal (within 4 Å) to the heme cofactor. Site-directed mutagenesis of NzeB demonstrated that active-site residues are not the sole drivers of site-, stereo-, and chemoselectivity. Tandem structural, biochemical, and computational approaches explored the viability of cationic and radical mechanisms in intermolecular C–C bond formation and provided evidence for a likely radical mechanism for intermolecular C–N bond formation. The structural, biochemical, and computational characterization of NzeB provides the first insight into the molecular basis for DKP dimerization. This new information opens the door to engineering NzeB and its functional homologues for improved total turnover number, broadened substrate scope, and new non-natural dimerization modes, furthering our efforts toward a catalyst-controlled method for DKP dimerization.

■ ASSOCIATED CONTENT

SI Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/jacs.0c06312>.

Full experimental and computational details, NMR spectra, tables, and figures (PDF)

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Notes

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■ REFERENCES

- (1) Hino, T.; Nakagawa, M., Chemistry and Reactions of Cyclic Tautomers of Tryptamines and Tryptophans. In *The Alkaloids: Chemistry and Pharmacology*; Brossi, A., Ed.; Academic Press: New York, 1989; Vol. 34, pp 1–75.
- (2) Anthoni, U.; Christophersen, C.; Nielsen, P. H., Naturally Occurring Cyclotryptophans and Cyclotryptamines. In *Alkaloids: Chemical and Biological Perspectives*; Pelletier, S. W., Ed.; Pergamon: London, 1999; Vol. 13, pp 163–236.
- (3) Borthwick, A. D. 2,5-Diketopiperazines: synthesis, reactions, medicinal chemistry, and bioactive natural products. *Chem. Rev.* **2012**, *112* (7), 3641–3716.
- (4) Tadano, S.; Ishikawa, H. Synthesis of tryptophan-based dimeric diketopiperazine alkaloids using bioinspired reactions. *Synlett* **2014**, *25* (2), 157–162.
- (5) Kim, J.; Movassaghi, M. Biogenetically-inspired total synthesis of epidithiodiketopiperazines and related alkaloids. *Acc. Chem. Res.* **2015**, *48* (4), 1159–1171.
- (6) Springer, J. P.; Büchi, G.; Kobbe, B.; Demain, A. L.; Clardy, J. The structure of ditryptophenaline - a new metabolite of *Aspergillus flavus*. *Tetrahedron Lett.* **1977**, *18* (28), 2403–2406.
- (7) Movassaghi, M.; Schmidt, M. A.; Ashenhurst, J. A. Concise total synthesis of (+)-WIN 64821 and (–)-ditryptophenaline. *Angew. Chem., Int. Ed.* **2008**, *47* (8), 1485–1487.
- (8) Raju, R.; Piggott, A. M.; Conte, M.; Aalbersberg, W. G.; Feussner, K.; Capon, R. J. Naseseazines A and B: a new dimeric diketopiperazine framework from a marine-derived actinomycete, *Streptomyces* sp. *Org. Lett.* **2009**, *11* (17), 3862–3865.
- (9) Kim, J.; Movassaghi, M. Concise total synthesis and stereochemical revision of (+)-naseseazines A and B: regioselective arylative dimerization of diketopiperazine alkaloids. *J. Am. Chem. Soc.* **2011**, *133* (38), 14940–14943.
- (10) Xiong, Z. Q.; Liu, Q. X.; Pan, Z. L.; Zhao, N.; Feng, Z. X.; Wang, Y. Diversity and bioprospecting of culturable actinomycetes from marine sediment of the Yellow Sea, China. *Arch. Microbiol.* **2015**, *197* (2), 299–309.
- (11) Buedenbender, L.; Grkovic, T.; Duffy, S.; Kurtboke, D. I.; Avery, V. M.; Carroll, A. R. Naseseazine C, a new anti-plasmodial

dimeric diketopiperazine from a marine sediment derived *Streptomyces* sp. *Tetrahedron Lett.* **2016**, 57 (52), 5893–5895.

(12) Cai, S. X.; Kong, X. L.; Wang, W.; Zhou, H. N.; Zhu, T. J.; Li, D. H.; Gu, Q. Q. Aspergilazine A, a diketopiperazine dimer with a rare N-1 to C-6 linkage, from a marine-derived fungus *Aspergillus taichungensis*. *Tetrahedron Lett.* **2012**, 53 (21), 2615–2617.

(13) Kieffer, M. E.; Chuang, K. V.; Reisman, S. E. Copper-catalyzed diastereoselective arylation of tryptophan derivatives: total synthesis of (+)-naseaeazines A and B. *J. Am. Chem. Soc.* **2013**, 135 (15), 5557–5560.

(14) Boyd, E. M.; Sperry, J. Total synthesis of (–)-aspergilazine A. *Org. Lett.* **2014**, 16 (19), 5056–5059.

(15) Chuang, K. V.; Kieffer, M. E.; Reisman, S. E. A mild and general Larock indolization protocol for the preparation of unnatural tryptophans. *Org. Lett.* **2016**, 18 (18), 4750–4753.

(16) Alonso, I.; Alvarez, R.; de Lera, A. R. Indole-indole Ullmann cross-coupling for CAr–N bond formation: Total synthesis of (–)-aspergilazine A. *Eur. J. Org. Chem.* **2017**, 2017 (33), 4948–4954.

(17) Loach, R. P.; Fenton, O. S.; Movassaghi, M. Concise total synthesis of (+)-asperazine, (+)-pestalazine A, and (+)-iso-pestalazine A. Structure revision of (+)-pestalazine A. *J. Am. Chem. Soc.* **2016**, 138 (3), 1057–1064.

(18) Nelson, B. M.; Loach, R. P.; Schiesser, S.; Movassaghi, M. Concise total synthesis of (+)-asperazine A and (+)-pestalazine B. *Org. Biomol. Chem.* **2018**, 16 (2), 202–207.

(19) Saruwatari, T.; Yagishita, F.; Mino, T.; Noguchi, H.; Hotta, K.; Watanabe, K. Cytochrome P450 as dimerization catalyst in diketopiperazine alkaloid biosynthesis. *ChemBioChem* **2014**, 15 (5), 656–659.

(20) While this paper was in proofs the following article was published which expanded the known repertoire of bacterial DKP dimer scaffolds: Liu, J.; Xie, X.; Li, S.-M. Increasing cytochrome P450 enzyme diversity by identification of two distinct cyclodipeptide dimers. *Chem. Commun.* **2020**, DOI: 10.1039/D0CC04772D.

(21) Tian, W.; Sun, C.; Zheng, M.; Harmer, J. R.; Yu, M.; Zhang, Y.; Peng, H.; Zhu, D.; Deng, Z.; Chen, S. L.; Mobli, M.; Jia, X.; Qu, X. Efficient biosynthesis of heterodimeric C(3)-aryl pyrroloindoline alkaloids. *Nat. Commun.* **2018**, 9 (1), 4428.

(22) Yu, H.; Li, S. M. Two cytochrome P450 enzymes from *Streptomyces* sp. NRRL S-1868 catalyze distinct dimerization of tryptophan-containing cyclodipeptides. *Org. Lett.* **2019**, 21 (17), 7094–7098.

(23) Due to the limited availability of NznB protein, we were only able to perform scale-up reactions for characterization of (+)-naseaeazine B. Analytical reactions were performed to detect formation of (+)-naseaeazine A, and its structure was assigned by analogy and via comparison of retention time to an analytical standard (Supporting Information, Supplementary Figure 11).

(24) Gerlt, J. A.; Bouvier, J. T.; Davidson, D. B.; Imker, H. J.; Sadkhin, B.; Slater, D. R.; Whalen, K. L. Enzyme Function Initiative-Enzyme Similarity Tool (EFI-EST): A web tool for generating protein sequence similarity networks. *Biochim. Biophys. Acta, Proteins Proteomics* **2015**, 1854 (8), 1019–1037.

(25) Alkhalaf, L. M.; Barry, S. M.; Lea, D.; Gallo, A.; Griffiths, D.; Lewandowski, J. R.; Fulop, V.; Challis, G. L. Binding of distinct substrate conformations enables hydroxylation of remote sites in thaxtomin D by cytochrome P450 TxtC. *J. Am. Chem. Soc.* **2019**, 141 (1), 216–222.

(26) Cryle, M. J.; Bell, S. G.; Schlichting, I. Structural and biochemical characterization of the cytochrome P450 CypX (CYP134A1) from *Bacillus subtilis*: a cyclo-L-leucyl-L-leucyl dipeptide oxidase. *Biochemistry* **2010**, 49 (34), 7282–7296.

(27) Yu, H.; Xie, X.; Li, S. M. Coupling of guanine with cyclo-L-Trp-L-Trp mediated by a cytochrome P450 homologue from *Streptomyces purpureus*. *Org. Lett.* **2018**, 20 (16), 4921–4925.

(28) Shi, J.; Xu, X.; Zhao, E. J.; Zhang, B.; Li, W.; Zhao, Y.; Jiao, R. H.; Tan, R. X.; Ge, H. M. Genome mining and enzymatic total biosynthesis of purincyclamide. *Org. Lett.* **2019**, 21 (17), 6825–6829.

(29) Yu, H.; Xie, X.; Li, S. M. Coupling of cyclo-L-Trp-L-Trp with hypoxanthine increases the structure diversity of guanitryptamycins. *Org. Lett.* **2019**, 21 (22), 9104–9108.

(30) Liu, J.; Xie, X.; Li, S. M. Guanitryptamycin biosynthetic pathways imply cytochrome P450 mediated regio- and stereospecific guaninyl-transfer reactions. *Angew. Chem., Int. Ed.* **2019**, 58 (33), 11534–11540.

(31) Patteson, J. B.; Cai, W.; Johnson, R. A.; Santa Maria, K. C.; Li, B. Identification of the biosynthetic pathway for the antibiotic bicyclomycin. *Biochemistry* **2018**, 57 (1), 61–65.

(32) Meng, S.; Han, W.; Zhao, J.; Jian, X. H.; Pan, H. X.; Tang, G. L. A six-oxidase cascade for tandem C–H bond activation revealed by reconstitution of bicyclomycin biosynthesis. *Angew. Chem., Int. Ed.* **2018**, 57 (3), 719–723.

(33) Vetting, M. W.; Hegde, S. S.; Blanchard, J. S. The structure and mechanism of the *Mycobacterium tuberculosis* cyclodityrosine synthetase. *Nat. Chem. Biol.* **2010**, 6 (11), 797–799.

(34) Belin, P.; Le Du, M. H.; Fielding, A.; Lequin, O.; Jacquet, M.; Charbonnier, J. B.; Lecoq, A.; Thai, R.; Courcon, M.; Masson, C.; Dugave, C.; Genet, R.; Pernodet, J. L.; Gondry, M. Identification and structural basis of the reaction catalyzed by CYP121, an essential cytochrome P450 in *Mycobacterium tuberculosis*. *Proc. Natl. Acad. Sci. U. S. A.* **2009**, 106 (18), 7426–7431.

(35) Holm, L. Benchmarking fold detection by DaliLite v.5. *Bioinformatics* **2019**, 35 (24), 5326–5327.

(36) Lautru, S.; Gondry, M.; Genet, R.; Pernodet, J. L. The albonoursin gene cluster of *S. noursei* biosynthesis of diketopiperazine metabolites independent of nonribosomal peptide synthetases. *Chem. Biol.* **2002**, 9 (12), 1355–1364.

(37) Yao, T.; Liu, J.; Liu, Z.; Li, T.; Li, H.; Che, Q.; Zhu, T.; Li, D.; Gu, Q.; Li, W. Genome mining of cyclodipeptide synthases unravels unusual tRNA-dependent diketopiperazine-terpene biosynthetic machinery. *Nat. Commun.* **2018**, 9 (1), 4091.

(38) Giessen, T. W.; von Tesmar, A. M.; Marahiel, M. A. A tRNA-dependent two-enzyme pathway for the generation of singly and doubly methylated ditryptophan 2,5-diketopiperazines. *Biochemistry* **2013**, 52 (24), 4274–4283.

(39) Podust, L. M.; Sherman, D. H. Diversity of P450 enzymes in the biosynthesis of natural products. *Nat. Prod. Rep.* **2012**, 29 (10), 1251–1266.

(40) Sherman, D. H.; Li, S.; Yermalitskaya, L. V.; Kim, Y.; Smith, J. A.; Waterman, M. R.; Podust, L. M. The structural basis for substrate anchoring, active site selectivity, and product formation by P450 PikC from *Streptomyces venezuelae*. *J. Biol. Chem.* **2006**, 281 (36), 26289–26297.

(41) Carlson, J. C.; Li, S.; Gunatilleke, S. S.; Anzai, Y.; Burr, D. A.; Podust, L. M.; Sherman, D. H. Tirandamycin biosynthesis is mediated by co-dependent oxidative enzymes. *Nat. Chem.* **2011**, 3 (8), 628–633.

(42) McIntosh, J. A.; Coelho, P. S.; Farwell, C. C.; Wang, Z. J.; Lewis, J. C.; Brown, T. R.; Arnold, F. H. Enantioselective intramolecular C–H amination catalyzed by engineered cytochrome P450 enzymes in vitro and in vivo. *Angew. Chem., Int. Ed.* **2013**, 52 (35), 9309–9312.

(43) Singh, R.; Bordeaux, M.; Fasan, R. P450-catalyzed intramolecular sp(3) C–H amination with arylsulfonyl azide substrates. *ACS Catal.* **2014**, 4 (2), 546–552.

(44) Baunach, M.; Ding, L.; Bruhn, T.; Bringmann, G.; Hertweck, C. Regiodivergent N–C and N–N aryl coupling reactions of indoloterpenes and cycloether formation mediated by a single bacterial flavoenzyme. *Angew. Chem., Int. Ed.* **2013**, 52 (34), 9040–9043.

(45) Tsutsumi, H.; Katsuyama, Y.; Izumikawa, M.; Takagi, M.; Fujie, M.; Satoh, N.; Shin-Ya, K.; Ohnishi, Y. Unprecedented cyclization catalyzed by a cytochrome P450 in benzastatin biosynthesis. *J. Am. Chem. Soc.* **2018**, 140 (21), 6631–6639.

(46) Tian, W.; Chen, C.; Lei, X.; Zhao, J.; Liang, J. CASTp 3.0: computed atlas of surface topography of proteins. *Nucleic Acids Res.* **2018**, 46 (W1), W363–W367.

- (47) Li, X. B.; Li, Y. L.; Zhou, J. C.; Yuan, H. Q.; Wang, X. N.; Lou, H. X. A new diketopiperazine heterodimer from an endophytic fungus *Aspergillus niger*. *J. Asian Nat. Prod. Res.* **2015**, *17* (2), 182–187.
- (48) Ding, G.; Jiang, L.; Guo, L.; Chen, X.; Zhang, H.; Che, Y. Pestalazines and pestalamides, bioactive metabolites from the plant pathogenic fungus *Pestalotiopsis theae*. *J. Nat. Prod.* **2008**, *71* (11), 1861–1865.
- (49) Varoglu, M.; Corbett, T. H.; Valeriote, F. A.; Crews, P. Asperazine, a selective cytotoxic alkaloid from a sponge-derived culture of *Aspergillus niger*. *J. Org. Chem.* **1997**, *62* (21), 7078–7079.
- (50) Noble, M. A.; Miles, C. S.; Chapman, S. K.; Lysek, D. A.; MacKay, A. C.; Reid, G. A.; Hanzlik, R. P.; Munro, A. W. Roles of key active-site residues in flavocytochrome P450 BM3. *Biochem. J.* **1999**, *339* (2), 371–379.
- (51) Tripathi, S.; Li, H.; Poulos, T. L. Structural basis for effector control and redox partner recognition in cytochrome P450. *Science* **2013**, *340* (6137), 1227–1230.
- (52) Makino, M.; Sugimoto, H.; Shiro, Y.; Asamizu, S.; Onaka, H.; Nagano, S. Crystal structures and catalytic mechanism of cytochrome P450 StaP that produces the indolocarbazole skeleton. *Proc. Natl. Acad. Sci. U. S. A.* **2007**, *104* (28), 11591–11596.
- (53) Zhao, B.; Guengerich, F. P.; Bellamine, A.; Lamb, D. C.; Izumikawa, M.; Lei, L.; Podust, L. M.; Sundaramoorthy, M.; Kalaitzis, J. A.; Reddy, L. M.; Kelly, S. L.; Moore, B. S.; Stec, D.; Voehler, M.; Falck, J. R.; Shimada, T.; Waterman, M. R. Binding of two flavin substrate molecules, oxidative coupling, and crystal structure of *Streptomyces coelicolor* A3(2) cytochrome P450 158A2. *J. Biol. Chem.* **2005**, *280* (12), 11599–11607.
- (54) Zhao, B.; Guengerich, F. P.; Voehler, M.; Waterman, M. R. Role of active site water molecules and substrate hydroxyl groups in oxygen activation by cytochrome P450 158A2: a new mechanism of proton transfer. *J. Biol. Chem.* **2005**, *280* (51), 42188–42197.
- (55) He, F.; Mori, T.; Morita, I.; Nakamura, H.; Alblova, M.; Hoshino, S.; Awakawa, T.; Abe, I. Molecular basis for the P450-catalyzed C–N bond formation in indolactam biosynthesis. *Nat. Chem. Biol.* **2019**, *15* (12), 1206–1213.
- (56) Dumas, V. G.; Defelipe, L. A.; Petruk, A. A.; Turjanski, A. G.; Marti, M. A. QM/MM study of the C–C coupling reaction mechanism of CYP121, an essential cytochrome p450 of *Mycobacterium tuberculosis*. *Proteins: Struct., Funct., Genet.* **2014**, *82* (6), 1004–1021.
- (57) Wang, Y.; Chen, H.; Makino, M.; Shiro, Y.; Nagano, S.; Asamizu, S.; Onaka, H.; Shaik, S. Theoretical and experimental studies of the conversion of chromopyrrolic acid to an antitumor derivative by cytochrome P450 StaP: the catalytic role of water molecules. *J. Am. Chem. Soc.* **2009**, *131* (19), 6748–6762.
- (58) Gebler, J. C.; Woodside, A. B.; Poulter, C. D. Dimethylallyl-tryptophan synthase - an enzyme-catalyzed electrophilic aromatic substitution. *J. Am. Chem. Soc.* **1992**, *114* (19), 7354–7360.