

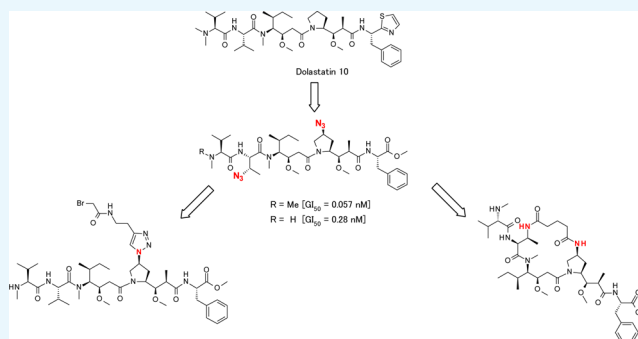
# Synthesis and Evaluation of Linear and Macrocyclic Dolastatin 10 Analogues Containing Pyrrolidine Ring Modifications

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## Supporting Information

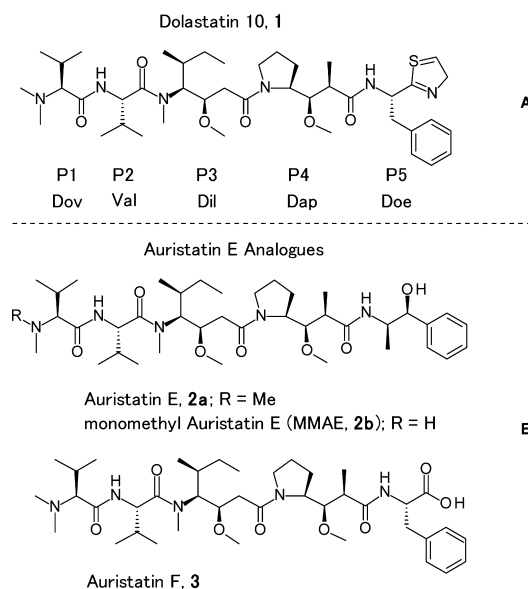
**ABSTRACT:** Because of their potent cytotoxic activity, members of the auristatin family (synthetic analogues of the naturally occurring dolastatin 10) have remained a target of significant research, most notably in the context of antibody drug conjugate payloads. Typically, modifications of the backbone scaffold of dolastatin 10 have focused on variations on the N-terminal (P1) and C-terminal (P5) subunits. Scant attention has been paid thus far to the P4 subunit in the scientific literature. In this paper, we introduce an azide functional group at the P4 subunit, resulting in potent cytotoxic activity seen in vitro. Another highly active compound in this study contained azide functional groups in both the P2 and P4 subunits and required dolavaline as the P1 subunit and a phenylalanine as the P5 subunit. Furthermore, these two azide groups served not only as modifiers of cytotoxicity but also as handles for linker attachment or as a tether for use in the synthesis of a macrocyclic analogue.



## INTRODUCTION

Naturally occurring peptides have been shown to be potent antibiotic agents, causing cell death at picomolar or low nanomolar concentrations.<sup>1</sup> Tubulin-binding natural products, especially vinca domain inhibitors, have played an important role in cancer research for decades.<sup>2–5</sup> One of them, the natural product dolastatin 10 (**1**), isolated from the sea hare *Dolabella auricularia*, was discovered by Pettit et al.<sup>6</sup> Dolastatin 10 (**1**) and its synthetic analogues (termed “auristatins”) (i.e., TZT-1027) demonstrated extraordinary cytotoxicity with subnanomolar activity in vitro toward a variety of cancer cell lines. However, when tested in the clinic, no significant activity was observed at the maximum tolerated dose.<sup>7–19</sup> Given these clinical results, and the likelihood that the lack of activity was due to the potency/toxicity of this class of compound as a small-molecule chemotherapeutic, many research groups have engaged in studies of synthetic dolastatin 10 derivatives as cytotoxic payloads in targeted therapies. The auristatin molecular class has found considerable application within the field of antibody drug conjugates (ADCs), with the goal to widen a clinically relevant therapeutic window. One targeted therapy approach has used an auristatin E (**2a**) derivative, monomethyl auristatin E (MMAE, **2b**), as the active payload in ADCETRIS<sup>20</sup> (Figure 1B). Other auristatin family members are under investigation as payloads in a variety of ADCs currently in clinical trials.<sup>21</sup>

Dolastatin 10 (**1**) consists of four amino acid building blocks, dolavaline (Dov, P1), valine (Val, P2), dolaisoleuine (Dil, P3), and dolaproine (Dap, P4), and the C-terminal amine dolaphenine (Doe, P5) (Figure 1A). Figure 1B illustrates the



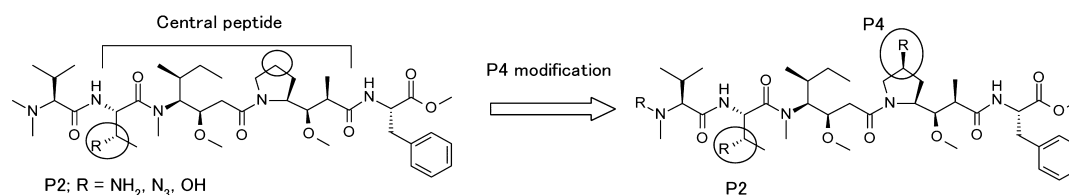
**Figure 1.** Structure of dolastatin 10 (**1**) (panel A) and auristatin analogues (**2a**, **2b**, and **3**) (panel B).

structure of auristatin analogues.<sup>22,23</sup> Many research groups have studied linear dolastatin 10 analogues and this area has been extensively reviewed.<sup>23</sup> In particular, numerous N-

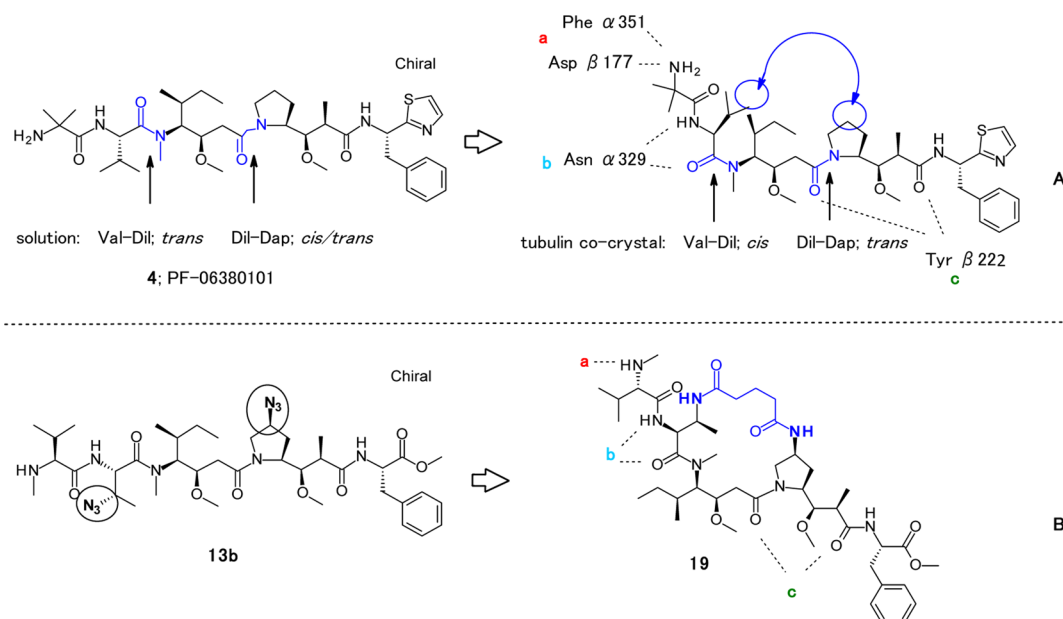
**Received:** February 5, 2018

**Accepted:** April 24, 2018

**Published:** May 15, 2018

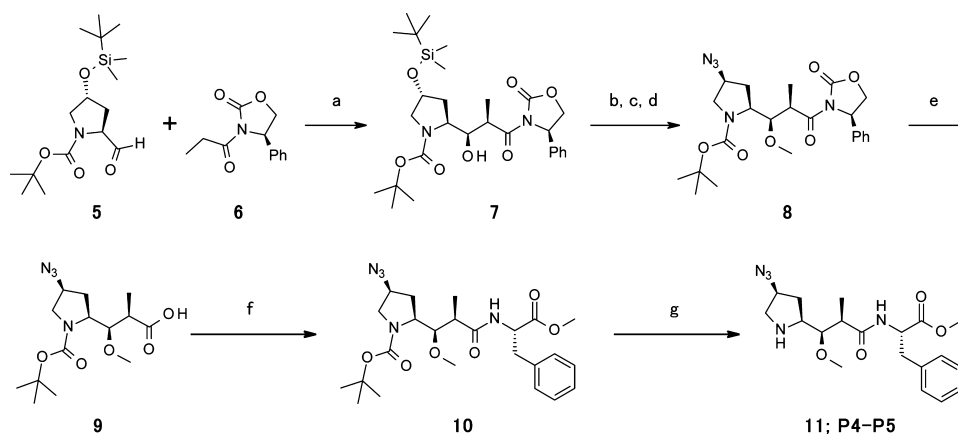


**Figure 2.** Modified central peptide strategy; structure of P2-modified and P4-modified dolastatin 10.



**Figure 3.** Critical binding spots of PF-06380101, 4 to tubulin (panel A), and design of macrocyclic analogue 19 (panel B).

### Scheme 1. Synthesis of the P4-Modified Dipeptide 11 (P4–P5)<sup>a</sup>



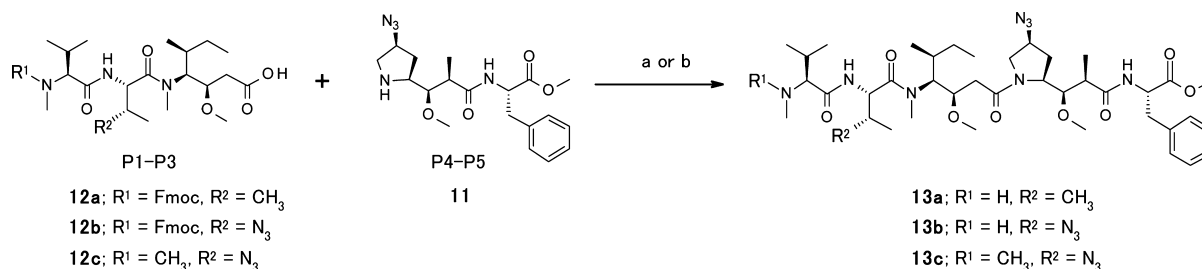
<sup>a</sup>(a) *n*-Bu<sub>2</sub>BOTf, *i*PrNEt<sub>3</sub>, CH<sub>2</sub>Cl<sub>2</sub>, −78 to 0 °C. (b) Me<sub>3</sub>OBf<sub>4</sub>, proton sponge, molecular sieves, 4 Å, EtOH, 0 °C to rt. (c) HF-Py, THF, 0 °C. (d) DPPA, DIAD, PPh<sub>3</sub>, THF, 0 °C to rt. (e) H<sub>2</sub>O<sub>2</sub>, LiOH, H<sub>2</sub>O, THF, 0 °C. (f) H-Phe-OMe, EDCl, HOBt, Et<sub>3</sub>N, CH<sub>2</sub>Cl<sub>2</sub>. (g) 4 M HCl–dioxane.

terminal P1 subunit and C-terminal P5 subunit modifications were performed by Pettit et al., Miyazaki et al., and other groups to investigate their structure–activity relationship (SAR).<sup>24–26</sup> In our recent report, we investigated the incorporation of heterocycles into the P5 subunit.<sup>27</sup> These P5 heterocycle modifications generated ADCs with more hydrophilic character and lower aggregation.

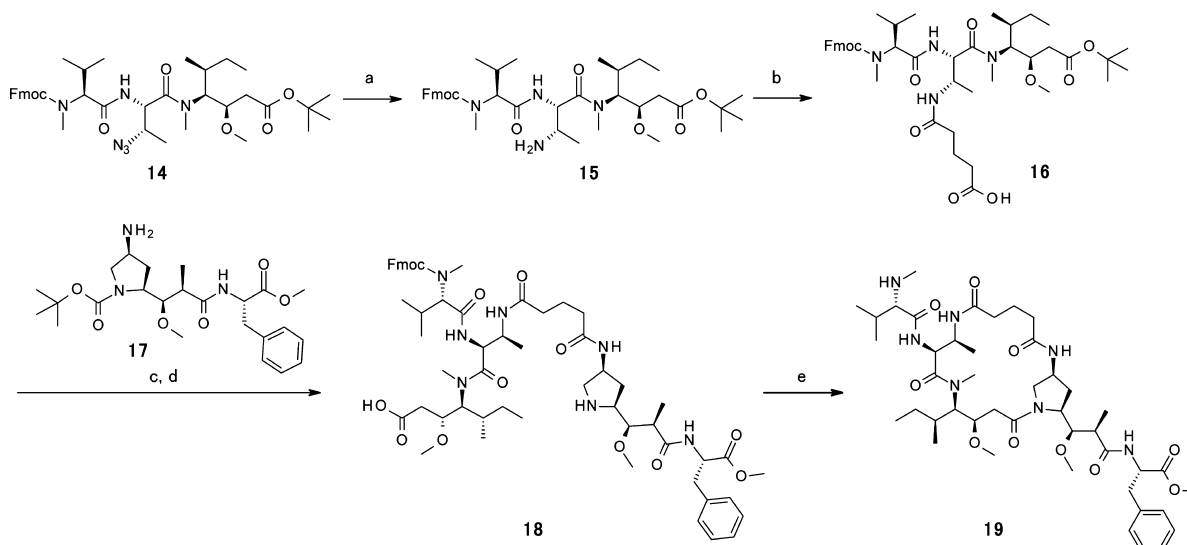
Modification of the core central peptides (P2–P4 subunits) has not been as extensively investigated because it has been reported that changes in these subunits result in attenuated compound potency.<sup>23</sup> In terms of the P2 subunit, substitution

of the natural valine residue with leucine or isoleucine is reported<sup>26</sup> and our group recently reported SAR results for analogues containing heteroatoms and other non-natural amino acids in the P2 position.<sup>28</sup> In contrast, regarding the P4 subunit, there are a few reports of analogues with mannose- and glucose-derived sugar amino acids as replacements for the Dap portion by Gajula et al.<sup>29,30</sup> and also hydroxyl, methoxy, and amino substituents on the pyrrolidine are described by Park et al.<sup>31</sup>

Discussed here are new linear dolastatin 10 analogues with an azide substituted P4 central amino acid and also in

Scheme 2. Synthesis of New Dolastatin Analogues 13a, 13b, and 13c<sup>a</sup>

<sup>a</sup>(a) EDCI, HOBT, Et<sub>3</sub>N, *N,N*-dimethylacetamide (DMAc). (b) EDCI, HOBT, Et<sub>3</sub>N, DMAc, then Et<sub>2</sub>NH.

Scheme 3. Synthesis of New Macrocylic Dolastatin Analogue 19<sup>a</sup>

<sup>a</sup>(a) H<sub>2</sub>, Pd/C, EtOH. (b) Glutaric anhydride, DMAc, 60 °C. (c) 1-[bis(dimethylamino)methylene]-1*H*-1,2,3-triazolo[4,5-*b*]pyridinium 3-oxid hexafluorophosphate (HATU), *i*PrNEt<sub>2</sub>, DMAc. (d) 4 M HCl–dioxane. (e) CMPI, *i*PrNEt<sub>2</sub>, EtOAc, then Et<sub>2</sub>NH.

combination with P2 azide substitution. These azide substitutions show changes in cytotoxicity and can serve as a handle for linker attachment or a tether for use in the synthesis of novel macrocyclic dolastatin 10 analogues (Figures 2 and 3).

## RESULTS AND DISCUSSION

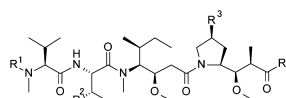
In our previous studies, we reported SAR results for auristatins containing specific P2 side chain modifications. We showed that P2-modified analogues, with the P2 units as 3*S*-amino-2*S*-aminobutyric acid (Abu (3-NH<sub>2</sub>)), 3*S*-azide-2*S*-aminobutyric acid (Abu (3-N<sub>3</sub>)), and 3*S*-hydroxyl-2*S*-aminobutyric acid (Abu (3-OH)), could display high *in vitro* cytotoxic activity (Figure 2).<sup>28</sup> Building from this previous work, we investigated whether central P4-modified analogues containing heteroatom substituents on the pyrrolidine ring retained cytotoxic potency with and without P2 heteroatom modifications. In addition, we examined the potency associated with a constrained macrocyclic dolastatin 10 analogue. The published PF-06380101 tubulin crystal data revealed a critical hydrogen bond network between Asp β177 and the amide carbonyl of Phe α351 with the protonated amino group of P1, a bifocal interaction of the N-2 valine with Asn α329, and also between the backbone amide of Tyr β222 with the terminal carbonyl groups of Dap and Doe (Figure 3A).<sup>32</sup> Moreover, the *cis* Val-Dil amide bond and *trans* Dil-Dap amide bond geometries were revealed in a cocrystal structure with tubulin (Figure 3A).<sup>32,33</sup> We presumed

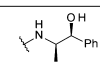
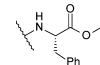
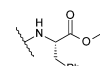
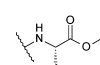
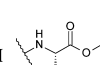
that a tether between the P2 and P4 subunits enables the formation of a macrocyclic structure that could force *cis/trans* geometry across these relevant bonds (Figure 3B).

The synthesis route of the azide-containing dipeptide intermediate is shown in Scheme 1. *syn*-Aldol product 7 was prepared from chiral proline aldehyde 5 and the boron enolate of chiral (*R*)-oxazolidinone 6 with moderate yield.<sup>31,34</sup> After methylation and *tert*-butyl dimethyl silyl deprotection, introduction of the azide group was performed by Mitsunobu-type chemistry using diphenylphosphoryl azide (DPPA) and diisopropyl azodicarboxylate (DIAD) to provide Boc-Dap (4-N<sub>3</sub>)-phenyloxazolidinone 8. Hydrolysis of the chiral auxiliary unit on compound 8 was accomplished using LiOH and H<sub>2</sub>O<sub>2</sub>, which led to the desired carboxylic acid Boc-Dap (4-N<sub>3</sub>)-OH 9. Boc-dipeptide 10 was formed using 1-ethyl-3-(3-dimethylaminopropyl) carbodiimide hydrochloride (EDCI) and 1-hydroxybenzotriazole hydrate (HOBT) to couple compound 9 with phenylalanine methyl ester (H-Phe-OMe). HCl-mediated Boc deprotection yielded compound 11.

Convergent syntheses of new linear dolastatin analogues are shown in Scheme 2. The P4–P5 intermediate, 11, and the separately prepared P1–P3 intermediates, 12a–c, were coupled in a modular manner to yield final dolastatin analogues 13a, 13b, and 13c.

The synthesis of a macrocyclic dolastatin derivative is shown in Scheme 3. The reduction of the azide group of the P1–P3

Table 1. Derivatives of P4 Central Modification with GI<sub>50</sub> Values and Properties


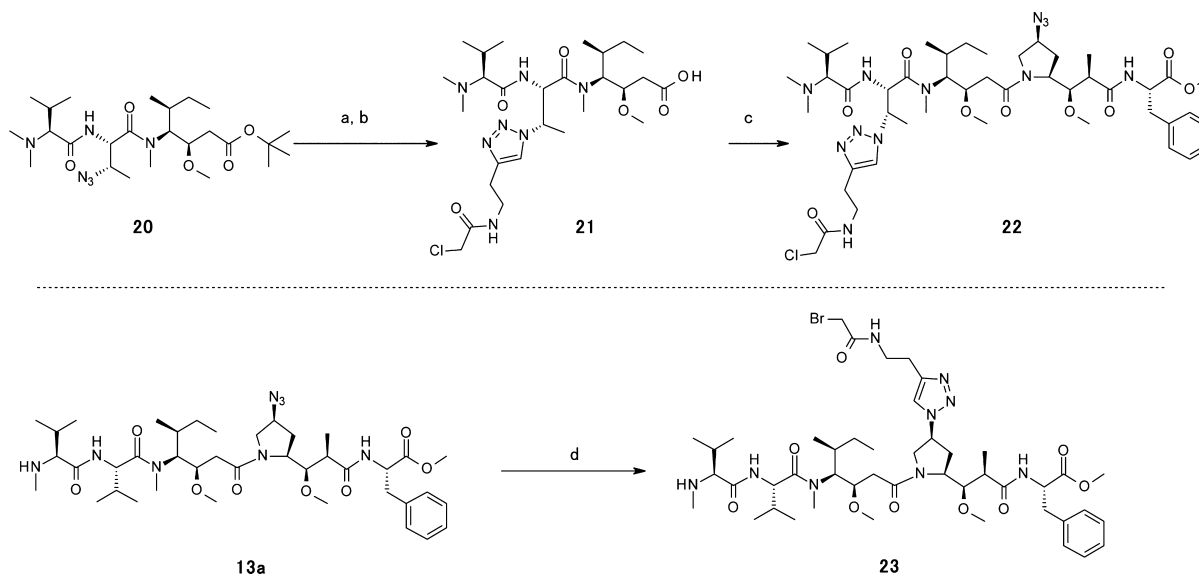
compound	R <sup>1</sup>	R <sup>2</sup> / R <sup>3</sup>	R <sup>4</sup>	GI <sub>50</sub> MOLM13 (nM) <sup>a</sup>	GI <sub>50</sub> SW780 (nM) <sup>a</sup>	Log D <sup>b</sup> pH 7.4 / 6.5	TPSA <sup>b</sup>
MMAE (2b)	H	CH <sub>3</sub> / H		0.274	0.343	2.03 / 1.15	149.5
13a	H	CH <sub>3</sub> / N <sub>3</sub>		0.224	1.90	2.69 / 1.81	180.3
13b	H	N <sub>3</sub> / N <sub>3</sub>		0.279	2.40	2.52 / 1.65	205.1
13c	CH <sub>3</sub>	N <sub>3</sub> / N <sub>3</sub>		0.057	0.365	4.28 / 3.65	196.3
19	H	NHCO(CH <sub>2</sub> ) <sub>3</sub> CONH		>1000	>1000	1.77 / 0.95	213.8

<sup>a</sup>Results are the average of two independent triplicate runs with compound purity above >90%. <sup>b</sup>log *D* and TPSA values were calculated with ACD/PhysChem Batch (version 12.01).

intermediate **14**, followed by the ring-opening reaction with glutaric anhydride, provided carboxylic intermediate **16**. The coupling reaction of carboxylic acid **16** and amine **17** followed by the deprotection of the Boc group and *tert*-Bu ester hydrolysis provided macrocyclic precursor **18**. Macrocyclization was achieved through reaction using the Mukaiyama condensation reagent, 2-chloro-1-methylpyridinium iodide (CMPI), under high dilution conditions (0.00014 M). Finally, the deprotection of the Fmoc group using Et<sub>3</sub>NH gave the desired macrocyclic analogue **19** in high yield (c–e, three steps; 61%).

Table 1 highlights the *in vitro* potency of each compound with respective P1, P2, and P4 unit modifications. Each compound was evaluated in an *in vitro* cell proliferation assay using an acute myeloid leukemia cell line (MOLM13) and a bladder cancer cell line (SW780) (see the Supporting Information). Compounds **2b**, **13a**, **13b**, and **13c** demonstrated similar GI<sub>50</sub> values (within 1 log) in both MOLM13 and SW780 cell lines; however, the SW780 cell line appeared less sensitive overall in this experiment. In our previous studies evaluating P2 modifications,<sup>28</sup> we determined that modification of the P2 side chain required a specific aromatic P5 subunit (containing an ester or amide functional group), which did not follow previously reported SAR trends for P2 valine-based auristatins.<sup>23</sup> On the basis of these results, we held the aromatic P5 ester units unchanged while examining the SAR of P4 azide introduction. Compound **13a**, containing an azide only in the P4 central peptide, demonstrated a similar activity *in vitro* when compared to MMAE (**2b**) in the MOLM13 cell line. Next, we explored combinations of both P4 and P2 azide modifications. Compound **13b**, with P2 Abu (3-N<sub>3</sub>) and a MeVal unit in the P1 position, showed a similar GI<sub>50</sub> value in the MOLM13 cell line compared to **13a** (0.279 and 0.224 nM, respectively). Interestingly, compound **13c**, with P2 and P4 azide modifications and Dov in the P1 position, showed enhanced cytotoxic activity (GI<sub>50</sub> = 0.057 nM) compared to compound **13b**. One possible explanation of the cytotoxicity is that

analogue **13c**, the pentapeptide containing two azide groups in the P2 and P4 positions, could take a more favorable conformation by electric and steric effects arising from the azide groups.<sup>35,36</sup> To date, it was known that structural changes in the dolastatin Dap P4 subunit did not significantly compromise the cytotoxic potency, even though the P1 unit was Dov.<sup>29–31</sup> However, our research revealed that the modification to the Dap moiety has shown to be beneficial in improving the potency. In contrast, the macrocyclic analogue **19** showed a low *in vitro* potency, and these differences in cytotoxicity may be explained by differences in cellular membrane permeability reflected in the compound calculated hydrophilicity measurements, log *D* pH 7.4/6.5 and topological polar surface area (TPSA) (Table 1). This inability for some molecules to cross the cell membrane changes when they are attached to an antibody as antibodies can transport the attached molecule across the cell membrane and have the ability to release the payload inside the cell. This has been shown for MMAF (monomethyl auristatin F)-derived ADCs, which are highly potent despite the low membrane permeability of MMAF (Figure 1).<sup>28,37</sup> Further investigation is needed to determine whether an ADC composed of a drug linker prepared from compound **19** demonstrates cytotoxicity. Additionally, <sup>1</sup>H NMR data gathered for macrocyclic analogue **19** did not show *cis/trans* amide signals found in the linear compound **13b** (see the Supporting Information), indicating a constrained amide bond geometry and relative enhancement of rigidity. Given the more rigid structure of the macrocyclic analogue compared to the linear analogue, it may also be possible that macrocyclic analogue **19** was forced into a less favorable conformation by the relatively rigid structure. Studies are ongoing to elucidate a cocrystal structure of tubulin with compound **13a** and with compound **19** to reveal the conformation of the Val-Dil and Dil-Dap amide bond geometries and the binding mode with tubulin to enable further design optimizations and investigation of tethers for preparation of potent macrocyclic dolastatin 10 analogues.

Scheme 4. Preparation of Drug Linkers 22 and 23 with P2 or P4 Linkage<sup>a</sup>

<sup>a</sup>(a) *N*-(But-3-yn-1-yl)-2-chloroacetamide, CuBr, *i*PrNEt<sub>2</sub>, dimethylformamide (DMF). (b) 4 M HCl–dioxane. (c) EDCl, HOBT, Et<sub>3</sub>N, DMAc. (d) *N*-(But-3-yn-1-yl)-2-bromoacetamide, CuBr, *i*PrNEt<sub>2</sub>, DMF.

Through this work, we have also confirmed that azide groups introduced into the P2 and P4 subunits demonstrate the possibility for serving as handles for linker attachment.

The synthesis of drug linkers using a P2 or P4 linkage is shown in Scheme 4. The click reaction<sup>38</sup> of compound 20 and *N*-(but-3-yn-1-yl)-2-chloroacetamide proceeded smoothly to yield compound 21. Deprotection of the Boc group and subsequent coupling to the P4–P5 dimer 11 yielded the desired drug linker 22. Payload 13a, containing the P4 azide group, gave drug linker 23 directly using the click reaction with *N*-(but-3-yn-1-yl)-2-bromoacetamide. These examples, using noncleavable linkers, suggest that payloads that incorporate azide groups could serve as linker attachment sites for other noncleavable or cleavable linkers. Thus, compounds 13a–c are promising payloads for preparation of novel drug linkers at the P2 or P4 position.

## CONCLUSIONS

We revealed that the modification of the Dap unit has proven to be beneficial in improving potency. Compounds with a P4-modified azide group on the pyrrolidine ring demonstrate cytotoxic activity *in vitro*. In particular, the dolastatin 10 analogues with azide modifications on both P2 and P4 showed enhanced cytotoxic activity ( $GI_{50} = 0.057$  nM) when compared to MMAE. On the contrary, the macrocyclic analogue showed poor potency. Possible explanations of these cytotoxicity values were due to hydrophilicity and conformation of these pentapeptides. Additional work is now under way to further reveal the amide bond geometries and the binding mode of these compounds. We also demonstrated that the analogue containing an azide group can serve as sites of linker attachment in the preparation of ADCs.

## EXPERIMENTAL SECTION

Synthesis strategies for assembling the compounds described herein as well as the purification strategies and analytical methods employed mirror a recent report by our research group.<sup>28</sup>

**General Methods.** NMR spectra were obtained on a Bruker AV 500, 400, or 300 MHz spectrometer at 25 °C. All NMR spectra were referenced to the DMSO-*d*<sub>6</sub> residual solvent peak (<sup>1</sup>H: 2.50 ppm; <sup>13</sup>C: 39.5 ppm).

High-resolution mass spectrometry (HRESIMS) samples were directly injected into a Dionex 3000-Orbitrap Velos LC-MS.

Flash column chromatography was carried out using prepacked Yamazen Universal columns on a Yamazen purification system. Preparative high-performance liquid chromatography (HPLC) was conducted with a Phenomenex Gemini-NX 10 μm, C18 110 Å column (150 × 30 mm<sup>2</sup>) using a 5–95% gradient of acetonitrile/0.05% aqueous trifluoroacetic acid (TFA) mixture over 13 min unless another column or solvent system is noted. Preparative HPLC-purified compounds were assumed to be salts containing one molecule of TFA.

Liquid chromatography–mass spectrometry (LC–MS) data was acquired using an Acquity UPLC BEH C8 1.7 μm 2.1 × 50 mm<sup>2</sup> column, 40 °C. 0–0.5 min: isocratic 85:5:10 H<sub>2</sub>O/MeCN/0.5% TFA in H<sub>2</sub>O; 0.5–1.6 min: linear gradient 85:5:10 H<sub>2</sub>O/MeCN/0.5% TFA in H<sub>2</sub>O to 98:2 MeCN/0.5% TFA in H<sub>2</sub>O; 1.60–1.9 min linear gradient 98:2 MeCN/0.5% TFA in H<sub>2</sub>O to 85:5:10 H<sub>2</sub>O/MeCN/0.5% TFA in H<sub>2</sub>O; 1.9–2.0 isocratic 85:5:10 H<sub>2</sub>O/MeCN/0.5% TFA in H<sub>2</sub>O.<sup>28</sup>

**Materials.** All solvents and reagents were purchased from commercial sources and used without further purification.

**Synthetic Procedures.** Reactions were typically carried out at ambient room temperature (rt) with exposure to air, unless otherwise noted.

**Compound 8.** To a stirred solution of (4*R*)-4-phenyl-3-propanoyl-1,3-oxazolidin-2-one (1.33 g, 6.06 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (36 mL) that was cooled to 0 °C, *n*-Bu<sub>2</sub>BOTf (1.78 mL, 8.27 mmol) and *N,N*-diisopropylethylamine (DIEA) (1.5 mL, 8.43 mmol) were added and then stirred for 45 min. The resulting solution was cooled to –78 °C, added dropwise of a solution of aldehyde 5 (1.82 g, 5.51 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (36 mL), and stirred for 1 h. Stirring was then further conducted at 0 °C for an additional 1 h. Analysis by LC–MS showed that the reaction

was complete (high diastereoselectivity; see the [Supporting Information](#)). The reaction was terminated with methanol and saturated sodium bicarbonate. The reaction mixture was extracted with  $\text{CH}_2\text{Cl}_2$ . The combined organic fractions were washed with brine, dried over a pad of magnesium sulfate, filtered, and concentrated in vacuo. The residue was purified by flash chromatography on silica gel (40  $\mu\text{m}$ , 60 Å, 3.0  $\times$  16.5 cm) using 0–5% MeOH in  $\text{CH}_2\text{Cl}_2$  as the eluent. A total of 1.4 g of compound **7** was obtained (2.55 mmol, 46%) as a white amorphous solid. LC–MS  $t_{\text{R}}$  = 1.83 min; ESIMS  $m/z$  549.33  $[\text{M} + \text{H}]^+$

To a stirred solution of compound **7** (0.114 g, 0.21 mmol), proton sponge (0.325 g, 1.51 mmol), and molecular sieves (4 Å) in  $\text{CH}_2\text{Cl}_2$  (3.5 mL) at 5 °C was added  $\text{Me}_3\text{OBF}_4$  (0.219 g, 1.48 mmol). The reaction mixture was stirred at rt. After 68 h, analysis by LC–MS showed that the reaction was complete. The reaction solution was filtered and concentrated in vacuo. The residue was purified by flash chromatography on silica gel (30  $\mu\text{m}$ , 60 Å, 2.3  $\times$  12.3  $\text{cm}^2$ ) using 2–25% EtOAc in hexanes as the eluent. A total of 0.073 g of Boc-Dap (4-OTBS)-phenyloxazolidinone was obtained (0.13 mmol, 63%) as a white amorphous solid.

To a stirred solution of Boc-Dap (4-OTBS)-phenyloxazolidinone (0.073 g, 0.13 mmol) in tetrahydrofuran (THF) (3 mL) was slowly added HF-Py (140  $\mu\text{L}$ , 1.56 mmol) and then stirred at rt. After 5 h, analysis by LC–MS showed that the reaction was complete. The reaction mixture was added to 30 mL of saturated sodium bicarbonate and then extracted with EtOAc. The organic layers were washed with 30 mL of 1 N HCl, dried over anhydrous magnesium sulfate, concentrated in vacuo, and dried further under high vacuum. A total of 58 mg of Boc-Dap (4-OH)-phenyloxazolidinone was obtained (0.13 mmol, quant.) as a white amorphous solid, which was used without further purification.

To a stirred solution of Boc-Dap (4-OH)-phenyloxazolidinone (58 mg, 0.13 mmol) in THF (2 mL) at 5 °C were added DPPA (42  $\mu\text{L}$ , 0.20 mmol), DIAD (77  $\mu\text{L}$ , 0.39 mmol), and  $\text{PPh}_3$  (103 mg, 0.20 mmol). After 15 h, analysis by LC–MS showed that the reaction was complete. The reaction solution was concentrated in vacuo, and the residue was purified by flash chromatography on silica gel (30  $\mu\text{m}$ , 60 Å, 2.3  $\times$  12.3  $\text{cm}^2$ ) using 2–50% EtOAc in hexanes as the eluent. A total of 37 mg of compound **8** was obtained (0.078 mmol, 59%) as a white amorphous solid.  $^1\text{H}$  NMR (400 MHz,  $\text{DMSO}-d_6$ ):  $\delta$  7.42–7.24 (m, 5H), 5.49 (dd,  $J$  = 8.6, 3.6 Hz, 1H), 4.73 (t,  $J$  = 8.7 Hz, 1H), 4.20–4.08 (m, 2H), 4.00–3.76 (m, 4H), 2.88 (s, 3H), 2.87–2.74 (m, 1H), 2.39–2.23 (m, 1H), 2.05–1.83 (m, 1H), 1.39 (s, 9H), 1.03 (d,  $J$  = 6.0 Hz, 3H);  $^{13}\text{C}$  NMR (101 MHz,  $\text{DMSO}-d_6$ ):  $\delta$  174.77, 153.97, 139.97, 129.02, 128.24, 126.21, 83.35, 79.61, 70.22, 60.54, 57.71, 57.60, 57.31, 51.07, 50.56, 46.39, 29.86, 28.46, 14.23; LC–MS  $t_{\text{R}}$  = 1.66 min; ESIMS  $m/z$ : 474.30  $[\text{M} + \text{H}]^+$ ; HRESIMS  $m/z$ : 474.2355  $[\text{M} + \text{H}]^+$  calcd for  $\text{C}_{23}\text{H}_{32}\text{N}_5\text{O}_6$ , 474.2347.

**Compound 10.** To a stirred solution of Boc-Dap (4- $\text{N}_3$ )-phenyloxazolidinone (241 mg, 0.51 mmol) in THF (10 mL) at 5 °C were added 30%  $\text{H}_2\text{O}_2$  (0.81 mL, 8.14 mmol) and 0.5 M LiOH (5 mL, 2.50 mmol). After 15 h, analysis by LC–MS showed that the reaction was complete. The reaction was quenched with 1 M sodium thiosulfate by stirring for 10 min and then extracted with saturated sodium bicarbonate and  $\text{CH}_2\text{Cl}_2$ , which was cooled to 0–5 °C previously. The aqueous layer was adjusted to pH 2, followed by extraction with EtOAc (three times). The combined organic layer was dried over

anhydrous magnesium sulfate and concentrated in vacuo to afford Boc-Dap (4- $\text{N}_3$ )-OH **9** as a white solid, which was used without further purification.

To a stirred suspension of H-Phe-OMe HCl salt (54 mg, 0.250 mmol), Boc-Dap (4- $\text{N}_3$ )-OH **9** (90 mg, 0.274 mmol), EDCI (75 mg, 0.391 mmol), and HOBT (40 mg, 0.261 mmol) in  $\text{CH}_2\text{Cl}_2$  (2 mL) was added  $\text{Et}_3\text{N}$  (60  $\mu\text{L}$ , 0.430 mmol). After 2 h, analysis by LC–MS showed that the reaction was complete. The mixture was purified by preparatory reversed-phase (RP)-HPLC with a Phenomenex Gemini-NX 10  $\mu\text{m}$ , C18 110 Å column (150  $\times$  30  $\text{mm}^2$ ) using 5–95% MeCN in 0.05% aqueous TFA solution as the eluent. A total of 50 mg of the title compound was obtained as the TFA salt (0.102 mmol, 41%) as a white amorphous solid.  $^1\text{H}$  NMR (400 MHz,  $\text{DMSO}-d_6$ ):  $\delta$  8.32 (d,  $J$  = 8.0 Hz, 1H), 7.31–7.16 (m, 5H), 4.49 (ddd,  $J$  = 9.7, 7.9, 5.3 Hz, 1H), 4.16–4.02 (m, 1H), 3.95–3.71 (m, 3H), 3.61 (s, 3H), 3.18 (s, 3H), 3.05 (dd,  $J$  = 13.7, 5.4 Hz, 1H), 2.95–2.71 (m, 2H), 2.33–2.12 (m, 2H), 1.95–1.75 (m, 1H), 1.41 (s, 9H), 0.73 (d,  $J$  = 6.8 Hz, 3H);  $^{13}\text{C}$  NMR (101 MHz,  $\text{DMSO}-d_6$ ):  $\delta$  174.33, 172.42, 153.28, 137.74, 129.55, 128.56, 126.90, 82.39, 79.50, 60.58, 57.57, 57.21, 53.72, 52.20, 50.94, 50.35, 43.15, 37.19, 29.79, 28.49, 14.63; LC–MS  $t_{\text{R}}$  = 1.61 min, ESIMS  $m/z$  490.45  $[\text{M} + \text{H}]^+$ ; HRESIMS  $m/z$ : 490.2671  $[\text{M} + \text{H}]^+$  calcd for  $\text{C}_{24}\text{H}_{36}\text{N}_5\text{O}_6$ , 490.2660.

**Compound 11.** To 4.0 M HCl in dioxane (1 mL, 4 mmol) solution was added Boc-Dap (4- $\text{N}_3$ )-Phe-OMe **10** (24 mg, 0.05 mmol) and stirred for 2 h until analysis by LC–MS showed that the boc-deprotection reaction was complete. The reaction solution was concentrated in vacuo and dried further under high vacuum. A total of 20.5 mg of the title compound (0.05 mmol, 98%) was obtained as the HCl salt as a pale brown solid, which was used without further purification.  $^1\text{H}$  NMR (400 MHz,  $\text{DMSO}-d_6$ ):  $\delta$  8.62 (d,  $J$  = 8.1 Hz, 1H), 8.55 (s, 1H), 7.33–7.17 (m, 5H), 4.60–4.43 (m, 2H), 3.78–3.57 (m, 4H), 3.46–3.35 (m, 1H), 3.33–3.25 (m, 4H), 3.16–3.05 (m, 2H), 2.88 (dd,  $J$  = 13.8, 9.9 Hz, 1H), 2.49–2.34 (m, 2H), 1.76 (ddd,  $J$  = 13.3, 9.9, 6.1 Hz, 1H), 0.73 (d,  $J$  = 7.0 Hz, 3H); LC–MS  $t_{\text{R}}$  = 1.03 min, ESIMS  $m/z$ : 390.32  $[\text{M} + \text{H}]^+$ ; HRESIMS  $m/z$ : 390.2146  $[\text{M} + \text{H}]^+$  calcd for  $\text{C}_{19}\text{H}_{28}\text{N}_5\text{O}_4$ , 390.2136.

**Compound 13a.** To a stirred solution of H-Dap (4- $\text{N}_3$ )-Phe-OMe **11** (7 mg, 0.02 mmol), Fmoc-MeVal-Val-Dil-OH **12a** (11 mg, 0.016 mmol), and  $\text{Et}_3\text{N}$  (8  $\mu\text{L}$ , 0.06 mmol) in DMAc (1 mL) were added EDCI (7 mg, 0.04 mmol) and HOBT (2 mg, 0.01 mmol), and the mixture was stirred for 15 h. To the mixture was added  $\text{Et}_2\text{NH}$  (40  $\mu\text{L}$ , 0.387 mmol). After 1 h, analysis by LC–MS showed that the reaction was complete. The reaction mixture was diluted with  $\text{H}_2\text{O}$  and DMAc, and then the mixture was purified by preparatory RP-HPLC with a Phenomenex Gemini-NX 10  $\mu\text{m}$ , C18 110 Å column (150  $\times$  30  $\text{mm}^2$ ) using 5–95% MeCN in 0.05% aqueous TFA solution as the eluent. A total of 6 mg of compound **13a** was obtained as the TFA salt (6.66  $\mu\text{mol}$ , 41%) as a white amorphous solid.  $^1\text{H}$  NMR (400 MHz,  $\text{DMSO}-d_6$ ): a complex spectrum was observed, presumably because of cis/trans conformational isomers:  $\delta$  8.90–8.75 (m, 2H), [8.42 (d,  $J$  = 7.9 Hz), 8.32 (d,  $J$  = 8.0 Hz) 1H], 7.31–7.17 (m, 5H), 4.72–4.63 (m, 1H), 4.59 (t,  $J$  = 8.5 Hz, 1H), 4.48 (ddd,  $J$  = 9.8, 8.0, 5.4 Hz, 1H), 4.14–4.02 (m, 2H), 4.00–3.63 (m, 5H), [3.61 (s), 3.60 (s) 3H], [3.19 (s), 3.17 (s) 3H], 3.12 (s, 3H), 3.07–2.96 (m, 4H), 2.88 (dd,  $J$  = 13.7, 9.7 Hz, 1H), 2.47 (t,  $J$  = 5.0 Hz, 3H), 2.32–2.16 (m, 3H), 2.12–1.94 (m, 2H), 1.92–1.66 (m, 2H), 1.34–1.19 (m, 1H), 0.98–0.83 (m, 17H), 0.80–0.70 (m, 6H);  $^{13}\text{C}$  NMR (75 MHz,  $\text{DMSO}$ ):  $\delta$  174.38, 173.90, 172.44, 169.46,

167.30, 166.43, 164.91, 137.73, 129.55, 128.98, 128.57, 128.30, 126.91, 85.77, 82.20, 80.61, 78.65, 65.99, 63.66, 60.12, 59.02, 57.89, 57.59, 57.35, 56.07, 55.30, 55.13, 53.72, 52.20, 50.97, 49.36, 43.32, 38.89, 37.48, 37.26, 32.46, 32.09, 32.01, 30.51, 29.90, 25.85, 25.13, 21.50, 20.63, 18.96, 18.90, 18.73, 18.05, 15.95, 14.67, 10.73; LC-MS  $t_R = 1.31$  min, ESIMS  $m/z$ : 787.72  $[M + H]^+$ ; HRESIMS  $m/z$ : 787.5096  $[M + H]^+$  calcd for  $C_{40}H_{67}N_8O_8$ , 787.5076.

**Compound 13b.** To a stirred solution of H-Dap ( $N_3$ )-Phe-OMe **11** (18 mg, 0.04 mmol), Fmoc-MeVal-Abu (3- $N_3$ )-Dil-OH **12b** (26 mg, 0.037 mmol), and  $Et_3N$  (20  $\mu$ L, 0.14 mmol) in DMAc (2 mL, 16.23 mmol) were added EDCI (18 mg, 0.09 mmol) and HOBt (4 mg, 0.03 mmol), and the mixture was stirred for 15 h. To the mixture was added  $Et_2NH$  (40  $\mu$ L, 0.386 mmol). After 3 h, analysis by LC-MS showed that the reaction was complete. The reaction mixture was diluted with  $H_2O$  and DMAc, and then the mixture was purified by preparatory RP-HPLC with a Phenomenex Gemini-NX 10  $\mu$ m, C18 110  $\text{\AA}$  column (150  $\times$  30 mm<sup>2</sup>) using 5–95% MeCN in 0.05% aqueous TFA solution as the eluent. A total of 22 mg of compound **13b** was obtained as a TFA salt (0.024 mmol, 64%) as a white amorphous solid. <sup>1</sup>H NMR (400 MHz, DMSO- $d_6$ ): a complex spectrum was observed, presumably because of cis/trans conformational isomers):  $\delta$  [9.10 (d,  $J = 8.6$  Hz), 8.71 (d,  $J = 9.1$  Hz) 1H], 8.90 (br-s, 1H), [8.41 (d,  $J = 7.9$  Hz), 8.33 (d,  $J = 8.0$  Hz) 1H], [7.49–7.33 (m), 7.32–7.07 (m), 5H], 4.86 (t,  $J = 8.3$  Hz, 1H), 4.59 (br-s, 1H), 4.48 (ddd,  $J = 9.7, 7.9, 5.4$  Hz, 1H), 4.33–3.86 (m, 6H), 3.78–3.66 (m, 1H), [3.61 (s), 3.60 (s) 3H], [3.21 (s), 3.18 (s), 3H], [3.13 (s), 3.09 (s), 3H], 3.09–3.01 (m, 2H), 2.98 (s, 3H), 2.93–2.81 (m, 1H), 2.49–2.43 (m, 3H), 2.36–2.17 (m, 3H), 2.09 (dq,  $J = 13.1, 6.5$  Hz, 1H), 1.94–1.64 (m, 2H), 1.36–1.21 (m, 4H), 0.99–0.83 (m, 11H), 0.83–0.66 (m, 6H); <sup>13</sup>C NMR (101 MHz, DMSO)  $\delta$  174.38, 172.43, 170.27, 169.45, 167.86, 166.84, 137.79, 137.73, 129.54, 128.80, 128.56, 126.90, 124.86, 80.61, 77.61, 65.98, 60.16, 58.23, 57.90, 57.81, 57.34, 53.71, 52.80, 52.18, 51.44, 43.31, 37.19, 32.67, 32.46, 29.82, 28.82, 25.85, 18.65, 18.22, 18.04, 16.08, 15.97, 14.65, 13.42, 10.94; LC-MS  $t_R = 1.32$  min, ESIMS  $m/z$ : 814.66  $[M + H]^+$ ; HRESIMS  $m/z$ : 814.4972  $[M + H]^+$  calcd for  $C_{39}H_{64}N_{11}O_8$ , 814.4934.

**Compound 13c.** To a stirred solution of H-Dap ( $N_3$ )-Phe-OMe **11** (20 mg, 0.04 mmol), Dov-Val-Dil-OH **12c** (20 mg, 0.041 mmol), and  $Et_3N$  (25  $\mu$ L, 0.18 mmol) in DMAc (2 mL) were added EDCI (15 mg, 0.08 mmol) and HOBt (5 mg, 0.03 mmol). After 15 h, analysis by LC-MS showed that the reaction was complete. The reaction mixture was diluted with  $H_2O$  and DMAc, and then the mixture was purified by preparatory RP-HPLC with a Phenomenex Gemini-NX 10  $\mu$ m, C18 110  $\text{\AA}$  column (150  $\times$  30 mm<sup>2</sup>) using 5–95% MeCN in 0.05% aqueous TFA solution as the eluent. A total of 22 mg of compound **13c** was obtained as the TFA salt (0.023 mmol, 58%) as a white amorphous solid. <sup>1</sup>H NMR (400 MHz, DMSO- $d_6$ ): a complex spectrum was observed, presumably because of cis/trans conformational isomers):  $\delta$  9.17 (d,  $J = 8.5$  Hz, 1H), [8.41 (d,  $J = 8.0$  Hz), 8.32 (d,  $J = 8.0$  Hz) 1H], 7.34–7.08 (m, 5H), 4.89 (t,  $J = 8.1$  Hz, 1H), 4.59 (br-s, 1H), 4.48 (ddd,  $J = 9.7, 7.9, 5.4$  Hz, 2H), 4.18–3.89 (m, 3H), 3.81–3.65 (m, 1H), [3.61 (s), 3.60 (s), 3H], 3.29–3.25 (m, 1H), [3.20 (s), 3.18 (s), 3H], [3.13 (s), 3.09 (s), 3H], 3.08–2.94 (m, 7H), 2.88 (dd,  $J = 13.7, 9.7$  Hz, 1H), [2.78 (s), 2.75 (s), 6H], 2.35–2.17 (m, 3H), 1.94–1.69 (m, 2H), 1.38–1.22 (m, 4H), 0.96 (d,  $J = 6.8$  Hz, 3H), 0.93–0.81 (m, 8H), 0.82–0.71 (m, 6H); <sup>13</sup>C NMR (126 MHz, DMSO):  $\delta$  174.39, 173.54, 172.45, 170.38,

169.45, 165.99, 137.74, 129.56, 128.58, 126.92, 80.59, 77.59, 71.82, 60.17, 58.38, 58.24, 58.15, 58.06, 57.90, 57.83, 57.75, 57.33, 56.90, 53.72, 52.70, 52.26, 52.21, 51.45, 46.15, 43.31, 42.17, 41.50, 37.41, 37.19, 32.52, 31.94, 28.82, 28.15, 27.00, 25.85, 19.63, 19.55, 16.91, 16.08, 16.00, 15.90, 14.83, 14.67, 14.45, 12.39, 11.03, 10.92, 9.05; LC-MS  $t_R = 1.33$  min, ESIMS  $m/z$ : 828.74  $[M + H]^+$ ; HRESIMS  $m/z$ : 828.5119  $[M + H]^+$  calcd for  $C_{40}H_{66}N_{11}O_8$ , 828.5090.

**Compound 15.** To a stirred solution of Fmoc-MeVal-Abu (3- $N_3$ )-Dil-OtBu (130 mg, 0.18 mmol) in EtOH (3 mL) was added Pd on carbon (150 mg, 0.070 mmol) under a nitrogen atmosphere. The reaction mixture was stirred under a hydrogen atmosphere. After 4 h, analysis by LC-MS indicated that the reaction was complete. After insoluble materials were removed by filtration, the filtrate was concentrated under reduced pressure. The residue was purified by preparatory RP-HPLC with a Phenomenex Gemini-NX 10  $\mu$ m, C18 110  $\text{\AA}$  column (150  $\times$  30 mm<sup>2</sup>) using 5–95% MeCN in 0.05% aqueous TFA solution as the eluent. A total of 90 mg of compound **15** was obtained as the TFA salt (0.13 mmol, 72%) as a white amorphous solid. <sup>1</sup>H NMR (400 MHz, DMSO- $d_6$ ):  $\delta$  8.66 (d,  $J = 8.3$  Hz, 1H), 7.94–7.81 (m, 4H), 7.64 (d,  $J = 7.6$  Hz, 2H), 7.47–7.38 (m, 2H), 7.37–7.26 (m, 2H), 4.86 (t,  $J = 7.5$  Hz, 1H), 4.56–4.19 (m, 4H), 3.88–3.77 (m, 1H), 3.57–3.44 (m, 1H), 3.28 (s, 3H), 2.99–2.74 (m, 6H), 2.60–2.53 (m, 1H), 2.28–2.00 (m, 2H), 1.79 (br-s, 1H), 1.48–1.27 (m, 10H), 1.22–1.08 (m, 3H), 0.97–0.66 (m, 14H); <sup>13</sup>C NMR (101 MHz, DMSO- $d_6$ ):  $\delta$  171.27, 170.78, 170.01, 156.49, 144.28, 144.18, 141.24, 128.10, 127.51, 125.41, 120.58, 80.51, 78.47, 67.18, 63.77, 57.96, 51.48, 48.05, 47.19, 38.67, 33.0, 30.08, 28.15, 28.07, 27.25, 25.69, 19.41, 19.19, 16.25, 14.90, 11.04; LC-MS  $t_R = 1.65$  min, ESIMS  $m/z$ : 695.65  $[M + H]^+$ ; HRESIMS  $m/z$ : 695.4350  $[M + H]^+$  calcd for  $C_{39}H_{59}N_4O_7$ , 695.4378.

**Compound 16.** To a stirred solution of Fmoc-MeVal-Abu (3- $NH_2$ )-Dil-OtBu (43 mg, 0.06 mmol) in DMAc (1 mL) was added glutaric anhydride (7 mg, 0.06 mmol). The reaction mixture was stirred at 60 °C. After 4 h, analysis by LC-MS showed the reaction was complete. The mixture was purified by preparatory RP-HPLC with a Phenomenex Gemini-NX 10  $\mu$ m, C18 110  $\text{\AA}$  column (150  $\times$  30 mm<sup>2</sup>) using 5–95% MeCN in aqueous TFA solution as the eluent. A total of 40 mg of compound **16** was obtained as the TFA salt (0.043 mmol, 70%) as a white amorphous solid. <sup>1</sup>H NMR (400 MHz, DMSO- $d_6$ ):  $\delta$  11.99 (s, 1H), 8.17 (d,  $J = 8.5$  Hz, 1H), 7.90 (d,  $J = 7.6$  Hz, 2H), 7.79–7.60 (m, 3H), 7.42 (td,  $J = 7.6, 1.2$  Hz, 2H), 7.32 (t,  $J = 7.3$  Hz, 2H), 4.98–4.86 (m, 1H), 4.56–4.18 (m, 5H), 4.14–4.01 (m, 2H), 3.90–3.63 (m, 1H), 3.22 (d,  $J = 8.7$  Hz, 3H), 2.99 (s, 3H), 2.78 (s, 3H), 2.21–2.13 (m, 2H), 2.19–1.89 (m, 4H), 1.87–1.60 (m, 4H), 1.40 (s, 9H), 1.37–1.29 (s, 2H), 1.08–0.57 (m, 14H); LC-MS  $t_R = 1.81$  min, ESIMS  $m/z$ : 809.70  $[M + H]^+$ .

**Compound 17.** To a stirred solution of Boc-Dap (4- $N_3$ )-Phe-OMe (35 mg, 0.06 mmol) in EtOH (2 mL) was added Pd on carbon (30 mg, 0.028 mmol) under a nitrogen atmosphere. The reaction mixture was stirred under a hydrogen atmosphere. After 4 h, analysis by LC-MS showed that the reaction was complete. After insoluble materials were removed by filtration, the filtrate was concentrated under reduced pressure.

A total of 26 mg of compound **17** was obtained (0.06 mmol, 97%) as a white amorphous solid, which was used without further purification. <sup>1</sup>H NMR (400 MHz, DMSO- $d_6$ ):  $\delta$  8.30 (d,  $J = 7.9$  Hz, 1H), 8.00 (s, 2H), 7.35–7.12 (m, 5H), 4.52

(ddd,  $J = 9.6, 7.9, 5.4$  Hz, 1H), 4.01–3.39 (m, 7H), 3.21 (s, 3H), 3.05 (dd,  $J = 13.8, 5.4$  Hz, 2H), 2.89 (dd,  $J = 13.7, 9.6$  Hz, 1H), 2.38–2.07 (m, 2H), 1.88 (br-s, 1H), 1.42 (s, 9H), 0.75 (d,  $J = 6.6$  Hz, 3H); LC–MS  $t_R = 1.13$  min, ESIMS  $m/z$ : 464.45  $[M + H]^+$ ; HRESIMS  $m/z$ : 464.2790  $[M + H]^+$  calcd for  $C_{24}H_{37}N_3O_6$ , 464.2755.

**Compound 18.** To a stirred solution of Fmoc-MeVal-Abu (3-NHCOCH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>COOH)-Dil-OtBu (61 mg, 0.08 mmol), Boc-Dap (4-NH<sub>2</sub>)-Phe-OMe (33 mg, 0.07 mmol), and DIEA (37  $\mu$ L, 0.21 mmol) in DMAc (1.5 mL) was added HATU (70 mg, 0.184 mmol). After 4 h, analysis by LC–MS showed that the reaction was complete. To the mixture was added 1 N HCl aq., and then the mixture was stirred for 1 h. After separation, the organic layer was washed with saturated sodium bicarbonate and brine and dried over anhydrous magnesium sulfate. The organic layer was concentrated under reduced pressure, and the residue was dried further under high vacuum. A total of 95 mg of enriched Fmoc-MeVal-Abu (3-NHCOCH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CO-[Boc-Dap (4-NH<sub>2</sub>)]-Phe-OMe)-Dil-OtBu was obtained as a white amorphous solid, which was used without further purification.

A solution of Fmoc-MeVal-Abu (3-NHCOCH<sub>2</sub>CH<sub>2</sub>CH<sub>2</sub>CO-[Boc-Dap (4-NH<sub>2</sub>)]-Phe-OMe)-Dil-OtBu (90 mg, 0.07 mmol) in 4 M HCl in dioxane (3 mL, 12 mmol) was stirred at rt. After 2 h, analysis by LC–MS showed that the reaction was complete. The crude reaction mixture was concentrated under reduced pressure and dried further under high vacuum. The residue was purified by preparatory RP-HPLC with a Phenomenex Gemini-NX 10  $\mu$ m, C18 110 Å column (150  $\times$  30 mm<sup>2</sup>) using 5–95% MeCN in 0.05% aqueous TFA solution as the eluent. A total of 81 mg of compound **18** was obtained as the TFA salt (0.07 mmol, 99%) as a white amorphous solid. <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>):  $\delta$  12.24 (s, 1H), 8.95 (s, 1H), 8.56 (d,  $J = 8.1$  Hz, 1H), 8.26–8.11 (m, 1H), 8.08 (d,  $J = 6.6$  Hz, 1H), 7.90 (d,  $J = 7.7$  Hz, 2H), 7.64 (d,  $J = 7.3$  Hz, 2H), 7.47–7.38 (m, 3H), 7.38–7.16 (m, 8H), 6.56–6.48 (m, 2H), 4.62–4.16 (m, 4H), 4.13–3.64 (m, 4H), 3.62 (s, 3H), 3.46–3.16 (m, 9H), 3.14–2.84 (m, 4H), 2.79 (s, 3H), 2.23–1.95 (m, 8H), 1.89–1.58 (m, 2H), 1.18–0.83 (m, 1H), 1.12–0.83 (m, 6H), 0.83–0.56 (m, 17H); LC–MS  $t_R = 1.54$  min, ESIMS  $m/z$ : 1099.03  $[M + H]^+$ ; HRESIMS  $m/z$ : 1098.6164  $[M + H]^+$  calcd for  $C_{59}H_{84}N_7O_{13}$ , 1098.6122.

**Compound 19.** To a stirred solution of **18** (81 mg, 0.0714 mmol) in EtOAc (500 mL) were added CMPI (100 mg, 0.391 mmol) and DIEA (62  $\mu$ L, 0.36 mmol). After 16 h, analysis by LC–MS showed that the macrocyclic condensation reaction was complete. The mixture was evaporated to give a yellow oil. To a stirred solution of the resulting oil in CH<sub>2</sub>Cl<sub>2</sub> (4 mL) was added Et<sub>2</sub>NH (500  $\mu$ L, 4.83 mmol). After 2 h, analysis by LC–MS showed that the deprotection reaction was complete. The reaction mixture was diluted with H<sub>2</sub>O and DMAc, and then the mixture was purified by preparatory RP-HPLC with a Phenomenex Gemini-NX 10  $\mu$ m, C18 110 Å column (150  $\times$  30 mm<sup>2</sup>) using 5–95% MeCN in 0.05% aqueous TFA solution as the eluent. A total of 40 mg of compound **19** was obtained as the TFA salt (0.041 mmol, 61%) as a white amorphous solid. <sup>1</sup>H NMR (500 MHz, DMSO-*d*<sub>6</sub>):  $\delta$  9.08 (d,  $J = 6.5$  Hz, 1H), 8.84 (s, 1H), 8.69 (s, 1H), 8.40 (d,  $J = 6.1$  Hz, 1H), 7.88–7.76 (m, 2H), 7.33–7.17 (m, 5H), 6.60–6.47 (m, 1H), 4.92–4.81 (m, 1H), 4.62 (d,  $J = 11.3$  Hz, 1H), 4.58–4.47 (m, 1H), 4.37 (d,  $J = 10.8$  Hz, 1H), 4.34–4.28 (m, 1H), 4.27–4.18 (m, 1H), 4.02 (d,  $J = 10.7$  Hz, 1H), 3.83–3.79 (m, 1H), 3.69 (d,  $J = 10.4$  Hz, 1H), 3.65–3.55 (m, 2H), 3.33–3.25 (m, 5H), 3.08 (s, 3H),

2.96 (s, 3H), 2.93–2.83 (m, 1H), 2.56–2.52 (m, 3H), 2.42–2.19 (m, 3H), 2.17–1.92 (m, 3H), 1.89–1.65 (m, 6H), 1.29–1.13 (m, 1H), 1.08–1.03 (m, 4H), 1.07–0.70 (m, 17H); <sup>13</sup>C NMR (126 MHz, DMSO):  $\delta$  174.35, 172.49, 171.91, 171.69, 171.34, 166.34, 137.74, 129.58, 128.59, 126.94, 80.33, 65.95, 60.79, 58.53, 57.65, 57.39, 54.78, 53.74, 53.03, 52.30, 47.53, 45.88, 44.05, 41.80, 40.88, 37.16, 36.38, 36.03, 32.59, 32.43, 32.02, 31.33, 29.92, 29.77, 29.11, 25.99, 23.75, 23.43, 18.91, 18.75, 18.19, 17.81, 15.92, 14.93, 14.75, 11.47, 10.14; LC–MS  $t_R = 1.11$  min,  $m/z$ : 858.76  $[M + H]^+$ ; HRESIMS  $m/z$ : 858.5345  $[M + H]^+$  calcd for  $C_{44}H_{71}N_7O_{10}$ , 858.5335.

**Compound 21.** To a stirred solution of Dov-Abu (3-N3)-Dil-OtBu (180 mg, 0.29 mmol) and *N*-(but-3-yn-1-yl)-2-chloroacetamide (75 mg, 0.52 mmol) in DMF (5 mL) was added CuBr (76 mg, 0.53 mmol). After 15 h, analysis by LC–MS showed that the reaction was complete. The crude suspension was diluted with 15 mL of DMF, 2 mL of 0.5 M ethylenediaminetetraacetic acid (EDTA), and 10 mL of water. The mixture was purified by preparatory RP-HPLC using MeCN in 0.05% aqueous TFA as the eluent. A total of 120 mg of Dov-Abu (3-triazole-CH<sub>2</sub>CH<sub>2</sub>NH-acetyl-Cl)-Dil-OtBu (0.016 mmol, 54%) was obtained as a white amorphous solid.

Dov-Abu (3-triazole-CH<sub>2</sub>CH<sub>2</sub>NH-acetyl-Cl)-Dil-OtBu (120 mg, 0.016 mmol) was stirred in 4.0 M HCl in dioxane (2.4 mL, 4 mmol). After 3 h, analysis by LC–MS showed that the reaction was complete. The crude reaction mixture was concentrated under reduced pressure. A total of 89 mg of compound **21** was obtained as the HCl salt (0.19 mmol, 99%) as an off-white amorphous solid. <sup>1</sup>H NMR (400 MHz, DMSO-*d*<sub>6</sub>): a complex spectrum was observed, presumably because of conformational isomers):  $\delta$  9.83 (s, 1H), [9.58 (d,  $J = 8.7$  Hz), 9.35 (d,  $J = 8.7$  Hz) 1H], 8.35 (t,  $J = 5.7$  Hz, 1H), [7.80 (s), 7.94 (s), 1H], 5.46 (t,  $J = 8.2$  Hz, 1H), 5.09–4.98 (m, 1H), 4.38 (br-s, 1H), [4.07 (s), 4.05 (s), 2H], 3.82 (d,  $J = 5.9$  Hz, 1H), 3.64 (br-s, 1H), 3.42–3.31 (m, 2H), 3.18 (s, 3H), [2.93 (s), 2.90 (s), 3H], 2.84–2.67 (m, 8H), 2.37–2.13 (m, 2H), 1.99 (dd,  $J = 15.7, 9.3$  Hz, 1H), 1.83–1.68 (m, 1H), 1.63–1.45 (m, 4H), 1.36–1.19 (m, 1H), 0.96 (d,  $J = 6.8$  Hz, 3H), 0.91–0.80 (m, 6H), 0.75 (t,  $J = 7.2$  Hz, 3H); <sup>13</sup>C NMR (101 MHz, DMSO-*d*<sub>6</sub>):  $\delta$  173.28, 169.78, 166.33, 165.96, 159.43, 159.06, 158.71, 158.35, 144.30, 122.75, 78.31, 72.33, 71.78, 57.54, 56.57, 53.24, 50.27, 43.07, 41.93, 41.60, 39.04, 38.23, 37.28, 32.72, 31.64, 28.92, 27.03, 25.82, 25.66, 24.01, 19.55, 16.91, 16.20, 10.90; LC–MS  $t_R = 0.94$  min,  $m/z$ : 602.49, 604.50  $[M + H]^+$ ; HRESIMS  $m/z$ : 602.3425  $[M + H]^+$  calcd for  $C_{27}H_{49}ClN_7O_6$ , 602.3427.

**Compound 22.** To a stirred solution of compound **21** (24 mg, 0.04 mmol) and H-Dap (N<sub>3</sub>)-Phe-OMe HCl salt **11** (15 mg, 0.04 mmol) in CH<sub>2</sub>Cl<sub>2</sub> (3 mL) were added DIEA (25  $\mu$ L, 0.140 mmol), EDCI (18 mg, 0.09 mmol), and HOBT (5 mg, 0.03 mmol). After 18 h, analysis by LC–MS showed that the reaction conversion was complete. The reaction mixture was diluted with H<sub>2</sub>O and DMAc, and then the mixture was purified by preparatory RP-HPLC with a Phenomenex Gemini-NX 10  $\mu$ m, C18 110 Å column (150  $\times$  30 mm<sup>2</sup>) using 5–95% MeCN in 0.05% aqueous TFA solution as the eluent. A total of 24 mg of compound **22** was recovered as the TFA salt (0.022 mmol, 59%) as a white amorphous solid. LC–MS  $t_R = 1.29$  min,  $m/z$ : 973.83, 975.85  $[M + H]^+$ ; HRESIMS  $m/z$ : 973.5387  $[M + H]^+$  calcd for  $C_{46}H_{74}ClN_{12}O_9$ , 973.5385.

**Compound 23.** To a stirred solution of compound **13a** (15 mg, 0.017 mmol) and *N*-(but-3-yn-1-yl)-2-bromoacetamide (5 mg, 0.026 mmol) in DMF (1 mL) was added CuBr (10 mg,



0.067 mmol). After 15 h, analysis by LC–MS showed that the reaction was complete. The crude suspension was diluted with 7 mL of DMAc, 2 mL of 0.5 M EDTA, and 5 mL of water. The mixture was purified by preparatory RP–HPLC using MeCN in 0.05% aqueous TFA as the eluent. A total of 5 mg of compound **23** (4.58  $\mu$ mol, 28%) was obtained as a white amorphous solid.  $^1\text{H}$  NMR (400 MHz, DMSO- $d_6$ ; a complex spectrum was observed, presumably because of cis/trans conformational isomers):  $\delta$  8.80 (d,  $J$  = 8.6 Hz, 2H), 8.65 (s, 1H), 8.38–8.29 (m, 1H), 8.03–7.92 (m, 1H), 7.32–7.14 (m, 5H), 5.12–4.90 (m, 1H), 4.67 (br-s, 1H), 4.57 (t,  $J$  = 8.7 Hz, 1H), 4.52–4.41 (m, 1H), 4.24 (t,  $J$  = 8.5 Hz, 1H), 4.09–3.94 (m, 2H), 3.89–3.39 (m, 9H), 3.35–3.24 (m, 2H), 3.23–3.11 (m, 6H), 3.08–2.94 (m, 4H), 2.92–2.56 (m, 3H), 2.45 (t,  $J$  = Hz, 3H), [2.35–2.30 (m), 2.29–2.14 (m), 2H], 2.12–1.92 (m, 2H), 1.87–1.67 (m, 2H), 1.35–1.18 (m, 1H), [1.11–1.05 (m), 1.02–0.83 (m), 17H], 0.81–0.66 (m, 6H); LC–MS  $t_{\text{R}}$  = 1.19 min,  $m/z$ : 976.79, 978.77 [ $M + H$ ] $^+$ ; HRESIMS  $m/z$ : 976.4874, 978.4873 [ $M + H$ ] $^+$  calcd for  $\text{C}_{46}\text{H}_{75}\text{BrN}_9\text{O}_9$ , 976.4866.

## ■ ASSOCIATED CONTENT

### Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acsomega.8b00093.

Detailed experimental procedures and NMR spectra ( $^1\text{H}$  and  $^{13}\text{C}$ ) for **8**, **10**, **11**, **13a**, **13b**, **13c**, **15**, **16**, **17**, **18**, **19**, **21**, **22**, and **23** (PDF)

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### Notes

The authors declare no competing financial interest.

## ■ ACKNOWLEDGMENTS

The authors acknowledge the work of C. Kemball and M. Wu for in vitro cytotoxicity work and B. Wiggins for mass spectrometry assistance. They would like to acknowledge D. Weller and K. Morrison for manuscript review and helpful discussions.

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