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Discovery of Unguisin J, a new cyclic peptide from *Aspergillus heteromorphus* CBS 117.55, and phylogeny-based bioinformatic analysis of UngA NRPS domains

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Abstract

Several under-explored *Aspergillus* sp. produce intriguing heptapeptides containing a γ -aminobutyric acid (GABA) residue with as yet unknown biological functions. In this study a new GABA-containing heptapeptide – unguisin J (**1**) – along with known unguisin B (**2**) were isolated from a solid culture of *Aspergillus heteromorphus* CBS 117.55. The structure of compound **1** was elucidated by extensive 1D and 2D NMR spectroscopic analysis including HSQC, HMBC, COSY, and 2D-NOESY as well as HRESIMS. The stereochemistry of **1** and **2** was determined by Marfey's method. A biosynthetic gene cluster (BGC) encoding unguisins B and J was compared to characterized BGCs in other *Aspergillus* sp. Since the unguisin family of heptapeptides incorporate different amino-acid residues at different positions of the peptide, the A and C domains of the UngA NRPS were analyzed in an attempt to understand the lack of substrate specificity observed.

Keywords

Unguisin biosynthesis; fungal non-ribosomal peptide synthetase; adenylation domain; condensation domain; heptapeptide

Introduction

Unguisins are a small family of fungal cyclic heptapeptides isolated predominantly from *Aspergillus* sp. [1-8]. Distinctive features of these cyclic peptides include the non-proteinogenic amino acid γ -aminobutyric acid (GABA) and the incorporation of up to five D-amino acids (Figure 1) [1-8]. The amino acids at positions 1 (D-Ala) and 7 (GABA) are conserved but there is considerable variability at positions 2 – 6, including the incorporation of additional non-proteinogenic amino acids β -methylphenylalanine (β MePhe) and kynurenine (Kyn) [3,4]. So far, no significant biological activities have been reported for these small peptides [3,4,9], however unguisin A has been shown to bind a series of anions [10].

Recently the biosynthesis of unguisins A and B from *Aspergillus violaceofuscus* CBS 115571 was reported [5]. A seven module non-ribosomal peptide synthetase (NRPS; UngA) was heterologously expressed in *Aspergillus oryzae* NSAR1 yielding both unguisins A and B, which differ by the incorporation of D-Phe and D-Leu at position 3 respectively. The highly conserved D-Ala at position 1 was shown to be synthesized from L-alanine via the PLP-dependent alanine racemase UngC [5].

NRPS enzymes are large multifunctional enzymes that often synthesize very important bioactive molecules [11,12]. These enzymes consist of several catalytic domains organized into modules. Typically, a module possesses: an adenylation (A) domain for selecting and activating amino- or ketoacids; a thiolation (T) domain for shuttling intermediates between catalytic domains; and a condensation (C) domain that catalyzes amide or ester bond formation. Additional common domains include: epimerization (E) domains for converting naturally occurring L-amino acids to D- amino acids; methyltransferase (MT) domains that typically methylate specific N atoms; and terminal condensation (C_T) domains which cyclize the growing peptide chain and

facilitate release from the NRPS. Of the fungal NRPS studied to date, many appear to have some tolerance for the range of amino acids incorporated by the A domains and the C domain has been highlighted as a gatekeeper [13].

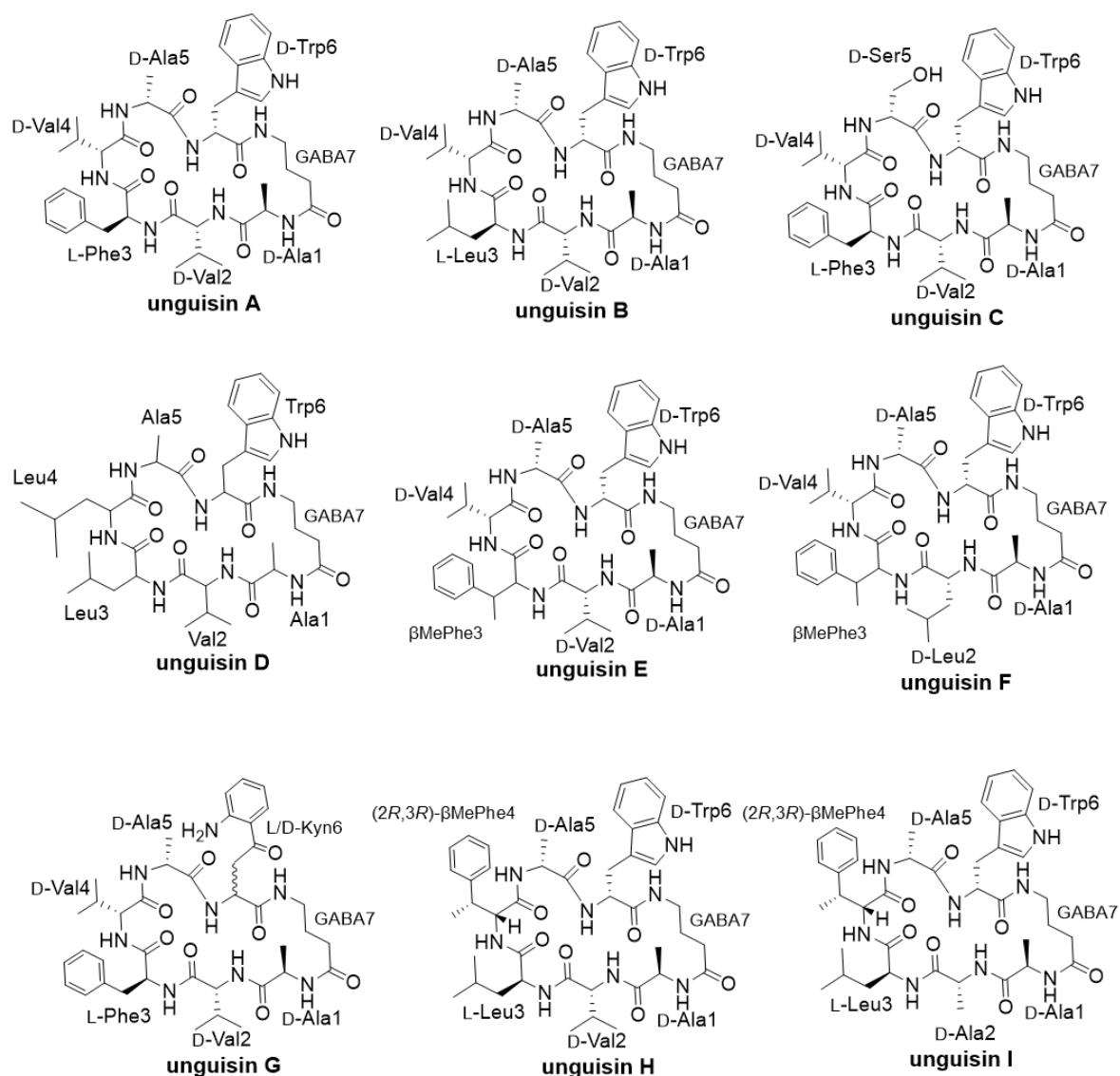


Figure 1: Structures of unguisins.

Here, we describe the isolation of unguisin B, and a new congener named unguisin J, from *Aspergillus heteromorphus* CBS 117.55. We also perform bioinformatic analysis of the A and C domains of the UngA NRPS enzymes involved

in their biosynthesis to try and rationalize the relaxed substrate specificity observed in this family of heptapeptides.

Results and Discussion

The cultivation of *A. heteromorphus* CBS 117.55 on rice solid medium yielded an organic-soluble extract, which was subjected to fractionation using preparative HPLC-PDA-ELSD and purification by semipreparative HPLC-PDA; this led to the isolation of a new cyclic peptide (**1**), along with unguisin B (**2**) (Figure 2). The structure of the new compound **1** was elucidated by 1D- and 2D-NMR and HRESIMS/MS. Unguisin B was identified by the ^1H and ^{13}C NMR data with the reported data [1,5].

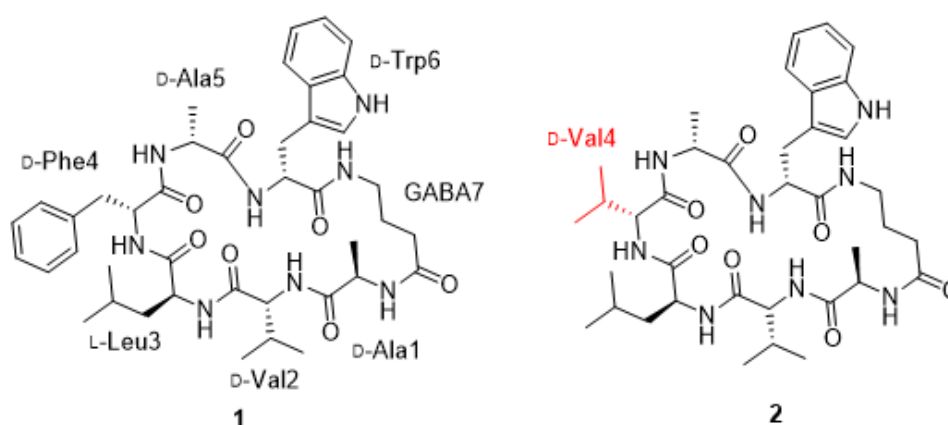


Figure 2: Chemical structures of unguisin J (**1**) and unguisin B (**2**).

Compound **1** was obtained as a white amorphous solid optically active, with $[\alpha]_D^{22} + 23.4$ (c 0.1; MeOH). Its molecular formula was established as $\text{C}_{41}\text{H}_{56}\text{N}_8\text{O}_7$ by HRMS ($[\text{M} + \text{H}]^+$ at m/z 773.4338, calculated for $\text{C}_{41}\text{H}_{57}\text{N}_8\text{O}_7^+$, m/z 773.4345, Δ 0.9 ppm; $[\text{M} + \text{Na}]^+$ at m/z 795.4162, calculated for $\text{C}_{41}\text{H}_{56}\text{N}_8\text{O}_7\text{Na}^+$, m/z 795.4164, Δ 0.3 ppm) and NMR data analysis, corresponding to seventeen indices of hydrogen deficiency. Its UV spectrum exhibited absorption maxima at λ_{max} 219 and 279 nm.

The ^1H and ^{13}C NMR spectra of **1** revealed the presence of seven amide NH signals between δ_{H} 7.43 and 8.35 ppm supported by the amide carbonyl signals at δ_{C} 173.1, 172.6, 172.6, 172.1, 172.1, 171.2 and 171.0 ppm (Table 1). An additional NH signal at δ_{H} 10.82 ppm and four aromatic signals at δ_{H} 7.50, 7.33, 7.07 and 6.97 ppm, exhibiting key ^1H - ^1H COSY and HMBC correlations, suggested a tryptophan aromatic amino acid portion (Figure 3). The other six amino acid residues were assigned based on 2D-NMR spectra (^1H - ^1H COSY, HSQC and HMBC) as Ala (2 eq.), Phe (1 eq.), Leu (1 eq.), Val (1 eq.), and γ -aminobutyric acid (GABA) (1 eq.). In addition to the COSY and HMBC correlations, the NOESY experiment showed important interactions between the NH signals corroborating with the peptide sequence defined to be Ala-1, Val-2, Leu-3, Phe-4, Ala-5, Trp-6, and GABA-7 (Figure 3).

Analysis of the NMR data of **1** allowed identifying characteristic ^1H and ^{13}C signals very similar to those of unguisin B (**2**) [1,5], the difference being the replacement of the Phe-4 in **1** by Val-4 in **2**. This assignment was confirmed by observation of HMBC correlations from δ_{H} 7.98 (NH) and δ_{H} 8.44 (NH) to C=O (δ_{C} 173.1) and from δ_{H} 2.93 and 3.01 ($\text{H}_{2-\beta}$) to C=O (δ_{C} 171.1) (Figure 3), together with key NOESY interactions between the NH signals at δ_{H} 7.98 \leftrightarrow 8.44 \leftrightarrow 8.14.

The absolute configuration of **1** was assigned by Marfey's method. [14] Comparison of the retention time by LC-MS between the derivatized **1** as well as the authentic amino acid samples determined the structure of **1** as cyclo-(D-alanine-D-valine-L-leucine-D-phenylalanine-D-alanine-D-tryptophan-GABA). Compound **1** was named as unguisin J.

Table 1: ^1H and ^{13}C NMR data for **1** (DMSO- d_6).

		1	
residue/position		δ_{C}	δ_{H} (mult., J in Hz)
Alanine	NH	-	8.35 (d, 4.2)
	C α	50.2	3.83 (m)
	C β	17.0	1.13 (d, 6.9)
	C=O	172.6	-
Valine	NH	-	7.43 (d 8.9)
	C α	58.4	3.98 (t 8.7)
	C β	30.0	1.98 (m)
	C γ	18.2	0.68 (d, 6.7)
	C γ	19.0	0.78 (d, 6.7)
	C=O	172.1	-
Leucine	NH	-	8.14 (d, 7.6)
	C α	51.8	3.88 (m)
	C β	39.4	1.36 (m)
	C γ	23.6	0.98 (m)
	C δ	20.8	0.56 (d, 6.6)
	C δ	23.3	0.69 (d, 6.6)
	C=O	172.1	-
	Phenylalanine	NH	-
C α		55.8	4.15 (m)
C β		36.1	2.93 (m), 3.01 (m)
C-1		137.2	-
C-2, C-6		129.2	7.16 (dd, 7.5, 1.3)
C-3, C-5		128.3	7.24 (d, 7.5)
C-4		126.4	7.19 (dd, 7.5, 1.3)
C=O		171.1	-
Alanine		NH	-
	C α	48.7	4.06 (m)
	C β	17.5	1.12 (d, 6.9)
	C=O	173.1	-
Tryptophan	NH	-	7.86 (d, 6.9)
	C α	55.3	4.01 (m)
	C β	25.0	3.22 (m)
	C-1	-	10.82 (d, 1.7)
	C-2	123.7	7.06 (br s)
	C-3	110.7	-
	C3a	127.2	-
	C-4	118.3	7.50 (br d, 8.0)
	C-5	118.4	6.97 (dt, 8.0, 0.9)
	C-6	121.1	7.07 (dt, 8.0, 1.0)
	C-7	111.5	7.33 (dt, 8.0, 0.9)
C-7a	136.4	-	
C=O	172.6	-	
GABA	NH	-	7.61 (dd, 5.4, 4.3)
	C α	38.8	3.07 (m), 2.99 (m)
	C β	25.6	1.69 (m), 1.60 (m)
	C γ	32.9	2.14 (m), 1.98 (m)
	C=O	171.0	-

A second peptide was isolated from the same culture of *A. heteromorphus* CBS 117.55 and identified as unguisin B (**2**) by comparison with literature data [1,5], further corroborating the identification of the new unguisin J (**1**). To the best of our knowledge these are the first metabolites reported from *A. heteromorphus* CBS 117.55.

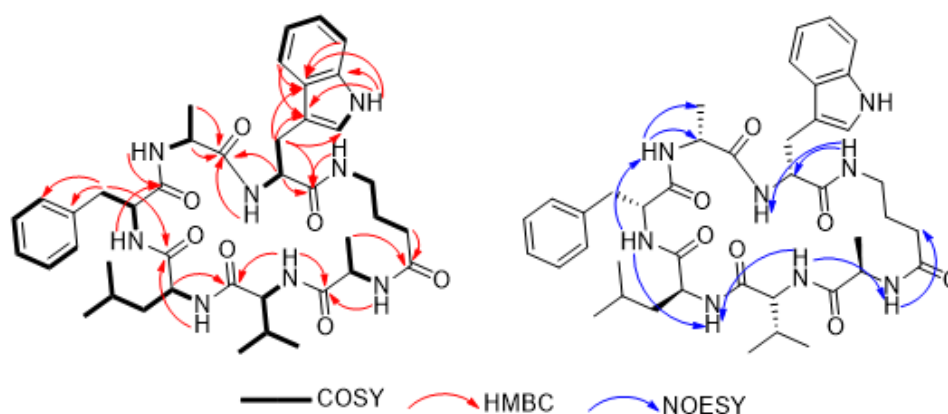


Figure 3: gHMBC and gCOSY correlations, and NOESY interactions of **1**.

The co-isolation of unguisins B and J indicates that module 4 of the NRPS is able to accept two different amino substrates and so may possess subtle differences to UngA from *A. violaceofuscus* CBS 115571 which has relaxed substrate specificity in module 3. We performed genome mining of the publicly available *A. heteromorphus* CBS 117.55 (accession number MSFL00000000.1) [15] using fungiSMASH and identified a four gene BGC encoding a seven module NRPS, an alanine-racemase, a hydrolase, and a transporter. We named this BGC *ung*'' to distinguish it from the *ung* BGC present in *A. violaceofuscus* and the *ung*' BGC in *A. campestris* IBT 28561 which encodes unguisins H and I [5]. Clinker analysis with the *ung* BGCs from *A. violaceofuscus* CBS 115571 and *A. campestris* IBT 28561 indicated a high level of homology (Figure 4). The biosynthesis of unguisins B and J therefore is proposed to arise from this single BGC, similar to the biosynthesis of unguisins A and B in *A. violaceofuscus* CBS 115571 (Scheme 1).

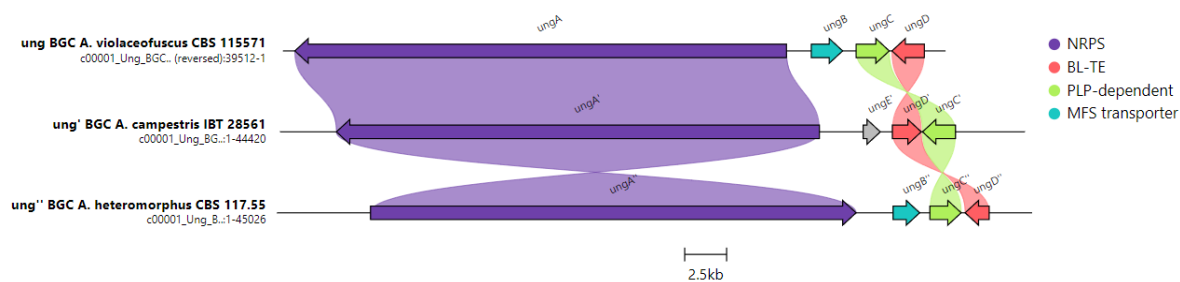
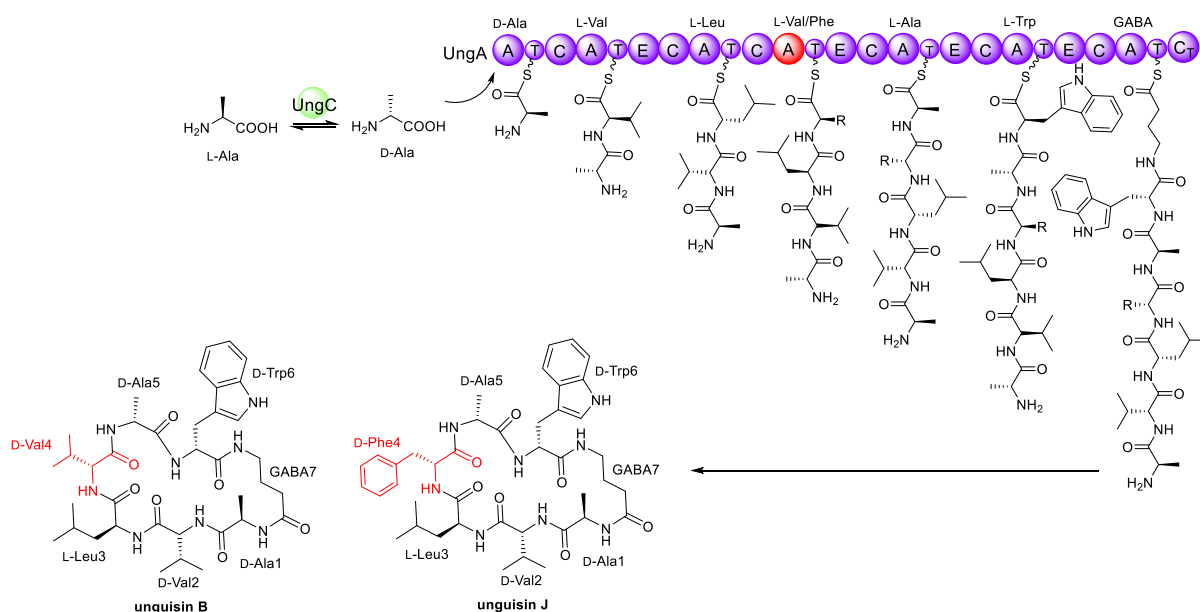


Figure 4: Clinker analysis of identified unguisin-encoding BGCs.



Scheme 1: Proposed biosynthesis of unguisins B and J in *Aspergillus heteromorphus* CBS 117.55.

Within the unguisin family, there is variability in the amino acids incorporated at positions 2 – 6 (Figure 1) however there are usually only one or two residue differences between molecules that are co-isolated from each source e.g. A and B from *A. violaceofuscus* CBS 115571 [5]; A, B, and C from *Emericella unguis* [1]; A, E, F and G from *Aspergillus candidus* NF2412 [4]; H and I from *A. campestris* IBT 28561 [5]; and B and J from *A. heteromorphus* CBS 117.55 (Figure 1). This implies that only one or two modules per NRPS possesses a noticeable level of relaxed substrate specificity.

To explore this observation the A and C domains were identified in UngA, UngA' and UngA'' and phylogenetic analysis of the A and C domains was performed (Figures 5 and 6).

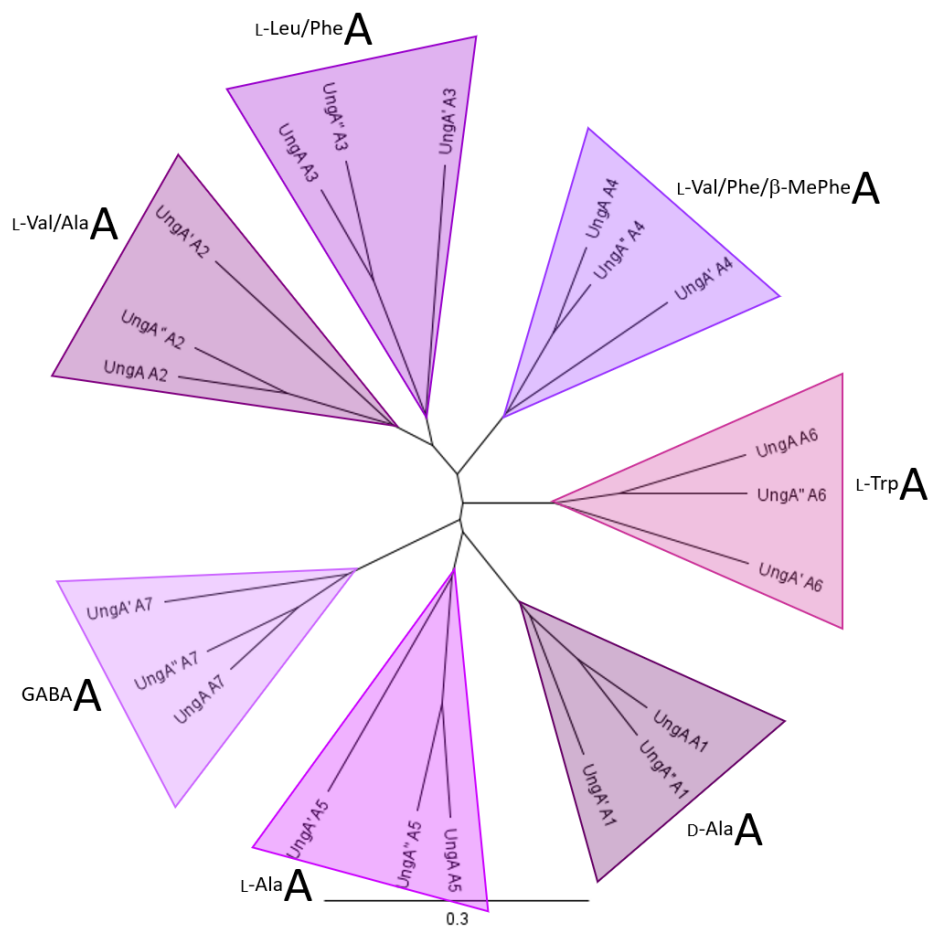


Figure 5: Phylogenetic analysis of A domains extracted from UngA NRPS. The substrate of the A domain is indicated for each clade.

The A domains do not clade according to substrate specificity – instead they clade according to which module they were extracted from. The A domains from modules 2, 3, and 4, which have relaxed substrate specificity, do appear to have evolved differently than A domains from modules 1, 5, 6, and 7 (Figure 5). Perhaps unsurprisingly the domains from UngA and UngA'' which both synthesize unguisin B, were more closely related than those from UngA' despite differences in substrate

specificity in modules 3 and 4. Previously Matsuda *et al.* had compared the putative non-ribosomal codes for the UngA and UngA' A domains and also observed that conventional approaches are inadequate to understand or predict the specificity of fungal A domains [5].

The clades formed by the C domains showed higher divergence than the A domains with the C_T domains forming their own branch and C domain from modules 1 and 3 clearly distinct to those from modules 2, 4, 5, and 6 (Figure 6). This separation of the non-terminal C domains could be due to modules 1 and 3 lacking an E domain. Again, the domains from UngA and UngA'' were more closely related than those from UngA' regardless of which two amino acids were condensed.

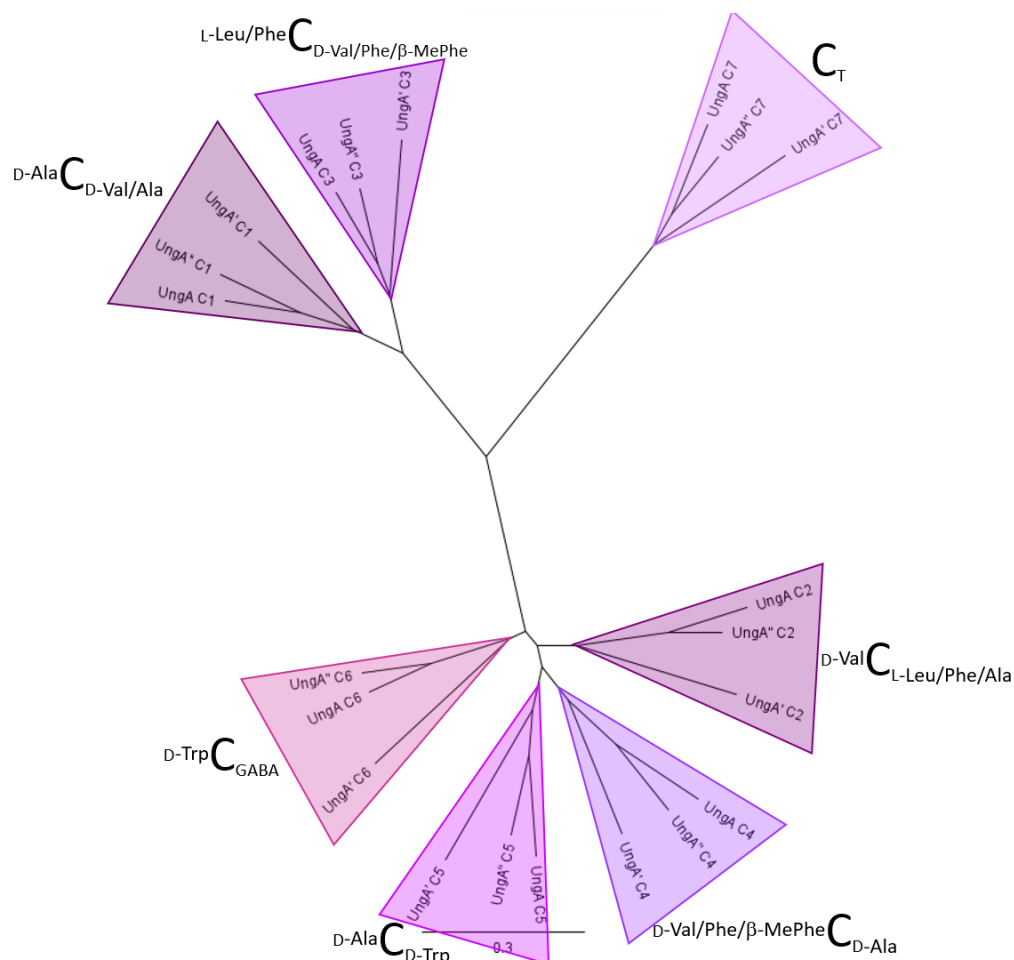


Figure 6: Phylogenetic analysis of C domains extracted from UngA NRPS. The substrates condensed by each C domain is indicated.

Conclusion

In this study unguisins B and J were isolated from *A. heteromorphus* CBS 117.55 which has not been extensively investigated for secondary metabolite production. A BGC encoding the unguisins was identified by genome mining with high homology to *ung* BGCs from other *Aspergillus* sp. Phylogenetic analysis of the A and C domains extracted from the UngA NRPS indicates that domains within modules are more closely related – even when substrate specificity differs - than domains within other modules that accept the same substrates.

Experimental

General experimental procedures

A. heteromorphus CBS 117.55 was purchased from the Westerdijk Fungal Biodiversity Institute. Rice solid medium was purchased from RiceSelect Organic Texmati. All solvents used for conducting LC analysis were purchased from Fisher Scientific. DMSO- d_6 NMR solvent was purchased from Sigma-Aldrich. N α -(5-Fluoro-2,4-dinitrophenyl)-D-leucinamide was purchased from TCI Chemicals. Authentic amino acid samples were purchased from Thermo Scientific. Sodium bicarbonate purchased from Fisher Chemical.

1D- and 2D-NMR experiments were recorded on a Varian INOVA 500 instrument (^1H : 500 MHz; ^{13}C : 125 MHz). The chemical shifts (δ) were expressed in

ppm and recorded with reference to solvent signals (^1H NMR: DMSO- d_6 2.50 ppm; ^{13}C NMR: DMSO- d_6 39.5 ppm). Optical rotation was measured on a Jasco P-2000 polarimeter with a path length of 100 mm. Analytical HPLC-PDA-MS system was a Shimadzu instrument (LC2030C 3D Plus Prominence) coupled to a Shimadzu LCMS-2020 mass spectrometer. Analyses were performed using a Phenomenex Kinetex RP₁₈ column (100 mm \times 4.6 mm i.d., 2.6 μm) along with the Security Guard RP₁₈ protective guard column (4.6 mm i.d.) and eluting with H₂O + 0.1% formic acid and MeCN + 0.1% formic acid using a gradient from 90:10 to 90:10 of H₂O-MeCN over 15 min, maintaining in 10:90 H₂O-MeCN for 3 min, from 10:90 to 90:5:5 in 1 min, and maintaining at 90:5:5 for 1 min, using a flow rate of 1.0 mL/min. The PDA detector scanned between λ 190 and 700 nm. The MS was optimized using the following conditions: interface voltage 4.5 kV; interface temperature 350 $^\circ\text{C}$; DL temperature 250 $^\circ\text{C}$; heat block 200 $^\circ\text{C}$; ESI mode, acquisition range 100 to 1000 Da; nebulizing gas 1.5 L min⁻¹; drying gas flow 15 L min⁻¹. The fractionation of the sample was performed on a Shimadzu LC-20AP preparative liquid chromatograph (SCL-40 System Controller Deliver and LH-40 Liquid Handler) coupled to a Shimadzu SPD-M40 Photo Diode Array Detector (PDA) system using a RP-18 column (Phenomenex, Kinetex 250 \times 30 mm i.d., 5 μm , flow rate of 18.0 mL/min). The purification of compounds was performed on a Shimadzu LC-20AD liquid chromatography (CBM-20A Communication Bus Module, CTO-20A column oven, DGU-20A Degassing Unit and SIL-20A AutoSampler) coupled to a Shimadzu SPD-20A UV/vis Detector system using a RP-18 column (Shimadzu, Premier 250 \times 10 mm i.d., 5 μm , flow rate of 3.0 mL/min). The high-resolution mass spectrometry (HRMS) was recorded on an ABSciex TripleTOF 6600+ mass spectrometer. Direct infusion of compounds **1** and **2** through the HRMS was performed using a flow rate of 10 $\mu\text{L}/\text{min}$ which the samples were diluted at 10 ppm with a solution of MeCN/H₂O (50:50; v/v) containing 0.1% formic

acid. The parameters such as declustering and entrance potentials remained constant for MS and MS/MS were set up at 150 V and 10 V, respectively. Collision energy for MS and MS² scan surveys was 10 V and 45 V, respectively, with a collision energy spread of 12 V for MS² scan survey. Precursor ion was impacted with three different collision energies (33, 45, 57 V), and the resulting MS² spectra were combined into one final MS² spectrum. The mass spectra were acquired using Turbo Spray Ionization set to 5.5 kV in positive ion mode with an accumulation time of 100 msec. The mass ranges for MS and MS² scan surveys were 500-800 amu and 30-800 amu, respectively. The curtain gas (nitrogen), nebulizing and heating gas were fixed at 25 psi, 20 psi and 15 psi, respectively. The temperature of the source was 25 °C. MS spectra were acquired and processed using Analyst TF 1.8.1 software.

Fungal growth and extraction

A. heteromorphus CBS 117.55 was cultivated in 2 Erlenmeyer flasks (500 mL), each containing 90 g of rice and 150 mL of H₂O [16]. The medium was autoclaved at 121°C for 20 min. After sterilization, the medium was inoculated with the spore solution of *A. heteromorphus* (1 mL) and incubated in static mode at 25°C for 21 days. The following day, the cultured mass in the flasks was ground and extracted with ethyl acetate (EtOAc, 3 × 100 mL). The EtOAc fraction was dried using a rotary evaporator and then dissolved in CH₃CN for defatting with hexane by partitioning. The CH₃CN fraction was evaporated, yielding 0.601 g of soluble-organic extract.

Fractionation and isolation of unguisin J and B

The soluble-organic extract was fractionated by preparative HPLC-PDA using Kinetex RP18 column (250 mm × 30 mm i.d., 5 μm) and UV detector at λ_{max} 254 nm.

The mobile phase consisted of H₂O+0.05% formic acid (eluent A) and MeCN+0.05% formic acid (eluent B), which was eluted of 20-100% of B with flow rate of 18 mL/min, yielding 20 fractions.

Fractions Fr13 and Fr15 were subjected to a purification by semipreparative HPLC-UV using a Premier RP₁₈ column (250 mm × 10 mm i.d., 5 μm) and UV detector at λ_{max} 210 nm. The gradient elution consisted of 65:35 to 35:65 of H₂O/MeCN over 20 min, using a flow rate of 3.0 mL/min). Fractions Fr13 and Fr15 resulted in the isolation of **1** (11.5 mg) and **2** (14.1 mg), respectively.

Unguisin J (**1**). Amorphous white powder, $[\alpha]_D^{22} + 23.4$ (c 0.1, MeOH); UV (photodiode array, MeCN:H₂O) λ_{max} 219 and 279 nm. HR-ESI-MS *m/z* 773.4338 [M+H]⁺ (calcd for C₄₁H₅₇N₈O₇⁺, *m/z* 773.4345), *m/z* 795.4162 [M+Na]⁺ (calcd for C₄₁H₅₆N₈O₇Na⁺, *m/z* 795.4164). For the ¹H and ¹³C NMR spectroscopic data, see Table 1.

Bioinformatics

The *A. heteromorphus* CBS 117.55 genome was initially screened using fungiSMASH to identify scaffolds / contigs encoding secondary metabolites. Scaffold MSFL01000005.1 was further investigated using FGENESH [17] to further refine gene boundaries, introns, and resulting protein sequence (Table S1). Comparative genomics / Clinker analysis was performed using Cagecat [18,19]. A and C domains were identified via Scan Prosite [20] and fungiSMASH [21]. Sequence alignments (Figure S1 and S2) and phylogenetic trees were generated using GeneiousTM. Functional domain motifs were visualized using Web Logo [22] and compared to literature motifs (Figure S3 and S4).

Marfey's method to determine the absolute configuration of **1** and **2**.

Samples **1** and **2** (0.3 mg each) were hydrolyzed in 0.5 mL of 6 N HCl at 115 °C for 20 h. After cooling, the reaction mixture was evaporated under nitrogen gas flow. The residue was dissolved in 0.5 mL of H₂O, and dried using speedvac to remove the residual HCl. The hydrolysates of **1** and **2** as well as authentic amino acid samples were derivatized with Marfey's reagent (N α -(5-Fluoro-2,4-dinitrophenyl)-D-leucinamide) and analyzed by LC-MS for comparison of the retention times.

Supporting Information

Supporting Information File 1:

File Name: BJOC ESI Unguisin J

File Format: pdf file

Title: Spectroscopic and spectrometric data of **1** and **2**. Bioinformatic data of the biosynthetic gene clusters.

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