Combining Mass Spectrometric Metabolic Profiling with Genomic Analysis: A Powerful Approach for Discovering Natural Products from Cyanobacteria

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S *Supporting Information*

ABSTRACT: An innovative approach was developed for the discovery of new natural products by combining mass spectrometric metabolic profiling with genomic analysis and resulted in the discovery of the columbamides, a new class of di- and trichlorinated acyl amides with cannabinomimetic activity. Three species of cultured marine cyanobacteria, *Moorea producens* 3L, *Moorea producens* JHB, and *Moorea bouillonii* PNG, were subjected to genome sequencing and analysis for their recognizable biosynthetic pathways, and this information was then compared with their respective metabolomes as detected by MS profiling. By genome analysis,

a presumed regulatory domain was identified upstream of several previously described biosynthetic gene clusters in two of these cyanobacteria, *M. producens* 3L and *M. producens* JHB. A similar regulatory domain was identified in the *M. bouillonii* PNG genome, and a corresponding downstream biosynthetic gene cluster was located and carefully analyzed. Subsequently, MS-based molecular networking identified a series of candidate products, and these were isolated and their structures rigorously established. On the basis of their distinctive acyl amide structure, the most prevalent metabolite was evaluated for cannabinomimetic properties and found to be moderate affinity ligands for CB$_1$.
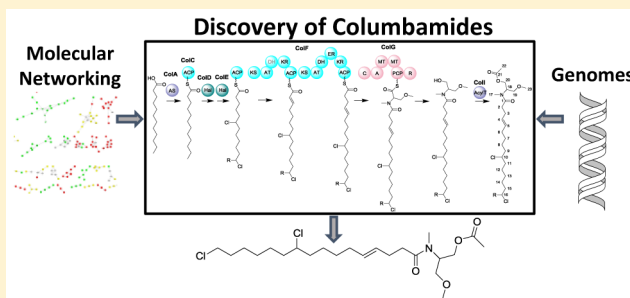
Marine cyanobacteria have emerged as a bountiful source of structurally diverse and biologically active natural products, some of which have inspired the development of new pharmaceutical agents.[1] Using the orthologous methods of genome mining and rapid mass spectrometric dereplication followed by careful structure elucidation, the discovery process of new secondary metabolites is becoming increasingly streamlined and efficient. The genomics approach provides information about the type of biosynthetic gene cluster present and, correspondingly, structural predictions about the natural products produced.[2−4] In cyanobacteria, polyketide synthases (PKS), nonribosomal peptide synthetases (NRPS), or hybrids of these two are most commonly encountered and are generally amenable to informatic-based deductions of structure.[5] With the mass spectrometric based metabolomics approach, deductions can be made about the number of compounds and compound classes present within a natural product extract. In addition, combining high-resolution mass spectrometry (HRMS) together with the molecular ion isotopic pattern and MS[2]-based fragmentation analyses, it is possible to develop tentative structural information about unknown compounds. Therefore, combining genomics and metabolomics makes possible the linkage of specific compounds to gene clusters

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Table 1. Moorea Species Studied in This Report, Their Origins and Reported Natural Products, and References to Previously Described Biosynthetic Gene Clusters[a]

strain	origin	compounds	cluster known?	NCBI accession number
Moorea producens 3L	Curaçao	barbamide,[19] dechlorobarbamide[20]	yes[13]	HQ696501
		carmabins A,B[21]	yes[11]	NZ_GL890970
		curacins A−C,[12] curazole[22]	yes[23]	HQ696500
Moorea producens JHB JHB22Aug96-1	Jamaica - Hector's Bay	hectochlorin[15]	yes[24]	AY974560
		jamaicamides A−C[14]	yes[14]	AY522504
Moorea bouillonii PNG PNG19May05-8	Papua New Guinea - Pigeon Island	apratoxins A−C[25]	yes[16]	
		lyngbyabellin A[17]	yes, unpublished	

[a]A phylogenetic analysis of these strains was previously published.[9,18]

and vice versa, and this information can be used to enhance the discovery and isolation of new natural products.[6,7]

Herein we describe the discovery of a new class of acyl amide based on genome comparisons and mass spectrometric metabolic profiling of three cyanobacterial strains of the genus *Moorea* (formerly known as *Lyngbya*[8]). This genus is known to produce many structurally diverse and biologically active natural products.[9] *Moorea producens* 3L, collected in Curaçao, produces the tubulin polymerization inhibitor curacin A, the molluscicide barbamide, and the antimalarial compound carmabin.[10−13] *Moorea producens* JHB, obtained from shallow coastal waters in Jamaica, is known for its production of the sodium channel blocker jamaicamide and the fungicide hectochlorin.[14,15] Complementing these, *Moorea bouillonii* PNG from Papua New Guinea produces the cytotoxic apratoxins A−C and lyngbyabellin A (Table 1).[16,17]

Improvements in whole genome sequencing and bioinformatic tools have resulted in a more facile identification of the biosynthetic gene clusters responsible for the formation of natural products.[26] In particular, the biosynthetic gene clusters encoding polyketides and nonribosomal peptides are readily detected, and subsequent structure predictions are possible.[27] Nevertheless, questions still remain whether an identified biosynthetic gene cluster is functionally expressed and if it is responsible for the production of a new or a known natural product. Due to the relative lack of molecular biology techniques, such as mutagenic gene knockouts and heterologous expression systems, for cyanobacterial *Moorea* strains as well as filamentous marine cyanobacteria in general, other methods must be used to unequivocally relate a given gene cluster to a specific natural product. For example, functional expression of distinctive biosynthetic enzymes from these clusters and characterization of their specificity and chemical reactivity has been used in several cases to confirm the connection between gene cluster and compound (e.g., barbamide, curacin A, jamaicamide A[14,28,29]). Another conceivable approach is to identify similar or nearly identical biosynthetic genes between different cyanobacterial genomes and to compare this information with that generated from parallel metabolomic studies. In the current study, the latter approach was taken in that all of the NRPS and PKS biosynthetic gene clusters of the *Moorea* strains listed above were identified in their respective genomic data sets, and this information was then juxtaposed with mass spectrometric profiles observed with the Molecular Networks algorithm[30] to identify a family of functionally expressed novel metabolites. Subsequently, these metabolites were isolated in high purity from laboratory cultures, and their structures rigorously determined as a series of acyl amides with unique positions

of chlorination. Owing to their structural relationship to anandamide and other cannibinomimetic compounds, the two major compounds columbamides A and B were evaluated for cannabinoid receptor CB_1 and CB_2 binding efficacy and found to be the most potent analogues yet isolated from the marine world.[31]

■ RESULTS AND DISCUSSION

Identification of Biosynthetic Gene Cluster. The genome of *M. producens* 3L was obtained by Sanger and 454 sequencing,[11] whereas those of *M. producens* JHB and *M. bouillonii* PNG were sequenced using Illumina Hiseq; in addition, for *M. bouillonii* PNG PACBIO sequencing was also performed. Assembly of all genomes utilized SPAdes 3.0 and 3.1[32,33] with BayesHammer [NN2] read error correction.[34] In each case, the source cyanobacteria were contaminated with heterotrophic bacteria that were associated with the filament sheath. As a result, a modest complexity metagenomic data set was obtained, and therefore the sequence data were binned to obtain the low GC cyanobacterial sequences (43.6% GC) separate from the higher GC content contaminants. Table 2

Table 2. QUAST (Quality Assessment Tool for Genome Assemblies) Results for the Three Moorea Strains

	3L	JHB 22Aug96-1	PNG 19May05-8
no. of contigs	161	960	579
largest contig	593 782	144 691	135 461
total length	8 478 612	9 501 724	8 576 967
N50	123 305	20 017	31 790
L50	19	145	77
GC (%)	43.68	43.67	43.65

depicts the quality of the three draft genomes using the Quality Assessment Tool (QUAST) for Genome Assemblies.[35] The published genome from *M. producens* 3L had 161 contigs, while the DNA sequences for *M. producens* JHB and *M. bouillonii* PNG were more fragmented, with 960 and 579 contigs, respectively. The approximate sizes of the *Moorea* genomes appear to be between 8 and 9.5 MB (Table 2).

Because the focus of this study was to detect biosynthetic gene clusters and their respective encoded compounds, we performed a manual assembly to close gaps (unpublished data) in cases where putative biosynthetic gene clusters were split between two contigs. Biosynthetic gene clusters were identified in the three draft genomes using common bioinformatics tools, including AntiSMASH and NaPDos.[27,36,37] The genome of *M. producens* 3L contains six PKS, NRPS, or hybrid biosynthetic pathways that include the curacin A, barbamide, and carmabin
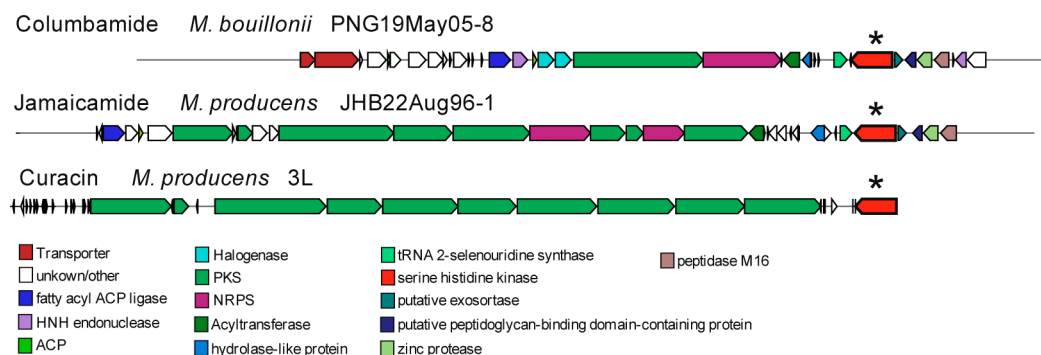
Figure 1. MultiGeneBlast result for the columbamide, jamaicamide A, and curacin A biosynthetic gene clusters. At the 3′ end of each pathway is an open reading frame for a putative regulatory serine histidine kinase (*).

pathways. The *M. producens* JHB strain has eight biosynthetic pathways in addition to those encoding for jamaicamide and hectochlorin production. In the genome of *M. bouillonii* PNG, seven PKS/NRPS-based biosynthetic pathways are present, including those for apratoxin and lyngbyabellin biosynthesis.

From LC-MS results as well as previous reports, *M. producens* 3L and *M. producens* JHB produce large amounts of the hybrid PKS-NRPS metabolites curacin A and jamaicamide A, respectively.[12,14] The biosynthetic gene clusters for both of these natural products are known, and intriguingly, they each are located adjacent to a regulatory serine histidine kinase gene.[38] This regulatory kinase is highly homologous between the two *Moorea* strains (96.1% identity based on protein sequence), suggesting the hypothesis that it is specifically associated with highly expressed natural product gene clusters. We therefore reasoned that searching for the gene encoding this regulatory enzyme within the *M. bouillonii* PNG genome sequence might identify new natural product biosynthetic gene clusters. Such a BLAST search was conducted using the entirety of the kinase protein sequence from *M. producens* 3L (WP_008191774.1) and revealed one highly homologous sequence in the *M. bouillonii* PNG genome. Investigation of the gene neighborhood for this kinase revealed a new and undescribed biosynthetic gene cluster with several unique features, as described below.[39] To evaluate the potential expression of novel metabolites by this gene cluster, the metabolic profile of each strain was analyzed by mass spectrometric molecular networking[30,40] and then utilized in a comparative metabolomics approach to complement the genomic analysis.

Metabolic Profiling of *Moorea* Strains. Cultured biomass of *M. producens* 3L, *M. producens* JHB, and *M. bouillonii* PNG was harvested and chemically extracted, and the metabolic profile of each was evaluated using molecular networking.[30] This mass spectrometry based tool uses fragmentation patterns to determine the relative similarity of different metabolites. Molecules with similar fragmentation patterns cluster together and are visualized as a series of nodes (=molecules) and edges (=similarities) in Cytoscape. In addition, by comparing MS²-fragmentation patterns against the natural product MS/MS library, the dereplication process is accelerated, and novel compound classes can be easily identified. In the current case, the extracts were analyzed using both ion trap (ITMS) and QTof mass spectrometers; this allowed production of a robustly annotated network by utilizing chemical standards generated from our in-house pure compound library[41] as well as accurate formula assignments from the HRMS data. Figure 2

depicts the molecular network obtained from combining the data sets generated from analysis of the extracts of *M. producens* 3L, *M. producens* JHB, and *M. bouillonii* PNG. The color of the node reflects the strain [*M. producens* 3L (red), *M. producens* JHB (yellow), and *M. bouillonii* PNG (green)]. The size of each individual node reflects the abundance of the compound based on the number of MS²-spectra scans present. The square-shaped nodes indicate matches to previously characterized compounds. Each node is labeled with its respective parent mass.

The major compounds produced by *M. producens* 3L are the curacins,[12] barbamide,[13] and carmabins,[21] whereas those of *Moorea producens* JHB are the hectochlorins[24] and jamaicamides.[14] The metabolic profile of *M. bouillonii* PNG reflects the biosynthesis of lyngbyabellin A[17] and the apratoxins.[16] In addition, two clusters (marked with black squares) were particularly interesting because the isotopic pattern of the parent masses indicated di- and trichlorinated species. In addition, the relatively large size of the nodes reflected the fact that these molecules were abundant in *M. bouillonii* PNG. Moreover, these features of size and chlorination were generally in accord with the prediction of the new biosynthetic gene cluster in *M. bouillonii* PNG. Consequently, the biosynthetic gene cluster was analyzed in greater detail, the encoded compounds were isolated, and their chemical structures were elucidated by NMR, resulting in the discovery of columbamides A (**1**), B (**2**), and C (**3**).

Analysis of the Col Biosynthetic Gene Cluster. Table 3 and Figure 3 summarize the deduced function of each gene and depict its involvement in the putative biosynthetic pathway. On the basis of the subsequent structure elucidation of the columbamides, the gene cluster is designated as "*col*". Natural product biosynthetic gene clusters are often found associated with transposases; in this case, two were identified at the presumed 5′-end boundary of the gene cluster (Orf1, Orf2).[14,24,42] These genetically mobile elements can play a role in the horizontal acquisition and distribution of natural product gene clusters between different strains.[43] The initial biosynthetic enzyme of the cluster, ColA, encodes for a putative acyl-CoA synthetase. Cyanobacterial biosynthetic gene clusters often contain acyl-ACP synthetase starting units (jamaicamide JamA, hectochlorin HctA),[14,24] and commonly hexanoic acid is loaded onto an acyl carrier protein. In this case, BlastP analysis revealed a 63% identity between *col*A and *puw*C of the lipopeptide puwainaphycin biosynthetic gene cluster and a 57% identity between *col*A and the fatty acyl ACP ligase of the olefin-synthase (OLS) pathway.[18,44] The OLS pathway is
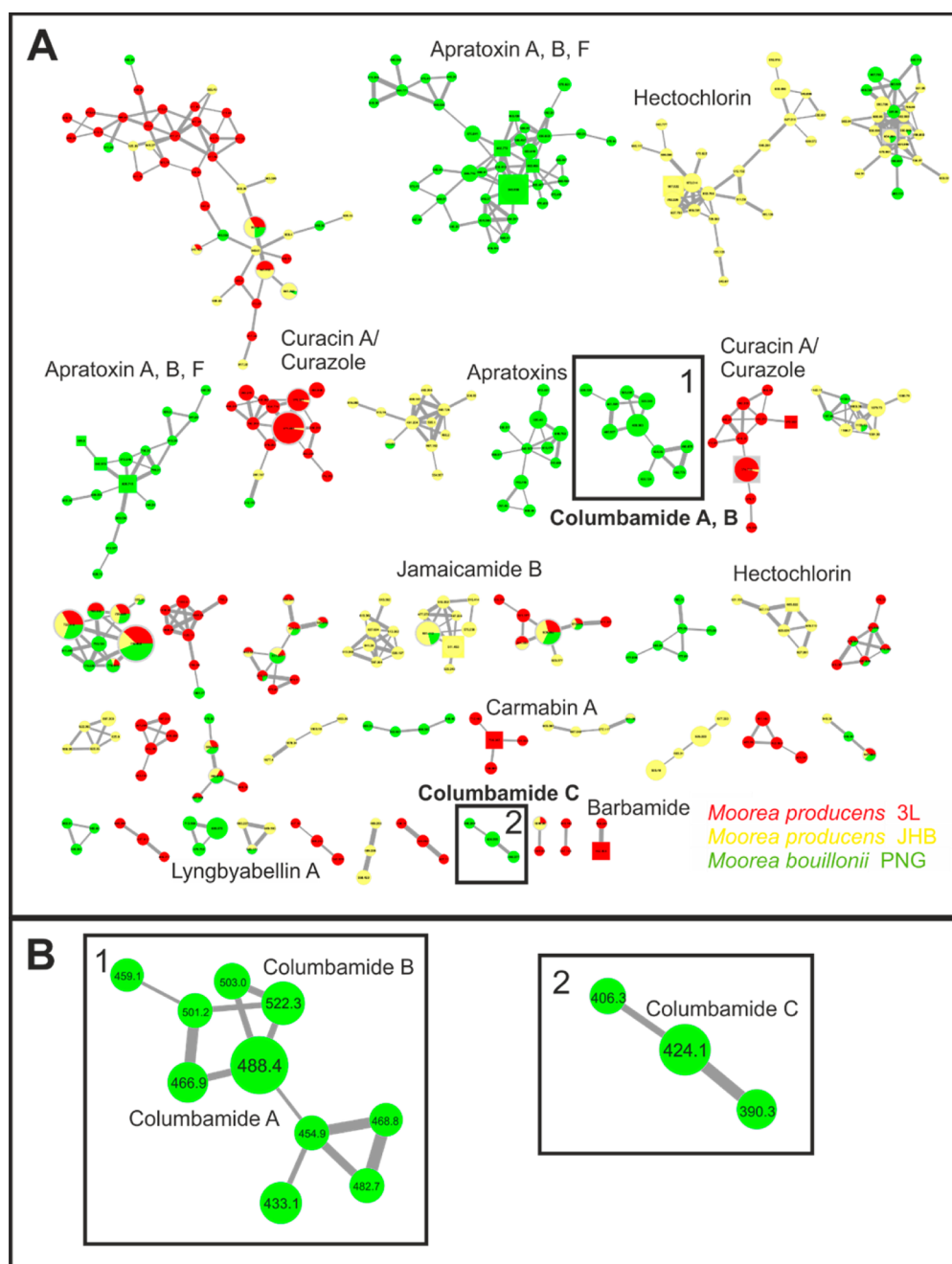
Figure 2. Molecular network (A) derived from mass spectrometric analysis of extracts of *M. producens* 3L (red), *M. producens* JHB (yellow), and *M. bouillonii* PNG (green). In the nodes, squares indicate consensus MS/MS spectra to compounds in a MS/MS library of known molecules. The respective compound name of an identified molecule is given next to the square node. The node size is representative of the numbers of MS^2 spectra obtained for that specific *m/z*. The thickness of the edges between nodes indicates the degree of similarity between their respective MS^2 spectra. Multiple clusters per compound derives from the fact that [M + H]$^+$ and [M + Na]$^+$ parent ions sometimes fragment differently. The black rectangles indicate the clusters of interest and are enlarged in B.

responsible for hydrocarbon production in cyanobacteria and utilizes long-chain fatty acids as initial substrates.[18] On the basis of subsequent structure elucidation of the compounds formed by this biosynthetic gene cluster, the *col*A gene loads dodecanoic acid onto the acyl carrier protein ColC.

The next gene in the cluster, *col*B, encodes for a putative HNH endonuclease [77% identity to HNH-endonuclease of *M. producens* 3L (WP_008186580.1)]. A functional role for ColB is not apparent at this time. The next two genes, *col*D and *col*E, most likely encode for a new type of halogenase, although the initial BLAST hits for these were for oxygenases. Cyanobacteria

have a variety of unique genes that encode for proteins involved in the halogenation of natural products. Examples include HctB in hectochlorin biosynthesis, BarB1 and BarB2 in barbamide biosynthesis, and the cryptic halogenase CurA in curacin A biosynthesis.[45−47] For ColD and ColE, the BlastP search yielded weak hits to *p*-aminobenzoate *N*-oxygenases AurF (ColD: 27% identity, 47% similarity, ColE: 30% identity, 44% similarity) and to the protein CylC (ColD: 48% identity, 66% similarity, ColE: 47% identity, 62% similarity), which is potentially a cryptic halogenase involved in the carbon−carbon bond activation of cylindrocyclophane biosynthesis.[48,49] The

Table 3. Deduced Functions of the Open Reading Frames in the Col Gene Cluster

protein	amino acids	proposed function	sequence similarity (protein, origin)	identity/ similarity	accession no.
ORF7	413		sucrose synthase, *Moorea producens*	97%, 98%	WP_008191839.1
ORF6	91		transposase, *Leptolyngbya* sp. PCC 7376	66%, 80%	WP_015134379.1
ORF5	51		glycyl-tRNA synthetase subunit alpha, *Streptococcus pneumoniae* GA17570	28%, 40%	EGI82761.1
ORF4	384		sucrose synthase, *Moorea producens*	98%, 98%	WP_008191839.1
ORF3	34				
ORF2	37		transposase, *Dolichospermum circinale*	61%, 75%	WP_028082114.1
ORF1	33		transposase, *Nostoc flagelliforme* str. Sunitezuoqi	74%,85%	ADO19008.1
ColA	583	dodecanoyl-ACP synthetase	PuwC, *Cylindrospermum alatosporum* CCALA 988	63%/79%	AIW82280.1
			fatty acyl ACP ligase, *Moorea bouillonii* PNG5-198	57%/71%	AHH34187.1
ColB	432	unknown	HNH endonuclease, *Moorea producens*	77%, 87%	WP_008186580.1
ColC	119	ACP	short-chain dehydrogenase family protein, *Mycobacterium ulcerans* str. Harvey	31%/50%	EUA89990.1
ColD	463	halogenase	CylC, *Cylindrospermum licheniforme* UTEX B 2014	48%/66%	AFV96137.1
			p-aminobenzoate *N*-oxygenase, *Leptolyngbya* sp. Heron Island J	27%/47%	ESA35271.1
ColE	471	halogenase	CylC, *Cylindrospermum licheniforme* UTEX B 2014	47%/62%	AFV96137.1
			p-aminobenzoate *N*-oxygenase AurF, *Oscillatoria acuminata*	30%, 44%	WP_015146466.1
ColF	3712	KS-AT-(DH)-KR-ACP-KS-AT-DH-ER-KR-ACP	CrpB, *Nostoc* sp. ATCC 53789	53%, 69%	ABM21570.1
ColG	2227	C-A$_{(Ser)}$-OMT-NMT-PCP-R	NcpB, *Nostoc* sp. ATCC 53789	51%, 66%	AAO23334.1
ColH	35	unknown	hypothetical protein, *Alicyclobacillus pohliae*	54%, 36%	WP_018133539.1
ColI	459	acyltransferase	Acyltransferase, *Mycobacterium tuberculosis*	16%, 32%	WP_031725540.1
ORF1′	723		choloylglycine hydrolase-like protein, *Lyngbya majuscula*	73%, 83%	AAS98791.1
ORF2′	41		hypothetical protein, *Lyngbya majuscula*	91%, 91%	AAS98793.1
ORF3′	33		hypothetical protein, *Gloeocapsa* sp. PCC 73106	81%, 93%	WP_006530731.1
ORF4′	49		hypothetical protein, *Moorea producens*	97%,96%	WP_008191272.1
ORF5′	355		tRNA 2-selenouridine synthase, *Moorea producens*	96%, 97%	WP_008191777.1
ORF6′	31		hypothetical protein, *Moorea producens*	84%, 84%	WP_008180717.1
ORF7′	1169	regulatory enzyme	guanylate cyclase, *Moorea producens*	94%, 96%	WP_008191774.1
ORF8′	220		putative exosortase, PEP-CTERM interaction domain protein, *Moorea producens*	80%, 88%	WP_008189692.1
ORF9′	269		putative peptidoglycan-binding domain-containing protein, *Moorea producens*	97%, 98%	WP_008189690.1
ORF10′	424		zinc protease, *Desulfotomaculum alkaliphilum*	26%, 47%	WP_031514442.1
ORF11′	428		peptidase M16, *Moorea producens*	97%, 98%	WP_008189686.1
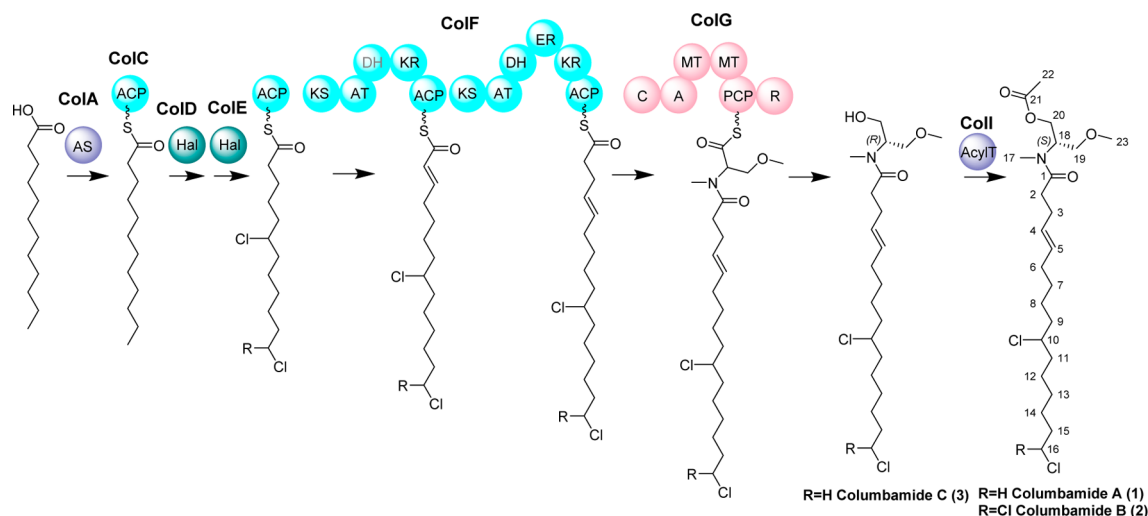
Figure 3. Proposed biosynthetic pathway of the columbamides in *M. bouillonii* PNG. AS: acyl synthetase, ACP: acyl carrier protein, Hal: halogenase, KS: β-ketoacyl-ACP synthase; AT: acyl transferase, DH: β-hydroxy-acyl-ACP dehydratase, KR: β-ketoacyl-ACP reductase, ER: enoyl reductase, C: condensation domain, A: adenylation domain, MT: methyltransferase, PCP: peptidyl carrier protein, R: reductase, AcylT: acyltransferase. Using multiple bioinformatic methods, the first DH domain in ColF appears to be absent and is therefore colored in light gray. As discussed in the text, the DH domain in the second PKS module of ColF is believed to provide this biochemical property.

Table 4. ^{1}H and ^{13}C NMR Spectroscopic Data for Columbamides A (1), B (2), and C (3) in CDCl$_3$ at 600 MHz for ^{1}H and 150 MHz for ^{13}C^a

	columbamide A (1)			columbamide B (2)			columbamide C (3)		
no.	δ_C, type	δ_H (J [Hz])	HMBC	δ_C	δ_H (J [Hz])	HMBC	δ_C	δ_H (J [Hz])	HMBC
1	173.5, C			173.1			174.2		
2	34.2, CH$_2$	2.40, m	1, 3, 4	33.9	2.40, m	1, 3, 4	34.2	2.41, m	1, 3, 4
3	28.6, CH$_2$	2.32, m	2, 4, 5	28.0	2.32, m	4, 5	28.0	2.33, m	2, 4, 5
4	129.3, CH	5.45, m	3, 6, 4, 5	129.3	5.45, m	6	129.0	5.46, m	3, 4, 5, 6
5	131.1, CH	5.45, m	3, 4, 5, 6	131.1	5.45, m	6	131.0	5.46, m	3, 4, 5, 6
6	32.5, CH$_2$	1.99, m	4, 5, 7, 8	32.3	1.99, m	4, 5, 7	32.0	1.99, m	4, 5, 7, 8
7	29.1, CH$_2$	1.35, m*	5, 6, 8, 9	28.6	1.33, m	6, 8	28.7	1.34, m	5, 6, 9
8	26.2, CH$_2$	1.35, m*	7, 9, 10	25.9	1.42, m	9	26.0	1.39, m	6, 7, 9, 10
9	38.5, CH$_2$	1.70, h (8.9, 7.7)	7, 8, 10	38.5	1.71, m	8, 10, 11	38.3	1.70, tp (10.5, 5.5, 4.3)	7, 8, 10, 11
10	64.3, CH	3.88, ddd (12.8, 8.1, 4.9)	8, 9, 11, 12	63.9	3.88, ddd (12.8, 8.1, 5.0)	8, 9, 11, 12	64.1	3.88, tt (8.2, 4.9)	8, 9, 11, 12
11	38.5, CH$_2$	1.70, h (8.9, 7.7)	9, 10, 12	38.5	1.71, dtd (12.1, 8.8, 8.1, 4.1)	9, 10, 12	38.3	1.70, tp (10.5, 5.5, 4.3)	8, 9, 10, 12, 13
12	26.5, CH$_2$	1.53, td (9.1, 5.0)	10, 11, 13, 14	25.9	1.43, m	11	26.0	1.54, ddt (23.2, 13.1, 6.1)	10, 11, 14
13	28.2, CH$_2$	1.35, m*	11, 12, 13, 14, 15	28.6	1.33, m	12, 14	28.6	1.33, m	11, 12, 14, 15
14	26.9, CH$_2$	1.45, dq (13.5, 6.7)	12, 13, 15, 16	25.8	1.55, m	12, 13, 15, 16	26.4	1.45, dq (13.0, 6.6, 5.9)	12, 13, 15, 16
15	32.6, CH$_2$	1.78, p (6.9)	13, 14, 16	43.5	2.2, m	13, 14, 16	32.4	1.78, p (6.9)	13, 14, 16
16	45.2, CH	3.53, t (6.7)	14, 15	73.5	5.75, t (6.0)	14	45.0	3.54, t (6.7)	14, 15
17	31.9 (27.6), CH$_3$	2.95, s (2.82, s)	1, 18 1, 18	31.6 (27.3)	2.96, s (2.82, s)	1, 18	33.8 (27.4)	3.02, s (2.84, s)	1, 18
18	52.1 (55.3), CH	4.90, tt (6.8, 6.8) (4.27, m)	1, 17, 19, 20	51.9 (55.1)	4.90, tt (5.5, 5.5) (4.27, m)		57.8 (57.8)	4.38, tt (5.2,5.2) (4.13, tt (6.3, 6.3)	1, 17, 19, 20
19	71.1, CH$_2$	3.58, dd (10.3, 6.9)	18, 20, 23	70.9 70.8 70.7	3.58, dd (10.3, 6.8)	18, 20, 23	70.9	3.57, dd (10.3, 4.9)	18, 20, 19
		3.48, dd (10.4, 5.3)			3.49, dd (10.3, 5.1)			3.67, dd (10.2, 7.4)	
20	62.2, CH$_2$	4.26, dd (11.5, 7.9)	1, 18, 19, 21	61.9	4.27, dd (11.5, 7.9)	18, 19, 21	62.3	3.79, d (3.9) 3.76, m	18, 19
		4.20, dd (11.9, 4.8)			4.20, dd (11.8, 4.9)				
21	170.9, C			170.6					
22	21.0, CH$_3$	2.04, s	21	20.7	2.04, s	21			
		2.05, s			2.05, s				
23	59.1, CH$_3$	3.32, s	19	59.0	3.32, s	19	59.1 (59.1)	3.34, s (3.35, s)	19
		3.34, s			3.34, s				

^{a}* represents overlapping signals; chemical shift values in parentheses are those resulting from minor conformer.

former gene, *aur*F, is involved in the biosynthesis of aureothin; however, the exact functioning and mechanism of the encoded enzyme are controversial. The Hertweck group identified AurF as a dimanganese enzyme, while the Zhao group reported AurF as a nonheme di-iron monoxygenase. Finally, the Bollinger group recently reported that it catalyzes a four-electron oxidation within a di-iron cluster.[50−52] The similarity in protein structure of ColD and ColE to AurF suggests that the halogenation mechanism at an unactivated center on an alkyl chain might be similar to the oxidation of aminoarenes to nitroarenes. Consistent with this deduction, the *cyl*C gene product is likely involved in the carbon−carbon bond activation of cylindrocyclophane, possibly through a cryptic chlorination event.[48,49] Ultimately, through isolation of the encoded compounds **1**−**3**, it was deduced that one putative halogenase introduces either one or two chlorine atoms at the terminal end of the alkyl chain, whereas the other introduces a chlorine atom at the ω-7 position of the fatty acid chain. Clearly, halogenation of these positions involves a highly unusual C−H bond activation and clearly warrants future investigation.

The next gene in the cluster, *col*F, matches a bimodular PKS motif. The protein sequence shows 53% similarity to CrpB, responsible for cryptophycin biosynthesis in *Nostoc* sp. ATCC 53789.[53] In this biosynthetic sequence, the acyl ACP is extended by two rounds of ketide extension by the two PKS modules found in ColF. Bioinformatics analysis in concert with the elucidated structure indicated the presence of the following domain structure: KS-AT-(DH)-KR-ACP-KS-AT-DH-ER-KR-ACP. On the basis of the structure of the isolated compound, the first polyketide module should contain a dehydratase domain. However, using common bioinformatics tools such as Delta BlastP, AntiSMASH, and 2metDB, a dehydratase domain was not detectable.[27,54,55] Interestingly, the curacin A pathway also has expected DH domains in each of the CurG, CurH, CurI, CurJ, and CurK proteins, but only three functional DH domains were evident.[23,56] It is believed that these functional DH domains provide the required biochemical activity for those modules in which the domain is absent; however, to date, the mechanism behind this presumed multiple use of DH domains remains unknown.
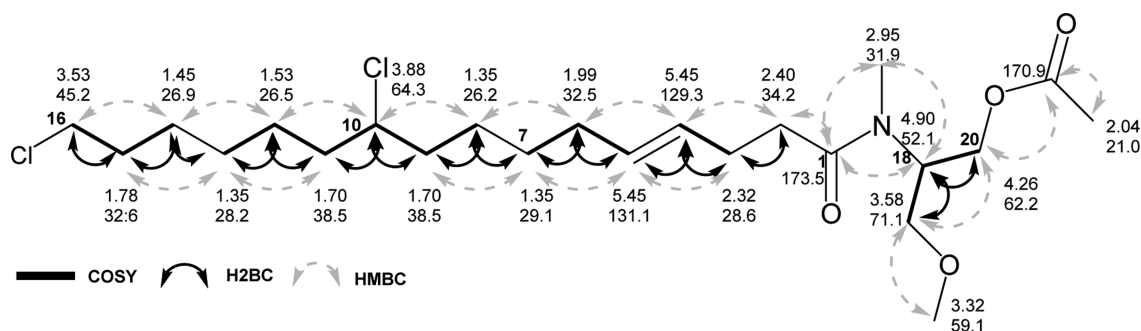
Figure 4. Selected 2D NMR data for columbamide A.

Next, the NRPS module (ColG) extends the acyl ACP by an amino acid. On the basis of the Stachelhaus predictions, the amino acid residues in the substrate-binding pocket of the A-domain are consistent with those encoding for the amino acid serine (Supporting Information Table S1).[57,58] The absence of an epimerase domain suggests that L-serine is incorporated into the molecule; this configuration was confirmed through subsequent hydrolysis and Marfey's analysis (Figure S10). Using the Delta BlastP function of the NCBI, two methyltransferase domains were identified, and subsequent structure elucidation confirmed the presence of one O-methyl and one N-methyl group. The sequences of the ColG methyltransferases were compared to known O- and N-methyltransferases (see Supporting Information for multiple sequence alignments, Tables S2 and S3). The N-methyltransferase domain of ColG shares an 80% identity to the N-methyltransferase domain of BarG. By contrast, the O-methyltransferase domain of ColG does not share a high sequence homology to any known O-methyltransferase, perhaps due to its unique positioning within an NRPS module. This is the first report for such an arrangement.[59] Lastly, the peptidyl carrier protein (PCP)-bound product is released by a reductase-releasing mechanism, presumably as a primary alcohol. The reductase domain shows a 48% identity and 66% similarity to the LtxA release domain of the lyngbyatoxin biosynthetic gene cluster, a terminating domain that has been demonstrated to catalyze an NADPH-dependent reductive cleavage to form a primary alcohol.[60] ColH encodes for a small hypothetical protein (35 amino acids), whose function is not clear at this time. In a final postmodular assembly step, ColI, a predicted acyltransferase, is thought to acetylate the primary alcohol to form columbamides A (**1**) and B (**2**).

Structure Elucidation of Columbamides. Cultured *M. bouillonii* PNG biomass was repetitively extracted with CH_2Cl_2/MeOH (2:1, v:v) and fractionated using normal-phase vacuum liquid chromatography (NP VLC). A fraction containing a relatively abundant series of lipid metabolites containing multiple halogen atoms was further purified using a reversed-phase C_{18} column followed by preparative HPLC purification. The structure elucidation of columbamide A was accomplished using an integrated approach that made use of traditional analytical data as well as bioinformatics from the gene cluster. For example, the molecular formula was determined considering the HRMS data, the isotopic pattern of the parent ion, and information deduced from the biosynthetic pathway. According to the latter, the formula should contain halogen atoms, one nitrogen atom, four oxygen atoms, two methyl groups, and a fatty acid chain. Using this information, the three molecular formulas were determined as $C_{23}H_{41}Cl_2NO_4$ for columbamide

A (**1**), $C_{23}H_{40}Cl_3NO_4$ for columbamide B (**2**), and $C_{21}H_{39}Cl_2NO_3$ for columbamide C (**3**).

The planar structures of each of the three analogues were deduced from 1D and 2D NMR data (Supporting Information, Figures S1−S9, S11−S21). The major metabolite, columbamide A (**1**), possessed an N-methyl amide as revealed from a three-proton singlet (distributed between δ_H 2.95 and δ_H 2.82 due to two rotamers around the CO−N bond[61]) that showed an HMBC correlation to the amide carbonyl carbon at δ_C 173.5. This carbon was connected by HMBC to a distinctive spin system composed of two methylenes (δ_H 4.26/4.20, δ_H 3.58/3.48) that bordered either side of a deshielded methine (δ_H 4.90/4.27). An O-methyl group, appearing as a three-proton singlet at δ_H 3.32/3.34, was attached to the more shielded of these two methylene groups, according to HMBC data. An acetoxy group was attached to the second methylene group (δ_H 4.26/4.20), as revealed by HMBC correlations from both a three-proton singlet at δ_H 2.04/2.05 and the deshielded methylene protons to an acetoxy carbonyl at δ_C 170.9. Location of this functionalized serinol fragment distal to the amide carbonyl was confirmed by HMBC correlations from H-18 to C-1 and from H_3-17 to both C-1 and C-18 (Table 4, Figure 4).

A 1,2-disubstituted alkene was evident from the two nearly identical alkene protons at δ_H 5.45 connected by HSQC to carbons with chemical shifts of δ_C 129.3 and 131.1. These protons showed COSY correlations to allylic methylene protons at δ_H 2.32 and 1.99. The former of these allylic protons was located adjacent to the C-2 protons at δ_H 2.40 by HMBC, thereby placing the olefin at C-4/C-5. The E-geometry of the double bond was indicated by characteristic ^{1}H and ^{13}C NMR chemical shifts for isolated olefins in alkyl chains (e.g., methyl elaidate and methyl oleate).[62] This was confirmed by measurement of the $^3J_{HH}$ coupling constant (14 Hz) between the two olefinic protons from the ^{13}C satellites observed in the HSQC spectrum (Figure S22).[63]

One chlorine atom was located attached to a terminal methylene group at δ_H 3.53 (δ_C 45.2, C-16) and, by its appearance as a triplet, indicated an adjacent second methylene group at δ_H 1.78 (δ_C 32.6, C-15). The latter methylene showed a COSY correlation to yet another methylene group at δ_H 1.45 (δ_C 26.9, C-14). An H2BC NMR experiment, providing correlations between protons and carbons specifically separated by two bonds, defined the methylene at δ_H 1.45 to be adjacent to a methylene carbon at δ_C 28.2 with associated protons at δ_H 1.35 (C-13).[64] H2BC data further placed this δ_C 28.2 carbon adjacent to a methylene at δ_H 1.53 (δ_C 26.5, C-12). By COSY, this methylene was adjacent to yet another CH_2 group at δ_H 1.70 (δ_C 38.5, C-11). Finally, the latter resonance was located by COSY and HMBC next to the second site of chlorination

(C-10) with a characteristic methine resonance at δ_H 3.88 (δ_C 64.3).

Confirmation that the second chlorine atom resided at the ω-7 position was obtained by locating this functional group at the C-10 position relative to the carboxylate of a hexadecenoic acid derivative. Despite overlap in both the proton and carbon dimensions between the methylenes to either side of C-10, an HMBC from the α-chloro methine proton to a carbon resonance at δ_C 26.2 allowed connectivity to the upstream portion of the carbon chain. This connectivity of C-10 to C-9 to C-8 could also be established from two sequential H2BC correlations (Figure 4 and Table 4). H2BC correlations from H$_2$-8 (δ_H 1.35) to C-7 (δ_C 29.1) were reinforced by HMBC correlations from H$_2$-9 to C-7. By COSY as well as H2BC, the C-7 methylene could be placed adjacent to the somewhat deshielded allylic methylene at δ_H 1.99 (H$_2$-6). As this methylene was already assigned in relationship to the carboxylate terminus, an (E)-10,16-dichlorohexadec-4-enoate residue was defined.

The only difference between columbamides A (**1**) and B (**2**) was the presence of an additional chlorine atom. The location of this addition was indicated by the triplet proton at δ_H 5.75 with its associated carbon shift of δ_C 73.5 ppm, indicative of a terminal *gem*-dichloro functionality. Indeed, these bands replaced those of the terminal chloro-methylene group in columbamide A (**1**) [see Supporting Information for stacked ^{1}H and 2D NMR spectra (Figures S1, S11−17) and Table 4]. In addition, the adjacent methylene group (C-15) was more deshielded (δ_H 2.2 and δ_C 43.5). The other portions of the molecule were identical to columbamide A.

In a similar fashion, the structure of columbamide C was easily deduced as being the same as columbamide A but lacking the acetoxy group at C-20 [see Supporting Information for stacked ^{1}H and 2D NMR spectra (Figures S1, S18−21) and Table 4]. The chemical shifts of the hexadecenoic acid portion of columbamide C were identical to columbamide A. The lack of the acetoxy group at C-20 resulted in a more shielded shift of the hydroxy-substituted methylene group at δ_H 3.79/3.76 (C-20) and the adjacent α-proton (C-18) at δ_H 4.38/4.13.

Next, the stereocenter of the dimethylated and acetylated serinol residue was established through Marfey's analysis. D- and L-N-methyl-O-methylserinol standards were synthesized according to standard protocols.[65] The two standards were derivatized with Marfey's reagent (L-FDAA) and compared to the derivatized hydrolysate of columbamide A. From the matching retention times, it was clear that this residue derived from L-serine, in accordance with the analysis of the biosynthetic pathway (Figure S10).

Biological Properties of Columbamides. The relative structural homology between the columbamides and the endogenous cannabinoid ligands, anandamide and 2-arachidonoyl glycerol, led us to evaluate the two most plentiful columbamides (**1**, **2**) for binding properties to the cannabinoid receptors CB$_1$ and CB$_2$. Columbamides A and B were found to be potent ligands for the CB$_1$ and CB$_2$ receptor [for CB$_1$ (±standard error): columbamide A: K_i = 0.59 ± 0.08 μM, columbamide B: K_i = 0.41 ± 0.06 μM; for CB$_2$: columbamide A: K_i = 1.03 ± 0.12 μM, columbamide B: K_i = 0.86 ± 0.09 μM], with columbamide B being slightly more active than columbamide A. Both compounds showed almost 2-fold greater affinity for CB$_1$ compared to CB$_2$. Previous studies have reported the isolation of other cyanobacterial-derived acyl amides, such as mooreamide A,[31] serinolamides A and B,[66,67]

grenadamide, and semiplenamides A, B, and G.[66] All of these compounds were shown to have binding activity with the cannabinoid receptors CB$_1$ and CB$_2$. Comparing K_i values, columbamide B (K_i = 0.41 μM) is slightly more active for CB$_1$ than the previously reported marine-derived CB$_1$ ligand mooreamide A (K_i = 0.47 μM). However, the columbamides are not as CB$_1$ receptor selective as was mooreamide A.[31] These two columbamides are the highest affinity ligands for CB$_1$ and CB$_2$ ever identified from marine natural products.[31]

■ CONCLUSION

Employing a combination of orthologous techniques, such as genome mining and mass spectrometric networking, led to the isolation and characterization of the structurally unique acyl amides columbamides A−C (**1−3**). The Global Natural Products Social Networking is an efficient tool for dereplicating known compounds as well as identifying new compound classes.[41] Using the Cytoscape platform, nodes are labeled by parent mass, colored by strain, shaped by library hits, and sized by number of spectra. Armed with this information, large numbers of metabolites can be easily visualized and assessed, and interesting new targets for isolation can be rapidly detected. In addition, the pie chart node application allows identification of which compounds are shared between *Moorea* strains and which compounds are unique to each strain.[68] In the current study, we found that the majority of secondary metabolites are strain specific. Further, we came to focus on the identification of several unknown products of *M. bouillonii* PNG. Through molecular networking the columbamides were detected, and this information was integrated with genomic information. However, the fragmentation pattern of columbamide C, due to the lack of the acetyl group, was different from the fragmentation pattern of columbamides A and B, and therefore, columbamide C was grouped into a different cluster (Figure 2). These new natural products are interesting for the concise integration of structural units and the uniqueness of the positions of chlorination. In this regard, this pathway, along with others such as that for cylindrocyclophane,[48,49] appears to possess a novel type of halogenase distinct from the 2-oxoglutarate-dependent radical halogenases we and others previously characterized as a part of the barbamide biosynthetic pathway.[28,46] These new metabolites are biologically significant as a result of their potent inhibition of the CB$_1$ and CB$_2$ receptors. Comparison of the genomes from *M. producens* JHB and 3L led to the identification of a highly homologous serine histidine kinase sensor closely associated with major secondary metabolite biosynthetic gene clusters (jamaicamide A in JHB and curacin A in 3L, Figure 1). The concept of a "genome hot spot" for secondary metabolite production is quite intriguing, as an aid in both informatically locating new biosynthetic gene clusters and potentially understanding their regulation. However, whether the locus of the biosynthetic gene cluster itself plays a crucial role or if the guanylate cyclase with its serine histidine kinase domain plays an important function in regulating secondary metabolite production remains unanswered.

■ EXPERIMENTAL SECTION

General Experimental Procedures. Optical rotations were measured on a Jasco P-2000 polarimeter. IR spectra were measured on a Thermo Electron Corporation Nicolet IR 100 FT-IR. ^{1}H and ^{13}C NMR spectra were recorded on a 600 MHz Bruker Avance III spectrometer with a 1.7 mm Bruker TXI cyroprobe. Spectra were

referenced to residual CDCl₃ solvent signals with resonances at $\delta_{H/C}$ 7.26/77.1. Semipreparative HPLC was performed on a Waters 515 pump system equipped with a Waters 996 PDA detector.

Cyanobacteria. *Moorea bouillonii* PNG05-198 was collected in May 2005 by scuba in 3−10 m water depth, where it was found growing tangled around the coral *Stylophora pistillata* off Pigeon Island, Papua New Guinea. *M. producens* 3L was originally collected near CARMABI Research Station in Curaçao, Netherlands Antilles. *Moorea producens* JHB 22Aug96-1 was collected in Hector's Bay, Jamaica. Live cultures have been maintained in SW BG-11 media under laboratory conditions.

Metabolic Profiling. Each strain was grown in SW BG-11 media. After harvest, the cells were freeze-dried and extracted with (2:1, v/v) CH₂Cl₂/MeOH. The extraction procedure was repeated until the supernatant was colorless. After removal of the solvent, the extracts were dissolved in CH₃CN. To remove compounds that would not elute from reversed-phase column material, each extract was purified using a disposable 100 mg reversed-phase C₁₈ catridge (Bond Elut-C₁₈ OH, 100 mg, 1 mL, Agilent Technologies). The column was conditioned with CH₃CN, and then an aliquot of extract was added to the column and eluted with 4 mL of CH₃CN. The solvent was removed, and the residue dissolved in CH₃CN to give a 10 mg/mL solution. The samples were analyzed via HPLC coupled to a Thermo Finnigan LCQ Advantage Max mass spectrometer or an Agilent 6530 Accurate Mass Q-TOF MS system. The Thermo Finnigan mass spectrometer was attached to a Thermo Finnigan Surveyor Autosampler-Plus, a LC-Pump-Plus, and a PDA-Plus system. ESI conditions were set to 325 °C capillary temperature, 5 kV source voltage, and 69 psi sheath gas flow rate. Four scan events were set up: positive total ion count of m/z 100−2000, followed by three data-dependent MS² scans of the first, second, and third most intense ions from the first scan event. The collision energy was 35%, the minimum intensity 10⁵ counts, the isolation width m/z 2, the dynamic exclusion count 5 with a repeat duration of 1 min, an exclusion list of 25, and an exclusion duration of 1.5 min.

The Agilent 6530 Accurate Mass Q-TOF MS system was attached to a 1290 Infinity Binary LC system (Agilent Technologies). ESI conditions were set to 300 °C capillary temperature, 3.5 kV VCap, and 10 L/min gas flow. The scan events were set up as follows: positive total ion count of m/z 100−1700, followed by data-dependent MS² scans. The collision energy was ramped with a 0.5 slope and a 30 offset; three precursors per cycle were fragmented with a MS abs threshold of 200, isolation width of m/z 1.3, and an active exclusion count of 5 with a repeat duration of 0.5 min.

The same gradient was used on both machines. The separation was performed using a Phenomenex Kinetex C-18 100 Å 100 × 4.6 mm column with a 700 μL/min flow and a linear gradient from 5% A (A: CH₃CN, B: 0.1% HCOOH in H₂O) for 3 min to 100% A in 45 min, which was maintained for 10 min. Afterward the column was equilibrated back to starting conditions.

The data were converted to mzXML format, a text-based format for mass spectrometry data by using MSConvert, part of the ProteoWizard package,[69] and uploaded to the Global Natural Products Social Molecular Networking Web site. The mass spectrometry data are available through MassIVE Public GNPS Data sets (MSV000079016, gnps.ucsd.edu). These data were analyzed using the molecular networking workflow.[41] The input data were searched against annotated reference spectra of the MS² library within GNPS. The GNPS MS² library contains ITMS and QTof reference spectra of previously isolated compounds of the Gerwick laboratory.[41]

Computationally, the algorithms compare MS² spectra by their similarity and assign similarity scores. The basic underlying algorithm was published in detail elsewhere.[30,40,41,68] For the network presented in this paper, the parent mass peak tolerance was set to 2 Da and the ion tolerance for mass fragments was set to 0.5 Da. The minimum cluster size was set to 1, and the cosine score was set to 0.7. For visualization, the created molecular networks were imported into the program Cytoscape 2.8.3.[70] Each node was labeled with their respective parent mass. The edges between nodes indicated the level of similarity between nodes, with thicker lines indicating higher

similarity. Hits to library inputs were marked as squares. The size of the node is representative of the number of MS² spectra present for each parent mass. To visualize which compounds are shared between the three *Moorea* strains, the pie-chart-creating tool (nodeCharts plugin for Cytoscape) was used.[68] Pie slices are proportional to the number of MS2 spectra for each parent mass and are therefore a proxy for its relative quantity. To simplify the network, nodes created by solvent background were removed from the network.

Isolation of Genomic DNA. Genomic DNA was isolated from live cultures of *M. producens* JHB and *M. bouillonii* PNG using a standard phenol/chloroform/isoamyl alcohol (PCI) extraction protocol. In brief, two grams (wet weight) of cultured biomass was rinsed with fresh SW BG-11 media, flash frozen in liquid nitrogen, and ground to a fine powder using a prechilled mortar and pestle. The powder was resuspended in 10 volumes of lysis buffer (10 mM Tris pH 8, 0.1 M EDTA pH 8, 0.5% w/v SDS, and 20 μg/mL RNase) and incubated at 37 °C for 30 min. Proteinase K was added (100 μg/mL final concentration), and samples were incubated at 50 °C for 1 h. After cooling the samples to room temperature (rt), one volume of equilibrated phenol (Life Technologies) was added and samples were mixed for 10 min. Phases were separated by centrifugation at 3300 rcf for 10 min, and the aqueous phase was removed to a new tube. The phenol extraction was repeated twice followed by a chloroform/ isoamyl alcohol (24:1) (Life Technologies) extraction. DNA was precipitated and purified by ethanol precipitation. Genomic DNA was quantified using a NanoDrop (Thermo Scientific) and qualified by gel electrophoresis. Genomic DNA was sequenced at the genomics core facility at the University of Michigan using Illumina HiSeq and at the University of California San Diego genomics facility using PACBIO (*M. bouillonii* PNG). Assembly was done using the SPAdes Genome Assember 3.0 followed by Opera to produce cyanobacterial scaffolds.[32] Further binning was performed to obtain cyanobacterial-specific scaffolds.

Bioinformatics Analysis. Biosynthetic gene clusters were identified using the genome mining software programs AntiSMASH, NaPDoS, and met2db.[27,37,54] NRPS adenylation domain substrate predictions were made using NRPSpredictor2.[71] Annotations were refined using Delta BlastP to identify conserved domains.[55] Phylogenetic analysis was performed by comparing the *N*- and *O*-methyltransferase domains using the Web-based tool at www. phylogeny.fr. Sequences of methyltransferases were obtained from SBSPKS.[72] The nucleotide sequence of the columbamide pathway genes has been submitted to GenBank under the accession number KP715425.

Structure Elucidation. About 5.5 g (wet weight) of *M. bouillonii* PNG was repetitively extracted with CH₂Cl₂/MeOH (2:1, v/v), yielding 389 mg of extract. Subsequently the extract was fractionated by silica gel vacuum liquid chromatography (VLC) using a stepwise gradient solvent system of increasing polarity beginning from 100% hexanes to 100% EtOAc to 100% MeOH, yielding nine subfractions. The fraction eluting with 40% EtOAc and 60% hexanes (fraction D, 3.8 mg) was separated further using a 100 mg reversed-phase C₁₈ solid-phase extraction cartridge (Bond Elut-C18 OH, 100 mg, 1 mL, Agilent Technologies) and a stepwise gradient solvent system of 25%, 50%, 75%, and 100% CH₃CN/H₂O, respectively. The 75% eluting fraction was further purified using reversed-phase HPLC: Phenomenex Kinetex C-18 100 Å 100 × 10 mm column with a 4 mL/min flow and a linear gradient from 75% A (A: CH₃CN, B: H₂O) for 3 min to 95% A in 24 min. Afterward, the column was equilibrated back to starting conditions. The yield of columbamide A was 2 mg, columbamide B 1 mg, and columbamide C 0.5 mg.

Columbamide A (1): $[\alpha]^{23}_D$ −11.3 (*c* 0.27, CHCl₃); IR (neat) ν_{max} 2927, 2856, 1621, 1452, 1124, 1089, 1046 cm⁻¹; ¹H and ¹³C NMR, Table 4 and Supporting Information; HRESIMS m/z 466.2478 [M + H]⁺ (calcd for C₂₃H₄₂Cl₂NO₄, 466.2485); MS/MS (CID 32.3%; [M + H]⁺) m/z (%) 370 (100), 398 (30), 334 (27), 102 (14), 362 (6).

Columbamide B (2): $[\alpha]^{23}_D$ 8.7 (*c* 0.77, CHCl₃); IR (neat) ν_{max} 2928, 2853, 1627, 1467 cm⁻¹; ¹H and ¹³C NMR, Table 4 and Supporting Information; HRESIMS m/z 500.2087 [M + H]⁺ (calcd

for $C_{23}H_{41}Cl_3NO_4$, 500.2096); MS/MS (CID 32.5%; $[M + H]^+$ m/z (%) 404 (100), 368 (99), 432 (35), 396 (30), 102 (17), 162 (7).

Columbamide C (3): $[\alpha]^{23}{}_D$ −27.5 (c 0.2, $CHCl_3$); IR (neat) ν_{max} 3401, 2931, 2859, 1627, 1465, 1124 cm^{-1}; ^{1}H and ^{13}C NMR, Table 4 and Supporting Information; HRESIMS m/z 424.2377 $[M + H]^+$ (calcd for $C_{21}H_{40}Cl_2NO_3$, 424.2380); MS/MS (CID 32.1%; $[M + H]^+$) m/z (%) 356 (100), 320 (46), 370 (17), 120 (4).

Marfey's Analysis of Columbamide A and Synthesis of Standards. To a solution of columbamide A (**1**) (0.4 mg) in MeOH (100 μL) was added 9 N HCl (200 μL), and the reaction stirred in a microwave reactor at 140 °C. After 20 min the reaction was quenched by 200 μL of saturated sodium bicarbonate solution, followed by addition of 45 μL of L-FDAA (0.5% solution in acetone). The solution was heated at 40 °C for 60 min, then quenched by addition of 150 μL of 2 N HCl. The mixture was diluted with 200 μL of CH_3CN, and 30 μL of the solution was analyzed by using the HPLC-ITMS described above.

The standard alcohols were synthesized according to the literature procedure.[65] Esterification of *O*-methylserine (**4**) followed by Boc protection gave product **5** in 82% yield over two steps. Reduction of ester **5** with $NaBH_4$ gave alcohol **6**, which was protected with TBDMS to give compound **7**. It was deprotected using TFA, and the crude product was reacted with L-FDAA Marfey's reagent, in the presence of excess Hünig's base, to prepare the standards. The *O*-methyl-L-serine gave the *R*-enantiomer of alcohol **8**, while the *O*-methyl-D-serine resulted in the *S*-enantiomer of compound **8**. The Marfey's derivatives L-FDAA of the hydrolysate and standards were analyzed with HPLC-ITMS by using a Phenomenex Luna 5 μm C_{18} column (4.6 × 250 mm). The HPLC conditions started with 10% CH_3CN and 90% HCOOH (0.1%) in H_2O followed by a gradual increase to 50% CH_3CN and 50% HCOOH in H_2O (0.1%) over 100 min at a flow rate of 0.4 mL/min with monitoring the total ion count from 100 to 2000 m/z. The retention time for the L-FDAA derivative of the *S*-alcohol was 72.4 min, while it was 64.3 min for the *R*-alcohol. The retention time of the L-FDAA derivative of the hydrolysate product gave a peak with a retention time of 64.7 min, corresponding to the *R*-alcohol from the L-serine.
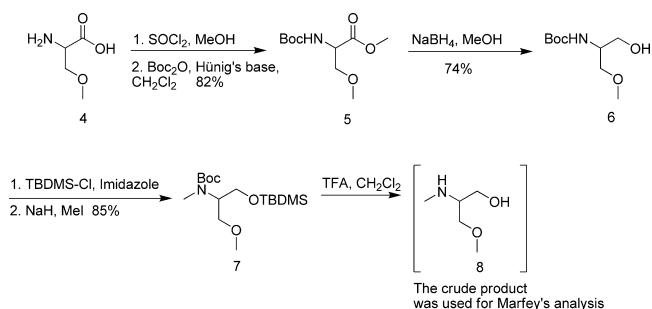
Synthesis of Methyl 2-(*tert*-Butoxycarbonylamino)-3-methoxypropanoate (5). To a solution of D- or L-*O*-methyl serine (0.15 g, 1.26 mmol) in MeOH (3 mL) at 0 °C was added thionyl chloride (1 mL) dropwise. The reaction mixture was warmed to room temperature and stirred overnight. The solvent was evaporated to give a yellow solid, which was used without further purification.

A mixture of *O*-methylserine methyl ester hydrochloride salt in CH_2Cl_2 (7 mL) at 0 °C was added to Hünig's base (0.66 mL, 3.78 mmol) and Boc-anhydride (275 mg, 1.26 mmol). The reaction mixture was warmed to rt and stirred overnight, after which it was diluted with CH_2Cl_2 (10 mL) and washed with 1 N HCl and brine. The combined organic layer was dried over anhydrous Na_2SO_4 and evaporated. It was purified by flash chromatography on silica gel in CH_2Cl_2/hexanes (75−100%) to give methyl 2-(*tert*-butoxycarbonyl)-3-methoxypropanoate (**5**) as a yellow oil (0.23 g, 82%). ^{1}H NMR (500 MHz, $CDCl_3$) δ 5.37 (d, J = 8.5 Hz, 1H), 4.41 (dd, J = 8.2, 3.8 Hz, 1H), 3.83−3.77 (m, 1H), 3.76 (s, 3H), 3.59 (dd, J = 9.5, 3.5 Hz, 1H), 3.35 (s, 3H), 1.45 (s, 9H); ^{13}C NMR (125 MHz, $CDCl_3$) δ 171.4, 155.7, 80.1, 72.7, 59.4, 54.0, 52.6, 28.4; LRESIMS m/z 234.17 $[M + H]^+$ (calcd for $C_{10}H_{20}NO_5$ 234.13).

Synthesis of *tert*-Butyl 1-Hydroxy-3-methoxypropan-2-yl-carbamate (6). To a stirred solution of ester **5** (75 mg, 0.32 mmol) in MeOH at 0 °C was added $NaBH_4$ in several portions. The reaction mixture was stirred at rt overnight, treated with water, and extracted with ethyl acetate. The combined organic layer was washed with brine, dried over anhydrous Na_2SO_4, and evaporated. The residue was purified by flash chromatography on silica gel in ethyl acetate/hexanes (10−30%) to afford **6** as a colorless oil (48 mg, 74%). ^{1}H NMR (**6**, 500 MHz, $CDCl_3$) δ 5.26 (br s, 1H), 3.77−3.73 (m, 2H), 3.68−3.64 (m, 1H), 3.56 (dd, J = 9.4, 3.8 Hz, 1H), 3.50 (dd, J = 9.5, 4.5 Hz, 1H), 3.37 (s, 3H), 1.45 (s, 9H); ^{13}C NMR (125 MHz, $CDCl_3$) δ 156.3, 79.8, 72.9, 63.5, 59.3, 51.6, 28.5; LRESIMS m/z 206.34 $[M + H]^+$ (calcd for $C_9H_{20}NO_4$ 206.14).

Synthesis of *tert*-Butyl 1-(*tert*-Butyldimethylsilyloxy)-3-methoxypropan-2-yl(methyl)carbamate (7). To a solution of alcohol **6** (34 mg, 0.17 mmol) in CH_2Cl_2 (2 mL) at 0 °C were added *tert*-butyldimethylsilyl chloride (26 mg, 0.17 mmol) and imidazole (13 mg, 0.19 mmol). The reaction was stirred overnight at rt and then quenched by addition of water. The organic layer was separated, washed with 2 N HCl, dried over anhydrous Na_2SO_4, and evaporated. The crude product was used in the next step without additional purification.

To a solution of *tert*-butyl 1-(*tert*-butyldimethylsilyloxy)-3-methoxypropan-2-ylcarbamate in THF (1 mL) was added NaH (60% dispersion in mineral oil, 22 mg, 0.47 mmol) at 0 °C. After stirring at the same temperature for 30 min, a solution of MeI (30 μL, 0.59 mmol) in DMF (30 μL) was added. The reaction mixture was warmed to rt and stirred overnight. It was quenched with H_2O, and the organic layer was separated. The aqueous layer was extracted with EtOAc, and the combined organic layer was dried over anhydrous Na_2SO_4 and concentrated. The residue was purified by flash chromatography on silica gel in CH_2Cl_2 to give product **7** (48 mg, 85%). $[\alpha]^{22}{}_D$ −12.6 (c 1.0, $CHCl_3$) for the *S*-enantiomer of **7** (derivative from *O*-methyl-L-serine). $[\alpha]_D{}^{22}$ +10.7 (c 1.0, $CHCl_3$) for the *R*-enantiomer of **7** (derivative from *O*-methyl-D-serine). ^{1}H NMR (**7**, 500 MHz, $CDCl_3$) δ 4.24 (br s, 1H), 3.75−3.66 (m, 2H), 3.61−3.57 (m, 1H), 3.53−3.47 (m, 2H), 3.33 (s, 3H), 2.84 (s, 3H), 1.46 (s, 9H), 0.89 (s, 9H), 0.15 (s, 6H); ^{13}C NMR (125 MHz, $CDCl_3$) δ 156.3, 79.7, 71.0, 62.1, 59.0, 57.2, 31.6, 28.8, 26.2, −5.3; LRESIMS m/z 334.18 $[M + H]^+$ (calcd for $C_{16}H_{36}NO_4Si$ 334.24).

CB_1 and CB_2 Receptor Binding Assays. CB_1 and CB_2 displacement assays were performed according to a previously reported protocol.[73] Membranes from HEK-293 cells stably transfected with human recombinant CB_1 receptor (B_{max} = 2.9 pmol/mg protein) or human recombinant CB_2 receptor (B_{max} = 6.0 pmol/mg protein) were incubated with the high-affinity ligand $[^3H]$-CP-55,940 (0.4 nM/K_d = 0.11 nM and 0.53 nM/K_d = 0.19 nM for CB_1 and CB_2, respectively). The radioactive ligand was displaced with 10 μM WIN 55212-2 as the heterologous competitor to determine nonspecific binding (K_i values 8.8 and 0.9 nM for CB_1 and CB_2, respectively). The compounds were tested according to the protocol described by the manufacturer (PerkinElmer). Displacement curves were obtained by incubating columbamides A and B with $[^3H]$-CP-55,940 for 90 min at 30 °C. K_i values were calculated by applying the Cheng-Prusoff equation to the IC_{50} values obtained from GraphPad for the displacement of the bound radioligand by increasing concentrations of the test compounds. Data are means of at least three replicates.

■ ASSOCIATED CONTENT

⑤ Supporting Information

Stachelhaus prediction, multiple sequence alignments of methyltransferases, NMR data, and Marfey's analysis chromatograms. The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.jnatprod.5b00301.

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Notes

The authors declare no competing financial interest.

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