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We report the characterization of the penilumamide biosynthetic cluster from *Aspergillus flavipes* CNL-338. *In vitro* reconstitution experiments demonstrated that three nonribosomal peptide synthetases are required for constructing the tripeptide and studies with dissected adenylation domains allowed for the first biochemical characterization of a domain that selects a pterin-derived building block.

In the post-genomic era, natural product discovery has been enhanced through various genome mining and bioinformatic platforms. Strains that are likely to generate unique chemistry are prioritized due to our ability to identify putative biosynthetic clusters and predict building blocks and potential downstream modifications. Natural products produced by marine-derived microorganisms continue to be a treasure trove of novel bioactive small molecules,^{1–3} including those of the well-established nonribosomal peptide class. Although the catalytic domains and mechanisms for peptide biosynthesis are similar between bacterial and fungal nonribosomal peptide synthetases (NRPSs), the ability to predict substrate specificity of building blocks in fungal systems is lacking compared to bacterial counterparts.⁴ Thus, to enhance our ability to accurately predict nonribosomal peptides in fungal genomes, additional NRPS machinery needs to be biochemically characterized, especially those with selectivity for unprecedented nonproteinogenic building blocks.

In recent years, unique lumazine-containing nonribosomal peptides have been isolated from marine-derived *Aspergillus* strains (Fig. 1). Penilumamide A (1), the first in its class, is a

Biosynthesis of the fungal nonribosomal peptide penilumamide A and biochemical characterization of a pterin-specific adenylation domain†

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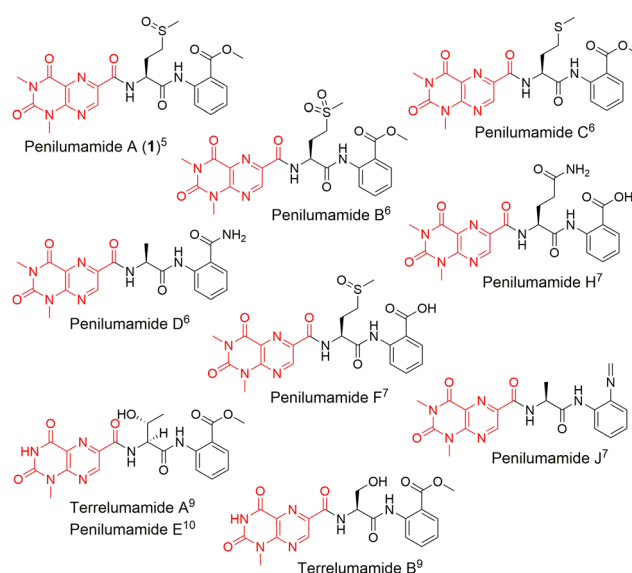


Fig. 1 Structures of penilumamide A and other related lumazine-containing peptides. The lumazine-derived moieties are highlighted in red.

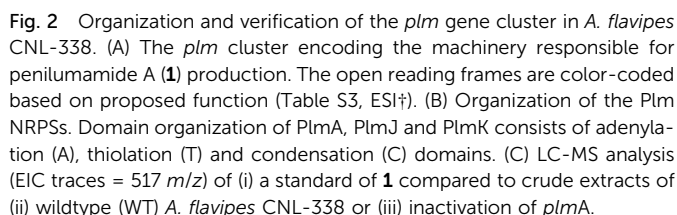
tripeptide containing a distinctive 1,3-dimethyl-lumazine-6-carboxylic acid functional group,⁵ and this unique pterin-derived moiety is found in penilumamide analogs, usually exhibiting either 1-*N*-methylation or *N,N*-dimethylation. Additional structural differences within this family of compounds includes variation in proteinogenic amino acid incorporation at the second position, as well as three different oxidation states of methionine, and incorporation of different aniline-derived C-terminal units such as anthranilic acid, methyl anthranilate, anthranilamide, or 2-aminophenyl isocyanide.^{6–11} Despite the number of lumazine-containing peptides that have been isolated, there have been no biosynthetic investigations of the respective nonribosomal peptide machinery, and more importantly, no reports on adenylation domains with preference for lumazine- or pterin-derived building blocks. Herein, we report the biosynthetic pathway for 1, detailed bioinformatic

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Inactivation of *plmA*, coding for a monomodular NRPS, confirmed the cluster's role in synthesizing **1** (Fig. 2C and Fig. S3, ESI[†]).^{18,19} Interestingly, while penilumamide A is a linear tripeptide, closer inspection of the three NRPSs, PlmA, PlmJ and PlmK, revealed a total of four modules, indicative of a tetrapeptide product. Thus, the activities of the monomodular PlmA and PlmK and dimodular PlmJ were reconstituted *in vitro* to verify if all three NRPSs were required for assembly of **1**. *plmA*, *plmJ* and *plmK* were solubly expressed as recombinant C-terminal hexahistidyl-tagged proteins from *Saccharomyces cerevisiae* BJ5464-NpgA²⁰ (Fig. S4, ESI[†]). Combinations of PlmA, PlmK and fractions enriched with PlmJ were incubated with pterine-6-carboxylic acid, L-methionine and anthranilic acid before analysis by liquid chromatography-mass spectrometry.



As the *in vitro* reconstitution assays showed that PlmA, PlmJ and PlmK were required for the production of 2, detailed bioinformatic investigations of the signature motifs in each domain of the three NRPSs was carried out to determine if any modules could be inactive (Fig. S8, ESI[†]). All four A domains and all four T domains in PlmA, PlmJ and PlmK contain the required catalytic residues and were predicted to be active (Tables S5 and S7, ESI[†]). Of the five C domains, all were predicted to be active with the exception of the N-terminal C domain in PlmA (PlmA C1), which is truncated, lacking key catalytic residues, and thus predicted to be inactive (Table S9, ESI[†]). As C domains have been posited as secondary gatekeepers to NRPS biosynthetic pathways,²¹ downstream of the primary selectivity of A domains, a maximum-likelihood phylogenetic tree was constructed to aid in predicting the biosynthetic function of the Plm C domains and perhaps module order (Table S8 and Fig. S10, ESI[†]). While the C domain in PlmK was shown to clade with terminal anthranilate-incorporating domains, the second C domain in PlmJ (PlmJ C2) clades with C domains that couple L-amino acid donors to

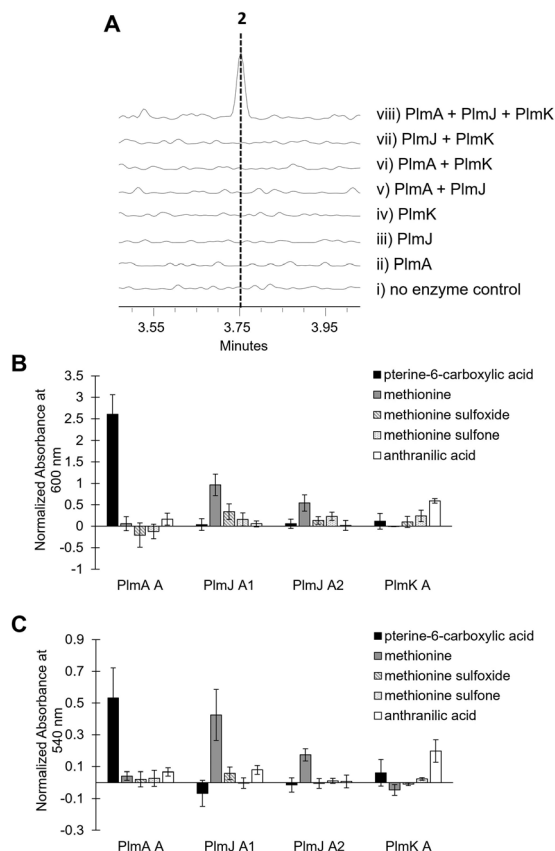


Fig. 3 *In vitro* reconstitution of demethyl-pterin-penilumamide (**2**) and adenylation assays to determine substrate specificity of the four dissected adenylation domains found in PlmA, PlmJ and PlmK. (A) LC-MS analysis (EIC = 458 *m/z*) of **2** when the pterine-6-carboxylic acid, L-methionine and anthranilic acid building blocks were (i) incubated together; (ii) incubated with PlmA; (iii) incubated with PlmJ; (iv) incubated with PlmK; (v) incubated with PlmA and PlmJ; (vi) incubated with PlmA and PlmK; (vii) incubated with PlmJ and PlmK; and (viii) incubated with PlmA, PlmJ and PlmK. (B) Adenylation activity was determined first through the malachite green/phosphate detection method.²² Activity was measured in triplicate as absorbance at 600 nm and normalized to boiled enzyme controls for each substrate. (C) Adenylation activity was verified with the hydroxylamine release/iron complex method.²³ Activity was measured in triplicate as absorbance at 540 nm and normalized to boiled enzyme controls for each substrate.

anthranilate acceptors. The *plm* biosynthetic cluster contains no thioesterase, reductive, or otherwise obvious offloading domain; however, due to the C-terminus of **1** being modified to the anthranilic acid methyl ester, it is likely that the C domain embedded in PlmK hydrolyzes **2** as the free carboxylic acid which is then methylated, as in crocacin biosynthesis.²⁴ Thus, the order of NRPSs in the biosynthesis of **1** likely goes from PlmA to PlmJ to PlmK. To validate this order, and as all four A domains were predicted to be active, we investigated the substrate specificities of the individual A domains.

Adenylation domain sequences were excised from all three Plm NRPSs based on predicted domain boundaries²⁵ and expressed as N-terminal octahistidine-tagged proteins from *Escherichia coli* BL21 (DE3) (Fig. S5, ESI†). As there are several

routine biochemical methods that can be used to probe adenylation activity, we chose two complementary colorimetric methods: a PPI release assay and a hydroxylamine-trapping assay. First, using an established phosphate detection method,²² A domain activation was indirectly measured using formic acid, anthranilic acid, pterine-6-carboxylic acid, L-methionine sulfoxide, L-methionine sulfone and the full panel of 20 proteinogenic amino acids (Fig. 3B and Fig. S6, ESI†). From these assays, the A domain in PlmA showed preference for pterine-6-carboxylic acid, whereas the second A domain in PlmJ (PlmJ A2) activated methionine, and the A domain in PlmK was specific for anthranilic acid. Unexpectedly, the first A domain of PlmJ (PlmJ A1) was also found to activate L-methionine with slightly higher activity than PlmJ A2, though only one methionine residue is present in **1**. This pattern of adenylation activity was confirmed using a well-established method involving direct detection of hydroxylamine-trapped aminoacyl-adenylates²³ (Fig. 3C and Fig. S7, ESI†). Altogether, these A domain results support the order of Plm NRPS modules predicted by C domain analysis.

As all four Plm A domains were biochemically characterized, the amino acid “specificity codes” of each domain were determined and compared to other A domains to interrogate patterns denoting substrate selectivity (Table S5, ESI†). It is well established in bacterial NRPSs that their substrate preferences can often be rationalized by 10 key amino acid residues that line the binding pocket of the A domain and act as a fingerprint for selectivity.^{26,27} This work is the first report of an A domain, PlmA A, activating a pterin-derived building block; its code DVMVLLMITK appears most similar to tryptophan-loading fungal A domains such as AnaPS A2 from acetylazonalenin biosynthesis.^{28,29} The A domain in PlmK, which was demonstrated through *in vitro* assays to activate anthranilic acid, is comparable to anthranilate-activating fungal A domains³⁰ such as PsyC A from psychrophilin biosynthesis.³¹ Both A domains of PlmJ were found to activate L-methionine, the first time this proteinogenic amino acid has been biochemically characterized in an Ascomycota-derived fungal NRPS system. Other native methionine-incorporating A domains, such as NpsP8 from napsamycin biosynthesis,³² SsaO from sansanmycin biosynthesis,³³ and JahA A3 from jahnellamide biosynthesis³⁴ have been predicted but lack biochemical confirmation. In malpinin biosynthesis, modules 1, 3 and 4 of MalA activate hydrophobic amino acids and were found to have relaxed substrate promiscuity towards L-methionine; however, methionine-containing malpinins were not identified as native natural products and were only observed upon media supplementation.¹²

The specificity codes for PlmJ A1 and PlmJ A2 were determined to be SIVIVTAGTK and DVVLLLSSTK, respectively. Due to the proclivity of pterins to photosensitize neighbouring amino acid residues,^{35,36} it is likely that PlmJ incorporates L-methionine which then spontaneously oxidizes to the sulf-oxide either while tethered as a dipeptide or after hydrolytic release of the tripeptide. This is supported by the preference for both PlmJ A1 and A2 to activate unoxidized methionine substrates (Fig. 3B and C). Of note, PlmJ A1 lacks the highly conserved aspartate in the first position of its code, which is

replaced with a serine. Interestingly, this change has been observed in fungal fumaric acid-activating A domains.^{37,38} However, penilumamide A does not contain this type of non-proteinogenic amino acid, nor any derivative isolated to date.

From detailed bioinformatic and biochemical investigations, the dimodular PlmJ contributes only one methionine residue to penilumamide A, suggesting the first module may be skipped or the C domain may be inactive. In previous studies on fungal NRPS module skipping,^{39,40} two possible mechanisms were suggested: either complete A-T-C module skipping *via* condensation by the upstream C domain, or chain transfer *via* the T domain of the skipped module (Fig. 4B). Based on our bioinformatic analyses (Fig. S8 and Table S9, ESI†), although PlmJ C1 maintains the required catalytic histidine, its predicted function as an anthranilate-donating C domain does not correlate with the preference of PlmJ A1 for L-methionine observed in the adenylation assays. Furthermore, the function of PlmJ A1 was incorrectly predicted, as it was shown to clade with a conserved group of anthranilate-activating A domains (Fig. S10, ESI†). Moving beyond primary sequence alignments to interrogate the structures of the four intact Plm C domains, homology models were generated using AlphaFold2⁴¹ and compared to the standalone C domain VibH.⁴² PlmJ C1 showed the lowest RMSD value when aligned with VibH (10.312), and when its HHxxxDR motif was modeled into VibH, the side chain of R131 protruded into the substrate channel, likely blocking it (Fig. S12, ESI†). Arginine has previously been observed in the motif of the Oxy enzyme-recruiting X domain Tcp12 C2⁴³ (Table S9, ESI†) as an H2R mutation, indicating PlmJ C1 might be involved in

the docking of tailoring enzymes such as the deaminase PlmD, or the methyltransferases PlmH or PlmI. All other modules and domains in PlmA, PlmJ and PlmK have multiple lines of evidence supporting their proposed roles, making the first module of PlmJ incongruous in the pathway. Therefore, we propose that the first module of PlmJ is not involved in chain extension and possibly skipped due to a structurally inactive C domain. However, it is not yet unequivocally determined which mechanism of module skipping is occurring, since PlmJ T1 maintains the active site serine that would be required for phosphopantetheinylation and subsequent chain transfer (Fig. 4B and Table S7, ESI†). Future work will aim to untangle the mechanistic possibilities, including independent mutagenesis of the PlmJ T1 and T2 domains, which may decrease yield of **1** or abolish production entirely.

Based on bioinformatic data and *in vitro* biochemical assays, we propose that PlmA, which contains a truncated and inactive C domain at its N-terminus, activates and incorporates the unique pterin-derived building block, which originates from GTP *via* the GMC oxidoreductase PlmB, the GTP cyclohydrolase I PlmC, the aldehyde dehydrogenase PlmE, the FAD-dependent oxidoreductase PlmF, and the dihydroneopterin aldolase PlmG (Fig. 4A). PlmJ then contributes a single L-methionine, with extension catalyzed by the C domain of PlmA, and anthranilic acid is loaded onto the T domain of PlmK before the terminal C domain of PlmJ couples it with the dipeptide (Fig. 4B(i)). The tripeptide **2** is then offloaded from PlmK *via* hydrolysis, followed by deamination and methylation reactions facilitated by PlmD, and PlmH and PlmI, respectively, to afford **1**. However, we can't rule out the possibility that deamination or *N,N*-dimethylation of pterin occurs while the peptide is tethered to a thiolation domain, perhaps mediated by protein-protein interactions with PlmJ C1. Interestingly, the Plm NRPSs do not contain annotated docking domains for inter-module communication as determined by HMMER. It is currently unknown how PlmA, PlmJ and PlmK associate together to generate the tripeptide, but this could provide an opportunity for engineering of NRPS assembly lines. Previous efforts to generate non-native peptides *in vivo* have revealed that the C domain-A domain interface provides a flexible region for insertion of didomains or entire modules.⁴⁴ Further, *in vitro* reconstitution of single NRPS modules has been successful when larger polypeptides are split at the C/A domain interface.⁴⁵ Because the Plm NRPSs lack this docking region and are already organized into A-T-C modules, they may be more tolerant of non-native protein-protein interactions.

In summary, we have identified and characterized the first biosynthetic cluster responsible for synthesizing a lumazine-containing natural product, penilumamide A, from the marine-derived fungus *A. flavipes* CNL-338. Using gene inactivation experiments and *in vitro* reconstitution assays, we have shown that all three Plm NRPSs, encoding four modules, are required for the biosynthesis of the tripeptide, suggesting potential module skipping. Through detailed *in vitro* biochemical characterization assays, we determined the substrate specificity of the four Plm NRPS adenylation domains, with bioinformatic

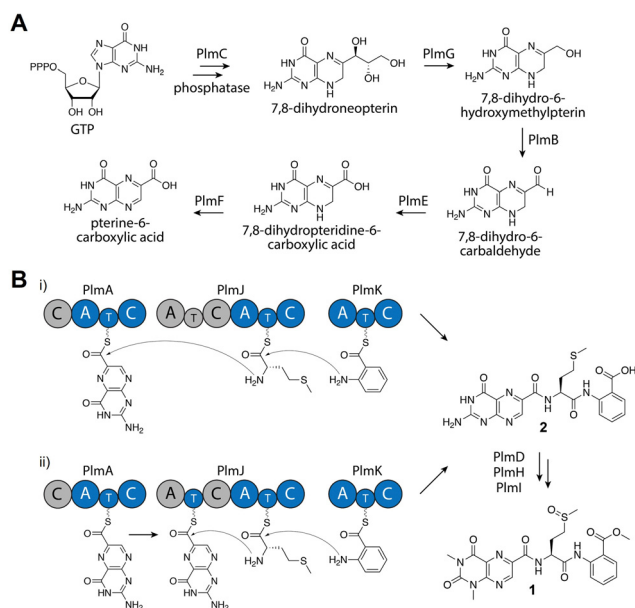


Fig. 4 Proposed biosynthesis of penilumamide A (**1**). (A) Biosynthesis of the pterine-6-carboxylic acid building block. (B) Assembly of the tripeptide can proceed through two possible mechanisms: entire A-T-C module skipping (i) or chain transfer *via* the T domain of the skipped module (ii). Based on bioinformatic analyses, gray domains are proposed to be inactive or skipped.

analyses revealing the first characterized “specificity codes” for native methionine- and pterin-activating A domains. Altogether, this knowledge can be applied to other fungal NRPS systems and assist with bioinformatic-based predictions. Furthermore, penilumamides exhibit promising therapeutic potential as insulin-sensitizing agents for the treatment of type II diabetes mellitus.^{8,9} Understanding how subtle variations in chemical structure affect bioactivity will improve our bioengineering toolkit and expedite the development of penilumamides for clinical use.

S. C. H. contributed to experiment conceptualization, investigation, validation, data analysis, visualization, and manuscript writing at all stages. K. L. D. contributed to experiment conceptualization, methodology, and helpful discussions. J. M. W. contributed to experiment conceptualization, data analysis, funding acquisition, project administration, resources, supervision, and manuscript writing at all stages.

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Conflicts of interest

The authors declare there are no conflicts of interest.

Notes and references

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§ The penilumamide biosynthetic cluster is deposited under the GenBank accession number ON297683. The sequenced fungal internal transcribed spacer region for *Aspergillus flavipes* CNL-338 is deposited under accession number MT579592.

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